NUCLEOBASE EDITORS AND USES THEREOF

GOVERNMENT SUPPORT

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RELATED APPLICTIONS

[0002] This application claims priority under 35 U.S.C. § 119(e) to U.S. provisional patent applications, U.S.S.N. 62/245,828 filed October 23, 2015, U.S.S.N. 62/279,346 filed January 15, 2016, U.S.S.N. 62/311,763 filed March 22, 2016, U.S.S.N. 62/322,178 filed April 13, 2016, U.S.S.N. 62/357,352 filed June 30, 2016, U.S.S.N. 62/370,700 filed August 3, 2016, U.S.S.N. 62/398,490 filed September 22, 2016, U.S.S.N. 62/408,686 filed October 14, 2016, and U.S.S.N. 62/357,332 filed June 30, 2016; each of which is incorporated herein by reference.

BACKGROUND OF THE INVENTION

[0003] Targeted editing of nucleic acid sequences, for example, the targeted cleavage or the targeted introduction of a specific modification into genomic DNA, is a highly promising approach for the study of gene function and also has the potential to provide new therapies for human genetic diseases.[1] An ideal nucleic acid editing technology possesses three characteristics: (1) high efficiency of installing the desired modification; (2) minimal off-target activity; and (3) the ability to be programmed to edit precisely any site in a given nucleic acid, *e.g.*, any site within the human genome.[2] Current genome engineering tools, including engineered zinc finger nucleases (ZFNs),[3] transcription activator like effector nucleases (TALENs),[4] and most recently, the RNA-guided DNA endonuclease Cas9,[5] effect sequence-specific DNA cleavage in a genome. This programmable cleavage can result in mutation of the DNA at the cleavage site via non-homologous end joining (NHEJ) or replacement of the DNA surrounding the cleavage site via homology-directed repair (HDR).[6,7]

[0004] One drawback to the current technologies is that both NHEJ and HDR are stochastic processes that typically result in modest gene editing efficiencies as well as unwanted gene alterations that can compete with the desired alteration.[8] Since many genetic diseases in principle can be treated by effecting a specific nucleotide change at a specific location in the genome (for example, a C to T change in a specific codon of a gene associated with a disease),[9] the development of a programmable way to achieve such precision gene editing would represent both a powerful new research tool, as well as a potential new approach to gene editing-based human therapeutics.

SUMMARY OF THE INVENTION

[0005] The clustered regularly interspaced short palindromic repeat (CRISPR) system is a recently discovered prokaryotic adaptive immune system[10] that has been modified to enable robust and general genome engineering in a variety of organisms and cell lines.[11] CRISPR-Cas (CRISPR associated) systems are protein-RNA complexes that use an RNA molecule (sgRNA) as a guide to localize the complex to a target DNA sequence *via* base-pairing.[12] In the natural systems, a Cas protein then acts as an endonuclease to cleave the targeted DNA sequence.[13] The target DNA sequence must be both complementary to the sgRNA, and also contain a "protospacer-adjacent motif" (PAM) at the 3′-end of the complementary region in order for the system to function.[14]

[0006] Among the known Cas proteins, *S. pyogenes* Cas9 has been mostly widely used as a tool for genome engineering.[15] This Cas9 protein is a large, multi-domain protein containing two distinct nuclease domains. Point mutations can be introduced into Cas9 to abolish nuclease activity, resulting in a dead Cas9 (dCas9) that still retains its ability to bind DNA in a sgRNA-programmed manner.[16] In principle, when fused to another protein or domain, dCas9 can target that protein to virtually any DNA sequence simply by co-expression with an appropriate sgRNA.

[0007] The potential of the dCas9 complex for genome engineering purposes is immense. Its unique ability to bring proteins to specific sites in a genome programmed by the sgRNA in theory can be developed into a variety of site-specific genome engineering tools beyond nucleases, including transcriptional activators, transcriptional repressors, histone-modifying proteins, integrases, and recombinases.[11] Some of these potential applications have recently been implemented through dCas9 fusions with transcriptional activators to afford RNA-guided transcriptional activators,[17,18] transcriptional repressors,[16,19,20] and chromatin

modification enzymes.[21] Simple co-expression of these fusions with a variety of sgRNAs results in specific expression of the target genes. These seminal studies have paved the way for the design and construction of readily programmable sequence-specific effectors for the precise manipulation of genomes.

[0008] Significantly, 80-90% of protein mutations responsible for human disease arise from the substitution, deletion, or insertion of only a single nucleotide.[6] Most current strategies for single-base gene correction include engineered nucleases (which rely on the creation of double-strand breaks, DSBs, followed by stochastic, inefficient homology-directed repair, HDR), and DNA-RNA chimeric oligonucleotides.[22] The latter strategy involves the design of a RNA/DNA sequence to base pair with a specific sequence in genomic DNA except at the nucleotide to be edited. The resulting mismatch is recognized by the cell's endogenous repair system and fixed, leading to a change in the sequence of either the chimera or the genome. Both of these strategies suffer from low gene editing efficiencies and unwanted gene alterations, as they are subject to both the stochasticity of HDR and the competition between HDR and non-homologous end-joining, NHEJ.[23-25] HDR efficiencies vary according to the location of the target gene within the genome,[26] the state of the cell cycle,[27] and the type of cell/tissue.[28] The development of a direct, programmable way to install a specific type of base modification at a precise location in genomic DNA with enzyme-like efficiency and no stochasticity therefore represents a powerful new approach to gene editing-based research tools and human therapeutics.

[0009] Some aspects of the disclosure are based on the recognition that certain configurations of a dCas9 domain, and a cytidine deaminase domain fused by a linker are useful for efficiently deaminating target cytidine residues. Other aspects of this disclosure relate to the recognition that a nucleobase editing fusion protein with a cytidine deaminase domain fused to the N-terminus of a nuclease inactive Cas9 (dCas9) via a linker was capable of efficiently deaminating target nucleic acids in a double stranded DNA target molecule. See for example, Examples 3 and 4 below, which demonstrate that the fusion proteins, which are also referred to herein as base editors, generate less indels and more efficiently deaminate target nucleic acids than other base editors, such as base editors without a UGI domain. In some embodiments, the fusion protein comprises a nuclease-inactive Cas9 (dCas9) domain and an apolipoprotein B mRNA-editing complex 1 (APOBEC1) deaminase domain, where the deaminase domain is fused to the N-terminus of the dCas9 domain via a linker comprising the amino acid sequence SGSETPGTSESATPES (SEQ ID NO: 7). In some embodiments, the nuclease-inactive Cas9 (dCas9) domain of comprises the amino acid

sequence set forth in SEQ ID NO: 263. In some embodiments, the deaminase is rat APOBEC1 (SEQ ID NO: 284). In some embodiments, the deaminase is human APOBEC1 (SEQ ID NO: 282). In some embodiments, the deaminase is pmCDA1 (SEQ ID NO: 5738). In some embodiments, the deaminase is human APOBEC3G (SEQ ID NO: 275). In some embodiments, the deaminase is a human APOBEC3G variant of any one of (SEQ ID NOs: 5739-5741).

[0010] Some aspects of the disclosure are based on the recognition that certain configurations of a dCas9 domain, and a cytidine deaminase domain fused by a linker are useful for efficiently deaminating target cytidine residues. Other aspects of this disclosure relate to the recognition that a nucleobase editing fusion protein with an apolipoprotein B mRNA-editing complex 1 (APOBEC1) deaminase domain fused to the N-terminus of a nuclease inactive Cas9 (dCas9) via a linker comprising the amino acid sequence SGSETPGTSESATPES (SEQ ID NO: 7) was capable of efficiently deaminating target nucleic acids in a double stranded DNA target molecule. In some embodiments, the fusion protein comprises a nuclease-inactive Cas9 (dCas9) domain and an apolipoprotein B mRNA-editing complex 1 (APOBEC1) deaminase domain, where the deaminase domain is fused to the N-terminus of the dCas9 domain via a linker comprising the amino acid sequence SGSETPGTSESATPES (SEQ ID NO: 7).

[0011] In some embodiments, the fusion protein comprises the amino acid residues 11-1629 of the amino acid sequence set forth in SEQ ID NO: 591. In some embodiments, the fusion protein comprises the amino acid sequence set forth in SEQ ID NO: 591. In some embodiments, the fusion protein comprises the amino acid sequence of any one of SEQ ID NOs: 5737, 5743, 5745, and 5746.

[0012] Some aspects of this disclosure provide strategies, systems, reagents, methods, and kits that are useful for the targeted editing of nucleic acids, including editing a single site within a subject's genome, *e.g.*, a human's genome. In some embodiments, fusion proteins of Cas9 (*e.g.*, dCas9, nuclease active Cas9, or Cas9 nickase) and deaminases or deaminase domains, are provided. In some embodiments, methods for targeted nucleic acid editing are provided. In some embodiments, reagents and kits for the generation of targeted nucleic acid editing proteins, *e.g.*, fusion proteins of Cas9 and deaminases or deaminase domains, are provided.

[0013] Some aspects of this disclosure provide fusion proteins comprising a Cas9 protein as provided herein that is fused to a second protein (*e.g.,* an enzymatic domain such as a cytidine deaminase domain), thus forming a fusion protein. In some embodiments, the

second protein comprises an enzymatic domain, or a binding domain. In some embodiments, the enzymatic domain is a nuclease, a nickase, a recombinase, a deaminase, a methyltransferase, a methylase, an acetylase, an acetyltransferase, a transcriptional activator, or a transcriptional repressor domain. In some embodiments, the enzymatic domain is a nucleic acid editing domain. In some embodiments, the nucleic acid editing domain is a deaminase domain. In some embodiments, the deaminase is a cytosine deaminase or a cytidine deaminase. In some embodiments, the deaminase is an apolipoprotein B mRNA-editing complex (APOBEC) family deaminase. In some embodiments, the deaminase is an APOBEC1 deaminase. In some embodiments, the deaminase is an APOBEC2 deaminase. In some embodiments, the deaminase is an APOBEC3 deaminase. In some embodiments, the deaminase is an APOBEC3A deaminase. In some embodiments, the deaminase is an APOBEC3B deaminase. In some embodiments, the deaminase is an APOBEC3C deaminase. In some embodiments, the deaminase is an APOBEC3D deaminase. In some embodiments, the deaminase is an APOBEC3E deaminase. In some embodiments, the deaminase is an APOBEC3F deaminase. In some embodiments, the deaminase is an APOBEC3G deaminase. In some embodiments, the deaminase is an APOBEC3H deaminase. In some embodiments, the deaminase is an APOBEC4 deaminase. In some embodiments, the deaminase is an activation-induced deaminase (AID). It should be appreciated that the deaminase may be from any suitable organism (*e.g.*, a human or a rat). In some embodiments, the deaminase is from a human, chimpanzee, gorilla, monkey, cow, dog, rat, or mouse. In some embodiments, the deaminase is rat APOBEC1 (SEQ ID NO: 284). In some embodiments, the deaminase is human APOBEC1 (SEQ ID NO: 282). In some embodiments, the deaminase is pmCDA1.

[0014] Some aspects of this disclosure provide fusion proteins comprising: (i) a nuclease-inactive Cas9 (dCas9) domain comprising the amino acid sequence of SEQ ID NO: 263; and (ii) an apolipoprotein B mRNA-editing complex 1 (APOBEC1) deaminase domain, wherein the deaminase domain is fused to the N-terminus of the dCas9 domain via a linker comprising the amino acid sequence of SGSETPGTSESATPES (SEQ ID NO: 7). In some embodiments, the deaminase is rat APOBEC1 (SEQ ID NO: 284). In some embodiments, the deaminase is human APOBEC1 (SEQ ID NO: 282). In some embodiments, the fusion protein comprises the amino acid sequence of SEQ ID NO: 591. In some embodiments, the fusion protein comprises the amino acid sequence of SEQ ID NO: 5737. In some embodiments, the deaminase is pmCDA1 (SEQ ID NO: 5738). In some embodiments, the deaminase is human APOBEC3G (SEQ ID NO: 275). In some embodiments, the deaminase is a human APOBEC3G variant of any one of SEQ ID NOs: 5739-5741.

[0015] Some aspects of this disclosure provide fusion proteins comprising: (i) a Cas9 nickase domain and (ii) an apolipoprotein B mRNA-editing complex 1 (APOBEC1) deaminase domain, wherein the deaminase domain is fused to the N-terminus of the Cas9 nickase domain.. In some embodiments, the Cas9 nickase domain comprises a D10X mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for D. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a D10A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a histidine at amino acid position 840 of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding amino acid position in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises the amino acid sequence as set forth in SEQ ID NO: 267. In some embodiments, the deaminase is rat APOBEC1 (SEQ ID NO: 284). In some embodiments, the deaminase is human APOBEC1 (SEQ ID NO: 282). In some embodiments, the deaminase is pmCDA1.

[0016] Some aspects of this disclosure provide fusion proteins comprising: (i) a Cas9 nickase domain and (ii) an apolipoprotein B mRNA-editing complex 1 (APOBEC1) deaminase domain, wherein the deaminase domain is fused to the N-terminus of the Cas9 nickase domain.. In some embodiments, the Cas9 nickase domain comprises a D10X mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for D. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a D10A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a histidine at amino acid position 840 of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding amino acid position in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises the amino acid sequence as set forth in SEQ ID NO: 267. In some embodiments, the deaminase is rat APOBEC1 (SEQ ID NO: 284). In some embodiments, the deaminase is human APOBEC1 (SEQ ID NO: 282). In some embodiments, the deaminase is pmCDA1.

[0017] Other aspects of this disclosure relate to the recognition that fusion proteins comprising a deaminase domain, a dCas9 domain and a uracil glycosylase inhibitor (UGI) domain demonstrate improved efficiency for deaminating target nucleotides in a nucleic acid molecule. Without wishing to be bound by any particular theory, cellular DNA-repair response to the presence of U:G heteroduplex DNA may be responsible for a decrease in nucleobase editing efficiency in cells. Uracil DNA glycosylase (UDG) catalyzes removal of U from DNA in cells, which may initiate base excision repair, with reversion of the U:G pair to a C:G pair as the most common outcome. As demonstrated herein, Uracil DNA Glycosylase Inhibitor (UGI) may inhibit human UDG activity. Without wishing to be bound by any particular theory, base excision repair may be inhibited by molecules that bind the single strand, block the edited base, inhibit UGI, inhibit base excision repair, protect the edited base, and/or promote "fixing" of the non-edited strand, *etc.*Thus, this disclosure contemplates fusion proteins comprising a dCas9-cytidine deaminase domain that is fused to a UGI domain.

[0018] In some embodiments, the fusion protein comprises a nuclease-inactive Cas9 (dCas9) domain; a nucleic acid editing domain; and a uracil glycosylase inhibitor (UGI) domain. In some embodiments, the dCas9 domain comprises a D10X mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for D. In some embodiments, the amino acid sequence of the dCas9 domain comprises a D10A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the dCas9 domain comprises an H840X mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for H. In some embodiments, the amino acid sequence of the dCas9 domain comprises an H840A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the dCas9 domain comprises the amino acid sequence as set forth in SEQ ID NO: 263.

[0019] Further aspects of this disclosure relate to the recognition that fusion proteins using a Cas9 nickase as the Cas9 domain demonstrate improved efficiency for editing nucleic acids. For example, aspects of this disclosure relate to the recognition that fusion proteins comprising a Cas9 nickase, a deaminase domain and a UGI domain demonstrate improved

efficiency for editing nucleic acids. For example, the improved efficiency for editing nucleotides is described below in the Examples section.

[0020] Some aspects of the disclosure are based on the recognition that any of the base editors provided herein are capable of modifying a specific nucleotide base without generating a significant proportion of indels. An "indel", as used herein, refers to the insertion or deletion of a nucleotide base within a nucleic acid. Such insertions or deletions can lead to frame shift mutations within a coding region of a gene. In some embodiments, it is desirable to generate base editors that efficiently modify (*e.g.* mutate or deaminate) a specific nucleotide within a nucleic acid, without generating a large number of insertions or deletions (*i.e.*, indels) in the nucleic acid. In certain embodiments, any of the base editors provided herein are capable of generating a greater proportion of intended modifications (*e.g.,* point mutations or deaminations) versus indels.

[0021] Some aspects of the disclosure are based on the recognition that any of the base editors provided herein are capable of efficiently generating an intended mutation, such as a point mutation, in a nucleic acid (*e.g.* a nucleic acid within a genome of a subject) without generating a significant number of unintended mutations, such as unintended point mutations.

[0022] In some embodiments, a fusion protein comprises a Cas9 nickase domain, a nucleic acid editing domain; and a uracil glycosylase inhibitor (UGI) domain. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a D10X mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for D. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a D10A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises a histidine at amino acid position 840 of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding amino acid position in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the amino acid sequence of the Cas9 nickase domain comprises the amino acid sequence as set forth in SEQ ID NO: 267.

[0023] In some embodiments, the deaminase domain of the fusion protein is fused to the N-terminus of the dCas9 domain or the Cas9 nickase. In some embodiments, the UGI domain is fused to the C-terminus of the dCas9 domain or the Cas9 nickase. In some embodiments, the dCas9 domain or the Cas9 nickase and the nucleic acid editing domain are fused via a linker.

In some embodiments, the dCas9 domain or the Cas9 nickase and the UGI domain are fused via a linker.

[0024] In certain embodiments, linkers may be used to link any of the peptides or peptide domains of the invention. The linker may be as simple as a covalent bond, or it may be a polymeric linker many atoms in length. In certain embodiments, the linker is a polpeptide or based on amino acids. In other embodiments, the linker is not peptide-like. In certain embodiments, the linker is a covalent bond (*e.g.*, a carbon-carbon bond, disulfide bond, carbon-heteroatom bond, *etc.*). In certain embodiments, the linker is a carbon-nitrogen bond of an amide linkage. In certain embodiments, the linker is a cyclic or acyclic, substituted or unsubstituted, branched or unbranched aliphatic or heteroaliphatic linker. In certain embodiments, the linker is polymeric (*e.g.*, polyethylene, polyethylene glycol, polyamide, polyester, *etc.*). In certain embodiments, the linker comprises a monomer, dimer, or polymer of aminoalkanoic acid. In certain embodiments, the linker comprises an aminoalkanoic acid (*e.g.*, glycine, ethanoic acid, alanine, beta-alanine, 3-aminopropanoic acid, 4-aminobutanoic acid, 5-pentanoic acid, *etc.*). In certain embodiments, the linker comprises a monomer, dimer, or polymer of aminohexanoic acid (Ahx). In certain embodiments, the linker is based on a carbocyclic moiety (*e.g.*, cyclopentane, cyclohexane). In other embodiments, the linker comprises a polyethylene glycol moiety (PEG). In other embodiments, the linker comprises amino acids. In certain embodiments, the linker comprises a peptide. In certain embodiments, the linker comprises an aryl or heteroaryl moiety. In certain embodiments, the linker is based on a phenyl ring. The linker may included funtionalized moieties to facilitate attachment of a nucleophile (*e.g.*, thiol, amino) from the peptide to the linker. Any electrophile may be used as part of the linker. Exemplary electrophiles include, but are not limited to, activated esters, activated amides, Michael acceptors, alkyl halides, aryl halides, acyl halides, and isothiocyanates.

[0025] In some embodiments, the linker comprises the amino acid sequence $(GGGGS)_n$ (SEQ ID NO: 5), $(G)_n$, $(EAAAK)_n$ (SEQ ID NO: 6), $(GGS)_n$, $(SGGS)_n$ (SEQ ID NO: 4288), SGSETPGTSESATPES (SEQ ID NO: 7), $(XP)_n$, or any combination thereof, wherein n is independently an integer between 1 and 30, and wherein X is any amino acid. In some embodiments, the linker comprises the amino acid sequence $(GGS)_n$, wherein n is 1, 3, or 7. In some embodiments, the linker comprises the amino acid sequence SGSETPGTSESATPES (SEQ ID NO: 7).

[0026] In some embodiments, the fusion protein comprises the structure [nucleic acid editing domain]-[optional linker sequence]-[dCas9 or Cas9 nickase]-[optional linker

sequence]-[UGI]. In some embodiments, the fusion protein comprises the structure [nucleic acid editing domain]-[optional linker sequence]-[UGI]-[optional linker sequence]-[dCas9 or Cas9 nickase]; [UGI]-[optional linker sequence]-[nucleic acid editing domain]-[optional linker sequence]-[dCas9 or Cas9 nickase]; [UGI]-[optional linker sequence]-[dCas9 or Cas9 nickase]-[optional linker sequence]-[nucleic acid editing domain]; [dCas9 or Cas9 nickase]-[optional linker sequence]-[UGI]-[optional linker sequence]-[nucleic acid editing domain]; or [dCas9 or Cas9 nickase]-[optional linker sequence]-[nucleic acid editing domain]-[optional linker sequence]-[UGI].

[0027] In some embodiments, the nucleic acid editing domain comprises a deaminase. In some embodiments, the nucleic acid editing domain comprises a deaminase. In some embodiments, the deaminase is a cytidine deaminase. In some embodiments, the deaminase is an apolipoprotein B mRNA-editing complex (APOBEC) family deaminase. In some embodiments, the deaminase is an APOBEC1 deaminase, an APOBEC2 deaminase, an APOBEC3A deaminase, an APOBEC3B deaminase, an APOBEC3C deaminase, an APOBEC3D deaminase, an APOBEC3F deaminase, an APOBEC3G deaminase, an APOBEC3H deaminase, or an APOBEC4 deaminase. In some embodiments, the deaminase is an activation-induced deaminase (AID). In some embodiments, the deaminase is a Lamprey CDA1 (pmCDA1) deaminase.

[0028] In some embodiments, the deaminase is from a human, chimpanzee, gorilla, monkey, cow, dog, rat, or mouse. In some embodiments, the deaminase is from a human. In some embodiments the deaminase is from a rat. In some embodiments, the deaminase is a rat APOBEC1 deaminase comprising the amino acid sequence set forth in (SEQ ID NO: 284). In some embodiments, the deaminase is a human APOBEC1 deaminase comprising the amino acid sequence set forth in (SEQ ID NO: 282). In some embodiments, the deaminase is pmCDA1 (SEQ ID NO: 5738). In some embodiments, the deaminase is human APOBEC3G (SEQ ID NO: 275). In some embodiments, the deaminase is a human APOBEC3G variant of any one of (SEQ ID NOs: 5739-5741). In some embodiments, the deaminase is at least 80%, at least 85%, at least 90%, at least 92%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to any one of the amino acid sequences set forth in SEQ ID NOs: 266-284 or 5725-5741.

[0029] In some embodiments, the UGI domain comprises an amino acid sequence that is at least 80%, at least 85%, at least 90%, at least 92%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to SEQ ID NO: 600. In some

embodiments, the UGI domain comprises the amino acid sequence as set forth in SEQ ID NO: 600.

[0030] Some aspects of this disclosure provide complexes comprising a Cas9 protein or a Cas9 fusion protein as provided herein, and a guide RNA bound to the Cas9 protein or the Cas9 fusion protein.

[0031] Some aspects of this disclosure provide methods of using the Cas9 proteins, fusion proteins, or complexes provided herein. For example, some aspects of this disclosure provide methods comprising contacting a DNA molecule (a) with a Cas9 protein or a fusion protein as provided herein and with a guide RNA, wherein the guide RNA is about 15-100 nucleotides long and comprises a sequence of at least 10 contiguous nucleotides that is complementary to a target sequence; or (b) with a Cas9 protein, a Cas9 fusion protein, or a Cas9 protein or fusion protein complex with a gRNA as provided herein.

[0032] Some aspects of this disclosure provide kits comprising a nucleic acid construct, comprising (a) a nucleotide sequence encoding a Cas9 protein or a Cas9 fusion protein as provided herein; and (b) a heterologous promoter that drives expression of the sequence of (a). In some embodiments, the kit further comprises an expression construct encoding a guide RNA backbone, wherein the construct comprises a cloning site positioned to allow the cloning of a nucleic acid sequence identical or complementary to a target sequence into the guide RNA backbone.

[0033] Some aspects of this disclosure provide polynucleotides encoding a Cas9 protein of a fusion protein as provided herein. Some aspects of this disclosure provide vectors comprising such polynucleotides. In some embodiments, the vector comprises a heterologous promoter driving expression of polynucleotide.

[0034] Some aspects of this disclosure provide cells comprising a Cas9 protein, a fusion protein, a nucleic acid molecule, and/or a vector as provided herein.

[0035] The description of exemplary embodiments of the reporter systems above is provided for illustration purposes only and not meant to be limiting. Additional reporter systems, *e.g.*, variations of the exemplary systems described in detail above, are also embraced by this disclosure.

[0036] The summary above is meant to illustrate, in a non-limiting manner, some of the embodiments, advantages, features, and uses of the technology disclosed herein. Other embodiments, advantages, features, and uses of the technology disclosed herein will be apparent from the Detailed Description, the Drawings, the Examples, and the Claims.

BRIEF DESCRIPTION OF THE DRAWINGS

[0037] **Figure 1** shows the deaminase activity of deaminases on single stranded DNA substrates. Single stranded DNA substrates using randomized PAM sequences (NNN PAM) were used as negative controls. Canonical PAM sequneces used (NGG PAM)

[0038] **Figure 2** shows activity of Cas9:deaminase fusion proteins on single stranded DNA substrates.

[0039] **Figure 3** illustrates double stranded DNA substrate binding by Cas9:deaminase:sgRNA complexes.

[0040] **Figure 4** illustrates a double stranded DNA deamination assay.

[0041] **Figure 5** demonstrates that Cas9 fusions can target positions 3-11 of double-stranded DNA target sequences (numbered according to the schematic in **Figure 5**). Upper Gel: 1 μM rAPOBEC1-GGS-dCas9, 125 nM dsDNA, 1 equivalent sgRNA. Mid Gel: 1 μM rAPOBEC1-$(GGS)_3$(SEQ ID NO: 596)-dCas9, 125 nM dsDNA, 1 equivalent sgRNA. Lower Gel: 1.85 μM rAPOBEC1-XTEN-dCas9, 125 nM dsDNA, 1 equivalent sgRNA.

[0042] **Figure 6** demonstrates that the correct guide RNA, *e.g.*, the correct sgRNA, is required for deaminase activity.

[0043] **Figure 7** illustrates the mechanism of target DNA binding of *in vivo* target sequences by deaminase-dCas9:sgRNA complexes.

[0044] **Figure 8** shows successful deamination of exemplary disease-associated target sequences.

[0045] **Figure 9** shows *in vitro* C→T editing efficiencies using His6-rAPOBEC1-XTEN-dCas9.

[0046] **Figure 10** shows C→T editing efficiencies in HEK293T cells is greatly enhanced by fusion with UGI.

[0047] **Figures 11A** to **11C** show NBE1 mediates specific, guide RNA-programmed C to U conversion *in vitro*. Figure 11A: Nucleobase editing strategy. DNA with a target C (red) at a locus specified by a guide RNA (green) is bound by dCas9 (blue), which mediates the local denaturation of the DNA substrate. Cytidine deamination by a tethered APOBEC1 enzyme (orange) converts the target C to U. The resulting G:U heteroduplex can be permanently converted to an A:T base pair following DNA replication or repair. If the U is in the template DNA strand, it will also result in an RNA transcript containing a G to A mutation following transcription. Figure 11B: Deamination assay showing an activity window of approximately five nucleotides. Following incubation of NBE1-sgRNA complexes with dsDNA substrates

at 37 °C for 2 h, the 5' fluorophore-labeled DNA was isolated and incubated with USER enzyme (uracil DNA glycosylase and endonuclease VIII) at 37 °C for 1 h to induce DNA cleavage at the site of any uracils. The resulting DNA was resolved on a denaturing polyacrylamide gel, and any fluorophore-linked strands were visualized. Each lane is labeled according to the position of the target C within the protospacer, or with "–" if no target C is present, counting the base distal from the PAM as position 1. Figure 11C: Deaminase assay showing the sequence specificity and sgRNA-dependence of NBE1. The DNA substrate with a target C at position 7 was incubated with NBE1 as in Figure 11B with either the correct sgRNA, a mismatched sgRNA, or no sgRNA. No C to U editing is observed with the mismatched sgRNA or with no sgRNA. The positive control sample contains a DNA sequence with a U synthetically incorporated at position 7.

[0048] **Figures 12A** to **12B** show effects of sequence context and target C position on nucleobase editing efficiency *in vitro*. Figure 12A: Effect of changing the sequence surrounding the target C on editing efficiency *in vitro*. The deamination yield of 80% of targeted strands (40% of total sequencing reads from both strands) for C_7 in the protospacer sequence 5′-TTATTTCGTGGATTTATTTA-3′(SEQ ID NO: 264) was defined as 1.0, and the relative deamination efficiencies of substrates containing all possible single-base mutations at positions 1-6 and 8-13 are shown. Values and error bars reflect the mean and standard deviation of two or more independent biological replicates performed on different days. Figure 12B: Positional effect of each NC motif on editing efficiency *in vitro*. Each NC target motif was varied from positions 1 to 8 within the protospacer as indicated in the sequences shown on the right (the PAM shown in red, the protospacer plus one base 5' to the protospacer are also shown). The percentage of total sequence reads containing T at each of the numbered target C positions following incubation with NBE1 is shown in the graph. Note that the maximum possible deamination yield *in vitro* is 50% of total sequencing reads (100% of targeted strands). Values and error bars reflect the mean and standard deviation of two or three independent biological replicates performed on different days. Figure 12B depicts SEQ ID NOs: 285 through 292 from top to bottom, respectively.

[0049] **Figures 13A** to **13C** show nucleobase editing in human cells. Figure 13A: Protospacer (black) and PAM (red) sequences of the six mammalian cell genomic loci targeted by nucleobase editors. Target Cs are indicated with subscripted numbers corresponding to their positions within the protospacer. Figure 13A depicts SEQ ID NOs: 293 through 298 from top to bottom, respectively. Figure 13B: HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE3 and an appropriate sgRNA. Three days

after transfection, genomic DNA was extracted and analyzed by high-throughput DNA sequencing at the six loci. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, are shown for NBE1, NBE2, and NBE3 at all six genomic loci, and for wt Cas9 with a donor HDR template at three of the six sites (EMX1, HEK293 site 3, and HEK293 site 4). Values and error bars reflect the mean and standard deviation of three independent biological replicates performed on different days. Figure 13C: Frequency of indel formation, calculated as described in the Methods, is shown following treatment of HEK293T cells with NBE2 and NBE3 for all six genomic loci, or with wt Cas9 and a single-stranded DNA template for HDR at three of the six sites (EMX1, HEK293 site 3, and HEK293 site 4). Values reflect the mean of at least three independent biological replicates performed on different days.

[0050] **Figures 14A** to **14C** show NBE2- and NBE3-mediated correction of three disease-relevant mutations in mammalian cells. For each site, the sequence of the protospacer is indicated to the right of the name of the mutation, with the PAM highlighted in green and the base responsible for the mutation indicated in bold with a subscripted number corresponding to its position within the protospacer. The amino acid sequence above each disease-associated allele is shown, together with the corrected amino acid sequence following nucleobase editing in red. Underneath each sequence are the percentages of total sequencing reads with the corresponding base. Cells were nucleofected with plasmids encoding NBE2 or NBE3 and an appropriate sgRNA. Two days after nucleofection, genomic DNA was extracted and analyzed by HTS to assess pathogenic mutation correction. Figure 14A: The Alzheimer's disease-associated *APOE4* allele is converted to *APOE3'* in mouse astrocytes by NBE3 in 11% of total reads (44% of nucleofected astrocytes). Two nearby Cs are also converted to Ts, but with no change to the predicted sequence of the resulting protein (SEQ ID NO: 299). Figure 14B The cancer-associated p53 N239D mutation is corrected by NBE2 in 11% of treated human lymphoma cells (12% of nucleofected cells) that are heterozygous for the mutation (SEQ ID NO: 300). Figure 14C The p53 Y163C mutation is corrected by NBE3 in 7.6% of nucleofected human breast cancer cells (SEQ ID NO: 301).

[0051] **Figures 15A** to **15D** show effects of deaminase–dCas9 linker length and composition on nucleobase editing. Gel-based deaminase assay showing the deamination window of nucleobase editors with deaminase–Cas9 linkers of GGS (Figure 15A), $(GGS)_3$ (SEQ ID NO: 596) (Figure 15B), XTEN (Figure 15C), or $(GGS)_7$ (SEQ ID NO: 597) (Figure 15D). Following incubation of 1.85 μM editor-sgRNA complexes with 125 nM dsDNA substrates at 37 °C for 2 h, the dye-conjugated DNA was isolated and incubated with USER

enzyme (uracil DNA glycosylase and endonuclease VIII) at 37 °C for an additional hour to cleave the DNA backbone at the site of any uracils. The resulting DNA was resolved on a denaturing polyacrylamide gel, and the dye-conjugated strand was imaged. Each lane is numbered according to the position of the target C within the protospacer, or with – if no target C is present. 8U is a positive control sequence with a U synthetically incorporated at position 8.

[0052] **Figures 16A** to **16B** show NBE1 is capable of correcting disease-relevant mutations *in vitro*. Figure 16A: Protospacer and PAM sequences (red) of seven disease-relevant mutations. The disease-associated target C in each case is indicated with a subscripted number reflecting its position within the protospacer. For all mutations except both *APOE4* SNPs, the target C resides in the template (non-coding) strand. Figure 16A depicts SEQ ID NOs: 302 through 308 from top to bottom, respectively. Figure 16B: Deaminase assay showing each dsDNA oligonucleotide before (–) and after (+) incubation with NBE1, DNA isolation, and incubation with USER enzymes to cleave DNA at positions containing U. Positive control lanes from incubation of synthetic oligonucleotides containing U at various positions within the protospacer with USER enzymes are shown with the corresponding number indicating the position of the U.

[0053] **Figure 17** shows processivity of NBE1. The protospacer and PAM (red) of a 60-mer DNA oligonucleotide containing eight consecutive Cs is shown at the top. The oligonucleotide (125 nM) was incubated with NBE1 (2 μM) for 2 h at 37 °C. The DNA was isolated and analyzed by high-throughput sequencing. Shown are the percent of total reads for the most frequent nine sequences observed. The vast majority of edited strands (>93%) have more than one C converted to T. This figure depicts SEQ ID NO: 309.

[0054] **Figures 18A** to **18H** show the effect of fusing UGI to NBE1 to generate NBE2. Figure 18A: Protospacer and PAM (red) sequences of the six mammalian cell genomic loci targeted with nucleobase editors. Editable Cs are indicated with labels corresponding to their positions within the protospacer. Figure 18A depicts SEQ ID NOs: 293 through 298 from top to bottom, respectively. Figures 18B to 18G: HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE1 and UGI, and an appropriate sgRNA. Three days after transfection, genomic DNA was extracted and analyzed by high-throughput DNA sequencing at the six loci. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, are shown for NBE1, NBE1 and UGI, and NBE2 at all six genomic loci. Figure 18H: C to T mutation rates at 510 Cs surrounding the protospacers of interest for NBE1, NBE1 plus UGI on a separate plasmid,

NBE2, and untreated cells are shown. The data show the results of 3,000,000 DNA sequencing reads from 1.5x106 cells. Values reflect the mean of at least two biological experiments conducted on different days.

[0055] **Figure 19** shows nucleobase editing efficiencies of NBE2 in U2OS and HEK293T cells. Cellular C to T conversion percentages by NBE2 are shown for each of the six targeted genomic loci in HEK293T cells and U2OS cells. HEK293T cells were transfected using lipofectamine 2000, and U2OS cells were nucleofected. U2OS nucleofection efficiency was 74%. Three days after plasmid delivery, genomic DNA was extracted and analyzed for nucleobase editing at the six genomic loci by HTS. Values and error bars reflect the mean and standard deviation of at least two biological experiments done on different days.

[0056] **Figure 20** shows nucleobase editing persists over multiple cell divisions. Cellular C to T conversion percentages by NBE2 are displayed at two genomic loci in HEK293T cells before and after passaging the cells. HEK293T cells were transfected using Lipofectamine 2000. Three days post transfection, the cells were harvested and split in half. One half was subjected to HTS analysis, and the other half was allowed to propagate for approximately five cell divisions, then harvested and subjected to HTS analysis.

[0057] **Figure 21** shows genetic variants from ClinVar that can be corrected in principle by nucleobase editing. The NCBI ClinVar database of human genetic variations and their corresponding phenotypes[68] was searched for genetic diseases that can be corrected by current nucleobase editing technologies. The results were filtered by imposing the successive restrictions listed on the left. The x-axis shows the number of occurrences satisfying that restriction and all above restrictions on a logarithmic scale.

[0058] **Figure 22** shows *in vitro* identification of editable Cs in six genomic loci. Synthetic 80-mers with sequences matching six different genomic sites were incubated with NBE1 then analyzed for nucleobase editing *via* HTS. For each site, the sequence of the protospacer is indicated to the right of the name of the site, with the PAM highlighted in red. Underneath each sequence are the percentages of total DNA sequencing reads with the corresponding base. A target C was considered as "editable" if the *in vitro* conversion efficiency is >10%. Note that maximum yields are 50% of total DNA sequencing reads since the non-targeted strand is not a substrate for nucleobase editing. This figure depicts SEQ ID NOs: 293 through 298 from top to bottom, respectively.

[0059] **Figure 23** shows activities of NBE1, NBE2, and NBE3 at EMX1 off-targets. HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE3 and a sgRNA matching the EMX1 sequence using Lipofectamine 2000. Three days after

transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus the top ten known Cas9 off-target loci for the EMX1 sgRNA, as previously determined using the GUIDE-seq method[55]. EMX1 off-target 5 locus did not amplify and is not shown. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for NBE1, NBE2, and NBE3. On the far right are displayed the total number of sequencing reads reported for each sequence. This figure depicts SEQ ID NOs: 293, and 310 through 318 from top to bottom, respectively.

[0060] **Figure 24** shows activities of NBE1, NBE2, and NBE3 at FANCF off-targets. HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE3 and a sgRNA matching the FANCF sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus all of the known Cas9 off-target loci for the FANCF sgRNA, as previously determined using the GUIDE-seq method[55]. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for NBE1, NBE2, and NBE3. On the far right are displayed the total number of sequencing reads reported for each sequence. This figure depicts SEQ ID NOs: 294 and 319 through 326 from top to bottom, respectively.

[0061] **Figure 25** shows activities of NBE1, NBE2, and NBE3 at HEK293 site 2 off-targets. HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE3 and a sgRNA matching the HEK293 site 2 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus all of the known Cas9 off-target loci for the HEK293 site 2 sgRNA, as previously determined using the GUIDE-seq method[55]. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for NBE1, NBE2, and NBE3. On the far right are displayed the total number of sequencing reads reported for each sequence. This figure depicts SEQ ID NOs: 295, 327, and 328 from top to bottom, respectively.

[0062] **Figure 26** shows activities of NBE1, NBE2, and NBE3 at HEK293 site 3 off-targets. HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE3 and a sgRNA matching the HEK293 site 3 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus all of the known Cas9 off-target loci for the HEK293 site 3 sgRNA, as previously determined using the GUIDE-seq method[55]. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for NBE1, NBE2, and NBE3. On the far right are displayed the total number of sequencing reads reported for each sequence. This figure depicts SEQ ID NOs: 296 and 659 through 663 from top to bottom, respectively.

[0063] **Figure 27** shows activities of NBE1, NBE2, and NBE3 at HEK293 site 4 off-targets. HEK293T cells were transfected with plasmids expressing NBE1, NBE2, or NBE3 and a sgRNA matching the HEK293 site 4 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus the top ten known Cas9 off-target loci for the HEK293 site 4 sgRNA, as previously determined using the GUIDE-seq method[55]. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for NBE1, NBE2, and NBE3. On the far right are displayed the total number of sequencing reads reported for each sequence. This figure depicts SEQ ID NOs: 297 and 664 through 673 from top to bottom, respectively.

[0064] **Figure 28** shows non-target C mutation rates. Shown here are the C to T mutation rates at 2,500 distinct cytosines surrounding the six on-target and 34 off-target loci tested, representing a total of 14,700,000 sequence reads derived from approximately 1.8x106 cells.

[0065] **Figures 29A to 29C** show base editing in human cells. Figure 29A shows possible base editing outcomes in mammalian cells. Initial editing resulted in a U:G mismatch. Recognition and excision of the U by uracil DNA glycosylase (UDG) initiated base excision repair (BER), which lead to reversion to the C:G starting state. BER was impeded by BE2 and BE3, which inhibited UDG. The U:G mismatch was also processed by mismatch repair (MMR), which preferentially repaired the nicked strand of a mismatch. BE3 nicked the non-edited strand containing the G, favoring resolution of the U:G mismatch to the desired U:A or

T:A outcome. Figure 29B shows HEK293T cells treated as described in the Materials and Methods in the Examples below. The percentage of total DNA sequencing read with Ts at the target positions indicated show treatment with BE1, BE2, or BE3, or for treatment with wt Cas9 with a donor HDR template. Figure 29C shows frequency of indel formation following the treatment in Figure 29B. Values are listed in Figure 34. For Figures 29B and 29C, values and error bars reflect the mean and s.d. of three independent biological replicates performed on different days.

[0066] **Figures 30A to 30B** show BE3-mediated correction of two disease-relevant mutations in mammalian cells. The sequence of the protospacer is shown to the right of the mutation, with the PAM in blue and the target base in red with a subscripted number indicating its position within the protospacer. Underneath each sequence are the percentages of total sequencing reads with the corresponding base. Cells were treated as described in the Materials and Methods. Figure 30A shows the Alzheimer's disease-associated APOE4 allele converted to APOE3r in mouse astrocytes by BE3 in 74.9% of total reads. Two nearby Cs were also converted to Ts, but with no change to the predicted sequence of the resulting protein. Identical treatment of these cells with wt Cas9 and donor ssDNA results in only 0.3% correction, with 26.1% indel formation. Figure 30B shows the cancer associated p53 Y163C mutation corrected by BE3 in 7.6% of nucleofected human breast cancer cells with 0.7% indel formation. Identical treatment of these cells with wt Cas9 and donor ssDNA results in no mutation correction with 6.1% indel formation. This figure depicts SEQ ID NOs: 675 to 680 from top to bottom, respectively.

[0067] **Figure 31** shows activities of BE1, BE2, and BE3 at HEK293 site 2 off-targets. HEK293T cells were transfected with plasmids expressing BE1, BE2, or BE3 and a sgRNA matching the HEK293 site 2 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus all of the known Cas9 and dCas9 off-target loci for the HEK293 site 2 sgRNA, as previously determined by Joung and coworkers using the GUIDE-seq method (63), and Adli and coworkers using chromatin immunoprecipitation high-throughput sequencing (ChIP-seq) experiments (18). Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for BE1, BE2, and BE3. On the far right are displayed the total number of sequencing reads

reported , and the ChIP-seq signal intensity reported for each sequence. This figure depicts SEQ ID NOs: 681 to 688 from top to bottom, respectively.

[0068] **Figure 32** shows activities of BE1, BE2, and BE3 at HEK293 site 3 off-targets. HEK293T cells were transfected with plasmids expressing BE1, BE2, or BE3 and a sgRNA matching the HEK293 site 3 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus all of the known Cas9 off-target loci and the top five known dCas9 off-target loci for the HEK293 site 3 sgRNA, as previously determined by Joung and coworkers using the GUIDE-seq method[54], and using chromatin immunoprecipitation high-throughput sequencing (ChIP-seq) experiments[61]. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for BE1, BE2, and BE3. On the far right are displayed the total number of sequencing reads reported, and the ChIP-seq signal intensity reported for each sequence. This figure depicts SEQ ID NOs: 689 to 699 from top to bottom, respectively.

[0069] **Figure 33** shows activities of BE1, BE2, and BE3 at HEK293 site 4 off-targets. HEK293T cells were transfected with plasmids expressing BE1, BE2, or BE3 and a sgRNA matching the HEK293 site 4 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci, plus the top ten known Cas9 off-target loci and the top five known dCas9 off-target loci for the HEK293 site 4 sgRNA, as previously determined using the GUIDE-seq method[54], and using chromatin immunoprecipitation high-throughput sequencing (ChIP-seq) experiments[61]. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for BE1, BE2, and BE3. On the far right are displayed the total number of sequencing reads reported, and the ChIP-seq signal intensity reported for each sequence. This figure depicts SEQ ID NOs: 700 to 712 from top to bottom, respectively.

[0070] **Figure 34** shows mutation rates of non-protospacer bases following BE3-mediated correction of the Alzheimer's disease-associated *APOE4* allele to *APOE3r* in mouse astrocytes. The DNA sequence of the 50 bases on either side of the protospacer from Figure 30A and Figure 34B is shown with each base's position relative to the protospacer. The side

of the protospacer distal to the PAM is designated with positive numbers, while the side that includes the PAM is designated with negative numbers, with the PAM shown in blue. Underneath each sequence are the percentages of total DNA sequencing reads with the corresponding base for untreated cells, for cells treated with BE3 and an sgRNA targeting the *APOE4* C158R mutation, or for cells treated with BE3 and an sgRNA targeting the *VEGFA* locus. Neither BE3-treated sample resulted in mutation rates above those of untreated controls. This figure depicts SEQ ID NOs: 713 to 716 from top to bottom, respectively.

[0071] **Figure 35** shows mutation rates of non-protospacer bases following BE3-mediated correction of the cancer-associated p53 Y163C mutation in HCC1954 human cells. The DNA sequence of the 50 bases on either side of the protospacer from Figure 30B and Figure 39Bis shown with each base's position relative to the protospacer. The side of the protospacer distal to the PAM is designated with positive numbers, while the side that includes the PAM is designated with negative numbers, with the PAM shown in blue. Underneath each sequence are the percentages of total sequencing reads with the corresponding base for untreated cells, for cells treated with BE3 and an sgRNA targeting the *TP53* Y163C mutation, or for cells treated with BE3 and an sgRNA targeting the *VEGFA* locus. Neither BE3-treated sample resulted in mutational rates above those of untreated controls. This figure depicts SEQ ID NOs: 717 to 720 from top to bottom, respectively.

[0072] **Figures 36A to 36F** show the effects of deaminase, linker length, and linker composition on base editing. Figure 36A shows a gel-based deaminase assay showing activity of rAPOBEC1, pmCDA1, hAID, hAPOBEC3G, rAPOBEC1-GGS-dCas9, rAPOBEC1- $(GGS)_3$(SEQ ID NO: 596)-dCas9, and dCas9-$(GGS)_3$(SEQ ID NO: 596)-rAPOBEC1 on ssDNA. Enzymes were expressed in a mammalian cell lysate-derived *in vitro* transcription-translation system and incubated with 1.8 μM dye-conjugated ssDNA and USER enzyme (uracil DNA glycosylase and endonuclease VIII) at 37 °C for 2 hours. The resulting DNA was resolved on a denaturing polyacrylamide gel and imaged. The positive control is a sequence with a U synthetically incorporated at the same position as the target C. Figure 36B shows coomassie-stained denaturing PAGE gel of the expressed and purified proteins used in Figures 36C to 36F. Figures 36C to 36F show gel-based deaminase assay showing the deamination window of base editors with deaminase–Cas9 linkers of GGS (Figure 36C), $(GGS)_3$ (SEQ ID NO: 596) (Figure 36D), XTEN (Figure 36E), or $(GGS)_7$ (SEQ ID NO: 597) (Figure 36F). Following incubation of 1.85 μM deaminase-dCas9 fusions complexed with sgRNA with 125 nM dsDNA substrates at 37 °C for 2 hours, the dye-conjugated DNA was isolated and incubated with USER enzyme at 37 °C for 1 hour to cleave

the DNA backbone at the site of any uracils. The resulting DNA was resolved on a denaturing polyacrylamide gel, and the dye-conjugated strand was imaged. Each lane is numbered according to the position of the target C within the protospacer, or with – if no target C is present. 8U is a positive control sequence with a U synthetically incorporated at position 8. .

[0073] **Figures 37A to 37C** show BE1 base editing efficiencies are dramatically decreased in mammalian cells. Figure 37A Protospacer (black and red) and PAM (blue) sequences of the six mammalian cell genomic loci targeted by base editors. Target Cs are indicated in red with subscripted numbers corresponding to their positions within the protospacer. Figure 37B shows synthetic 80-mers with sequences matching six different genomic sites were incubated with BE1 then analyzed for base editing by HTS. For each site, the sequence of the protospacer is indicated to the right of the name of the site, with the PAM highlighted in blue. Underneath each sequence are the percentages of total DNA sequencing reads with the corresponding base. We considered a target C as "editable" if the *in vitro* conversion efficiency is >10%. Note that maximum yields are 50% of total DNA sequencing reads since the non-targeted strand is unaffected by BE1. Values are shown from a single experiment. Figure 37C shows HEK293T cells were transfected with plasmids expressing BE1 and an appropriate sgRNA. Three days after transfection, genomic DNA was extracted and analyzed by high-throughput DNA sequencing at the six loci. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, are shown for BE1 at all six genomic loci. Values and error bars of all data from HEK293T cells reflect the mean and standard deviation of three independent biological replicates performed on different days. Figure 37A depicts SEQ ID NOs: 721 to 726 from top to bottom, respectively. Figure 37B depicts SEQ ID NOs: 727 to 732 from top to bottom, respectively.

[0074] **Figure 38** shows base editing persists over multiple cell divisions. Cellular C to T conversion percentages by BE2 and BE3 are shown for HEK293 sites 3 and 4 in HEK293T cells before and after passaging the cells. HEK293T cells were nucleofected with plasmids expressing BE2 or BE3 and an sgRNA targeting HEK293 site 3 or 4. Three days after nucleofection, the cells were harvested and split in half. One half was subjected to HTS analysis, and the other half was allowed to propagate for approximately five cell divisions, then harvested and subjected to HTS analysis. Values and error bars reflect the mean and standard deviation of at least two biological experiments.

[0075] **Figures 39A to 39C** show non-target C/G mutation rates. Shown here are the C to T and G to A mutation rates at 2,500 distinct cytosines and guanines surrounding the six on-target and 34 off-target loci tested, representing a total of 14,700,000 sequence reads derived from approximately $1.8x10^6$ cells. Figures 39A and 39B show cellular non-target C to T and G to A conversion percentages by BE1, BE2, and BE3 are plotted individually against their positions relative to a protospacer for all 2,500 cytosines/guanines. The side of the protospacer distal to the PAM is designated with positive numbers, while the side that includes the PAM is designated with negative numbers. Figure 39C shows average non-target cellular C to T and G to A conversion percentages by BE1, BE2, and BE3 are shown, as well as the highest and lowest individual conversion percentages.

[0076] **Figures 40A to 40B** show additional data sets of BE3-mediated correction of two disease-relevant mutations in mammalian cells**.** For each site, the sequence of the protospacer is indicated to the right of the name of the mutation, with the PAM highlighted in blue and the base responsible for the mutation indicated in red bold with a subscripted number corresponding to its position within the protospacer. The amino acid sequence above each disease-associated allele is shown, together with the corrected amino acid sequence following base editing in green. Underneath each sequence are the percentages of total sequencing reads with the corresponding base. Cells were nucleofected with plasmids encoding BE3 and an appropriate sgRNA. Two days after nucleofection, genomic DNA was extracted from the nucleofected cells and analyzed by HTS to assess pathogenic mutation correction. Figure 40A shows the Alzheimer's disease-associated *APOE4* allele is converted to *APOE3r* in mouse astrocytes by BE3 in 58.3% of total reads only when treated with the correct sgRNA. Two nearby Cs are also converted to Ts, but with no change to the predicted sequence of the resulting protein. Identical treatment of these cells with wt Cas9 and donor ssDNA results in 0.2% correction, with 26.7% indel formation. Figure 40B shows the cancer-associated p53 Y163C mutation is corrected by BE3 in 3.3% of nucleofected human breast cancer cells only when treated with the correct sgRNA. Identical treatment of these cells with wt Cas9 and donor ssDNA results in no detectable mutation correction with 8.0% indel formation. Figures 40A to 40B depict SEQ ID NOs: 733 to 740 from top to bottom, respectively.

[0077] **Figure 41** shows a schematic representation of an exemplary USER (Uracil-Specific Excision Reagent) Enzyme-based assay, which may be used to test the activity of various deaminases on single-stranded DNA (ssDNA) substrates.

[0078] **Figure 42** is a schematic of the pmCDA-nCas9-UGI-NLS construct and its activity at the HeK-3 site relative to the base editor (rAPOBEC1) and the negative control (untreated).

[0079] **Figure 43** is a schematic of the pmCDA1-XTEN-nCas9-UGI-NLS construct and its activity at the HeK-3 site relative to the base editor (rAPOBEC1) and the negative control (untreated).

[0080] **Figure 44** shows the percent of total sequencing reads with target C converted to T using cytidine deaminases (CDA) or APOBEC.

[0081] **Figure 45** shows the percent of total sequencing reads with target C converted to A using deaminases (CDA) or APOBEC.

[0082] **Figure 46** shows the percent of total sequencing reads with target C converted to G using deaminases (CDA) or APOBEC.

[0083] **Figure 47** is a schematic of the huAPOBEC3G-XTEN-nCas9-UGI-NLS construct and its activity at the HeK-2 site relative to a mutated form (huAPOBEC3G*(D316R_D317R)-XTEN-nCas9-UGI-NLS, the base editor (rAPOBEC1) and the negative control (untreated).

[0084] **Figure 48** shows the schematic of the LacZ construct used in the selection assay of Example 7.

[0085] **Figure 49** shows reversion data from different plasmids and constructs.

[0086] **Figure 50** shows the verification of lacZ reversion and the purification of reverted clones.

[0087] **Figure 51** is a schematic depicting a deamination selection plasmid used in Example 7.

[0088] **Figure 52** shows the results of a chloramphenicol reversion assay (pmCDA1 fusion).

[0089] **Figures 53A** to **53B** demonstrated DNA correction induction of two constructs.

[0090] **Figure 54** shows the results of a chloramphenicol reversion assay (huAPOBEC3G fusion).

[0091] **Figure 55** shows the activities of BE3 and HF-BE3 at EMX1 off-targets. The sequences, from top to bottom, correspond to SEQ ID NOs: 286-292, 299-301.

[0092] **Figure 56** shows on-target base editing efficiencies of BE3 and HF-BE3.

[0093] **Figure 57** is a graph demonstrating that mutations affect cytidine deamination with varying degrees. Combinations of mutations that each slightly impairs catalysis allow

selective deamination at one position over others. The FANCF site was GGAATC$_6$C$_7$C$_8$TTC$_{11}$TGCAGCACCTGG (SEQ ID NO: 303).

[0094] **Figure 58** is a schematic depicting next generation base editors.

[0095] **Figure 59** is a schematic illustrating new base editors made from Cas9 variants.

[0096] **Figure 60** shows the base-edited percentage of different NGA PAM sites.

[0097] **Figure 61** shows the base-edited percentage of cytidines using NGCG PAM EMX (VRER BE3) and the C$_1$TC$_3$C$_4$C$_5$ATC$_8$AC$_{10}$ATCAACCGGT (SEQ ID NO: 304) spacer.

[0098] **Figure 62** shows the based-edited percentages resulting from different NNGRRT PAM sites.

[0099] **Figure 63** shows the based-edited percentages resulting from different NNHRRT PAM sites.

[00100] **Figures 64A** to **64C** show the base-edited percentages resulting from different TTTN PAM sites using Cpf1 BE2. The spacers used were: TTTCCTC$_3$C$_4$C$_5$C$_6$C$_7$C$_8$C$_9$AC$_{11}$AGGTAGAACAT (Figure 64A, SEQ ID NO: 305), TTTCC$_1$C$_2$TC$_4$TGTC$_8$C$_9$AC$_{11}$ACCCTCATCCTG (Figure 64B, SEQ ID NO: 306), and TTTCC$_1$C$_2$C$_3$AGTC$_7$C$_8$TC$_{10}$C$_{11}$AC$_{13}$AC$_{15}$C$_{16}$C$_{17}$TGAAAC (Figure 64C, SEQ ID NO: 307).

[00101] **Figure 65** is a schematic depicting selective deamination as achieved through kinetic modulation of cytidine deaminase point mutagenesis.

[00102] **Figure 66** is a graph showing the effect of various mutations on the deamination window probed in cell culture with multiple cytidines in the spacer. The spacer used was: TGC$_3$C$_4$C$_5$C$_6$TC$_8$C$_9$C$_{10}$TC$_{12}$C$_{13}$C$_{14}$TGGCCC (SEQ ID NO: 308).

[00103] **Figure 67** is a graph showing the effect of various mutations on the deamination window probed in cell culture with multiple cytidines in the spacer. The spacer used was: AGAGC$_5$C$_6$C$_7$C$_8$C$_9$C$_{10}$C$_{11}$TC$_{13}$AAAGAGA (SEQ ID NO: 309).

[00104] **Figure 68** is a graph showing the effect of various mutations on the FANCF site with a limited number of cytidines. The spacer used was: GGAATC$_6$C$_7$C$_8$TTC$_{11}$TGCAGCACCTGG (SEQ ID NO: 303). Note that the triple mutant (W90Y, R126E, R132E) preferentially edits the cytidine at the sixth position.

[00105] **Figure 69** is a graph showing the effect of various mutations on the HEK3 site with a limited number of cytidines. The spacer used was: GGCC$_4$C$_5$AGACTGAGCACGTGATGG (SEQ ID NO: 310). Note that the double and triple mutants preferentially edit the cytidine at the fifth position over the cytidine in the fourth position.

[00106] **Figure 70** is a graph showing the effect of various mutations on the EMX1 site with a limited number of cytidines. The spacer used was: GAGTC$_5$C$_6$GAGCAGAAGAAGAAGGG (SEQ ID NO: 311). Note that the triple mutant only edits the cytidine at the fifth position, not the sixth.

[00107] **Figure 71** is a graph showing the effect of various mutations on the HEK2 site with a limited number of cytidines. The spacer used was: GAAC$_4$AC$_6$AAAGCATAGACTGCGGG (SEQ ID NO: 312).

[00108] **Figure 72** shows on-target base editing efficiencies of BE3 and BE3 comprising mutations W90Y R132E in immortalized astrocytes.

[00109] **Figure 73** depicts a schematic of three Cpf1 fusion constructs.

[00110] **Figures 74** shows a comparison of plasmid delivery of BE3 and HF-BE3 (EMX1, FANCF, and RNF2).

[00111] **Figure 75** shows a comparison of plasmid delivery of BE3 and HF-BE3 (HEK3 and HEK 4).

[00112] **Figure 76** shows off-target editing of EMX-1 at all 10 sites.

[00113] **Figure 77** shows deaminase protein lipofection to HEK cells using a GAGTCCGAGCAGAAGAAGAAG (SEQ ID NO: 313) spacer. The EMX-1 on-target and EMX-1 off target site 2 were examined.

[00114] **Figure 78** shows deaminase protein lipofection to HEK cells using a GGAATCCCTTCTGCAGCACCTGG (SEQ ID NO: 314) spacer. The FANCF on target and FANCF off target site 1 were examined.

[00115] **Figure 79** shows deaminase protein lipofection to HEK cells using a GGCCCAGACTGAGCACGTGA (SEQ ID NO: 315) spacer. The HEK-3 on target site was examined.

[00116] **Figure 80** shows deaminase protein lipofection to HEK cells using a GGCACTGCGGCTGGAGGTGGGGG (SEQ ID NO: 316) spacer. The HEK-4 on target, off target site 1, site 3, and site 4.

[00117] **Figure 81** shows the results of an *in vitro* assay for sgRNA activity for sgHR_13 (GTCAGGTCGAGGGTTCTGTC (SEQ ID NO: 317) spacer; C8 target: G51 to STOP), sgHR_14 (GGGCCGCAGTATCCTCACTC (SEQ ID NO: 318) spacer; C7 target; C7 target: Q68 to STOP), and sgHR_15 (CCGCCAGTCCCAGTACGGGA (SEQ ID NO: 319) spacer; C10 and C11 are targets: W239 or W237 to STOP).

[00118] **Figure 82** shows the results of an *in vitro* assay for sgHR_17 (CAACCACTGCTCAAAGATGC (SEQ ID NO: 320) spacer; C4 and C5 are targets: W410

to STOP), and sgHR_16 (CTTCCAGGATGAGAACACAG (SEQ ID NO: 321) spacer; C4 and C5 are targets: W273 to STOP).

[00119] **Figure 83** shows the direct injection of BE3 protein complexed with sgHR_13 in zebrafish embryos.

[00120] **Figure 84** shows the direct injection of BE3 protein complexed with sgHR_16 in zebrafish embryos.

[00121] **Figure 85** shows the direct injection of BE3 protein complexed with sgHR_17 in zebrafish embryos.

[00122] **Figure 86** shows exemplary nucleic acid changes that may be made using base editors that are capable of making a cytosine to thymine change.

[00123] **Figure 87** shows an illustration of apolipoprotein E (APOE) isoforms, demonstrating how a base editor (*e.g.,* BE3) may be used to edit one APOE isoform (*e.g.,* APOE4) into another APOE isoform (*e.g.,* APOE3r) that is associated with a decreased risk of Alzheimer's disease.

[00124] **Figure 88** shows base editing of APOE4 to APOE3r in mouse astrocytes.

[00125] **Figure 89** shows base editing of PRNP to cause early truncation of the protein at arginine residue 37.

[00126] **Figure 90** shows that knocking out UDG (which UGI inhibits) dramatically improves the cleanliness of efficiency of C to T base editing.

[00127] **Figure 91** shows that use of a base editor with the nickase but without UGI leads to a mixture of outcomes, with very high indel rates.

[00128] **Figures 92A** to **92G** show that SaBE3, SaKKH-BE3, VQR-BE3, EQR-BE3, and VRER-BE3 mediate efficient base editing at target sites containing non-NGG PAMs in human cells. Figure 92A shows base editor architectures using *S. pyogenes* and *S. aureus* Cas9. Figure 92B shows recently characterized Cas9 variants with alternate or relaxed PAM requirements. Figures 92C and 92D show HEK293T cells treated with the base editor variants shown as described in Example 12. The percentage of total DNA sequencing reads (with no enrichment for transfected cells) with C converted to T at the target positions indicated are shown. The PAM sequence of each target tested is shown below the X-axis. The charts show the results for SaBE3 and SaKKH-BE3 at genomic loci with NNGRRT PAMs (Figure 92C), SaBE3 and SaKKH-BE3 at genomic loci with NNNRRT PAMs (Figure 92D), VQR-BE3 and EQR-BE3 at genomic loci with NGAG PAMs (Figure 92E), and with NGAH PAMs (Figure 92F), and VRER-BE3 at genomic loci with NGCG PAMs (Figure 92G).

Values and error bars reflect the mean and standard deviation of at least two biological replicates.

[00129] **Figures 93A** to **93C** demonstrate that base editors with mutations in the cytidine deaminase domain exhibit narrowed editing windows. Figures 93A to 93C show HEK293T cells transfected with plasmids expressing mutant base editors and an appropriate sgRNA. Three days after transfection, genomic DNA was extracted and analyzed by high-throughput DNA sequencing at the indicated loci. The percentage of total DNA sequencing reads (without enrichment for transfected cells) with C changed to T at the target positions indicated are shown for the EMX1 site, HEK293 site 3, FANCF site, HEK293 site 2, site A, and site B loci. Figure 93A illustrates certain cytidine deaminase mutations which narrow the base editing window. See Figure 98 for the characterization of additional mutations. Figure 93B shows the effect of cytidine deaminase mutations which effect the editing window width on genomic loci. Combining beneficial mutations has an additive effect on narrowing the editing window. Figure 93C shows that YE1-BE3, YE2-BE3, EE-BE3, and YEE-BE3 effect the product distribution of base editing, producing predominantly singly-modified products in contrast with BE3. Values and error bars reflect the mean and standard deviation of at least two biological replicates.

[00130] **Figures 94A** and **94B** show genetic variants from ClinVar that in principle can be corrected by the base editors developed in this work. The NCBI ClinVar database of human genetic variations and their corresponding phenotypes was searched for genetic diseases that in theory can be corrected by base editing. Figure 94A demonstrates improvement in base editing targeting scope among all pathogenic T→C mutations in the ClinVar database through the use of base editors with altered PAM specificities. The white fractions denote the proportion of pathogenic T→C mutations accessible on the basis of the PAM requirements of either BE3, or BE3 together with the five modified-PAM base editors developed in this work. Figure 94B shows improvement in base editing targeting scope among all pathogenic T→C mutations in the ClinVar database through the use of base editors with narrowed activity windows. BE3 was assumed to edit Cs in positions 4-8 with comparable efficiency as shown in Figures 93A to 93C. YEE-BE3 was assumed to edit with C5>C6>C7>others preference within its activity window. The white fractions denote the proportion of pathogenic T→C mutations that can be edited BE3 without comparable editing of other Cs (left), or that can be edited BE3 or YEE-BE3 without comparable editing of other Cs (right).

[00131] **Figures 95A** to **95C** show the effect of truncated guide RNAs on base editing window width. HEK293T cells were transfected with plasmids expressing BE3 and sgRNAs

of different 5' truncation lengths. The treated cells were analyzed as described in the Examples. **Figure 95A** shows protospacer and PAM sequence (top, SEQ ID NO: 4270) and cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, at a site within the EMX1 genomic locus. At this site, the base editing window was altered through the use of a 17-nt truncated gRNA. **Figure 95B** shows protospacer and PAM sequences (top, SEQ ID NO: 4270) and cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, at sites within the HEK site 3 and site 4 genomic loci. At these sites, no change in the base editing window was observed, but a linear decrease in editing efficiency for all substrate bases as the sgRNA is truncated was noted.

[00132] **Figure 96** shows the effect of APOBEC1-Cas9 linker lengths on base editing window width. HEK293T cells were transfected with plasmids expressing base editors with rAPOBEC1–Cas9 linkers of XTEN, GGS, $(GGS)_3$ (SEQ ID NO: 596), $(GGS)_5$ (SEQ ID NO: 4271), or $(GGS)_7$ (SEQ ID NO: 597) and an sgRNA. The treated cells were analyzed as described in the Examples. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, are shown for the various base editors with different linkers.

[00133] **Figures 97A** to **97C** show the effect of rAPOBEC mutations on base editing window width. **Figure 97C** shows HEK293T cells transfected with plasmids expressing an sgRNA targeting either Site A or Site B and the BE3 point mutants indicated. The treated cells were analyzed as described in the Examples. All C's in the protospacer and within three basepairs of the protospacer are displayed and the cellular C to T conversion percentages are shown. The 'editing window widths', defined as the calculated number of nucleotides within which editing efficiency exceeds the half-maximal value, are displayed for all tested mutants.

[00134] **Figure 98** shows the effect of APOBEC1 mutation son product distributions of base editing in mammalian cells. HEK293T cells were transfected with plasmids expressing BE3 or its mutants and an appropriate sgRNAs. The treated cells were analyzed as described in the Examples. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, are shown (left). Percent of total sequencing reads containing the C to T conversion is shown on the right. The BE3 point mutants do not significantly affect base editing efficiencies at HEK site 4, a site with only one target cytidine.

[00135] **Figure 99** shows a comparison of on-target editing plasma delivery in BE3 and HF-BE3.

[00136] **Figure 100** shows a comparison of on-target editing in protein and plasma delivery of BE3.

[00137] **Figure 101** shows a comparison of on-target editing in protein and plasma devliery of HF-BE3.

[00138] **Figure 102** shows that both lipofection and installing HF mutations decrease off-target deamination events. The diamond indicates no off targets were detected and the specificity ratio was set to 100.

[00139] **Figure 103** shows *in vitro* C to T editing on a synthetic substrate with Cs placed at even positions in the protospacer ($NNNNTC_2TC_4TC_6TC_8TC_{10}TC_{12}TC_{14}TC_{16}TC_{18}TC_{20}NGG$, SEQ ID NO: 4272).

[00140] **Figure 104** shows *in vitro* C to T editing on a synthetic substrate with Cs placed at odd positions in the protospacer ($NNNNTC_2TC_4TC_6TC_8TC_{10}TC_{12}TC_{14}TC_{16}TC_{18}TC_{20}NGG$, SEQ ID NO: 4272).

[00141] **Figure 105** includes two graphs depicting the specificity ratio of base editing with plasmid vs. protein delivery.

[00142] **Figures 106A** to **106B** shows BE3 activity on non-NGG PAM sites. HEK293T cells were transfected with plasmids expressing BE3 and appropriate sgRNA. The treated cells were analyzed as described in the Examples. **Figure 106A** shows BE3 activity on sites can be efficiently targeted by SaBE3 or SaKKH-BE3. BE3 shows low but significant activity on the NAG PAM. **Figure 106B** shows BE3 has significantly reduced editing at sites with NGA or NGCG PAMs, in contrast to VQR-BE3 or VRER-BE3.

[00143] **Figures 107A** to **107B** show the effect of APOBEC1 mutations on VQR-BE3 and SaKKH-BE3. HEK293T cells were transfected with plasmids expressing VQR-BE3, SaKKH-BE3 or its mutants and an appropriate sgRNAs. The treated cells were analyzed as described in the Methods. Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with Ts at the target positions indicated, are shown. **Figure 107A** shows that the window-modulating mutations can be applied to VQR-BE3 to enable selective base editing at sites targetable by NGA PAM. **Figure 107B** shows that, when applied to SaKKH-BE3, the mutations cause overall decrease in base editing efficiency without conferring base selectivity within the target window.

[00144] **Figure 108** shows a schematic representation of nucleotide editing. The following abbreviations are used: (MMR) - mismatch repair, (BE3 Nickase) - refers to base editor 3, which comprises a Cas9 nickase domain, (UGI) - uracil glycosylase inhibitor, UDG) - uracil DNA glycosylase, (APOBEC) – refers to an APOBEC cytidine deaminase.

DEFINITIONS

[00145] As used herein and in the claims, the singular forms "a," "an," and "the" include the singular and the plural reference unless the context clearly indicates otherwise. Thus, for example, a reference to "an agent" includes a single agent and a plurality of such agents.

[00146] The term "Cas9" or "Cas9 nuclease" refers to an RNA-guided nuclease comprising a Cas9 protein, or a fragment thereof (*e.g.*, a protein comprising an active, inactive, or partially active DNA cleavage domain of Cas9, and/or the gRNA binding domain of Cas9). A Cas9 nuclease is also referred to sometimes as a casn1 nuclease or a CRISPR (clustered regularly interspaced short palindromic repeat)-associated nuclease. CRISPR is an adaptive immune system that provides protection against mobile genetic elements (viruses, transposable elements and conjugative plasmids). CRISPR clusters contain spacers, sequences complementary to antecedent mobile elements, and target invading nucleic acids. CRISPR clusters are transcribed and processed into CRISPR RNA (crRNA). In type II CRISPR systems correct processing of pre-crRNA requires a trans-encoded small RNA (tracrRNA), endogenous ribonuclease 3 (rnc) and a Cas9 protein. The tracrRNA serves as a guide for ribonuclease 3-aided processing of pre-crRNA. Subsequently, Cas9/crRNA/tracrRNA endonucleolytically cleaves linear or circular dsDNA target complementary to the spacer. The target strand not complementary to crRNA is first cut endonucleolytically, then trimmed 3′-5′ exonucleolytically. In nature, DNA-binding and cleavage typically requires protein and both RNAs. However, single guide RNAs ("sgRNA", or simply "gNRA") can be engineered so as to incorporate aspects of both the crRNA and tracrRNA into a single RNA species. See, *e.g.*, Jinek M., Chylinski K., Fonfara I., Hauer M., Doudna J.A., Charpentier E. *Science* 337:816-821(2012), the entire contents of which is hereby incorporated by reference. Cas9 recognizes a short motif in the CRISPR repeat sequences (the PAM or protospacer adjacent motif) to help distinguish self versus non-self. Cas9 nuclease sequences and structures are well known to those of skill in the art (see, *e.g.*, "Complete genome sequence of an M1 strain of Streptococcus pyogenes." Ferretti *et al.*, J.J., McShan W.M., Ajdic D.J., Savic D.J., Savic G., Lyon K., Primeaux C., Sezate S., Suvorov A.N., Kenton S., Lai H.S., Lin S.P., Qian Y., Jia H.G., Najar F.Z., Ren Q., Zhu H., Song L., White J., Yuan X., Clifton S.W., Roe B.A., McLaughlin R.E., Proc. Natl. Acad. Sci. U.S.A. 98:4658-4663(2001); "CRISPR RNA maturation by trans-encoded small RNA and host factor RNase III." Deltcheva E., Chylinski K., Sharma C.M., Gonzales K., Chao Y., Pirzada Z.A., Eckert M.R., Vogel J., Charpentier E., Nature 471:602-607(2011); and "A

programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity." Jinek M., Chylinski K., Fonfara I., Hauer M., Doudna J.A., Charpentier E. *Science* 337:816-821(2012), the entire contents of each of which are incorporated herein by reference). Cas9 orthologs have been described in various species, including, but not limited to, *S. pyogenes* and *S. thermophilus*. Additional suitable Cas9 nucleases and sequences will be apparent to those of skill in the art based on this disclosure, and such Cas9 nucleases and sequences include Cas9 sequences from the organisms and loci disclosed in Chylinski, Rhun, and Charpentier, "The tracrRNA and Cas9 families of type II CRISPR-Cas immunity systems" (2013) RNA Biology 10:5, 726-737; the entire contents of which are incorporated herein by reference. In some embodiments, a Cas9 nuclease has an inactive (*e.g.*, an inactivated) DNA cleavage domain, that is, the Cas9 is a nickase.

[00147] A nuclease-inactivated Cas9 protein may interchangeably be referred to as a "dCas9" protein (for nuclease-"dead" Cas9). Methods for generating a Cas9 protein (or a fragment thereof) having an inactive DNA cleavage domain are known (See, *e.g.*, Jinek *et al., Science.* 337:816-821(2012); Qi *et al.,* "Repurposing CRISPR as an RNA-Guided Platform for Sequence-Specific Control of Gene Expression" (2013) *Cell.* 28;152(5):1173-83, the entire contents of each of which are incorporated herein by reference). For example, the DNA cleavage domain of Cas9 is known to include two subdomains, the HNH nuclease subdomain and the RuvC1 subdomain. The HNH subdomain cleaves the strand complementary to the gRNA, whereas the RuvC1 subdomain cleaves the non-complementary strand. Mutations within these subdomains can silence the nuclease activity of Cas9. For example, the mutations D10A and H840A completely inactivate the nuclease activity of *S. pyogenes* Cas9 (Jinek *et al., Science.* 337:816-821(2012); Qi *et al., Cell.* 28;152(5):1173-83 (2013)). In some embodiments, proteins comprising fragments of Cas9 are provided. For example, in some embodiments, a protein comprises one of two Cas9 domains: (1) the gRNA binding domain of Cas9; or (2) the DNA cleavage domain of Cas9. In some embodiments, proteins comprising Cas9 or fragments thereof are referred to as "Cas9 variants." A Cas9 variant shares homology to Cas9, or a fragment thereof. For example a Cas9 variant is at least about 70% identical, at least about 80% identical, at least about 90% identical, at least about 95% identical, at least about 96% identical, at least about 97% identical, at least about 98% identical, at least about 99% identical, at least about 99.5% identical, or at least about 99.9% identical to wild type Cas9. In some embodiments, the Cas9 variant may have 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 21, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50 or more amino acid

changes compared to wild type Cas9. In some embodiments, the Cas9 variant comprises a fragment of Cas9 (*e.g.*, a gRNA binding domain or a DNA-cleavage domain), such that the fragment is at least about 70% identical, at least about 80% identical, at least about 90% identical, at least about 95% identical, at least about 96% identical, at least about 97% identical, at least about 98% identical, at least about 99% identical, at least about 99.5% identical, or at least about 99.9% identical to the corresponding fragment of wild type Cas9. In some embodiments, the fragment is is at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95% identical, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% of the amino acid length of a corresponding wild type Cas9.

[00148] In some embodiments, the fragment is at least 100 amino acids in length. In some embodiments, the fragment is at least 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, or at least 1300 amino acids in length. In some embodiments, wild type Cas9 corresponds to Cas9 from *Streptococcus pyogenes* (NCBI Reference Sequence: NC_017053.1, SEQ ID NO:1 (nucleotide); SEQ ID NO:2 (amino acid)).

ATGGATAAGAAATACTCAATAGGCTTAGATATCGGCACAAATAGCGTCGGATGG
GCGGTGATCACTGATGATTATAAGGTTCCGTCTAAAAAGTTCAAGGTTCTGGGAA
ATACAGACCGCCACAGTATCAAAAAAAATCTTATAGGGGCTCTTTTATTTGGCAG
TGGAGAGACAGCGGAAGCGACTCGTCTCAAACGGACAGCTCGTAGAAGGTATAC
ACGTCGGAAGAATCGTATTTGTTATCTACAGGAGATTTTTTCAAATGAGATGGCG
AAAGTAGATGATAGTTTCTTTCATCGACTTGAAGAGTCTTTTTTGGTGGAAGAAG
ACAAGAAGCATGAACGTCATCCTATTTTTGGAAATATAGTAGATGAAGTTGCTTA
TCATGAGAAATATCCAACTATCTATCATCTGCGAAAAAAATTGGCAGATTCTACT
GATAAAGCGGATTTGCGCTTAATCTATTTGGCCTTAGCGCATATGATTAAGTTTC
GTGGTCATTTTTTGATTGAGGGAGATTTAAATCCTGATAATAGTGATGTGGACAA
ACTATTTATCCAGTTGGTACAAATCTACAATCAATTATTTGAAGAAAACCCTATT
AACGCAAGTAGAGTAGATGCTAAAGCGATTCTTTCTGCACGATTGAGTAAATCA
AGACGATTAGAAAATCTCATTGCTCAGCTCCCCGGTGAGAAGAGAAATGGCTTG
TTTGGGAATCTCATTGCTTTGTCATTGGGATTGACCCCTAATTTTAAATCAAATTT
TGATTTGGCAGAAGATGCTAAATTACAGCTTTCAAAAGATACTTACGATGATGAT
TTAGATAATTTATTGGCGCAAATTGGAGATCAATATGCTGATTTGTTTTTGGCAG
CTAAGAATTTATCAGATGCTATTTTACTTTCAGATATCCTAAGAGTAAATAGTGA
AATAACTAAGGCTCCCCTATCAGCTTCAATGATTAAGCGCTACGATGAACATCAT
CAAGACTTGACTCTTTTAAAAGCTTTAGTTCGACAACAACTTCCAGAAAAGTATA
AAGAAATCTTTTTTGATCAATCAAAAAACGGATATGCAGGTTATATTGATGGGGG
AGCTAGCCAAGAAGAATTTTATAAATTTATCAAACCAATTTTAGAAAAAATGGAT
GGTACTGAGGAATTATTGGTGAAACTAAATCGTGAAGATTTGCTGCGCAAGCAA
CGGACCTTTGACAACGGCTCTATTCCCCATCAAATTCACTTGGGTGAGCTGCATG
CTATTTTGAGAAGACAAGAAGACTTTTATCCATTTTTAAAAGACAATCGTGAGAA
GATTGAAAAAATCTTGACTTTTCGAATTCCTTATTATGTTGGTCCATTGGCGCGTG

GCAATAGTCGTTTTGCATGGATGACTCGGAAGTCTGAAGAAACAATTACCCCATG
GAATTTTGAAGAAGTTGTCGATAAAGGTGCTTCAGCTCAATCATTTATTGAACGC
ATGACAAACTTTGATAAAAATCTTCCAAATGAAAAAGTACTACCAAAACATAGT
TTGCTTTATGAGTATTTTACGGTTTATAACGAATTGACAAAGGTCAAATATGTTA
CTGAGGGAATGCGAAAACCAGCATTTCTTTCAGGTGAACAGAAGAAAGCCATTG
TTGATTTACTCTTCAAAACAAATCGAAAAGTAACCGTTAAGCAATTAAAAGAAG
ATTATTTCAAAAAAATAGAATGTTTTGATAGTGTTGAAATTTCAGGAGTTGAAGA
TAGATTTAATGCTTCATTAGGCGCCTACCATGATTTGCTAAAAATTATTAAAGAT
AAAGATTTTTTGGATAATGAAGAAAATGAAGATATCTTAGAGGATATTGTTTTAA
CATTGACCTTATTTGAAGATAGGGGGATGATTGAGGAAAGACTTAAAACATATG
CTCACCTCTTTGATGATAAGGTGATGAAACAGCTTAAACGTCGCCGTTATACTGG
TTGGGGACGTTTGTCTCGAAAATTGATTAATGGTATTAGGGATAAGCAATCTGGC
AAAACAATATTAGATTTTTTGAAATCAGATGGTTTTGCCAATCGCAATTTTATGC
AGCTGATCCATGATGATAGTTTGACATTTAAAGAAGATATTCAAAAAGCACAGG
TGTCTGGACAAGGCCATAGTTTACATGAACAGATTGCTAACTTAGCTGGCAGTCC
TGCTATTAAAAAAGGTATTTTACAGACTGTAAAAATTGTTGATGAACTGGTCAAA
GTAATGGGGCATAAGCCAGAAAATATCGTTATTGAAATGGCACGTGAAAATCAG
ACAACTCAAAAGGGCCAGAAAAATTCGCGAGAGCGTATGAAACGAATCGAAGA
AGGTATCAAAGAATTAGGAAGTCAGATTCTTAAAGAGCATCCTGTTGAAAATAC
TCAATTGCAAAATGAAAAGCTCTATCTCTATTATCTACAAAATGGAAGAGACATG
TATGTGGACCAAGAATTAGATATTAATCGTTTAAGTGATTATGATGTCGATCACA
TTGTTCCACAAAGTTTCATTAAAGACGATTCAATAGACAATAAGGTACTAACGCG
TTCTGATAAAAATCGTGGTAAATCGGATAACGTTCCAAGTGAAGAAGTAGTCAA
AAAGATGAAAAACTATTGGAGACAACTTCTAAACGCCAAGTTAATCACTCAACG
TAAGTTTGATAATTTAACGAAAGCTGAACGTGGAGGTTTGAGTGAACTTGATAAA
GCTGGTTTTATCAAACGCCAATTGGTTGAAACTCGCCAAATCACTAAGCATGTGG
CACAAATTTTGGATAGTCGCATGAATACTAAATACGATGAAAATGATAAACTTAT
TCGAGAGGTTAAAGTGATTACCTTAAAATCTAAATTAGTTTCTGACTTCCGAAAA
GATTTCCAATTCTATAAAGTACGTGAGATTAACAATTACCATCATGCCCATGATG
CGTATCTAAATGCCGTCGTTGGAACTGCTTTGATTAAGAAATATCCAAAACTTGA
ATCGGAGTTTGTCTATGGTGATTATAAAGTTTATGATGTTCGTAAAATGATTGCT
AAGTCTGAGCAAGAAATAGGCAAAGCAACCGCAAAATATTTCTTTTACTCTAATA
TCATGAACTTCTTCAAAACAGAAATTACACTTGCAAATGGAGAGATTCGCAAAC
GCCCTCTAATCGAAACTAATGGGGAAACTGGAGAAATTGTCTGGGATAAAGGGC
GAGATTTTGCCACAGTGCGCAAAGTATTGTCCATGCCCCAAGTCAATATTGTCAA
GAAAACAGAAGTACAGACAGGCGGATTCTCCAAGGAGTCAATTTTACCAAAAAG
AAATTCGGACAAGCTTATTGCTCGTAAAAAAGACTGGGATCCAAAAAAATATGG
TGGTTTTGATAGTCCAACGGTAGCTTATTCAGTCCTAGTGGTTGCTAAGGTGGAA
AAAGGGAAATCGAAGAAGTTAAAATCCGTTAAAGAGTTACTAGGGATCACAATT
ATGGAAAGAAGTTCCTTTGAAAAAAATCCGATTGACTTTTTAGAAGCTAAAGGAT
ATAAGGAAGTTAAAAAAGACTTAATCATTAAACTACCTAAATATAGTCTTTTTGA
GTTAGAAAACGGTCGTAAACGGATGCTGGCTAGTGCCGGAGAATTACAAAAAGG
AAATGAGCTGGCTCTGCCAAGCAAATATGTGAATTTTTTATATTTAGCTAGTCAT
TATGAAAAGTTGAAGGGTAGTCCAGAAGATAACGAACAAAAACAATTGTTTGTG
GAGCAGCATAAGCATTATTTAGATGAGATTATTGAGCAAATCAGTGAATTTTCTA
AGCGTGTTATTTTAGCAGATGCCAATTTAGATAAAGTTCTTAGTGCATATAACAA
ACATAGAGACAAACCAATACGTGAACAAGCAGAAAATATTATTCATTTATTTAC
GTTGACGAATCTTGGAGCTCCCGCTGCTTTTAAATATTTTGATACAACAATTGATC
GTAAACGATATACGTCTACAAAAGAAGTTTTAGATGCCACTCTTATCCATCAATC

CATCACTGGTCTTTATGAAACACGCATTGATTTGAGTCAGCTAGGAGGTGACTGA (SEQ ID NO:1)

MDKKYSIGLDIGTNSVGWAVITDDYKVPSKKFKVLGNTDRHSIKKNLIGALLFGSGE TAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHE RHPIFGNIVDEVAYHEKYPTIYHLRKKLADSTDKADLRLIYLALAHMIKFRGHFLIEG DLNPDNSDVDKLFIQLVQIYNQLFEENPINASRVDAKAILSARLSKSRRLENLIAQLPG EKRNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYAD LFLAAKNLSDAILLSDILRVNSEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEK YKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRT FDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFA WMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTV YNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFD SVEISGVEDRFNASLGAYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDRGMIEER LKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANR NFMQLIHDDSLTFKEDIQKAQVSGQGHSLHEQIANLAGSPAIKKGILQTVKIVDELVK VMGHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQ NEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFIKDDSIDNKVLTRSDKNR GKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQ LVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREI NNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKAT AKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQ VNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAK VEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELE NGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHK HYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPA AFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO:2) (single underline: HNH domain; double underline: RuvC domain)

[00149] In some embodiments, wild type Cas9 corresponds to, or comprises SEQ ID NO:3 (nucleotide) and/or SEQ ID NO: 4 (amino acid):

ATGGATAAAAAGTATTCTATTGGTTTAGACATCGGCACTAATTCCGTTGGATGGG CTGTCATAACCGATGAATACAAAGTACCTTCAAAGAAATTTAAGGTGTTGGGGA ACACAGACCGTCATTCGATTAAAAAGAATCTTATCGGTGCCCTCCTATTCGATAG TGGCGAAACGGCAGAGGCGACTCGCCTGAAACGAACCGCTCGGAGAAGGTATAC ACGTCGCAAGAACCGAATATGTTACTTACAAGAAATTTTTAGCAATGAGATGGCC AAAGTTGACGATTCTTTCTTTCACCGTTTGGAAGAGTCCTTCCTTGTCGAAGAGG ACAAGAAACATGAACGGCACCCCATCTTTGGAAACATAGTAGATGAGGTGGCAT ATCATGAAAAGTACCCAACGATTTATCACCTCAGAAAAAAGCTAGTTGACTCAA CTGATAAAGCGGACCTGAGGTTAATCTACTTGGCTCTTGCCCATATGATAAAGTT CCGTGGGCACTTTCTCATTGAGGGTGATCTAAATCCGGACAACTCGGATGTCGAC AAACTGTTCATCCAGTTAGTACAAACCTATAATCAGTTGTTTGAAGAGAACCCTA TAAATGCAAGTGGCGTGGATGCGAAGGCTATTCTTAGCGCCCGCCTCTCTAAATC CCGACGGCTAGAAAACCTGATCGCACAATTACCCGGAGAGAAGAAAAATGGGTT GTTCGGTAACCTTATAGCGCTCTCACTAGGCCTGACACCAAATTTTAAGTCGAAC TTCGACTTAGCTGAAGATGCCAAATTGCAGCTTAGTAAGGACACGTACGATGAC GATCTCGACAATCTACTGGCACAAATTGGAGATCAGTATGCGGACTTATTTTTGG CTGCCAAAAACCTTAGCGATGCAATCCTCCTATCTGACATACTGAGAGTTAATAC TGAGATTACCAAGGCGCCGTTATCCGCTTCAATGATCAAAAGGTACGATGAACAT

CACCAAGACTTGACACTTCTCAAGGCCCTAGTCCGTCAGCAACTGCCTGAGAAAT
ATAAGGAAATATTCTTTGATCAGTCGAAAAACGGGTACGCAGGTTATATTGACG
GCGGAGCGAGTCAAGAGGAATTCTACAAGTTTATCAAACCCATATTAGAGAAGA
TGGATGGGACGGAAGAGTTGCTTGTAAAACTCAATCGCGAAGATCTACTGCGAA
AGCAGCGGACTTTCGACAACGGTAGCATTCCACATCAAATCCACTTAGGCGAATT
GCATGCTATACTTAGAAGGCAGGAGGATTTTTATCCGTTCCTCAAAGACAATCGT
GAAAAGATTGAGAAAATCCTAACCTTTCGCATACCTTACTATGTGGGACCCCTGG
CCCGAGGGAACTCTCGGTTCGCATGGATGACAAGAAAGTCCGAAGAAACGATTA
CTCCATGGAATTTTGAGGAAGTTGTCGATAAAGGTGCGTCAGCTCAATCGTTCAT
CGAGAGGATGACCAACTTTGACAAGAATTTACCGAACGAAAAAGTATTGCCTAA
GCACAGTTTACTTTACGAGTATTTCACAGTGTACAATGAACTCACGAAAGTTAAG
TATGTCACTGAGGGCATGCGTAAACCCGCCTTTCTAAGCGGAGAACAGAAGAAA
GCAATAGTAGATCTGTTATTCAAGACCAACCGCAAAGTGACAGTTAAGCAATTG
AAAGAGGACTACTTTAAGAAAATTGAATGCTTCGATTCTGTCGAGATCTCCGGGG
TAGAAGATCGATTTAATGCGTCACTTGGTACGTATCATGACCTCCTAAAGATAAT
TAAAGATAAGGACTTCCTGGATAACGAAGAGAATGAAGATATCTTAGAAGATAT
AGTGTTGACTCTTACCCTCTTTGAAGATCGGGAAATGATTGAGGAAAGACTAAAA
ACATACGCTCACCTGTTCGACGATAAGGTTATGAAACAGTTAAAGAGGCGTCGCT
ATACGGGCTGGGGACGATTGTCGCGGAAACTTATCAACGGGATAAGAGACAAGC
AAAGTGGTAAAACTATTCTCGATTTTCTAAAGAGCGACGGCTTCGCCAATAGGAA
CTTTATGCAGCTGATCCATGATGACTCTTTAACCTTCAAAGAGGATATACAAAAG
GCACAGGTTTCCGGACAAGGGGACTCATTGCACGAACATATTGCGAATCTTGCTG
GTTCGCCAGCCATCAAAAAGGGCATACTCCAGACAGTCAAAGTAGTGGATGAGC
TAGTTAAGGTCATGGGACGTCACAAACCGGAAAACATTGTAATCGAGATGGCAC
GCGAAAATCAAACGACTCAGAAGGGGCAAAAAAACAGTCGAGAGCGGATGAAG
AGAATAGAAGAGGGTATTAAAGAACTGGGCAGCCAGATCTTAAAGGAGCATCCT
GTGGAAAATACCCAATTGCAGAACGAGAAACTTTACCTCTATTACCTACAAAATG
GAAGGGACATGTATGTTGATCAGGAACTGGACATAAACCGTTTATCTGATTACGA
CGTCGATCACATTGTACCCCAATCCTTTTTGAAGGACGATTCAATCGACAATAAA
GTGCTTACACGCTCGGATAAGAACCGAGGGAAAAGTGACAATGTTCCAAGCGAG
GAAGTCGTAAAGAAAATGAAGAACTATTGGCGGCAGCTCCTAAATGCGAAACTG
ATAACGCAAAGAAAGTTCGATAACTTAACTAAAGCTGAGAGGGGTGGCTTGTCT
GAACTTGACAAGGCCGGATTTATTAAACGTCAGCTCGTGGAAACCCGCCAAATC
ACAAAGCATGTTGCACAGATACTAGATTCCCGAATGAATACGAAATACGACGAG
AACGATAAGCTGATTCGGGAAGTCAAAGTAATCACTTTAAAGTCAAAATTGGTG
TCGGACTTCAGAAAGGATTTTCAATTCTATAAAGTTAGGGAGATAAATAACTACC
ACCATGCGCACGACGCTTATCTTAATGCCGTCGTAGGGACCGCACTCATTAAGAA
ATACCCGAAGCTAGAAAGTGAGTTTGTGTATGGTGATTACAAAGTTTATGACGTC
CGTAAGATGATCGCGAAAAGCGAACAGGAGATAGGCAAGGCTACAGCCAAATA
CTTCTTTTATTCTAACATTATGAATTTCTTTAAGACGGAAATCACTCTGGCAAACG
GAGAGATACGCAAACGACCTTTAATTGAAACCAATGGGGAGACAGGTGAAATCG
TATGGGATAAGGGCCGGGACTTCGCGACGGTGAGAAAAGTTTTGTCCATGCCCC
AAGTCAACATAGTAAAGAAAACTGAGGTGCAGACCGGAGGGTTTTCAAAGGAAT
CGATTCTTCCAAAAAGGAATAGTGATAAGCTCATCGCTCGTAAAAAGGACTGGG
ACCCGAAAAAGTACGGTGGCTTCGATAGCCCTACAGTTGCCTATTCTGTCCTAGT
AGTGGCAAAAGTTGAGAAGGGAAAATCCAAGAAACTGAAGTCAGTCAAAGAAT
TATTGGGGATAACGATTATGGAGCGCTCGTCTTTTGAAAAGAACCCCATCGACTT
CCTTGAGGCGAAAGGTTACAAGGAAGTAAAAAAGGATCTCATAATTAAACTACC
AAAGTATAGTCTGTTTGAGTTAGAAAATGGCCGAAAACGGATGTTGGCTAGCGC
CGGAGAGCTTCAAAAGGGGAACGAACTCGCACTACCGTCTAAATACGTGAATTT

CCTGTATTTAGCGTCCCATTACGAGAAGTTGAAAGGTTCACCTGAAGATAACGAA
CAGAAGCAACTTTTTGTTGAGCAGCACAAACATTATCTCGACGAAATCATAGAGC
AAATTTCGGAATTCAGTAAGAGAGTCATCCTAGCTGATGCCAATCTGGACAAAGT
ATTAAGCGCATACAACAAGCACAGGGATAAACCCATACGTGAGCAGGCGGAAA
ATATTATCCATTTGTTTACTCTTACCAACCTCGGCGCTCCAGCCGCATTCAAGTAT
TTTGACACAACGATAGATCGCAAACGATACACTTCTACCAAGGAGGTGCTAGAC
GCGACACTGATTCACCAATCCATCACGGGATTATATGAAACTCGGATAGATTTGT
CACAGCTTGGGGGTGACGGATCCCCCAAGAAGAAGAGGAAAGTCTCGAGCGACT
ACAAAGACCATGACGGTGATTATAAAGATCATGACATCGATTACAAGGATGACG
ATGACAAGGCTGCAGGA (SEQ ID NO:3)

MDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGE
TAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHE
RHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEG
DLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLP
GEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYA
DLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPE
KYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQR
TFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFA
WMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTV
YNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFD
SVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO:4)
(single underline: HNH domain; double underline: RuvC domain)

[00150] In some embodiments, wild type Cas9 corresponds to Cas9 from *Streptococcus pyogenes* (NCBI Reference Sequence: NC_002737.2, SEQ ID NO: 8 (nucleotide); and Uniport Reference Sequence: Q99ZW2, SEQ ID NO: 10 (amino acid).

ATGGATAAGAAATACTCAATAGGCTTAGATATCGGCACAAATAGCGTCGGATGG
GCGGTGATCACTGATGAATATAAGGTTCCGTCTAAAAAGTTCAAGGTTCTGGGAA
ATACAGACCGCCACAGTATCAAAAAAAATCTTATAGGGGCTCTTTTATTTGACAG
TGGAGAGACAGCGGAAGCGACTCGTCTCAAACGGACAGCTCGTAGAAGGTATAC
ACGTCGGAAGAATCGTATTTGTTATCTACAGGAGATTTTTTCAAATGAGATGGCG
AAAGTAGATGATAGTTTCTTTCATCGACTTGAAGAGTCTTTTTTGGTGGAAGAAG
ACAAGAAGCATGAACGTCATCCTATTTTTGGAAATATAGTAGATGAAGTTGCTTA
TCATGAGAAATATCCAACTATCTATCATCTGCGAAAAAAATTGGTAGATTCTACT

GATAAAGCGGATTTGCGCTTAATCTATTTGGCCTTAGCGCATATGATTAAGTTTC
GTGGTCATTTTTTGATTGAGGGAGATTTAAATCCTGATAATAGTGATGTGGACAA
ACTATTTATCCAGTTGGTACAAACCTACAATCAATTATTTGAAGAAAACCCTATT
AACGCAAGTGGAGTAGATGCTAAAGCGATTCTTTCTGCACGATTGAGTAAATCA
AGACGATTAGAAAATCTCATTGCTCAGCTCCCCGGTGAGAAGAAAAATGGCTTA
TTTGGGAATCTCATTGCTTTGTCATTGGGTTTGACCCCTAATTTTAAATCAAATTT
TGATTTGGCAGAAGATGCTAAATTACAGCTTTCAAAAGATACTTACGATGATGAT
TTAGATAATTTATTGGCGCAAATTGGAGATCAATATGCTGATTTGTTTTTGGCAG
CTAAGAATTTATCAGATGCTATTTTACTTTCAGATATCCTAAGAGTAAATACTGA
AATAACTAAGGCTCCCCTATCAGCTTCAATGATTAAACGCTACGATGAACATCAT
CAAGACTTGACTCTTTTAAAAGCTTTAGTTCGACAACAACTTCCAGAAAAGTATA
AAGAAATCTTTTTTGATCAATCAAAAAACGGATATGCAGGTTATATTGATGGGGG
AGCTAGCCAAGAAGAATTTTATAAATTTATCAAACCAATTTTAGAAAAAATGGAT
GGTACTGAGGAATTATTGGTGAAACTAAATCGTGAAGATTTGCTGCGCAAGCAA
CGGACCTTTGACAACGGCTCTATTCCCCATCAAATTCACTTGGGTGAGCTGCATG
CTATTTTGAGAAGACAAGAAGACTTTTATCCATTTTTAAAAGACAATCGTGAGAA
GATTGAAAAAATCTTGACTTTTCGAATTCCTTATTATGTTGGTCCATTGGCGCGTG
GCAATAGTCGTTTTGCATGGATGACTCGGAAGTCTGAAGAAACAATTACCCCATG
GAATTTTGAAGAAGTTGTCGATAAAGGTGCTTCAGCTCAATCATTTATTGAACGC
ATGACAAACTTTGATAAAAATCTTCCAAATGAAAAAGTACTACCAAAACATAGT
TTGCTTTATGAGTATTTTACGGTTTATAACGAATTGACAAAGGTCAAATATGTTA
CTGAAGGAATGCGAAAACCAGCATTTCTTTCAGGTGAACAGAAGAAAGCCATTG
TTGATTTACTCTTCAAAACAAATCGAAAAGTAACCGTTAAGCAATTAAAAGAAG
ATTATTTCAAAAAAATAGAATGTTTTGATAGTGTTGAAATTTCAGGAGTTGAAGA
TAGATTTAATGCTTCATTAGGTACCTACCATGATTTGCTAAAAATTATTAAAGAT
AAAGATTTTTTGGATAATGAAGAAAATGAAGATATCTTAGAGGATATTGTTTTAA
CATTGACCTTATTTGAAGATAGGGAGATGATTGAGGAAAGACTTAAAACATATG
CTCACCTCTTTGATGATAAGGTGATGAAACAGCTTAAACGTCGCCGTTATACTGG
TTGGGGACGTTTGTCTCGAAAATTGATTAATGGTATTAGGGATAAGCAATCTGGC
AAAACAATATTAGATTTTTTGAAATCAGATGGTTTTGCCAATCGCAATTTTATGC
AGCTGATCCATGATGATAGTTTGACATTTAAAGAAGACATTCAAAAAGCACAAG
TGTCTGGACAAGGCGATAGTTTACATGAACATATTGCAAATTTAGCTGGTAGCCC
TGCTATTAAAAAAGGTATTTTACAGACTGTAAAAGTTGTTGATGAATTGGTCAAA
GTAATGGGGCGGCATAAGCCAGAAAATATCGTTATTGAAATGGCACGTGAAAAT
CAGACAACTCAAAAGGGCCAGAAAAATTCGCGAGAGCGTATGAAACGAATCGA
AGAAGGTATCAAAGAATTAGGAAGTCAGATTCTTAAAGAGCATCCTGTTGAAAA
TACTCAATTGCAAAATGAAAAGCTCTATCTCTATTATCTCCAAAATGGAAGAGAC
ATGTATGTGGACCAAGAATTAGATATTAATCGTTTAAGTGATTATGATGTCGATC
ACATTGTTCCACAAAGTTTCCTTAAAGACGATTCAATAGACAATAAGGTCTTAAC
GCGTTCTGATAAAAATCGTGGTAAATCGGATAACGTTCCAAGTGAAGAAGTAGT
CAAAAAGATGAAAAACTATTGGAGACAACTTCTAAACGCCAAGTTAATCACTCA
ACGTAAGTTTGATAATTTAACGAAAGCTGAACGTGGAGGTTTGAGTGAACTTGAT
AAAGCTGGTTTTATCAAACGCCAATTGGTTGAAACTCGCCAAATCACTAAGCATG
TGGCACAAATTTTGGATAGTCGCATGAATACTAAATACGATGAAAATGATAAAC

TTATTCGAGAGGTTAAAGTGATTACCTTAAAATCTAAATTAGTTTCTGACTTCCG
AAAAGATTTCCAATTCTATAAAGTACGTGAGATTAACAATTACCATCATGCCCAT
GATGCGTATCTAAATGCCGTCGTTGGAACTGCTTTGATTAAGAAATATCCAAAAC
TTGAATCGGAGTTTGTCTATGGTGATTATAAAGTTTATGATGTTCGTAAAATGATT
GCTAAGTCTGAGCAAGAAATAGGCAAAGCAACCGCAAAATATTTCTTTTACTCTA
ATATCATGAACTTCTTCAAAACAGAAATTACACTTGCAAATGGAGAGATTCGCAA
ACGCCCTCTAATCGAAACTAATGGGGAAACTGGAGAAATTGTCTGGGATAAAGG
GCGAGATTTTGCCACAGTGCGCAAAGTATTGTCCATGCCCCAAGTCAATATTGTC
AAGAAAACAGAAGTACAGACAGGCGGATTCTCCAAGGAGTCAATTTTACCAAAA
AGAAATTCGGACAAGCTTATTGCTCGTAAAAAAGACTGGGATCCAAAAAAATAT
GGTGGTTTTGATAGTCCAACGGTAGCTTATTCAGTCCTAGTGGTTGCTAAGGTGG
AAAAAGGGAAATCGAAGAAGTTAAAATCCGTTAAAGAGTTACTAGGGATCACAA
TTATGGAAAGAAGTTCCTTTGAAAAAAATCCGATTGACTTTTTAGAAGCTAAAGG
ATATAAGGAAGTTAAAAAAGACTTAATCATTAAACTACCTAAATATAGTCTTTTT
GAGTTAGAAAACGGTCGTAAACGGATGCTGGCTAGTGCCGGAGAATTACAAAAA
GGAAATGAGCTGGCTCTGCCAAGCAAATATGTGAATTTTTTATATTTAGCTAGTC
ATTATGAAAAGTTGAAGGGTAGTCCAGAAGATAACGAACAAAAACAATTGTTTG
TGGAGCAGCATAAGCATTATTTAGATGAGATTATTGAGCAAATCAGTGAATTTTC
TAAGCGTGTTATTTTAGCAGATGCCAATTTAGATAAAGTTCTTAGTGCATATAAC
AAACATAGAGACAAACCAATACGTGAACAAGCAGAAAATATTATTCATTTATTT
ACGTTGACGAATCTTGGAGCTCCCGCTGCTTTTAAATATTTTGATACAACAATTG
ATCGTAAACGATATACGTCTACAAAAGAAGTTTTAGATGCCACTCTTATCCATCA
ATCCATCACTGGTCTTTATGAAACACGCATTGATTTGAGTCAGCTAGGAGGTGAC
TGA (SEQ ID NO: 8)

MDKKYSIGLDIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGE
TAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHE
RHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEG
DLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLP
GEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYA
DLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPE
KYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQR
TFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFA
WMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTV
YNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFD
SVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK

ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO:
10) (single underline: HNH domain; double underline: RuvC domain)

[00151] In some embodiments, Cas9 refers to Cas9 from: *Corynebacterium ulcerans* (NCBI Refs: NC_015683.1, NC_017317.1); *Corynebacterium diphtheria* (NCBI Refs: NC_016782.1, NC_016786.1); *Spiroplasma syrphidicola* (NCBI Ref: NC_021284.1); *Prevotella intermedia* (NCBI Ref: NC_017861.1); *Spiroplasma taiwanense* (NCBI Ref: NC_021846.1); *Streptococcus iniae* (NCBI Ref: NC_021314.1); *Belliella baltica* (NCBI Ref: NC_018010.1); *Psychroflexus torquisI* (NCBI Ref: NC_018721.1); *Streptococcus thermophilus* (NCBI Ref: YP_820832.1), *Listeria innocua* (NCBI Ref: NP_472073.1), *Campylobacter jejuni* (NCBI Ref: YP_002344900.1) or *Neisseria. meningitidis* (NCBI Ref: YP_002342100.1) or to a Cas9 from any of the organisms listed in Example 5.

[00152] In some embodiments, dCas9 corresponds to, or comprises in part or in whole, a Cas9 amino acid sequence having one or more mutations that inactivate the Cas9 nuclease activity. For example, in some embodiments, a dCas9 domain comprises D10A and/or H840A mutation.

dCas9 (D10A and H840A):

MDKK**YSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDS**
GETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKK
HERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLI
EGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQ
LPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQ
YADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQL
PEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRK
QRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSR
FAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYF
TVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIEC
FDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEE
RLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFAN
RNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDEL
VKVMGRHKPENIVIEMARENQTTQK**GQKNSRERMKRIEEGIKELGSQILKEHPVE**
NTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKV
LTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLS
ELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFR

KDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMI
AKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDF
ATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSP
TVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLI
IKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDN
EQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIH
LFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD
(SEQ ID NO: 9) (single underline: HNH domain; double underline: RuvC domain).

[00153] In some embodiments, the Cas9 domain comprises a D10A mutation, while the residue at position 840 remains a histidine in the amino acid sequence provided in SEQ ID NO: 10, or at corresponding positions in any of the amino acid sequences provided in SEQ ID NOs: 11-260. Without wishing to be bound by any particular theory, the presence of the catalytic residue H840 restores the acvitity of the Cas9 to cleave the non-edited (*e.g.,* non-deaminated) strand containing a G opposite the targeted C. Restoration of H840 (*e.g.,* from A840) does not result in the cleavage of the target strand containing the C. Such Cas9 variants are able to generate a single-strand DNA break (nick) at a specific location based on the gRNA-defined target sequence, leading to repair of the non-edited strand, ultimately resulting in a G to A change on the non-edited strand. A schematic representation of this process is shown in Figure 108. Briefly, the C of a C-G basepair can be deaminated to a U by a deaminase, *e.g.,* an APOBEC deamonase. Nicking the non-edited strand, having the G, facilitates removal of the G via mismatch repair mechanisms. UGI inhibits UDG, which prevents removal of the U.

[00154] In other embodiments, dCas9 variants having mutations other than D10A and H840A are provided, which, *e.g.*, result in nuclease inactivated Cas9 (dCas9). Such mutations, by way of example, include other amino acid substitutions at D10 and H820, or other substitutions within the nuclease domains of Cas9 (*e.g.*, substitutions in the HNH nuclease subdomain and/or the RuvC1 subdomain). In some embodiments, variants or homologues of dCas9 (*e.g.*, variants of SEQ ID NO: 10) are provided which are at least about 70% identical, at least about 80% identical, at least about 90% identical, at least about 95% identical, at least about 98% identical, at least about 99% identical, at least about 99.5% identical, or at least about 99.9% identical to SEQ ID NO: 10. In some embodiments, variants of dCas9 (*e.g.*, variants of SEQ ID NO: 10) are provided having amino acid sequences which are shorter, or longer than SEQ ID NO: 10, by about 5 amino acids, by about 10 amino acids, by about 15 amino acids, by about 20 amino acids, by about 25 amino

acids, by about 30 amino acids, by about 40 amino acids, by about 50 amino acids, by about 75 amino acids, by about 100 amino acids or more.

[00155] In some embodiments, Cas9 fusion proteins as provided herein comprise the full-length amino acid sequence of a Cas9 protein, *e.g.*, one of the Cas9 sequences provided herein. In other embodiments, however, fusion proteins as provided herein do not comprise a full-length Cas9 sequence, but only a fragment thereof. For example, in some embodiments, a Cas9 fusion protein provided herein comprises a Cas9 fragment, wherein the fragment binds crRNA and tracrRNA or sgRNA, but does not comprise a functional nuclease domain, *e.g.*, in that it comprises only a truncated version of a nuclease domain or no nuclease domain at all. Exemplary amino acid sequences of suitable Cas9 domains and Cas9 fragments are provided herein, and additional suitable sequences of Cas9 domains and fragments will be apparent to those of skill in the art.

[00156] In some embodiments, Cas9 refers to Cas9 from: *Corynebacterium ulcerans* (NCBI Refs: NC_015683.1, NC_017317.1); *Corynebacterium diphtheria* (NCBI Refs: NC_016782.1, NC_016786.1); *Spiroplasma syrphidicola* (NCBI Ref: NC_021284.1); *Prevotella intermedia* (NCBI Ref: NC_017861.1); *Spiroplasma taiwanense* (NCBI Ref: NC_021846.1); *Streptococcus iniae* (NCBI Ref: NC_021314.1); *Belliella baltica* (NCBI Ref: NC_018010.1); *Psychroflexus torquisI* (NCBI Ref: NC_018721.1); *Streptococcus thermophilus* (NCBI Ref: YP_820832.1); *Listeria innocua* (NCBI Ref: NP_472073.1); *Campylobacter jejuni* (NCBI Ref: YP_002344900.1); or *Neisseria. meningitidis* (NCBI Ref: YP_002342100.1).

[00157] The term "deaminase" or "deaminase domain," as used herein, refers to a protein or enzyme that catalyzes a deamination reaction. In some embodiments, the deaminase or deaminase domain is a cytidine deaminase, catalyzing the hydrolytic deamination of cytidine or deoxycytidine to uridine or deoxyuridine, respectively. In some embodiments, the deaminase or deaminase domain is a cytidine deaminase domain, catalyzing the hydrolytic deamination of cytosine to uracil. In some embodiments, the deaminase or deaminase domain is a naturally-occuring deaminase from an organism, such as a human, chimpanzee, gorilla, monkey, cow, dog, rat, or mouse. In some embodiments, the deaminase or deaminase domain is a variant of a naturally-occuring deaminase from an organism, that does not occur in nature. For example, in some embodiments, the deaminase or deaminase domain is at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75% at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to a naturally-occuring deaminase from an organism.

[00158] The term "effective amount," as used herein, refers to an amount of a biologically active agent that is sufficient to elicit a desired biological response. For example, in some embodiments, an effective amount of a nuclease may refer to the amount of the nuclease that is sufficient to induce cleavage of a target site specifically bound and cleaved by the nuclease. In some embodiments, an effective amount of a fusion protein provided herein, *e.g.*, of a fusion protein comprising a nuclease-inactive Cas9 domain and a nucleic acid editing domain (*e.g.*, a deaminase domain) may refer to the amount of the fusion protein that is sufficient to induce editing of a target site specifically bound and edited by the fusion protein. As will be appreciated by the skilled artisan, the effective amount of an agent, *e.g.*, a fusion protein, a nuclease, a deaminase, a recombinase, a hybrid protein, a protein dimer, a complex of a protein (or protein dimer) and a polynucleotide, or a polynucleotide, may vary depending on various factors as, for example, on the desired biological response, *e.g.*, on the specific allele, genome, or target site to be edited, on the cell or tissue being targeted, and on the agent being used.

[00159] The term "linker," as used herein, refers to a chemical group or a molecule linking two molecules or moieties, *e.g.*, two domains of a fusion protein, such as, for example, a nuclease-inactive Cas9 domain and a nucleic acid editing domain (*e.g.*, a deaminase domain). In some embodiments, a linker joins a gRNA binding domain of an RNA-programmable nuclease, including a Cas9 nuclease domain, and the catalytic domain of anucleic-acid editing protein. In some embodiments, a linker joins a dCas9 and a nucleic-acid editing protein. Typically, the linker is positioned between, or flanked by, two groups, molecules, or other moieties and connected to each one via a covalent bond, thus connecting the two. In some embodiments, the linker is an amino acid or a plurality of amino acids (*e.g.*, a peptide or protein). In some embodiments, the linker is an organic molecule, group, polymer, or chemical moiety. In some embodiments, the linker is 5-100 amino acids in length, for example, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 30-35, 35-40, 40-45, 45-50, 50-60, 60-70, 70-80, 80-90, 90-100, 100-150, or 150-200 amino acids in length. Longer or shorter linkers are also contemplated.

[00160] The term "mutation," as used herein, refers to a substitution of a residue within a sequence, *e.g.*, a nucleic acid or amino acid sequence, with another residue, or a deletion or insertion of one or more residues within a sequence. Mutations are typically described herein by identifying the original residue followed by the position of the residue within the sequence and by the identity of the newly substituted residue. Various methods for making the amino acid substitutions (mutations) provided herein are well known in the art, and are provided by,

for example, Green and Sambrook, *Molecular Cloning: A Laboratory Manual* (4th ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (2012)).

[00161] The terms "nucleic acid" and "nucleic acid molecule," as used herein, refer to a compound comprising a nucleobase and an acidic moiety, *e.g.*, a nucleoside, a nucleotide, or a polymer of nucleotides. Typically, polymeric nucleic acids, *e.g.*, nucleic acid molecules comprising three or more nucleotides are linear molecules, in which adjacent nucleotides are linked to each other via a phosphodiester linkage. In some embodiments, "nucleic acid" refers to individual nucleic acid residues (*e.g.* nucleotides and/or nucleosides). In some embodiments, "nucleic acid" refers to an oligonucleotide chain comprising three or more individual nucleotide residues. As used herein, the terms "oligonucleotide" and "polynucleotide" can be used interchangeably to refer to a polymer of nucleotides (*e.g.*, a string of at least three nucleotides). In some embodiments, "nucleic acid" encompasses RNA as well as single and/or double-stranded DNA. Nucleic acids may be naturally occurring, for example, in the context of a genome, a transcript, an mRNA, tRNA, rRNA, siRNA, snRNA, a plasmid, cosmid, chromosome, chromatid, or other naturally occurring nucleic acid molecule. On the other hand, a nucleic acid molecule may be a non-naturally occurring molecule, *e.g.*, a recombinant DNA or RNA, an artificial chromosome, an engineered genome, or fragment thereof, or a synthetic DNA, RNA, DNA/RNA hybrid, or including non-naturally occurring nucleotides or nucleosides. Furthermore, the terms "nucleic acid," "DNA," "RNA," and/or similar terms include nucleic acid analogs, *e.g.,* analogs having other than a phosphodiester backbone. Nucleic acids can be purified from natural sources, produced using recombinant expression systems and optionally purified, chemically synthesized, *etc.* Where appropriate, *e.g.*, in the case of chemically synthesized molecules, nucleic acids can comprise nucleoside analogs such as analogs having chemically modified bases or sugars, and backbone modifications. A nucleic acid sequence is presented in the 5′ to 3′ direction unless otherwise indicated. In some embodiments, a nucleic acid is or comprises natural nucleosides (*e.g.* adenosine, thymidine, guanosine, cytidine, uridine, deoxyadenosine, deoxythymidine, deoxyguanosine, and deoxycytidine); nucleoside analogs (*e.g.*, 2-aminoadenosine, 2-thiothymidine, inosine, pyrrolo-pyrimidine, 3-methyl adenosine, 5-methylcytidine, 2-aminoadenosine, C5-bromouridine, C5-fluorouridine, C5-iodouridine, C5-propynyl-uridine, C5-propynyl-cytidine, C5-methylcytidine, 2-aminoadenosine, 7-deazaadenosine, 7-deazaguanosine, 8-oxoadenosine, 8-oxoguanosine, O(6)-methylguanine, and 2-thiocytidine); chemically modified bases; biologically modified bases (*e.g.*, methylated bases); intercalated bases; modified sugars (*e.g.*, 2′-fluororibose, ribose, 2′-deoxyribose,

arabinose, and hexose); and/or modified phosphate groups (*e.g.*, phosphorothioates and 5′-*N*-phosphoramidite linkages).

[00162] The term "nucleic acid editing domain," as used herein refers to a protein or enzyme capable of making one or more modifications (*e.g.,* deamination of a cytidine residue) to a nucleic acid (*e.g.,* DNA or RNA). Exemplary nucleic acid editing domains include, but are not limited to a deaminase, a nuclease, a nickase, a recombinase, a methyltransferase, a methylase, an acetylase, an acetyltransferase, a transcriptional activator, or a transcriptional repressor domain. In some embodiments the nucleic acid editing domain is a deaminase (*e.g.,* a cytidine deaminase, such as an APOBEC or an AID deaminase).

[00163] The term "proliferative disease," as used herein, refers to any disease in which cell or tissue homeostasis is disturbed in that a cell or cell population exhibits an abnormally elevated proliferation rate. Proliferative diseases include hyperproliferative diseases, such as pre-neoplastic hyperplastic conditions and neoplastic diseases. Neoplastic diseases are characterized by an abnormal proliferation of cells and include both benign and malignant neoplasias. Malignant neoplasia is also referred to as cancer.

[00164] The terms "protein," "peptide," and "polypeptide" are used interchangeably herein, and refer to a polymer of amino acid residues linked together by peptide (amide) bonds. The terms refer to a protein, peptide, or polypeptide of any size, structure, or function. Typically, a protein, peptide, or polypeptide will be at least three amino acids long. A protein, peptide, or polypeptide may refer to an individual protein or a collection of proteins. One or more of the amino acids in a protein, peptide, or polypeptide may be modified, for example, by the addition of a chemical entity such as a carbohydrate group, a hydroxyl group, a phosphate group, a farnesyl group, an isofarnesyl group, a fatty acid group, a linker for conjugation, functionalization, or other modification, *etc.* A protein, peptide, or polypeptide may also be a single molecule or may be a multi-molecular complex. A protein, peptide, or polypeptide may be just a fragment of a naturally occurring protein or peptide. A protein, peptide, or polypeptide may be naturally occurring, recombinant, or synthetic, or any combination thereof. The term "fusion protein" as used herein refers to a hybrid polypeptide which comprises protein domains from at least two different proteins. One protein may be located at the amino-terminal (N-terminal) portion of the fusion protein or at the carboxy-terminal (C-terminal) protein thus forming an "amino-terminal fusion protein" or a "carboxy-terminal fusion protein," respectively. A protein may comprise different domains, for example, a nucleic acid binding domain (*e.g.*, the gRNA binding domain of Cas9 that directs the binding of the protein to a target site) and a nucleic acid cleavage domain or a catalytic domain of a

nucleic-acid editing protein. In some embodiments, a protein comprises a proteinaceous part, *e.g.*, an amino acid sequence constituting a nucleic acid binding domain, and an organic compound, *e.g.*, a compound that can act as a nucleic acid cleavage agent. In some embodiments, a protein is in a complex with, or is in association with, a nucleic acid, *e.g.*, RNA. Any of the proteins provided herein may be produced by any method known in the art. For example, the proteins provided herein may be produced via recombinant protein expression and purification, which is especially suited for fusion proteins comprising a peptide linker. Methods for recombinant protein expression and purification are well known, and include those described by Green and Sambrook, *Molecular Cloning: A Laboratory Manual* (4th ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (2012)), the entire contents of which are incorporated herein by reference.

[00165] The term "RNA-programmable nuclease," and "RNA-guided nuclease" are used interchangeably herein and refer to a nuclease that forms a complex with (*e.g.*, binds or associates with) one or more RNA that is not a target for cleavage. In some embodiments, an RNA-programmable nuclease, when in a complex with an RNA, may be referred to as a nuclease:RNA complex. Typically, the bound RNA(s) is referred to as a guide RNA (gRNA). gRNAs can exist as a complex of two or more RNAs, or as a single RNA molecule. gRNAs that exist as a single RNA molecule may be referred to as single-guide RNAs (sgRNAs), though "gRNA" is used interchangeably to refer to guide RNAs that exist as either single molecules or as a complex of two or more molecules. Typically, gRNAs that exist as single RNA species comprise two domains: (1) a domain that shares homology to a target nucleic acid (*e.g.*, and directs binding of a Cas9 complex to the target); and (2) a domain that binds a Cas9 protein. In some embodiments, domain (2) corresponds to a sequence known as a tracrRNA, and comprises a stem-loop structure. For example, in some embodiments, domain (2) is identical or homologous to a tracrRNA as provided in Jinek *et al.*, *Science* 337:816-821(2012), the entire contents of which is incorporated herein by reference. Other examples of gRNAs (*e.g.*, those including domain 2) can be found in U.S. Provisional Patent Application, U.S.S.N. 61/874,682, filed September 6, 2013, entitled "Switchable Cas9 Nucleases And Uses Thereof," and U.S. Provisional Patent Application, U.S.S.N. 61/874,746, filed September 6, 2013, entitled "Delivery System For Functional Nucleases," the entire contents of each are hereby incorporated by reference in their entirety. In some embodiments, a gRNA comprises two or more of domains (1) and (2), and may be referred to as an "extended gRNA." For example, an extended gRNA will, *e.g.*, bind two or more Cas9 proteins and bind a target nucleic acid at two or more distinct regions, as

described herein. The gRNA comprises a nucleotide sequence that complements a target site, which mediates binding of the nuclease/RNA complex to said target site, providing the sequence specificity of the nuclease:RNA complex. In some embodiments, the RNA-programmable nuclease is the (CRISPR-associated system) Cas9 endonuclease, for example Cas9 (Csn1) from *Streptococcus pyogenes* (see, *e.g.*, "Complete genome sequence of an M1 strain of *Streptococcus pyogenes*." Ferretti J.J., McShan W.M., Ajdic D.J., Savic D.J., Savic G., Lyon K., Primeaux C., Sezate S., Suvorov A.N., Kenton S., Lai H.S., Lin S.P., Qian Y., Jia H.G., Najar F.Z., Ren Q., Zhu H., Song L., White J., Yuan X., Clifton S.W., Roe B.A., McLaughlin R.E., Proc. Natl. Acad. Sci. U.S.A. 98:4658-4663(2001); "CRISPR RNA maturation by trans-encoded small RNA and host factor RNase III." Deltcheva E., Chylinski K., Sharma C.M., Gonzales K., Chao Y., Pirzada Z.A., Eckert M.R., Vogel J., Charpentier E., Nature 471:602-607(2011); and "A programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity." Jinek M., Chylinski K., Fonfara I., Hauer M., Doudna J.A., Charpentier E. Science 337:816-821(2012), the entire contents of each of which are incorporated herein by reference.

[00166] Because RNA-programmable nucleases (*e.g.*, Cas9) use RNA:DNA hybridization to target DNA cleavage sites, these proteins are able to be targeted, in principle, to any sequence specified by the guide RNA. Methods of using RNA-programmable nucleases, such as Cas9, for site-specific cleavage (*e.g.*, to modify a genome) are known in the art (see *e.g.*, Cong, L. *et al.* Multiplex genome engineering using CRISPR/Cas systems. *Science* **339**, 819-823 (2013); Mali, P. *et al.* RNA-guided human genome engineering via Cas9. *Science* **339**, 823-826 (2013); Hwang, W.Y. *et al.* Efficient genome editing in zebrafish using a CRISPR-Cas system. *Nature biotechnology* **31**, 227-229 (2013); Jinek, M. *et al.* RNA-programmed genome editing in human cells. *eLife* **2**, e00471 (2013); Dicarlo, J.E. *et al.* Genome engineering in *Saccharomyces cerevisiae* using CRISPR-Cas systems. *Nucleic acids research* (2013); Jiang, W. *et al.* RNA-guided editing of bacterial genomes using CRISPR-Cas systems. *Nature biotechnology* **31**, 233-239 (2013); the entire contents of each of which are incorporated herein by reference).

[00167] The term "subject," as used herein, refers to an individual organism, for example, an individual mammal. In some embodiments, the subject is a human. In some embodiments, the subject is a non-human mammal. In some embodiments, the subject is a non-human primate. In some embodiments, the subject is a rodent. In some embodiments, the subject is a sheep, a goat, a cattle, a cat, or a dog. In some embodiments, the subject is a vertebrate, an amphibian, a reptile, a fish, an insect, a fly, or a nematode. In some

embodiments, the subject is a research animal. In some embodiments, the subject is genetically engineered, *e.g.*, a genetically engineered non-human subject. The subject may be of either sex and at any stage of development.

[00168] The term "target site" refers to a sequence within a nucleic acid molecule that is deaminated by a deaminase or a fusion protein comprising a deaminase, (*e.g.*, a dCas9-deaminase fusion protein provided herein).

[00169] The terms "treatment," "treat," and "treating," refer to a clinical intervention aimed to reverse, alleviate, delay the onset of, or inhibit the progress of a disease or disorder, or one or more symptoms thereof, as described herein. As used herein, the terms "treatment," "treat," and "treating" refer to a clinical intervention aimed to reverse, alleviate, delay the onset of, or inhibit the progress of a disease or disorder, or one or more symptoms thereof, as described herein. In some embodiments, treatment may be administered after one or more symptoms have developed and/or after a disease has been diagnosed. In other embodiments, treatment may be administered in the absence of symptoms, *e.g.*, to prevent or delay onset of a symptom or inhibit onset or progression of a disease. For example, treatment may be administered to a susceptible individual prior to the onset of symptoms (*e.g.*, in light of a history of symptoms and/or in light of genetic or other susceptibility factors). Treatment may also be continued after symptoms have resolved, for example, to prevent or delay their recurrence.

[00170] The term "recombinant" as used herein in the context of proteins or nucleic acids refers to proteins or nucleic acids that do not occur in nature, but are the product of human engineering. For example, in some embodiments, a recombinant protein or nucleic acid molecule comprises an amino acid or nucleotide sequence that comprises at least one, at least two, at least three, at least four, at least five, at least six, or at least seven mutations as compared to any naturally occurring sequence.

[00171] The term "nucleobase editors (NBEs)" or "base editors (BEs)," as used herein, refers to the Cas9 fusion proteins described herein. In some embodiments, the fusion protein comprises a nuclease-inactive Cas9 (dCas9) fused to a deaminase. In some embodiments, the fusion protein comprises a Cas9 nickase fused to a deaminase. In some embodiments, the fusion protein comprises a nuclease-inactive Cas9 fused to a deaminase and further fused to a UGI domain. In some embodiments, the fusion protein comprises a Cas9 nickase fused to a deaminase and further fused to a UGI domain. In some embodiments, the dCas9 of the fusion protein comprises a D10A and a H840A mutation of SEQ ID NO: 10, or a corresponding mutation in any of SEQ ID NOs: 11-260, which inactivates nuclease activity

of the Cas9 protein. In some embodiments, the fusion protein comprises a D10A mutation and comprises a histidine at residue 840 of SEQ ID NO: 10, or a corresponding mutation in any of SEQ ID NOs: 11-260, which renders Cas9 capable of cleaving only one strand of a nucleic acid duplex. An example of a Cas9 nickase is shown below in SEQ ID NO: 674. The terms "nucleobase editors (NBEs)" and "base editors (BEs)" may be used interchangeably.

[00172] The term "uracil glycosylase inhibitor" or "UGI," as used herein, refers to a protein that is capable of inhibiting a uracil-DNA glycosylase base-excision repair enzyme.

[00173] The term "Cas9 nickase," as used herein, refers to a Cas9 protein that is capable of cleaving only one strand of a duplexed nucleic acid molecule (*e.g.,* a duplexed DNA molecule). In some embodiments, a Cas9 nickase comprises a D10A mutation and has a histidine at position H840 of SEQ ID NO: 10, or a corresponding mutation in any of SEQ ID NOs: 11-260. For example, a Cas9 nickase may comprise the amino acid sequence as set forth in SEQ ID NO: 674. Such a Cas9 nickase has an active HNH nuclease domain and is able to cleave the non-targeted strand of DNA, *i.e.*, the strand bound by the gRNA. Further, such a Cas9 nickase has an inactive RuvC nuclease domain and is not able to cleave the targeted strand of the DNA, *i.e.*, the strand where base editing is desired.

[00174] Exemplary Cas9 nickase (Cloning vector pPlatTET-gRNA2; Accession No. BAV54124).

MDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGE
TAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHE
RHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEG
DLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLP
GEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYA
DLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPE
KYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQR
TFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFA
WMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTV
YNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFD
SVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK

NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVV AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 674)

DETAILED DESCRIPTION OF CERTAIN EMBODIMENTS OF THE INVENTION

[00175] Some aspects of this disclosure provide fusion proteins that comprise a domain capable of binding to a nucleotide sequence (*e.g.*, a Cas9, or a Cpf1 protein) and an enzyme domain, for example, a DNA-editing domain, such as, *e.g.*, a deaminase domain. The deamination of a nucleobase by a deaminase can lead to a point mutation at the respective residue, which is referred to herein as nucleic acid editing. Fusion proteins comprising a Cas9 variant or domain and a DNA editing domain can thus be used for the targeted editing of nucleic acid sequences. Such fusion proteins are useful for targeted editing of DNA *in vitro*, *e.g.*, for the generation of mutant cells or animals; for the introduction of targeted mutations, *e.g.*, for the correction of genetic defects in cells *ex vivo*, *e.g.*, in cells obtained from a subject that are subsequently re-introduced into the same or another subject; and for the introduction of targeted mutations, *e.g.*, the correction of genetic defects or the introduction of deactivating mutations in disease-associated genes in a subject. Typically, the Cas9 domain of the fusion proteins described herein does not have any nuclease activity but instead is a Cas9 fragment or a dCas9 protein or domain. Methods for the use of Cas9 fusion proteins as described herein are also provided.

Cas9 domains of Nucleobase Editors

[00176] Non-limiting, exemplary Cas9 domains are provided herein. The Cas9 domain may be a nuclease active Cas9 domain, a nucleasae inactive Cas9 domain, or a Cas9 nickase. In some embodiments, the Cas9 domain is a nuclease active domain. For example, the Cas9 domain may be a Cas9 domain that cuts both strands of a duplexed nucleic acid (*e.g.*, both strands of a duplexed DNA molecule). In some embodiments, the Cas9 domain comprises

any one of the amino acid sequences as set forth in SEQ ID NOs: 10-263. In some embodiments the Cas9 domain comprises an amino acid sequence that is at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to any one of the amino acid sequences set forth in SEQ ID NOs: 10-263. In some embodiments, the Cas9 domain comprises an amino acid sequence that has 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 21, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50 or more or more mutations compared to any one of the amino acid sequences set forth in SEQ ID NOs: 10-263. In some embodiments, the Cas9 domain comprises an amino acid sequence that has at least 10, at least 15, at least 20, at least 30, at leat 40, at least 50, at least 60, at least 70, at least 80, at least 90, at least 100, at least 150, at least 200, at least 250, at least 300, at least 350, at least 400, at least 500, at least 600, at least 700, at least 800, at least 900, at least 1000, at least 1100, or at least 1200 identical contiguous amino acid residues as compared to any one of the amino acid sequences set forth in SEQ ID NOs: 10-263.

[00177] In some embodiments, the Cas9 domain is a nuclease-inactive Cas9 domain (dCas9). For example, the dCas9 domain may bind to a duplexed nucleic acid molecule (*e.g.*, via a gRNA molecule) without cleaving either strand of the duplexed nucleic acid molecule. In some embodiments, the nuclease-inactive dCas9 domain comprises a D10X mutation and a H840X mutation of the amino acid sequence set forth in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid change. In some embodiments, the nuclease-inactive dCas9 domain comprises a D10A mutation and a H840A mutation of the amino acid sequence set forth in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. As one example, a nuclease-inactive Cas9 domain comprises the amino acid sequence set forth in SEQ ID NO: 263 (Cloning vector pPlatTET-gRNA2, Accession No. BAV54124).

MDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGE
TAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHE
RHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEG
DLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLP
GEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYA
DLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPE

KYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQR
TFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFA
WMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTV
YNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFD
SVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO:
263; see, *e.g.*, Qi *et al.*, Repurposing CRISPR as an RNA-guided platform for sequence-specific control of gene expression. *Cell.* 2013; 152(5):1173-83, the entire contents of which are incorporated herein by reference).

[00178] Additional suitable nuclease-inactive dCas9 domains will be apparent to those of skill in the art based on this disclosure and knowledge in the field, and are within the scope of this disclosure. Such additional exemplary suitable nuclease-inactive Cas9 domains include, but are not limited to, D10A/H840A, D10A/D839A/H840A, and D10A/D839A/H840A/N863A mutant domains (See, *e.g.*, Prashant *et al.*, CAS9 transcriptional activators for target specificity screening and paired nickases for cooperative genome engineering. *Nature Biotechnology.* 2013; 31(9): 833-838, the entire contents of which are incorporated herein by reference). In some embodiments the dCas9 domain comprises an amino acid sequence that is at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to any one of the dCas9 domains provided herein. In some embodiments, the Cas9 domain comprises an amino acid sequences that has 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 21, 24, 25, 26, 27, 28,

29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50 or more or more mutations compared to any one of the amino acid sequences set forth in SEQ ID NOs: 10-263. In some embodiments, the Cas9 domain comprises an amino acid sequence that has at least 10, at least 15, at least 20, at least 30, at leat 40, at least 50, at least 60, at least 70, at least 80, at least 90, at least 100, at least 150, at least 200, at least 250, at least 300, at least 350, at least 400, at least 500, at least 600, at least 700, at least 800, at least 900, at least 1000, at least 1100, or at least 1200 identical contiguous amino acid residues as compared to any one of the amino acid sequences set forth in SEQ ID NOs: 10-263.

[00179] In some embodiments, the Cas9 domain is a Cas9 nickase. The Cas9 nickase may be a Cas9 protein that is capable of cleaving only one strand of a duplexed nucleic acid molecule (*e.g.,* a duplexed DNA molecule). In some embodiments the Cas9 nickase cleaves the target strand of a duplexed nucleic acid molecule, meaning that the Cas9 nickase cleaves the strand that is base paired to (complementary to) a gRNA (*e.g.,* an sgRNA) that is bound to the Cas9. In some embodiments, a Cas9 nickase comprises a D10A mutation and has a histidine at position 840 of SEQ ID NO: 10, or a mutation in any of SEQ ID NOs: 11-260. For example, a Cas9 nickase may comprise the amino acid sequence as set forth in SEQ ID NO: 674. In some embodiments the Cas9 nickase cleaves the non-target, non-base-edited strand of a duplexed nucleic acid molecule, meaning that the Cas9 nickase cleaves the strand that is not base paired to a gRNA (*e.g.,* an sgRNA) that is bound to the Cas9. In some embodiments, a Cas9 nickase comprises an H840A mutation and has an aspartic acid residue at position 10 of SEQ ID NO: 10, or a corresponding mutation in any of SEQ ID NOs: 11-260. In some embodiments the Cas9 nickase comprises an amino acid sequence that is at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to any one of the Cas9 nickases provided herein. Additional suitable Cas9 nickases will be apparent to those of skill in the art based on this disclosure and knowledge in the field, and are within the scope of this disclosure.

Cas9 Domains with Reduced PAM Exclusivity

[00180] Some aspects of the disclosure provide Cas9 domains that have different PAM specificities. Typically, Cas9 proteins, such as Cas9 from *S. pyogenes* (spCas9), require a canonical NGG PAM sequence to bind a particular nucleic acid region. This may limit the ability to edit desired bases within a genome. In some embodiments, the base editing fusion

proteins provided herein may need to be placed at a precise location, for example where a target base is placed within a 4 base region (*e.g.*, a "deamination window"), which is approximately 15 bases upstream of the PAM. See Komor, A.C., *et al.*, "Programmable editing of a target base in genomic DNA without double-stranded DNA cleavage" *Nature* 533, 420-424 (2016), the entire contents of which are hereby incorporated by reference. Accordingly, in some embodiments, any of the fusion proteins provided herein may contain a Cas9 domain that is capable of binding a nucleotide sequence that does not contain a canonical (*e.g.*, NGG) PAM sequence. Cas9 domains that bind to non-canonical PAM sequences have been described in the art and would be apparent to the skilled artisan. For example, Cas9 domains that bind non-canonical PAM sequences have been described in Kleinstiver, B. P., *et al.*, "Engineered CRISPR-Cas9 nucleases with altered PAM specificities" *Nature* 523, 481-485 (2015); and Kleinstiver, B. P., *et al.*, "Broadening the targeting range of *Staphylococcus aureus* CRISPR-Cas9 by modifying PAM recognition" *Nature Biotechnology* 33, 1293-1298 (2015); the entire contents of each are hereby incorporated by reference.

[00181] In some embodiments, the Cas9 domain is a Cas9 domain from *Staphylococcus aureus* (SaCas9). In some embodiments, the SaCas9 domain is a nuclease active SaCas9, a nuclease inactive SaCas9 (SaCas9d), or a SaCas9 nickase (SaCas9n). In some embodiments, the SaCas9 comprises the amino acid sequence SEQ ID NO: 4273. In some embodiments, the SaCas9 comprises a N579X mutation of SEQ ID NO: 4273, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for N. In some embodiments, the SaCas9 comprises a N579A mutation of SEQ ID NO: 4273, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SaCas9 domain, the SaCas9d domain, or the SaCas9n domain can bind to a nucleic acid seuqnce having a non-canonical PAM. In some embodiments, the SaCas9 domain, the SaCas9d domain, or the SaCas9n domain can bind to a nucleic acid sequence having a NNGRRT PAM sequence. In some embodiments, the SaCas9 domain comprises one or more of a E781X, a N967X, and a R1014X mutation of SEQ ID NO: 4273, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid. In some embodiments, the SaCas9 domain comprises one or more of a E781K, a N967K, and a R1014H mutation of SEQ ID NO: 4273, or one or more corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SaCas9 domain comprises a

E781K, a N967K, or a R1014H mutation of SEQ ID NO: 4273, or corresponding mutations in any of the amino acid sequences provided in SEQ ID NOs: 11-260.

[00182] In some embodiments, the Cas9 domain of any of the fusion proteins provided herein comprises an amino acid sequence that is at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to any one of SEQ ID NOs: 4273-4275. In some embodiments, the Cas9 domain of any of the fusion proteins provided herein comprises the amino acid sequence of any one of SEQ ID NOs: 4273-4275. In some embodiments, the Cas9 domain of any of the fusion proteins provided herein consists of the amino acid sequence of any one of SEQ ID NOs: 4273-4275.

Exemplary SaCas9 sequence

KRNYILGLDIGITSVGYGIIDYETRDVIDAGVRLFKEANVENNEGRRSKRGARRLKRR
RRHRIQRVKKLLFDYNLLTDHSELSGINPYEARVKGLSQKLSEEEFSAALLHLAKRRG
VHNVNEVEEDTGNELSTKEQISRNSKALEEKYVAELQLERLKKDGEVRGSINRFKTS
DYVKEAKQLLKVQKAYHQLDQSFIDTYIDLLETRRTYYEGPGEGSPFGWKDIKEWY
EMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITRDENEKLEYYEKFQIIENV
FKQKKKPTLKQIAKEILVNEEDIKGYRVTSTGKPEFTNLKVYHDIKDITARKEIIENAE
LLDQIAKILTIYQSSEDIQEELTNLNSELTQEEIEQISNLKGYTGTHNLSLKAINLILDEL
WHTNDNQIAIFNRLKLVPKKVDLSQQKEIPTTLVDDFILSPVVKRSFIQSIKVINAIIKK
YGLPNDIIIELAREKNSKDAQKMINEMQKRNRQTNERIEEIIRTTGKENAKYLIEKIKL
HDMQEGKCLYSLEAIPLEDLLNNPFNYEVDHIIPRSVSFDNSFNNKVLVKQEE**N**SKKG
NRTPFQYLSSSDSKISYETFKKHILNLAKGKGRISKTKKEYLLEERDINRFSVQKDFIN
RNLVDTRYATRGLMNLLRSYFRVNNLDVKVKSINGGFTSFLRRKWKFKKERNKGY
KHHAEDALIIANADFIFKEWKKLDKAKKVMENQMFEEKQAESMPEIETEQEYKEIFIT
PHQIKHIKDFKDYKYSHRVDKKPNRELINDTLYSTRKDDKGNTLIVNNLNGLYDKDN
DKLKKLINKSPEKLLMYHHDPQTYQKLKLIMEQYGDEKNPLYKYYEETGNYLTKYS
KKDNGPVIKKIKYYGNKLNAHLDITDDYPNSRNKVVKLSLKPYRFDVYLDNGVYKF
VTVKNLDVIKKENYYEVNSKCYEEAKKLKKISNQAEFIASFYNNDLIKINGELYRVIG
VNNDLLNRIEVNMIDITYREYLENMNDKRPPRIIKTIASKTQSIKKYSTDILGNLYEVK
SKKHPQIIKKG (SEQ ID NO: 4273)

Residue N579 of SEQ ID NO: 4273, which is underlined and in bold, may be mutated (*e.g.*, to a A579) to yield a SaCas9 nickase.

Exemplary SaCas9n sequence

KRNYILGLDIGITSVGYGIIDYETRDVIDAGVRLFKEANVENNEGRRSKRGARRLKRR
RRHRIQRVKKLLFDYNLLTDHSELSGINPYEARVKGLSQKLSEEEFSAALLHLAKRRG
VHNVNEVEEDTGNELSTKEQISRNSKALEEKYVAELQLERLKKDGEVRGSINRFKTS
DYVKEAKQLLKVQKAYHQLDQSFIDTYIDLLETRRTYYEGPGEGSPFGWKDIKEWY
EMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITRDENEKLEYYEKFQIIENV
FKQKKKPTLKQIAKEILVNEEDIKGYRVTSTGKPEFTNLKVYHDIKDITARKEIIENAE
LLDQIAKILTIYQSSEDIQEELTNLNSELTQEEIEQISNLKGYTGTHNLSLKAINLILDEL
WHTNDNQIAIFNRLKLVPKKVDLSQQKEIPTTLVDDFILSPVVKRSFIQSIKVINAIIKK
YGLPNDIIIELAREKNSKDAQKMINEMQKRNRQTNERIEEIIRTTGKENAKYLIEKIKL
HDMQEGKCLYSLEAIPLEDLLNNPFNYEVDHIIPRSVSFDNSFNNKVLVKQEE**A**SKKG
NRTPFQYLSSSDSKISYETFKKHILNLAKGKGRISKTKKEYLLEERDINRFSVQKDFIN
RNLVDTRYATRGLMNLLRSYFRVNNLDVKVKSINGGFTSFLRRKWKFKKERNKGY
KHHAEDALIIANADFIFKEWKKLDKAKKVMENQMFEEKQAESMPEIETEQEYKEIFIT
PHQIKHIKDFKDYKYSHRVDKKPNRELINDTLYSTRKDDKGNTLIVNNLNGLYDKDN
DKLKKLINKSPEKLLMYHHDPQTYQKLKLIMEQYGDEKNPLYKYYEETGNYLTKYS
KKDNGPVIKKIKYYGNKLNAHLDITDDYPNSRNKVVKLSLKPYRFDVYLDNGVYKF
VTVKNLDVIKKENYYEVNSKCYEEAKKLKKISNQAEFIASFYNNDLIKINGELYRVIG
VNNDLLNRIEVNMIDITYREYLENMNDKRPPRIIKTIASKTQSIKKYSTDILGNLYEVK
SKKHPQIIKKG (SEQ ID NO: 4274).

Residue A579 of SEQ ID NO: xx, which can be mutated from N579 of SEQ ID NO: 4274 to yield a SaCas9 nickase, is underlined and in bold.

Exemplary SaKKH Cas9

KRNYILGLDIGITSVGYGIIDYETRDVIDAGVRLFKEANVENNEGRRSKRGARRLKRR
RRHRIQRVKKLLFDYNLLTDHSELSGINPYEARVKGLSQKLSEEEFSAALLHLAKRRG
VHNVNEVEEDTGNELSTKEQISRNSKALEEKYVAELQLERLKKDGEVRGSINRFKTS
DYVKEAKQLLKVQKAYHQLDQSFIDTYIDLLETRRTYYEGPGEGSPFGWKDIKEWY
EMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITRDENEKLEYYEKFQIIENV
FKQKKKPTLKQIAKEILVNEEDIKGYRVTSTGKPEFTNLKVYHDIKDITARKEIIENAE
LLDQIAKILTIYQSSEDIQEELTNLNSELTQEEIEQISNLKGYTGTHNLSLKAINLILDEL
WHTNDNQIAIFNRLKLVPKKVDLSQQKEIPTTLVDDFILSPVVKRSFIQSIKVINAIIKK
YGLPNDIIIELAREKNSKDAQKMINEMQKRNRQTNERIEEIIRTTGKENAKYLIEKIKL
HDMQEGKCLYSLEAIPLEDLLNNPFNYEVDHIIPRSVSFDNSFNNKVLVKQEE**A**SKKG

NRTPFQYLSSSDSKISYETFKKHILNLAKGKGRISKTKKEYLLEERDINRFSVQKDFIN
RNLVDTRYATRGLMNLLRSYFRVNNLDVKVKSINGGFTSFLRRKWKFKKERNKGY
KHHAEDALIIANADFIFKEWKKLDKAKKVMENQMFEEKQAESMPEIETEQEYKEIFIT
PHQIKHIKDFKDYKYSHRVDKKPNR*K*LINDTLYSTRKDDKGNTLIVNNLNGLYDKD
NDKLKKLINKSPEKLLMYHHDPQTYQKLKLIMEQYGDEKNPLYKYYEETGNYLTKY
SKKDNGPVIKKIKYYGNKLNAHLDITDDYPNSRNKVVKLSLKPYRFDVYLDNGVYK
FVTVKNLDVIKKENYYEVNSKCYEEAKKLKKISNQAEFIASFY*K*NDLIKINGELYRVI
GVNNDLLNRIEVNMIDITYREYLENMNDKRPP*H*IIKTIASKTQSIKKYSTDILGNLYEV
KSKKHPQIIKKG (SEQ ID NO: 4275).

Residue A579 of SEQ ID NO: 4275, which can be mutated from N579 of SEQ ID NO: 4275 to yield a SaCas9 nickase, is underlined and in bold. Residues K781, K967, and H1014 of SEQ ID NO: 4275, which can be mutated from E781, N967, and R1014 of SEQ ID NO: 4275 to yield a SaKKH Cas9 are underlined and in italics.

[00183] In some embodiments, the Cas9 domain is a Cas9 domain from *Streptococcus pyogenes* (SpCas9). In some embodiments, the SpCas9 domain is a nuclease active SpCas9, a nuclease inactive SpCas9 (SpCas9d), or a SpCas9 nickase (SpCas9n). In some embodiments, the SpCas9 comprises the amino acid sequence SEQ ID NO: 4276. In some embodiments, the SpCas9 comprises a D9X mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid except for D. In some embodiments, the SpCas9 comprises a D9A mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SpCas9 domain, the SpCas9d domain, or the SpCas9n domain can bind to a nucleic acid seuqnce having a non-canonical PAM. In some embodiments, the SpCas9 domain, the SpCas9d domain, or the SpCas9n domain can bind to a nucleic acid sequence having a NGG, a NGA, or a NGCG PAM sequence. In some embodiments, the SpCas9 domain comprises one or more of a D1134X, a R1334X, and a T1336X mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid. In some embodiments, the SpCas9 domain comprises one or more of a D1134E, R1334Q, and T1336R mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SpCas9 domain comprises a D1134E, a R1334Q, and a T1336R mutation of SEQ ID NO: 4276, or corresponding mutations in any of the amino acid

sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SpCas9 domain comprises one or more of a D1134X, a R1334X, and a T1336X mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid. In some embodiments, the SpCas9 domain comprises one or more of a D1134V, a R1334Q, and a T1336R mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SpCas9 domain comprises a D1134V, a R1334Q, and a T1336R mutation of SEQ ID NO: 4276, or corresponding mutations in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SpCas9 domain comprises one or more of a D1134X, a G1217X, a R1334X, and a T1336X mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid. In some embodiments, the SpCas9 domain comprises one or more of a D1134V, a G1217R, a R1334Q, and a T1336R mutation of SEQ ID NO: 4276, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the SpCas9 domain comprises a D1134V, a G1217R, a R1334Q, and a T1336R mutation of SEQ ID NO: 4276, or corresponding mutations in any of the amino acid sequences provided in SEQ ID NOs: 11-260.

[00184] In some embodiments, the Cas9 domain of any of the fusion proteins provided herein comprises an amino acid sequence that is at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to any one of SEQ ID NOs: 4276-4280. In some embodiments, the Cas9 domain of any of the fusion proteins provided herein comprises the amino acid sequence of any one of SEQ ID NOs: 4276-4280. In some embodiments, the Cas9 domain of any of the fusion proteins provided herein consists of the amino acid sequence of any one of SEQ ID NOs: 4276-4280.

Exemplary SpCas9

DKKYSIGLDIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY

KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTF
DNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAW
MTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVY
NELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDS
VEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO:
4276)

Exemplary SpCas9n

DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTF
DNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAW
MTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVY
NELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDS
VEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK

VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO:
4277)

Exemplary SpEQR Cas9

DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTF
DNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAW
MTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVY
NELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDS
VEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGF**E**SPTVAYSVLVV

AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRK**Q**Y**R**STKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 4278)

Residues E1134, Q1334, and R1336 of SEQ ID NO: 4278, which can be mutated from D1134, R1334, and T1336 of SEQ ID NO: 4278 to yield a SpEQR Cas9, are underlined and in bold.

Exemplary SpVQR Cas9

DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTF
DNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAW
MTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVY
NELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDS
VEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGF**V**SPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA

PAAFKYFDTTIDRK**Q**Y**R**STKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 4279)

Residues V1134, Q1334, and R1336 of SEQ ID NO: 4279, which can be mutated from D1134, R1334, and T1336 of SEQ ID NO: 4279 to yield a SpVQR Cas9, are underlined and in bold.

Exemplary SpVRER Cas9

DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTF
DNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAW
MTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVY
NELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDS
VEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERL
KTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRN
FMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVK
VMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQL
QNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGK
ATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSM
PQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGF**V**SPTVAYSVLVV
AKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFE
LENGRKRMLASA**R**ELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQ
HKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRK**E**Y**R**STKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 4280)

Residues V1134, R1217, Q1334, and R1336 of SEQ ID NO: 4280, which can be mutated from D1134, G1217, R1334, and T1336 of SEQ ID NO: 4280 to yield a SpVRER Cas9, are underlined and in bold.

[00185] The following are exemplary fusion proteins (*e.g.*, base editing proteins) capable of binding to a nucleic acid sequence having a non-canonical (*e.g.*, a non-NGG) PAM sequence:

Exemplary SaBE3 (rAPOBEC1-XTEN-SaCas9n-UGI-NLS)
MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKSGSET
PGTSESATPESKRNYILGLDIGITSVGYGIIDYETRDVIDAGVRLFKEANVENNEGRRS
KRGARRLKRRRRHRIQRVKKLLFDYNLLTDHSELSGINPYEARVKGLSQKLSEEEFS
AALLHLAKRRGVHNVNEVEEDTGNELSTKEQISRNSKALEEKYVAELQLERLKKDG
EVRGSINRFKTSDYVKEAKQLLKVQKAYHQLDQSFIDTYIDLLETRRTYYEGPGEGSP
FGWKDIKEWYEMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITRDENEKL
EYYEKFQIIENVFKQKKKPTLKQIAKEILVNEEDIKGYRVTSTGKPEFTNLKVYHDIK
DITARKEIIENAELLDQIAKILTIYQSSEDIQEELTNLNSELTQEEIEQISNLKGYTGTHN
LSLKAINLILDELWHTNDNQIAIFNRLKLVPKKVDLSQQKEIPTTLVDDFILSPVVKRS
FIQSIKVINAIIKKYGLPNDIIIELAREKNSKDAQKMINEMQKRNRQTNERIEEIIRTTGK
ENAKYLIEKIKLHDMQEGKCLYSLEAIPLEDLLNNPFNYEVDHIIPRSVSFDNSFNNKV
LVKQEE**A**SKKGNRTPFQYLSSSDSKISYETFKKHILNLAKGKGRISKTKKEYLLEERDI
NRFSVQKDFINRNLVDTRYATRGLMNLLRSYFRVNNLDVKVKSINGGFTSFLRRKW
KFKKERNKGYKHHAEDALIIANADFIFKEWKKLDKAKKVMENQMFEEKQAESMPEI
ETEQEYKEIFITPHQIKHIKDFKDYKYSHRVDKKPNRELINDTLYSTRKDDKGNTLIV
NNLNGLYDKDNDKLKKLINKSPEKLLMYHHDPQTYQKLKLIMEQYGDEKNPLYKY
YEETGNYLTKYSKKDNGPVIKKIKYYGNKLNAHLDITDDYPNSRNKVVKLSLKPYRF
DVYLDNGVYKFVTVKNLDVIKKENYYEVNSKCYEEAKKLKKISNQAEFIASFYNND
LIKINGELYRVIGVNNDLLNRIEVNMIDITYREYLENMNDKRPPRIIKTIASKTQSIKKY
STDILGNLYEVKSKKHPQIIKKGSGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIG
NKPESDILVHTAYDESTDENVMLLTSDAPEYKPWALVIQDSNGENKIKMLSGGSPKK
KRKV (SEQ ID NO: 4281)

Exemplary SaKKH-BE3 (rAPOBEC1-XTEN-SaCas9n-UGI-NLS)
MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKSGSET
PGTSESATPESKRNYILGLDIGITSVGYGIIDYETRDVIDAGVRLFKEANVENNEGRRS
KRGARRLKRRRRHRIQRVKKLLFDYNLLTDHSELSGINPYEARVKGLSQKLSEEEFS
AALLHLAKRRGVHNVNEVEEDTGNELSTKEQISRNSKALEEKYVAELQLERLKKDG
EVRGSINRFKTSDYVKEAKQLLKVQKAYHQLDQSFIDTYIDLLETRRTYYEGPGEGSP
FGWKDIKEWYEMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITRDENEKL
EYYEKFQIIENVFKQKKKPTLKQIAKEILVNEEDIKGYRVTSTGKPEFTNLKVYHDIK
DITARKEIIENAELLDQIAKILTIYQSSEDIQEELTNLNSELTQEEIEQISNLKGYTGTHN
LSLKAINLILDELWHTNDNQIAIFNRLKLVPKKVDLSQQKEIPTTLVDDFILSPVVKRS
FIQSIKVINAIIKKYGLPNDIIIELAREKNSKDAQKMINEMQKRNRQTNERIEEIIRTTGK

ENAKYLIEKIKLHDMQEGKCLYSLEAIPLEDLLNNPFNYEVDHIIPRSVSFDNSFNNKV
LVKQEEASKKGNRTPFQYLSSSDSKISYETFKKHILNLAKGKGRISKTKKEYLLEERDI
NRFSVQKDFINRNLVDTRYATRGLMNLLRSYFRVNNLDVKVKSINGGFTSFLRRKW
KFKKERNKGYKHHAEDALIIANADFIFKEWKKLDKAKKVMENQMFEEKQAESMPEI
ETEQEYKEIFITPHQIKHIKDFKDYKYSHRVDKKPNR*K*LINDTLYSTRKDDKGNTLIV
NNLNGLYDKDNDKLKKLINKSPEKLLMYHHDPQTYQKLKLIMEQYGDEKNPLYKY
YEETGNYLTKYSKKDNGPVIKKIKYYGNKLNAHLDITDDYPNSRNKVVKLSLKPYRF
DVYLDNGVYKFVTVKNLDVIKKENYYEVNSKCYEEAKKLKKISNQAEFIASFY*K*ND
LIKINGELYRVIGVNNDLLNRIEVNMIDITYREYLENMNDKRPP*H*IIKTIASKTQSIKKY
STDILGNLYEVKSKKHPQIIKKGSGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIG
NKPESDILVHTAYDESTDENVMLLTSDAPEYKPWALVIQDSNGENKIKMLSGGSPKK
KRKV (SEQ ID NO: 4282)

Exemplary EQR-BE3 (rAPOBEC1-XTEN-Cas9n-UGI-NLS)
MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKSGSET
PGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLI
GALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFL
VEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIK
FRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRR
LENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNL
LAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLK
ALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLN
REDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGP
LARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPK
HSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKE
DYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLF
EDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFL
KSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTV
KVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKE
HPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDN
KVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLS
ELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFR
KDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMI
AKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFA
TVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGF*E*SPT
VAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLII
KLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDN
EQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIH
LFTLTNLGAPAAFKYFDTTIDRK*Q*Y*R*STKEVLDATLIHQSITGLYETRIDLSQLGGDSG
GSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLL
TSDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV (SEQ ID NO: 4283)

VQR-BE3 (rAPOBEC1-XTEN-Cas9n-UGI-NLS)
MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKSGSET

PGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLI
GALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFL
VEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIK
FRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRR
LENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNL
LAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLK
ALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLN
REDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGP
LARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPK
HSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKE
DYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLF
EDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFL
KSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTV
KVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKE
HPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDN
KVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLS
ELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFR
KDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMI
AKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFA
TVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFVSPT
VAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLII
KLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDN
EQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIH
LFTLTNLGAPAAFKYFDTTIDRKQYRSTKEVLDATLIHQSITGLYETRIDLSQLGGDSG
GSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLL
TSDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV (SEQ ID NO: 4284)

VRER-BE3 (rAPOBEC1-XTEN-Cas9n-UGI-NLS)

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKSGSET
PGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLI
GALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFL
VEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIK
FRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRR
LENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNL
LAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLK
ALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLN
REDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGP
LARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPK
HSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKE
DYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLF
EDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFL
KSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTV
KVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKE
HPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDN
KVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLS
ELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFR
KDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMI

AKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFA TVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGF*V*SPT VAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLII KLPKYSLFELENGRKRMLASA*R*ELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNE QKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHL FTLTNLGAPAAFKYFDTTIDRK*EYR*STKEVLDATLIHQSITGLYETRIDLSQLGGDSGG STNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLLTS DAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV (SEQ ID NO: 4285)

High Fidelity Base Editors

[00186] Some aspects of the disclosure provide Cas9 fusion proteins (*e.g.*, any of the fusion proteins provided herein) comprising a Cas9 domain that has high fidelity. Additional aspects of the disclosure provide Cas9 fusion proteins (*e.g.*, any of the fusion proteins provided herein) comprising a Cas9 domain with decreased electrostatic interactions between the Cas9 domain and a sugar-phosphate backbone of a DNA, as compared to a wild-type Cas9 domain. In some embodiments, a Cas9 domain (*e.g.*, a wild type Cas9 domain) comprises one or more mutations that decreases the association between the Cas9 domain and a sugar-phosphate backbone of a DNA. In some embodiments, any of the Cas9 fusion proteins provided herein comprise one or more of a N497X, a R661X, a Q695X, and/or a Q926X mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, wherein X is any amino acid. In some embodiments, any of the Cas9 fusion proteins provided herein comprise one or more of a N497A, a R661A, a Q695A, and/or a Q926A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the Cas9 domain comprises a D10A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the Cas9 domain (*e.g.*, of any of the fusion proteins provided herein) comprises the amino acid sequence as set forth in SEQ ID NO: 325. In some embodiments, the fusion protein comprises the amino acid sequence as set forth in SEQ ID NO: 285. Cas9 domains with high fidelity are known in the art and would be apparent to the skilled artisan. For example, Cas9 domains with high fidelity have been described in Kleinstiver, B.P., *et al.* "High-fidelity CRISPR-Cas9 nucleases with no detectable genome-wide off-target effects." *Nature* 529, 490-495 (2016); and Slaymaker, I.M., *et al.* "Rationally engineered Cas9 nucleases with improved specificity." *Science* 351, 84-88 (2015); the entire contents of each are incorporated herein by reference.

[00187] It should be appreciated that the base editors provided herein, for example base editor 2 (BE2) or base editor 3 (BE3), may be converted into high fidelity base editors by modifyint the Cas9 domain as described herein to generate high fidelity base editors, for example high fidelity base editor 2 (HF-BE2) or high fidelity base editor 3 (HF-BE3). In some embodiments, base editor 2 (BE2) comprises a deaminase domain, a dCas9, and a UGI domain. In some embodiments, base editor 3 (BE3) comprises a deaminase domain an nCas9 domain and a UGI domain.

Cas9 domain where mutations relative to Cas9 of SEQ ID NO: 10 are shown in bold and underlines

DKKYSIGL**A**IGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARR
RYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRK
KLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAK
AILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLL
AQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYK
EIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGE
LHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASA
QSFIERMT**A**FDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTN
RKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFE
DREMIEERLKTYAHLFDDKVMKQLKRRRYTGWG**A**LSRKLINGIRDKQSGKTILDFLKSDGFANRNFM
ALIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEM
ARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDIN
RLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFD
NLTKAERGGLSELDKAGFIKRQLVETR**A**ITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFR
KDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATA
KYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTG
GFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERS
SFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASH
YEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIH
LFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 325)

HF-BE3

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKHVEVNFIEKF
TTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGLRDLISSGVTI
QIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQ
SCHYQRLPPHILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLG
NTDRHSIKKNLIGALLFDSGETAEATRLKRTARRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFL
VEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLN

PDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALS
LGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITK
APLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKM
DGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGP
LARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTAFDKNLPNEKVLPKHSLLYEYFTVYN
ELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASL
GTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWG
ALSRKLINGIRDKQSGKTILDFLKSDGFANRNFMALIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGS
PAIKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEH
PVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGKS
DNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRAITKHVAQIL
DSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPK
LESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIV
WDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVA
YSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGR
KRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKR
VILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIH
QSITGLYETRIDLSQLGGD (SEQ ID NO: 285)

Cas9 fusion proteins

[00188] Any of the Cas9 domains (e.g., a nuclease active Cas9 protein, a nuclease-inactive dCas9 protein, or a Cas9 nickase protein) disclosed herein may be fused to a second protein, thus fusion proteins provided herein comprise a Cas9 domain as provided herein and a second protein, or a "fusion partner". In some embodiments, the second protein is fused to the N-terminus of the Cas9 domain. However, in other embodiments, the second protein is fused to the C-terminus of the Cas9 domain. In some embodiments, the second protein that is fused to the Cas9 domain is a nucleic acid editing domain. In some embodiments, the Cas9 domain and the nucleic acid editing domain are fused via a linker, while in other embodiments the Cas9 domain and the nucleic acid editing domain are fused directly to one another. In some embodiments, the linker comprises $(GGGS)_n$ (SEQ ID NO: 265), $(GGGGS)_n$ (SEQ ID NO: 5), $(G)_n$, $(EAAAK)_n$ (SEQ ID NO: 6), $(GGS)_n$, $(SGGS)_n$ (SEQ ID NO: 4288), SGSETPGTSESATPES (SEQ ID NO: 7), or $(XP)_n$ motif, or a combination of any of these, wherein n is independently an integer between 1 and 30, and wherein X is any amino acid. In some embodiments, the linker comprises a $(GGS)_n$ motif, wherein n is 1, 3, or 7. In some embodiments, the linker comprises a $(GGS)_n$ motif, wherein n is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, or 15. In some embodiments, the linker comprises an amino acid sequence of

SGSETPGTSESATPES (SEQ ID NO: 7),also referred to as the XTEN linker in the Examples). The length of the linker can influence the base to be edited, as illustrated in the Examples. For example, a linker of 3-amino-acid long (*e.g.*, $(GGS)_1$) may give a 2-5, 2-4, 2-3, 3-4 base editing window relative to the PAM sequence, while a 9-amino-acid linker (*e.g.*, $(GGS)_3$ (SEQ ID NO: 596)) may give a 2-6, 2-5, 2-4, 2-3, 3-6, 3-5, 3-4, 4-6, 4-5, 5-6 base editing window relative to the PAM sequence. A 16-amino-acid linker (*e.g.*, the XTEN linker) may give a 2-7, 2-6, 2-5, 2-4, 2-3, 3-7, 3-6, 3-5, 3-4, 4-7, 4-6, 4-5, 5-7, 5-6, 6-7 base window relative to the PAM sequence with exceptionally strong activity, and a 21-amino-acid linker (*e.g.*, $(GGS)_7$ (SEQ ID NO: 597)) may give a 3-8, 3-7, 3-6, 3-5, 3-4, 4-8, 4-7, 4-6, 4-5, 5-8, 5-7, 5-6, 6-8, 6-7, 7-8 base editing window relative to the PAM sequence. The novel finding that varying linker length may allow the dCas9 fusion proteins of the disclosure to edit nucleobases different distances from the PAM sequence affords siginicant clinical importance, since a PAM sequence may be of varying distance to the disease-causing mutation to be corrected in a gene. It is to be understood that the linker lengths described as examples here are not meant to be limiting.

[00189] In some embodiments, the second protein comprises an enzymatic domain. In some embodiments, the enzymatic domain is a nucleic acid editing domain. Such a nucleic acid editing domain may be, without limitation, a nuclease, a nickase, a recombinase, a deaminase, a methyltransferase, a methylase, an acetylase, or an acetyltransferase. Non-limiting exemplary binding domains that may be used in accordance with this disclosure include transcriptional activator domains and transcriptional repressor domains.

Deaminase Domains

[00190] In some embodiments, second protein comprises a nucleic acid editing domain. In some embodiments, the nucleic acid editing domain can catalyze a C to U base change. In some embodiments, the nucleic acid editing domain is a deaminase domain. In some embodiments, the deaminase is a cytidine deaminase or a cytidine deaminase. In some embodiments, the deaminase is an apolipoprotein B mRNA-editing complex (APOBEC) family deaminase. In some embodiments, the deaminase is an APOBEC1 deaminase. In some embodiments, the deaminase is an APOBEC2 deaminase. In some embodiments, the deaminase is an APOBEC3 deaminase. In some embodiments, the deaminase is an APOBEC3A deaminase. In some embodiments, the deaminase is an APOBEC3B deaminase. In some embodiments, the deaminase is an APOBEC3C deaminase. In some embodiments, the deaminase is an APOBEC3D deaminase. In some embodiments, the deaminase is an

APOBEC3E deaminase. In some embodiments, the deaminase is an APOBEC3F deaminase. In some embodiments, the deaminase is an APOBEC3G deaminase. In some embodiments, the deaminase is an APOBEC3H deaminase. In some embodiments, the deaminase is an APOBEC4 deaminase. In some embodiments, the deaminase is an activation-induced deaminase (AID). In some embodiments, the deaminase is a vertebrate deaminase. In some embodiments, the deaminase is an invertebrate deaminase. In some embodiments, the deaminase is a human, chimpanzee, gorilla, monkey, cow, dog, rat, or mouse deaminase. In some embodiments, the deaminase is a human deaminase. In some embodiments, the deaminase is a rat deaminase, *e.g.,* rAPOBEC1. In some embodiments, the deaminase is a *Petromyzon marinus* cytidine deaminase 1 (pmCDA1). In some embodiments, the deminase is a human APOBEC3G (SEQ ID NO: 275). In some embodiments, the deaminase is a fragment of the human APOBEC3G (SEQ ID NO: 5740). In some embodiments, the deaminase is a human APOBEC3G variant comprising a D316R_D317R mutation (SEQ ID NO: 5739). In some embodiments, the deaminase is a frantment of the human APOBEC3G and comprising mutations corresponding to the D316R_D317R mutations in SEQ ID NO: 275 (SEQ ID NO: 5741).

[00191] In some embodiments, the nucleic acid editing domain is at least 80%, at least 85%, at least 90%, at least 92%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the deaminase domain of any one of SEQ ID NOs: 266-284, 607-610, 5724-5736, or 5738-5741. In some embodiments, the nucleic acid editing domain comprises the amino acid sequence of any one of SEQ ID NOs: 266-284, 607-610, 5724-5736, or 5738-5741.

Deaminase Domains that Modulate the Editing Window of Base Editors

[00192] Some aspects of the disclosure are based on the recognition that modulating the deaminase domain catalytic activity of any of the fusion proteins provided herein, for example by making point mutations in the deaminase domain, affect the processivity of the fusion proteins (*e.g.,* base editors). For example, mutations that reduce, but do not eliminate, the catalytic activity of a deaminase domain within a base editing fusion protein can make it less likely that the deaminase domain will catalyze the deamination of a residue adjacent to a target residue, thereby narrowing the deamination window. The ability to narrow the deaminataion window may prevent unwanted deamination of residues adjacent of specific target residues, which may decrease or prevent off-target effects.

[00193] In some embodiments, any of the fusion proteins provided herein comprise a deaminase domain (*e.g.,* a cytidine deaminase domain) that has reduced catalytic deaminase activity. In some embodiments, any of the fusion proteins provided herein comprise a deaminase domain (*e.g.,* a cytidine deaminase domain) that has a reduced catalytic deaminase activity as compared to an appropriate control. For example, the appropriate control may be the deaminase activity of the deaminase prior to introducing one or more mutations into the deaminase. In other embodiments, the appropriate control may be a wild-type deaminase. In some embodiments, the appropriate control is a wild-type apolipoprotein B mRNA-editing complex (APOBEC) family deaminase. In some embodiments, the appropriate control is an APOBEC1 deaminase, an APOBEC2 deaminase, an APOBEC3A deaminase, an APOBEC3B deaminase, an APOBEC3C deaminase, an APOBEC3D deaminase, an APOBEC3F deaminase, an APOBEC3G deaminase, or an APOBEC3H deaminase. In some embodiments, the appropriate control is an activation induced deaminase (AID). In some embodiments, the appropriate control is a cytidine deaminase 1 from *Petromyzon marinu*s (pmCDA1). In some embodiments, the deaminse domain may be a deaminase domain that has at least 1%, at least 5%, at least 15%, at least 20%, at least 25%, at least 30%, at least 40%, at least 50%, at least 60%, at lest 70%, at least 80%, at least 90%, or at least 95% less catalytic deaminase activity as compared to an appropriate control.

[00194] In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising one or more mutations selected from the group consisting of H121X, H122X, R126X, R126X, R118X, W90X, W90X, and R132X of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase, wherin X is any amino acid. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising one or more mutations selected from the group consisting of H121R, H122R, R126A, R126E, R118A, W90A, W90Y, and R132E of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase.

[00195] In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising one or more mutations selected from the group consisting of D316X, D317X, R320X, R320X, R313X, W285X, W285X, R326X of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase, wherin X is any amino acid. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising one or more mutations selected from the group consisting of D316R, D317R, R320A, R320E, R313A, W285A, W285Y, R326E of

hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase.

[00196] In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a H121R and a H122Rmutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R126A mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R126E mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R118A mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W90A mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W90Y mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R132E mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W90Y and a R126E mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R126E and a R132E mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W90Y and a R132E mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W90Y, R126E, and R132E mutation of rAPOBEC1 (SEQ ID NO: 284), or one or more corresponding mutations in another APOBEC deaminase.

[00197] In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a D316R and a D317R mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R320A mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R320E mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R313A mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W285A mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W285Y mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase.In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R326E mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W285Y and a R320E mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a R320E and a R326E mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W285Y and a R326E mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase. In some embodiments, any of the fusion proteins provided herein comprise an APOBEC deaminase comprising a W285Y, R320E, and R326E mutation of hAPOBEC3G (SEQ ID NO: 275), or one or more corresponding mutations in another APOBEC deaminase.

[00198] Some aspects of this disclosure provide fusion proteins comprising (i) a nuclease-inactive Cas9 domain; and (ii) a nucleic acid editing domain. In some embodiments, a

nuclease-inactive Cas9 domain (dCas9), comprises an amino acid sequence that is at least 80%, at least 85%, at least 90%, at least 92%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the amino acid sequence of a Cas9 as provided by any one of SEQ ID NOs: 10-263, and comprises mutations that inactivate the nuclease activity of Cas9. Mutations that render the nuclease domains of Cas9 inactive are well-known in the art. For example, the DNA cleavage domain of Cas9 is known to include two subdomains, the HNH nuclease subdomain and the RuvC1 subdomain. The HNH subdomain cleaves the strand complementary to the gRNA, whereas the RuvC1 subdomain cleaves the non-complementary strand. Mutations within these subdomains can silence the nuclease activity of Cas9. For example, the mutations D10A and H840A completely inactivate the nuclease activity of *S. pyogenes* Cas9 (Jinek *et al., Science.* 337:816-821(2012); Qi *et al., Cell.* 28;152(5):1173-83 (2013)). In some embodiments, the dCas9 of this disclosure comprises a D10A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the dCas9 of this disclosure comprises a H840A mutation of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the dCas9 of this disclosure comprises both D10A and H840A mutations of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. In some embodiments, the Cas9 further comprises a histidine residue at position 840 of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. The presence of the catalytic residue H840 restores the acvitity of the Cas9 to cleave the non-edited strand containing a G opposite the targeted C. Restoration of H840 does not result in the cleavage of the target strand containing the C. In some embodiments, the dCas9 comprises an amino acid sequence of SEQ ID NO: 263. It is to be understood that other mutations that inactivate the nuclease domains of Cas9 may also be included in the dCas9 of this disclosure.

[00199] The Cas9 or dCas9 domains comprising the mutations disclosed herein, may be a full-length Cas9, or a fragment thereof. In some embodiments, proteins comprising Cas9, or fragments thereof, are referred to as "Cas9 variants." A Cas9 variant shares homology to Cas9, or a fragment thereof. For example a Cas9 variant is at least about 70% identical, at least about 80% identical, at least about 90% identical, at least about 95% identical, at least about 96% identical, at least about 97% identical, at least about 98% identical, at least about

99% identical, at least about 99.5% identical, or at least about 99.9% to wild type Cas9. In some embodiments, the Cas9 variant comprises a fragment of Cas9 (*e.g.*, a gRNA binding domain or a DNA-cleavage domain), such that the fragment is at least about 70% identical, at least about 80% identical, at least about 90% identical, at least about 95% identical, at least about 96% identical, at least about 97% identical, at least about 98% identical, at least about 99% identical, at least about 99.5% identical, or at least about 99.9% identical to the corresponding fragment of wild type Cas9, *e.g.*, a Cas9 comprising the amino acid sequence of SEQ ID NO: 10.

[00200] Any of the Cas9 fusion proteins of this disclosure may further comprise a nucleic acid editing domain (*e.g.,* an enzyme that is capable of modifying nucleic acid, such as a deaminase). In some embodiments, the nucleic acid editing domain is a DNA-editing domain. In some embodiments, the nucleic acid editing domain has deaminase activity. In some embodiments, the nucleic acid editing domain comprises or consists of a deaminase or deaminase domain. In some embodiments, the deaminase is a cytidine deaminase. In some embodiments, the deaminase is an apolipoprotein B mRNA-editing complex (APOBEC) family deaminase. In some embodiments, the deaminase is an APOBEC1 family deaminase. In some embodiments, the deaminase is an activation-induced cytidine deaminase (AID). Some nucleic-acid editing domains as well as Cas9 fusion proteins including such domains are described in detail herein. Additional suitable nucleic acid editing domains will be apparent to the skilled artisan based on this disclosure and knowledge in the field.

[00201] Some aspects of the disclosure provide a fusion protein comprising a Cas9 domain fused to a nucleic acid editing domain, wherein the nucleic acid editing domain is fused to the N-terminus of the Cas9 domain. In some embodiments, the Cas9 domain and the nucleic acid editing-editing domain are fused via a linker. In some embodiments, the linker comprises a $(GGGS)_n$ (SEQ ID NO: 265), a $(GGGGS)_n$ (SEQ ID NO: 5), a $(G)_n$, an $(EAAAK)_n$ (SEQ ID NO: 6), a $(GGS)_n$, $(SGGS)_n$ (SEQ ID NO: 4288), an SGSETPGTSESATPES (SEQ ID NO: 7) motif (see, *e.g.*, Guilinger JP, Thompson DB, Liu DR. Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. *Nat. Biotechnol.* 2014**;** 32(6): 577-82; the entire contents are incorporated herein by reference), or an $(XP)_n$ motif, or a combination of any of these, wherein n is independently an integer between 1 and 30. In some embodiments, n is independently 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, or 30, or, if more than one linker or more than one linker motif is present, any combination thereof. In some embodiments, the linker comprises a $(GGS)_n$ motif,

wherein n is 1,2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15. In some embodiments, the linker comprises a $(GGS)_n$ motif, wherein n is 1, 3, or 7. In some embodiments, the linker comprises the amino acid sequence SGSETPGTSESATPES (SEQ ID NO: 7). Additional suitable linker motifs and linker configurations will be apparent to those of skill in the art. In some embodiments, suitable linker motifs and configurations include those described in Chen *et al.*, Fusion protein linkers: property, design and functionality. *Adv Drug Deliv Rev.* 2013; 65(10):1357-69, the entire contents of which are incorporated herein by reference. Additional suitable linker sequences will be apparent to those of skill in the art based on the instant disclosure. In some embodiments, the general architecture of exemplary Cas9 fusion proteins provided herein comprises the structure:

$[NH_2]$-[nucleic acid editing domain]-[Cas9]-[COOH] or

$[NH_2]$-[nucleic acid editing domain]-[linker]-[Cas9]-[COOH],

wherein NH_2 is the N-terminus of the fusion protein, and COOH is the C-terminus of the fusion protein.

[00202] The fusion proteins of the present disclosure may comprise one or more additional features. For example, in some embodiments, the fusion protein comprises a nuclear localization sequence (NLS). In some embodiments, the NLS of the fusion protein is localized between the nucleic acid editing domain and the Cas9 domain. In some embodiments, the NLS of the fusion protein is localized C-terminal to the Cas9 domain.

[00203] Other exemplary features that may be present are localization sequences, such as cytoplasmic localization sequences, export sequences, such as nuclear export sequences, or other localization sequences, as well as sequence tags that are useful for solubilization, purification, or detection of the fusion proteins. Suitable protein tags provided herein include, but are not limited to, biotin carboxylase carrier protein (BCCP) tags, myc-tags, calmodulin-tags, FLAG-tags, hemagglutinin (HA)-tags, polyhistidine tags, also referred to as histidine tags or His-tags, maltose binding protein (MBP)-tags, nus-tags, glutathione-S-transferase (GST)-tags, green fluorescent protein (GFP)-tags, thioredoxin-tags, S-tags, Softags (*e.g.*, Softag 1, Softag 3), strep-tags , biotin ligase tags, FlAsH tags, V5 tags, and SBP-tags. Additional suitable sequences will be apparent to those of skill in the art. In some embodiments, the fusion protein comprises one or more His tags.

[00204] In some embodiments, the nucleic acid editing domain is a deaminase. For example, in some embodiments, the general architecture of exemplary Cas9 fusion proteins with a deaminase domain comprises the structure:

$[NH_2]$-[NLS]-[deaminase]-[Cas9]-[COOH],

$[NH_2]$-[Cas9]-[deaminase]-[COOH],

$[NH_2]$-[deaminase]-[Cas9]-[COOH], or

$[NH_2]$-[deaminase]-[Cas9]-[NLS]-[COOH]

wherein NLS is a nuclear localization sequence, NH_2 is the N-terminus of the fusion protein, and COOH is the C-terminus of the fusion protein. Nuclear localization sequences are known in the art and would be apparent to the skilled artisan. For example, NLS sequences are described in Plank *et al.*, PCT/EP2000/011690, the contents of which are incorporated herein by reference for their disclosure of exemplary nuclear localization sequences. In some embodiments, a NLS comprises the amino acid sequence PKKKRKV (SEQ ID NO: 741) or MDSLLMNRRKFLYQFKNVRWAKGRRETYLC (SEQ ID NO: 742). In some embodiments, a linker is inserted between the Cas9 and the deaminase. In some embodiments, the NLS is located C-terminal of the Cas9 domain. In some embodiments, the NLS is located N-terminal of the Cas9 domain. In some embodiments, the NLS is located between the deaminase and the Cas9 domain. In some embodiments, the NLS is located N-terminal of the deaminase domain. In some embodiments, the NLS is located C-terminal of the deaminase domain.

[00205] One exemplary suitable type of nucleic acid editing domain is a cytidine deaminase, for example, of the APOBEC family. The apolipoprotein B mRNA-editing complex (APOBEC) family of cytidine deaminase enzymes encompasses eleven proteins that serve to initiate mutagenesis in a controlled and beneficial manner.[29] One family member, activation-induced cytidine deaminase (AID), is responsible for the maturation of antibodies by converting cytosines in ssDNA to uracils in a transcription-dependent, strand-biased fashion.[30] The apolipoprotein B editing complex 3 (APOBEC3) enzyme provides protection to human cells against a certain HIV-1 strain via the deamination of cytosines in reverse-transcribed viral ssDNA.[31] These proteins all require a Zn^{2+}-coordinating motif (His-X-Glu-X_{23-26}-Pro-Cys-X_{2-4}-Cys; SEQ ID NO: 598) and bound water molecule for catalytic activity. The Glu residue acts to activate the water molecule to a zinc hydroxide for nucleophilic attack in the deamination reaction. Each family member preferentially deaminates at its own particular "hotspot", ranging from WRC (W is A or T, R is A or G) for hAID, to TTC for hAPOBEC3F.[32] A recent crystal structure of the catalytic domain of APOBEC3G revealed a secondary structure comprised of a five-stranded β-sheet core flanked by six α-helices, which is believed to be conserved across the entire family.[33] The active center loops have been shown to be responsible for both ssDNA binding and in determining "hotspot" identity.[34]

Overexpression of these enzymes has been linked to genomic instability and cancer, thus highlighting the importance of sequence-specific targeting.[35]

[00206] Some aspects of this disclosure relate to the recognition that the activity of cytidine deaminase enzymes such as APOBEC enzymes can be directed to a specific site in genomic DNA. Without wishing to be bound by any particular theory, advantages of using Cas9 as a recognition agent include (1) the sequence specificity of Cas9 can be easily altered by simply changing the sgRNA sequence; and (2) Cas9 binds to its target sequence by denaturing the dsDNA, resulting in a stretch of DNA that is single-stranded and therefore a viable substrate for the deaminase. It should be understood that other catalytic domains, or catalytic domains from other deaminases, can also be used to generate fusion proteins with Cas9, and that the disclosure is not limited in this regard.

[00207] Some aspects of this disclosure are based on the recognition that Cas9:deaminase fusion proteins can efficiently deaminate nucleotides at positions 3-11 according to the numbering scheme in Figure 3. In view of the results provided herein regarding the nucleotides that can be targeted by Cas9:deaminase fusion proteins, a person of skill in the art will be able to design suitable guide RNAs to target the fusion proteins to a target sequence that comprises a nucleotide to be deaminated.

[00208] In some embodiments, the deaminase domain and the Cas9 domain are fused to each other via a linker. Various linker lengths and flexibilities between the deaminase domain (*e.g.*, AID) and the Cas9 domain can be employed (*e.g.*, ranging from very flexible linkers of the form $(GGGGS)_n$ (SEQ ID NO: 5), $(GGS)_n$, and $(G)_n$ to more rigid linkers of the form $(EAAAK)_n$ (SEQ ID NO: 6), $(SGGS)_n$ (SEQ ID NO: 4288), SGSETPGTSESATPES (SEQ ID NO: 7) (see, *e.g.*, Guilinger JP, Thompson DB, Liu DR. Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. *Nat. Biotechnol.* 2014; 32(6): 577-82; the entire contents are incorporated herein by reference) and $(XP)_n$)[36] in order to achieve the optimal length for deaminase activity for the specific application. In some embodiments, the linker comprises a $(GGS)_n$ motif, wherein n is 1, 3, or 7. In some embodiments, the linker comprises a (an SGSETPGTSESATPES (SEQ ID NO: 7) motif.

[00209] Some exemplary suitable nucleic-acid editing domains, *e.g.*, deaminases and deaminase domains, that can be fused to Cas9 domains according to aspects of this disclosure are provided below. It should be understood that, in some embodiments, the active domain of the respective sequence can be used, *e.g.*, the domain without a localizing signal (nuclear localization sequence, without nuclear export signal, cytoplasmic localizing signal).

[00210] Human AID:

MDSLLMNRRKFLYQFKNVRWAKGRRETYLCYVVKRRDSATSFSLDFGYLRNKNGC HVELLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVADFLRGNPNLSLRIFTAR LYFCEDRKAEPEGLRRLHRAGVQIAIMTFKDYFYCWNTFVENHERTFKAWEGLHEN SVRLSRQLRRILLPLYEVDDLRDAFRTLGL (SEQ ID NO: 266)

(underline: nuclear localization sequence; double underline: nuclear export signal)

[00211] Mouse AID:

MDSLLMKQKKFLYHFKNVRWAKGRHETYLCYVVKRRDSATSCSLDFGHLRNKSGC HVELLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVAEFLRWNPNLSLRIFTAR LYFCEDRKAEPEGLRRLHRAGVQIGIMTFKDYFYCWNTFVENRERTFKAWEGLHEN SVRLTRQLRRILLPLYEVDDLRDAFRMLGF (SEQ ID NO: 267)

(underline: nuclear localization sequence; double underline: nuclear export signal)

[00212] Dog AID:

MDSLLMKQRKFLYHFKNVRWAKGRHETYLCYVVKRRDSATSFSLDFGHLRNKSGC HVELLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVADFLRGYPNLSLRIFAAR LYFCEDRKAEPEGLRRLHRAGVQIAIMTFKDYFYCWNTFVENREKTFKAWEGLHEN SVRLSRQLRRILLPLYEVDDLRDAFRTLGL (SEQ ID NO: 268)

(underline: nuclear localization sequence; double underline: nuclear export signal)

[00213] Bovine AID:

MDSLLKKQRQFLYQFKNVRWAKGRHETYLCYVVKRRDSPTSFSLDFGHLRNKAGC HVELLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVADFLRGYPNLSLRIFTAR LYFCDKERKAEPEGLRRLHRAGVQIAIMTFKDYFYCWNTFVENHERTFKAWEGLHE NSVRLSRQLRRILLPLYEVDDLRDAFRTLGL (SEQ ID NO: 269)

(underline: nuclear localization sequence; double underline: nuclear export signal)

[00214] Rat AID

MAVGSKPKAALVGPHWERERIWCFLCSTGLGTQQTGQTSRWLRPAATQDPVSPPRS LLMKQRKFLYHFKNVRWAKGRHETYLCYVVKRRDSATSFSLDFGYLRNKSGCHVE LLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVADFLRGNPNLSLRIFTARLTG WGALPAGLMSPARPSDYFYCWNTFVENHERTFKAWEGLHENSVRLSRRLRRILLPL YEVDDLRDAFRTLGL (SEQ ID NO: 5725)

(underline: nuclear localization sequence; double underline: nuclear export signal)

[00215] Mouse APOBEC-3:

MGPFCLGCSHRKCYSPIRNLISQETFKFHFKNLGYAKGRKDTFLCYEVTRKDCDSPVS LHHGVFKNKDNI*HAEICFLYWFHDKVLKVLSPREEFKITWYMSWSPCFEC*AEQIVRFLA

THHNLSLDIFSSRLYNVQDPETQQNLCRLVQEGAQVAAMDLYEFKKCWKKFVDNG
GRRFRPWKRLLTNFRYQDSKLQEILRPCYIPVPSSSSSTLSNICLTKGLPETRFCVEGR
RMDPLSEEEFYSQFYNQRVKHLCYYHRMKPYLCYQLEQFNGQAPLKGCLLSEKGKQ
*HAEILFLDKIRSMELSQVTITCYLTWSPCPNC*AWQLAAFKRDRPDLILHIYTSRLYFHWK
RPFQKGLCSLWQSGILVDVMDLPQFTDCWTNFVNPKRPFWPWKGLEIISRRTQRRLR
RIKESWGLQDLVNDFGNLQLGPPMS (SEQ ID NO: 270)

(italic: nucleic acid editing domain)

[00216] Rat APOBEC-3:

MGPFCLGCSHRKCYSPIRNLISQETFKFHFKNLRYAIDRKDTFLCYEVTRKDCDSPVS
LHHGVFKNKDNI*HAEICFLYWFHDKVLKVLSPREEFKITWYMSWSPCFEC*AEQVLRFLA
THHNLSLDIFSSRLYNIRDPENQQNLCRLVQEGAQVAAMDLYEFKKCWKKFVDNGG
RRFRPWKKLLTNFRYQDSKLQEILRPCYIPVPSSSSSTLSNICLTKGLPETRFCVERRR
VHLLSEEEFYSQFYNQRVKHLCYYHGVKPYLCYQLEQFNGQAPLKGCLLSEKGKQ*H*
*AEILFLDKIRSMELSQVIITCYLTWSPCPNC*AWQLAAFKRDRPDLILHIYTSRLYFHWKR
PFQKGLCSLWQSGILVDVMDLPQFTDCWTNFVNPKRPFWPWKGLEIISRRTQRRLHR
IKESWGLQDLVNDFGNLQLGPPMS (SEQ ID NO: 271)

(italic: nucleic acid editing domain)

[00217] Rhesus macaque APOBEC-3G:

MVEPMDPRTFVSNFNNRPILSGLNTVWLCCEVKTKDPSGPPLDAKIFQGKVYSKAKY
*HPEMRFLRWFHKWRQLHHDQEYKVTWYVSWSPCTRC*ANSVATFLAKDPKVTLTIFVA
RLYYFWKPDYQQALRILCQKRGGPHATMKIMNYNEFQDCWNKFVDGRGKPFKPRN
NLPKHYTLLQATLGELLRHLMDPGTFTSNFNNKPWVSGQHETYLCYKVERLHNDT
WVPLNQHRGFLRNQAPNIHGFPKGR*HAELCFLDLIPFWKLDGQQYRVTCFTSWSPCFS*
*C*AQEMAKFISNNEHVSLCIFAARIYDDQGRYQEGLRALHRDGAKIAMMNYSEFEYC
WDTFVDRQGRPFQPWDGLDEHSQALSGRLRAI (SEQ ID NO: 272)

(italic: nucleic acid editing domain; underline: cytoplasmic localization signal)

[00218] Chimpanzee APOBEC-3G:

MKPHFRNPVERMYQDTFSDNFYNRPILSHRNTVWLCYEVKTKGPSRPPLDAKIFRGQ
VYSKLKY*HPEMRFFHWFSKWRKLHRDQEYEVTWYISWSPCTKC*TRDVATFLAEDPKV
TLTIFVARLYYFWDPDYQEALRSLCQKRDGPRATMKIMNYDEFQHCWSKFVYSQRE
LFEPWNNLPKYYILLHIMLGEILRHSMDPPTFTSNFNNELWVRGRHETYLCYEVERL
HNDTWVLLNQRRGFLCNQAPHKHGFLEGR*HAELCFLDVIPFWKLDLHQDYRVTCFTS*
*WSPCFSC*AQEMAKFISNNKHVSLCIFAARIYDDQGRCQEGLRTLAKAGAKISIMTYSE
FKHCWDTFVDHQGCPFQPWDGLEEHSQALSGRLRAILQNQGN (SEQ ID NO: 273)

(italic: nucleic acid editing domain; underline: cytoplasmic localization signal)

[00219] Green monkey APOBEC-3G:

MNPQIRNMVEQMEPDIFVYYFNNRPILSGRNTVWLCYEVKTKDPSGPPLDANIFQGK
LYPEAKD*HPEMKFLHWFRKWRQLHRDQEYEVTWYVSWSPCTRC*ANSVATFLAEDPKV
TLTIFVARLYYFWKPDYQQALRILCQERGGPHATMKIMNYNEFQHCWNEFVDGQG
KPFKPRKNLPKHYTLLHATLGELLRHVMDPGTFTSNFNNKPWVSGQRETYLCYKVE
RSHNDTWVLLNQHRGFLRNQAPDRHGFPKGR*HAELCFLDLIPFWKLDDQQYRVTCFT*
*SWSPCFSC*AQKMAKFISNNKHVSLCIFAARIYDDQGRCQEGLRTLHRDGAKIAVMNY
SEFEYCWDTFVDRQGRPFQPWDGLDEHSQALSGRLRAI (SEQ ID NO: 274)

(italic: nucleic acid editing domain; underline: cytoplasmic localization signal)

[00220] Human APOBEC-3G:

MKPHFRNTVERMYRDTFSYNFYNRPILSRRNTVWLCYEVKTKGPSRPPLDAKIFRGQ
VYSELKY*HPEMRFFHWFSKWRKLHRDQEYEVTWYISWSPCTKC*TRDMATFLAEDPKV
TLTIFVARLYYFWDPDYQEALRSLCQKRDGPRATMKIMNYDEFQHCWSKFVYSQRE
LFEPWNNLPKYYILLHIMLGEILRHSMDPPTFTFNFNNEPWVRGRHETYLCYEVERM
HNDTWVLLNQRRGFLCNQAPHKHGFLEGR*HAELCFLDVIPFWKLDLDQDYRVTCFTS*
*WSPCFSC*AQEMAKFISKNKHVSLCIFTARIYDDQGRCQEGLRTLAEAGAKISIMTYSE
FKHCWDTFVDHQGCPFQPWDGLDEHSQDLSGRLRAILQNQEN (SEQ ID NO: 275)

(italic: nucleic acid editing domain; underline: cytoplasmic localization signal)

[00221] Human APOBEC-3F:

MKPHFRNTVERMYRDTFSYNFYNRPILSRRNTVWLCYEVKTKGPSRPRLDAKIFRGQ
VYSQPEH*HAEMCFLSWFCGNQLPAYKCFQITWFVSWTPCPDC*VAKLAEFLAEHPNVTL
TISAARLYYYWERDYRRALCRLSQAGARVKIMDDEEFAYCWENFVYSEGQPFMPW
YKFDDNYAFLHRTLKEILRNPMEAMYPHIFYFHFKNLRKAYGRNESWLCFTMEVVK
HHSPVSWKRGVFRNQVDPETHC*HAERCFLSWFCDDILSPNTNYEVTWYTSWSPCPEC*A
GEVAEFLARHSNVNLTIFTARLYYFWDTDYQEGLRSLSQEGASVEIMGYKDFKYCW
ENFVYNDDEPFKPWKGLKYNFLFLDSKLQEILE (SEQ ID NO: 276)

(italic: nucleic acid editing domain)

[00222] Human APOBEC-3B:

MNPQIRNPMERMYRDTFYDNFENEPILYGRSYTWLCYEVKIKRGRSNLLWDTGVFR
GQVYFKPQY*HAEMCFLSWFCGNQLPAYKCFQITWFVSWTPCPDC*VAKLAEFLSEHPN
VTLTISAARLYYYWERDYRRALCRLSQAGARVTIMDYEEFAYCWENFVYNEGQQF
MPWYKFDENYAFLHRTLKEILRYLMDPDTFTFNFNNDPLVLRRRQTYLCYEVERLD

NGTWVLMDQHMGFLCNEAKNLLCGFY*GRHAELRFLDLVPSLQLDPAQIYRVTWFISWS PCFSWGC*AGEVRAFLQENTHVRLRIFAARIYDYDPLYKEALQMLRDAGAQVSIMTY DEFEYCWDTFVYRQGCPFQPWDGLEEHSQALSGRLRAILQNQGN (SEQ ID NO: 277) (italic: nucleic acid editing domain)

[00223] Rat APOBEC-3B:

MQPQGLGPNAGMGPVCLGCSHRRPYSPIRNPLKKLYQQTFYFHFKNVRYAWGRKN NFLCYEVNGMDCALPVPLRQGVFRKQGHIHAELCFIYWFHDKVLRVLSPMEEFKVT WYMSWSPCSKCAEQVARFLAAHRNLSLAIFSSRLYYYLRNPNYQQKLCRLIQEGVH VAAMDLPEFKKCWNKFVDNDGQPFRPWMRLRINFSFYDCKLQEIFSRMNLLREDVF YLQFNNSHRVKPVQNRYYRRKSYLCYQLERANGQEPLKGYLLYKKGEQHVEILFLE KMRSMELSQVRITCYLTWSPCPNCARQLAAFKKDHPDLILRIYTSRLYFYWRKKFQK GLCTLWRSGIHVDVMDLPQFADCWTNFVNPQRPFRPWNELEKNSWRIQRRLRRIKE SWGL (SEQ ID NO: 5729)

[00224] Bovine APOBEC-3B:

DGWEVAFRSGTVLKAGVLGVSMTEGWAGSGHPGQGACVWTPGTRNTMNLLREVL FKQQFGNQPRVPAPYYRRKTYLCYQLKQRNDLTLDRGCFRNKKQRHAEIRFIDKINS LDLNPSQSYKIICYITWSPCPNCANELVNFITRNNHLKLEIFASRLYFHWIKSFKMGLQ DLQNAGISVAVMTHTEFEDCWEQFVDNQSRPFQPWDKLEQYSASIRRRLQRILTAPI (SEQ ID NO: 5730)

[00225] Chimpanzee APOBEC-3B:

MNPQIRNPMEWMYQRTFYYNFENEPILYGRSYTWLCYEVKIRRGHSNLLWDTGVFR GQMYSQPEHHAEMCFLSWFCGNQLSAYKCFQITWFVSWTPCPDCVAKLAKFLAEHP NVTLTISAARLYYYWERDYRRALCRLSQAGARVKIMDDEEFAYCWENFVYNEGQPF MPWYKFDDNYAFLHRTLKEIIRHLMDPDTFTFNFNNDPLVLRRHQTYLCYEVERLD NGTWVLMDQHMGFLCNEAKNLLCGFYGRHAELRFLDLVPSLQLDPAQIYRVTWFIS WSPCFSWGCAGQVRAFLQENTHVRLRIFAARIYDYDPLYKEALQMLRDAGAQVSIM TYDEFEYCWDTFVYRQGCPFQPWDGLEEHSQALSGRLRAILQVRASSLCMVPHRPPP PPQSPGPCLPLCSEPPLGSLLPTGRPAPSLPFLLTASFSFPPPASLPPLPSLSLSPGHLPVP SFHSLTSCSIQPPCSSRIRETEGWASVSKEGRDLG (SEQ ID NO: 5731)

[00226] Human APOBEC-3C:

MNPQIRNPMKAMYPGTFYFQFKNLWEANDRNETWLCFTVEGIKRRSVVSWKTGVF RNQVDSETH*CHAERCFLSWFCDDILSPNTKYQVTWYTSWSPCPDC*AGEVAEFLARHSN VNLTIFTARLYYFQYPCYQEGLRSLSQEGVAVEIMDYEDFKYCWENFVYNDNEPFKP WKGLKTNFRLLKRRLRESLQ (SEQ ID NO: 278)

(italic: nucleic acid editing domain)

[00227] Gorilla APOBEC3C

MNPQIRNPMKAMYPGTFYFQFKNLWEANDRNETWLCFTVEGIKRRSVVSWKTGVF RNQVDSETH*CHAERCFLSWFCDDILSPNTNYQVTWYTSWSPCPEC*AGEVAEFLARHSN VNLTIFTARLYYFQDTDYQEGLRSLSQEGVAVKIMDYKDFKYCWENFVYNDDEPFK PWKGLKYNFRFLKRRLQEILE (SEQ ID NO: 5726)

(italic: nucleic acid editing domain)

[00228] Human APOBEC-3A:

MEASPASGPRHLMDPHIFTSNFNNGIGRHKTYLCYEVERLDNGTSVKMDQHRGFLH NQAKNLLCGFYGR*HAELRFLDLVPSLQLDPAQIYRVTWFISWSPCFSWGC*AGEVRAFLQ ENTHVRLRIFAARIYDYDPLYKEALQMLRDAGAQVSIMTYDEFKHCWDTFVDHQGC PFQPWDGLDEHSQALSGRLRAILQNQGN (SEQ ID NO: 279)

(italic: nucleic acid editing domain)

[00229] Rhesus macaque APOBEC-3A:

MDGSPASRPRHLMDPNTFTFNFNNDLSVRGRHQTYLCYEVERLDNGTWVPMDERR GFLCNKAKNVPCGDYGC*HVELRFLCEVPSWQLDPAQTYRVTWFISWSPC*FRRGCAGQ VRVFLQENKHVRLRIFAARIYDYDPLYQEALRTLRDAGAQVSIMTYEEFKHCWDTF VDRQGRPFQPWDGLDEHSQALSGRLRAILQNQGN (SEQ ID NO: 5727)

(italic: nucleic acid editing domain)

[00230] Bovine APOBEC-3A:

MDEYTFTENFNNQGWPSKTYLCYEMERLDGDATIPLDEYKGFVRNKGLDQPEKPC*H AELYFLGKIHSWNLDRNQHYRLTCFISWSPC*YDCAQKLTTFLKENHHISLHILASRIYTH NRFGCHQSGLCELQAAGARITIMTFEDFKHCWETFVDHKGKPFQPWEGLNVKSQAL CTELQAILKTQQN (SEQ ID NO: 5728)

(italic: nucleic acid editing domain)

[00231] Human APOBEC-3H:

MALLTAETFRLQFNNKRRLRRPYYPRKALLCYQLTPQNGSTPTRGYFENKKKC*HAEI CFINEIKSMGLDETQCYQVTCYLTWSPCSSC*AWELVDFIKAHDHLNLGIFASRLYYHWC KPQQKGLRLLCGSQVPVEVMGFPKFADCWENFVDHEKPLSFNPYKMLEELDKNSRA IKRRLERIKIPGVRAQGRYMDILCDAEV (SEQ ID NO: 280)

(italic: nucleic acid editing domain)

[00232] Rhesus macaque APOBEC-3H:

MALLTAKTFSLQFNNKRRVNKPYYPRKALLCYQLTPQNGSTPTRGHLKNKKKDHAE IRFINKIKSMGLDETQCYQVTCYLTWSPCPSCAGELVDFIKAHRHLNLRIFASRLYYH

WRPNYQEGLLLLCGSQVPVEVMGLPEFTDCWENFVDHKEPPSFNPSEKLEELDKNS QAIKRRLERIKSRSVDVLENGLRSLQLGPVTPSSSIRNSR (SEQ ID NO: 5732)

[00233] Human APOBEC-3D:

MNPQIRNPMERMYRDTFYDNFENEPILYGRSYTWLCYEVKIKRGRSNLLWDTGVFR GPVLPKRQSNHRQEVYFRFEN*HAEMCFLSWFCGNRLPANRRFQITWFVSWNPCLPC*VV KVTKFLAEHPNVTLTISAARLYYYRDRDWRWVLLRLHKAGARVKIMDYEDFAYCW ENFVCNEGQPFMPWYKFDDNYASLHRTLKEILRNPMEAMYPHIFYFHFKNLLKACG RNESWLCFTMEVTKHHSAVFRKRGVFRNQVDPETHC*HAERCFLSWFCDDILSPNTNY EVTWYTSWSPCPEC*AGEVAEFLARHSNVNLTIFTARLCYFWDTDYQEGLCSLSQEGAS VKIMGYKDFVSCWKNFVYSDDEPFKPWKGLQTNFRLLKRRLREILQ (SEQ ID NO: 281)

(italic: nucleic acid editing domain)

[00234] Human APOBEC-1:

MTSEKGPSTGDPTLRRRIEPWEFDVFYDPRELRKEACLLYEIKWGMSRKIWRSSGKN TTNHVEVNFIKKFTSERDFHPSMSCSITWFLSWSPCWECSQAIREFLSRHPGVTLVIYV ARLFWHMDQQNRQGLRDLVNSGVTIQIMRASEYYHCWRNFVNYPPGDEAHWPQYP PLWMMLYALELHCIILSLPPCLKISRRWQNHLTFFRLHLQNCHYQTIPPHILLATGLIH PSVAWR (SEQ ID NO: 282)

[00235] Mouse APOBEC-1:

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSVWRHTSQN TSNHVEVNFLEKFTTERYFRPNTRCSITWFLSWSPCGECSRAITEFLSRHPYVTLFIYIA RLYHHTDQRNRQGLRDLISSGVTIQIMTEQEYCYCWRNFVNYPPSNEAYWPRYPHL WVKLYVLELYCIILGLPPCLKILRRKQPQLTFFTITLQTCHYQRIPPHLLWATGLK (SEQ ID NO: 283)

[00236] Rat APOBEC-1:

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK (SEQ ID NO: 284)

[00237] Human APOBEC-2:

MAQKEEAAVATEAASQNGEDLENLDDPEKLKELIELPPFEIVTGERLPANFFKFQFRN VE

YSSGRNKTFLCYVVEAQGKGGQVQASRGYLEDEHAAAHAEEAFFNTILPAFDPALR YNVTWYVSSSPCAACADRIIKTLSKTKNLRLLILVGRLFMWEEPEIQAALKKLKEAG CKLRIMKPQDFEYVWQNFVEQEEGESKAFQPWEDIQENFLYYEEKLADILK (SEQ ID NO: 5733)

[00238] Mouse APOBEC-2:

MAQKEEAAEAAAPASQNGDDLENLEDPEKLKELIDLPPFEIVTGVRLPVNFFKFQFR NVEYSSGRNKTFLCYVVEVQSKGGQAQATQGYLEDEHAGAHAEEAFFNTILPAFDP ALKYNVTWYVSSSPCAACADRILKTLSKTKNLRLLILVSRLFMWEEPEVQAALKKLK EAGCKLRIMKPQDFEYIWQNFVEQEEGESKAFEPWEDIQENFLYYEEKLADILK (SEQ ID NO: 5734)

[00239] Rat APOBEC-2:

MAQKEEAAEAAAPASQNGDDLENLEDPEKLKELIDLPPFEIVTGVRLPVNFFKFQFR NVEYSSGRNKTFLCYVVEAQSKGGQVQATQGYLEDEHAGAHAEEAFFNTILPAFDP ALKYNVTWYVSSSPCAACADRILKTLSKTKNLRLLILVSRLFMWEEPEVQAALKKLK EAGCKLRIMKPQDFEYLWQNFVEQEEGESKAFEPWEDIQENFLYYEEKLADILK (SEQ ID NO: 5735)

[00240] Bovine APOBEC-2:

MAQKEEAAAAAEPASQNGEEVENLEDPEKLKELIELPPFEIVTGERLPAHYFKFQFRN VE

YSSGRNKTFLCYVVEAQSKGGQVQASRGYLEDEHATNHAEEAFFNSIMPTFDPALR YMVTWYVSSSPCAACADRIVKTLNKTKNLRLLILVGRLFMWEEPEIQAALRKLKEA GCRLRIMKPQDFEYIWQNFVEQEEGESKAFEPWEDIQENFLYYEEKLADILK (SEQ ID NO: 5736)

[00241] Petromyzon marinus CDA1 (pmCDA1)

MTDAEYVRIHEKLDIYTFKKQFFNNKKSVSHRCYVLFELKRRGERRACFWGYAVNK PQSGTERGIHAEIFSIRKVEEYLRDNPGQFTINWYSSWSPCADCAEKILEWYNQELRG NGHTLKIWACKLYYEKNARNQIGLWNLRDNGVGLNVMVSEHYQCCRKIFIQSSHNQ LNENRWLEKTLKRAEKRRSELSIMIQVKILHTTKSPAV (SEQ ID NO: 5738)

[00242] Human APOBEC3G D316R_D317R

MKPHFRNTVERMYRDTFSYNFYNRPILSRRNTVWLCYEVKTKGPSRPPLDAKIFRGQ VYSELKYHPEMRFFHWFSKWRKLHRDQEYEVTWYISWSPCTKCTRDMATFLAEDP KVTLTIFVARLYYFWDPDYQEALRSLCQKRDGPRATMKIMNYDEFQHCWSKFVYSQ RELFEPWNNLPKYYILLHIMLGEILRHSMDPPTFTFNFNNEPWVRGRHETYLCYEVER

MHNDTWVLLNQRRGFLCNQAPHKHGFLEGRHAELCFLDVIPFWKLDLDQDYRVTC FTSWSPCFSCAQEMAKFISKNKHVSLCIFTARIYRRQGRCQEGLRTLAEAGAKISIMT YSEFKHCWDTFVDHQGCPFQPWDGLDEHSQDLSGRLRAILQNQEN (SEQ ID NO: 5739)

[00243] Human APOBEC3G chain A MDPPTFTFNFNNEPWVRGRHETYLCYEVERMHNDTWVLLNQRRGFLCNQAPHKHG FLEGRHAELCFLDVIPFWKLDLDQDYRVTCFTSWSPCFSCAQEMAKFISKNKHVSLCI FTARIYDDQGRCQEGLRTLAEAGAKISIMTYSEFKHCWDTFVDHQGCPFQPWDGLD EHSQDLSGRLRAILQ (SEQ ID NO: 5740)

[00244] Human APOBEC3G chain A D120R_D121R MDPPTFTFNFNNEPWVRGRHETYLCYEVERMHNDTWVLLNQRRGFLCNQAPHKHG FLEGRHAELCFLDVIPFWKLDLDQDYRVTCFTSWSPCFSCAQEMAKFISKNKHVSLCI FTARIYRRQGRCQEGLRTLAEAGAKISIMTYSEFKHCWDTFVDHQGCPFQPWDGLDE HSQDLSGRLRAILQ (SEQ ID NO: 5741)

[00245] In some embodiments, fusion proteins as provided herein comprise the full-length amino acid of a nucleic acid editing enzyme, *e.g.*, one of the sequences provided above. In other embodiments, however, fusion proteins as provided herein do not comprise a full-length sequence of a nucleic acid editing enzyme, but only a fragment thereof. For example, in some embodiments, a fusion protein provided herein comprises a Cas9 domain and a fragment of a nucleic acid editing enzyme, *e.g.*, wherein the fragment comprises a nucleic acid editing domain. Exemplary amino acid sequences of nucleic acid editing domains are shown in the sequences above as italicized letters, and additional suitable sequences of such domains will be apparent to those of skill in the art.

[00246] Additional suitable nucleic-acid editing enzyme sequences, *e.g.*, deaminase enzyme and domain sequences, that can be used according to aspects of this invention, *e.g.*, that can be fused to a nuclease-inactive Cas9 domain, will be apparent to those of skill in the art based on this disclosure. In some embodiments, such additional enzyme sequences include deaminase enzyme or deaminase domain sequences that are at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, or at least 99% similar to the sequences provided herein. Additional suitable Cas9 domains, variants, and sequences will also be apparent to those of skill in the art. Examples of such additional suitable Cas9 domains include, but are not limited to, D10A,

D10A/D839A/H840A, and D10A/D839A/H840A/N863A mutant domains (See, *e.g.*, Prashant *et al.*, CAS9 transcriptional activators for target specificity screening and paired nickases for cooperative genome engineering. Nature Biotechnology. 2013; 31(9): 833-838 the entire contents of which are incorporated herein by reference). In some embodiments, the Cas9 comprises a histidine residue at position 840 of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260. The presence of the catalytic residue H840 restores the acvitity of the Cas9 to cleave the non-edited strand containing a G opposite the targeted C. Restoration of H840 does not result in the cleavage of the target strand containing the C.

[00247] Additional suitable strategies for generating fusion proteins comprising a Cas9 domain and a deaminase domain will be apparent to those of skill in the art based on this disclosure in combination with the general knowledge in the art. Suitable strategies for generating fusion proteins according to aspects of this disclosure using linkers or without the use of linkers will also be apparent to those of skill in the art in view of the instant disclosure and the knowledge in the art. For example, Gilbert *et al.*, CRISPR-mediated modular RNA-guided regulation of transcription in eukaryotes. *Cell.* **2013**; 154(2):442-51, showed that C-terminal fusions of Cas9 with VP64 using 2 NLS's as a linker (SPKKKRKVEAS, SEQ ID NO: 599), can be employed for transcriptional activation. Mali *et al.*, CAS9 transcriptional activators for target specificity screening and paired nickases for cooperative genome engineering. *Nat Biotechnol.* **2013**; 31(9):833-8, reported that C-terminal fusions with VP64 without linker can be employed for transcriptional activation. And Maeder *et al.*, CRISPR RNA-guided activation of endogenous human genes. *Nat Methods.* **2013**; 10: 977-979, reported that C-terminal fusions with VP64 using a Gly_4Ser (SEQ ID NO: 5) linker can be used as transcriptional activators. Recently, dCas9- FokI nuclease fusions have successfully been generated and exhibit improved enzymatic specificity as compared to the parental Cas9 enzyme (In Guilinger JP, Thompson DB, Liu DR. Fusion of catalytically inactive Cas9 to FokI nuclease improves the specificity of genome modification. *Nat. Biotechnol.* 2014; 32(6): 577-82, and in Tsai SQ, Wyvekens N, Khayter C, Foden JA, Thapar V, Reyon D, Goodwin MJ, Aryee MJ, Joung JK. Dimeric CRISPR RNA-guided FokI nucleases for highly specific genome editing. *Nat Biotechnol.* 2014; 32(6):569-76. PMID: 24770325 a SGSETPGTSESATPES (SEQ ID NO: 7) or a GGGGS (SEQ ID NO: 5) linker was used in FokI-dCas9 fusion proteins, respectively).

[00248] Some aspects of this disclosure provide fusion proteins comprising (i) a Cas9 enzyme or domain (e.g., a first protein); and (ii) a nucleic acid-editing enzyme or domain

(e.g., a second protein). In some aspects, the fusion proteins provided herein further include (iii) a programmable DNA-binding protein, for example, a zinc-finger domain, a TALE, or a second Cas9 protein (e.g., a third protein). Without wishing to be bound by any particular theory, fusing a programmable DNA-binding protein (e.g., a second Cas9 protein) to a fusion protein comprising (i) a Cas9 enzyme or domain (e.g., a first protein); and (ii) a nucleic acid-editing enzyme or domain (e.g., a second protein) may be useful for improving specificity of the fusion protein to a target nucleic acid sequence, or for improving specificity or binding affinity of the fusion protein to bind target nucleic acid sequence that does not contain a canonical PAM (NGG) sequence. In some embodiments, the third protein is a Cas9 protein (e.g, a second Cas9 protein). In some embodiments, the third protein is any of the Cas9 proteins provided herein. In some embodiments, the third protein is fused to the fusion protein N-terminal to the Cas9 protein (e.g., the first protein). In some embodiments, the third protein is fused to the fusion protein C-terminal to the Cas9 protein (e.g., the first protein). In some embodiments, the Cas9 domain (e.g., the first protein) and the third protein (e.g., a second Cas9 protein) are fused via a linker (e.g., a second linker). In some embodiments, the linker comprises a (GGGGS)n (SEQ ID NO: 5), a (G)n, an (EAAAK)n (SEQ ID NO: 6), a (GGS)n, $(SGGS)_n$ (SEQ ID NO: 4288), an SGSETPGTSESATPES (SEQ ID NO: 7), or an (XP)n motif, or a combination of any of these, wherein n is independently an integer between 1 and 30. In some embodiments, the general architecture of exemplary Cas9 fusion proteins provided herein comprises the structure:

[NH2]-[nucleic acid-editing enzyme or domain]-[Cas9]-[third protein]-[COOH];

[NH2]-[third protein]-[Cas9]-[nucleic acid-editing enzyme or domain]-[COOH];

[NH2]-[Cas9]-[nucleic acid-editing enzyme or domain]-[third protein]-[COOH];

[NH2]-[third protein]-[nucleic acid-editing enzyme or domain]-[Cas9]-[COOH];

[NH2]-[UGI]-[nucleic acid-editing enzyme or domain]-[Cas9]-[third protein]-[COOH];

[NH2]-[UGI]-[third protein]-[Cas9]-[nucleic acid-editing enzyme or domain]-[COOH];

[NH2]-[UGI]-[Cas9]-[nucleic acid-editing enzyme or domain]-[third protein]-[COOH];

[NH2]-[UGI]-[third protein]-[nucleic acid-editing enzyme or domain]-[Cas9]-[COOH];

[NH2]-[nucleic acid-editing enzyme or domain]-[Cas9]-[third protein]-[UGI]-[COOH];

[NH2]-[third protein]-[Cas9]-[nucleic acid-editing enzyme or domain]-[UGI]-[COOH];

[NH2]-[Cas9]-[nucleic acid-editing enzyme or domain]-[third protein]-[UGI]-[COOH]; or

[NH2]-[third protein]-[nucleic acid-editing enzyme or domain]-[Cas9]-[UGI]-[COOH];

wherein NH2 is the N-terminus of the fusion protein, and COOH is the C-terminus of the fusion protein. In some embodiments, the "]-[" used in the general architecture above indicates the presence of an optional linker sequence. In other examples, the general architecture of exemplary Cas9 fusion proteins provided herein comprises the structure:

[NH2]-[nucleic acid-editing enzyme or domain]-[Cas9]-[second Cas9 protein]-[COOH];

[NH2]-[second Cas9 protein]-[Cas9]-[nucleic acid-editing enzyme or domain]-[COOH];

[NH2]-[Cas9]-[nucleic acid-editing enzyme or domain]-[second Cas9 protein]-[COOH];

[NH2]-[second Cas9 protein]-[nucleic acid-editing enzyme or domain]-[Cas9]-[COOH];

[NH2]-[UGI]-[nucleic acid-editing enzyme or domain]-[Cas9]-[second Cas9 protein]-[COOH],

[NH2]-[UGI]-[second Cas9 protein]-[Cas9]-[nucleic acid-editing enzyme or domain]-[COOH];

[NH2]-[UGI]-[Cas9]-[nucleic acid-editing enzyme or domain]-[second Cas9 protein]-[COOH];

[NH2]-[UGI]-[second Cas9 protein]-[nucleic acid-editing enzyme or domain]-[Cas9]-[COOH];

[NH2]-[nucleic acid-editing enzyme or domain]-[Cas9]-[second Cas9 protein]-[UGI]-[COOH];

[NH2]-[second Cas9 protein]-[Cas9]-[nucleic acid-editing enzyme or domain]-[UGI]-[COOH];

[NH2]-[Cas9]-[nucleic acid-editing enzyme or domain]-[second Cas9 protein]-[UGI]-[COOH]; or

[NH2]-[second Cas9 protein]-[nucleic acid-editing enzyme or domain]-[Cas9]-[UGI]-[COOH];

wherein NH_2 is the N-terminus of the fusion protein, and COOH is the C-terminus of the fusion protein. In some embodiments, the "]-[" used in the general architecture above indicates the presence of an optional linker sequence. In some embodiments, the second Cas9 is a dCas9 protein. In some examples, the general architecture of exemplary Cas9 fusion proteins provided herein comprises a structure as shown in Figure 3. It should be appreciated that any of the proteins provided in any of the general architectures of exemplary Cas9 fusion proteins may be connected by one or more of the linkers provided herein. In some embodiments, the linkers are the same. In some embodiments, the linkers are different. In some embodiments, one or more of the proteins provided in any of the general architectures of exemplary Cas9 fusion proteins are not fused via a linker. In some embodiments, the fusion proteins further comprise a nuclear targeting sequence, for example a nuclear localization sequence. In some embodiments, fusion proteins provided herein further comprise a nuclear localization sequence (NLS). In some embodiments, the NLS is fused to the N-terminus of the fusion protein. In some embodiments, the NLS is fused to the C-terminus of the fusion protein. In some embodiments, the NLS is fused to the N-terminus of the third protein. In some embodiments, the NLS is fused to the C-terminus of the third protein. In some embodiments, the NLS is fused to the N-terminus of the Cas9 protein. In some embodiments, the NLS is fused to the C-terminus of the Cas9 protein. In some embodiments, the NLS is fused to the N-terminus of the nucleic acid-editing enzyme or domain. In some embodiments, the NLS is fused to the C-terminus of the nucleic acid-editing enzyme or domain. In some embodiments, the NLS is fused to the N-terminus of the UGI protein. In some embodiments, the NLS is fused to the C-terminus of the UGI protein. In some embodiments, the NLS is fused to the fusion protein via one or more linkers. In some embodiments, the NLS is fused to the fusioin protein without a linker

Uracil glycosylase inhibitor fusion proteins

[00249] Some aspects of the disclosure relate to fusion proteins that comprise a uracil glycosylase inhibitor (UGI) domain. In some embodiments, any of the fusion proteins provided herein that comprise a Cas9 domain (*e.g.,* a nuclease active Cas9 domain, a nuclease inactive dCas9 domain, or a Cas9 nickase) may be further fused to a UGI domain either directly or via a linker. Some aspects of this disclosure provide deaminase-dCas9 fusion proteins, deaminase-nuclease active Cas9 fusion proteins and deaminase-Cas9 nickase fusion proteins with increased nucleobase editing efficiency. Without wishing to be bound by any particular theory, cellular DNA-repair response to the presence of U:G heteroduplex DNA

may be responsible for the decrease in nucleobase editing efficiency in cells. For example, uracil DNA glycosylase (UDG) catalyzes removal of U from DNA in cells, which may initiate base excision repair, with reversion of the U:G pair to a C:G pair as the most common outcome. As demonstrated in the Examples below, Uracil DNA Glycosylase Inhibitor (UGI) may inhibit human UDG activity. Thus, this disclosure contemplates a fusion protein comprising dCas9-nucleic acid editing domain futher fused to a UGI domain . This disclosure also contemplates a fusion protein comprising a Cas9 nickase-nucleic acid editing domain further fused to a UGI domain. It should be understood that the use of a UGI domain may increase the editing efficiency of a nucleic acid editing domain that is capable of catalyzing a C to U change. For example, fusion proteins comprising a UGI domain may be more efficient in deaminating C residues. In some embodiments, the fusion protein comprises the structure:

[deaminase]-[optional linker sequence]-[dCas9]-[optional linker sequence]-[UGI];
[deaminase]-[optional linker sequence]-[UGI]-[optional linker sequence]-[dCas9];
[UGI]-[optional linker sequence]-[deaminase]-[optional linker sequence]-[dCas9];
[UGI]-[optional linker sequence]-[dCas9]-[optional linker sequence]-[deaminase];
[dCas9]-[optional linker sequence]-[deaminase]-[optional linker sequence]-[UGI]; or
[dCas9]-[optional linker sequence]-[UGI]-[optional linker sequence]-[deaminase].

In other embodiments, the fusion protein comprises the structure:

[deaminase]-[optional linker sequence]-[Cas9 nickase]-[optional linker sequence]-[UGI];

[deaminase]-[optional linker sequence]-[UGI]-[optional linker sequence]-[Cas9 nickase];

[UGI]-[optional linker sequence]-[deaminase]-[optional linker sequence]-[Cas9 nickase];

[UGI]-[optional linker sequence]-[Cas9 nickase]-[optional linker sequence]-[deaminase];

[Cas9 nickase]-[optional linker sequence]-[deaminase]-[optional linker sequence]-[UGI]; or

[Cas9 nickase]-[optional linker sequence]-[UGI]-[optional linker sequence]-[deaminase].

[00250] In some embodiments, the fusion proteins provided herein do not comprise a linker sequence. In some embodiments, one or both of the optional linker sequences are present.

[00251] In some embodiments, the "-" used in the general architecture above indicates the presence of an optional linker sequence. In some embodiments, the fusion proteins comprising a UGI further comprise a nuclear targeting sequence, for example a nuclear localization sequence. In some embodiments, fusion proteins provided herein further comprise a nuclear localization sequence (NLS). In some embodiments, the NLS is fused to the N-terminus of the fusion protein. In some embodiments, the NLS is fused to the C-terminus of the fusion protein. In some embodiments, the NLS is fused to the N-terminus of the UGI protein. In some embodiments, the NLS is fused to the C-terminus of the UGI protein. In some embodiments, the NLS is fused to the N-terminus of the Cas9 protein. In some embodiments, the NLS is fused to the C-terminus of the Cas9 protein. In some embodiments, the NLS is fused to the N-terminus of the deaminase. In some embodiments, the NLS is fused to the C-terminus of the deaminase. In some embodiments, the NLS is fused to the N-terminus of the second Cas9. In some embodiments, the NLS is fused to the C-terminus of the second Cas9. In some embodiments, the NLS is fused to the fusion protein via one or more linkers. In some embodiments, the NLS is fused to the fusioin protein without a linker. In some embodiments, the NLS comprises an amino acid sequence of any one of the NLS sequences provided or referenced herein. In some embodiments, the NLS comprises an amino acid sequence as set forth in SEQ ID NO: 741or SEQ ID NO: 742.

[00252] In some embodiments, a UGI domain comprises a wild-type UGI or a UGI as set forth in SEQ ID NO: 600. In some embodiments, the UGI proteins provided herein include fragments of UGI and proteins homologous to a UGI or a UGI fragment. For example, in some embodiments, a UGI domain comprises a fragment of the amino acid sequence set forth in SEQ ID NO: 600. In some embodiments, a UGI fragment comprises an amino acid sequence that comprises at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% of the amino acid sequence as set forth in SEQ ID NO: 600. In some embodiments, a UGI comprises an amino acid sequence homologous to the amino acid sequence set forth in SEQ ID NO: 600 or an amino acid sequence homologous to a fragment of the amino acid sequence set forth in SEQ ID NO: 600. In some embodiments, proteins comprising UGI or fragments of UGI or homologs of UGI or UGI fragments are referred to as "UGI variants." A UGI variant shares homology to UGI, or a fragment thereof. For example a UGI variant is at least 70% identical, at least 75% identical, at least 80% identical, at least 85% identical, at least 90% identical, at least 95% identical, at least 96% identical, at least 97% identical, at least 98% identical, at least 99% identical, at least 99.5% identical, or

at least 99.9% identical to a wild type UGI or a UGI as set forth in SEQ ID NO: 600. In some embodiments, the UGI variant comprises a fragment of UGI, such that the fragment is at least 70% identical, at least 80% identical, at least 90% identical, at least 95% identical, at least 96% identical, at least 97% identical, at least 98% identical, at least 99% identical, at least 99.5% identical, or at least 99.9% to the corresponding fragment of wild-type UGI or a UGI as set forth in SEQ ID NO: 600. In some embodiments, the UGI comprises the following amino acid sequence:
>sp|P14739|UNGI_BPPB2 Uracil-DNA glycosylase inhibitor
MTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLLT
SDAPEYKPWALVIQDSNGENKIKML (SEQ ID NO: 600)

[00253] Suitable UGI protein and nucleotide sequences are provided herein and additional suitable UGI sequences are known to those in the art, and include, for example, those published in Wang et al., Uracil-DNA glycosylase inhibitor gene of bacteriophage PBS2 encodes a binding protein specific for uracil-DNA glycosylase. J. Biol. Chem. 264:1163-1171(1989); Lundquist et al., Site-directed mutagenesis and characterization of uracil-DNA glycosylase inhibitor protein. Role of specific carboxylic amino acids in complex formation with Escherichia coli uracil-DNA glycosylase. J. Biol. Chem. 272:21408-21419(1997); Ravishankar et al., X-ray analysis of a complex of Escherichia coli uracil DNA glycosylase (EcUDG) with a proteinaceous inhibitor. The structure elucidation of a prokaryotic UDG. Nucleic Acids Res. 26:4880-4887(1998); and Putnam et al., Protein mimicry of DNA from crystal structures of the uracil-DNA glycosylase inhibitor protein and its complex with Escherichia coli uracil-DNA glycosylase. J. Mol. Biol. 287:331-346(1999), the entire contents of each are incorporated herein by reference.

[00254] It should be appreciated that additional proteins may be uracil glycosylase inhibitors. For example, other proteins that are capable of inhibiting (*e.g.*, sterically blocking) a uracil-DNA glycosylase base-excision repair enzyme are within the scope of this disclosure. Additionally, any proteins that block or inhibit base-excision repair as also within the scope of this disclosure. In some embodiments, a protein that binds DNA is used. In another embodiment, a substitute for UGI is used. In some embodiments, a uracil glycosylase inhibitor is a protein that binds single-stranded DNA. For example, a uracil glycosylase inhibitor may be a Erwinia tasmaniensis single-stranded binding protein. In some embodiments, the single-stranded binding protein comprises the amino acid sequence (SEQ ID NO: 322). In some embodiments, a uracil glycosylase inhibitor is a protein that

binds uracil. In some embodiments, a uracil glycosylase inhibitor is a protein that binds uracil in DNA. In some embodiments, a uracil glycosylase inhibitor is a catalytically inactive uracil DNA-glycosylase protein. In some embodiments, a uracil glycosylase inhibitor is a catalytically inactive uracil DNA-glycosylase protein that does not excise uracil from the DNA. For example, a uracil glycosylase inhibitor is a UdgX. In some embodiments, the UdgX comprises the amino acid sequence (SEQ ID NO: 323). As another example, a uracil glycosylase inhibitor is a catalytically inactive UDG. In some embodiments, a catalytically inactive UDG comprises the amino acid sequence (SEQ ID NO: 324). It should be appreciated that other uracil glycosylase inhibitors would be apparent to the skilled artisan and are within the scope of this disclosure. In some embodiments, a uracil glycosylase inhibitor is a protein that is homologous to any one of SEQ ID NOs: 322-324.. In some embodiments, a uracil glycosylase inhibitor is a protein that is at least 50% identical, at least 55% identical at least 60% identical, at least 65% identical, at least 70% identical, at least 75% identical, at least 80% identical at least 85% identical, at least 90% identical, at least 95% identical, at least 96% identical, at least 98% identical, at least 99% identical, or at least 99.5% identical to any one of SEQ ID NOs: 322-324.

Erwinia tasmaniensis SSB (themostable single-stranded DNA binding protein)
MASRGVNKVILVGNLGQDPEVRYMPNGGAVANITLATSESWRDKQTGETKEKTEW
HRVVLFGKLAEVAGEYLRKGSQVYIEGALQTRKWTDQAGVEKYTTEVVVNVGGT
MQMLGGRSQGGGASAGGQNGGSNNGWGQPQQPQGGNQFSGGAQQQARPQQQPQ
QNNAPANNEPPIDFDDDIP (SEQ ID NO: 322)

UdgX (binds to Uracil in DNA but does not excise)
MAGAQDFVPHTADLAELAAAAGECRGCGLYRDATQAVFGAGGRSARIMMIGEQPG
DKEDLAGLPFVGPAGRLLDRALEAADIDRDALYVTNAVKHFKFTRAAGGKRRIHKT
PSRTEVVACRPWLIAEMTSVEPDVVVLLGATAAKALLGNDFRVTQHRGEVLHVDDV
PGDPALVATVHPSSLLRGPKEERESAFAGLVDDLRVAADVRP (SEQ ID NO: 323)

UDG (catalytically inactive human UDG, binds to Uracil in DNA but does not excise)
MIGQKTLYSFFSPSPARKRHAPSPEPAVQGTGVAGVPEESGDAAAIPAKKAPAGQEEP
GTPPSSPLSAEQLDRIQRNKAAALLRLAARNVPVGFGESWKKHLSGEFGKPYFIKLM
GFVAEERKHYTVYPPPHQVFTWTQMCDIKDVKVVILGQEPYHGPNQAHGLCFSVQR
PVPPPPSLENIYKELSTDIEDFVHPGHGDLSGWAKQGVLLLNAVLTVRAHQANSHKE

RGWEQFTDAVVSWLNQNSNGLVFLLWGSYAQKKGSAIDRKRHHVLQTAHPSPLSV YRGFFGCRHFSKTNELLQKSGKKPIDWKEL (SEQ ID NO: 324)

[00255] In some embodiments, the nucleic acid editing domain is a deaminase domain. In some embodiments, the deaminase is a cytosine deaminase or a cytidine deaminase. In some embodiments, the deaminase is an apolipoprotein B mRNA-editing complex (APOBEC) family deaminase. In some embodiments, the deaminase is an APOBEC1 deaminase. In some embodiments, the deaminase is an APOBEC2 deaminase. In some embodiments, the deaminase is an APOBEC3 deaminase. In some embodiments, the deaminase is an APOBEC3A deaminase. In some embodiments, the deaminase is an APOBEC3B deaminase. In some embodiments, the deaminase is an APOBEC3C deaminase. In some embodiments, the deaminase is an APOBEC3D deaminase. In some embodiments, the deaminase is an APOBEC3E deaminase. In some embodiments, the deaminase is an APOBEC3F deaminase. In some embodiments, the deaminase is an APOBEC3G deaminase. In some embodiments, the deaminase is an APOBEC3H deaminase. In some embodiments, the deaminase is an APOBEC4 deaminase. In some embodiments, the deaminase is an activation-induced deaminase (AID). In some embodiments, the demianse is a rat APOBEC1 (SEQ ID NO: 282). In some embodiments, the deminase is a human APOBEC1 (SEQ ID No: 284). In some embodiments, the deaminase is a *Petromyzon marinus* cytidine deaminase 1 (pmCDA1). In some embodiments, the deminase is a human APOBEC3G (SEQ ID NO: 275). In some embodiments, the deaminase is a fragment of the human APOBEC3G (SEQ ID NO: 5740). In some embodiments, the deaminase is a human APOBEC3G variant comprising a D316R_D317R mutation (SEQ ID NO: 5739). In some embodiments, the deaminase is a frantment of the human APOBEC3G and comprising mutations corresponding to the D316R_D317R mutations in SEQ ID NO: 275 (SEQ ID NO: 5741).

[00256] In some embodiments, the linker comprises a $(GGGS)_n$ (SEQ ID NO: 265), $(GGGGS)_n$ (SEQ ID NO: 5), a $(G)_n$, an $(EAAAK)_n$ (SEQ ID NO: 6), a $(GGS)_n$, an SGSETPGTSESATPES (SEQ ID NO: 7), or an $(XP)_n$ motif, or a combination of any of these, wherein n is independently an integer between 1 and 30.

[00257] Suitable UGI protein and nucleotide sequences are provided herein and additional suitable UGI sequences are known to those in the art, and include, for example, those published in Wang *et al.*, Uracil-DNA glycosylase inhibitor gene of bacteriophage PBS2 encodes a binding protein specific for uracil-DNA glycosylase. *J. Biol. Chem.* 264:1163-

1171(1989); Lundquist *et al.*, Site-directed mutagenesis and characterization of uracil-DNA glycosylase inhibitor protein. Role of specific carboxylic amino acids in complex formation with Escherichia coli uracil-DNA glycosylase. *J. Biol. Chem.* 272:21408-21419(1997); Ravishankar *et al.*, X-ray analysis of a complex of Escherichia coli uracil DNA glycosylase (EcUDG) with a proteinaceous inhibitor. The structure elucidation of a prokaryotic UDG. *Nucleic Acids Res.* 26:4880-4887(1998); and Putnam *et al.*, Protein mimicry of DNA from crystal structures of the uracil-DNA glycosylase inhibitor protein and its complex with Escherichia coli uracil-DNA glycosylase. *J. Mol. Biol.* 287:331-346(1999), the entire contents of which are incorporated herein by reference. In some embodiments, the optional linker comprises a $(GGS)_n$ motif, wherein n is 1, 2, 3, 4, 5, 6, 7, 8, 9, 19, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20. In some embodiments, the optional linker comprises a (GGS)n motif, wherein n is 1, 3, or 7. In some embodiments, the optional linker comprises the amino acid sequence SGSETPGTSESATPES (SEQ ID NO: 7), which is also referred to as the XTEN linker in the Examples.

[00258] In some embodiments, a Cas9 nickase may further facilitate the removal of a base on the non-edited strand in an organism whose genome is edited *in vivo*. The Cas9 nickase, as described herein, may comprise a D10A mutation in SEQ ID NO: 10, or a corresponding mutation in any of SEQ ID NOs: 11-260. In some embodiments, the Cas9 nickase of this disclosure may comprise a histidine at mutation 840 of SEQ ID NO: 10, or a corresponding residue in any of SEQ ID NOs: 11-260. Such fusion proteins comprising the Cas9 nickase, can cleave a single strand of the target DNA sequence, *e.g.*, the strand that is not being edited. Without wishing to be bound by any particular theory, this cleavage may inhibit mis-match repair mechanisms that reverse a C to U edit made by the deaminase.

Cas9 complexes with guide RNAs

[00259] Some aspects of this disclosure provide complexes comprising any of the fusion proteins provided herein, and a guide RNA bound to a Cas9 domain (*e.g.*, a dCas9, a nuclease active Cas9, or a Cas9 nickase) of fusion protein.

[00260] In some embodiments, the guide RNA is from 15-100 nucleotides long and comprises a sequence of at least 10 contiguous nucleotides that is complementary to a target sequence. In some embodiments, the guide RNA is 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, or 50 nucleotides long. In some embodiments, the guide RNA comprises a sequence of 15, 16,

17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, or 40 contiguous nucleotides that is complementary to a target sequence. In some embodiments, the target sequence is a DNA sequence. In some embodiments, the target sequence is a sequence in the genome of a mammal. In some embodiments, the target sequence is a sequence in the genome of a human. In some embodiments, the 3′ end of the target sequence is immediately adjacent to a canonical PAM sequence (NGG). In some embodiments, the guide RNA is complementary to a sequence associated with a disease or disorder. In some embodiments, the guide RNA is complementary to a sequence associated with a disease or disorder having a mutation in a gene selected from the genes disclosed in any one of Tables 1-3. In some embodiments, the guide RNA comprises a nucleotide sequence of any one of the guide sequences provided in Table 2 or Table 3. Exemplary sequences in the human genome that may be targeted by the complexes of this disclosure are provided herein in Tables 1-3.

Methods of using Cas9 fusion proteins

[00261] Some aspects of this disclosure provide methods of using the Cas9 proteins, fusion proteins, or complexes provided herein. For example, some aspects of this disclosure provide methods comprising contacting a DNA molecule (a) with any of the the Cas9 proteins or fusion proteins provided herein, and with at least one guide RNA, wherein the guide RNA is about 15-100 nucleotides long and comprises a sequence of at least 10 contiguous nucleotides that is complementary to a target sequence; or (b) with a Cas9 protein, a Cas9 fusion protein, or a Cas9 protein or fusion protein complex with at least one gRNA as provided herein. In some embodiments, the 3' end of the target sequence is not immediately adjacent to a canonical PAM sequence (NGG). In some embodiments, the 3' end of the target sequence is immediately adjacent to an AGC, GAG, TTT, GTG, or CAA sequence.

[00262] In some embodiments, the target DNA sequence comprises a sequence associated with a disease or disorder. In some embodiments, the target DNA sequence comprises a point mutation associated with a disease or disorder. In some embodiments, the activity of the Cas9 protein, the Cas9 fusion protein, or the complex results in a correction of the point mutation. In some embodiments, the target DNA sequence comprises a T→C point mutation associated with a disease or disorder, and wherein the deamination of the mutant C base results in a sequence that is not associated with a disease or disorder. In some embodiments, the target DNA sequence encodes a protein and wherein the point mutation is in a codon and results in a change in the amino acid encoded by the mutant codon as compared to the wild-type codon. In some embodiments, the deamination of the mutant C results in a change of the amino acid

encoded by the mutant codon. In some embodiments, the deamination of the mutant C results in the codon encoding the wild-type amino acid. In some embodiments, the contacting is *in vivo* in a subject. In some embodiments, the subject has or has been diagnosed with a disease or disorder. In some embodiments, the disease or disorder is cystic fibrosis, phenylketonuria, epidermolytic hyperkeratosis (EHK), Charcot-Marie-Toot disease type 4J, neuroblastoma (NB), von Willebrand disease (vWD), myotonia congenital, hereditary renal amyloidosis, dilated cardiomyopathy (DCM), hereditary lymphedema, familial Alzheimer's disease, HIV, Prion disease, chronic infantile neurologic cutaneous articular syndrome (CINCA), desmin-related myopathy (DRM), a neoplastic disease associated with a mutant PI3KCA protein, a mutant CTNNB1 protein, a mutant HRAS protein, or a mutant p53 protein.

[00263] Some embodiments provide methods for using the Cas9 DNA editing fusion proteins provided herein. In some embodiments, the fusion protein is used to introduce a point mutation into a nucleic acid by deaminating a target nucleobase, *e.g.*, a C residue. In some embodiments, the deamination of the target nucleobase results in the correction of a genetic defect, *e.g.*, in the correction of a point mutation that leads to a loss of function in a gene product. In some embodiments, the genetic defect is associated with a disease or disorder, *e.g.*, a lysosomal storage disorder or a metabolic disease, such as, for example, type I diabetes. In some embodiments, the methods provided herein are used to introduce a deactivating point mutation into a gene or allele that encodes a gene product that is associated with a disease or disorder. For example, in some embodiments, methods are provided herein that employ a Cas9 DNA editing fusion protein to introduce a deactivating point mutation into an oncogene (*e.g.*, in the treatment of a proliferative disease). A deactivating mutation may, in some embodiments, generate a premature stop codon in a coding sequence, which results in the expression of a truncated gene product, *e.g.*, a truncated protein lacking the function of the full-length protein.

[00264] In some embodiments, the purpose of the methods provide herein is to restore the function of a dysfunctional gene via genome editing. The Cas9 deaminase fusion proteins provided herein can be validated for gene editing-based human therapeutics *in vitro*, *e.g.*, by correcting a disease-associated mutation in human cell culture. It will be understood by the skilled artisan that the fusion proteins provided herein, *e.g.*, the fusion proteins comprising a Cas9 domain and a nucleic acid deaminase domain can be used to correct any single point T -> C or A -> G mutation. In the first case, deamination of the mutant C back to U corrects the mutation, and in the latter case, deamination of the C that is base-paired with the mutant G, followed by a round of replication, corrects the mutation.

[00265] An exemplary disease-relevant mutation that can be corrected by the provided fusion proteins *in vitro* or *in vivo* is the H1047R (A3140G) polymorphism in the PI3KCA protein. The phosphoinositide-3-kinase, catalytic alpha subunit (PI3KCA) protein acts to phosphorylate the 3-OH group of the inositol ring of phosphatidylinositol. The PI3KCA gene has been found to be mutated in many different carcinomas, and thus it is considered to be a potent oncogene.[37] In fact, the A3140G mutation is present in several NCI-60 cancer cell lines, such as, for example, the HCT116, SKOV3, and T47D cell lines, which are readily available from the American Type Culture Collection (ATCC).[38]

[00266] In some embodiments, a cell carrying a mutation to be corrected, *e.g.*, a cell carrying a point mutation, *e.g.*, an A3140G point mutation in exon 20 of the PI3KCA gene, resulting in a H1047R substitution in the PI3KCA protein, is contacted with an expression construct encoding a Cas9 deaminase fusion protein and an appropriately designed sgRNA targeting the fusion protein to the respective mutation site in the encoding PI3KCA gene. Control experiments can be performed where the sgRNAs are designed to target the fusion enzymes to non-C residues that are within the PI3KCA gene. Genomic DNA of the treated cells can be extracted, and the relevant sequence of the PI3KCA genes PCR amplified and sequenced to assess the activities of the fusion proteins in human cell culture.

[00267] It will be understood that the example of correcting point mutations in PI3KCA is provided for illustration purposes and is not meant to limit the instant disclosure. The skilled artisan will understand that the instantly disclosed DNA-editing fusion proteins can be used to correct other point mutations and mutations associated with other cancers and with diseases other than cancer including other proliferative diseases.

[00268] The successful correction of point mutations in disease-associated genes and alleles opens up new strategies for gene correction with applications in therapeutics and basic research. Site-specific single-base modification systems like the disclosed fusions of Cas9 and deaminase enzymes or domains also have applications in "reverse" gene therapy, where certain gene functions are purposely suppressed or abolished. In these cases, site-specifically mutating Trp (TGG), Gln (CAA and CAG), or Arg (CGA) residues to premature stop codons (TAA, TAG, TGA) can be used to abolish protein function *in vitro*, *ex vivo*, or *in vivo*.

[00269] The instant disclosure provides methods for the treatment of a subject diagnosed with a disease associated with or caused by a point mutation that can be corrected by a Cas9 DNA editing fusion protein provided herein. For example, in some embodiments, a method is provided that comprises administering to a subject having such a disease, *e.g.*, a cancer associated with a PI3KCA point mutation as described above, an effective amount of a Cas9

deaminase fusion protein that corrects the point mutation or introduces a deactivating mutation into the disease-associated gene. In some embodiments, the disease is a proliferative disease. In some embodiments, the disease is a genetic disease. In some embodiments, the disease is a neoplastic disease. In some embodiments, the disease is a metabolic disease. In some embodiments, the disease is a lysosomal storage disease. Other diseases that can be treated by correcting a point mutation or introducing a deactivating mutation into a disease-associated gene will be known to those of skill in the art, and the disclosure is not limited in this respect.

[00270] The instant disclosure provides methods for the treatment of additional diseases or disorders, *e.g.*, diseases or disorders that are associated or caused by a point mutation that can be corrected by deaminase-mediated gene editing. Some such diseases are described herein, and additional suitable diseases that can be treated with the strategies and fusion proteins provided herein will be apparent to those of skill in the art based on the instant disclosure. Exemplary suitable diseases and disorders are listed below. It will be understood that the numbering of the specific positions or residues in the respective sequences depends on the particular protein and numbering scheme used. Numbering might be different, *e.g.*, in precursors of a mature protein and the mature protein itself, and differences in sequences from species to species may affect numbering. One of skill in the art will be able to identify the respective residue in any homologous protein and in the respective encoding nucleic acid by methods well known in the art, *e.g.*, by sequence alignment and determination of homologous residues. Exemplary suitable diseases and disorders include, without limitation, cystic fibrosis (see, *e.g.*, Schwank *et al.*, Functional repair of CFTR by CRISPR/Cas9 in intestinal stem cell organoids of cystic fibrosis patients. *Cell stem cell.* **2013**; 13: 653-658; and Wu et. al., Correction of a genetic disease in mouse via use of CRISPR-Cas9. *Cell stem cell.* **2013;** 13: 659-662, neither of which uses a deaminase fusion protein to correct the genetic defect); phenylketonuria – *e.g.*, phenylalanine to serine mutation at position 835 (mouse) or 240 (human) or a homologous residue in phenylalanine hydroxylase gene (T>C mutation) – see, *e.g.*, McDonald *et al.*, *Genomics.* **1997;** 39:402-405; Bernard-Soulier syndrome (BSS) – *e.g.*, phenylalanine to serine mutation at position 55 or a homologous residue, or cysteine to arginine at residue 24 or a homologous residue in the platelet membrane glycoprotein IX (T>C mutation) – see, *e.g.*, Noris *et al.*, *British Journal of Haematology.* **1997;** 97: 312-320, and Ali *et al.*, *Hematol.* **2014;** 93: 381-384; epidermolytic hyperkeratosis (EHK) – *e.g.*, leucine to proline mutation at position 160 or 161 (if counting the initiator methionine) or a homologous residue in keratin 1 (T>C mutation) – see, *e.g.*,

Chipev *et al.*, *Cell.* **1992;** 70: 821-828, see also accession number P04264 in the UNIPROT database at www[dot]uniprot[dot]org; chronic obstructive pulmonary disease (COPD) – *e.g.*, leucine to proline mutation at position 54 or 55 (if counting the initiator methionine) or a homologous residue in the processed form of α_1-antitrypsin or residue 78 in the unprocessed form or a homologous residue (T>C mutation) – see, *e.g.*, Poller *et al.*, *Genomics.* **1993;** 17: 740-743, see also accession number P01011 in the UNIPROT database; Charcot-Marie-Toot disease type 4J – *e.g.*, isoleucine to threonine mutation at position 41 or a homologous residue in FIG4 (T>C mutation) – see, *e.g.*, Lenk *et al.*, PLoS Genetics. 2011; 7: e1002104; neuroblastoma (NB) – *e.g.*, leucine to proline mutation at position 197 or a homologous residue in Caspase-9 (T>C mutation) – see, *e.g.*, Kundu *et al.*, *3 Biotech.* **2013, 3:225-234**; von Willebrand disease (vWD) – *e.g.*, cysteine to arginine mutation at position 509 or a homologous residue in the processed form of von Willebrand factor, or at position 1272 or a homologous residue in the unprocessed form of von Willebrand factor (T>C mutation) – see, *e.g.*, Lavergne *et al.*, *Br. J. Haematol.* **1992**, see also accession number P04275 in the UNIPROT database**;** 82: 66-72; myotonia congenital – *e.g.*, cysteine to arginine mutation at position 277 or a homologous residue in the muscle chloride channel gene CLCN1 (T>C mutation) – see, *e.g.*, Weinberger *et al.*, *The J. of Physiology.* **2012;** 590: 3449-3464; hereditary renal amyloidosis – *e.g.*, stop codon to arginine mutation at position 78 or a homologous residue in the processed form of apolipoprotein AII or at position 101 or a homologous residue in the unprocessed form (T>C mutation) – see, *e.g.*, Yazaki *et al.*, *Kidney Int.* **2003;** 64: 11-16; dilated cardiomyopathy (DCM) – *e.g.*, tryptophan to Arginine mutation at position 148 or a homologous residue in the FOXD4 gene (T>C mutation), see, *e.g.*, Minoretti et. al., *Int. J. of Mol. Med.* **2007;** 19: 369-372; hereditary lymphedema – *e.g.*, histidine to arginine mutation at position 1035 or a homologous residue in VEGFR3 tyrosine kinase (A>G mutation), see, *e.g.*, Irrthum *et al.*, *Am. J. Hum. Genet.* **2000;** 67: 295-301; familial Alzheimer's disease – *e.g.*, isoleucine to valine mutation at position 143 or a homologous residue in presenilin1 (A>G mutation), see, *e.g.*, Gallo et. al., *J. Alzheimer's disease.* **2011;** 25: 425-431; Prion disease – *e.g.*, methionine to valine mutation at position 129 or a homologous residue in prion protein (A>G mutation) – see, *e.g.*, Lewis et. al., *J. of General Virology.* **2006;** 87: 2443-2449; chronic infantile neurologic cutaneous articular syndrome (CINCA) – *e.g.*, Tyrosine to Cysteine mutation at position 570 or a homologous residue in cryopyrin (A>G mutation) – see, *e.g.*, Fujisawa et. al. *Blood.* **2007;** 109: 2903-2911; and desmin-related myopathy (DRM) – *e.g.*, arginine to glycine mutation at position 120 or a homologous residue in αβ crystallin (A>G mutation) – see, *e.g.*, Kumar *et al.*, *J.*

Biol. Chem. **1999;** 274: 24137-24141. The entire contents of all references and database entries is incorporated herein by reference.

[00271] The instant disclosure provides lists of genes comprising pathogenic T>C or A>G mutations. Provided herein, are the names of these genes, their respective SEQ ID NOs, their gene IDs, and sequences flanking the mutation site. (Tables 2 and 3). In some instances, the gRNA sequences that can be used to correct the mutations in these genes are disclosed (Tables 2 and 3).

[00272] In some embodiments, a Cas9-deaminase fusion protein recognizes canonical PAMs and therefore can correct the pathogenic T>C or A>G mutations with canonical PAMs, e.g., NGG (listed in Tables 2 and 3, SEQ ID NOs: 2540-2702 and 5084-5260), respectively, in the flanking sequences. For example, the Cas9 proteins that recognize canonical PAMs comprise an amino acid sequence that is at least 90% identical to the amino acid sequence of *Streptococcus pyogenes* Cas9 as provided by SEQ ID NO: 10, or to a fragment thereof comprising the RuvC and HNH domains of SEQ ID NO: 10.

[00273] It will be apparent to those of skill in the art that in order to target a Cas9:nucleic acid editing enzyme/domain fusion protein as disclosed herein to a target site, *e.g.*, a site comprising a point mutation to be edited, it is typically necessary to co-express the Cas9:nucleic acid editing enzyme/domain fusion protein together with a guide RNA, *e.g.*, an sgRNA. As explained in more detail elsewhere herein, a guide RNA typically comprises a tracrRNA framework allowing for Cas9 binding, and a guide sequence, which confers sequence specificity to the Cas9:nucleic acid editing enzyme/domain fusion protein. In some embodiments, the guide RNA comprises a structure 5'-[guide sequence]-guuuuagagcuagaaauagcaaguuaaaauaaaggcuaguccguuaucaacuugaaaaaguggcaccgagucggugcuuuuu-3' (SEQ ID NO: 601), wherein the guide sequence comprises a sequence that is complementary to the target sequence. The guide sequence is typically 20 nucleotides long. The sequences of suitable guide RNAs for targeting Cas9:nucleic acid editing enzyme/domain fusion proteins to specific genomic target sites will be apparent to those of skill in the art based on the instant disclosure. Such suitable guide RNA sequences typically comprise guide sequences that are complementary to a nucleic sequence within 50 nucleotides upstream or downstream of the target nucleotide to be edited. Some exemplary guide RNA sequences suitable for targeting Cas9:nucleic acid editing enzyme/domain fusion proteins to specific target sequences are provided below.

Base Editor Efficiency

[00274] Some aspects of the disclosure are based on the recognition that any of the base editors provided herein are capable of modifying a specific nucleotide base without generating a significant proportion of indels. An "indel", as used herein, refers to the insertion or deletion of a nucleotide base within a nucleic acid. Such insertions or deletions can lead to frame shift mutations within a coding region of a gene. In some embodiments, it is desirable to generate base editors that efficiently modify (*e.g.* mutate or deaminate) a specific nucleotide within a nucleic acid, without generating a large number of insertions or deletions (*i.e.*, indels) in the nucleic acid. In certain embodiments, any of the base editors provided herein are capable of generating a greater proportion of intended modifications (*e.g.*, point mutations or deaminations) versus indels. In some embodiments, the base editors provided herein are capable of generating a ratio of intended point mutations to indels that is greater than 1:1. In some embodiments, the base editors provided herein are capable of generating a ratio of intended point mutations to indels that is at least 1.5:1, at least 2:1, at least 2.5:1, at least 3:1, at least 3.5:1, at least 4:1, at least 4.5:1, at least 5:1, at least 5.5:1, at least 6:1, at least 6.5:1, at least 7:1, at least 7.5:1, at least 8:1, at least 10:1, at least 12:1, at least 15:1, at least 20:1, at least 25:1, at least 30:1, at least 40:1, at least 50:1, at least 100:1, at least 200:1, at least 300:1, at least 400:1, at least 500:1, at least 600:1, at least 700:1, at least 800:1, at least 900:1, or at least 1000:1, or more. The number of intended mutations and indels may be determined using any suitable method, for example the methods used in the below Examples.

[00275] In some embodiments, the base editors provided herein are capable of limiting formation of indels in a region of a nucleic acid. In some embodiments, the region is at a nucleotide targeted by a base editor or a region within 2, 3, 4, 5, 6, 7, 8, 9, or 10 nucleotides of a nucleotide targeted by a base editor. In some embodiments, any of the base editors provided herein are capable of limiting the formation of indels at a region of a nucleic acid to less than 1%, less than 1.5%, less than 2%, less than 2.5%, less than 3%, less than 3.5%, less than 4%, less than 4.5%, less than 5%, less than 6%, less than 7%, less than 8%, less than 9%, less than 10%, less than 12%, less than 15%, or less than 20%. The number of indels formed at a nucleic acid region may depend on the amount of time a nucleic acid (*e.g.*, a nucleic acid within the genome of a cell) is exposed to a base editor. In some embodiments, an number or proportion of indels is determined after at least 1 hour, at least 2 hours, at least 6 hours, at least 12 hours, at least 24 hours, at least 36 hours, at least 48 hours, at least 3 days, at least 4 days, at least 5 days, at least 7 days, at least 10 days, or at least 14 days of exposing a nucleic acid (*e.g.*, a nucleic acid within the genome of a cell) to a base editor.

[00276] Some aspects of the disclosure are based on the recognition that any of the base editors provided herein are capable of efficiently generating an intended mutation, such as a point mutation, in a nucleic acid (*e.g.* a nucleic acid within a genome of a subject) without generating a significant number of unintended mutations, such as unintended point mutations. In some embodiments, a intended mutation is a mutation that is generated by a specific base editor bound to a gRNA, specifically designed to generate the intended mutation. In some embodiments, the intended mutation is a mutation associated with a disease or disorder. In some embodiments, the intended mutation is a cytosine (C) to thymine (T) point mutation associated with a disease or disorder. In some embodiments, the intended mutation is a guanine (G) to adenine (A) point mutation associated with a disease or disorder. In some embodiments, the intended mutation is a cytosine (C) to thymine (T) point mutation within the coding region of a gene. In some embodiments, the intended mutation is a guanine (G) to adenine (A) point mutation within the coding region of a gene. In some embodiments, the intended mutation is a point mutation that generates a stop codon, for example, a premature stop codon within the coding region of a gene. In some embodiments, the intended mutation is a mutation that eliminates a stop codon. In some embodiments, the intended mutation is a mutation that alters the splicing of a gene. In some embodiments, the intended mutation is a mutation that alters the regulatory sequence of a gene (*e.g.*, a gene promotor or gene repressor). In some embodiments, any of the base editors provided herein are capable of generating a ratio of intended mutations to unintended mutations (*e.g.,* intended point mutations:unintended point mutations) that is greater than 1:1. In some embodiments, any of the base editors provided herein are capable of generating a ratio of intended mutations to unintended mutations (*e.g.,* intended point mutations:unintended point mutations) that is at least 1.5:1, at least 2:1, at least 2.5:1, at least 3:1, at least 3.5:1, at least 4:1, at least 4.5:1, at least 5:1, at least 5.5:1, at least 6:1, at least 6.5:1, at least 7:1, at least 7.5:1, at least 8:1, at least 10:1, at least 12:1, at least 15:1, at least 20:1, at least 25:1, at least 30:1, at least 40:1, at least 50:1, at least 100:1, at least 150:1, at least 200:1, at least 250:1, at least 500:1, or at least 1000:1, or more. It should be appreciated that the characterstics of the base editors described in the "*Base Editor Efficiency*" section, herein, may be applied to any of the fusion proteins, or methods of using the fusion proteins provided herein.

Methods for Editing Nucleic Acids

[00277] Some aspects of the disclosure provide methods for editing a nucleic acid. In some embodiments, the method is a method for editing a nucleobase of a nucleic acid (*e.g.*, a

base pair of a double-stranded DNA sequence). In some embodiments, the method comprises the steps of: a) contacting a target region of a nucleic acid (*e.g.*, a double-stranded DNA sequence) with a complex comprising a base editor (*e.g.*, a Cas9 domain fused to a cytidine deaminase domain) and a guide nucleic acid (*e.g.*, gRNA), wherein the target region comprises a targeted nucleobase pair, b) inducing strand separation of said target region, c)converting a first nucleobase of said target nucleobase pair in a single strand of the target region to a second nucleobase, and d) cutting no more than one strand of said target region, where a third nucleobase complementary to the first nucleobase base is replaced by a fourth nucleobase complementary to the second nucleobase; and the method results in less than 20% indel formation in the nucleic acid. It should be appreciated that in some embodiments, step b is omitted. In some embodiments, the first nucleobase is a cytosine. In some embodiments, the second nucleobase is a deaminated cytosine, or a uracil. In some embodiments, the third nucleobase is a guanine. In some embodiments, the fourth nucleobase is an adenine. In some embodiments, the first nucleobase is a cytosine, the second nucleobase is a deaminated cytosine, or a uracil, the third nucleobase is a guanine, and the fourth nucleobase is an adenine.In some embodiments, the method results in less than 19%, 18%, 16%, 14%, 12%, 10%, 8%, 6%, 4%, 2%, 1%, 0.5%, 0.2%, or less than 0.1% indel formation. In some embodiments, the method further comprises replacing the second nucleobase with a fifth nucleobase that is complementary to the fourth nucleobase, thereby generating an intended edited base pair (*e.g.*, C:G -> T:A). In some embodiments, the fifth nucleobase is a thymine. In some embodiments, at least 5% of the intended basepaires are edited. In some embodiments, at least 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, or 50% of the intended basepaires are edited.

[00278] In some embodiments, the ratio of intended products to unintended products in the target nucleotide is at least 2:1, 5:1, 10:1, 20:1, 30:1, 40:1, 50:1, 60:1, 70:1, 80:1, 90:1, 100:1, or 200:1, or more. In some embodiments, the ratio of intended point mutation to indel formation is greater than 1:1, 10:1, 50:1, 100:1, 500:1, or 1000:1, or more. In some embodiments, the cut single strand (nicked strand) is hybridized to the guide nucleic acid. In some embodiments, the cut single strand is opposite to the strand comprising the first nucleobase. In some embodiments, the base editor comprises a Cas9 domain. In some embodiments, the first base is cytosine, and the second base is not a G, C, A, or T. In some embodiments, the second base is uracil. In some embodiments, the first base is cytosine. In some embodiments, the second base is not a G, C, A, or T. In some embodiments, the second base is uracil. In some embodiments, the base editor inhibits base escision repair of the

edited strand. In some embodiments, the base editor protects or binds the non-edited strand. In some embodiments, the base editor comprises UGI activity. In some embodiments, the base editor comprises nickase activity. In some embodiments, the intended edited basepair is upstream of a PAM site. In some embodiments, the intended edited base pair is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides upstream of the PAM site. In some embodiments, the intended edited basepair is downstream of a PAM site. In some embodiments, the intended edited base pair is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides downstream stream of the PAM site. In some embodiments, the method does not require a canonical (*e.g.,* NGG) PAM site. In some embodiments, the nucleobase editor comprises a linker. In some embodiments, the linker is 1-25 amino acids in length. In some embodiments, the linker is 5-20 amino acids in length. In some embodiments, linker is 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 amino acids in length. In some embodiments, the target region comprises a target window, wherein the target window comprises the target nucleobase pair. In some embodiments, the target window comprises 1-10 nucleotides. In some embodiments, the target window is 1-9, 1-8, 1-7, 1-6, 1-5, 1-4, 1-3, 1-2, or 1 nucleotides in length. In some embodiments, the target window is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides in length. In some embodiments, the intended edited base pair is within the target window. In some embodiments, the target window comprises the intended edited base pair. In some embodiments, the method is performed using any of the base editors provided herein. In some embodiments, a target windo is a deamination window

[00279] In some embodiments, the disclosure provides methods for editing a nucleotide. In some embodiments, the disclosure provides a method for editing a nucleobase pair of a double-stranded DNA sequence. In some embodiments, the method comprises a) contacting a target region of the double-stranded DNA sequence with a complex comprising a base editor and a guide nucleic acid (*e.g.*, gRNA), where the target region comprises a target nucleobase pair, b) inducing strand separation of said target region, c) converting a first nucleobase of said target nucleobase pair in a single strand of the target region to a second nucleobase, d) cutting no more than one strand of said target region, wherein a third nucleobase complementary to the first nucleobase base is replaced by a fourth nucleobase complementary to the second nucleobase, and the second nucleobase is replaced with a fifth nucleobase that is complementary to the fourth nucleobase, thereby generating an intended edited basepair, wherein the efficiency of generating the intended edited basepair is at least 5%. It should be appreciated that in some embodiments, step b is omitted. In some

embodiments, at least 5% of the intended basepaires are edited. In some embodiments, at least 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, or 50% of the intended basepaires are edited. In some embodiments, the method causes less than 19%, 18%, 16%, 14%, 12%, 10%, 8%, 6%, 4%, 2%, 1%, 0.5%, 0.2%, or less than 0.1% indel formation. In some embodiments, the ratio of intended product to unintended products at the target nucleotide is at least 2:1, 5:1, 10:1, 20:1, 30:1, 40:1, 50:1, 60:1, 70:1, 80:1, 90:1, 100:1, or 200:1, or more. In some embodiments, the ratio of intended point mutation to indel formation is greater than 1:1, 10:1, 50:1, 100:1, 500:1, or 1000:1, or more. In some embodiments, the cut single strand is hybridized to the guide nucleic acid. In some embodiments, the cut single strand is opposite to the strand comprising the first nucleobase. In some embodiments, the first base is cytosine. In some embodiments, the second nucleobase is not G, C, A, or T. In some embodiments, the second base is uracil. In some embodiments, the base editor inhibits base escision repair of the edited strand. In some embodiments, the base editor protects or binds the non-edited strand. In some embodiments, the nucleobase editor comprises UGI activity. In some embodiments, the nucleobase edit comprises nickase activity. In some embodiments, the intended edited basepair is upstream of a PAM site. In some embodiments, the intended edited base pair is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides upstream of the PAM site. In some embodiments, the intended edited basepair is downstream of a PAM site. In some embodiments, the intended edited base pair is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides downstream stream of the PAM site. In some embodiments, the method does not require a canonical (e.g., NGG) PAM site. In some embodiments, the nucleobase editor comprises a linker. In some embodiments, the linker is 1-25 amino acids in length. In some embodiments, the linker is 5-20 amino acids in length. In some embodiments, the linker is 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 amino acids in length. In some embodiments, the target region comprises a target window, wherein the target window comprises the target nucleobase pair. In some embodiments, the target window comprises 1-10 nucleotides. In some embodiments, the target window is 1-9, 1-8, 1-7, 1-6, 1-5, 1-4, 1-3, 1-2, or 1 nucleotides in length. In some embodiments, the target window is 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides in length. In some embodiments, the intended edited base pair occurs within the target window. In some embodiments, the target window comprises the intended edited base pair. In some embodiments, the nucleobase editor is any one of the base editors provided herein.

[00280]

Kits, vectors, cells

[00281] Some aspects of this disclosure provide kits comprising a nucleic acid construct, comprising (a) a nucleotide sequence encoding a Cas9 protein or a Cas9 fusion protein as provided herein; and (b) a heterologous promoter that drives expression of the sequence of (a). In some embodiments, the kit further comprises an expression construct encoding a guide RNA backbone, wherein the construct comprises a cloning site positioned to allow the cloning of a nucleic acid sequence identical or complementary to a target sequence into the guide RNA backbone.

[00282] Some aspects of this disclosure provide polynucleotides encoding a Cas9 protein of a fusion protein as provided herein. Some aspects of this disclosure provide vectors comprising such polynucleotides. In some embodiments, the vector comprises a heterologous promoter driving expression of polynucleotide.

[00283] Some aspects of this disclosure provide cells comprising a Cas9 protein, a fusion protein, a nucleic acid molecule encoding the fusion protein, a complex comprise the Cas9 protein and the gRNA, and/or a vector as provided herein.

[00284] The description of exemplary embodiments of the reporter systems above is provided for illustration purposes only and not meant to be limiting. Additional reporter systems, *e.g.*, variations of the exemplary systems described in detail above, are also embraced by this disclosure.

EXAMPLES

EXAMPLE 1: Cas9 Deaminase Fusion Proteins

[00285] A number of Cas9:Deaminase fusion proteins were generated and deaminase activity of the generated fusions was characterized. The following deaminases were tested:

Human AID (hAID):
MDSLLMNRRKFLYQFKNVRWAKGRRETYLCYVVKRRDSATSFSLDFGYLRNKNGC
HVELLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVADFLRGNPYLSLRIFTAR
LYFCEDRKAEPEGLRRLHRAGVQIAIMTFKDYFYCWNTFVENHERTFKAWEGLHEN
SVRLSRQLRRILLPLYEVDDLRDAFRTLGLLD (SEQ ID NO: 607)

Human AID-DC (hAID-DC, truncated version of hAID with 7-fold increased activity):
MDSLLMNRRKFLYQFKNVRWAKGRRETYLCYVVKRRDSATSFSLDFGYLRNKNGC
HVELLFLRYISDWDLDPGRCYRVTWFTSWSPCYDCARHVADFLRGNPNLSLRIFTAR
LYFCEDRKAEPEGLRRLHRAGVQIAIMTFKDYFYCWNTFVENHERTFKAWEGLHEN

SVRLSRQLRRILL (SEQ ID NO: 608)

Rat APOBEC1 (rAPOBEC1):
MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK (SEQ
ID NO: 284)

Human APOBEC1 (hAPOBEC1)
MTSEKGPSTGDPTLRRRIEPWEFDVFYDPRELRKEACLLYEIKWGMSRKIWRSSGKN
TTNHVEVNFIKKFTSERDFHPSMSCSITWFLSWSPCWECSQAIREFLSRHPGVTLVIYV
ARLFWHMDQQNRQGLRDLVNSGVTIQIMRASEYYHCWRNFVNYPPGDEAHWPQYP
PLWMMLYALELHCIILSLPPCLKISRRWQNHLTFFRLHLQNCHYQTIPPHILLATGLIH
PSVAWR (SEQ ID NO: 5724)

Petromyzon marinus (Lamprey) CDA1 (pmCDA1):
MTDAEYVRIHEKLDIYTFKKQFFNNKKSVSHRCYVLFELKRRGERRACFWGYAVNK
PQSGTERGIHAEIFSIRKVEEYLRDNPGQFTINWYSSWSPCADCAEKILEWYNQELRG
NGHTLKIWACKLYYEKNARNQIGLWNLRDNGVGLNVMVSEHYQCCRKIFIQSSHNQ
LNENRWLEKTLKRAEKRRSELSIMIQVKILHTTKSPAV (SEQ ID NO: 609)

Human APOBEC3G (hAPOBEC3G):
MELKYHPEMRFFHWFSKWRKLHRDQEYEVTWYISWSPCTKCTRDMATFLAEDPKV
TLTIFVARLYYFWDPDYQEALRSLCQKRDGPRATMKIMNYDEFQHCWSKFVYSQRE
LFEPWNNLPKYYILLHIMLGEILRHSMDPPTFTFNFNNEPWVRGRHETYLCYEVERM
HNDTWVLLNQRRGFLCNQAPHKHGFLEGRHAELCFLDVIPFWKLDLDQDYRVTCFT
SWSPCFSCAQEMAKFISKNKHVSLCIFTARIYDDQGRCQEGLRTLAEAGAKISIMTYS
EFKHCWDTFVDHQGCPFQPWDGLDEHSQDLSGRLRAILQNQEN (SEQ ID NO: 610)

[00286] Deaminase Activity on ssDNA. A USER (Uracil-Specific Excision Reagent) Enzyme-based assay for deamination was employed to test the activity of various deaminases on single-stranded DNA (ssDNA) substrates. USER Enzyme was obtained from New England Biolabs. An ssDNA substrate was provided with a target cytosine residue at different positions. Deamination of the ssDNA cytosine target residue results in conversion of the target cytosine to a uracil. The USER Enzyme excises the uracil base and cleaves the ssDNA backbone at that position, cutting the ssDNA substrate into two shorter fragments of DNA. In some assays, the ssDNA substrate is labeled on one end with a dye, *e.g.*, with a 5' Cy3 label (the * in the scheme below). Upon deamination, excision, and cleavage of the strand, the substrate can be subjected to electrophoresis, and the substrate and any fragment released from it can be visualized by detecting the label. Where Cy5 is images, only the fragment with the label will be visible via imaging.

[00287] In one USER Enzyme assay, ssDNA substrates were used that matched the target sequences of the various deaminases tested. Expression cassettes encoding the deaminases

tested were inserted into a CMV backbone plasmid that has been used previously in the lab (Addgene plasmid 52970). The deaminase proteins were expressed using a TNT Quick Coupled Transcription/Translation System (Promega) according to the manufacturers recommendations. After 90 min of incubation, 5 mL of lysate was incubated with 5' Cy3-labeled ssDNA substrate and 1 unit of USER Enzyme (NEB) for 3 hours. The DNA was resolved on a 10% TBE PAGE gel and the DNA was imaged using Cy-dye imaging. A schematic reparesentation of the USER Enzyme assay is shown in Figure 41.

[00288] **Figure 1** shows the deaminase activity of the tested deaminases on ssDNA substrates, such as Doench 1, Doench 2, G7′ and VEGF Target 2. The rAPOBEC1 enzyme exhibited a substantial amount of deamination on the single-stranded DNA substrate with a canonical NGG PAM, but not with a negative control non-canonical NNN PAM.

Cas9 fusion proteins with APOBEC family deaminases were generated. The following fusion architectures were constructed and tested on ssDNA:

rAPOBEC1-GGS-*dCas9* primary sequence

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKGGS*D*
KKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAE
ATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHP
IFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLN
PDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEK
KNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLF
LAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYK
EIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTF
DNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFA
WMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFT
VYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFD
SVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEE
RLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANR
NFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELV
KVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQ
LQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDK
NRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIK
RQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVR
EINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKAT
AKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMP
QVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAK
VEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELE
NGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHK
HYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPA
AFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 611)

rAPOBEC1-(GGS)$_3$-*dCas9* primary sequence

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW
VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK GGSG
GSGGS *MDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLF*
DSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEED
KKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGH
FLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLI
AQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIG
DQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQ
QLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLL
RKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLAR
GNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSL
LYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFK
KIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDR
EMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSD
GFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKV
VDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHP
VENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVL
TRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDK
AGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQ
FYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQE
IGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKV
LSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVL
VVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSL
FELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFV
EQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNL
GAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD (SEQ ID NO: 612)

dCas9-GGS-rAPOBEC1

DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRT
FDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRF
AWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYF
TVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECF
DSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIE
ERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFAN
RNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDEL
VKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENT
QLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSD
KNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFI

KRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKA
TAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMP
QVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAK
VEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELE
NGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHK
HYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPA
*AFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD*GGSMSSETGPVA
VDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKHVEVNFI
EKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPR
NRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLEL
YCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK (SEQ ID NO: 613)

dCas9-GGS_3-rAPOBEC1

DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETA
EATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERH
PIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGE
KKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADL
FLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKY
KEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRT
FDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRF
AWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYF
TVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECF
DSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIE
ERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFAN
RNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDEL
VKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENT
QLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSD
KNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFI
KRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKV
REINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKA
TAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMP
QVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAK
VEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELE
NGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHK
HYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPA
*AFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD*GGSGGSGGSMSS
ETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKH
VEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYH
HADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRL
YVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK (SEQ ID NO: 614)

rAPOBEC1-XTEN-*dCas9* primary sequence

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLW

VRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLKSGSE TPGTSESATPES*DKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLI GALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESF LVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIK FRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRR LENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNL LAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLK ALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLN REDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYV GPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVL PKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLK EDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLT LFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILD FLKSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGIL QTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQI LKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSI DNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGL SELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDF RKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIA KSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFA TVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPT VAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIK LPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQ KQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLF TLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD* (SEQ ID NO: 615)

[00289] **Figure 2** shows that the N-terminal deaminase fusions showed significant activity on the single stranded DNA substrates. For this reason, only the N-terminal architecture was chosen for further experiments.

[00290] **Figure 3** illustrates double stranded DNA substrate binding by deaminase-dCas9:sgRNA complexes. A number of double stranded deaminase substrate sequences were generated. The sequences are provided below. The structures according to **Figure 3** are identified in these sequences (36bp: underlined, sgRNA target sequence: bold; PAM: boxed; 21bp: italicized). All substrates were labeled with a 5'-Cy3 label:

2:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGTC**CCGCGGATTTATTTATTT AA**TGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 616)
3:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCT**TCCGCGGATTTATTTATT TA**TGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 617)
4:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TTCCGCGGATTTATTTAT TA**TGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 618)
5:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTCCGCGGATTTATTTA TT**TGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 619)
6:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TATTCCGCGGATTTATTT AT**TGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 620)

7:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TTATTCCGCGGATTTATT**
TATGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 621)
8:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTATTCCGCGGATTTAT**
TTTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 622)
9:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TATTATTCCGCGGATTTA**
TTTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 623)
10:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTATATTCCGCGGATTT**
ATTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 624)
11:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TATTATATTCCGCGGATT**
TATGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 625)
12:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TTATTATATTCCGCGGAT**
TTTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 626)
13:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTATTATATTCCGCGGA**
TTTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 627)
14:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**TATTATTATATTCCGCGG**
ATTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 628)
15:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTATTATTATTACCGCG**
GATGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 629)
18:GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTATTATTATTATTACC**
GCTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 630)
"-" :
GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGTA**ATATTAATTTATTTATTTAA**
TGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 631)
8U:GTAGGTAGTTAGGATGAATGGAAGGTTGGTGTAG**ATTATTATCUGCGGATTTA**
TTGG*ATGACCTCTGGATCCATGGACAT*-3' (SEQ ID NO: 632)

*In all substrates except for "8U", the top strand in Figure 3 is the complement of the sequence specified here. In the case of "8U", there is a "G" opposite the U.

[00291] **Figure 4** shows the results of a double stranded DNA Deamination Assay. The fusions were expressed and purified with an N-terminal His6 tag via both Ni-NTA and sepharose chromatography. In order to assess deamination on dsDNA substrates, the various dsDNA substrates shown on the previous slide were incubated at a 1:8 dsDNA:fusion protein ratio and incubated at 37 °C for 2 hours. Once the dCas9 portion of the fusion binds to the DNA it blocks access of the USER enzyme to the DNA. Therefore, the fusion proteins were denatured following the incubation and the dsDNA was purified on a spin column, followed by incubation for 45 min with the USER Enzyme and resolution of the resulting DNA substrate and substrate fragments on a 10% TBE-urea gel.

[00292] **Figure 5** demonstrates that Cas9 fusions can target positions 3-11 of double-stranded DNA target sequences (numbered according to the schematic in Figure 3). Upper Gel: 1 μM rAPOBEC1-GGS-dCas9, 125 nM dsDNA, 1 eq sgRNA. Mid Gel: 1 μM

rAPOBEC1-$(GGS)_3$-dCas9, 125 nM dsDNA, 1 eq sgRNA. Lower Gel: 1.85 μM rAPOBEC1-XTEN-dCas9, 125 nM dsDNA, 1 eq sgRNA. Based on the data from these gels, positions 3-11 (according to the numbering in Figure 3) are sufficiently exposed to the activity of the deaminase to be targeted by the fusion proteins tested. Access of the deaminase to other positions is most likely blocked by the dCas9 protein.

[00293] The data further indicates that a linker of only 3 amino acids (GGS) is not optimal for allowing the deaminase to access the single stranded portion of the DNA. The 9 amino acid linker [$(GGS)_3$] (SEQ ID NO: 596) and the more structured 16 amino acid linker (XTEN) allow for more efficient deamination.

[00294] **Figure 6** demonstrates that the correct guide RNA, *e.g.*, the correct sgRNA, is required for deaminase activity. The gel shows that fusing the deaminase to dCas9, the deaminase enzyme becomes sequence specific (*e.g.*, using the fusion with an eGFP sgRNA results in no deamination), and also confers the capacity to the deaminase to deaminate dsDNA. The native substrate of the deaminase enzyme is ssDNA, and no deamination occurred when no sgRNA was added. This is consistent with reported knowledge that APOBEC deaminase by itself does not deaminate dsDNA. The data indicates that Cas9 opens the double-stranded DNA helix within a short window, exposing single-stranded DNA that is then accessible to the APOBEC deaminase for cytidine deamination. The sgRNA sequences used are provided below. sequences (36bp: underlined, sgRNA target sequence: bold; PAM: boxed; 21bp: italicized)

DNA sequence 8:

5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTATAGCC**ATTATTCCGCGGATTTATT**
TTGG*ATGACCTCTGGATCCATGGAC*-3' (SEQ ID NO: 633)

Correct sgRNA sequence (partial 3' sequence):
5'-**AUUAUUCCGCGGAUUUAUUU**GUUUUAGAGCUAG...-3' (SEQ ID NO: 634)

eGFP sgRNA sequence (partial 3'-sequence):
5'-**CGUAGGCCAGGGUGGUCACG**GUUUUAGAGCUAG...-3' (SEQ ID NO: 635)

EXAMPLE 2: Deamination of DNA target sequence

[00295] **Exemplary deamination targets.** The dCas9:deaminase fusion proteins described herein can be delivered to a cell in vitro or ex vivo or to a subject *in vivo* and can be used to effect C to T or G to A transitions when the target nucleotide is in positions 3-11 with respect to a PAM. Exemplary deamination targets include, without limitation, the following: CCR5

truncations: any of the codons encoding **Q93, Q102, Q186, R225, W86,** or **Q261** of CCR5 can be deaminated to generate a STOP codon, which results in a nonfunctional truncation of CCR5 with applications in HIV treatment. APOE4 mutations: mutant codons encoding **C11R** and **C57R** mutant APOE4 proteins can be deaminated to revert to the wild-type amino acid with applications in Alzheimer's treatment. eGFP truncations: any of the codons encoding **Q158, Q184, Q185** can be deaminated to generate a STOP codon, or the codon encoding **M1** can be deaminated to encode I, all of which result in loss of eGFP fluorescence, with applications in reporter systems. eGFP restoration: a mutant codon encoding **T65A** or **Y66C** mutant GFP, which does not exhibit substantial fluorescence, can be deaminated to restore the wild-type amino acid and confer fluorescence. PIK3CA mutation: a mutant codon encoding **K111E** mutant PIK3CA can be deaminated to restore the wild-type amino acid residue with applications in cancer. CTNNB1 mutation: a mutant codon encoding **T41A** mutant CTNNB1 can be deaminated to restore the wild-type amino acid residue with applications in cancer. HRAS mutation: a mutant codon encoding **Q61R** mutant HRAS can be deaminated to restore the wild-type amino acid residue with applications in cancer. P53 mutations: any of the mutant codons encoding **Y163C, Y236C**, or **N239D** mutant p53 can be deaminated to encode the wild type amino acid sequence with applications in cancer.

The feasibility of deaminating these target sequences in double-stranded DNA is demonstrated in **Figures** 7 and **8**. **Figure** 7 illustrates the mechanism of target DNA binding of *in vivo* target sequences by deaminase-dCas9:sgRNA complexes.

[00296] **Figure 8** shows successful deamination of exemplary disease-associated target sequences. Upper Gel: CCR5 Q93: coding strand target in pos. 10 (potential off-targets at positions 2, **5**, 6, 8, 9); CCR5 Q102: coding strand target in pos. **9** (potential off-targets at positions 1, 12, 14); CCR5 Q186: coding strand target in pos. 9 (potential off-targets at positions 1, **5**, 15); CCR5 R225: coding strand target in pos. **6** (no potential off-targets); eGFP Q158: coding strand target in pos. **5** (potential off-targets at positions 1, 13, 16); eGFP Q184 /185: coding strand target in pos. **4** and 7 (potential off-targets at positions 3, 12, 14, 15, 16, 17, 18); eGFP M1: template strand target in pos. 12 (potential off-targets at positions 2, **3**, 7, 9, 11) (targets positions 7 and 9 to small degree); eGFP T65A: template strand target in pos. 7 (potential off-targets at positions 1, **8**, 17); PIK3CA K111E: template strand target in pos. 2 (potential off-targets at positions **5**, **8**, 10, 16, 17); PIK3CA K111E: template strand target in pos. 13 (potential off-targets at positions 11, 16, 19) **X.** Lower Gel: CCR5 W86: template strand target in pos. 2 and 3 (potential off-targets at positions 1, 13) **X;** APOE4 C11R: coding strand target in pos. 11 (potential off-targets at positions 7, 13, 16, 17); APOE4 C57R: coding

strand target in pos. **5**) (potential off-targets at positions 7, 8, 12); eGFP Y66C: template strand target in pos. 11 (potential off-targets at positions 1, 4 ,**6** , 8, 9, 16); eGFP Y66C: template strand target in pos. 3 (potential off-targets at positions 1, **8**, 17); CCR5 Q261: coding strand target in pos. 10 (potential off-targets at positions 3, **5**, **6**, 9, 18); CTNNB1 T41A: template strand target in pos. 7 (potential off-targets at positions 1, 13, 15, 16) **X;** HRAS Q61R: template strand target in pos. 6 (potential off-targets at positions 1, 2, 4, **5**, 9, 10, 13); p53 Y163C: template strand target in pos. **6** (potential off-targets at positions 2, 13, 14); p53 Y236C: template strand target in pos. 8 (potential off-targets at positions 2, **4**); p53 N239D: template strand target in pos. **4** (potential off-targets at positions 6, 8). Exemplary DNA sequences of disease targets are provided below (PAMs (5'-NGG-3') and target positions are boxed):

CCR5 **Q93**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAACTAT**GCTGCCGCC**
CAGTGGGACTT[TGG]*AAATACAATGTGTCAACTCTT*-3' (SEQ ID NO: 636)
CCR5 **Q102**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAAAATA**CAATGTGT**
CAACTCTTGACA[GGG]*CTCTATTTTATAGGCTTCTTC*-3' (SEQ ID NO: 637)
CCR5 **Q186**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTATTTTC**CATACAGT**
CAGTATCAATTC[TGG]*AAGAATTTCCAGACATTAAAG*-3' (SEQ ID NO: 638)
CCR5 **R225**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGCTTC**GGTGT**[C]**GA**
AATGAGAAGAAG[AGG]*CACAGGGCTGTGAGGCTTATC*-3' (SEQ ID NO: 639)
CCR5 **W86**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGTGAG**C**[**CC**]**AGAAGG**
GGACAGTAAGA[AGG]*AAAAACAGGTCAGAGATGGCC*-3' (SEQ ID NO: 640)
CCR5 **Q261**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTATCCTG**AACACCTT**
C[**C**]**AGGAATTCTT**[TGG]*CCTGAATAATTGCAGTAGCTC*-3' (SEQ ID NO: 641)
APOE4 **C11R**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGACAT**GGAGGAC**
GTG[C]**GCGGCCGCC**[TGG]*TGCAGTACCGCGGCGAGGTGC*-3' (SEQ ID NO: 642)
APOE4 **C57R**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTACTGCA**GAAG**[C]**GC**
CTGGCAGTGTACC[AGG]*CCGGGGCCCGCGAGGGCGCCG*-3' (SEQ ID NO: 643)
eGFP **Q158**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGCCGA**CAAG**[C]**AGA**
AGAACGGCATCA[AGG]*TGAACTTCAAGATCCGCCACA*-3' (SEQ ID NO: 644)
eGFP **Q184/185**: 5'-Cy3-GTAGGTAGTTAGGATGAATGGAAGGTTGGTAACCAC**TAC**[C]
AG[C]**AGAACACCCCCAT**[CGG]*CGACGGCCCCGTGCTGCTGCC*-3' (SEQ ID NO: 645)
eGFP **M1**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTACCTCG**CCCTTGCTCA**
C[**C**]**ATCTCGAGT**[CGG]*CCGCCAGTGTGATGGATATCT*-3' (SEQ ID NO: 646)

eGFP **T65A**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTACACGC**GTAGGCCA**
GGGTGGTCACGAGG*GTGGGCCAGGGCACGGGCAGC*-3' (SEQ ID NO: 647)
eGFP **Y66C**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAAAGCA**CTGCACTC**
CGCAGGTCAGGGTGG*TCACGAGGGTTGGCCAGGGCA*-3' (SEQ ID NO: 648)
eGFP **Y66C**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTACACTC**CGCAGGTC**
AGGGTGGTCACGAGG*GTTGGCCAGGGCACGGGCAGG*-3' (SEQ ID NO: 649)
PIK3CA **K111E**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGGATC**TCTTC**
TTCACGGTTGCCTACTGG*TTCAATTACTTTTAAAAATGG*-3' (SEQ ID NO: 650)
PIK3CA **K111E**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTATTCTC**GATTG**
AGGATCTCTTCTTCACGG*TTGCCTACTGGTTCAATTACT*-3' (SEQ ID NO: 651)
CTNNB1 **T41A**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAAGGAG**CTGTGG**
CAGTGGCACCAGAATGG*ATTCCAGAGTCCAGGTAAGAC*-3' (SEQ ID NO: 652)
HRAS **Q61R**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGTACT**CCTCCCGG**
CCGGCGGTATCCAGG*ATGTCCAACAGGCACGTCTCC*-3' (SEQ ID NO: 653)
p53 **Y163C**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTATGACT**GCTTGCAG**
ATGGCCATGGCGCGG*ACGCGGGTGCCGGGCGGGGGT*-3' (SEQ ID NO: 654)
p53 **Y236C**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTACTGTT**ACACATGC**
AGTTGTAGTGGATGG*TGGTACAGTCAGAGCCAACCT*-3' (SEQ ID NO: 655)
p53 **N239D**: 5'-Cy3-
GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGGAAC**TGTCACAC**
ATGTAGTTGTAGTGG*ATGGTGGTACAGTCAGAGCCA*-3' (SEQ ID NO: 656)

EXAMPLE 3: Uracil Glycosylase Inhibitor Fusion Improves Deamination Efficiency

[00297] Direct programmable nucleobase editing efficiencies in mammalian cells by dCas9:deaminase fusion proteins can be improved significantly by fusing a uracil glycosylase inhibitor (UGI) to the dCas9:deaminase fusion protein.

[00298] **Figure 9** shows *in vitro* C→T editing efficiencies in human HEK293 cells using rAPOBEC1-XTEN-dCas9:

rAPOBEC1-**XTEN**-dCas9-*NLS* primary sequence

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWV
RLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK**SGSETP**
GTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIG
ALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLV

EEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFR
GHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLE
NLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLA
QIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALV
RQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNRED
LLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLAR
GNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLL
YEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFK
KIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDRE
MIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDG
FANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVD
ELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVE
NTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLT
RSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDK
AGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQ
FYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQ
EIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKV
LSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSV
LVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYS
LFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLF
VEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNL
GAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGS*PKKKR*
KV (SEQ ID NO: 657)

Protospacer sequences were as follows:

EMX1: 5'- GAGT**C**$_5$**C**$_6$GAG**C**$_{10}$AGAAGAAGAAGGG -3' (SEQ ID NO: 293)

FANCF: 5'- GGAAT**C**$_6$**C**$_7$**C**$_8$TT**C**$_{11}$TGCAGCACCTGG -3' (SEQ ID NO: 294)

HEK293 site 2: 5'- GAA**C**$_4$A**C**$_6$AAAG**C**$_{11}$ATAGACTGCGGG -3' (SEQ ID NO: 295)

HEK293 site 3: 5'- GG**C**$_3$**C**$_4$**C**$_5$AGA**C**$_9$TGAGCACGTGATGG -3' (SEQ ID NO: 296)

HEK293 site 4: 5'- GG**C**$_3$A**C**$_5$TG**C**$_8$GG**C**$_{11}$TGGAGGTGGGGG -3' (SEQ ID NO: 297)

RNF2: 5'- GT**C**$_3$AT**C**$_6$TTAGTCATTACCTGAGG -3' (SEQ ID NO: 298)

*PAMs are boxed, C residues within target window (positions 3-11) are numbered and bolded.

[00299] **Figure 10** demonstrates that C→T editing efficiencies on the same protospacer sequences in HEK293T cells are greatly enhanced when a UGI domain is fused to the rAPOBEC1:dCas9 fusion protein.

rAPOBEC1-**XTEN**-dCas9-**UGI**-*NLS* primary sequence

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNT
NKHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIAR
LYHHADPRNRQGLRDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWV
RLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSCHYQRLPPHILWATGLK**SGSETP**

GTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIG
ALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLV
EEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFR
GHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLE
NLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLA
QIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALV
RQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNRED
LLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLAR
GNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLL
YEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFK
KIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDRE
MIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDG
FANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVD
ELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVE
NTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLT
RSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDK
AGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQ
FYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQ
EIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKV
LSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSV
LVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYS
LFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLF
VEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNL
GAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGS**TNLSD
IIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLLTSDAP
EYKPWALVIQDSNGENKIKML**SGGS*PKKKRKV* (SEQ ID NO: 658)

[00300] The percentages in Figures 9 and 10 are shown from sequencing both strands of the target sequence. Because only one of the strands is a substrate for deamination, the maximum possible deamination value in this assay is 50%. Accordingly, the deamination efficiency is double the percentages shown in the tables. *E.g.*, a value of 50% relates to deamination of 100% of double-stranded target sequences.

When a uracil glycosylase inhibitor (UGI) was fused to the dCas9:deaminase fusion protein (*e.g.*, rAPOBEC1-XTEN-dCas9-[UGI]-NLS), a significant increase in editing efficiency in cells was observed. This result indicates that in mammalian cells, the DNA repair machinery that cuts out the uracil base in a U:G base pair is a rate-limiting process in DNA editing. Tethering UGI to the dVas9:deaminase fusion proteins greatly increases editing yields.

[00301] Without UGI, typical editing efficiencies in human cells were in the ~2-14% yield range (Figure 9 and Figure 10, "XTEN" entries). With UGI (Figure 10, "UGI" entries) the editing was observed in the ~6-40% range. Using a UGI fusion is thus more efficient than the current alternative method of correcting point mutations via HDR, which also creates an

excess of indels in addition to correcting the point mutation. No indels resulting from treatment with the cas9:deaminase:UGI fusions were observed.

EXAMPLE 4: Direct, programmable conversion of a target nucleotide in genomic DNA without double-stranded DNA cleavage

[00302] Current genome-editing technologies introduce double-stranded DNA breaks at a target locus of interest as the first step to gene correction.[39,40] Although most genetic diseases arise from mutation of a single nucleobase to a different nucleobase, current approaches to revert such changes are very inefficient and typically induce an abundance of random insertions and deletions (indels) at the target locus as a consequence of the cellular response to double-stranded DNA breaks.[39,40] Reported herein is the development of nucleobase editing, a new strategy for genome editing that enables the direct conversion of one target nucleobase into another in a programmable manner, without requiring double-stranded DNA backbone cleavage. Fusions of CRISPR/Cas9 were engineered and the cytidine deaminase enzyme APOBEC1 that retain the ability to be programmed with a guide RNA, do not induce double-stranded DNA breaks, and mediate the direct conversion of cytidine to uracil, thereby effecting a C→T (or G→A) substitution following DNA replication, DNA repair, or transcription if the template strand is targeted. The resulting "nucleobase editors" convert cytidines within a window of approximately five nucleotides, and can efficiently correct a variety of point mutations relevant to human disease *in vitro*. In four transformed human and murine cell lines, second- and third-generation nucleobase editors that fuse uracil glycosylase inhibitor (UGI), and that use a Cas9 nickase targeting the non-edited strand, respectively, can overcome the cellular DNA repair response to nucleobase editing, resulting in permanent correction of up to 37% or (~15-75%) of total cellular DNA in human cells with minimal (typically ≤ 1%) indel formation. In contrast, canonical Cas9-mediated HDR on the same targets yielded an average of 0.7% correction with 4% indel formation. Nucleobase editors were used to revert two oncogenic *p53* mutations into wild-type alleles in human breast cancer and lymphoma cells, and to convert an Alzheimer's Disease associated Arg codon in *ApoE4* into a non-disease-associated Cys codon in mouse astrocytes. Base editing expands the scope and efficiency of genome editing of point mutations.

[00303] The clustered regularly interspaced short palindromic repeat (CRISPR) system is a prokaryotic adaptive immune system that has been adapted to mediate genome engineering in a variety of organisms and cell lines.[41] CRISPR/Cas9 protein-RNA complexes localize to a

target DNA sequence through base pairing with a guide RNA, and natively create a DNA double-stranded break (DSB) at the locus specified by the guide RNA. In response to DSBs, endogenous DNA repair processes mostly result in random insertions or deletions (indels) at the site of DNA cleavage through non-homologous end joining (NHEJ). In the presence of a homologous DNA template, the DNA surrounding the cleavage site can be replaced through homology-directed repair (HDR). When simple disruption of a disease-associated gene is sufficient (for example, to treat some gain-of-function diseases), targeted DNA cleavage followed by indel formation can be effective. For most known genetic diseases, however, correction of a point mutation in the target locus, rather than stochastic disruption of the gene, is needed to address or study the underlying cause of the disease.[68]

[00304] Motivated by this need, researchers have invested intense effort to increase the efficiency of HDR and suppress NHEJ. For example, a small-molecule inhibitor of ligase IV, an essential enzyme in the NHEJ pathway, has been shown to increase HDR efficiency.[42,43] However, this strategy is challenging in post-mitotic cells, which typically down-regulate HDR, and its therapeutic relevance is limited by the potential risks of inhibiting ligase IV in non-target cells. Enhanced HDR efficiency can also be achieved by the timed delivery of Cas9-guide RNA complexes into chemically synchronized cells, as HDR efficiency is highly cell-cycle dependent.[44] Such an approach, however, is limited to research applications in cell culture since synchronizing cells is highly disruptive. Despite these developments, current strategies to replace point mutations using HDR in most contexts are very inefficient (typically ~0.1 to 5%),[42,43,45,46, 75] especially in unmodified, non-dividing cells. In addition, HDR competes with NHEJ during the resolution of double-stranded breaks, and indels are generally more abundant outcomes than gene replacement. These observations highlight the need to develop alternative approaches to install specific modifications in genomic DNA that do not rely on creating double-stranded DNA breaks. A small-molecule inhibitor of ligase IV, an essential enzyme in the NHEJ pathway, has been shown to increase HDR efficiency.[42,43] However, this strategy is challenging in post-mitotic cells, which typically down-regulate HDR, and its therapeutic relevance is limited by the potential risks of inhibiting ligase IV in non-target cells. Enhanced HDR efficiency can also be achieved by the timed delivery of Cas9-guide RNA complexes into chemically synchronized cells, as HDR efficiency is highly cell-cycle dependent.[44] Such an approach, however, is limited to research applications in cell culture since synchronizing cells is highly disruptive. In some cases, it is possible to design HDR templates such that the product of successful HDR contains mutations in the PAM sequence and therefore is no longer a substrate for subsequent Cas9 modification, increasing

the overall yield of HDR products,[75] although such an approach imposes constraints on the product sequences. Recently, this strategy has been coupled to the use of ssDNA donors that are complementary to the non-target strand and high-efficiency ribonucleoprotein (RNP) delivery to substantially increase the efficiency of HDR, but even in these cases the ratio of HDR to NHEJ outcomes is relatively low (< 2).[83]

[00305] It was envisioned that direct catalysis of the conversion of one nucleobase to another at a programmable target locus without requiring DNA backbone cleavage could increase the efficiency of gene correction relative to HDR without introducing undesired random indels at the locus of interest. Catalytically dead Cas9 (dCas9), which contains Asp10Ala and His840Ala mutations that inactivate its nuclease activity, retains its ability to bind DNA in a guide RNA-programmed manner but does not cleave the DNA backbone.[16,47] In principle, conjugation of dCas9 with an enzymatic or chemical catalyst that mediates the direct conversion of one nucleobase to another could enable RNA-programmed nucleobase editing. The deamination of cytosine (C) is catalyzed by cytidine deaminases[29] and results in uracil (U), which has the base pairing properties of thymine (T). dCas9 was fused to cytidine deaminase enzymes in order to test their ability to convert C to U at a guide RNA-specified DNA locus. Most known cytidine deaminases operate on RNA, and the few examples that are known to accept DNA require single-stranded DNA.[48] Recent studies on the dCas9-target DNA complex reveal that at least nine nucleotides of the displaced DNA strand are unpaired upon formation of the Cas9:guide RNA:DNA "R-loop" complex.[12] Indeed, in the structure of the Cas9 R- loop complex the first 11 nucleotides of the protospacer on the displaced DNA strand are disordered, suggesting that their movement is not highly restricted.[76] It has also been speculated that Cas9 nickase-induced mutations at cytosines in the non-template strand might arise from their accessibility by cellular cytidine deaminase enzymes.[77] Recent studies on the dCas9-target DNA complex have revealed that at least 26 bases on the non-template strand are unpaired when Cas9 binds to its target DNA sequence.[49] It was reasoned that a subset of this stretch of single-stranded DNA in the R-loop might serve as a substrate for a dCas9-tethered cytidine deaminase to effect direct, programmable conversion of C to U in DNA (Figure 11A).

[00306] Four different cytidine deaminase enzymes (hAID, hAPOBEC3G, rAPOBEC1, and pmCDA1) were expressed in a mammalian cell lysate-derived *in vitro* transcription-translation system and evaluated for ssDNA deamination. Of the four enzymes, rAPOBEC1 showed the highest deaminase activity under the tested conditions and was chosen for dCas9 fusion experiments (Figure 36A). Although appending rAPOBEC1 to the C-terminus of

dCas9 abolishes deaminase activity, fusion to the N-terminus of dCas9 preserves deaminase activity on ssDNA at a level comparable to that of the unfused enzyme. Four rAPOBEC1-dCas9 fusions were expressed and purified with linkers of different length and composition (Figure 36B), and evaluated each fusion for single guide RNA (sgRNA)-programmed dsDNA deamination *in vitro* (Figures 11A to 11C and Figures 15A to 15D).

Efficient, sequence-specific, sgRNA-dependent C to U conversion was observed *in vitro* (Figures 11A to 11C). Conversion efficiency was greatest using rAPOBEC1-dCas9 linkers over nine amino acids in length. The number of positions susceptible to deamination (the deamination "activity window") increases with linker length was extended from three to 21 amino acids (Figures 36C to 36F15A to 15D). The 16-residue XTEN linker[50] was found to offer a promising balance between these two characteristics, with an efficient deamination window of approximately five nucleotides, from positions 4 to 8 within the protospacer, counting the end distal to the protospacer-adjacent motif (PAM) as position 1. The rAPOBEC1-XTEN-dCas9 protein served as the first-generation nucleobase editor (NBE1).

[00307] Elected were seven mutations relevant to human disease that in theory could be corrected by C to T nucleobase editing, synthesized double-stranded DNA 80-mers of the corresponding sequences, and assessed the ability of NBE1 to correct these mutations *in vitro* (Figures 16A to 16B). NBE1 yielded products consistent with efficient editing of the target C, or of at least one C within the activity window when multiple Cs were present, in six of these seven targets *in vitro*, with an average apparent editing efficiency of 44% (Figures 16A to 16B). In the three cases in which multiple Cs were present within the deamination window, evidence of deamination of some or all of these cytosines was observed. In only one of the seven cases tested were substantial yields of edited product observed (Figures 16A to 16B). Although the preferred sequence context for APOBEC1 substrates is reported to be CC or TC,[51] it was anticipated that the increased effective molarity of the deaminase and its single-stranded DNA substrate mediated by dCas9 binding to the target locus may relax this restriction. To illuminate the sequence context generality of NBE1, its ability to edit a 60-mer double-stranded DNA oligonucleotide containing a single fixed C at position 7 within the protospacer was assayed, as well as all 36 singly mutated variants in which protospacer bases 1-6 and 8-13 were individually varied to each of the other three bases. Each of these 37 sequences were treated with 1.9 μM NBE1, 1.9 μM of the corresponding sgRNA, and 125 nM DNA for 2 h, similar to standard conditions for *in vitro* Cas9 assays[52]. High-throughput DNA sequencing (HTS) revealed 50 to 80% C to U conversion of targeted strands (25 to 40% of total sequence reads arising from both DNA strands, one of which is not a substrate for

NBE1) (Figure 12A). The nucleotides surrounding the target C had little effect on editing efficiency was independent of sequence context unless the base immediately 5' of the target C is a G, in which case editing efficiency was substantially lower (Figures 12A to 12B). NBE1 activity *in vitro* was assessed on all four NC motifs at positions 1 through 8 within the protospacer (Figures 12A to 12B). In general NBE1 activity on substrates was observed to follow the order **T**C ≥ **C**C ≥ A**C** >G**C**, with maximum editing efficiency achieved when the target C is at or near position 7. In addition, it was observed that the nucleobase editor is highly processive, and will efficiently convert most of all Cs to Us on the same DNA strand within the 5-base activity window (Figure 17).

[00308] While BE1 efficiently processes substrates in a test tube, in cells a tree of possible DNA repair outcomes determines the fate of the initial U:G product of base editing (Figure 29A). To test the effectiveness of nucleobase editing in human cells, NBE1 codon usage was optimized for mammalian expression, appended a C-terminal nuclear localization sequence (NLS),[53] and assayed its ability to convert C to T in human cells on 14Cs in six well-studied target sites throughout the human genome (Figure 37A).[54] The editable Cs were confirmed within each protospacer *in vitro* by incubating NBE1 with synthetic 80-mers that correspond to the six different genomic sites, followed by HTS (Figures 13A to 13C, Figure 29B and Figure 25). Next, HEK293T cells were transfected with plasmids encoding NBE1 and one of the six target sgRNAs, allowed three days for nucleobase editing to occur, extracted genomic DNA from the cells, and analyzed the loci by HTS. Although C to T editing in cells at the target locus was observed for all six cases, the efficiency of nucleobase editing was 1.1% to 6.3% or 0.8%-7.7% of total DNA sequences (corresponding to 2.2% to 12.6% of targeted strands), a 6.3-fold to 37-fold or 5-fold to 36-fold decrease in efficiency compared to that of *in vitro* nucleobase editing (Figures 13A to 13C, Figure 29B and Figure 25). It was observed that some base editing outside of the typical window of positions 4 to 8 when the substrate C is preceded by a T, which we attribute to the unusually high activity of APOBEC1 for TC substrates.[48]

[00309] It was asked whether the cellular DNA repair response to the presence of U:G heteroduplex DNA was responsible for the large decrease in nucleobase editing efficiency in cells (Figure 29A). Uracil DNA glycosylase (UDG) catalyzes removal of U from DNA in cells and initiates base excision repair (BER), with reversion of the U:G pair to a C:G pair as the most common outcome (Figure 29A).[55] Uracil DNA glycosylase inhibitor (UGI), an 83-residue protein from *B. subtilis* bacteriophage PBS1, potently blocks human UDG activity (IC_{50} = 12 pM).[56] UGI was fused to the C-terminus of NBE1 to create the second-generation

nucleobase editor NBE2 and repeated editing assays on all six genomic loci. Editing efficiencies in human cells were on average 3-fold higher with NBE2 than with NBE1, resulting in gene conversion efficiencies of up to 22.8% of total DNA sequenced (up to 45.6% of targeted strands) (Figures 13A to 13C and Figure 29B). To test base editing in human cells, BE1 codon usage was optimized for mammalian expression and appended a C-terminal nuclear localization sequence (NLS).[53]

[00310] Similar editing efficiencies were observed when a separate plasmid overexpressing UGI was co-transfected with NBE1 (Figures 18A to 18H). However, while the direct fusion of UGI to NBE1 resulted in no significant increase in C to T mutations at monitored non-targeted genomic locations, overexpression of unfused UGI detectably increased the frequency of C to T mutations elsewhere in the genome (Figures 18A to 18H). The generality of NBE2-mediated nucleobase editing was confirmed by assessing editing efficiencies on the same six genomic targets in U2OS cells, and observed similar results with those in HEK293T cells (Figure 19). Importantly, NBE2 typically did not result in any detectable indels (Figure 13C and Figure 29C), consistent with the known mechanistic dependence of NHEJ on double-stranded DNA breaks.[57, 78] Together, these results indicate that conjugating UGI to NBE1 can greatly increase the efficiency of nucleobase editing in human cells.

[00311] The permanence of nucleobase editing in human cells was confirmed by monitoring editing efficiencies over multiple cell divisions in HEK293T cells at two of the tested genomic loci. Genomic DNA was harvested at two time points: three days after transfection with plasmids expressing NBE2 and appropriate sgRNAs, and after passaging the cells and growing them for four additional days (approximately five subsequent cell divisions). No significant change in editing efficiency was observed between the non-passaged cells (editing observed in 4.6% to 6.6% of targeted strands for three different target Cs) and passaged cells (editing observed in 4.6% to 6.4% of targeted strands for the same three target Cs), confirming that the nucleobase edits became permanent following cell division (Figure 20). Indels will on rare occasion arise from the processing of U:G lesions by cellular repair processes, which involve single-strand break intermediates that are known to lead to indels.[84] Given that several hundred endogenous U:G lesions are generated every day per human cell from spontaneous cytidine deaminase,[85] it was anticipate that the total indel frequency from U:G lesion repair is unlikely to increase from BE1 or BE2 activity at a single target locus.

[00312] To further increase the efficiency of nucleobase editing in cells, it was anticipated that nicking the non-edited strand may result in a smaller fraction of edited Us being removed

by the cell, since eukaryotic mismatch repair machinery uses strand discontinuity to direct DNA repair to any broken strand of a mismatched duplex (Figure 29A).[58, 79, 80] The catalytic His residue was restored at position 840 in the Cas9 HNH domain,[47,59] resulting in the third-generation nucleobase editor NBE3 that nicks the non-edited strand containing a G opposite the targeted C, but does not cleave the target strand containing the C. Because NBE3 still contains the Asp10Ala mutation in Cas9, it does not induce double-stranded DNA cleavage. This strategy of nicking the non-edited strand augmented nucleobase editing efficiency in human cells by an additional 1.4- to 4.8-fold relative to NBE2, resulting in up to 36.3% of total DNA sequences containing the targeted C to T conversion on the same six human genomic targets in HEK293T cells (Figures 13A to 13C and Figure 29B). Importantly, only a small frequency of indels, averaging 0.8% (ranging from 0.2% to 1.6% for the six different loci), was observed from NBE3 treatment (Figure 13C, Figure 29C, and Figure 34). In contrast, when cells were treated with wild-type Cas9, sgRNA, and a single-stranded DNA donor template to mediate HDR at three of these loci C to T conversion efficiencies averaging only 0.7% were observed, with much higher relative indel formation averaging 3.9% (Figures 13A to 13C and Figure 29C). The ratio of allele conversion to NHEJ outcomes averaged >1,000 for BE2, 23 for BE3, and 0.17 for wild-type Cas9 (Fig. 3c). We confirmed the permanence of base editing in human cells by monitoring editing efficiencies over multiple cell divisions in HEK293T cells at the HEK293 site 3 and 4 genomic loci (Figure 38). These results collectively establish that nucleobase editing can effect much more efficient targeted single-base editing in human cells than Cas9-mediated HDR, and with much less (NBE3) or no (NBE2) indel formation.

[00313] Next, the off-target activity of NBE1, NBE2, and NBE3 in human cells was evaluated. The off-target activities of Cas9, dCas9, and Cas9 nickase have been extensively studied (Figures 23 to 24 and 31 to 33).[54,60-62] Because the sequence preference of rAPOBEC1 has been shown to be independent of DNA bases more than one base from the target C,[63] consistent with the sequence context independence observed in Figures 12A to 12B, it was assumed that potential off-target activity of nucleobase editors arises from off-target Cas9 binding. Since only a fraction of Cas9 off-target sites will have a C within the active window for nucleobase editing, off-target nucleobase editing sites should be a subset of the off-target sites of canonical Cas9 variants. For each of the six sites studied, the top ten known Cas9 off-target loci in human cells that were previously determined using the GUIDE-seq method were sequenced (Figures 23 to 27 and 31 to 33).[54, 61] Detectable off-target nucleobase editing at only a subset (16/34, 47% for NBE1 and NBE2, and 17/34, 50% for

NBE3) of known dCas9 off-target loci was observed. In all cases, the off-target base-editing substrates contained a C within the five-base target window. In general, off-target C to T conversion paralleled off-target Cas9 nuclease-mediated genome modification frequencies (Figures 23 to 27). Also monitored were C to T conversions at 2,500 distinct cytosines surrounding the six on-target and 34 off-target loci tested, representing a total of 14,700,000 sequence reads derived from approximately $1.8x10^6$ cells, and observed no detectable increase in C to T conversions at any of these other sites upon NBE1, NBE2, or NBE3 treatment compared to that of untreated cells (Figure 28). Taken together, these findings suggest that off-target substrates of nucleobase editors include a subset of Cas9 off-target substrates, and that nucleobase editors in human cells do not induce untargeted C to T conversion throughout the genome at levels that can be detected by the methods used here. No substantial change was observed in editing efficiency between non-passaged HEK293T cells (editing observed in 1.8% to 2.6% of sequenced strands for the three target Cs with BE2, and 6.2% to 14.3% with BE3) and cells that had undergone approximately five cell divisions after base editing (editing observed in 1.9% to 2.3% of sequenced strands for the same target Cs with BE2, and 6.4% to 14.5% with BE3), confirming that base edits in these cells are durable (Extended Data Fig. 6).

[00314] Finally, the potential of nucleobase editing to correct three disease-relevant mutations in mammalian cells was tested. The apolipoprotein E gene variant *APOE4* encodes two Arg residues at amino acid positions 112 and 158, and is the largest and most common genetic risk factor for late-onset Alzheimer's disease.[64] ApoE variants with Cys residues in positions 112 or 158, including *APOE2* (Cys112/Cys158), *APOE3* (Cys112/Arg158), and *APOE3'* (Arg112/Cys158) have been shown[65] or are presumed[81] to confer substantially lower Alzheimer's disease risk than *APOE4*. Encouraged by the ability of NBE1 to convert *APOE4* to *APOE3' in vitro* (Figures 16A to 16B), this conversion was attempted in immortalized mouse astrocytes in which the endogenous murine *APOE* gene has been replaced by human *APOE4* (Taconic). DNA encoding NBE3 and an appropriate sgRNA was delivered into these astrocytes by nucleofection (nucleofection efficiency of 25%), extracted genomic DNA from all treated cells two days later, and measured editing efficiency by HTS. Conversion of Arg158 to Cys158 was observed in 58-75% of total DNA sequencing reads (44% of nucleofected astrocytes) (Figures 14A to 14C and Figures 30A). Also observed was 36-50% editing of total DNA at the third position of codon 158 and 38-55% editing of total DNA at the first position of Leu159, as expected since all three of these Cs are within the active nucleobase editing window. However, neither of the other two C→T conversions results in a

change in the amino acid sequence of the ApoE3' protein since both TGC and TGT encode Cys, and both CTG and TTG encode Leu. From > 1,500,000 sequencing reads derived from $1x10^6$ cells evidence of 1.7% indels at the targeted locus following NBE3 treatment was observed (Figure 35). In contrast, identical treatment of astrocytes with wt Cas9 and donor ssDNA resulted in 0.1-0.3% APOE4 correction and 26-40% indels at the targeted locus, efficiencies consistent with previous reports of single-base correction using Cas9 and HDR[45,75] (Figure 30A and Figure 40A). Astrocytes treated identically but with an sgRNA targeting the VEGFA locus displayed no evidence of APOE4 base editing (Figure 34 and Figure 40A). These results demonstrate how nucleobase editors can effect precise, single-amino acid changes in the coding sequence of a protein as the major product of editing, even when their processivity results in more than one nucleotide change in genomic DNA. The off-target activities of Cas9, dCas9, and Cas9 nickase have been extensively studied.[54, 60-62] In general, off-target C to T conversions by BE1, BE2, and BE3 paralleled off-target Cas9 nuclease-mediated genome modification frequencies.

[00315] The dominant-negative p53 mutations Tyr163Cys and Asn239Asp are strongly associated with several types of cancer.[66-67] Both of these mutations can be corrected by a C to T conversion on the template strand (Figures 16A to 16B). A human breast cancer cell line homozygous for the p53 Tyr163Cys mutation (HCC1954 cells) was nucleofected with DNA encoding NBE3 and an sgRNA programmed to correct Tyr163Cys. Because the nucleofection efficiency of HCC1954 cells was < 10%, a plasmid expressing IRFP was co-nucleofected into these cells to enable isolation of nucleofected cells by fluorescence-activated cell sorting two days after treatment. HTS of genomic DNA revealed correction of the Tyr163Cys mutation in 7.6% of nucleofected HCC1954 cells (Figure 30B and Figure 40A to 40B). Also nucleofected was a human lymphoma cell line that is heterozygous for p53 Asn239Asp (ST486 cells) with DNA encoding NBE2 and an sgRNA programmed to correct Asn239Asp with 92% nucleofection efficiency). Correction of the Asn239Asp mutation was observed in 11% of treated ST486 cells (12% of nucleofected ST486 cells). Consistent with the findings in HEK cells, no indels were observed from the treatment of ST486 cells with NBE2, and 0.6% indel formation from the treatment of HCC1954 cells with NBE3. No other DNA changes within at least 50 base pairs of both sides of the protospacer were detected at frequencies above that of untreated controls out of > 2,000,000 sequencing reads derived from $2x10^5$ cells (Figures 14A to 14C, Figure 30B and Table 1). These results collectively represent the conversion of three disease-associated alleles in genomic DNA into their wild-

type forms with an efficiency and lack of other genome modification events that is, to our knowledge, not currently achievable using other methods.

[00316] To illuminate the potential relevance of nucleobase editors to address human genetic diseases, the NCBI ClinVar database[68] was searched for known genetic diseases that could in principle be corrected by this approach. ClinVar was filtered by first examining only single nucleotide polymorphisms (SNPs), then removing any nonpathogenic variants. Out of the 24,670 pathogenic SNPs, 3,956 are caused by either a T to C, or an A to G, substitution. This list was further filtered to only include variants with a nearby NGG PAM that would position the SNP within the deamination activity window, resulting in 1,089 clinically relevant pathogenic gene variants that could in principle be corrected by the nucleobase editors described here (Figure 21 and Table 1). To illuminate the potential relevance of base editors to address human genetic diseases, the NCBI ClinVar database[68] was searched for known genetic diseases that could in principle be corrected by this approach. ClinVar was filtered by first examining only single nucleotide polymorphisms (SNPs), then removing any non-pathogenic variants. Out of the 24,670 pathogenic SNPs, 3,956 are caused by either a T to C, or an A to G, substitution. This list was further filtered to only include variants with a nearby NGG PAM that would position the SNP within the deamination activity window, resulting in 911 clinically relevant pathogenic gene variants that could in principle be corrected by the base editors described here. Of these, 284 contain only one C within the base editing activity window. A detailed list of these pathogenic mutations can be found in Table 1.

[00317] Table 1. List of 911 base-editable gene variants associated with human disease with an NGG PAM (SEQ ID NOs: 747 to 1868 appear from top to bottom below, respectively). The "Y" in the protospacer and PAM sequences indicates the base to be edited, e.g., C. (SEQ ID NOs: 747 to 1868 appear from top to bottom below, respectively)

dbSNP #	Genotype	Protospacer and PAM sequence(s)	Associated genetic disease
755445790	NM_000391.3(TPP1):c.887-10A>G	TTTYTTTTTTTTTTTTTTTGAGG	Ceroid lipofuscinosis, neuronal, 2
113994167	NM_000018.3(ACADVL):c.848T>C (p.Val283Ala)	TTTGYGGTGGAGAGGGGCTTCGG, TTGYGGTGGAGAGGGGCTTCGGG	Very long chain acyl-CoA dehydrogenase deficiency
119470018	NM_024996.5(GFM1):c.521A>G (p.Asn174Ser)	TTGYTAATAAAAGTTAGAAACGG	Combined oxidative phosphorylation deficiency 1
115650537	NM_000426.3(LAMA2):c.8282T>C (p.Ile2761Thr)	TTGAYAGGGAGCAAGCAGTTCGG, TGAYAGGGAGCAAGCAGTTCGGG	Merosin deficient congenital muscular dystrophy
587777752	NM_014946.3(SPAST):c.1688-	TTCYGTAAAACATAAAAGTCAGG	Spastic paraplegia 4, autosomal dominant

794726821	NM_001165963.1(SCN1A):c.4055T>C (p.Leu1352Pro)	TTCYGGTTTGTCTTATATTCTGG	Severe myoclonic epilepsy in infancy
397514745	NM_001130089.1(KARS):c.517T>C (p.Tyr173His)	CTTCYATGATCTTCGAGGAGAGG, TTCYATGATCTTCGAGGAGAGG G	Deafness, autosomal recessive 89
376960358	NM_001202.3(BMP4):c.362A>G (p.His121Arg)	TTCGTGGYGGAAGCTCCTCACGG	Microphthalmia syndromic 6
606231280	NM_001287223.1(SCN11A):c.1142T>C (p.Ile381Thr)	CTTCAYTGTGGTCATTTTCCTGG, TTCAYTGTGGTCATTTTCCTGG G	Episodic pain syndrome, familial, 3
387906735	m.608A>G	TTCAGYGTATTGCTTTGAGGAGG	
199474663	m.3260A>G	TTAAGTTYTATGCGATTACCGGG	Cardiomyopathy with or without skeletal myopathy
104894962	NM_003413.3(ZIC3):c.1213A>G (p.Lys405Glu)	TGTGTTYGCGCAGGGAGCTCGGG, ATGTGTTYGCGCAGGGAGCTCG G	Heterotaxy, visceral, X-linked
796053181	NM_021007.2(SCN2A):c.1271T>C (p.Val424Ala)	TGTGGYGGCCATGGCCTATGAGG	not provided
267606788	NM_000129.3(F13A1):c.728T>C (p.Met243Thr)	TGTGAYGGACAGAGCACAAATGG	Factor xiii, a subunit, deficiency of
397514503	NM_003863.3(DPM2):c.68A>G (p.Tyr23Cys)	TGTAGYAGGTGAAGATGATCAGG	Congenital disorder of glycosylation type 1u
104893973	NM_000416.2(IFNGR1):c.260T>C (p.Ile87Thr)	TGTAATAYTTCTGATCATGTTGG	Disseminated atypical mycobacterial infection, Mycobacterium tuberculosis, susceptibility to
121908466	NM_005682.6(ADGRG1):c.263A>G (p.Tyr88Cys)	TGGYAGAGGCCCCTGGGGTCAGG	Polymicrogyria, bilateral frontoparietal
147952488	NM_002437.4(MPV17):c.186+2T>C	TGGYAAGTTCTCCCCTCAACAGG	Navajo neurohepatopathy
121909537	NM_001145.4(ANG):c.121A>G (p.Lys41Glu)	TGGTTYGGCATCATAGTGCTGGG, GTGGTTYGGCATCATAGTGCTG G	Amyotrophic lateral sclerosis type 9
121918489	NM_000141.4(FGFR2):c.1018T>C (p.Tyr340His)	TGGGGAAYATACGTGCTTGGCGG, GGGGAAYATACGTGCTTGGCGGG	Crouzon syndrome
121434463	m.12320A>G	GAGTYGCACCAAAATTTTTGGGG, GGAGTYGCACCAAAATTTTTGGG, TGGAGTYGCACCAAAATTTTTG G	Mitochondrial myopathy
121908046	NM_000403.3(GALE):c.101A>G (p.Asn34Ser)	TGGAAGYTATCGATGACCACAGG	UDPglucose-4-epimerase deficiency
431905512	NM_003764.3(STX11):c.173T>C (p.Leu58Pro)	TGCYGGTGGCCGACGTGAAGCGG	Hemophagocytic lymphohistiocytosis, familial, 4
121917905	NM_000124.3(ERCC6):c.2960T>C (p.Leu987Pro)	TGCYAAAAGACCCAAAACAAAGG	Cerebro-oculo-facio-skeletal syndrome
121918500	NM_000141.4(FGFR2):c.874A>G (p.Lys292Glu)	TGCTYGATCCACTGGATGTGGGG, GTGCTYGATCCACTGGATGTGGG, CGTGCTYGATCCACTGGATGTG G	Crouzon syndrome
60431989	NM_000053.3(ATP7B):c.3443T>C (p.Ile1148Thr)	TGCTGAYTGGAAACCGTGAGTGG	Wilson disease
78950939	NM_000250.1(MPO):c.518A>G (p.Tyr173Cys)	GTGCGGYATTTGTCCTGCTCCGG, TGCGGYATTTGTCCTGCTCCGG G	Myeloperoxidase deficiency
115677373	NM_201631.3(TGM5):c.763T>C (p.Trp255Arg)	TGCGGAGYGGACGGGCAGCGTGG	Peeling skin syndrome, acral type
5030804	NM_000551.3(VHL):c.233A>G (p.Asn78Ser)	GCGAYTGCAGAAGATGACCTGGG, TGCGAYTGCAGAAGATGACCTG G	Von Hippel-Lindau syndrome
397508328	NM_000492.3(CFTR):c.1A>G (p.Met1Val)	GCAYGGTCTCTCGGGCGCTGGGG, TGCAYGGTCTCTCGGGCGCTGGG , CTGCAYGGTCTCTCGGGCGCTGG	Cystic fibrosis

137853299	NM_000362.4(TIMP3):c.572A>G (p.Tyr191Cys)	TGCAGYAGCCGCCCTTCTGCCGG	Sorsby fundus dystrophy
121908549	NM_000334.4(SCN4A):c.3478A>G (p.Ile1160Val)	TGAYGGAGGGGATGGCGCCTAGG	
121909337	NM_001451.2(FOXF1):c.1138T>C (p.Ter380Arg)	TGATGYGAGGCTGCCGCCGCAGG	Alveolar capillary dysplasia with misalignment of pulmonary veins
281875320	NM_005359.5(SMAD4):c.1500A>G (p.Ile500Met)	TGAGYATGCATAAGCGACGAAGG	Myhre syndrome
730880132	NM_170707.3(LMNA):c.710T>C (p.Phe237Ser)	TGAGTYTGAGAGCCGGCTGGCGG	Primary dilated cardiomyopathy
281875322	NM_005359.5(SMAD4):c.1498A>G (p.Ile500Val)	TGAGTAYGCATAAGCGACGAAGG	Hereditary cancer-predisposing syndrome, Myhre syndrome
72556283	NM_000531.5(OTC):c.527A>G (p.Tyr176Cys)	TGAGGYAATCAGCCAGGATCTGG	not provided
74315311	NM_020435.3(GJC2):c.857T>C (p.Met286Thr)	TGAGAYGGCCCACCTGGGCTTGG, GAGAYGGCCCACCTGGGCTTGGG	Leukodystrophy, hypomyelinating, 2
121912495	NM_170707.3(LMNA):c.1139T>C (p.Leu380Ser)	TCTYGGAGGGCGAGGAGGAGAGG	Congenital muscular dystrophy, LMNA-related
128620184	NM_000061.2(BTK):c.1288A>G (p.Lys430Glu)	TCTYGATGGCCACGTCGTACTGG	X-linked agammaglobulinemia
118192252	NM_004519.3(KCNQ3):c.1403A>G (p.Asn468Ser)	TCTTTAYTGTTTAAGCCAACAGG	Benign familial neonatal seizures 2, not specified
121909142	NM_001300.5(KLF6):c.190T>C (p.Trp64Arg)	TCTGYGGACCAAAATCATTCTGG	
104895503	NM_001127255.1(NLRP7):c.2738A>G (p.Asn913Ser)	TCTGGYTGATACTCAAGTCCAGG	Hydatidiform mole
587783035	NM_000038.5(APC):c.1744-2A>G	TCCYAGTAAGAAACAGAATATGG	Familial adenomatous polyposis 1
72556289	NM_000531.5(OTC):c.541-2A>G	TCCYAAAAGGCACGGGATGAAGG	not provided
28937313	NM_005502.3(ABCA1):c.2804A>G (p.Asn935Ser)	TCCAYTGTGGCCCAGGAAGGAGG, CGCTCCAYTGTGGCCCAGGAAGG	Tangier disease
143246552	NM_001003811.1(TEX11):c.511A>G (p.Met171Val)	TCCAYGGTCAAGTCAGCCTCAGG, CCAYGGTCAAGTCAGCCTCAGGG	Spermatogenic failure, X-linked, 2
587776451	NM_002049.3(GATA1):c.2T>C (p.Met1Thr)	CTCCAYGGAGTTCCCTGGCCTGG, TCCAYGGAGTTCCCTGGCCTGGG, CCAYGGAGTTCCCTGGCCTGGGG	GATA-1-related thrombocytopenia with dyserythropoiesis
121908403	NM_021102.3(SPINT2):c.488A>G (p.Tyr163Cys)	TCCAYAGATGAAGTTATTGCAGG	Diarrhea 3, secretory sodium, congenital, syndromic
281874738	NM_000495.4(COL4A5):c.438+2T>C	CTCCAGYAAGTTATAAAATTTGG, TCCAGYAAGTTATAAAATTTGGG	Alport syndrome, X-linked recessive
730880279	NM_030653.3(DDX11):c.2271+2T>C	TCCAGGYGCGGGCGTCATGCTGG, CCAGGYGCGGGCGTCATGCTGGG	Warsaw breakage syndrome
28940272	NM_017890.4(VPS13B):c.8978A>G (p.Asn2993Ser)	TCAYTGATAAGCAGGGCCCAGGG, TTCAYTGATAAGCAGGGCCCAGG	Cohen syndrome, not specified
137852375	NM_000132.3(F8):c.5372T>C (p.Met1791Thr)	TCAYGGTGAGTTAAGGACAGTGG	Hereditary factor VIII deficiency disease
11567847	NM_021961.5(TEAD1):c.1261T>C (p.Tyr?His)	TCATATTYACAGGCTTGTAAAGG	
786203989	NM_016069.9(PAM16):c.226A>G (p.Asn76Asp)	CATAGTYCTGCAGAGGAGAGGGG, TCATAGTYCTGCAGAGGAGAGGG	Chondrodysplasia, megarbane-dagher-melki type
587776437	NC_012920.1:m.9478T>C	TCAGAAGYTTTTTTCTTCGCAGG	Leigh disease
121912474	NM_000424.3(KRT5):c.20T>C (p.Val7Ala)	TCAAGTGYGTCCTTCCGGAGCGG, CAAGTGYGTCCTTCCGGAGCGGG, AAGTGYGTCCTTCCGGAGCGGGG, AGTGYGTCCTTCCGGAGCGGGGG	Epidermolysis bullosa simplex, Koebner type
104886461	NM_020533.2(MCOLN1):c.406-2A>G	TACYGTGGGCAGAGAAGGGGAGG, AGGTACYGTGGGCAGAGAAGGGG, CAGGTACYGTGGGCAGAGAAGGG	Ganglioside sialidase deficiency

104894275	NM_000317.2(PTS):c.155A>G (p.Asn52Ser)	TAAYTGTGCCCATGGCCATTTGG	6-pyruvoyl-tetrahydropterin synthase deficiency
587777562	NM_015599.2(PGM3):c.737A>G (p.Asn246Ser)	TAAATGAYTGAGTTTGCCCTTGG	Immunodeficiency 23
121964906	NM_000027.3(AGA):c.916T>C (p.Cys306Arg)	GTTATAYGTGCCAATGTGACTGG	Aspartylglycosaminuria
28941769	NM_000356.3(TCOF1):c.149A>G (p.Tyr50Cys)	GTGTGTAYAGATGTCCAGAAGGG	Treacher collins syndrome 1
121434464	m.12297T>C	GTCYTAGGCCCCAAAAATTTTGG	Cardiomyopathy, mitochondrial
121908407	NM_054027.4(ANKH):c.143T>C (p.Met48Thr)	GTCGAGAYGCTGGCCAGCTACGG, TCGAGAYGCTGGCCAGCTACGGG	Chondrocalcinosis 2
59151893	NM_000422.2(KRT17):c.275A>G (p.Asn92Ser)	GTCAYTGAGGTTCTGCATGGTGG, GCGGTCAYTGAGGTTCTGCATGG	Pachyonychia congenita type 2
121909499	NM_002427.3(MMP13):c.272T>C (p.Met91Thr)	GTCAYGAAAAAGCCAAGATGCGG, TCAYGAAAAAGCCAAGATGCGG G	
61748478	NM_000552.3(VWF):c.2384A>G (p.Tyr795Cys)	GTCAYAGTTCTGGCACGTTTTGG	von Willebrand disease type 2N

387906889	NM_006796.2(AFG3L2):c.1847A>G (p.Tyr616Cys)	GTAYAGAGGTATTGTTCTTTTGG	Spastic ataxia 5, autosomal recessive
118203907	NM_000130.4(F5):c.5189A>G (p.Tyr1730Cys)	GTAGYAGGCCCAAGCCCGACAGG	Factor V deficiency
118203945	NM_013319.2(UBIAD1):c.305A>G (p.Asn102Ser)	GTAAGTGYTGACCAAATTACCGG	Schnyder crystalline corneal dystrophy
267607080	NM_005633.3(SOS1):c.1294T>C (p.Trp432Arg)	GGTYGGGAGGGAAAAGACATTGG	Noonan syndrome 4, Rasopathy
137852953	NM_012464.4(TLL1):c.1885A>G (p.Ile629Val)	GGTTAYGGTGCCGTTAAGTTTGG	Atrial septal defect 6
118203949	NM_013319.2(UBIAD1):c.695A>G (p.Asn232Ser)	GGTGTTGYTGGAATGGAGAATGG	Schnyder crystalline corneal dystrophy
137852952	NM_012464.4(TLL1):c.713T>C (p.Val238Ala)	GGGATTGYTGTTCATGAATTGGG	Atrial septal defect 6
41460449	m.3394T>C	GGCYATATACAACTACGCAAAGG	Leber optic atrophy
80357281	NM_007294.3(BRCA1):c.5291T>C (p.Leu1764Pro)	GGGCYAGAAATCTGTTGCTATGG, GGCYAGAAATCTGTTGCTATGGG	Familial cancer of breast, Breast-ovarian cancer, familial 1
5030764	NM_000174.4(GP9):c.182A>G (p.Asn61Ser)	GGCTGYTGTTGGCCAGCAGAAGG	Bernard-Soulier syndrome type C
72556282	NM_000531.5(OTC):c.526T>C (p.Tyr176His)	GGCTGATYACCTCACGCTCCAGG, GATYACCTCACGCTCCAGGTTGG	not provided
121913594	NM_000530.6(MPZ):c.242A>G (p.His81Arg)	GGCATAGYGGAAGATCTATGAGG	Charcot-Marie-Tooth disease type 1B
587777736	NM_017617.3(NOTCH1):c.1285T>C (p.Cys429Arg)	GGCAAGYGCATCAACACGCTGGG, GGGCAAGYGCATCAACACGCTGG	Adams-Oliver syndrome 1, Adams-Oliver syndrome 5
63750912	NM_016835.4(MAPT):c.1839T>C (p.Asn613=)	GGATAAYATCAAACACGTCCCGG, GATAAYATCAAACACGTCCCGG G	Frontotemporal dementia
121918075	NM_000371.3(TTR):c.401A>G (p.Tyr134Cys)	GGAGYAGGGGCTCAGCAGGGCGG, ATAGGAGYAGGGGCTCAGCAGGG	Amyloidogenic transthyretin amyloidosis
730882063	NM_004523.3(KIF11):c.2547+2T>C	GGAGGYAATAACTTTGTAAGTGG	Microcephaly with or without chorioretinopathy, lymphedema, or mental retardation
397516156	NM_000257.3(MYH7):c.2546T>C (p.Met849Thr)	GGAGAYGGCCTCCATGAAGGAGG	Primary familial hypertrophic cardiomyopathy,
118204430	NM_000035.3(ALDOB):c.442T>C (p.Trp148Arg)	GGAAGYGGCGTGCTGTGCTGAGG	Hereditary fructosuria
200198778	NM_013382.5(POMT2):c.1997A>G (p.Tyr666Cys)	GGAAGYAGTGGTGGAAGTAGAGG	Congenital muscular dystrophy, Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies, type A2, Muscular dystrophy, Congenital muscular dystrophy-dystroglycanopathy with mental retardation, type B2
754896795	NM_004006.2(DMD):c.6982A>T (p.Lys2328Ter)	GCTTTTYTTCAAGCTGCCCAAGG	Duchenne muscular dystrophy, Becker muscular dystrophy, Dilated cardiomyopathy 3B
148924904	NM_000546.5(TP53):c.488A>G (p.Tyr163Cys)	GCTTGYAGATGGCCATGGCGCGG	Hereditary cancer-predisposing syndrome

786204770	NM_016035.4(COQ4):c.155T>C (p.Leu52Ser)	GCTGTYGGCCGCCGGCTCCGCGG	COENZYME Q10 DEFICIENCY, PRIMARY, 7
121909520	NM_001100.3(ACTA1):c.350A>G (p.Asn117Ser)	CGGYTGGCCTTGGGATTGAGGGG, GCGGYTGGCCTTGGGATTGAGGG, CGCGGYTGGCCTTGGGATTGAGG	Nemaline myopathy 3
587776879	NM_004656.3(BAP1):c.438-2A>G	GCCYGGGGAAAAACAGAGTCAGG	Tumor predisposition syndrome
727504434	NM_000501.3(ELN):c.890-2A>G	GCCYGAAAACACAGCCACAGAGG	Supravalvar aortic stenosis
119455953	NM_000391.3(TPP1):c.1093T>C (p.Cys365Arg)	GCCGGGYGTTGGTCTGTCTCTGG	Ceroid lipofuscinosis, neuronal, 2
121964983	NM_000481.3(AMT):c.125A>G (p.His42Arg)	GCCAGGYGGAAGTCATAGAGCGG	Non-ketotic hyperglycinemia
121908300	NM_001005741.2(GBA):c.751T>C (p.Tyr251His)	GCCAGAYACTTTGTGAAGTAAGG, CCAGAYACTTTGTGAAGTAAGG	Gaucher disease, type 1
786205083	NM_003494.3(DYSF):c.3443-33A>G	GCCAGAGYGAGTGGCTGGAGTGG	Limb-girdle muscular dystrophy, type 2B
121908133	NM_175073.2(APTX):c.602A>G (p.His201Arg)	GCCAAYGGTAACGGGCCTTTGGG, AGCCAAYGGTAACGGGCCTTTGG	Adult onset ataxia with oculomotor apraxia
587777195	NM_005017.3(PCYT1A):c.571T>C (p.Phe191Leu)	GCATGYTTGCTCCAACACAGAGG	Spondylometaphyseal dysplasia with cone-rod dystrophy
431905520	NM_014714.3(IFT140):c.4078T>C (p.Cys1360Arg)	CAAGCAGYGTGAGCTGCTCCTGG, GCAGYGTGAGCTGCTCCTGGAGG	Renal dysplasia, retinal pigmentary dystrophy, cerebellar ataxia and skeletal dysplasia
121912889	NM_001844.4(COL2A1):c.4172A>G (p.Tyr1391Cys)	GCAGTGGYAGGTGATGTTCTGGG	Spondyloperipheral dysplasia, Platyspondylic lethal skeletal dysplasia Torrance type
137854492	NM_001363.4(DKC1):c.1069A>G (p.Thr357Ala)	GCAGGYAGAGATGACCGCTGTGG	Dyskeratosis congenita X-linked
121434362	NM_152783.4(D2HGDH):c.1315A>G (p.Asn439Asp)	GCAGGTYACCATCTCCTGGAGGG, TGCAGGTYACCATCTCCTGGAGG	D-2-hydroxyglutaric aciduria 1
80338732	NM_002764.3(PRPS1):c.344T>C (p.Met115Thr)	GCAAATAYGCTATCTGTAGCAGG	Charcot-Marie-Tooth disease, X-linked recessive, type 5
387906675	NM_000313.3(PROS1):c.701A>G (p.Tyr234Cys)	GATTAYATCTGTAGCCTTCGGGG, AGATTAYATCTGTAGCCTTCGGG, GAGATTAYATCTGTAGCCTTCGG	Thrombophilia due to protein S deficiency, autosomal recessive

28935478	NM_000061.2(BTK):c.1082A>G (p.Tyr361Cys)	GATGGYAGTTAATGAGCTCAGGG, TGATGGYAGTTAATGAGCTCAGG	
201777056	NM_005050.3(ABCD4):c.956A>G (p.Tyr319Cys)	GATGAGGYAGATGCACACAAAGG	METHYLMALONIC ACIDURIA AND HOMOCYSTINURIA, cblJ
121918528	NM_000098.2(CPT2):c.359A>G (p.Tyr120Cys)	GATAGGYACATATCAAACCAGGG, AGATAGGYACATATCAAACCAGG	Carnitine palmitoyltransferase II deficiency, infantile
267607014	NM_002942.4(ROBO2):c.2834T>C (p.Ile945Thr)	GAGAYTGGAAATTTTGGCCGTGG	Vesicoureteral reflux 2
281865192	NM_025114.3(CEP290):c.2991+1655A>G	GATAYTCACAATTACAACTGGGG, AGATAYTCACAATTACAACTGGG, GAGATAYTCACAATTACAACTG	Leber congenital amaurosis 10
386833492	NM_000112.3(SLC26A2):c.-26+2T>C	GAGAGGYGAGAAGAGGGAAGCGG	Diastrophic dysplasia
587779773	NM_001101.3(ACTB):c.356T>C (p.Met119Thr)	GAGAAGAYGACCCAGGTGAGTGG	Baraitser-Winter syndrome 1
121913512	NM_000222.2(KIT):c.1924A>G (p.Lys642Glu)	GACTTYGAGTTCAGACATGAGGG, GGACTTYGAGTTCAGACATGAGG	
28939072	NM_006329.3(FBLN5):c.506T>C (p.Ile169Thr)	GACAYTGATGAATGTCGCTATGG	Age-related macular degeneration 3
104894248	NM_000525.3(KCNJ11):c.776A>G (p.His259Arg)	GACAYGGTAGATGATCAGCGGGG, TGACAYGGTAGATGATCAGCGGG, ATGACAYGGTAGATGATCAGCGG	Islet cell hyperplasia
387907132	NM_016464.4(TMEM138):c.287A>G (p.His96Arg)	GACAYGAAGGGAGATGCTGAGGG, AGACAYGAAGGGAGATGCTGAGG	Joubert syndrome 16
121918170	NM_000275.2(OCA2):c.1465A>G (p.Asn489Asp)	GACATYTGGAGGGTCCCCGATGG	Tyrosinase-positive oculocutaneous albinism
122467173	NM_014009.3(FOXP3):c.970T>C (p.Phe324Leu)	GACAGAGYTCCTCCACAACATGG	Insulin-dependent diabetes mellitus secretory diarrhea syndrome
137852268	NM_000133.3(F9):c.1328T>C (p.Ile443Thr)	GAAYATATACCAAGGTATCCCGG	Hereditary factor IX deficiency disease
149054177	NM_001999.3(FBN2):c.3740T>C (p.Met1247Thr)	GAATGTAYGATAATGAACGGAGG	not specified, Macular degeneration, early-onset

137854488	NM_212482.1(FN1):c.2918A>G (p.Tyr973Cys)	GAAGTAAYAGGTGACCCCAGGGG	Glomerulopathy with fibronectin deposits 2
786204027	NM_005957.4(MTHFR):c.1530+2T >C	GAAGGYGTGGTAGGGAGGCACGG, AAGGYGTGGTAGGGAGGCACGGG, AGGYGTGGTAGGGAGGCACGGGG	Homocysteinemia due to MTHFR deficiency
104894223	NM_012193.3(FZD4):c.766A>G (p.Ile256Val)	GAAATAYGATGGGGCGCTCAGGG, AGAAATAYGATGGGGCGCTCAGG	Retinopathy of prematurity
137854474	NM_000138.4(FBN1):c.3793T>C (p.Cys1265Arg)	CTTGYGTTATGATGGATTCATGG	Marfan syndrome
587784418	NM_006306.3(SMC1A):c.3254A>G (p.Tyr1085Cys)	CTTAYAGATCTCATCAATGTTGG	Congenital muscular hypertrophy-cerebral syndrome
81002805	NM_000059.3(BRCA2):c.316+2T> C	CTTAGGYAAGTAATGCAATATGG	Familial cancer of breast, Breast-ovarian cancer, familial 2, Hereditary cancer-predisposing syndrome
121909653	NM_182925.4(FLT4):c.3104A>G (p.His1035Arg)	CTGYGGATGCACTGGGGTGCGGG, TCTGYGGATGCACTGGGGTGCGG	
786205107	NM_031226.2(CYP19A1):c.743+2T >C	CTGTGYAAGTAATACAACTTTGG	Aromatase deficiency
587777037	NM_001283009.1(RTEL1):c.3730T>C (p.Cys1244Arg)	CTGTGTGYGCCAGGGCTGTGGGG	Dyskeratosis congenita, autosomal recessive, 5
794728380	NM_000238.3(KCNH2):c.1945+6T >C	CTGTGAGYGTGCCCAGGGGCGGG, TGAGYGTGCCCAGGGGCGGGCGG	Cardiac arrhythmia
267607987	NM_000251.2(MSH2):c.2005+2T> C	CTGGYAAAAAACCTGGTTTTTGG, TGGYAAAAAACCTGGTTTTTGG G	Hereditary Nonpolyposis Colorectal Neoplasms
397509397	NM_006876.2(B4GAT1):c.1168A>G (p.Asn390Asp)	TGATYTTCAGCCTCCTTTTGGGG, CTGATYTTCAGCCTCCTTTTGGG, GCTGATYTTCAGCCTCCTTTTGG	Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies, type A13
121918381	NM_000040.1(APOC3):c.280A>G (p.Thr94Ala)	CTGAAGYTGGTCTGACCTCAGGG, GCTGAAGYTGGTCTGACCTCAGG	
104894919	NM_001015877.1(PHF6):c.769A>G (p.Arg257Gly)	CTCYTGATGTTGTTGTGAGCTGG	Borjeson-Forssman-Lehmann syndrome
267606869	NM_005144.4(HR):c.-218A>G	CTCYAGGGCCGCAGGTTGGAGGG, GCTCYAGGGCCGCAGGTTGGAGG, GGCGCTCYAGGGCCGCAGGTTGG	Marie Unna hereditary hypotrichosis 1
139732572	NM_000146.3(FTL):c.1A>G (p.Met1Val)	CTCAYGGTTGGTTGGCAAGAAGG	L-ferritin deficiency
397515418	NM_018486.2(HDAC8):c.1001A>G (p.His334Arg)	CTCAYGATCTGGGATCTCAGAGG	Cornelia de Lange syndrome 5
372395294	NM_198056.2(SCN5A):c.1247A>G (p.Tyr416Cys)	CTCAYAGGCCATTGCGACCACGG	not provided
104895304	NM_000431.3(MVK):c.803T>C (p.Ile268Thr)	CTCAAYAGATGCCATCTCCCTGG	Hyperimmunoglobulin D with periodic fever, Mevalonic aciduria
587777188	NM_001165899.1(PDE4D):c.1850T>C (p.Ile617Thr)	CTATAYTGTTCATCCCCTCTGGG, ACTATAYTGTTCATCCCCTCTGG	Acrodysostosis 2, with or without hormone resistance
398123026	NM_003867.3(FGF17):c.560A>G (p.Asn187Ser)	CGTGGYTGGGGAAGGGCAGCTGG	Hypogonadotropic hypogonadism 20 with or without anosmia

121964924	NM_001385.2(DPYS):c.1078T>C (p.Trp360Arg)	CGTAATAYGGGAAAAAGGCGTGG, AATAYGGGAAAAAGGCGTGGTGG, ATAYGGGAAAAAGGCGTGGTGGG	Dihydropyrimidinase deficiency
587777301	NM_199189.2(MATR3):c.1864A>G (p.Thr622Ala)	CGGYTGAACTCTCAGTCTTCTGG	Myopathy, distal, 2
200238879	NM_000527.4(LDLR):c.694+2T>C	ACTGCGGYATGGGCGGGGCCAGG, CTGCGGYATGGGCGGGGCCAGGG, CGGYATGGGCGGGGCCAGGGTGG	Familial hypercholesterolemia
142951029	NM_145046.4(CALR3):c.245A>G (p.Lys82Arg)	CGGTYTGAAGCGTGCAGAGATGG	Arrhythmogenic right ventricular cardiomyopathy, Familial hypertrophic cardiomyopathy 19, Hypertrophic cardiomyopathy
786200953	NM_006785.3(MALT1):c.1019-2A>G	CGCYTTGAAAAAAAAAGAAAGGG, TCGCYTTGAAAAAAAAAGAAAG	Combined immunodeficiency
120074192	NM_000218.2(KCNQ1):c.418A>G (p.Ser140Gly)	CGCYGAAGATGAGGCAGACCAGG	Atrial fibrillation, familial, 3, Atrial fibrillation
267606887	NM_005957.4(MTHFR):c.971A>G (p.Asn324Ser)	CGCGGYTGAGGGTGTAGAAGTGG	Homocystinuria due to MTHFR deficiency
118192117	NM_000540.2(RYR1):c.1205T>C (p.Met402Thr)	CGCAYGATCCACAGCACCAATGG	Congenital myopathy with fiber type disproportion, Central core disease
199473625	NM_198056.2(SCN5A):c.4978A>G (p.Ile1660Val)	CGAYGTTGAAGAGGGCAGGCAGG, AGCCCGAYGTTGAAGAGGGCAGG	Brugada syndrome
794726865	NM_000921.4(PDE3A):c.1333A>G (p.Thr445Ala)	CGAGGYGGTGGTGGTCCAAGTGG	Brachydactyly with hypertension
606231254	NM_005740.2(DNAL4):c.153+2T>C	CGAGGYATTGCCAGCAGTGCAGG	Mirror movements 3
786204826	NM_004771.3(MMP20):c.611A>G (p.His204Arg)	CGAAAYGTGTATCTCCTCCCAGG	Amelogenesis imperfecta, hypomaturation type, IIA2

796053139	NM_021007.2(SCN2A):c.4308+2T>C	CGAAATGYAAGTCTAGTTAGAGG, GAAATGYAAGTCTAGTTAGAGG	not provided
137854494	NM_005502.3(ABCA1):c.4429T>C (p.Cys1477Arg)	CCTGTGYGTCCCCCAGGGGCAGG, CTGTGYGTCCCCCAGGGGCAGGG, TGTGYGTCCCCCAGGGGCAGGGG, GTGYGTCCCCCAGGGGCAGGGGG	Tangier disease
786205144	NM_001103.3(ACTN2):c.683T>C (p.Met228Thr)	CCTAAAAYGTTGGATGCTGAAGG	Dilated cardiomyopathy 1AA
199919568	NM_007254.3(PNKP):c.1029+2T>C	CCGGYGAGGCCCTGGGGCGGGGG, TCCGGYGAGGCCCTGGGGCGGGG, ATCCGGYGAGGCCCTGGGGCGGG, GATCCGGYGAGGCCCTGGGGCGG	not provided
28939079	NM_018965.3(TREM2):c.401A>G (p.Asp134Gly)	TGAYCCAGGGGGTCTATGGGAGG, CGGTGAYCCAGGGGGTCTATGGG, CCGGTGAYCCAGGGGGTCTATGG	Polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy
193302855	NM_032520.4(GNPTG):c.610-2A>G	CCCYGAAGGTGGAGGATGCAGGG, GCCCYGAAGGTGGAGGATGCAGG	Mucolipidosis III Gamma
111033708	NM_000155.3(GALT):c.499T>C (p.Trp167Arg)	CCCTYGGGTGCAGGTTTGTGAGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
28933378	NM_000174.4(GP9):c.70T>C (p.Cys24Arg)	CCCAYGTACCTGCCGCGCCCTGG	Bernard Soulier syndrome, Bernard-Soulier syndrome type C
364897	NM_000157.3(GBA):c.680A>G (p.Asn227Ser)	CCAYTGGTCTTGAGCCAAGTGGG, TCCAYTGGTCTTGAGCCAAGTGG	Gaucher disease, Subacute neuronopathic Gaucher disease, Gaucher disease, type 1
796052551	NM_000833.4(GRIN2A):c.2449A>G (p.Met817Val)	CCAYGTTGTCAATGTCCAGCTGG	not provided
63751006	NM_002087.3(GRN):c.2T>C (p.Met1Thr)	CCAYGTGGACCCTGGTGAGCTGG	Frontotemporal dementia, ubiquitin-positive
786203997	NM_001031.4(RPS28):c.1A>G (p.Met1Val)	TGTCCAYGATGGCGGCGCGGCGG, CCAYGATGGCGGCGCGGCGGCGG	Diamond-Blackfan anemia with microtia and cleft palate
121908595	NM_002755.3(MAP2K1):c.389A>G (p.Tyr130Cys)	CCAYAGAAGCCCACGATGTACGG	Cardiofaciocutaneous syndrome 3, Rasopathy
398122910	NM_000431.3(MVK):c.1039+2T>C	CCAGGYATCCCGGGGGTAGGTGG, CAGGYATCCCGGGGGTAGGTGGG	Porokeratosis, disseminated superficial actinic 1
119474039	NM_020365.4(EIF2B3):c.1037T>C (p.Ile346Thr)	CCAGAYTGTCAGCAAACACCTGG	Leukoencephalopathy with vanishing white matter
587777866	NM_000076.2(CDKN1C):c.*5+2T>C	CCAAGYGAGTACAGCGCACCTGG, CAAGYGAGTACAGCGCACCTGGG, AAGYGAGTACAGCGCACCTGGGG	Beckwith-Wiedemann syndrome
121918530	NM_005587.2(MEF2A):c.788A>G (p.Asn263Ser)	AGAYTACCACCACCTGGTGGAGG, CCAAGAYTACCACCACCTGGTGG	
483352818	NM_000211.4(ITGB2):c.1877+2T>C	CATGYGAGTGCAGGCGGAGCAGG	Leukocyte adhesion deficiency type 1
460184	NM_000186.3(CFH):c.3590T>C (p.Val1197Ala)	CAGYTGAATTTGTGTGTAAACGG	Atypical hemolytic-uremic syndrome 1
121908423	NM_004795.3(KL):c.578A>G (p.His193Arg)	CAGYGGTACAGGGTGACCACGGG, CCAGYGGTACAGGGTGACCACGG	
281860300	NM_005247.2(FGF3):c.146A>G (p.Tyr49Cys)	CAGYAGAGCTTGCGGCGCCGGGG, GCAGYAGAGCTTGCGGCGCCGGG, CGCAGYAGAGCTTGCGGCGCCGG	Deafness with labyrinthine aplasia microtia and microdontia (LAMM)
28935488	NM_000169.2(GLA):c.806T>C (p.Val269Ala)	CAGTTAGYGATTGGCAACTTTGG	Fabry disease

587776514	NM_173560.3(RFX6):c.380+2T>C	CAGTGGYGAGACTCGCCCGCAGG, AGTGGYGAGACTCGCCCGCAGGG	Mitchell-Riley syndrome
104894117	NM_178138.4(LHX3):c.332A>G (p.Tyr111Cys)	CAGGTGGYACACGAAGTCCTGGG	Pituitary hormone deficiency, combined 3
34878913	NM_000184.2(HBG2):c.125T>C (p.Phe42Ser)	CAGAGGTYCTTTGACAGCTTTGG	Cyanosis, transient neonatal
120074124	NM_000543.4(SMPD1):c.911T>C (p.Leu304Pro)	AGCACYTGTGAGGAAGTTCCTGG, GCACYTGTGAGGAAGTTCCTGGG, CACYTGTGAGGAAGTTCCTGGGG	Sphingomyelin/cholesterol lipidosis, Niemann-Pick disease, type A, Niemann-Pick disease, type B
281860272	NM_005211.3(CSF1R):c.2320-2A>G	CACYGAGGGAAAGCACTGCAGGG, GCACYGAGGGAAAGCACTGCAGG	Hereditary diffuse leukoencephalopathy with spheroids
128624216	NM_000033.3(ABCD1):c.443A>G (p.Asn148Ser)	CACTGYTGACGAAGGTAGCAGGG, GCACTGYTGACGAAGGTAGCAGG	Adrenoleukodystrophy
398124257	NM_012463.3(ATP6V0A2):c.825+2T>C	CACTGYGAGTAAGCTGGAAGTGG	Cutis laxa with osteodystrophy
267606679	NM_004183.3(BEST1):c.704T>C (p.Val235Ala)	CACTGGYGTATACACAGGTGAGG	Vitreoretinochoroidopathy dominant
397514518	NM_000344.3(SMN1):c.388T>C (p.Tyr130His)	CACTGGAYATGGAAATAGAGAGG	Kugelberg-Welander disease

143946794	NM_001946.3(DUSP6):c.566A>G (p.Asn189Ser)	CACTAYTGGGGTCTCGGTCAAGG	Hypogonadotropic hypogonadism 19 with or without anosmia
397516076	NM_000256.3(MYBPC3):c.821+2T >C	GCACGYGAGTGGCCATCCTCAGG, CACGYGAGTGGCCATCCTCAGGG	Familial hypertrophic cardiomyopathy 4, not specified
149977726	NM_001257988.1(TYMP):c.665A>G (p.Lys222Arg)	CACGAGTYTCTTACTGAGAATGG, GAGTYTCTTACTGAGAATGGAGG	
121917770	NM_003361.3(UMOD):c.383A>G (p.Asn128Ser)	CACAYTGACACATGTGGCCAGGG, CCACAYTGACACATGTGGCCAGG	Familial juvenile gout
121909008	NM_000492.3(CFTR):c.2738A>G (p.Tyr913Cys)	CACATAAYACGAACTGGTGCTGG	Cystic fibrosis
137852819	NM_003688.3(CASK):c.2740T>C (p.Trp914Arg)	CACAGYGGGTCCCTGTCTCCTGG, ACAGYGGGTCCCTGTCTCCTGGG	FG syndrome 4
74315320	NM_024009.2(GJB3):c.421A>G (p.Ile141Val)	CAAYGATGAGCTTGAAGATGAGG	Deafness, autosomal recessive
80356747	NM_001701.3(BAAT):c.967A>G (p.Ile323Val)	CAAYGAAGAGGAATTGCCCCTGG	Atypical hemolytic-uremic syndrome 1
180177324	NM_012203.1(GRHPR):c.934A>G (p.Asn312Asp)	CAAGTYGTTAGCTGCCAACAAGG	Primary hyperoxaluria, type II
281860274	NM_005211.3(CSF1R):c.2381T>C (p.Ile794Thr)	CAAGAYTGGGGACTTCGGGCTGG	Hereditary diffuse leukoencephalopathy with spheroids
398122908	NM_005334.2(HCFC1):c.-970T>C	CAAGAYGGCGGCTCCCAGGGAGG	Mental retardation 3, X-linked
548076633	NM_002693.2(POLG):c.3470A>G (p.Asn1157Ser)	CAAGAGGYTGGTGATCTGCAAGG	not provided
120074146	NM_000019.3(ACAT1):c.935T>C (p.Ile312Thr)	CAAGAAYAGTAGGTAAGGCCAGG	Deficiency of acetyl-CoA acetyltransferase
397514489	NM_005340.6(HINT1):c.250T>C (p.Cys84Arg)	CAAGAAAYGTGCTGCTGATCTGG, AAGAAAYGTGCTGCTGATCTGGG	Gamstorp-Wohlfart syndrome
587783539	NM_178151.2(DCX):c.2T>C (p.Met1Thr)	CAAAATAYGGAACTTGATTTTGG	Heterotopia
104894765	NM_005448.2(BMP15):c.704A>G (p.Tyr235Cys)	ATTGAAAYAGAGTAACAAGAAGG	Ovarian dysgenesis 2
137852429	NM_000132.3(F8):c.1892A>G (p.Asn631Ser)	ATGYTGGAGGCTTGGAACTCTGG	Hereditary factor VIII deficiency disease
72558441	NM_000531.5(OTC):c.779T>C (p.Leu260Ser)	ATGTATYAATTACAGACACTTGG	not provided
398123765	NM_003494.3(DYSF):c.1284+2T>C	ATGGYAAGGAGCAAGGGAGCAGG	Limb-girdle muscular dystrophy, type 2B
387906924	NM_020191.2(MRPS22):c.644T>C (p.Leu215Pro)	ATCYTAGGGTAAGGTGACTTAGG	Combined oxidative phosphorylation deficiency 5
397518039	NM_206933.2(USH2A):c.8559-2A>G	ATCYAAAGCAAAAGACAAGCAGG	Retinitis pigmentosa, Usher syndrome, type 2A
5742905	NM_000071.2(CBS):c.833T>C (p.Ile278Thr)	ATCAYTGGGGTGGATCCCGAAGG, TCAYTGGGGTGGATCCCGAAGGG	Homocystinuria due to CBS deficiency, Homocystinuria, pyridoxine-responsive
397507473	NM_004333.4(BRAF):c.1403T>C (p.Phe468Ser)	ATCATYTGGAACAGTCTACAAGG, TCATYTGGAACAGTCTACAAGG	Cardiofaciocutaneous syndrome, Rasopathy
786204056	NM_000264.3(PTCH1):c.3168+2T>C	ATCATTGYGAGTGTATTATAAGG, TCATTGYGAGTGTATTATAAGGG, CATTGYGAGTGTATTATAAGGG	Gorlin syndrome
72558484	NM_000531.5(OTC):c.1005+2T>C	ATCATGGYAAGCAAGAAACAAGG	not provided
199473074	NM_000335.4(SCN5A):c.688A>G (p.Ile230Val)	ATAYAGTTTTCAGGGCCCGGAGG, CTGATAYAGTTTTCAGGGCCCGG	Brugada syndrome
111033273	NM_206933.2(USH2A):c.1606T>C (p.Cys536Arg)	ATATAGAYGCCTCTGCTCCCAGG	Usher syndrome, type 2A
72556290	NM_000531.5(OTC):c.542A>G (p.Glu181Gly)	ATAGTGTYCCTAAAAGGCACGGG	not provided
121918711	NM_004612.3(TGFBR1):c.1199A>G (p.Asp400Gly)	ATAGATGYCAGCACGTTTGAAGG	Loeys-Dietz syndrome 1
104886288	NM_000495.4(COL4A5):c.4699T>C (p.Cys1567Arg)	AGTAYGTGAAGCTCCAGCTGTGG	Alport syndrome, X-linked recessive

144637717	NM_016725.2(FOLR1):c.493+2T>C	CTTCAGGYGAGGGCTGGGGTGGG, AGGYGAGGGCTGGGGTGGGCAGG	not provided
72558492	NM_000531.5(OTC):c.1034A>G (p.Tyr345Cys)	AGGTGAGYAATCTGTCAGCAGGG	not provided
62638745	NM_000121.3(EPOR):c.1460A>G (p.Asn487Ser)	AGGGYTGGAGTAGGGGCCATCGG	Acute myeloid leukemia, M6 type, Familial erythrocytosis, 1

387907021	NM_031427.3(DNAL1):c.449A>G (p.Asn150Ser)	AGGGAYTGCCTACAAACACCAGG	Kartagener syndrome, Ciliary dyskinesia, primary, 16
397514488	NM_001161581.1(POC1A):c.398T>C (p.Leu133Pro)	AGCYGTGGGACAAGAGCAGCCGG	Short stature, onychodysplasia, facial dysmorphism, and hypotrichosis
154774633	NM_017882.2(CLN6):c.200T>C (p.Leu67Pro)	AGCYGGTATTCCCTCTCGAGTGG	Adult neuronal ceroid lipofuscinosis
111033700	NM_000155.3(GALT):c.482T>C (p.Leu161Pro)	AGCYGGGTGCCCAGTACCCTTGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
128621198	NM_000061.2(BTK):c.1223T>C (p.Leu408Pro)	GAGCYGGGGACTGGACAATTTGG, AGCYGGGGACTGGACAATTTGGG	X-linked agammaglobulinemia
137852611	NM_000211.4(ITGB2):c.446T>C (p.Leu149Pro)	AGCYAGGTGGCGACCTGCTCCGG	Leukocyte adhesion deficiency
121908838	NM_003722.4(TP63):c.697A>G (p.Lys233Glu)	AGCTTYTTTGTAGACAGGCATGG	Split-hand/foot malformation 4
397515869	NM_000169.2(GLA):c.1153A>G (p.Thr385Ala)	AGCTGTGYGATGAAGCAGGCAGG	not specified
118204064	NM_000237.2(LPL):c.548A>G (p.Asp183Gly)	GCTGGAYCGAGGCCTTAAAAGGG, AGCTGGAYCGAGGCCTTAAAAGG	Hyperlipoproteinemia, type I
128620186	NM_000061.2(BTK):c.2T>C (p.Met1Thr)	AGCTAYGGCCGCAGTGATTCTGG	X-linked agammaglobulinemia
786204132	NM_014946.3(SPAST):c.1165A>G (p.Thr389Ala)	ATTGYCTTCCCATTCCCAGGTGG, AGCATTGYCTTCCCATTCCCAGG	Spastic paraplegia 4, autosomal dominant
199473661	NM_000218.2(KCNQ1):c.550T>C (p.Tyr184His)	CAGCAAGBACGTGGGCCTCTGGG, AGCAAGBACGTGGGCCTCTGGGG, GCAAGBACGTGGGCCTCTGGGGG	Congenital long QT syndrome, Cardiac arrhythmia
387907129	NM_024599.5(RHBDF2):c.557T>C (p.Ile186Thr)	AGAYTGTGGATCCGCTGGCCCGG	Howel-Evans syndrome
387906702	NM_006306.3(SMC1A):c.2351T>C (p.Ile784Thr)	AGAYTGGTGTGCGCAACATCCGG	Congenital muscular hypertrophy-cerebral syndrome
193929348	NM_000525.3(KCNJ11):c.544A>G (p.Ile182Val)	AGAYGAGGGTCTCAGCCCTGCGG	Permanent neonatal diabetes mellitus
121908934	NM_004086.2(COCH):c.1535T>C (p.Met512Thr)	AGATAYGGCTTCTAAACCGAAGG	Deafness, autosomal dominant 9
397514377	NM_000060.3(BTD):c.641A>G (p.Asn214Ser)	AGAGGYTGTGTTTACGGTAGCGG	Biotinidase deficiency
72552295	NM_000531.5(OTC):c.2T>C (p.Met1Thr)	AGAAGAYGCTGTTTAATCTGAGG	not provided
201893545	NM_016247.3(IMPG2):c.370T>C (p.Phe124Leu)	ACTYTTTGGGATCGACTTCCTGG	Macular dystrophy, vitelliform, 5
121434469	m.4290T>C	ACTYTGATAGAGTAAATAATAGG	
121918733	NM_006920.4(SCN1A):c.269T>C (p.Phe90Ser)	ACTTYTATAGTATTGAATAAAGG, CTTYTATAGTATTGAATAAAGG G	Severe myoclonic epilepsy in infancy
121434471	m.4291T>C	ACTTYGATAGAGTAAATAATAGG	Hypertension, hypercholesterolemia, and hypomagnesemia, mitochondrial
606231289	NM_001302946.1(TRNT1):c.497T>C (p.Leu166Ser)	ACTTYATTTGACTACTTTAATGG	Sideroblastic anemia with B-cell immunodeficiency, periodic fevers, and developmental delay
63750067	NM_000517.4(HBA2):c.*92A>G	CTTYATTCAAAGACCAGGAAGGG, ACTTYATTCAAAGACCAGGAAG G	Hemoglobin H disease, nondeletional
121918734	NM_006920.4(SCN1A):c.272T>C (p.Ile91Thr)	ACTTTTAYAGTATTGAATAAAGG, CTTTTAYAGTATTGAATAAAGG G	Severe myoclonic epilepsy in infancy
137854557	NM_000267.3(NF1):c.1466A>G (p.Tyr489Cys)	ACTTAYAGCTTCTTGTCTCCAGG	Neurofibromatosis, type 1
397514626	NM_018344.5(SLC29A3):c.607T>C (p.Ser203Pro)	ACTGATAYCAGGTGAGAGCCAGG, CTGATAYCAGGTGAGAGCCAGGG	Histiocytosis-lymphadenopathy plus syndrome
118204440	NM_000512.4(GALNS):c.1460A>G (p.Asn487Ser)	ACGYTGAGCTGGGGCTGCGCGGG, CACGYTGAGCTGGGGCTGCGCGG	Mucopolysaccharidosis, MPS-IV-A
587776843	NG_012088.1:g.2209A>G	ACCYTATGATCCGCCCGCCTTGG	
137853033	NM_001080463.1(DYNC2H1):c.4610A>G (p.Gln1537Arg)	ACCYGTGAAGGGAACAGAGATGG	Short-rib thoracic dysplasia 3 with or without polydactyly
28933698	NM_000435.2(NOTCH3):c.1363T>C (p.Cys455Arg)	TTCACCYGTATCTGTATGGCAGG, ACCYGTATCTGTATGGCAGGTGG	Cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy
587776766	NM_000463.2(UGT1A1):c.1085-2A>G	ACCYGAGATGCAAAATAGGGAGG, GTGACCYGAGATGCAAAATAGGG, GGTGACCYGAGATGCAAAATAGG	Crigler Najjar syndrome, type 1
587781628	NM_001128425.1(MUTYH):c.1187-2A>G	ACCYGAGAGGGAGGGCAGCCAGG	Hereditary cancer-predisposing syndrome, Carcinoma of colon
61755817	NM_000322.4(PRPH2):c.736T>C (p.Trp246Arg)	ACCTGYGGGTGCGTGGCTGCAGG, CCTGYGGGTGCGTGGCTGCAGGG	Retinitis pigmentosa

121909184	NM_001089.2(ABCA3):c.1702A>G (p.Asn568Asp)	ACCGTYGTGGCCCAGCAGGACGG	Surfactant metabolism dysfunction, pulmonary, 3
121434466	m.4269A>G	ACAYATTTCTTAGGTTTGAGGGG, GACAYATTTCTTAGGTTTGAGGG, AGACAYATTTCTTAGGTTTGAGG	
794726768	NM_001165963.1(SCN1A):c.1048A>G (p.Met350Val)	ACAYATATCCCTCTGGACATTGG	Severe myoclonic epilepsy in infancy
28934876	NM_001382.3(DPAGT1):c.509A>G (p.Tyr170Cys)	ACAYAGTACAGGATTCCTGCGGG, GACAYAGTACAGGATTCCTGCGG	Congenital disorder of glycosylation type 1J
104894749	NM_000054.4(AVPR2):c.614A>G (p.Tyr205Cys)	ACAYAGGTGCGACGGCCCCAGGG, GACAYAGGTGCGACGGCCCCAGG	Nephrogenic diabetes insipidus, Nephrogenic diabetes insipidus, X-linked
128621205	NM_000061.2(BTK):c.1741T>C (p.Trp581Arg)	ACATTYGGGCTTTTGGTAAGTGG	X-linked agammaglobulinemia
28940892	NM_000529.2(MC2R):c.761A>G (p.Tyr254Cys)	ACATGYAGCAGGCGCAGTAGGGG, GACATGYAGCAGGCGCAGTAGGG, AGACATGYAGCAGGCGCAGTAGG	ACTH resistance
794726844	NM_001165963.1(SCN1A):c.1046A>G (p.Tyr349Cys)	ACATAYATCCCTCTGGACATTGG	Severe myoclonic epilepsy in infancy
587783083	NM_003159.2(CDKL5):c.449A>G (p.Lys150Arg)	ACAGTYTTAGGACATCATTGTGG	not provided
397514651	NM_000108.4(DLD):c.140T>C (p.Ile47Thr)	ACAGTTAYAGGTTCTGGTCCTGG, GTTAYAGGTTCTGGTCCTGGAGG	Maple syrup urine disease, type 3
794727060	NM_001848.2(COL6A1):c.957+2T >C	ACAAGGYGAGCGTGGGCTGCTGG, CAAGGYGAGCGTGGGCTGCTGGG	Ullrich congenital muscular dystrophy, Bethlem myopathy
72554346	NM_000531.5(OTC):c.284T>C (p.Leu95Ser)	ACAAGATYGTCTACAGAAACAGG	not provided
483353031	NM_002136.2(HNRNPA1):c.841T>C (p.Phe281Leu)	AATYTTGGAGGCAGAAGCTCTGG	Chronic progressive multiple sclerosis
104894271	NM_000315.2(PTH):c.52T>C (p.Cys18Arg)	AATTYGTTTTCTTACAAAATCGG	Hypoparathyroidism familial isolated
267608260	NM_015599.2(PGM3):c.248T>C (p.Leu83Ser)	AATGTYGGCACCATCCTGGGAGG	Immunodeficiency 23
267606900	NM_018109.3(MTPAP):c.1432A>G (p.Asn478Asp)	AATGGATYCTGAATGTACAGAGG	Ataxia, spastic, 4, autosomal recessive
796053169	NM_021007.2(SCN2A):c.387-2A>G	AATAAAGYAGAATATCGTCAAGG	not provided
104894937	NM_000116.4(TAZ):c.352T>C (p.Cys118Arg)	AAGYGTGTGCCTGTGTGCCGAGG	3-Methylglutaconic aciduria type 2
104893911	NM_001018077.1(NR3C1):c.1712T>C (p.Val571Ala)	AAGYGATTGCAGCAGTGAAATGG	Pseudohermaphroditism, female, with hypokalemia, due to glucocorticoid resistance
397514472	NM_004813.2(PEX16):c.992A>G (p.Tyr331Cys)	AAGYAGATTTTCTGCCAGGTGGG, GAAGYAGATTTTCTGCCAGGTGG, GTAGAAGYAGATTTTCTGCCAGG	Peroxisome biogenesis disorder 8B
121918407	NM_001083112.2(GPD2):c.1904T>C (p.Phe635Ser)	AAGTYTGATGCAGACCAGAAAGG	Diabetes mellitus type 2
63751110	NM_000251.2(MSH2):c.595T>C (p.Cys199Arg)	AAGGAAYGTGTTTTACCCGGAGG	Hereditary Nonpolyposis Colorectal Neoplasms
119450945	NM_000026.2(ADSL):c.674T>C (p.Met225Thr)	AAGAYGGTGACAGAAAAGGCAGG	Adenylosuccinate lyase deficiency
113993988	NM_002863.4(PYGL):c.2461T>C (p.Tyr821His)	AAGAAYATGCCCAAAACATCTGG	Glycogen storage disease, type VI
119485091	NM_022041.3(GAN):c.1268T>C (p.Ile423Thr)	AAGAAAAYCTACGCCATGGGTGG, AAAAYCTACGCCATGGGTGGAGG	Giant axonal neuropathy
137852419	NM_000132.3(F8):c.1660A>G (p.Ser554Gly)	AACYAGAGTAATAGCGGGTCAGG	Hereditary factor VIII deficiency disease
121964967	NM_000071.2(CBS):c.1150A>G (p.Lys384Glu)	AACTYGGTCCTGCGGGATGGGGG, GAACTYGGTCCTGCGGGATGGGG, GGAACTYGGTCCTGCGGGATGGG, AGGAACTYGGTCCTGCGGGATGG	Homocystinuria, pyridoxine-responsive
137852376	NM_000132.3(F8):c.1754T>C (p.Ile585Thr)	AACAGAYAATGTCAGACAAGAGG	Hereditary factor VIII deficiency disease
121917930	NM_006920.4(SCN1A):c.3577T>C (p.Trp1193Arg)	AACAAYGGTGGAACCTGAGAAGG	Generalized epilepsy with febrile seizures plus, type 1, Generalized epilepsy with febrile seizures plus, type 2
28939717	NM_003907.2(EIF2B5):c.271A>G (p.Thr91Ala)	AAATGYTTCCTGTACACCTGTGG	Leukoencephalopathy with vanishing white matter
80357276	NM_007294.3(BRCA1):c.122A>G (p.His41Arg)	AAATATGYGGTCACACTTTGTGG	Familial cancer of breast, Breast-ovarian cancer, familial 1

397515897	NM_000256.3(MYBPC3):c.1351+2T>C	AAAGGYGGGCCTGGGACCTGAGG	Familial hypertrophic cardiomyopathy 4, Cardiomyopathy
397514491	NM_005340.6(HINT1):c.152A>G (p.His51Arg)	AAAAYGTGTTGGTGCTTGAGGGG, GAAAAYGTGTTGGTGCTTGAGGG, AGAAAAYGTGTTGGTGCTTGAGG	Gamstorp-Wohlfart syndrome
387907164	NM_020894.2(UVSSA):c.94T>C (p.Cys32Arg)	AAAATTYGCAAGTATGTCTTAGG, AAATTYGCAAGTATGTCTTAGGG	UV-sensitive syndrome 3
118161496	NM_025152.2(NUBPL):c.815-27T>C	TGGTTCYAATGGATGTCTGCTGG, GGTTCYAATGGATGTCTGCTGGG	Mitochondrial complex I deficiency
764313717	NM_005609.2(PYGM):c.425_528del	TGGCTGYCAGGGACCCAGCAAGG, CTGYCAGGGACCCAGCAAGGAGG	
28934568	NM_003242.5(TGFBR2):c.923T>C (p.Leu308Pro)	AGTTCCYGACGGCTGAGGAGCGG	Loeys-Dietz syndrome 2

121913461	NM_007313.2(ABL1):c.814T>C (p.Tyr272His)	CCAGYACGGGGAGGTGTACGAGG, CAGYACGGGGAGGTGTACGAGGG	
377750405	NM_173551.4(ANKS6):c.1322A>G (p.Gln441Arg)	AGGGCYGTCGGACCTTCGAGTGG, GGGCYGTCGGACCTTCGAGTGGG, GGCYGTCGGACCTTCGAGTGGGG	Nephronophthisis 16
57639980	NM_001927.3(DES):c.1034T>C (p.Leu345Pro)	ATTCCCYGATGAGGCAGATGCGG, TTCCCYGATGAGGCAGATGCGGG	Myofibrillar myopathy 1
147391618	NM_020320.3(RARS2):c.35A>G (p.Gln12Arg)	ATACCYGGCAAGCAATAGCGCGG	Pontocerebellar hypoplasia type 6
182650126	NM_002977.3(SCN9A):c.2215A>G (p.Ile739Val)	GTAAYTGCAAGATCTACAAAAGG	Small fiber neuropathy
80358278	NM_004700.3(KCNQ4):c.842T>C (p.Leu281Ser)	ACATYGACAACCATCGGCTATGG	DFNA 2 Nonsyndromic Hearing Loss
786204012	NM_005957.4(MTHFR):c.388T>C (p.Cys130Arg)	GACCYGCTGCCGTCAGCGCCTGG	Homocysteinemia due to MTHFR deficiency
786204037	NM_005957.4(MTHFR):c.1883T>C (p.Leu628Pro)	TCCCACYGGACAACTGCCTCTGG	Homocysteinemia due to MTHFR deficiency
202147607	NM_000140.3(FECH):c.1137+3A>G	GTAGAYACCTTAGAGAACAATGG	Erythropoietic protoporphyria
122456136	NM_005183.3(CACNA1F):c.2267T>C (p.Ile756Thr)	TGCCAYTGCTGTGGACAACCTGG	
786204851	NM_007374.2(SIX6):c.110T>C (p.Leu37Pro)	GTCGCYGCCCGTGGCCCCTGCGG	Cataract, microphthalmia and nystagmus
794728167	NM_000138.4(FBN1):c.1468+2T>C	ATTGGYACGTGATCCATCCTAGG	Thoracic aortic aneurysms and aortic dissections
121964909	NM_000027.3(AGA):c.214T>C (p.Ser72Pro)	GACGGCYCTGTAGGCTTTGGAGG	Aspartylglycosaminuria
121964978	NM_000170.2(GLDC):c.2T>C (p.Met1Thr)	CGGCCAYGCAGTCCTGTGCCAGG, GGCCAYGCAGTCCTGTGCCAGGG	Non-ketotic hyperglycinemia
121965008	NM_000398.6(CYB5R3):c.446T>C (p.Leu149Pro)	CTGCYGGTCTACCAGGGCAAAGG	METHEMOGLOBINEMIA, TYPE I
121965064	NM_000128.3(F11):c.901T>C (p.Phe301Leu)	TGATYTCTTGGGAGAAGAACTGG	Hereditary factor XI deficiency disease
45517398	NM_000548.3(TSC2):c.5150T>C (p.Leu1717Pro)	GCCCYGCACGCAAATGTGAGTGG, CCCYGCACGCAAATGTGAGTGGG	Tuberous sclerosis syndrome
786205857	NM_015662.2(IFT172):c.770T>C (p.Leu257Pro)	TTGTGCYAGGAAGTTATGACAGG	RETINITIS PIGMENTOSA 71
786205904	NM_001135669.1(XPR1):c.653T>C (p.Leu218Ser)	GCGTTYACGTGTCCCCCTTTGG, CGTTYACGTGTCCCCCTTTGGG	BASAL GANGLIA CALCIFICATION,
104893704	NM_000388.3(CASR):c.2641T>C (p.Phe881Leu)	ACGCTYTCAAGGTGGCTGCCCGG, CGCTYTCAAGGTGGCTGCCCGGG	Hypercalciuric hypercalcemia
104893747	NM_198159.2(MITF):c.1195T>C (p.Ser399Pro)	ACTTYCCCTTATTCCATCCACGG, CTTYCCCTTATTCCATCCACGGG	Waardenburg syndrome type 2A
104893770	NM_000539.3(RHO):c.133T>C (p.Phe45Leu)	CATGYTTCTGCTGATCGTGCTGG, ATGYTTCTGCTGATCGTGCTGGG	Retinitis pigmentosa 4
28937596	NM_003907.2(EIF2B5):c.1882T>C (p.Trp628Arg)	AGGCCYGGAGCCCTGTTTTTAGG	Leukoencephalopathy with vanishing white matter
104893876	NM_001151.3(SLC25A4):c.293T>C (p.Leu98Pro)	GCAGCYCTTCTTAGGGGGTGTGG	Autosomal dominant progressive external ophthalmoplegia with mitochondrial DNA deletions 2
104893883	NM_006005.3(WFS1):c.2486T>C (p.Leu829Pro)	ACCATCCYGGAGGGCCGCCTGGG	WFS1-Related Disorders
104893962	NM_000165.4(GJA1):c.52T>C (p.Ser18Pro)	CTACYCAACTGCTGGAGGGAAGG	Oculodentodigital dysplasia

104893978	NM_000434.3(NEU1):c.718T>C (p.Trp240Arg)	GCCTCCYGGCGCTACGGAAGTGG, CCTCCYGGCGCTACGGAAGTGGG, CTCCYGGCGCTACGGAAGTGGGG	Sialidosis, type II
104894092	NM_002546.3(TNFRSF11B):c.349T>C (p.Phe117Leu)	TAGAGYTCTGCTTGAAACATAGG	Hyperphosphatasemia with bone disease
104894135	NM_000102.3(CYP17A1):c.316T>C (p.Ser106Pro)	CATCGCGYCCAACAACCGTAAGG, ATCGCGYCCAACAACCGTAAGGG	Complete combined 17-alpha-hydroxylase/17,20-lyase
104894151	NM_000102.3(CYP17A1):c.1358T>C (p.Phe453Ser)	AGCTCTYCCTCATCATGGCCTGG	Combined partial 17-alpha-hydroxylase/17,20-lyase deficiency
36015961	NM_000518.4(HBB):c.344T>C (p.Leu115Pro)	TGTGTGCYGGCCCATCACTTTGG	Beta thalassemia intermedia
104894472	NM_152443.2(RDH12):c.523T>C (p.Ser175Pro)	TCCYCGGTGGCTCACCACATTGG	Leber congenital amaurosis 13
104894587	NM_004870.3(MPDU1):c.356T>C (p.Leu119Pro)	TTCCYGGTCATGCACTACAGAGG	Congenital disorder of glycosylation type 1F
104894588	NM_004870.3(MPDU1):c.2T>C (p.Met1Thr)	AATAYGGCGGCCGAGGCGGACGG	Congenital disorder of glycosylation type 1F
104894626	NM_000304.3(PMP22):c.82T>C (p.Trp28Arg)	TAGCAAYGGATCGTGGGCAATGG	Charcot-Marie-Tooth disease, type IE
104894631	NM_018129.3(PNPO):c.784T>C (p.Ter262Gln)	ACCTYAACTCTGGGACCTGCTGG	"Pyridoxal 5-phosphate-dependent epilepsy"
104894703	NM_032551.4(KISS1R):c.305T>C (p.Leu102Pro)	GCCCTGCYGTACCCGCTGCCCGG, TGCYGTACCCGCTGCCCGGCTGG	
104894826	NM_000166.5(GJB1):c.407T>C (p.Val136Ala)	ATGYCATCAGCGTGGTGTTCCGG	Dejerine-Sottas disease, X-linked hereditary motor and sensory neuropathy

104894859	NM_001122606.1(LAMP2):c.961T>C (p.Trp321Arg)	CAGCTACYGGGATGCCCCCCTGG, AGCTACYGGGATGCCCCCCTGGG	Danon disease
104894931	NM_006517.4(SLC16A2):c.1313T>C (p.Leu438Pro)	TGAGCYGGTGGGCCCAATGCAGG	Allan-Herndon-Dudley syndrome
104894935	NM_000330.3(RS1):c.38T>C (p.Leu13Pro)	TTACTTCYCTTTGGCTATGAAGG	Juvenile retinoschisis
104895217	NM_001065.3(TNFRSF1A):c.175T>C (p.Cys59Arg)	TGCYGTACCAAGTGCCACAAAGG	TNF receptor-associated periodic fever syndrome (TRAPS)
143889283	NM_003793.3(CTSF):c.692A>G (p.Tyr231Cys)	CTCCAYACTGAGCTGTGCCACGG	Ceroid lipofuscinosis, neuronal, 13
122459147	NM_001159702.2(FHL1):c.310T>C (p.Cys104Arg)	GGGGYGCTTCAAGGCCATTGTGG	Myopathy, reducing body, X-linked, childhood- onset
74552543	NM_020184.3(CNNM4):c.971T>C (p.Leu324Pro)	AAGCTCCYGGACTTTTTTCTGGG	Cone-rod dystrophy amelogenesis imperfecta
199476117	m.10158T>C	AAAYCCACCCCTTACGAGTGCGG	Leigh disease, Leigh syndrome due to mitochondrial complex I deficiency, Mitochondrial complex I deficiency
794727808	NM_020451.2(SEPN1):c.872+2T>C	TTCCGGYGAGTGGGCCACACTGG	Congenital myopathy with fiber type disproportion, Eichsfeld type congenital muscular dystrophy
140547520	NM_005022.3(PFN1):c.350A>G (p.Glu117Gly)	CACCTYCTTTGCCCATCAGCAGG	Amyotrophic lateral sclerosis 18
397514359	NM_000060.3(BTD):c.445T>C (p.Phe149Leu)	TCACCGCYTCAATGACACAGAGG	Biotinidase deficiency
207460001	m.15197T>C	CTAYCCGCCATCCCATACATTGG	Exercise intolerance
397514406	NM_000060.3(BTD):c.1214T>C (p.Leu405Pro)	TTCACCCYGGTCCCTGTCTGGGG	Biotinidase deficiency
397514516	NM_006177.3(NRL):c.287T>C (p.Met96Thr)	GAGGCCAYGGAGCTGCTGCAGGG	Retinitis pigmentosa 27
72554312	NM_000531.5(OTC):c.134T>C (p.Leu45Pro)	CTCACTCYAAAAAACTTTACCGG	Ornithine carbamoyltransferase deficiency
397514569	NM_178012.4(TUBB2B):c.350T>C (p.Leu117Pro)	GGTCCYGGATGTGGTGAGGAAGG	Polymicrogyria, asymmetric
397514571	NM_000431.3(MVK):c.122T>C (p.Leu41Pro)	CGGCYTCAACCCCACAGCAATGG, GGCYTCAACCCCACAGCAATGGG	Porokeratosis, disseminated superficial actinic 1
794728390	NM_000238.3(KCNH2):c.2396T>C (p.Leu799Pro)	GCCATCCYGGGTATGGGGTGGGG, CCATCCYGGGTATGGGGTGGGGG, CATCCYGGGTATGGGGTGGGGGG	Cardiac arrhythmia
397514713	NM_001199107.1(TBC1D24):c.686T>C (p.Phe229Ser)	GGTCTYTGACGTCTTCCTGGTGG	Early infantile epileptic encephalopathy 16
397514719	NM_080605.3(B3GALT6):c.193A>G (p.Ser65Gly)	CGCYGGCCACCAGCACTGCCAGG	Spondyloepimetaphyseal dysplasia with joint laxity
730880608	NM_000256.3(MYBPC3):c.3796T>C (p.Cys1266Arg)	GAGYGCCGCCTGGAGGTGCGAGG	Cardiomyopathy
397515329	NM_001382.3(DPAGT1):c.503T>C (p.Leu168Pro)	AATCCYGTACTATGTCTACATGG, ATCCYGTACTATGTCTACATGGG, TCCYGTACTATGTCTACATGGGG	Congenital disorder of glycosylation type 1J

397515465	NM_018127.6(ELAC2):c.460T>C (p.Phe154Leu)	ATAYTTTCTGGTCCATTGAAAGG	Combined oxidative phosphorylation deficiency 17
397515557	NM_005211.3(CSF1R):c.2483T>C (p.Phe828Ser)	CATCTYTGACTGTGTCTACACGG	Hereditary diffuse leukoencephalopathy with spheroids
397515599	NM_194248.2(OTOF):c.3413T>C (p.Leu1138Pro)	AGGTGCYGTTCTGGGGCCTACGG, GGTGCYGTTCTGGGGCCTACGGG	Deafness, autosomal recessive 9
397515766	NM_000138.4(FBN1):c.2341T>C (p.Cys781Arg)	GGACAAYGTAGAAATACTCCTGG	Marfan syndrome
565779970	NM_001429.3(EP300):c.3573T>A (p.Tyr1191Ter)	CTTAYTACAGTTACCAGAACAGG	Rubinstein-Taybi syndrome 2
786200938	NM_080605.3(B3GALT6):c.1A>G (p.Met1Val)	AGCTTCAYGGCGCCCGCGCCGGG, TCAYGGCGCCCGCGCCGGGCCGG	Spondyloepimetaphyseal dysplasia with joint laxity
28942087	NM_000229.1(LCAT):c.698T>C (p.Leu233Pro)	ATCTCTCYTGGGGCTCCCTGGGG, TCTCYTGGGGCTCCCTGGGGTGG	Norum disease
128621203	NM_000061.2(BTK):c.1625T>C (p.Leu542Pro)	TCGGCCYGTCCAGGTGAGTGTGG	X-linked agammaglobulinemia with growth hormone deficiency
397515412	NM_006383.3(CIB2):c.368T>C (p.Ile123Thr)	CTTCAYCTGCAAGGAGGACCTGG	Deafness, autosomal recessive 48
193929364	NM_000352.4(ABCC8):c.404T>C (p.Leu135Pro)	AAGCYGCTAATTGGTAGGTGAGG	Permanent neonatal diabetes mellitus
730880872	NM_000257.3(MYH7):c.1400T>C (p.Ile467Thr)	TCGAGAYCTTCGATGTGAGTTGG, CGAGAYCTTCGATGTGAGTTGGG	Cardiomyopathy
80356474	NM_002977.3(SCN9A):c.2543T>C (p.Ile848Thr)	AAGATCAYTGGTAACTCAGTAGG, AGATCAYTGGTAACTCAGTAGGG, GATCAYTGGTAACTCAGTAGGGG	Primary erythromelalgia
80356489	NM_001164277.1(SLC37A4):c.352T>C (p.Trp118Arg)	GGGCYGGCCCCCATGTGGGAAGG	Glucose-6-phosphate transport defect
80356536	NM_152296.4(ATP1A3):c.2338T>C (p.Phe780Leu)	GCCCYTCCTGCTGTTCATCATGG	Dystonia 12
80356596	NM_194248.2(OTOF):c.3032T>C (p.Leu1011Pro)	GATGCYGGTGTTCGACAACCTGG	Deafness, autosomal recessive 9, Auditory neuropathy, autosomal recessive, 1

80356689	NM_000083.2(CLCN1):c.857T>C (p.Val286Ala)	AGGAGYGCTATTTAGCATCGAGG	Myotonia congenita
118203884	m.4409T>C	AGGYCAGCTAAATAAGCTATCGG	Mitochondrial myopathy
587777625	NM_173596.2(SLC39A5):c.911T>C (p.Met304Thr)	AGAACAYGCTGGGGCTTTTGCGG	Myopia 24, autosomal dominant
587783087	NM_003159.2(CDKL5):c.602T>C (p.Leu201Pro)	ATTCYTGGGGAGCTTAGCGATGG	not provided
118203951	NM_013319.2(UBIAD1):c.511T>C (p.Ser171Pro)	TCTGGCYCCTTTCTCTACACAGG, GGCYCCTTTCTCTACACAGGAGG	Schnyder crystalline corneal dystrophy
118204017	NM_000018.3(ACADVL):c.1372T>C (p.Phe458Leu)	TCGCATCYTCCGGATCTTTGAGG, CGCATCYTCCGGATCTTTGAGGG, GCATCYTCCGGATCTTTGAGGGG	Very long chain acyl-CoA dehydrogenase deficiency
397518466	NM_000833.4(GRIN2A):c.2T>C (p.Met1Thr)	CTAYGGGCAGAGTGGGCTATTGG	Focal epilepsy with speech disorder with or without mental retardation
118204069	NM_000237.2(LPL):c.337T>C (p.Trp113Arg)	GGACYGGCTGTCACGGGCTCAGG	Hyperlipoproteinemia, type I
118204080	NM_000237.2(LPL):c.755T>C (p.Ile252Thr)	GTGAYTGCAGAGAGAGGACTTGG	Hyperlipoproteinemia, type I
118204111	NM_000190.3(HMBS):c.739T>C (p.Cys247Arg)	GCTTCGCYGCATCGCTGAAAGGG	Acute intermittent porphyria
80357438	NM_007294.3(BRCA1):c.65T>C (p.Leu22Ser)	AAATCTYAGAGTGTCCCATCTGG	Familial cancer of breast, Breast-ovarian cancer, familial 1, Hereditary cancer-predisposing syndrome
139877390	NM_001040431.2(COA3):c.215A>G (p.Tyr72Cys)	CCAYCTGGGGAGGTAGGTTCAGG	
793888527	NM_005859.4(PURA):c.563T>C (p.Ile188Thr)	GACCAYTGCGCTGCCCGCGCAGG, ACCAYTGCGCTGCCCGCGCAGGG, CCAYTGCGCTGCCCGCGCAGGGG	not provided, Mental retardation, autosomal dominant 31
561425038	NM_002878.3(RAD51D):c.1A>G (p.Met1Val)	CGCCCAYGTTCCCCGCAGGCCGG	Hereditary cancer-predisposing syndrome
121907934	NM_024105.3(ALG12):c.473T>C (p.Leu158Pro)	TCCYGCTGGCCCTCGCGGCCTGG	Congenital disorder of glycosylation type 1G
80358207	NM_153212.2(GJB4):c.409T>C (p.Phe137Leu)	CCTCATCYTCAAGGCCGCCGTGG	Erythrokeratodermia variabilis
80358228	NM_002353.2(TACSTD2):c.557T>C (p.Leu186Pro)	TCGGCYGCACCCCAAGTTCGTGG	Lattice corneal dystrophy Type III
121908076	NM_138691.2(TMC1):c.1543T>C (p.Cys515Arg)	AGGACCTYGCTGGGAAACAATGG, ACCTYGCTGGGAAACAATGGTGG, CCTYGCTGGGAAACAATGGTGGG	Deafness, autosomal recessive 7

121908089	NM_017838.3(NHP2):c.415T>C (p.Tyr139His)	GGAGGCTYACGATGAGTGCCTGG, GGCTYACGATGAGTGCCTGGAGG	Dyskeratosis congenita autosomal recessive 1, Dyskeratosis congenita, autosomal recessive 2
121908154	NM_001243133.1(NLRP3):c.926T>C (p.Phe309Ser)	GGTGCCTYTGACGAGCACATAGG	Familial cold urticaria, Chronic infantile neurological, cutaneous and articular syndrome
121908158	NM_001033855.2(DCLRE1C):c.2T>C (p.Met1Thr)	GGCGCTAYGAGTTCTTTCGAGGG, GCGCTAYGAGTTCTTTCGAGGGG	Histiocytic medullary reticulosis
796052870	NM_018129.3(PNPO):c.2T>C (p.Met1Thr)	CCCCCAYGACGTGCTGGCTGCGG, CCCCAYGACGTGCTGGCTGCGGG, CCCAYGACGTGCTGGCTGCGGGG	not provided
121908318	NM_020427.2(SLURP1):c.43T>C (p.Trp15Arg)	GCAGCCYGGAGCATGGGCTGTGG	Acroerythrokeratoderma
121908352	NM_022124.5(CDH23):c.5663T>C (p.Phe1888Ser)	CTCACCTYCAACATCACTGCGGG	Deafness, autosomal recessive 12
121908520	NM_000030.2(AGXT):c.613T>C (p.Ser205Pro)	CCTGTACYCGGGCTCCCAGAAGG	Primary hyperoxaluria, type I
121908618	NM_004273.4(CHST3):c.920T>C (p.Leu307Pro)	CGTGCYGGCCTCGCGCATGGTGG	Spondyloepiphyseal dysplasia with congenital joint dislocations
11694	NM_006432.3(NPC2):c.199T>C (p.Ser67Pro)	TATTCAGYCTAAAAGCAGCAAGG	Niemann-Pick disease type C2
121908739	NM_000022.2(ADA):c.320T>C (p.Leu107Pro)	CCTGCYGGCCAACTCCAAAGTGG	Severe combined immunodeficiency due to ADA deficiency
80359022	NM_000059.3(BRCA2):c.7958T>C (p.Leu2653Pro)	TGCYTCTTCAACTAAAATACAGG	Familial cancer of breast, Breast-ovarian cancer, familial 2
121908902	NM_003880.3(WISP3):c.232T>C (p.Cys78Arg)	AAAATCYGTGCCAAGCAACCAGG, AAATCYGTGCCAAGCAACCAGGG, AATCYGTGCCAAGCAACCAGGGG	Progressive pseudorheumatoid dysplasia
121908947	NM_006892.3(DNMT3B):c.808T>C (p.Ser270Pro)	CAAGTTCYCCGAGGTGAGTCCGG, AAGTTCYCCGAGGTGAGTCCGGG, AGTTCYCCGAGGTGAGTCCGGGG	Centromeric instability of chromosomes 1,9 and 16 and immunodeficiency
121909028	NM_000492.3(CFTR):c.3857T>C (p.Phe1286Ser)	AGCCTYTGGAGTGATACCACAGG	Cystic fibrosis
121909135	NM_000085.4(CLCNKB):c.1294T>C (p.Tyr432His)	CTTTGTCYATGGTGAGTCTGGGG	Bartter syndrome type 3
121909143	NM_001300.5(KLF6):c.506T>C (p.Leu169Pro)	GGAGCYGCCCTCGCCAGGGAAGG	
121909182	NM_001089.2(ABCA3):c.302T>C (p.Leu101Pro)	GCACYTGTGATCAACATGCGAGG	Surfactant metabolism dysfunction, pulmonary, 3
121909200	NM_000503.5(EYA1):c.1459T>C (p.Ser487Pro)	CACTCYCGCTCATTCACTCCCGG	Melnick-Fraser syndrome

121909247	NM_004970.2(IGFALS):c.1618T>C (p.Cys540Arg)	GGACYGTGGCTGCCCTCTCAAGG	Acid-labile subunit deficiency
121909253	NM_005570.3(LMAN1):c.2T>C (p.Met1Thr)	AGAYGGCGGGATCCAGGCAAAGG	Combined deficiency of factor V and factor VIII, 1
121909385	NM_000339.2(SLC12A3):c.1868T>C (p.Leu623Pro)	CAACCYGGCCCTCAGCTACTCGG	Familial hypokalemia-hypomagnesemia
121909497	NM_002427.3(MMP13):c.224T>C (p.Phe75Ser)	TTCTYCGGCTTAGAGGTGACTGG	Spondyloepimetaphyseal dysplasia, Missouri type
121909508	NM_000751.2(CHRND):c.188T>C (p.Leu63Pro)	AACCYCATCTCCCTGGTGAGAGG	MYASTHENIC SYNDROME, CONGENITAL, 3B, FAST-CHANNEL
121909519	NM_001100.3(ACTA1):c.287T>C (p.Leu96Pro)	CGAGCYTCGCGTGGCTCCCGAGG	Nemaline myopathy 3
121909572	NM_000488.3(SERPINC1):c.667T>C (p.Ser223Pro)	TGGGTGYCCAATAAGACCGAAGG	Antithrombin III deficiency
121909677	NM_000821.6(GGCX):c.896T>C (p.Phe299Ser)	TATGTYCTCCTACGTCATGCTGG	Pseudoxanthoma elasticum-like disorder with multiple coagulation factor deficiency
121909727	NM_001018077.1(NR3C1):c.2209T>C (p.Phe737Leu)	CTATTGCYTCCAAACATTTTTGG	Glucocorticoid resistance, generalized
139573311	NM_000492.3(CFTR):c.1400T>C (p.Leu467Pro)	TTCACYTCTAATGGTGATTATGG, TCACYTCTAATGGTGATTATGGG	Cystic fibrosis
121912441	NM_000454.4(SOD1):c.341T>C (p.Ile114Thr)	CATCAYTGGCCGCACACTGGTGG	Amyotrophic lateral sclerosis type 1
121912446	NM_000454.4(SOD1):c.434T>C (p.Leu145Ser)	CGTTYGGCTTGTGGTGTAATTGG, GTTYGGCTTGTGGTGTAATTGGG	Amyotrophic lateral sclerosis type 1
121912463	NM_000213.3(ITGB4):c.1684T>C (p.Cys562Arg)	GGCCAGYGTGTGTGTGAGCCTGG	Epidermolysis bullosa with pyloric atresia
121912492	NM_002292.3(LAMB2):c.961T>C (p.Cys321Arg)	CCTCAACYGCGAGCAGTGTCAGG	Nephrotic syndrome, type 5, with or without ocular abnormalities
397516659	NM_001399.4(EDA):c.2T>C (p.Met1Thr)	GGCCAYGGGCTACCCGGAGGTGG	Hypohidrotic X-linked ectodermal dysplasia

111033589	NM_021044.2(DHH):c.485T>C (p.Leu162Pro)	GTTGCYGGCGCGCCTCGCAGTGG	46,XY gonadal dysgenesis, complete, dhh-related
111033622	NM_000206.2(IL2RG):c.343T>C (p.Cys115Arg)	TGGCYGTCAGTTGCAAAAAAAGG	X-linked severe combined immunodeficiency
121912613	NM_001041.3(SI):c.1859T>C (p.Leu620Pro)	ATGCYGGAGTTCAGTTTGTTTGG	Sucrase-isomaltase deficiency
121912619	NM_016180.4(SLC45A2):c.1082T>C (p.Leu361Pro)	GAGTTTCYCATCTACGAAAGAGG	Oculocutaneous albinism type 4
61750581	NM_000552.3(VWF):c.4837T>C (p.Ser1613Pro)	CTGCCYCTGATGAGATCAAGAGG	von Willebrand disease, type 2a
121912653	NM_000546.5(TP53):c.755T>C (p.Leu252Pro)	CATCCYCACCATCATCACACTGG	Li-Fraumeni syndrome 1
111033683	NM_000155.3(GALT):c.386T>C (p.Met129Thr)	AGGTCAYGTGCTTCCACCCCTGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
111033752	NM_000155.3(GALT):c.677T>C (p.Leu226Pro)	CAGGAGCYACTCAGGAAGGTGGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
121912729	NM_000039.1(APOA1):c.593T>C (p.Leu198Ser)	GCGCTYGGCCGCGCGCCTTGAGG	Familial visceral amyloidosis, Ostertag type
769452	NM_000041.3(APOE):c.137T>C (p.Leu46Pro)	AACYGGCACTGGGTCGCTTTTGG	
121912762	NM_016124.4(RHD):c.329T>C (p.Leu110Pro)	ACACYGTTCAGGTATTGGGATGG	
111033824	NM_000155.3(GALT):c.1138T>C (p.Ter380Arg)	CGCCYGACCACGCCGACCACAGG, GCCYGACCACGCCGACCACAGGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
111033832	NM_000155.3(GALT):c.980T>C (p.Leu327Pro)	TCCYGCGCTCTGCCACTGTCCGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
730881974	NM_000455.4(STK11):c.545T>C (p.Leu182Pro)	GGGAACCYGCTGCTCACCACCGG, AACCYGCTGCTCACCACCGGTGG	Hereditary cancer-predisposing syndrome
1064644	NM_000157.3(GBA):c.703T>C (p.Ser235Pro)	GGGYCACTCAAGGGACAGCCCGG	Gaucher disease
796052090	NM_138413.3(HOGA1):c.533T>C (p.Leu178Pro)	GGACCYGCCTGTGGATGCAGTGG	Primary hyperoxaluria, type III
121913141	NM_000208.2(INSR):c.779T>C (p.Leu260Pro)	CTACCYGGACGGCAGGTGTGTGG	Leprechaunism syndrome
121913272	NM_006218.2(PIK3CA):c.1258T>C (p.Cys420Arg)	GGAACACYGTCCATTGGCATGGG, GAACACYGTCCATTGGCATGGGG	Congenital lipomatous overgrowth, vascular malformations, and epidermal nevi, Neoplasm of ovary, PIK3CA Related Overgrowth Spectrum
61751310	NM_000552.3(VWF):c.8317T>C (p.Cys2773Arg)	GCTCCYGCTGCTCTCCGACACGG	von Willebrand disease, type 2a
312262799	NM_024408.3(NOTCH2):c.1438T>C (p.Cys480Arg)	TTCACAYGTCTGTGCATGCCAGG	Alagille syndrome 2
121913570	NM_000426.3(LAMA2):c.7691T>C (p.Leu2564Pro)	ATCATTCYTTTGGGAAGTGGAGG, TCATTCYTTTGGGAAGTGGAGGG	Merosin deficient congenital muscular dystrophy
121913640	NM_000257.3(MYH7):c.1046T>C (p.Met349Thr)	AACTCCAYGTATAAGCTGACAGG	Familial hypertrophic cardiomyopathy 1, Cardiomyopathy
121913642	NM_000257.3(MYH7):c.1594T>C (p.Ser532Pro)	CATCATGYCCATCCTGGAAGAGG	Dilated cardiomyopathy 1S

119463996	NM_001079802.1(FKTN):c.527T>C (p.Phe176Ser)	GTAGTCTYTCATGAGAGGAGTGG	Limb-girdle muscular dystrophy-
587776456	NM_002049.3(GATA1):c.1240T>C (p.Ter414Arg)	GCTCAYGAGGGCACAGAGCATGG	GATA-1-related thrombocytopenia with dyserythropoiesis
63750654	NM_000184.2(HBG2):c.-228T>C	ATGCAAAYATCTGTCTGAAACGG	Fetal hemoglobin quantitative trait locus 1
587776519	NM_001999.3(FBN2):c.3725-15A>G	AGCAYTGCAACCACATTGTCAGG	Congenital contractural arachnodactyly
78365220	NM_000402.4(G6PD):c.473T>C (p.Leu158Pro)	TGCCCYCCACCTGGGGTCACAGG	Anemia, nonspherocytic hemolytic, due to G6PD deficiency
63750741	NM_000179.2(MSH6):c.1346T>C (p.Leu449Pro)	CTGGGGCYGGTATTCATGAAAGG	Hereditary Nonpolyposis Colorectal Neoplasms
587776914	NM_017565.3(FAM20A):c.590-2A>G	GTAATCYGCAAAGGAGGAGAAGG, TAATCYGCAAAGGAGGAGAAGG	Enamel-renal syndrome
5030809	NM_000551.3(VHL):c.292T>C (p.Tyr98His)	CCCYACCCAACGCTGCCGCCTGG	Von Hippel-Lindau syndrome, Hereditary cancer-predisposing syndrome
199476132	m.5728T>C	CAATCYACTTCTCCCGCCGCCGG, AATCYACTTCTCCCGCCGCCGGG	Cytochrome-c oxidase deficiency, Mitochondrial complex I deficiency

62637012	NM_014336.4(AIPL1):c.715T>C (p.Cys239Arg)	CTGCCAGYGCCTGCTGAAGAAGG, CCAGYGCCTGCTGAAGAAGGAGG	Leber congenital amaurosis 4
199476199	NM_207352.3(CYP4V2):c.1021T>C (p.Ser341Pro)	AAACTGGYCCTTATACCTGTTGG, AACTGGYCCTTATACCTGTTGGG	Bietti crystalline corneoretinal dystrophy
587777183	NM_006702.4(PNPLA6):c.3053T>C (p.Phe1018Ser)	CCTYTAACCGCAGCATCCATCGG	Boucher Neuhauser syndrome
199476389	NM_000487.5(ARSA):c.899T>C (p.Leu300Ser)	GGTCTCTYGCGGTGTGGAAAGGG	Metachromatic leukodystrophy
199476398	NM_016599.4(MYOZ2):c.142T>C (p.Ser48Pro)	TTAYCCCATCTCAGTAACCGTGG	Familial hypertrophic cardiomyopathy 16
119456967	NM_001037633.1(SIL1):c.1370T>C (p.Leu457Pro)	TTGCYGAAGGAGCTGAGATGAGG	Marinesco-Sj\xc3\xb6grensyndrome
730882253	NM_006888.4(CALM1):c.268T>C (p.Phe90Leu)	GGCAYTCCGAGTCTTTGACAAGG	Long QT syndrome 14
587777283	NM_012338.3(TSPAN12):c.413A>G (p.Tyr138Cys)	TAATCCAYAATTTGTCATCCTGG	Exudative vitreoretinopathy 5
587777306	NM_015884.3(MBTPS2):c.1391T>C (p.Phe464Ser)	GCTYTGCTTTGGATGGACAATGG	Palmoplantar keratoderma, mutilating, with periorificial keratotic plaques, X-linked
56378716	NM_000250.1(MPO):c.752T>C (p.Met251Thr)	TCACTCAYGTTCATGCAATGGGG	Myeloperoxidase deficiency
587777390	NM_005026.3(PIK3CD):c.1246T>C (p.Cys416Arg)	GCAGGACYGCCCCATTGCCTGGG	Activated PI3K-delta syndrome
587777480	NM_003108.3(SOX11):c.178T>C (p.Ser60Pro)	TATGGYCCAAGATCGAACGCAGG	Mental retardation, autosomal dominant 27
587777663	NM_001288767.1(ARMC5):c.1379T>C (p.Leu460Pro)	GCCCGACYGCGGGATGCTGGTGG	Acth-independent macronodular adrenal hyperplasia 2
61753033	NM_000350.2(ABCA4):c.5819T>C (p.Leu1940Pro)	AAGGCYACATGAACTAACCAAGG	Stargardt disease, Stargardt disease 1, Cone-rod dystrophy 3
200488568	NM_002972.3(SBF1):c.4768A>G (p.Thr1590Ala)	CAGGCGYCCTCTTGCTCAGCCGG	Charcot-Marie-Tooth disease, type 4B3
132630274	NM_000377.2(WAS):c.809T>C (p.Leu270Pro)	CGGAGTCYGTTCTCCAGGGCAGG	Severe congenital neutropenia X-linked
132630308	NM_001399.4(EDA):c.181T>C (p.Tyr61His)	CTGCYACCTAGAGTTGCGCTCGG	Hypohidrotic X-linked ectodermal dysplasia
60934003	NM_170707.3(LMNA):c.1589T>C (p.Leu530Pro)	ACGGCTCYCATCAACTCCACTGG, CGGCTCYCATCAACTCCACTGGG, GGCTCYCATCAACTCCACTGGGG	Benign scapuloperoneal muscular dystrophy with cardiomyopathy
180177160	NM_000030.2(AGXT):c.1076T>C (p.Leu359Pro)	GGTGCYGCGGATCGGCCTGCTGG, GTGCYGCGGATCGGCCTGCTGGG	Primary hyperoxaluria, type I
180177222	NM_000030.2(AGXT):c.449T>C (p.Leu150Pro)	GTGCYGCTGTTCTTAACCCACGG, TGCYGCTGTTCTTAACCCACGGG	Primary hyperoxaluria, type I
180177254	NM_000030.2(AGXT):c.661T>C (p.Ser221Pro)	GCTCATCYCCTTCAGTGACAAGG	Primary hyperoxaluria, type I
180177264	NM_000030.2(AGXT):c.757T>C (p.Cys253Arg)	GGGGCYGTGACGACCAGCCCAGG	Primary hyperoxaluria, type I
180177293	NM_000030.2(AGXT):c.893T>C (p.Leu298Pro)	GTATCYGCATGGGCGCCTGCAGG	Primary hyperoxaluria, type I
376785840	NM_001282227.1(CECR1):c.1232A>G (p.Tyr411Cys)	GAAATCAYAGGACAAGCCTTTGG	Polyarteritis nodosa
587779393	NM_000257.3(MYH7):c.4937T>C (p.Leu1646Pro)	GAGCCYCCAGAGCTTGTTGAAGG	Myopathy, distal, 1
587779410	NM_012434.4(SLC17A5):c.500T>C (p.Leu167Pro)	ATTGTACYCAGAGCACTAGAAGG	Sialic acid storage disease, severe infantile type
587779513	NM_000090.3(COL3A1):c.2337+2T>C (p.Gly762_Lys779del)	AGGYAACCCTTAATACTACCTGG	Ehlers-Danlos syndrome, type 4
777539013	NM_020376.3(PNPLA2):c.757+2T>C	GAACGGYGCGCGGACCCGGGCGG, AACGGYGCGCGGACCCGGGCGGG	Neutral lipid storage disease with myopathy
34557412	NM_012452.2(TNFRSF13B):c.310T>C (p.Cys104Arg)	ACTTCYGTGAGAACAAGCTCAGG	Immunoglobulin A deficiency 2, Common variable
796052970	NM_001165963.1(SCN1A):c.1094T>C (p.Phe365Ser)	CAAGCTYTGATACCTTCAGTTGG, AAGCTYTGATACCTTCAGTTGGG	not provided

724159989	NC_012920.1:m.7505T>C	CCTCCAYGACTTTTTCAAAAAGG	Deafness, nonsyndromic sensorineural, mitochondrial
796053222	NM_014191.3(SCN8A):c.4889T>C (p.Leu1630Pro)	CGTCYGATCAAAGGCGCCAAAGG, GTCYGATCAAAGGCGCCAAAGGG	not provided
118192127	NM_000540.2(RYR1):c.10817T>C (p.Leu3606Pro)	TACTACCYGGACCAGGTGGGTGG, ACTACCYGGACCAGGTGGGTGGG, CTACCYGGACCAGGTGGGTGGGG	Central core disease
118192170	NM_000540.2(RYR1):c.14693T>C (p.Ile4898Thr)	AGGCAYTGGGGACGAGATCGAGG	Malignant hyperthermia susceptibility type 1, Central core disease

121917703	NM_005247.2(FGF3):c.466T>C (p.Ser156Pro)	GTACGTGYCTGTGAACGGCAAGG, TACGTGYCTGTGAACGGCAAGGG	Deafness with labyrinthine aplasia microtia and microdontia (LAMM)
690016549	NM_005211.3(CSF1R):c.2450T>C (p.Leu817Pro)	CCGCCYGCCTGTGAAGTGGATGG	Hereditary diffuse leukoencephalopathy with spheroids
690016552	NM_005211.3(CSF1R):c.2566T>C (p.Tyr856His)	GAATCCCYACCCTGGCATCCTGG	Hereditary diffuse leukoencephalopathy with spheroids
121917738	NM_001098668.2(SFTPA2):c.593T>C (p.Phe198Ser)	GGAGACTYCCGCTACTCAGATGG, GAGACTYCCGCTACTCAGATGGG	Idiopathic fibrosing alveolitis, chronic form
690016559	NM_005211.3(CSF1R):c.1957T>C (p.Cys653Arg)	AGCCYGTACCCATGGAGGTAAGG, GCCYGTACCCATGGAGGTAAGGG	Hereditary diffuse leukoencephalopathy with spheroids
690016560	NM_005211.3(CSF1R):c.2717T>C (p.Ile906Thr)	GCAGAYCTGCTCCTTCCTTCAGG	Hereditary diffuse leukoencephalopathy with spheroids
121917769	NM_003361.3(UMOD):c.376T>C (p.Cys126Arg)	GGCCACAYGTGTCAATGTGGTGG, GCCACAYGTGTCAATGTGGTGGG	Familial juvenile gout
121917773	NM_003361.3(UMOD):c.943T>C (p.Cys315Arg)	ATGGCACYGCCAGTGCAAACAGG	Glomerulocystic kidney disease with hyperuricemia and isosthenuria
121917818	NM_007255.2(B4GALT7):c.617T>C (p.Leu206Pro)	TGCYCTCCAAGCAGCACTACCGG	Ehlers-Danlos syndrome progeroid type
121917824	NM_021615.4(CHST6):c.827T>C (p.Leu276Pro)	GGACCYGGCGCGGGAGCCGCTGG	Macular corneal dystrophy Type I
121917848	NM_000452.2(SLC10A2):c.728T>C (p.Leu243Pro)	TTTCYTCTGGCTAGAATTGCTGG	Bile acid malabsorption, primary
121918006	NM_000478.4(ALPL):c.1306T>C (p.Tyr436His)	TGGACYATGGTGAGACCTCCAGG	Infantile hypophosphatasia
121918010	NM_000478.4(ALPL):c.979T>C (p.Phe327Leu)	CAAAGGCYTCTTCTTGCTGGTGG, GGCYTCTTCTTGCTGGTGGAAGG	Infantile hypophosphatasia
121918088	NM_000371.3(TTR):c.400T>C (p.Tyr134His)	CCCCYACTCCTATTCCACCACGG	
121918110	NM_001042465.1(PSAP):c.1055T>C (p.Leu352Pro)	GAAGCYGCCGAAGTCCCTGTCGG	Gaucher disease, atypical, due to saposin C deficiency
121918137	NM_003730.4(RNASET2):c.550T>C (p.Cys184Arg)	CCAGYGCCTTCCACCAAGCCAGG	Leukoencephalopathy, cystic, without megalencephaly
121918191	NM_001127628.1(FBP1):c.581T>C (p.Phe194Ser)	GGAGTYCATTTTGGTGGACAAGG	Fructose-biphosphatase deficiency
121918306	NM_006946.2(SPTBN2):c.758T>C (p.Leu253Pro)	ACCAAGCYGCTGGATCCCGAAGG, AAGCYGCTGGATCCCGAAGGTGG, AGCYGCTGGATCCCGAAGGTGGG	Spinocerebellar ataxia 5
121918505	NM_000141.4(FGFR2):c.799T>C (p.Ser267Pro)	AATGCCYCCACAGTGGTCGGAGG	Pfeiffer syndrome, Neoplasm of stomach
121918643	NM_003126.2(SPTA1):c.620T>C (p.Leu207Pro)	GTGGAGCYGGTAGCTAAAGAAGG, TGGAGCYGGTAGCTAAAGAAGGG	Hereditary pyropoikilocytosis, Elliptocytosis 2
121918646	NM_001024858.2(SPTB):c.604T>C (p.Trp202Arg)	CTCCAGCYGGAAGGATGGCTTGG	Spherocytosis type 2
121918648	NM_001024858.2(SPTB):c.6055T>C (p.Ser2019Pro)	ATGCCYCTGTGGCTGAGGCGTGG	
727504166	NM_000543.4(SMPD1):c.475T>C (p.Cys159Arg)	TGAGGCCYGTGGCCTGCTCCTGG, GAGGCCYGTGGCCTGCTCCTGGG	Niemann-Pick disease, type A, Niemann-Pick disease, type B
193922915	NM_000434.3(NEU1):c.1088T>C (p.Leu363Pro)	CAGCYATGGCCAGGCCCCAGTGG	Sialidosis, type II
727504419	NM_000501.3(ELN):c.889+2T>C	CAGGYAACATCTGTCCCAGCAGG, AGGYAACATCTGTCCCAGCAGGG	Supravalvar aortic stenosis
376395543	NM_000256.3(MYBPC3):c.26-2A>G	GAGACYGAAGGGCCAGGTGGAGG	Primary familial hypertrophic cardiomyopathy, Familial hypertrophic cardiomyopathy 4, Cardiomyopathy
1169305	NM_000545.6(HNF1A):c.1720G>A (p.Gly574Ser)	GATGCYGGCAGGGTCCTGGCTGG, ATGCYGGCAGGGTCCTGGCTGGG, TGCYGGCAGGGTCCTGGCTGGGG	Maturity-onset diabetes of the young, type 3
730880130	NM_000527.4(LDLR):c.1468T>C (p.Trp490Arg)	CTACYGGACCGACTCTGTCCTGG, TACYGGACCGACTCTGTCCTGGG	Familial hypercholesterolemia
281860286	NM_018713.2(SLC30A10):c.500T>C (p.Phe167Ser)	GGCGCTTYCGGGGGGCCTCAGGG	Hypermanganesemia with dystonia, polycythemia and cirrhosis
730880306	NM_145693.2(LPIN1):c.1441+2T>C	AAGGYACCGCGGGCCTCGCGCGG, AGGYACCGCGGGCCTCGCGCGGG	Myoglobinuria, acute recurrent, autosomal recessive
74315452	NM_000454.4(SOD1):c.338T>C (p.Ile113Thr)	TTGCAYCATTGGCCGCACACTGG	Amyotrophic lateral sclerosis type 1
730880455	NM_000169.2(GLA):c.41T>C (p.Leu14Pro)	CGCGCYTGCGCTTCGCTTCCTGG	not provided
267606656	NM_054027.4(ANKH):c.1015T>C (p.Cys339Arg)	AGCTCYGTTTCGTGATGTTTTGG	Craniometaphyseal dysplasia, autosomal dominant

267606687	NM_033409.3(SLC52A3):c.1238T>C (p.Val413Ala)	AGTTACGYCAAGGTGATGCTGGG	Brown-Vialetto-Van laere syndrome
267606721	NM_001928.2(CFD):c.640T>C (p.Cys214Arg)	GGTGYGCGGGGGCGTGCTCGAGG, GTGYGCGGGGGCGTGCTCGAGGG	Complement factor d deficiency
267606747	NM_001849.3(COL6A2):c.2329T>C (p.Cys777Arg)	CGCCYGCGACAAGCCACAGCAGG	Ullrich congenital muscular dystrophy
431905515	NM_001044.4(SLC6A3):c.671T>C (p.Leu224Pro)	CTGCACCYCCACCAGAGCCATGG	Infantile Parkinsonism-dystonia
267606857	NM_000180.3(GUCY2D):c.2846T>C (p.Ile949Thr)	AGAGAYCGCCAACATGTCACTGG	Cone-rod dystrophy 6
267606880	NM_022489.3(INF2):c.125T>C (p.Leu42Pro)	GCTGCYCCAGATGCCCTCTGTGG	Focal segmental glomerulosclerosis 5
515726191	NM_015713.4(RRM2B):c.581A>G (p.Glu194Gly)	AACTCCTYCTACAGCAGCAAAGG	RRM2B-related mitochondrial disease
267606917	NM_004646.3(NPHS1):c.793T>C (p.Cys265Arg)	GCTGCCGYGCGTGGCCCGAGGGG, CTGCCGYGCGTGGCCCGAGGGGG	Finnish congenital nephrotic syndrome
267607104	NM_001199107.1(TBC1D24):c.751T>C (p.Phe251Leu)	CAAGTTCYTCCACAAGGTGAGGG, TTCYTCCACAAGGTGAGGGCCGG	Myoclonic epilepsy, familial infantile
267607182	NM_144631.5(ZNF513):c.1015T>C (p.Cys339Arg)	TGGGCGCYGCATGCGAGGAGAGG, CGCYGCATGCGAGGAGAGGCTGG	Retinitis pigmentosa 58
267607211	NM_000229.1(LCAT):c.508T>C (p.Trp170Arg)	TATGACYGGCGGCTGGAGCCCGG	Norum disease
267607215	NM_016269.4(LEF1):c.181T>C (p.Ser61Pro)	GAACGAGYCTGAAATCATCCCGG	Sebaceous tumors, somatic
587783580	NM_178151.2(DCX):c.683T>C (p.Leu228Pro)	AAAAAACYCTACACTCTGGATGG	Heterotopia
587783644	NM_004004.5(GJB2):c.107T>C (p.Leu36Pro)	GATCCYCGTTGTGGCTGCAAAGG	Hearing impairment
587783653	NM_005682.6(ADGRG1):c.1460T>C (p.Leu487Pro)	CCCTGCYCACCTGCCTTTCCTGG	Polymicrogyria, bilateral frontoparietal
587783863	NM_000252.2(MTM1):c.958T>C (p.Ser320Pro)	GGAAYCTTTAAAAAAAGTGAAGG	Severe X-linked myotubular myopathy
267607751	NM_000249.3(MLH1):c.453+2T>C	ATCACGGYAAGAATGGTACATGG, TCACGGYAAGAATGGTACATGGG	Hereditary Nonpolyposis Colorectal Neoplasms
119103227	NM_000411.6(HLCS):c.710T>C (p.Leu237Pro)	CTATCYTTCTCAGGGAGGGAAGG	Holocarboxylase synthetase deficiency
119103237	NM_005787.5(ALG3):c.211T>C (p.Trp71Arg)	GATTGACYGGAAGGCCTACATGG	Congenital disorder of glycosylation type 1D
398122806	NM_003172.3(SURF1):c.679T>C (p.Trp227Arg)	CCACYGGCATTATCGAGACCTGG	Congenital myasthenic syndrome, acetazolamide-responsive
80338747	NM_004525.2(LRP2):c.7564T>C (p.Tyr2522His)	GTACCTGYACTGGGCTGACTGGG	Donnai Barrow syndrome
398122838	NM_001271723.1(FBXO38):c.616T>C (p.Cys206Arg)	TTCCTYGTATCCCAATGCTAAGG	Distal hereditary motor neuronopathy 2D
398122989	NM_014495.3(ANGPTL3):c.883T>C (p.Phe295Leu)	ACAAAACYTCAATGAAACGTGGG	Hypobetalipoproteinemia, familial, 2
80338945	NM_004004.5(GJB2):c.269T>C (p.Leu90Pro)	GCTCCYAGTGGCCATGCACGTGG	Deafness, autosomal recessive 1A, Hearing impairment
80338956	NM_000334.4(SCN4A):c.2078T>C (p.Ile693Thr)	AAGATCAYTGGCAATTCAGTGGG, AGATCAYTGGCAATTCAGTGGGG, GATCAYTGGCAATTCAGTGGGGG	Hyperkalemic Periodic Paralysis Type 1, Paramyotonia congenita of von Eulenburg
267608131	NM_000179.2(MSH6):c.4001+2T>C	CGGYAACTAACTAACTATAATGG	Hereditary Nonpolyposis Colorectal Neoplasms
587784573	NM_004963.3(GUCY2C):c.2782T>C (p.Cys928Arg)	TCCCYGTGCTGCTGGAGTTGTGG, CCCYGTGCTGCTGGAGTTGTGGG	Meconium ileus
267608511	NM_003159.2(CDKL5):c.659T>C (p.Leu220Pro)	CCAACYTTTTACTATTCAGAAGG	Early infantile epileptic encephalopathy 2
373842615	NM_000118.3(ENG):c.1273-2A>G	CCGCCYGCGGGGATAAAGCCAGG, CGCCYGCGGGGATAAAGCCAGGG	Haemorrhagic telangiectasia 1
185492581	NM_000335.4(SCN5A):c.376A>G (p.Lys126Glu)	GAATCTYCACAGCCGCTCTCCGG	Brugada syndrome
200533370	NM_133499.2(SYN1):c.1699A>G (p.Thr567Ala)	GATGYCTGACGGGTAGCCTGTGG, ATGYCTGACGGGTAGCCTGTGGG	Epilepsy, X-linked, with variable learning disabilities and behavior disorders, not specified
118203981	NM_148960.2(CLDN19):c.269T>C (p.Leu90Pro)	GCTCCYGGGCTTCGTGGCCATGG	Hypomagnesemia 5, renal, with ocular involvement
137853892	NM_001235.3(SERPINH1):c.233T>C (p.Leu78Pro)	GTCGCYAGGGCTCGTGTCGCTGG, TCGCYAGGGCTCGTGTCGCTGGG	Osteogenesis imperfecta type 10
118204024	NM_000263.3(NAGLU):c.142T>C (p.Phe48Leu)	GGCCGACYTCTCCGTGTCGGTGG	Mucopolysaccharidosis, MPS-III-B
690016563	NM_005211.3(CSF1R):c.1745T>C (p.Leu582Pro)	CAACCYGCAGTTTGGTGAGATGG	Hereditary diffuse leukoencephalopathy with spheroids
58380626	NM_000526.4(KRT14):c.1243T>C (p.Tyr415His)	CGCCACCYACCGCCGCCTGCTGG, CACCYACCGCCGCCTGCTGGAGG, ACCYACCGCCGCCTGCTGGAGGG	Epidermolysis bullosa herpetiformis, Dowling- Meara

113994151	NM_207346.2(TSEN54):c.277T>C (p.Ser93Pro)	TTGAAGYCTCCCGCGGTGAGCGG, AAGYCTCCCGCGGTGAGCGGCGG	Pontocerebellar hypoplasia type 4
113994206	NM_004937.2(CTNS):c.473T>C (p.Leu158Pro)	TGGTCYGAGCTTCGACTTCGTGG	Cystinosis

62516109	NM_000277.1(PAH):c.638T>C (p.Leu213Pro)	CCACTTCYTGAAAAGTACTGTGG	Phenylketonuria
370011798	NM_001302946.1(TRNT1):c.668T>C (p.Ile223Thr)	GCAAYTGCAGAAAATGCAAAAGG	Sideroblastic anemia with B-cell immunodeficiency, periodic fevers, and developmental delay
62517167	NM_000277.1(PAH):c.293T>C (p.Leu98Ser)	AAGATCTYGAGGCATGACATTGG	Mild non-PKU hyperphenylalanemia
12021720	NM_001918.3(DBT):c.1150G>A (p.Gly384Ser)	GACYCACAGAGCCCAATTTCTGG	Intermediate maple syrup urine disease type 2
104886289	NM_000495.4(COL4A5):c.4756T>C (p.Cys1586Arg)	TCCCCATYGTCCTCAGGGATGGG	Alport syndrome, X-linked recessive
370471013	NC_012920.1:m.5559A>G	CAACYTACTGAGGGCTTTGAAGG	Leigh disease
121434215	NM_000487.5(ARSA):c.410T>C (p.Leu137Pro)	GCCTTCCYGCCCCCCCATCAGGG	Metachromatic leukodystrophy, adult type
386134128	NM_000096.3(CP):c.1123T>C (p.Tyr375His)	ACACTACYACATTGCCGCTGAGG	Deficiency of ferroxidase
121434275	NM_001127328.2(ACADM):c.1136T>C (p.Ile379Thr)	GTGCAGAYACTTGGAGGCAATGG	Medium-chain acyl-coenzyme A dehydrogenase deficiency
121434276	NM_001127328.2(ACADM):c.742T>C (p.Cys248Arg)	CAGCGAYGTTCAGATACTAGAGG	Medium-chain acyl-coenzyme A dehydrogenase deficiency
121434284	NM_002225.3(IVD):c.134T>C (p.Leu45Pro)	ATGGGCYAAGCGAGGAGCAGAGG	ISOVALERIC ACIDEMIA, TYPE I
121434334	NM_005908.3(MANBA):c.1513T>C (p.Ser505Pro)	ATTACGYCCAGTCCTACAAATGG, TTACGYCCAGTCCTACAAATGGG, TACGYCCAGTCCTACAAATGGGG	Beta-D-mannosidosis
121434366	NM_000159.3(GCDH):c.883T>C (p.Tyr295His)	CGCCCGGYACGGCATCGCGTGGG, GCCCGGYACGGCATCGCGTGGGG	Glutaric aciduria, type 1
60715293	NM_000424.3(KRT5):c.541T>C (p.Ser181Pro)	GTTTGCCYCCTTCATCGACAAGG	Epidermolysis bullosa herpetiformis, Dowling- Meara
121434409	NM_001003722.1(GLE1):c.2051T>C (p.Ile684Thr)	AAGGACAYTCCTGTCCCCAAGGG	Lethal arthrogryposis with anterior horn cell disease
121434434	NM_001287.5(CLCN7):c.2297T>C (p.Leu766Pro)	GGGCCYGCGGCACCTGGTGGTGG	Osteopetrosis autosomal recessive 4
121434455	NM_000466.2(PEX1):c.1991T>C (p.Leu664Pro)	GATGACCYTGACCTCATTGCTGG	Zellweger syndrome
199422317	NM_001099274.1(TINF2):c.862T>C (p.Phe288Leu)	CTGYTTCCCTTTAGGAATCTCGG	Aplastic anemia
104895221	NM_001065.3(TNFRSF1A):c.349T>C (p.Cys117Arg)	CTCTTCTYGCACAGTGGACCGGG	TNF receptor-associated periodic fever syndrome (TRAPS)
137854459	NM_000138.4(FBN1):c.4987T>C (p.Cys1663Arg)	GGGACAYGTTACAACACCGTTGG	Marfan syndrome
387907075	NM_024027.4(COLEC11):c.505T>C (p.Ser169Pro)	CAGCTGYCCTGCCAGGGCCGCGG, AGCTGYCCTGCCAGGGCCGCGGG, GCTGYCCTGCCAGGGCCGCGGGG, CTGYCCTGCCAGGGCCGCGGGGG	Carnevale syndrome
1048095	NM_000352.4(ABCC8):c.674T>C (p.Leu225Pro)	TGCYGTCCAAAGGCACCTACTGG	Permanent neonatal diabetes mellitus
796065347	NM_019074.3(DLL4):c.1168T>C (p.Cys390Arg)	GAAYGTCCCCCCAACTTCACCGG	Adams-Oliver syndrome, ADAMS-OLIVER SYNDROME 6
137852347	NM_000402.4(G6PD):c.1054T>C (p.Tyr352His)	AGGGYACCTGGACGACCCCACGG	Anemia, nonspherocytic hemolytic, due to G6PD deficiency
74315327	NM_213653.3(HFE2):c.302T>C (p.Leu101Pro)	GGACCYCGCCTTCCATTCGGCGG	Hemochromatosis type 2A
137852579	NM_000044.3(AR):c.2033T>C (p.Leu678Pro)	GTCCYGGAAGCCATTGAGCCAGG	
137852636	NM_001166107.1(HMGCS2):c.520T>C (p.Phe174Leu)	CCCTCYTCAATGCTGCCAACTGG	mitochondrial 3-hydroxy-3-methylglutaryl-CoA synthase deficiency
137852661	NM_033163.3(FGF8):c.118T>C (p.Phe40Leu)	TTCCCTGYTCCGGGCTGGCCGGG	Kallmann syndrome 6
121912967	NM_005215.3(DCC):c.503T>C (p.Met168Thr)	AGCCCAYGCCAACAATCCACTGG	
137852806	NM_001039523.2(CHRNA1):c.901T>C (p.Phe301Leu)	TGTGYTCCTTCTGGTCATCGTGG	Myasthenic syndrome, congenital, fast-channel
137852850	NM_182760.3(SUMF1):c.463T>C (p.Ser155Pro)	GGCGACYCCTTTGTCTTTGAAGG	Multiple sulfatase deficiency

137852886	NM_000158.3(GBE1):c.671T>C (p.Leu224Pro)	AATGTACYACCAAGAATCAAAGG	Glycogen storage disease, type IV, GLYCOGEN STORAGE DISEASE IV, NONPROGRESSIVE HEPATIC
137852911	NM_000419.3(ITGA2B):c.641T>C (p.Leu214Pro)	CTGGTGCYTGGGGCTCCTGGCGG	Glanzmann thrombasthenia
137852948	NM_138694.3(PKHD1):c.10658T>C (p.Ile3553Thr)	GAGCCCAYTGAAATACGCTCAGG	Polycystic kidney disease, infantile type
137852964	NM_024960.4(PANK2):c.178T>C (p.Ser60Pro)	ATTGACYCAGTCGGATTCAATGG	
137853020	NM_006899.3(IDH3B):c.395T>C (p.Leu132Pro)	TGCGGCYGAGGTAGGTGGTCTGG, GCGGCYGAGGTAGGTGGTCTGGG	Retinitis pigmentosa 46
137853249	NM_033500.2(HK1):c.1550T>C (p.Leu517Ser)	GACTTCTYGGCCCTGGATCTTGG, TTCTYGGCCCTGGATCTTGGAGG	Hemolytic anemia due to hexokinase deficiency

137853270	NM_000444.5(PHEX):c.1664T>C (p.Leu555Pro)	AGCYCCAGAAGCCTTTCTTTTGG	Familial X-linked hypophosphatemic vitamin D refractory rickets
137853325	NM_003639.4(IKBKG):c.1249T>C (p.Cys417Arg)	TGGAGYGCATTGAGTAGGGCCGG	Hypohidrotic ectodermal dysplasia with immune deficiency, Hyper-IgM immunodeficiency, X-linked, with hypohidrotic ectodermal dysplasia
28932769	NM_002055.4(GFAP):c.1055T>C (p.Leu352Pro)	GGACCYGCTCAATGTCAAGCTGG	Alexander disease
397507439	NM_002769.4(PRSS1):c.116T>C (p.Val39Ala)	TACCAGGYGTCCCTGAATTCTGG	Hereditary pancreatitis
387906446	NM_000132.3(F8):c.1729T>C (p.Ser577Pro)	AAAGAAYCTGTAGATCAAAGAGG	Hereditary factor VIII deficiency disease
387906482	NM_000133.3(F9):c.1031T>C (p.Ile344Thr)	ACGAACAYCTTCCTCAAATTTGG	Hereditary factor IX deficiency disease
387906508	NM_000131.4(F7):c.983T>C (p.Phe328Ser)	GACGTYCTCTGAGAGGACGCTGG	Factor VII deficiency
387906532	NM_001040113.1(MYH11):c.3791T>C (p.Leu1264Pro)	GAAGCYGGAGGCGCAGGTGCAGG	Aortic aneurysm, familial thoracic 4
387906658	NM_002465.3(MYBPC1):c.2566T>C (p.Tyr856His)	CAAACCYATATCCGCAGAGTTGG	Distal arthrogryposis type 1B
387906701	NM_003491.3(NAA10):c.109T>C (p.Ser37Pro)	TGGCCTTYCCTGGCCCCAGGTGG, GGCCTTYCCTGGCCCCAGGTGGG	N-terminal acetyltransferase deficiency
387906717	NM_000377.2(WAS):c.881T>C (p.Ile294Thr)	GACTTCAYTGAGGACCAGGGTGG, ACTTCAYTGAGGACCAGGGTGGG	Severe congenital neutropenia X-linked
387906809	NM_000287.3(PEX6):c.1601T>C (p.Leu534Pro)	CTTCYGGGCCGGGACCGTGATGG, TTCYGGGCCGGGACCGTGATGGG	Peroxisome biogenesis disorder 4B
387906965	NM_024513.3(FYCO1):c.4127T>C (p.Leu1376Pro)	CAGCCYGATCCCCATCACTGTGG	Cataract, autosomal recessive congenital 2
387906967	NM_006147.3(IRF6):c.65T>C (p.Leu22Pro)	GCCYCTACCCTGGGCTCATCTGG	Van der Woude syndrome, Popliteal pterygium syndrome
387906982	NM_025132.3(WDR19):c.20T>C (p.Leu7Pro)	TCTCACYGCTAGAAAAGACTTGG	Asphyxiating thoracic dystrophy 5
387907072	NM_032446.2(MEGF10):c.2320T>C (p.Cys774Arg)	GGGCAGYGTACTTGCCGCACTGG	Myopathy, areflexia, respiratory distress, and dysphagia, early-onset, Myopathy, areflexia, respiratory distress, and dysphagia, early-onset, mild variant
137854499	NM_005502.3(ABCA1):c.6026T>C (p.Phe2009Ser)	GAGTYCTTTGCCCTTTTGAGAGG	Familial hypoalphalipoproteinemia
387907117	NM_000196.3(HSD11B2):c.1012T>C (p.Tyr338His)	CCGCCGCYATTACCCCGGCCAGG, CGCCGCYATTACCCCGGCCAGGG	Apparent mineralocorticoid excess
387907170	NM_004453.3(ETFDH):c.1130T>C (p.Leu377Pro)	CCAAAACYCACCTTTCCTGGTGG	
387907205	NM_033360.3(KRAS):c.211T>C (p.Tyr71His)	GGACCAGYACATGAGGACTGGGG, CCAGYACATGAGGACTGGGGAGG, CAGYACATGAGGACTGGGGAGGG	Cardiofaciocutaneous syndrome 2
387907240	NM_024110.4(CARD14):c.467T>C (p.Leu156Pro)	CAGCAGCYGCAGGAGCACCTGGG	Pityriasis rubra pilaris
387907282	NM_152296.4(ATP1A3):c.2431T>C (p.Ser811Pro)	TGCCATCYCACTGGCGTACGAGG	Alternating hemiplegia of childhood 2
387907361	NM_005120.2(MED12):c.3493T>C (p.Ser1165Pro)	AGGACYCTGAGCCAGGGGCCCGG	Ohdo syndrome, X-linked
28933970	NM_006194.3(PAX9):c.62T>C (p.Leu21Pro)	GGCCGCYGCCCAACGCCATCCGG	Tooth agenesis, selective, 3
137854472	NM_000138.4(FBN1):c.3128A>G (p.Lys1043Arg)	TGCACYTGCCGTGGGTGCAGAGG	
727504261	NM_000257.3(MYH7):c.2708A>G (p.Glu903Gly)	AGCGCYCCTCAGCATCTGCCAGG	Cardiomyopathy, not specified
81002853	NM_000059.3(BRCA2):c.476-2A>G	ACCACYGGGGGTAAAAAAAGGGG, TACCACYGGGGGTAAAAAAAGGG,	Familial cancer of breast, Breast-ovarian cancer, familial 2, Hereditary cancer-predisposing syndrome

119473032	NM_021020.3(LZTS1):c.355A>G (p.Lys119Glu)	CCCTYCTCGGAGCCCTGTAGAGG	
193922801	NM_000540.2(RYR1):c.7043A>G (p.Glu2348Gly)	TTCYCCTCCACGCTCTCGCCTGG	not provided
36210419	NM_000218.2(KCNQ1):c.652A>G (p.Lys218Glu)	GCCCCTYGGAGCCCACGCAGAGG	Torsades de pointes, Cardiac arrhythmia
121964989	NM_000108.4(DLD):c.1483A>G (p.Arg495Gly)	TTCTCYAAAAGCTTCTGATAAGG	Maple syrup urine disease, type 3
28936669	NM_000095.2(COMP):c.1418A>G (p.Asp473Gly)	ATTGYCGTCGTCGTCGTCGCAGG	
28936696	NM_018488.2(TBX4):c.1592A>G (p.Gln531Arg)	GTACYGTAAGGAAGATTCTCGGG, GGTACYGTAAGGAAGATTCTCGG	Ischiopatellar dysplasia
121965077	NM_000137.2(FAH):c.1141A>G (p.Arg381Gly)	TCCYGGTCTGACCATTCCCCAGG	Tyrosinemia type I
794728203	NM_000138.4(FBN1):c.3344A>G (p.Asp1115Gly)	ACTCAYCAATATCTGCAAAATGG	Thoracic aortic aneurysms and aortic dissections
786205436	NM_003002.3(SDHD):c.275A>G (p.Asp92Gly)	GAATAGYCCATCGCAGAGCAAGG	Fatal infantile mitochondrial cardiomyopathy
72551317	NM_000784.3(CYP27A1):c.776A>G (p.Lys259Arg)	AGTCCACYTGGGGAGGAAGGTGG	Cholestanol storage disease

786205687	NM_016218.2(POLK):c.1385A>G (p.Asn462Ser)	ATTCACAYTCTTCAACTTAATGG	Malignant tumor of prostate
794728280	NM_000138.4(FBN1):c.7916A>G (p.Tyr2639Cys)	TGTTCAYACTGGAAGCCGGCGGG, CTGTTCAYACTGGAAGCCGGCGG	Thoracic aortic aneurysms and aortic dissections
28937317	NM_000335.4(SCN5A):c.3971A>G (p.Asn1324Ser)	GCAYTGACCACCACCTCAAGTGG	Long QT syndrome 3, Congenital long QT syndrome
786205854	NM_144499.2(GNAT1):c.386A>G (p.Asp129Gly)	CGGAGYCCTTCCACAGCCGCTGG	NIGHT BLINDNESS, CONGENITAL
104893776	NM_000539.3(RHO):c.533A>G (p.Tyr178Cys)	GGATGYACCTGAGGACAGGCAGG	Retinitis pigmentosa 4
28937590	NM_001257342.1(BCS1L):c.232A>G (p.Ser78Gly)	GACACYGAGGTGCTGAGTACGGG, CGACACYGAGGTGCTGAGTACGG	GRACILE syndrome
104893866	NM_000320.2(QDPR):c.449A>G (p.Tyr150Cys)	TGCCGYACCCGATCATACCTGGG, ATGCCGYACCCGATCATACCTGG	Dihydropteridine reductase deficiency
587776590	NM_015629.3(PRPF31):c.527+3A>G	GACAYACCCCTGGGTGGTGGAGG, GCGGACAYACCCCTGGGTGGTGG	Retinitis pigmentosa 11
104894015	NM_000162.3(GCK):c.641A>G (p.Tyr214Cys)	GTAGYAGCAGGAGATCATCGTGG	Hyperinsulinemic hypoglycemia familial 3
202247823	NM_000532.4(PCCB):c.1606A>G (p.Asn536Asp)	ATATYTGCATGTTTTCTCCAAGG	Propionic acidemia
104894199	NM_000073.2(CD3G):c.1A>G (p.Met1Val)	CCAYGTCAGTCTCTGTCCTCCGG	Immunodeficiency 17
104894208	NM_001814.4(CTSC):c.857A>G (p.Gln286Arg)	CTCCYGAGGGCTTAGGATTGGGG, CCTCCYGAGGGCTTAGGATTGGG, ACCTCCYGAGGGCTTAGGATTGG	Papillon-Lef\xc3\xa8vre syndrome, Haim-Munk syndrome
104894211	NM_001814.4(CTSC):c.1040A>G (p.Tyr347Cys)	TCCTACAYAGTGGTACTCAGAGG	Papillon-Lef\xc3\xa8vre syndrome, Periodontitis,
104894290	NM_000448.2(RAG1):c.2735A>G (p.Tyr912Cys)	CTGYACTGGCAGAGGGATTCTGG	Histiocytic medullary reticulosis
104894354	NM_000217.2(KCNA1):c.676A>G (p.Thr226Ala)	GCGYTTCCACGATGAAGAAGGGG, AGCGYTTCCACGATGAAGAAGGG, CAGCGYTTCCACGATGAAGAAGG	Episodic ataxia type 1
104894425	NM_014239.3(EIF2B2):c.638A>G (p.Glu213Gly)	AGTTGTCYCAATACCTGCTTTGG	Leukoencephalopathy with vanishing white matter, Ovarioleukodystrophy
104894450	NM_000270.3(PNP):c.383A>G (p.Asp128Gly)	ATAYCTCCAACCTCAAACTTGGG, GATAYCTCCAACCTCAAACTTGG	Purine-nucleoside phosphorylase deficiency
147394623	NM_024887.3(DHDDS):c.124A>G (p.Lys42Glu)	GGCACTYCTTGGCATAGCGACGG	Retinitis pigmentosa 59
60723330	NM_005557.3(KRT16):c.374A>G (p.Asn125Ser)	GCGGTCAYTGAGGTTCTGCATGG	Pachyonychia congenita, type 1, Palmoplantar keratoderma, nonepidermolytic, focal
104894634	NM_030665.3(RAI1):c.4685A>G (p.Gln1562Arg)	CTGCTGCYGTCGTCGTCGCTTGG	Smith-Magenis syndrome
104894730	NM_000363.4(TNNI3):c.532A>G (p.Lys178Glu)	CCTYCTTCACCTGCTTGAGGTGG, CCTCCTYCTTCACCTGCTTGAGG	Familial restrictive cardiomyopathy 1
104894816	NM_002049.3(GATA1):c.653A>G (p.Asp218Gly)	GTCCTGYCCCTCCGCCACAGTGG	GATA-1-related thrombocytopenia with dyserythropoiesis
794726773	NM_001165963.1(SCN1A):c.1662+3A>G	GTGCCAYACCTGGTGTGGGGAGG	Severe myoclonic epilepsy in infancy

104894861	NM_000202.6(IDS):c.404A>G (p.Lys135Arg)	AAAGACTYTTCCCACCGACATGG	Mucopolysaccharidosis, MPS-II
104894874	NM_000266.3(NDP):c.125A>G (p.His42Arg)	TGGYGCCTCATGCAGCGTCGAGG	
191205969	NM_002420.5(TRPM1):c.296T>C (p.Leu99Pro)	AAGCYCTTAATATCTGTGCATGG	Congenital stationary night blindness, type 1C
794727073	NM_019109.4(ALG1):c.1188-2A>G	TAAACYGCAGAGAGAACCAAGGG, GTAAACYGCAGAGAGAACCAAGG	Congenital disorder of glycosylation type 1K
281875236	NM_001004334.3(GPR179):c.659A>G (p.Tyr220Cys)	CCCACAYATCCATCTGCCTGCGG	Congenital stationary night blindness, type 1E
28939094	NM_015915.4(ATL1):c.1222A>G (p.Met408Val)	CACCCAYCTTCTTCACCCCTCGG	Spastic paraplegia 3
281875324	NM_005359.5(SMAD4):c.989A>G (p.Glu330Gly)	ATCCATTYCAAAGTAAGCAATGG	Juvenile polyposis syndrome, Hereditary cancer-predisposing syndrome
77173848	NM_000037.3(ANK1):c.-[illegible]	GGGCCYGGCCCGCACGTCACAGG	Spherocytosis, type 1, autosomal recessive
150181226	NM_001159772.1(CANT1):c.671T>C (p.Leu224Pro)	CGTCYGTACGTGGGCGGCCTGGG, GCGTCYGTACGTGGGCGGCCTGG	Desbuquois syndrome
397514253	NM_000041.3(APOE):c.237-2A>G	CGCCCYGCGGCCGAGAGGGCGGG, GCGCCCYGCGGCCGAGAGGGCGG	Familial type 3 hyperlipoproteinemia
397514348	NM_000060.3(BTD):c.278A>G (p.Tyr93Cys)	GTTCAYAGATGTCAAGGTTCTGG	Biotinidase deficiency
397514415	NM_000060.3(BTD):c.1313A>G (p.Tyr438Cys)	GGCAYACAGCTCTTTGGATAAGG	Biotinidase deficiency
397514501	NM_007171.3(POMT1):c.430A>G (p.Asn144Asp)	GAGCATYCTCTGTTTCAAAGAGG	Limb-girdle muscular dystrophy-
370382601	NM_174917.4(ACSF3):c.1A>G (p.Met1Val)	GGCAGCAYTGCACTGACAGGCGG	not provided
72554332	NM_000531.5(OTC):c.238A>G (p.Lys80Glu)	AAGGACTYCCCTTGCAATAAAGG	Ornithine carbamoyltransferase deficiency
397514599	NM_033109.4(PNPT1):c.1424A>G	GACTYCAGATGTAACTCTTATGG	Deafness, autosomal recessive 70

	(p.Glu475Gly)		
397514650	NM_000108.4(DLD):c.1444A>G (p.Arg482Gly)	GACTCYAGCTATATCTTCACAGG	Maple syrup urine disease, type 3
397514675	NM_003156.3(STIM1):c.251A>G (p.Asp84Gly)	TTCCACAYCCACATCACCATTGG	Myopathy with tubular aggregates
794728378	NM_000238.3(KCNH2):c.1913A>G (p.Lys638Arg)	ATCYTCTCTGAGTTGGTGTTGGG, GATCYTCTCTGAGTTGGTGTTGG	Cardiac arrhythmia
397514711	NM_002163.2(IRF8):c.238A>G (p.Thr80Ala)	AACCTCGYCTTCCAAGTGGCTGG	Autosomal dominant CD11C+/CD1C+ dendritic cell deficiency
397514729	NM_000388.3(CASR):c.85A>G (p.Lys29Glu)	CCCCCTYCTTTTGGGCTCGCTGG	Hypocalcemia, autosomal dominant 1, with bartter syndrome
397514743	NM_022114.3(PRDM16):c.2447A>G (p.Asn816Ser)	GCCGCCGYTTTGGCTGGCACGGG	Left ventricular noncompaction 8
397514757	NM_005689.2(ABCB6):c.508A>G (p.Ser170Gly)	TGGGCYGTTCCAAGACACCAGGG, GTGGGCYGTTCCAAGACACCAGG	Dyschromatosis universalis hereditaria 3
28940313	NM_152443.2(RDH12):c.677A>G (p.Tyr226Cys)	CACTGCGYAGGTGGTGACCCCGG	Leber congenital amaurosis 13
794728538	NM_000218.2(KCNQ1):c.1787A>G (p.Glu596Gly)	GTCTYCTACTCGGTTCAGGCGGG, TGTCTYCTACTCGGTTCAGGCGG	Cardiac arrhythmia
794728569	NM_000218.2(KCNQ1):c.605A>G (p.Asp202Gly)	AGGYCTGTGGAGTGCAGGAGAGG	Cardiac arrhythmia
794728573	NM_000218.2(KCNQ1):c.1515-2A>G	GCCYGCAGTGGAGAGAGGAGAGG	Cardiac arrhythmia
370874727	NM_003494.3(DYSF):c.3349-2A>G	CCGCCCYGGAGACACGAAGCTGG	Limb-girdle muscular dystrophy, type 2B
794728859	NM_198056.2(SCN5A):c.2788-2A>G	ACCYGTCGAGATAATGGGTCAGG	not provided
794728887	NM_198056.2(SCN5A):c.4462A>G (p.Thr1488Ala)	CCTCTGYCATGAAGATGTCCTGG	not provided
28940878	NM_000372.4(TYR):c.125A>G (p.Asp42Gly)	CTCCTGYCCCCGCTCCACGGTGG	Tyrosinase-negative oculocutaneous albinism
397515420	NM_172107.2(KCNQ2):c.1636A>G (p.Met546Val)	GCAYGACACTGCAGGGGGGTGGG, CGCAYGACACTGCAGGGGGGTGG, AACCGCAYGACACTGCAGGGGGG	Early infantile epileptic encephalopathy 7

397515428	NM_001410.2(MEGF8):c.7099A>G (p.Ser2367Gly)	GACYCCCGTGAAATGATTCCCGG	Carpenter syndrome 2
143601447	NM_201631.3(TGM5):c.122T>C (p.Leu41Pro)	TCAACCYCACCCTGTACTTCAGG	Peeling skin syndrome, acral type
397515519	NM_000207.2(INS):c.*59A>G	GGGCYTTATTCCATCTCTCTCGG	Permanent neonatal diabetes mellitus
397515523	NM_000370.3(TTPA):c.191A>G (p.Asp64Gly)	CAGGYCCAGATCGAAATCCCGGG, CCAGGYCCAGATCGAAATCCCGG	Ataxia with vitamin E deficiency
397515891	NM_000256.3(MYBPC3):c.1224-2A>G	TACTTGCYGTAGAACAGAAGGGG	Familial hypertrophic cardiomyopathy 4, Cardiomyopathy
397516082	NM_000256.3(MYBPC3):c.927-2A>G	GTCCCYGTGTCCCGCAGTCTAGG	Familial hypertrophic cardiomyopathy 4, Cardiomyopathy
397516138	NM_000257.3(MYH7):c.2206A>G (p.Ile736Val)	TATCAAYGAACTGTCCCTCAGGG, CTATCAAYGAACTGTCCCTCAGG	Familial hypertrophic cardiomyopathy 1, Cardiomyopathy, not specified
1154510	NM_002150.2(HPD):c.97G>A (p.Ala33Thr)	ATGACGYGGCCTGAATCACAGGG, AATGACGYGGCCTGAATCACAGG	4-Alpha-hydroxyphenylpyruvate hydroxylase deficiency
397516330	NM_000260.3(MYO7A):c.6439-2A>G	ATATCCYGGGGGAGCAGAAAGGG, GATATCCYGGGGGAGCAGAAAGG	Usher syndrome, type 1
72556271	NM_000531.5(OTC):c.482A>G (p.Asn161Ser)	CAGCCCAYTGATAATTGGGATGG	not provided
606231260	NM_023073.3(C5orf42):c.3290-2A>G	ATCYATCAAATACAAAAATTTGG	Orofaciodigital syndrome 6
587777521	NM_004817.3(TJP2):c.1992-2A>G	CAGCTCYGAGAAGAAACCACGGG, TCAGCTCYGAGAAGAAACCACGG	Progressive familial intrahepatic cholestasis 4
730880846	NM_000257.3(MYH7):c.617A>G (p.Lys206Arg)	CTTCYTGCTGCGGTCCCCAATGG	Cardiomyopathy
397517978	NM_206933.2(USH2A):c.12067-2A>G	TTCCCYGTAAGAAAATTAACAGG	Usher syndrome, type 2A, Retinitis pigmentosa 39
606231409	NM_000216.2(ANOS1):c.1A>G (p.Met1Val)	GCACCAYGGCTGCGGGTCGAGGG, GGCACCAYGGCTGCGGGTCGAGG	Kallmann syndrome 1
80356546	NM_003334.3(UBA1):c.1639A>G (p.Ser547Gly)	TGGCYTGTCACCCGGATATGTGG	Arthrogryposis multiplex congenita, distal, X- linked
80356584	NM_194248.2(OTOF):c.766-2A>G	GACCYGCAGGCAGGAGAAGGGGG, TGACCYGCAGGCAGGAGAAGGGG, CTGACCYGCAGGCAGGAGAAGGG, GCTGACCYGCAGGCAGGAGAAGG	Deafness, autosomal recessive 9
730880930	NM_000257.3(MYH7):c.1615A>G (p.Met539Val)	GGAACAYGCACTCCTCTTCCAGG	Cardiomyopathy
118203947	NM_013319.2(UBIAD1):c.355A>G (p.Arg119Gly)	TCCYGTCATCACTCTTTTTGTGG	Schnyder crystalline corneal dystrophy
60171927	NM_000526.4(KRT14):c.368A>G (p.Asn123Ser)	GCGGTCAYTGAGGTTCTGCATGG	Epidermolysis bullosa herpetiformis, Dowling- Meara
199422248	NM_001363.4(DKC1):c.941A>G (p.Lys314Arg)	AATCYTGGCCCCATAGCAGATGG	Dyskeratosis congenita X-linked

72558467	NM_000531.5(OTC):c.929A>G (p.Glu310Gly)	TCCACTYCTTCTGGCTTTCTGGG, ATCCACTYCTTCTGGCTTTCTGG	not provided
72558478	NM_000531.5(OTC):c.988A>G (p.Arg330Gly)	ACTTTCYGTTTTCTGCCTCTGGG, CACTTTCYGTTTTCTGCCTCTGG	not provided
118204455	NM_000505.3(F12):c.158A>G (p.Tyr53Cys)	GGTGGYACTGGAAGGGGAAGTGG	
80357477	NM_007294.3(BRCA1):c.5453A>G (p.Asp1818Gly)	TTGYCCTCTGTCCAGGCATCTGG	Familial cancer of breast, Breast-ovarian cancer, familial 1
121907908	NM_024426.4(WT1):c.1021A>G (p.Ser341Gly)	CGCYCTCGTACCCTGTGCTGTGG	Mesothelioma
121907926	NM_000280.4(PAX6):c.1171A>G (p.Thr391Ala)	GTGGYGCCCGAGGTGCCCATTGG	Optic nerve aplasia, bilateral
121908023	NM_024740.2(ALG9):c.860A>G (p.Tyr287Cys)	TTAYACAAAACAATGTTGAGTGG	Congenital disorder of glycosylation type 1L
121908148	NM_001243133.1(NLRP3):c.1880A>G (p.Glu627Gly)	ACAATYCCAGCTGGCTGGGCTGG	Familial cold urticaria
121908166	NM_006492.2(ALX3):c.608A>G (p.Asn203Ser)	CGGYTCTGGAACCAGACCTGGGG, GCGGYTCTGGAACCAGACCTGGG, TGCGGYTCTGGAACCAGACCTGG	Frontonasal dysplasia 1

121908184	NM_020451.2(SEPN1):c.1A>G (p.Met1Val)	CCCAYGGCTGCGGCTGGCGGCGG, CGGCCCAYGGCTGCGGCTGGCGG	Eichsfeld type congenital muscular dystrophy
121908258	NM_130468.3(CHST14):c.878A>G (p.Tyr293Cys)	AAGTCAYAGTGCACGGCACAAGG	Ehlers-Danlos syndrome, musculocontractural type
121908383	NM_001128425.1(MUTYH):c.1241A>G (p.Gln414Arg)	AAGCYGCTCTGAGGGCTCCCAGG	Neoplasm of stomach
121908580	NM_004328.4(BCS1L):c.148A>G (p.Thr50Ala)	GTGYGATCATGTAATGGCGCCGG	Mitochondrial complex III deficiency
121908584	NM_016417.2(GLRX5):c.294A>G (p.Gln98=)	CCTGACCYTGTCGGAGCTCCGGG	Anemia, sideroblastic, pyridoxine-refractory, autosomal recessive
121908635	NM_022817.2(PER2):c.1984A>G (p.Ser662Gly)	GCCACACYCTCTGCCTTGCCCGG	Advanced sleep phase syndrome, familial
121908655	NM_003839.3(TNFRSF11A):c.508A>G (p.Arg170Gly)	GGGTCYGCATTTGTCCGTGGAGG	Osteopetrosis autosomal recessive 7
29001653	NM_000539.3(RHO):c.886A>G (p.Lys296Glu)	CGCTCTYGGCAAAGAACGCTGGG, GCGCTCTYGGCAAAGAACGCTGG	Retinitis pigmentosa 4
56307355	NM_006502.2(POLH):c.1603A>G (p.Lys535Glu)	AGACTTTYCTGCTTAAAGAAGGG	Xeroderma pigmentosum, variant type
121908919	NM_002977.3(SCN9A):c.1964A>G (p.Lys655Arg)	CCTTTTCYTGTGTATTTGATTGG	Generalized epilepsy with febrile seizures plus, type 7, not specified
121908939	NM_006892.3(DNMT3B):c.2450A>G (p.Asp817Gly)	GACACGYCTGTGTAGTGCACAGG	Centromeric instability of chromosomes 1,9 and 16 and immunodeficiency
121909088	NM_001005360.2(DNM2):c.1684A>G (p.Lys562Glu)	ACTYCTTCTCTTTCTCCTGAGGG, TACTYCTTCTCTTTCTCCTGAGG	Charcot-Marie-Tooth disease, dominant intermediate b, with neutropenia
120074112	NM_000483.4(APOC2):c.1A>G (p.Met1Val)	GCCCAYAGTGTCCAGAGACCTGG	Apolipoprotein C2 deficiency
121909239	NM_000314.6(PTEN):c.755A>G (p.Asp252Gly)	ATAYCACCACACACAGGTAACGG	Macrocephaly/autism syndrome
121909251	NM_198217.2(ING1):c.515A>G (p.Asn172Ser)	TGGYTGCACAGACAGTACGTGGG, CTGGYTGCACAGACAGTACGTGG	Squamous cell carcinoma of the head and neck
121909396	NM_001174089.1(SLC4A11):c.2518A>G (p.Met840Val)	GATCAYCTTCATGTAGGGCAGGG, AGATCAYCTTCATGTAGGGCAGG	Corneal dystrophy and perceptive deafness
121909533	NM_000034.3(ALDOA):c.386A>G (p.Asp129Gly)	CCAYCCAACCCTAAGAGAAGAGG	HNSHA due to aldolase A deficiency
128627255	NM_004006.2(DMD):c.835A>G (p.Thr279Ala)	TGACCGYGATCTGCAGAGAAGGG, CTGACCGYGATCTGCAGAGAAGG	Dilated cardiomyopathy 3B
116929575	NM_001085.4(SERPINA3):c.1240A>G (p.Met414Val)	GCTCAYGAAGAAGATGTTCTGGG, TGCTCAYGAAGAAGATGTTCTGG	
61748392	NM_004992.3(MECP2):c.410A>G (p.Glu137Gly)	CAACYCCACTTTAGAGCGAAAGG	Mental retardation, X-linked, syndromic 13
61748906	NM_001005741.2(GBA):c.667T>C (p.Trp223Arg)	CCCACTYGGCTCAAGACCAATGG	Gaucher disease, type 1
199473024	NM_000238.3(KCNH2):c.3118A>G (p.Ser1040Gly)	CTGCYCTCCACGTCGCCCCGGGG, CCTGCYCTCCACGTCGCCCCGGG, GCCTGCYCTCCACGTCGCCCCGG	Sudden infant death syndrome
794728365	NM_000238.3(KCNH2):c.1129-2A>G	GGACCYGCACCCGGGGAAGGCGG	Cardiac arrhythmia
72556293	NM_000531.5(OTC):c.548A>G (p.Tyr183Cys)	AGAGCTAYAGTGTTCCTAAAAGG	not provided
111033244	NM_000441.1(SLC26A4):c.1151A>G (p.Glu384Gly)	TGAATYCCTAAGGAAGAGACTGG	Pendred syndrome, Enlarged vestibular aqueduct syndrome
111033415	NM_000260.3(MYO7A):c.1344-2A>G	AGCYGCAGGGGCACAGGGATGGG, AAGCYGCAGGGGCACAGGGATGG	Usher syndrome, type 1
121912439	NM_000454.4(SOD1):c.302A>G (p.Glu101Gly)	AGAATCTYCAATAGACACATCGG	Amyotrophic lateral sclerosis type 1
111033567	NM_002769.4(PRSS1):c.68A>G (p.Lys23Arg)	ATCYTGTCATCATCATCAAAGGG, GATCYTGTCATCATCATCAAAGG	Hereditary pancreatitis
121912565	NM_000901.4(NR3C2):c.2327A>G (p.Gln776Arg)	TCATCYGTTTGCCTGCTAAGCGG	Pseudohypoaldosteronism type 1 autosomal dominant

121912574	NM_000901.4(NR3C2):c.2915A>G (p.Glu972Gly)	CCGACYCCACCTTGGGCAGCTGG	Pseudohypoaldosteronism type 1 autosomal dominant
121912589	NM_001173464.1(KIF21A):c.2839A>G (p.Met947Val)	ATTCAYATCTGCCTCCATGTTGG	Fibrosis of extraocular muscles, congenital, 1
111033661	NM_000155.3(GALT):c.253-2A>G	ATTCACCYACCGACAAGGATAGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
111033669	NM_000155.3(GALT):c.290A>G (p.Asn97Ser)	GAAGTCGYTGTCAAACAGGAAGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase

111033682	NM_000155.3(GALT):c.379A>G (p.Lys127Glu)	TGACCTYACTGGGTGGTGACGGG, ATGACCTYACTGGGTGGTGACGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
111033786	NM_000155.3(GALT):c.950A>G (p.Gln317Arg)	CAGCYGCCAATGGTTCCAGTTGG	Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase
121912765	NM_001202.3(BMP4):c.278A>G (p.Glu93Gly)	CCTCCYCCCCAGACTGAAGCCGG	Microphthalmia syndromic 6
121912856	NM_000094.3(COL7A1):c.425A>G (p.Lys142Arg)	CACCYTGGGGACACCAGGTCGGG, TCACCYTGGGGACACCAGGTCGG	Epidermolysis bullosa dystrophica inversa, autosomal recessive
199474715	NM_152263.3(TPM3):c.505A>G (p.Lys169Glu)	CCAACTYACGAGCCACCTACAGG	Congenital myopathy with fiber type disproportion
199474718	NM_152263.3(TPM3):c.733A>G (p.Arg245Gly)	ATCYCTCAGCAAACTCAGCACGG	Congenital myopathy with fiber type disproportion
121912895	NM_001844.4(COL2A1):c.2974A>G (p.Arg992Gly)	CCTCYCTCACCACGTTGCCCAGG	Spondyloepimetaphyseal dysplasia Strudwick type
121913074	NM_000129.3(F13A1):c.851A>G (p.Tyr284Cys)	ATAGGCAYAGATATTGTCCCAGG	Factor xiii, a subunit, deficiency of
121913145	NM_000208.2(INSR):c.707A>G (p.His236Arg)	GCTGYGGCAACAGAGGCCTTCGG	Leprechaunism syndrome
312262745	NM_025137.3(SPG11):c.2608A>G (p.Ile870Val)	ACTTAYCCTGGGGAGAAGGATGG	Spastic paraplegia 11, autosomal recessive
121913682	NM_000222.2(KIT):c.2459A>G (p.Asp820Gly)	AGAAYCATTCTTGATGTCTCTGG	Mast cell disease, systemic
587776757	NM_000151.3(G6PC):c.230+4A>G	GTTCYTACCACTTAAAGACGAGG	Glycogen storage disease type 1A
61752063	NM_000330.3(RS1):c.286T>C (p.Trp96Arg)	TTCTTCGYGGACTGCAAACAAGG	Juvenile retinoschisis
367543065	NM_024549.5(TCTN1):c.221-2A>G	AGCAACYGCAGAAAAAAGAGGGG, CAGCAACYGCAGAAAAAAGAGGG	Joubert syndrome 13
5030773	NM_000894.2(LHB):c.221A>G (p.Gln74Arg)	CCACCYGAGGCAGGGGCGGCAGG	Isolated lutropin deficiency
199476092	NM_000264.3(PTCH1):c.2479A>G (p.Ser827Gly)	CGTTACYGAAACTCCTGTGTAGG	Gorlin syndrome, Holoprosencephaly 7, not specified
398123158	NM_000117.2(EMD):c.450-2A>G	CGTTCCCYGAGGCAAAAGAGGGG	not provided
199476103	RMRP:n.71A>G	ACTTYCCCCTAGGCGGAAAGGGG, GACTTYCCCCTAGGCGGAAAGGG, GGACTTYCCCCTAGGCGGAAAGG	Metaphyseal chondrodysplasia, McKusick type, Metaphyseal dysplasia without hypotrichosis
5030856	NM_000277.1(PAH):c.1169A>G (p.Glu390Gly)	CTCYCTGCCACGTAATACAGGGG, ACTCYCTGCCACGTAATACAGGG, AACTCYCTGCCACGTAATACAGG	Phenylketonuria, Hyperphenylalaninemia, non-pku
5030860	NM_000277.1(PAH):c.1241A>G (p.Tyr414Cys)	GGGTCGYAGCGAACTGAGAAGGG, TGGGTCGYAGCGAACTGAGAAGG	Phenylketonuria, Hyperphenylalaninemia, non-pku
587777055	NM_020988.2(GNAO1):c.521A>G (p.Asp174Gly)	GGATGYCCTGCTCGGTGGGCTGG	Early infantile epileptic encephalopathy 17
587777223	NM_024301.4(FKRP):c.1A>G (p.Met1Val)	CCGCAYGGGGCCGAAGTCTGGGG, GCCGCAYGGGGCCGAAGTCTGGG, AGCCGCAYGGGGCCGAAGTCTGG	Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies type A5
587777479	NM_003108.3(SOX11):c.347A>G (p.Tyr116Cys)	GTACTTGYAGTCGGGGTAGTCGG	Mental retardation, autosomal dominant 27
587777496	NM_020435.3(GJC2):c.-170A>G	TTGYTCCCCCCTCGGCCTCAGGG, ATTGYTCCCCCCTCGGCCTCAGG	Leukodystrophy, hypomyelinating, 2
587777507	NM_022552.4(DNMT3A):c.1943T>C (p.Leu648Pro)	CTCCYGGTGCTGAAGGACTTGGG, GCTCCYGGTGCTGAAGGACTTGG	Tatton-Brown-rahman syndrome
587777557	NM_018400.3(SCN3B):c.482T>C (p.Met161Thr)	AATCAYGATGTACATCCTTCTGG	Atrial fibrillation, familial, 16
587777569	NM_001030001.2(RPS29):c.149T>C (p.Ile50Thr)	GATAYCGGTTTCATTAAGGTAGG	Diamond-Blackfan anemia 13
587777657	NM_153334.6(SCARF2):c.190T>C (p.Cys64Arg)	CCACGYGCTGCGCTGGCTGGAGG	Marden Walker like syndrome
587777689	NM_005726.5(TSFM):c.57+4A>G	ACTTCYCACCGGGTAGCTCCCGG	Combined oxidative phosphorylation deficiency 3
796052005	NM_000255.3(MUT):c.329A>G (p.Tyr110Cys)	GCAYACTGGCGGATGGTCCAGGG, AGCAYACTGGCGGATGGTCCAGG	not provided
587777809	NM_144596.3(TTC8):c.115-2A>G	GTTCCYGGAAAGCATTAAGAAGG	Retinitis pigmentosa 51
587777878	NM_000166.5(GJB1):c.580A>G (p.Met194Val)	TAGCAYGAAGACGGTGAAGACGG	X-linked hereditary motor and sensory neuropathy

74315420	NM_001029871.3(RSPO4):c.194A>G (p.Gln65Arg)	CGTACYGGCGGATGCCTTCCCGG	Anonychia
180177219	NM_000030.2(AGXT):c.424-2A>G (p.Gly_142Gln145del)	AGGCCCYGAGGAAGCAGGGACGG	Primary hyperoxaluria, type I

367610201	NM_002693.2(POLG):c.1808T>C (p.Met603Thr)	CTCAYGGCACTTACCTGGGATGG	not provided
180177319	NM_012203.1(GRHPR):c.84-2A>G	TCACAGCYGCGGGGAAAGGGAGG	Primary hyperoxaluria, type II
796052068	NM_000030.2(AGXT):c.777-2A>G	GGTACCYGGAAGACACGAGGGGG, TGGTACCYGGAAGACACGAGGGG	Primary hyperoxaluria, type I
61754010	NM_000552.3(VWF):c.1583A>G (p.Asn528Ser)	TGCCAYTGTAATTCCCACACAGG	von Willebrand disease, type 2a
587778866	NM_000321.2(RB1):c.1927A>G (p.Lys643Glu)	ATTYCAATGGCTTCTGGGTCTGG	Retinoblastoma
74435397	NM_006331.7(EMG1):c.257A>G (p.Asp86Gly)	ATAYCTGGCCGCGCTTCCCCAGG	Bowen-Conradi syndrome
796052527	NM_000156.5(GAMT):c.1A>G (p.Met1Val)	CGCTCAYGCTGCAGGCTGGACGG	not provided
796052637	NM_172107.2(KCNQ2):c.848A>G (p.Lys283Arg)	GTACYTGTCCCCGTAGCCAATGG	not provided
724159963	NM_032228.5(FAR1):c.1094A>G (p.Asp365Gly)	GATAYCATACAGGAATGCTGGGG, AGATAYCATACAGGAATGCTGGG, TAGATAYCATACAGGAATGCTGG	Peroxisomal fatty acyl-coa reductase 1 disorder
587779722	NM_000090.3(COL3A1):c.1762-2A>G (p.Gly588_Gln605del)	CACCCYAAAGAAGAAGTGGTCGG	Ehlers-Danlos syndrome, type 4
118192102	m.8296A>G	TTTACAGYGGGCTCTAGAGGGGG	Diabetes-deafness syndrome maternally transmitted
727502787	NM_001077494.3(NFKB2):c.2594A>G (p.Asp865Gly)	CTGYCTTCCTTCACCTCTGCTGG	Common variable immunodeficiency 10
727503036	NM_000117.2(EMD):c.266-2A>G	AGCCYTGGGAAGGGGGGCAGCGG	Emery-Dreifuss muscular dystrophy 1, X-linked
690016544	NM_005861.3(STUB1):c.194A>G (p.Asn65Ser)	GGCCCGGYTGGTGTAATACACGG	Spinocerebellar ataxia, autosomal recessive 16
690016554	NM_005211.3(CSF1R):c.2655-2A>G	GTATCYGGGAGATAGGACAGAGG	Hereditary diffuse leukoencephalopathy with spheroids
118192185	NM_172107.2(KCNQ2):c.1A>G (p.Met1Val)	GCACCAYGGTGCCTGGCGGGAGG	Benign familial neonatal seizures 1
121917869	NM_012064.3(MIP):c.401A>G (p.Glu134Gly)	AGATCYCCACTGTGGTTGCCTGG	Cataract 15, multiple types
121918014	NM_000478.4(ALPL):c.1250A>G (p.Asn417Ser)	AGGCCCAYTGCCATACAGGATGG	Infantile hypophosphatasia
121918036	NM_000174.4(GP9):c.110A>G (p.Asp37Gly)	GCAGYCCACCCACAGCCCCATGG	Bernard-Soulier syndrome type C
121918089	NM_000371.3(TTR):c.379A>G (p.Ile127Val)	CGGCAAYGGTGTAGCGGCGGGGG, GCGGCAAYGGTGTAGCGGCGGGG	Amyloidogenic transthyretin amyloidosis
121918121	NM_000823.3(GHRHR):c.985A>G (p.Lys329Glu)	CGACTYGGAGAGACGCCTGCAGG	Isolated growth hormone deficiency type 1B
121918333	NM_015335.4(MED13L):c.6068A>G (p.Asp2023Gly)	ATATCAYCTAGAGGGAAGGGGGG, CATATCAYCTAGAGGGAAGGGGG	Transposition of great arteries
121918605	NM_001035.2(RYR2):c.12602A>G (p.Gln4201Arg)	CGCCAGCYGCATTTCAAAGATGG	Catecholaminergic polymorphic ventricular tachycardia
587781262	NM_002764.3(PRPS1):c.343A>G (p.Met115Val)	TAGCAYATTTGCAACAAGCTTGG	Charcot-Marie-Tooth disease, X-linked recessive, type 5, Deafness, high-frequency sensorineural, X-linked
121918608	NM_001161766.1(AHCY):c.344A>G (p.Tyr115Cys)	GCGGGYACTTGGTGTGGATGAGG	Hypermethioninemia with s-adenosylhomocysteine hydrolase deficiency
121918613	NM_000702.3(ATP1A2):c.1033A>G (p.Thr345Ala)	CTGYCAGGGTCAGGCACACCTGG	Familial hemiplegic migraine type 2
587781339	NM_000535.5(PMS2):c.904-2A>G	GCAGACCYGCACAAAATACAAGG	Hereditary cancer-predisposing syndrome
121918691	NM_001128177.1(THRB):c.1324A>G (p.Met442Val)	CTTCAYGTGCAGGAAGCGGCTGG	Thyroid hormone resistance, generalized, autosomal dominant
121918692	NM_001128177.1(THRB):c.1327A>G (p.Lys443Glu)	CCACCTYCATGTGCAGGAAGCGG	Thyroid hormone resistance, generalized, autosomal dominant
727504333	NM_000256.3(MYBPC3):c.2906-2A>G	CCGTTCYGTGGGTATAGAGTGGG, GCCGTTCYGTGGGTATAGAGTGG	Familial hypertrophic cardiomyopathy 4
730880805	NM_006204.3(PDE6C):c.1483-2A>G	CTTTCYGTTGAAATAAGGATGGG, TCTTTCYGTTGAAATAAGGATGG	Achromatopsia 5

281860296	NM_000551.3(VHL):c.586A>T (p.Lys196Ter)	GGTCTTYCTGCACATTTGGGTGG	Von Hippel-Lindau syndrome
730880444	NM_000169.2(GLA):c.370-2A>G	GTGAACCYGAAATGAGAGGGAGG	not provided
756328339	NM_000256.3(MYBPC3):c.1227-2A>G	GTACCYGGGTGGGGGCCGCAGGG, TGTACCYGGGTGGGGGCCGCAGG	Familial hypertrophic cardiomyopathy 4, Cardiomyopathy
267606643	NM_013411.4(AK2):c.494A>G (p.Asp165Gly)	TCAYCTTTCATGGGCTCTTTTGG	Reticular dysgenesis
267606705	NM_005188.3(CBL):c.1144A>G (p.Lys382Glu)	TATTTYACATAGTTGGAATGTGG	Noonan syndrome-like disorder with or without juvenile myelomonocytic leukemia
62642934	NM_000277.1(PAH):c.916A>G (p.Ile306Val)	GGCCAAYTTCCTGTAATTGGGGG, AGGCCAAYTTCCTGTAATTGGGG	Phenylketonuria, Hyperphenylalaninemia, non-pku
267606782	NM_000117.2(EMD):c.1A>G (p.Met1Val)	TCCAYGGCGGGTGCGGGCTCAGG	Emery-Dreifuss muscular dystrophy, X-linked
267606820	NM_014053.3(FLVCR1):c.361A>G (p.Asn121Asp)	AGGCGTYGACCAGCGAGTACAGG	Posterior column ataxia with retinitis pigmentosa

[00318] In some embodiments, any of the base editors provided herein may be used to treat a disease or disorder. For example, any base editors provided herein may be used to correct one or more mutations associated with any of the diseases or disorders provided herein. Exemplary diseases or disorders that may be treated include, without limitation, 3-Methylglutaconic aciduria type 2, 46,XY gonadal dysgenesis, 4-Alpha-hydroxyphenylpyruvate hydroxylase deficiency, 6-pyruvoyl-tetrahydropterin synthase deficiency, achromatopsia, Acid-labile subunit deficiency, Acrodysostosis, acroerythrokeratoderma, ACTH resistance, ACTH-independent macronodular adrenal hyperplasia, Activated PI3K-delta syndrome, Acute intermittent porphyria, Acute myeloid leukemia, Adams-Oliver syndrome 1/5/6, Adenylosuccinate lyase deficiency, Adrenoleukodystrophy, Adult neuronal ceroid lipofuscinosis, Adult onset ataxia with oculomotor apraxia, Advanced sleep phase syndrome, Age-related macular degeneration, Alagille syndrome, Alexander disease, Allan-Herndon-Dudley syndrome, Alport syndrome, X-linked recessive, Alternating hemiplegia of childhood, Alveolar capillary dysplasia with misalignment of pulmonary veins, Amelogenesis imperfecta, Amyloidogenic transthyretin amyloidosis, Amyotrophic lateral sclerosis, Anemia (nonspherocytic hemolytic, due to G6PD deficiency), Anemia (sideroblastic, pyridoxine-refractory, autosomal recessive), Anonychia, Antithrombin III deficiency, Aortic aneurysm, Aplastic anemia, Apolipoprotein C2 deficiency, Apparent mineralocorticoid excess, Aromatase deficiency, Arrhythmogenic right ventricular cardiomyopathy, Familial hypertrophic cardiomyopathy, Hypertrophic cardiomyopathy, Arthrogryposis multiplex congenital, Aspartylglycosaminuria, Asphyxiating thoracic dystrophy, Ataxia with vitamin E deficiency, Ataxia (spastic), Atrial fibrillation, Atrial septal defect, atypical hemolytic-uremic syndrome, autosomal dominant CD11C+/CD1C+ dendritic cell deficiency, Autosomal dominant progressive external ophthalmoplegia with mitochondrial DNA deletions, Baraitser-Winter syndrome, Bartter syndrome, Basa ganglia calcification, Beckwith-Wiedemann syndrome, Benign familial neonatal seizures, Benign scapuloperoneal muscular dystrophy, Bernard Soulier syndrome, Beta thalassemia intermedia, Beta-D-mannosidosis, Bietti crystalline corneoretinal dystrophy, Bile acid malabsorption, Biotinidase deficiency, Borjeson-Forssman-Lehmann syndrome, Boucher Neuhauser syndrome, Bowen-Conradi syndrome, Brachydactyly, Brown-Vialetto-Van laere syndrome, Brugada syndrome, Cardiac arrhythmia, Cardiofaciocutaneous syndrome, Cardiomyopathy, Carnevale syndrome, Carnitine palmitoyltransferase II deficiency, Carpenter syndrome, Cataract, Catecholaminergic polymorphic

ventricular tachycardia, Central core disease, Centromeric instability of chromosomes 1,9 and 16 and immunodeficiency, Cerebral autosomal dominant arteriopathy, Cerebro-oculo-facio-skeletal syndrome, Ceroid lipofuscinosis, Charcot-Marie-Tooth disease, Cholestanol storage disease, Chondrocalcinosis, Chondrodysplasia, Chronic progressive multiple sclerosis, Coenzyme Q10 deficiency, Cohen syndrome, Combined deficiency of factor V and factor VIII, Combined immunodeficiency, Combined oxidative phosphorylation deficiency, Combined partial 17-alpha-hydroxylase/17,20- lyase deficiency, Complement factor d deficiency, Complete combined 17-alpha- hydroxylase/17,20-lyase deficiency, Cone-rod dystrophy, Congenital contractural arachnodactyly, Congenital disorder of glycosylation, Congenital lipomatous overgrowth, Neoplasm of ovary, PIK3CA Related Overgrowth Spectrum, Congenital long QT syndrome, Congenital muscular dystrophy, Congenital muscular hypertrophy-cerebral syndrome, Congenital myasthenic syndrome, Congenital myopathy with fiber type disproportion, Eichsfeld type congenital muscular dystrophy, Congenital stationary night blindness, Corneal dystrophy, Cornelia de Lange syndrome, Craniometaphyseal dysplasia, Crigler Najjar syndrome, Crouzon syndrome, Cutis laxa with osteodystrophy, Cyanosis, Cystic fibrosis, Cystinosis, Cytochrome-c oxidase deficiency, Mitochondrial complex I deficiency, D-2-hydroxyglutaric aciduria, Danon disease, Deafness with labyrinthine aplasia microtia and microdontia (LAMM), Deafness, Deficiency of acetyl-CoA acetyltransferase, Deficiency of ferroxidase, Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase, Dejerine-Sottas disease, Desbuquois syndrome, DFNA, Diabetes mellitus type 2, Diabetes-deafness syndrome, Diamond-Blackfan anemia, Diastrophic dysplasia, Dihydropteridine reductase deficiency, Dihydropyrimidinase deficiency, Dilated cardiomyopathy, Disseminated atypical mycobacterial infection, Distal arthrogryposis, Distal hereditary motor neuronopathy, Donnai Barrow syndrome, Duchenne muscular dystrophy, Becker muscular dystrophy, Dyschromatosis universalis hereditaria, Dyskeratosis congenital, Dystonia, Early infantile epileptic encephalopathy, Ehlers-Danlos syndrome, Eichsfeld type congenital muscular dystrophy, Emery-Dreifuss muscular dystrophy, Enamel-renal syndrome, Epidermolysis bullosa dystrophica inversa, Epidermolysis bullosa herpetiformis, Epilepsy, Episodic ataxia, Erythrokeratodermia variabilis, Erythropoietic protoporphyria, Exercise intolerance, Exudative vitreoretinopathy, Fabry disease, Factor V deficiency, Factor VII deficiency, Factor xiii deficiency, Familial adenomatous polyposis, breast cancer, ovarian cancer, cold urticarial, chronic infantile neurological, cutaneous and articular syndrome, hemiplegic migraine, hypercholesterolemia, hypertrophic cardiomyopathy,

hypoalphalipoproteinemia, hypokalemia-hypomagnesemia, juvenile gout, hyperlipoproteinemia, visceral amyloidosis, hypophosphatemic vitamin D refractory rickets, FG syndrome, Fibrosis of extraocular muscles, Finnish congenital nephrotic syndrome, focal epilepsy, Focal segmental glomerulosclerosis, Frontonasal dysplasia, Frontotemporal dementia, Fructose-biphosphatase deficiency, Gamstorp-Wohlfart syndrome, Ganglioside sialidase deficiency, GATA-1-related thrombocytopenia, Gaucher disease, Giant axonal neuropathy, Glanzmann thrombasthenia, Glomerulocystic kidney disease, Glomerulopathy, Glucocorticoid resistance, Glucose-6-phosphate transport defect, Glutaric aciduria, Glycogen storage disease, Gorlin syndrome, Holoprosencephaly, GRACILE syndrome, Haemorrhagic telangiectasia, Hemochromatosis, Hemoglobin H disease, Hemolytic anemia, Hemophagocytic lymphohistiocytosis, Carcinoma of colon, Myhre syndrome, leukoencephalopathy, Hereditary factor IX deficiency disease, Hereditary factor VIII deficiency disease, Hereditary factor XI deficiency disease, Hereditary fructosuria, Hereditary Nonpolyposis Colorectal Neoplasm, Hereditary pancreatitis, Hereditary pyropoikilocytosis, Elliptocytosis, Heterotaxy, Heterotopia, Histiocytic medullary reticulosis, Histiocytosis-lymphadenopathy plus syndrome, HNSHA due to aldolase A deficiency, Holocarboxylase synthetase deficiency, Homocysteinemia, Howel-Evans syndrome, Hydatidiform mole, Hypercalciuric hypercalcemia, Hyperimmunoglobulin D, Mevalonic aciduria, Hyperinsulinemic hypoglycemia, Hyperkalemic Periodic Paralysis, Paramyotonia congenita of von Eulenburg, Hyperlipoproteinemia, Hypermanganesemia, Hypermethioninemia, Hyperphosphatasemia, Hypertension, hypomagnesemia, Hypobetalipoproteinemia, Hypocalcemia, Hypogonadotropic hypogonadism, Hypogonadotropic hypogonadism, Hypohidrotic ectodermal dysplasia, Hyper-IgM immunodeficiency, Hypohidrotic X-linked ectodermal dysplasia, Hypomagnesemia, Hypoparathyroidism, Idiopathic fibrosing alveolitis, Immunodeficiency, Immunoglobulin A deficiency, Infantile hypophosphatasia, Infantile Parkinsonism-dystonia, Insulin-dependent diabetes mellitus, Intermediate maple syrup urine disease, Ischiopatellar dysplasia, Islet cell hyperplasia, Isolated growth hormone deficiency, Isolated lutropin deficiency, Isovaleric acidemia, Joubert syndrome, Juvenile polyposis syndrome, Juvenile retinoschisis, Kallmann syndrome, Kartagener syndrome, Kugelberg-Welander disease, Lattice corneal dystrophy, Leber congenital amaurosis, Leber optic atrophy, Left ventricular noncompaction, Leigh disease, Mitochondrial complex I deficiency, Leprechaunism syndrome, Arthrogryposis, Anterior horn cell disease, Leukocyte adhesion deficiency, Leukodystrophy, Leukoencephalopathy, Ovarioleukodystrophy, L-ferritin deficiency, Li-Fraumeni syndrome,

Limb-girdle muscular dystrophy- dystroglycanopathy, Loeys-Dietz syndrome, Long QT syndrome, Macrocephaly/autism syndrome, Macular corneal dystrophy, Macular dystrophy, Malignant hyperthermia susceptibility, Malignant tumor of prostate, Maple syrup urine disease, Marden Walker like syndrome, Marfan syndrome, Marie Unna hereditary hypotrichosis, Mast cell disease, Meconium ileus, Medium-chain acyl-coenzyme A dehydrogenase deficiency, Melnick-Fraser syndrome, Mental retardation, Merosin deficient congenital muscular dystrophy, Mesothelioma, Metachromatic leukodystrophy, Metaphyseal chondrodysplasia, Methemoglobinemia, methylmalonic aciduria, homocystinuria, Microcephaly, chorioretinopathy, lymphedema, Microphthalmia, Mild non-PKU hyperphenylalanemia, Mitchell-Riley syndrome, mitochondrial 3-hydroxy-3-methylglutaryl-CoA synthase deficiency, Mitochondrial complex I deficiency, Mitochondrial complex III deficiency, Mitochondrial myopathy, Mucolipidosis III, Mucopolysaccharidosis, Multiple sulfatase deficiency, Myasthenic syndrome, Mycobacterium tuberculosis, Myeloperoxidase deficiency, Myhre syndrome, Myoclonic epilepsy, Myofibrillar myopathy, Myoglobinuria, Myopathy, Myopia, Myotonia congenital, Navajo neurohepatopathy, Nemaline myopathy, Neoplasm of stomach, Nephrogenic diabetes insipidus, Nephronophthisis, Nephrotic syndrome, Neurofibromatosis, Neutral lipid storage disease, Niemann-Pick disease, Non-ketotic hyperglycinemia, Noonan syndrome, Noonan syndrome-like disorder, Norum disease, Macular degeneration, N-terminal acetyltransferase deficiency, Oculocutaneous albinism, Oculodentodigital dysplasia, Ohdo syndrome, Optic nerve aplasia, Ornithine carbamoyltransferase deficiency, Orofaciodigital syndrome, Osteogenesis imperfecta, Osteopetrosis, Ovarian dysgenesis, Pachyonychia, Palmoplantar keratoderma, nonepidermolytic, Papillon-Lef\xc3\xa8vre syndrome, Haim-Munk syndrome, Periodontitis, Peeling skin syndrome, Pendred syndrome, Peroxisomal fatty acyl-coa reductase 1 disorder, Peroxisome biogenesis disorder, Pfeiffer syndrome, Phenylketonuria, Phenylketonuria, Hyperphenylalaninemia, non-PKU, Pituitary hormone deficiency, Pityriasis rubra pilaris, Polyarteritis nodosa, Polycystic kidney disease, Polycystic lipomembranous osteodysplasia, Polymicrogyria, Pontocerebellar hypoplasia, Porokeratosis, Posterior column ataxia, Primary erythromelalgia, hyperoxaluria, Progressive familial intrahepatic cholestasis, Progressive pseudorheumatoid dysplasia, Propionic acidemia, Pseudohermaphroditism, Pseudohypoaldosteronism, Pseudoxanthoma elasticum-like disorder, Purine-nucleoside phosphorylase deficiency, Pyridoxal 5-phosphate-dependent epilepsy, Renal dysplasia, retinal pigmentary dystrophy, cerebellar ataxia, skeletal dysplasia, Reticular dysgenesis, Retinitis pigmentosa, Usher syndrome, Retinoblastoma, Retinopathy, RRM2B-related

mitochondrial disease, Rubinstein-Taybi syndrome, Schnyder crystalline corneal dystrophy, Sebaceous tumor, Severe congenital neutropenia, Severe myoclonic epilepsy in infancy, Severe X-linked myotubular myopathy, onychodysplasia, facial dysmorphism, hypotrichosis, Short-rib thoracic dysplasia, Sialic acid storage disease, Sialidosis, Sideroblastic anemia, Small fiber neuropathy, Smith-Magenis syndrome, Sorsby fundus dystrophy, Spastic ataxia, Spastic paraplegia, Spermatogenic failure, Spherocytosis, Sphingomyelin/cholesterol lipidosis, Spinocerebellar ataxia, Split-hand/foot malformation, Spondyloepimetaphyseal dysplasia, Platyspondylic lethal skeletal dysplasia, Squamous cell carcinoma of the head and neck, Stargardt disease, Sucrase-isomaltase deficiency, Sudden infant death syndrome, Supravalvar aortic stenosis, Surfactant metabolism dysfunction, Tangier disease, Tatton-Brown-rahman syndrome, Thoracic aortic aneurysms and aortic dissections, Thrombophilia, Thyroid hormone resistance, TNF receptor-associated periodic fever syndrome (TRAPS), Tooth agenesis, Torsades de pointes, Transposition of great arteries, Treacher Collins syndrome, Tuberous sclerosis syndrome, Tyrosinase-negative oculocutaneous albinism, Tyrosinase-positive oculocutaneous albinism, Tyrosinemia, UDPglucose-4-epimerase deficiency, Ullrich congenital muscular dystrophy, Bethlem myopathy Usher syndrome, UV-sensitive syndrome, Van der Woude syndrome, popliteal pterygium syndrome, Very long chain acyl-CoA dehydrogenase deficiency, Vesicoureteral reflux, Vitreoretinochoroidopathy, Von Hippel-Lindau syndrome, von Willebrand disease, Waardenburg syndrome, Warsaw breakage syndrome, WFS1-Related Disorders, Wilson disease, Xeroderma pigmentosum, X-linked agammaglobulinemia, X-linked hereditary motor and sensory neuropathy, X-linked severe combined immunodeficiency, and Zellweger syndrome.

[00319] The development of nucleobase editing advances both the scope and effectiveness of genome editing. The nucleobase editors described here offer researchers a choice of editing with virtually no indel formation (NBE2), or more efficient editing with a low frequency (here, typically $\leq$ 1%) of indel formation (NBE3). That the product of base editing is, by definition, no longer a substrate likely contributes to editing efficiency by preventing subsequent product transformation, which can hamper traditional Cas9 applications. By removing the reliance on double-stranded DNA cleavage and stochastic DNA repair processes that vary greatly by cell state and cell type, nucleobase editing has the potential to expand the type of genome modifications that can be cleanly installed, the efficiency of these modifications, and the type of cells that are amenable to editing. It is likely that recent engineered Cas9 variants[69,70,82] or delivery methods[71]

with improved DNA specificity, as well as Cas9 variants with altered PAM specificities,[72] can be integrated into this strategy to provide additional nucleobase editors with improved DNA specificity or that can target an even wider range of disease-associated mutations. These findings also suggest that engineering additional fusions of dCas9 with enzymes that catalyze additional nucleobase transformations will increase the fraction of the possible DNA base changes that can be made through nucleobase editing. These results also suggest architectures for the fusion of other DNA-modifying enzymes, including methylases and demathylases, that mau enable additional types of programmable genome and epigenome base editing.

Materials and Methods

[00320] Cloning. DNA sequences of all constructs and primers used in this paper are listed in the Supplementary Sequences. Plasmids containing genes encoding NBE1, NBE2, and NBE3 will be available from Addgene. PCR was performed using VeraSeq ULtra DNA polymerase (Enzymatics), or Q5 Hot Start High-Fidelity DNA Polymerase (New England Biolabs). NBE plasmids were constructed using USER cloning (New England Biolabs). Deaminase genes were synthesized as gBlocks Gene Fragments (Integrated DNA Technologies), and Cas9 genes were obtained from previously reported plasmids.[18] Deaminase and fusion genes were cloned into pCMV (mammalian codon-optimized) or pET28b (*E. coli* codon-optimized) backbones. sgRNA expression plasmids were constructed using site-directed mutagenesis. Briefly, the primers listed in the Supplementary Sequences were 5' phosphorylated using T4 Polynucleotide Kinase (New England Biolabs) according to the manufacturer's instructions. Next, PCR was performed using Q5 Hot Start High-Fidelity Polymerase (New England Biolabs) with the phosphorylated primers and the plasmid pFYF1320 (EGFP sgRNA expression plasmid) as a template according to the manufacturer's instructions. PCR products were incubated with DpnI (20 U, New England Biolabs) at 37 °C for 1 h, purified on a QIAprep spin column (Qiagen), and ligated using QuickLigase (New England Biolabs) according to the manufacturer's instructions. DNA vector amplification was carried out using Mach1 competent cells (ThermoFisher Scientific).

[00321] *In vitro* deaminase assay on ssDNA. Sequences of all ssDNA substrates are listed in the Supplementary Sequences. All Cy3-labelled substrates were obtained from Integrated DNA Technologies (IDT). Deaminases were expressed *in vitro* using the TNT T7 Quick Coupled Transcription/Translation Kit (Promega) according to the manufacturer's instructions using 1 μg of plasmid. Following protein expression, 5 μL of lysate was combined with 35 μL of ssDNA (1.8

μM) and USER enzyme (1 unit) in CutSmart buffer (New England Biolabs) (50 mM potassium acetate, 29 mM Trisacetate, 10 mM magnesium acetate, 100 ug/mL BSA, pH 7.9) and incubated at 37 °C for 2 h. Cleaved U-containing substrates were resolved from full-length unmodified substrates on a 10% TBE-urea gel (Bio-Rad).

[00322] Expression and purification of His_6-rAPOBEC1-linker-dCas9 fusions. *E. Coli* BL21 STAR (DE3)-competent cells (ThermoFisher Scientific) were transformed with plasmids encoding pET28b-His_6-rAPOBEC-linker-dCas9 with GGS, $(GGS)_3$, (SEQ ID NO: 596) XTEN, or $(GGS)_7$ (SEQ ID NO: 597) linkers. The resulting expression strains were grown overnight in Luria-Bertani (LB) broth containing 100 μg/mL of kanamycin at 37 °C. The cells were diluted 1:100 into the same growth medium and grown at 37 °C to OD_{600} = ~0.6. The culture was cooled to 4 °C over a period of 2 h, and isopropyl -β-D-1- thiogalactopyranoside (IPTG) was added at 0.5 mM to induce protein expression. After ~16 h, the cells were collected by centrifugation at 4,000 *g* and resuspended in lysis buffer (50 mM tris(hydroxymethyl)-aminomethane (Tris)-HCl, pH 7.0, 1 M NaCl, 20% glycerol, 10 mM tris(2-carboxyethyl)phosphine (TCEP, Soltec Ventures)). The cells were lysed by sonication (20 s pulse-on, 20 s pulse-off for 8 min total at 6 W output) and the lysate supernatant was isolated following centrifugation at 25,000 *g* for 15 min. The lysate was incubated with His-Pur nickel-nitriloacetic acid (nickel-NTA) resin (ThermoFisher Scientific) at 4 °C for 1 h to capture the His-tagged fusion protein. The resin was transferred to a column and washed with 40 mL of lysis buffer. The His-tagged fusion protein was eluted in lysis buffer supplemented with 285 mM imidazole, and concentrated by ultrafiltration (Amicon-Millipore, 100-kDa molecular weight cut-off) to 1 mL total volume. The protein was diluted to 20 mL in low-salt purification buffer containing 50 mM tris(hydroxymethyl)-aminomethane (Tris)-HCl, pH 7.0, 0.1 M NaCl, 20% glycerol, 10 mM TCEP and loaded onto SP Sepharose Fast Flow resin (GE Life Sciences). The resin was washed with 40 mL of this low-salt buffer, and the protein eluted with 5 mL of activity buffer containing 50 mM tris(hydroxymethyl)-aminomethane (Tris)-HCl, pH 7.0, 0.5 M NaCl, 20% glycerol, 10 mM TCEP. The eluted proteins were quantified on a SDSPAGE gel.

[00323] *In vitro* transcription of sgRNAs. Linear DNA fragments containing the T7 promoter followed by the 20-bp sgRNA target sequence were transcribed *in vitro* using the primers listed in the Supplementary Sequences with the TranscriptAid T7 High Yield Transcription Kit (ThermoFisher Scientific) according to the manufacturer's instructions. sgRNA products were

purified using the MEGAclear Kit (ThermoFisher Scientific) according to the manufacturer's instructions and quantified by UV absorbance.

[00324] Preparation of Cy3-conjugated dsDNA substrates. Sequences of 80-nucleotide unlabeled strands are listed in the Supplementary Sequences and were ordered as PAGE-purified oligonucleotides from IDT. The 25-nt Cy3-labeled primer listed in the Supplementary Sequences is complementary to the 3' end of each 80-nt substrate. This primer was ordered as an HPLC-purified oligonucleotide from IDT. To generate the Cy3-labeled dsDNA substrates, the 80-nt strands (5 μL of a 100 μM solution) were combined with the Cy3-labeled primer (5 μL of a 100 μM solution) in NEBuffer 2 (38.25 μL of a 50 mM NaCl, 10 mMTris-HCl, 10 mM $MgCl_2$, 1 mM DTT, pH 7.9 solution, New England Biolabs) with dNTPs (0.75 μL of a 100 mM solution) and heated to 95 °C for 5 min, followed by a gradual cooling to 45 °C at a rate of 0.1 °C/s. After this annealing period, Klenow exo^- (5 U, New England Biolabs) was added and the reaction was incubated at 37 °C for 1 h. The solution was diluted with Buffer PB (250 μL, Qiagen) and isopropanol (50 μL) and purified on a QIAprep spin column (Qiagen), eluting with 50 μL of Tris buffer.

[00325] Deaminase assay on dsDNA. The purified fusion protein (20 μL of 1.9 μM in activity buffer) was combined with 1 equivalent of appropriate sgRNA and incubated at ambient temperature for 5 min. The Cy3-labeled dsDNA substrate was added to final concentration of 125 nM and the resulting solution was incubated at 37 °C for 2 h. The dsDNA was separated from the fusion by the addition of Buffer PB (100 μL, Qiagen) and isopropanol (25 μL) and purified on a EconoSpin micro spin column (Epoch Life Science), eluting with 20 μL of CutSmart buffer (New England Biolabs). USER enzyme (1 U, New England Biolabs) was added to the purified, edited dsDNA and incubated at 37 °C for 1 h. The Cy3-labeled strand was fully denatured from its complement by combining 5 μL of the reaction solution with 15 μL of a DMSO-based loading buffer (5 mM Tris, 0.5 mM EDTA, 12.5% glycerol, 0.02% bromophenol blue, 0.02% xylene cyan, 80% DMSO). The full-length C-containing substrate was separated from any cleaved, U-containing edited substrates on a 10% TBE-urea gel (Bio-Rad) and imaged on a GE Amersham Typhoon imager.

[00326] Preparation of *in vitro*-edited dsDNA for high-throughput sequencing (HTS). The oligonucleotides listed in the Supplementary Sequences were obtained from IDT. Complementary sequences were combined (5 μL of a 100 μM solution) in Tris buffer and annealed by heating to 95 °C for 5 min, followed by a gradual cooling to 45 °C at a rate of 0.1 °C/s to generate 60-bp

dsDNA substrates. Purified fusion protein (20 μL of 1.9 μM in activity buffer) was combined with 1 equivalent of appropriate sgRNA and incubated at ambient temperature for 5 min. The 60-mer dsDNA substrate was added to final concentration of 125 nM and the resulting solution was incubated at 37 °C for 2 h. The dsDNA was separated from the fusion by the addition of Buffer PB (100 μL, Qiagen) and isopropanol (25 μL) and purified on a EconoSpin micro spin column (Epoch Life Science), eluting with 20 μL of Tris buffer. The resulting edited DNA (1 μL was used as a template) was amplified by PCR using the HTS primer pairs specified in the Supplementary Sequences and VeraSeq Ultra (Enzymatics) according to the manufacturer's instructions with 13 cycles of amplification. PCR reaction products were purified using RapidTips (Diffinity Genomics), and the purified DNA was amplified by PCR with primers containing sequencing adapters, purified, and sequenced on a MiSeq high-throughput DNA sequencer (Illumina) as previously described.[73]

[00327] Cell culture. HEK293T (ATCC CRL-3216), U2OS (ATCC-HTB-96) and ST486 cells (ATCC) were maintained in Dulbecco's Modified Eagle's Medium plus GlutaMax (ThermoFisher) supplemented with 10% (v/v) fetal bovine serum (FBS) and penicillin/streptomycin (1x, Amresco), at 37 °C with 5% CO_2. HCC1954 cells (ATCC CRL-2338) were maintained in RPMI-1640 medium (ThermoFisher Scientific) supplemented as described above. Immortalized rat astrocytes containing the ApoE4 isoform of the *APOE* gene (Taconic Biosciences) were cultured in Dulbecco's Modified Eagle's Medium plus GlutaMax (ThermoFisher Scientific) supplemented with 10% (v/v) fetal bovine serum (FBS) and 200 μg/mL Geneticin (ThermoFisher Scientific).

[00328] Transfections. HEK293T cells were seeded on 48-well collagen-coated BioCoat plates (Corning) and transfected at approximately 85% confluency. Briefly, 750 ng of NBE and 250 ng of sgRNA expression plasmids were transfected using 1.5 μl of Lipofectamine 2000 (ThermoFisher Scientific) per well according to the manufacturer's protocol. Astrocytes, U2OS, HCC1954, HEK293T and ST486 cells were transfected using appropriate AMAXA NUCLEOFECTOR™ II programs according to manufacturer's instructions. 40 ng of infrared RFP (Addgene plasmid 45457)[74] was added to the nucleofection solution to assess nucleofection efficiencies in these cell lines. For astrocytes, U2OS, and ST486 cells, nucleofection efficiencies were 25%, 74%, and 92%, respectively. For HCC1954 cells, nucleofection efficiency was <10%. Therefore, following trypsinization, the HCC1954 cells were filtered through a 40 micron strainer (Fisher Scientific), and the nucleofected HCC1954 cells were collected on a Beckman Coulter

MoFlo XDP Cell Sorter using the iRFP signal (abs 643 nm, em 670 nm). The other cells were used without enrichment of nucleofected cells.

[00329] High-throughput DNA sequencing of genomic DNA samples. Transfected cells were harvested after 3 d and the genomic DNA was isolated using the Agencourt DNAdvance Genomic DNA Isolation Kit (Beckman Coulter) according to the manufacturer's instructions. On-target and off-target genomic regions of interest were amplified by PCR with flanking HTS primer pairs listed in the Supplementary Sequences. PCR amplification was carried out with Phusion high-fidelity DNA polymerase (ThermoFisher) according to the manufacturer's instructions using 5 ng of genomic DNA as a template. Cycle numbers were determined separately for each primer pair as to ensure the reaction was stopped in the linear range of amplification (30, 28, 28, 28, 32, and 32 cycles for EMX1, FANCF, HEK293 site 2, HEK293 site 3, HEK293 site 4, and RNF2 primers, respectively). PCR products were purified using RapidTips (Diffinity Genomics). Purified DNA was amplified by PCR with primers containing sequencing adaptors. The products were gel-purified and quantified using the QUANT-IT™ PicoGreen dsDNA Assay Kit (ThermoFisher) and KAPA Library Quantification Kit-Illumina (KAPA Biosystems). Samples were sequenced on an Illumina MiSeq as previously described.[73]

[00330] Data analysis. Sequencing reads were automatically demultiplexed using MiSeq Reporter (Illumina), and individual FASTQ files were analyzed with a custom Matlab script provided in the Supplementary Notes. Each read was pairwise aligned to the appropriate reference sequence using the Smith-Waterman algorithm. Base calls with a Q-score below 31 were replaced with N's and were thus excluded in calculating nucleotide frequencies. This treatment yields an expected MiSeq base-calling error rate of approximately 1 in 1,000. Aligned sequences in which the read and reference sequence contained no gaps were stored in an alignment table from which base frequencies could be tabulated for each locus.

[00331] Indel frequencies were quantified with a custom Matlab script shown in the Supplementary Notes using previously described criteria[71]. Sequencing reads were scanned for exact matches to two 10-bp sequences that flank both sides of a window in which indels might occur. If no exact matches were located, the read was excluded from analysis. If the length of this indel window exactly matched the reference sequence the read was classified as not containing an indel. If the indel window was two or more bases longer or shorter than the reference sequence, then the sequencing read was classified as an insertion or deletion, respectively.

[00332] All publications, patents, patent applications, publication, and database entries (*e.g.*, sequence database entries) mentioned herein, *e.g.*, in the Background, Summary, Detailed Description, Examples, and/or References sections, are hereby incorporated by reference in their entirety as if each individual publication, patent, patent application, publication, and database entry was specifically and individually incorporated herein by reference. In case of conflict, the present application, including any definitions herein, will control.

Supplementary Sequences

[00333] Primers used for generating sgRNA transfection plasmids. rev_sgRNA_plasmid was used in all cases. The pFYF1320 plasmid was used as template as noted in Materials and Methods section. SEQ ID NOs: 329-338 appear from top to bottom below, respectively.

rev_sgRNA_plasmid	GGTGTTTCGTCCTTTCCACAAG
fwd _p53 _Y163C	GCTTGCAGATGGCCATGGCGGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_p53_N239D	TGTCACACATGTAGTTGTAGGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_APOE4 _C158R	GAAGCGCCTGGCAGTGTACCGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_EMX1	GAGTCCGAGCAGAAGAAGAAGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_FANCF	GGAATCCCTTCTGCAGCACCGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_HEK293_2	GAACACAAAGCATAGACTGCGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_HEK293_3	GGCCCAGACTGAGCACGTGAGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_HEK293_4	GGCACTGCGGCTGGAGGTGGGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC
fwd_RNF2	GTCATCTTAGTCATTACCTGGTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGC

[00334] Sequences of all ssDNA substrates used in in vitro deaminase assays. SEQ ID NOs: 339-341 appear from top to bottom below, respectively.

rAPOBEC1 substrate	Cy3-ATTATTATTATTCCGCGGATTTATTTATTTATTTATTTATTT
hAID/pmCDA1 substrate	Cy3-ATTATTATTATTAGCTATTTATTTATTTATTTATTTATTT
hAPOBEC3G substrate	Cy3-ATTATTATTATTCCCGGATTTATTTATTTATTTATTTATTT

[00335] Primers used for generating PCR products to serve as substrates for T7 transcription of sgRNAs for gel-based deaminase assay. rev_gRNA_T7 was used in all cases.

The pFYF1320 plasmid was used as template as noted in Materials and Methods section. SEQ ID NOs: 342-365 appear from top to bottom below, respectively.

rev_sgRNA_T7	AAAAAAAGCACCGACTCGGTG
fwd_sgRNA_T7_dsDNA_2	TAATACGACTCACTATAGGCCGCGGATTTATTTATTTAAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_3	TAATACGACTCACTATAGGTCCGCGGATTTATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_4	TAATACGACTCACTATAGGTTCCGCGGATTTATTTATTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_5	TAATACGACTCACTATAGGATTCCGCGGATTTATTTATTGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_6	TAATACGACTCACTATAGGTATTCCGCGGATTTATTTATGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_7	TAATACGACTCACTATAGGTTATTCCGCGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_8	TAATACGACTCACTATAGGATTATTCCGCGGATTTATTTGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_9	TAATACGACTCACTATAGGTATTATTCCGCGGATTTATTGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_10	TAATACGACTCACTATAGGATTATTATCCGCGGATTTATGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_11	TAATACGACTCACTATAGGTATTATATTCCGCGGATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_12	TAATACGACTCACTATAGGTTATTATATTCCGCGGATTTGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_13	TAATACGACTCACTATAGGATTATTATATTCCGCGGATTGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_14	TAATACGACTCACTATAGGTATTATTATATTCCGCGGATGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_15	TAATACGACTCACTATAGGATTATTATTATTACCGCGGAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_18	TAATACGACTCACTATAGGATTATTATTATTATTACCGCGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_noC	TAATACGACTCACTATAGGATATTAATTTATTTATTTAAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_APOE4_C112R	TAATACGACTCACTATAGGGGAGGACGTGCGCGGCCGCCGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_APOE4_C158R	TAATACGACTCACTATAGGGAAGCGCCTGGCAGTGTACCGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_CTNNB1_T41A	TAATACGACTCACTATAGGCTGTGGCAGTGGCACCAGAAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_HRAS_Q61R	TAATACGACTCACTATAGGCCTCCCGGCCGGCGGTATCCGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_53_Y163C	TAATACGACTCACTATAGGGCTTGCAGATGGCCATGGCGGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_53_Y236C	TAATACGACTCACTATAGGACACATGCAGTTGTAGTGGAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_dsDNA_53_N239D	TAATACGACTCACTATAGGTGTCACACATGTAGTTGTAGGTTTTAGAGCTAGAAATAGCA

[00336] Sequences of 80-nucleotide unlabeled strands and Cy3-labeled universal primer used in gel-based dsDNA deaminase assays. SEQ ID NOs: 366-390 appear from top to bottom below, respectively.

Cy3-primer	Cy3-GTAGGTAGTTAGGATGAATGGAAGGTTGGTA
dsDNA_2	GTCCATGGATCCAGAGGTCATCCATTAAATAAATAAATCCGCGGGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_3	GTCCATGGATCCAGAGGTCATCCATAAATAAATAAATCCGCGGAAGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_4	GTCCATGGATCCAGAGGTCATCCATAATAAATAAATCCGCGGAAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_5	GTCCATGGATCCAGAGGTCATCCAAATAAATAAATCCGCGGAATGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_6	GTCCATGGATCCAGAGGTCATCCAATAAATAAATCCGCGGAATAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_7	GTCCATGGATCCAGAGGTCATCCATAAATAAATCCGCGGAATAAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_8	GTCCATGGATCCAGAGGTCATCCAAAATAAATCCGCGGAATAATGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_9	GTCCATGGATCCAGAGGTCATCCAAATAAATCCGCGGAATAATAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_10	GTCCATGGATCCAGAGGTCATCCAATAAATCCGCGGATAATAATGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_11	GTCCATGGATCCAGAGGTCATCCATAAATCCGCGGAATATAATAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_12	GTCCATGGATCCAGAGGTCATCCAAAATCCGCGGAATATAATAAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_13	GTCCATGGATCCAGAGGTCATCCAAATCCGCGGAATATAATAATGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_14	GTCCATGGATCCAGAGGTCATCCAATCCGCGGAATATAATAATAGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_15	GTCCATGGATCCAGAGGTCATCCATCCGCGGTAATAATAATAATGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_16	GTCCATGGATCCAGAGGTCATCCAGCGGTAATAATAATAATAATGGCTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_noC	GTCCATGGATCCAGAGGTCATCCATTAAATAAATAAATTAATATTACTATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_8U	5Cy3-GTAGGTAGTTAGGATGAATGGAAGGTTGGTGTAGATTATTATGUGCGGATTTATTGGATGACCTCTGGATCCATGGACAT
dsDNA_APOE_C112R	GCACCTCGCCGCGGTACTGCACCAGGCGGCCGCGCACGTCCTCCATGTCTACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_APOE_C158R	CGGCGCCCTCGCGGGCCCCGGCCTGGTACACTGCCAGGCGCTTCTGCAGTACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_CTNNB1_T41A	GTCTTACCTGGACTCTGGAATCCATTCTGGTGCCACTGCCACAGCTCCTTACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_HRAS_Q61R	GGAGACGTGCCTGTTGGACATCCTGGATACCGCCGGCCGGGAGGAGTACTACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_p53_Y163C	ACCCCCGCCCGGCACCCGCGTCCGCGCCATGGCCATCTGCAAGCAGTCATACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_p53_Y236C	AGGTTGGCTCTGACTGTACCACCATCCACTACAACTGCATGTGTAACAGTACCAACCTTCCATTCATCCTAACTACCTAC
dsDNA_p53_N239D	TGGCTCTGACTGTACCACCATCCACTACAACTACATGTGTGACAGTTCCTACCAACCTTCCATTCATCCTAACTACCTAC

[00337] Primers used for generating PCR products to serve as substrates for T7 transcription of sgRNAs for high-throughput sequencing. rev_gRNA_T7 (above) was used in all cases. The pFYF1320 plasmid was used as template as noted in Materials and Methods section. SEQ ID NOs: 391-442 appear from top to bottom below, respectively.

fwd_sgRNA_T7_HTS_ base	TAATACGACTCACTATAGGTTATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 1A	TAATACGACTCACTATAGGATATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 1C	TAATACGACTCACTATAGGCTATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 1G	TAATACGACTCACTATAGGGTATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 2A	TAATACGACTCACTATAGGTAATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 2C	TAATACGACTCACTATAGGTCATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 2G	TAATACGACTCACTATAGGTGATTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 3T	TAATACGACTCACTATAGGTTTTTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 3C	TAATACGACTCACTATAGGTTCTTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 3G	TAATACGACTCACTATAGGTTGTTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 4A	TAATACGACTCACTATAGGTTAATTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 4C	TAATACGACTCACTATAGGTTACTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 4G	TAATACGACTCACTATAGGTTAGTTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 5A	TAATACGACTCACTATAGGTTATATCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 5C	TAATACGACTCACTATAGGTTATCTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 5G	TAATACGACTCACTATAGGTTATGTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 6A	TAATACGACTCACTATAGGTTATTACGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 6C	TAATACGACTCACTATAGGTTATTCCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 6G	TAATACGACTCACTATAGGTTATTGCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 8A	TAATACGACTCACTATAGGTTATTTCATGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 8T	TAATACGACTCACTATAGGTTATTTCTTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 8C	TAATACGACTCACTATAGGTTATTTCCTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 9A	TAATACGACTCACTATAGGTTATTTCGAGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 9C	TAATACGACTCACTATAGGTTATTTCGCGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 9G	TAATACGACTCACTATAGGTTATTTCGGGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 10A	TAATACGACTCACTATAGGTTATTTCGTAGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 10T	TAATACGACTCACTATAGGTTATTTCGTTGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 10C	TAATACGACTCACTATAGGTTATTTCGTCGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 11A	TAATACGACTCACTATAGGTTATTTCGTGAATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 11T	TAATACGACTCACTATAGGTTATTTCGTGTATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 11C	TAATACGACTCACTATAGGTTATTTCGTGCATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 12T	TAATACGACTCACTATAGGTTATTTCGTGGTTTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 12C	TAATACGACTCACTATAGGTTATTTCGTGGCTTTATTTAGTTTTAGAGCTAGAAATAGCA

fwd_sgRNA_T7_HTS_ 12G	TAATACGACTCACTATAGGTTATTTCGTGGGTTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 13A	TAATACGACTCACTATAGGTTATTTCGTGGAATTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 13C	TAATACGACTCACTATAGGTTATTTCGTGGACTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ 13G	TAATACGACTCACTATAGGTTATTTCGTGGAGTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ multiC	TAATACGACTCACTATAGGTTCCCCCCCCGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ TCGCACCC_odd	TAATACGACTCACTATAGGCGCACCCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ CCTCGCAC_odd	TAATACGACTCACTATAGGCTCGCACGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ ACCCTCGC_odd	TAATACGACTCACTATAGGCCCTCGCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ GCACCCTC_odd	TAATACGACTCACTATAGGCACCCTCGTGGATTTATTTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ TCGCACCC_even	TAATACGACTCACTATAGGTCGCACCCGTGGATTTATTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ CCTCGCAC_even	TAATACGACTCACTATAGGCCTCGCACGTGGATTTATTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ ACCCTCGC_even	TAATACGACTCACTATAGGACCCTCGCGTGGATTTATTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ GCACCCTC_even	TAATACGACTCACTATAGGGCACCCTCGTGGATTTATTAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ EMX1	TAATACGACTCACTATAGGGAGTCCGAGCAGAAGAAGAAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ FANCF	TAATACGACTCACTATAGGGGAATCCCTTCTGCAGCACCGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ HEK293_site2	TAATACGACTCACTATAGGGAACACAAAGCATAGACTGCGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ HEK293_site3	TAATACGACTCACTATAGGGGCCCAGACTGAGCACGTGAGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ HEK293_site4	TAATACGACTCACTATAGGGGCACTGCGGCTGGAGGTGGGTTTTAGAGCTAGAAATAGCA
fwd_sgRNA_T7_HTS_ RNF2	TAATACGACTCACTATAGGGTCATCTTAGTCATTACCTGGTTTTAGAGCTAGAAATAGCA

[00338] Sequences of *in vitro*-edited dsDNA for high-throughput sequencing (HTS). Shown are the sequences of edited strands. Reverse complements of all sequences shown were also obtained. dsDNA substrates were obtained by annealing complementary strands as described in Materials and Methods. Oligonucleotides representing the EMX1, FANCF, HEK293 site 2, HEK293 site 3, HEK293 site 4, and RNF2 loci were originally designed for use in the gel-based deaminase assay and therefore have the same 25-nt sequence on their 5'- ends (matching that of the Cy3-primer). SEQ ID NOs: 443-494 appear from top to bottom below, respectively.

Base sequence	ACGTAAACGGCCACAAGTTCTTATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
1A	ACGTAAACGGCCACAAGTTCATATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
1C	ACGTAAACGGCCACAAGTTCCTATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
1G	ACGTAAACGGCCACAAGTTCGTATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
2A	ACGTAAACGGCCACAAGTTCTAATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
2C	ACGTAAACGGCCACAAGTTCTCATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
2G	ACGTAAACGGCCACAAGTTCTGATTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
3T	ACGTAAACGGCCACAAGTTCTTTTTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
3C	ACGTAAACGGCCACAAGTTCTTCTTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
3G	ACGTAAACGGCCACAAGTTCTTGTTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
4A	ACGTAAACGGCCACAAGTTCTTAATTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
4C	ACGTAAACGGCCACAAGTTCTTACTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
4G	ACGTAAACGGCCACAAGTTCTTAGTTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
5A	ACGTAAACGGCCACAAGTTCTTATATCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
5C	ACGTAAACGGCCACAAGTTCTTATCTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
5G	ACGTAAACGGCCACAAGTTCTTATGTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
6A	ACGTAAACGGCCACAAGTTCTTATTACGTGGATTTATTTATGGCATCTTCTTCAAGGACG
6C	ACGTAAACGGCCACAAGTTCTTATTCCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
6G	ACGTAAACGGCCACAAGTTCTTATTGCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
8A	ACGTAAACGGCCACAAGTTCTTATTTCATGGATTTATTTATGGCATCTTCTTCAAGGACG
8T	ACGTAAACGGCCACAAGTTCTTATTTCTTGGATTTATTTATGGCATCTTCTTCAAGGACG
8C	ACGTAAACGGCCACAAGTTCTTATTTCCTGGATTTATTTATGGCATCTTCTTCAAGGACG
9A	ACGTAAACGGCCACAAGTTCTTATTTCGAGGATTTATTTATGGCATCTTCTTCAAGGACG
9C	ACGTAAACGGCCACAAGTTCTTATTTCGCGGATTTATTTATGGCATCTTCTTCAAGGACG
9G	ACGTAAACGGCCACAAGTTCTTATTTCGGGGATTTATTTATGGCATCTTCTTCAAGGACG
10A	ACGTAAACGGCCACAAGTTCTTATTTCGTAGATTTATTTATGGCATCTTCTTCAAGGACG
10T	ACGTAAACGGCCACAAGTTCTTATTTCGTTGATTTATTTATGGCATCTTCTTCAAGGACG
10C	ACGTAAACGGCCACAAGTTCTTATTTCGTCGATTTATTTATGGCATCTTCTTCAAGGACG
11A	ACGTAAACGGCCACAAGTTCTTATTTCGTGAATTTATTTATGGCATCTTCTTCAAGGACG
11T	ACGTAAACGGCCACAAGTTCTTATTTCGTGTATTTATTTATGGCATCTTCTTCAAGGACG
11C	ACGTAAACGGCCACAAGTTCTTATTTCGTGCATTTATTTATGGCATCTTCTTCAAGGACG

12T	ACGTAAACGGCCACAAGTTCTTATTTCGTGGTTTTATTTATGGCATCTTCTTCAAGGACG
12C	ACGTAAACGGCCACAAGTTCTTATTTCGTGGCTTTATTTATGGCATCTTCTTCAAGGACG
12G	ACGTAAACGGCCACAAGTTCTTATTTCGTGGGTTTATTTATGGCATCTTCTTCAAGGACG
13A	ACGTAAACGGCCACAAGTTCTTATTTCGTGGAATTATTTATGGCATCTTCTTCAAGGACG
13C	ACGTAAACGGCCACAAGTTCTTATTTCGTGGACTTATTTATGGCATCTTCTTCAAGGACG
13G	ACGTAAACGGCCACAAGTTCTTATTTCGTGGAGTTATTTATGGCATCTTCTTCAAGGACG
multiC	ACGTAAACGGCCACAAGTTCTTCCCCCCCCGATTTATTTATGGCATCTTCTTCAAGGACG
TCGCACCC_odd	ACGTAAACGGCCACAAGTTTCGCACCCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
CCTCGCAC_odd	ACGTAAACGGCCACAAGTTCCTCGCACGTGGATTTATTTATGGCATCTTCTTCAAGGACG
ACCCTCGC_odd	ACGTAAACGGCCACAAGTTACCCTCGCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
GCACCCTC_odd	ACGTAAACGGCCACAAGTTGCACCCTCGTGGATTTATTTATGGCATCTTCTTCAAGGACG
TCGCACCC_even	ACGTAAACGGCCACAAGTATTCGCACCCGTGGATTTATTATGGCATCTTCTTCAAGGACG
CCTCGCAC_even	ACGTAAACGGCCACAAGTATCCTCGCACGTGGATTTATTATGGCATCTTCTTCAAGGACG
ACCCTCGC_even	ACGTAAACGGCCACAAGTATACCCTCGCGTGGATTTATTATGGCATCTTCTTCAAGGACG
GCACCCTC_even	ACGTAAACGGCCACAAGTATGCACCCTCGTGGATTTATTATGGCATCTTCTTCAAGGACG
EMX1_invitro	GTAGGTAGTTAGGATGAATGGAAGGTTGGTAGGCCTGAGTCCGAGCAGAAGAAGAAGGGCTCCCATCACATCAACCGGTG
FANCF_invitro	GTAGGTAGTTAGGATGAATGGAAGGTTGGTACTCATGGAATCCCTTCTGCAGCACCTGGATCGCTTTTCCGAGCTTCTGG
HEK293_site2_invitro	GTAGGTAGTTAGGATGAATGGAAGGTTGGTAAACTGGAACACAAAGCATAGACTGCGGGGCGGGCCAGCCTGAATAGCTG
HEK293_site3_invitro	GTAGGTAGTTAGGATGAATGGAAGGTTGGTACTTGGGGCCCAGACTGAGCACGTGATGGCAGAGGAAAGGAAGCCCTGCT
HEK293_site4_invitro	GTAGGTAGTTAGGATGAATGGAAGGTTGGTACCGGTGGCACTGCGGCTGGAGGTGGGGGTTAAAGCGGAGACTCTGGTGC
RNF2_invitro	GTAGGTAGTTAGGATGAATGGAAGGTTGGTATGGCAGTCATCTTAGTCATTACCTGAGGTGTTCGTTGTAACTCATATAA

[00339] Primers for HTS of *in vitro* edited dsDNA. SEQ ID NOs: 495-503 appear from top to bottom below, respectively.

fwd_invitro_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNACGTAAACGGCCACAA
rev_invitro_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCGTCCTTGAAGAAGATGC
fwd_invitro_HEK_targets	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGTAGGTAGTTAGGATGAATGGAA
rev_EMX1_invitro	TGGAGTTCAGACGTGTGCTCTTCCGATCTCACCGGTTGATGTGATGG
rev_FANCF_invitro	TGGAGTTCAGACGTGTGCTCTTCCGATCTCCAGAAGCTCGGAAAAGC
rev_HEK293_site2_invitro	TGGAGTTCAGACGTGTGCTCTTCCGATCTCAGCTATTCAGGCTGGC
rev_HEK293_site3_invitro	TGGAGTTCAGACGTGTGCTCTTCCGATCTAGCAGGGCTTCCTTTC
rev_HEK293_site4_invitro	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCACCAGAGTCTCCG
rev_RNF2_invitro	TGGAGTTCAGACGTGTGCTCTTCCGATCTTTATATGAGTTACAACGAACACC

[00340] Primers for HTS of on-target and off-target sites from all mammalian cell culture experiements. SEQ ID NOs: 504-579 and 1869-1900 appear from top to bottom below, respectively.

fwd_EMX1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCAGCTCAGCCTGAGTGTTGA
rev_EMX1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCTCGTGGGTTTGTGGTTGC
fwd_FANCF_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCATTGCAGAGAGGCGTATCA
rev_FANCF_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGGGGTCCCAGGTGCTGAC
fwd_HEK293_site2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCAGCCCCATCTGTCAAACT
rev_HEK293_site2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTGAATGGATTCCTTGGAAACAATGA
fwd_HEK293_site3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNATGTGGGCTGCCTAGAAAGG
rev_HEK293_site3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCCCAGCCAAACTTGTCAACC
fwd_HEK293_site4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGAACCCAGGTAGCCAGAGAC
rev_HEK293_site4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTCCTTTCAACCCGAACGGAG
fwd_RNF2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCTCTTCTTTATTTCCAGCAATGT
rev_RNF2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGTTTTCATGTTCTAAAAATGTATCCCA
fwd_p53_Y163C_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTACAGTACTCCCCTGCCCTC
rev_p53_Y163C_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCTGCTCACCATCGCTATCT
fwd_p53_N239D_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCTCATCTTGGGCCTGTGTT
rev_p53_N239D_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTAAATCGGTAAGAGGTGGGCC
fwd_APOE4_C158R_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGCGGACATGGAGGACGTG
rev_APOE4_C158R_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCTGTTCCACCAGGGGCCC
fwd_EMX1_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTGCCCAATCATTGATGCTTTT
rev_EMX1_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTAGAAACATTTACCATAGACTATCACCT
fwd_EMX1_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNAGTAGCCTCTTTCTCAATGTGC
rev_EMX1_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCTTTCACAAGGATGCAGTCT
fwd_EMX1_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGAGCTAGACTCCGAGGGGA
rev_EMX1_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTCCTCGTCCTGCTCTCACTT
fwd_EMX1_off4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNAGAGGCTGAAGAGGAAGACCA
rev_EMX1_off4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGGCCCAGCTGTGCATTCTAT

fwd_EMX1_off5_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCAAGAGGGCCAAGTCCTG
rev_EMX1_off6_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCAGCGAGGAGTGACAGCC
fwd_EMX1_off7_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCACTCCACCTGATCTCGGGG
rev_EMX1_off7_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCGAGGAGGGAGGGAGCAG
fwd_EMX1_off8_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNACCACAAATGCCCAAGAGAC
rev_EMX1_off8_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGACACAGTCAAGGGCCGG
fwd_EMX1_off9_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCCACCTTTGAGGAGGCAAA
rev_EMX1_off9_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTTCCATCTGAGAAGAGAGTGGT
fwd_EMX1_off10_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGTCATACCTTGGCCCTTCCT
rev_EMX1_off10_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTCCCTAGGCCCACACCAG
fwd_FANCF_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNAACCCACTGAAGAAGCAGGG
rev_FANCF_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGGTGCTTAATCCGGCTCCAT
fwd_FANCF_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTCCAGTGTTTCCATCCCGAA
rev_FANCF_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCCTCTGACCTCCACAACTCT
fwd_FANCF_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCTGGGTACAGTTCTGCGTGT
rev_FANCF_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTCACTCTGAGCATCGCCAAG
fwd_FANCF_off4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGGTTTAGAGCCAGTGAACTAGAG
rev_FANCF_off4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCAAGACAAAATCCTCTTTATACTTTG
fwd_FANCF_off5_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGGGAGGGGACGGCCTTAC
rev_FANCF_off5_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCCTCTGGCGAACATGGC
fwd_FANCF_off6_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTCCTGGTTAAGAGCATGGGC
rev_FANCF_off6_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGATTGAGTCCCCACAGCACA
fwd_FANCF_off7_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCAGTGTTTCCCATCCCCAA
rev_FANCF_off7_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTGACCTCCACAACTGGAAAAT
fwd_FANCF_off8_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGCTTCCAGACCCACCTGAAG
rev_FANCF_off8_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTACCGAGGAAAATTGCTTGTCG
fwd_HEK293_site2_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGTGTGGAGAGTGAGTAAGCCA
rev_HEK293_site2_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTACGGTAGGATGATTTCAGGCA
fwd_HEK293_site2_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCACAAAGCAGTGTAGCTCAGG
rev_HEK293_site2_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTTTTTTGGTACTCGAGTGTTATTCAG
fwd_HEK293_site3_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTCCCCTGTTGACCTGGAGAA
rev_HEK293_site3_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCACTGTACTTGCCCTGACCA
fwd_HEK293_site3_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTTGGTGTTGACAGGGAGCAA
rev_HEK293_site3_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCTGAGATGTGGGCAGAAGGG
fwd_HEK293_site3_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTGAGAGGGAACAGAAGGGCT
rev_HEK293_site3_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGTCCAAAGGCCCAAGAACCT
fwd_HEK293_site3_off4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTCCTAGCACTTTGGAAGGTCG
rev_HEK293_site3_off4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCTCATCTTAATCTGCTCAGCC
fwd_HEK293_site3_off5_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNAAAGGAGCAGCTCTTCCTGG
rev_HEK293_site3_off5_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGTCTGCACCATCTCCCACAA

fwd_HEK293_site4_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGGCATGGCTTCTGAGACTCA
rev_HEK293_site4_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGTCTCCCTTGCACTCCCTGTCTTT
fwd_HEK293_site4_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTTTGGCAATGGAGGCATTGG
rev_HEK293_site4_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGAAGAGGCTGCCCATGAGAG
fwd_HEK293_site4_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGGTCTGAGGCTCGAATCCTG
rev_HEK293_site4_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCTGTGGCCTCCATATCCCTG
fwd_HEK293_site4_off4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTTTCCACCAGAACTCAGCCC
rev_HEK293_site4_off4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCCTCGGTTCCTCCACAACAC
fwd_HEK293_site4_off5_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCACGGGAAGGACAGGAGAAG
rev_HEK293_site4_off5_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCAGGGGAGGGATAAAGCAG
fwd_HEK293_site4_off6_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCACGGGAGATGGCTTATGT
rev_HEK293_site4_off6_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCACATCCTCACTGTGCCACT
fwd_HEK293_site4_off7_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGTCAGTCTCGGCCCCTCA
rev_HEK293_site4_off7_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCCACTGTAAAGCTCTTGGG
fwd_HEK293_site4_off8_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNAGGGTAGAGGGACAGAGCTG
rev_HEK293_site4_off8_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGGACCCCACATAGTCAGTGC
fwd_HEK293_site4_off9_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGCTGTCAGCCCTATCTCCATC
rev_HEK293_site4_off9_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTGGGCAATTAGGACAGGGAC
fwd_HEK293_site4_off10_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGCAGCGGAGGAGGTAGATTG
rev_HEK293_site4_off10_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCTCAGTACCTGGAGTCCCGA
fwd_HEK2_ChIP_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGACAGGCTCAGGAAAGCTGT
rev_HEK2_ChIP_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTACACAAGCCTTTCTCCAGGG
fwd_HEK2_ChIP_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNAATAGGGGGTGAGACTGGGG
rev_HEK2_ChIP_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTGCCTCAGACGAGACTTGAGG
fwd_HEK2_ChIP_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGGCCAGCAGGAAAGGAATCT
rev_HEK2_ChIP_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTGACTGCACCTGTAGCCATG
fwd_HEK2_ChIP_off4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTCAAGGAAATCACCCTGCCC
rev_HEK2_ChIP_off4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTAACTTCCTTGGTGTGCAGCT
fwd_HEK2_ChIP_off5_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNATGGGCTCAGCTACGTCATG
rev_HEK2_ChIP_off5_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTAATAGCAGTGTGGTGGGCAA

fwd_HEK3_ChIP_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCGCACATCCCTTGTCTCTCT
rev_HEK3_ChIP_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCTACTGGAGCACACCCCAAG
fwd_HEK3_ChIP_off2_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNTGGGTCACGTAGCTTTGGTC
rev_HEK3_ChIP_off2_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTGGTGGCCATGTGCAACTAA
fwd_HEK3_ChIP_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCTACTACGTGCCAGCTCAGG
rev_HEK3_ChIP_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTACCTCCCCTCCTCACTAACC
fwd_HEK3_ChIP_off4_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGCCTCAGCTCCATTTCCTGT
rev_HEK3_ChIP_off4_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTAACCTTTATGGCACCAGGGG
fwd_HEK3_ChIP_off5_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGAGCTCAGCATTAGCAGGCT
rev_HEK3_ChIP_off5_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTTTCCTGGCTTTCCGATTCCC
fwd_HEK4_ChIP_off1_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNGTGCAATTGGAGGAGGAGCT
rev_HEK4_ChIP_off1_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCACCAGCTACAGGCAGAACA
fwd_HEK4_ChIP_off3_HTS	ACACTCTTTCCCTACACGACGCTCTTCCGATCTNNNNCCTACCCCCAACACAGATGG
rev_HEK4_ChIP_off3_HTS	TGGAGTTCAGACGTGTGCTCTTCCGATCTCCACACAACTCAGGTCCTCC

[00341] Sequences of single-stranded oligonucleotide donor templates (ssODNs) used in HDR studies.

EMX1 sense (SEQ ID NO: 580)

TCATCTGTGCCCCTCCCTCCCTGGCCCAGGTGAAGGTGTGGTTCCAGAACCGGAGGACAAAGTACA
AACGGCAGAAGCTGGAGGAGGAAGGGCCTGAGTTTGAGCAGAAGAAGAAGGGCTCCCATCACATC
AACCGGTGGCGCATTGCCACGAAGCAGGCCAATGGGGAGGACATCGATGTCACCTCCAATGACTAG
GGT

EMX1 antisense (SEQ ID NO: 581)

ACCCTAGTCATTGGAGGTGACATCGATGTCCTCCCCATTGGCCTGCTTCGTGGCAATGCGCCACCG
GTTGATGTGATGGGAGCCCTTCTTCTTCTGCTCAAACTCAGGCCCTTCCTCCTCCAGCTTCTGCCGT
TTGTACTTTGTCCTCCGGTTCTGGAACCACACCTTCACCTGGGCCAGGGAGGGAGGGGCACAGATG
A

HEK293 site 3 sense (SEQ ID NO: 582)

CATGCAATTAGTCTATTTCTGCTGCAAGTAAGCATGCATTTGTAGGCTTGATGCTTTTTTTCTGCTTCT
CCAGCCCTGGCCTGGGTCAATCCTTGGGGCTTAGACTGAGCACGTGATGGCAGAGGAAAGGAAGC
CCTGCTTCCTCCAGAGGGCGTCGCAGGACAGCTTTTCCTAGACAGGGGCTAGTATGTGCAGCTCCT

HEK293 site 3 antisense (SEQ ID NO: 583)

AGGAGCTGCACATACTAGCCCCTGTCTAGGAAAAGCTGTCCTGCGACGCCCTCTGGAGGAAGCAGG
GCTTCCTTTCCTCTGCCATCACGTGCTCAGTCTAAGCCCCAAGGATTGACCCAGGCCAGGGCTGGA
GAAGCAGAAAAAAAGCATCAAGCCTACAAATGCATGCTTACTTGCAGCAGAAATAGACTAATTGCATG

HEK site 4 sense (SEQ ID NO: 584)

GGCTGACAAAGGCCGGGCTGGGTGGAAGGAAGGGAGGAAGGGCGAGGCAGAGGGTCCAAAGCAG
GATGACAGGCAGGGGCACCGCGGCGCCCCGGTGGCATTGCGGCTGGAGGTGGGGGTTAAAGCGG
AGACTCTGGTGCTGTGTGACTACAGTGGGGGCCCTGCCCTCTCTGAGCCCCCGCCTCCAGGCCTGT
GTGTGT

HEK site 4 antisense (SEQ ID NO: 585)

ACACACACAGGCCTGGAGGCGGGGGCTCAGAGAGGGCAGGGCCCCCACTGTAGTCACACAGCACC
AGAGTCTCCGCTTTAACCCCCACCTCCAGCCGCAATGCCACCGGGGCGCCGCGGTGCCCCTGCCT
GTCATCCTGCTTTGGACCCTCTGCCTCGCCCTTCCTCCCTTCCTTCCACCCAGCCCGGCCTTTGTCA
GCC

APOE4 sense (SEQ ID NO: 743)

AGCACCGAGGAGCTGCGGGTGCGCCTCGCCTCCCACCTGCGCAAGCTGCGTAAGCGGCTCCTCCG
CGATGCCGATGACCTGCAGAAGTGCCTGGCAGTGTACCAGGCCGGGGCCCGCGAGGGCGCCGAG

CGCGGCCTCAGCGCCATCCGCGAGCGCCTGGGGCCCCTGGTGGAACAGGGCCGCGTGCGGGCCG
CCACTGT

APOE4 antisense (SEQ ID NO: 744)

ACAGTGGCGGCCCGCACGCGGCCCTGTTCCACCAGGGGCCCCAGGCGCTCGCGGATGGCGCTGA
GGCCGCGCTCGGCGCCCTCGCGGGCCCCGGCCTGGTACACTGCCAGGCACTTCTGCAGGTCATCG
GCATCGCGGAGGAGCCGCTTACGCAGCTTGCGCAGGTGGGAGGCGAGGCGCACCCGCAGCTCCT
CGGTGCT

p53 Y163C sense (SEQ ID NO: 745)

ACTCCCCTGCCCTCAACAAGATGTTTTGCCAACTGGCCAAGACCTGCCCTGTGCAGCTGTGGGTTGA
TTCCACACCCCCGCCCGGCACCCGCGTCCGCGCCATGGCCATCTACAAGCAGTCACAGCACATGAC
GGAGGTTGTGAGGCGCTGCCCCCACCATGAGCGCTGCTCAGATAGCGATGGTGAGCAGCTGGGGC
TG

p53 Y163C antisense (SEQ ID NO: 746)

CAGCCCCAGCTGCTCACCATCGCTATCTGAGCAGCGCTCATGGTGGGGGCAGCGCCTCACAACCTC
CGTCATGTGCTGTGACTGCTTGTAGATGGCCATGGCGCGGACGCGGGTGCCGGGCGGGGGTGTGG
AATCAACCCACAGCTGCACAGGGCAGGTCTTGGCCAGTTGGCAAAACATCTTGTTGAGGGCAGGGG
AGT

[00342] Deaminase gene gBlocks Gene Fragments

hAID (SEQ ID NO: 586)

CATCCTTGGTACCGAGCTCGGATCCAGCCACCATGGATAGCCTCTTGATGAATAGACGCAAGTTCCT
GTATCAGTTTAAAAACGTGAGATGGGCAAAAGGCCGACGAGAGACATATCTGTGCTATGTCGTTAAG
CGCAGAGATTCAGCCACCAGTTTCTCTCTCGACTTCGGCTACCTGCGGAACAAGAATGGTTGCCATG
TTGAGCTCCTGTTCCTGAGGTATATCAGCGACTGGGATTTGGACCCAGGGCGGTGCTATAGGGTGA
CATGGTTTACCTCCTGGTCACCTTGTTATGACTGCGCGCGGCATGTTGCCGATTTTCTGAGAGGGAA
CCCTAACCTGTCTCTGAGGATCTTCACCGCGCGACTGTACTTCTGTGAGGACCGGAAAGCCGAACC
CGAGGGACTGAGACGCCTCCACAGAGCGGGTGTGCAGATTGCCATAATGACCTTTAAGGACTACTT
CTACTGCTGGAACACCTTCGTCGAAAATCACGAGCGGACTTTCAAGGCTTGGGAAGGATTGCATGAA
AACAGCGTCAGGCTTTCCAGGCAGCTTCGCCGCATTCTTCTCCCGTTGTACGAGGTTGATGACCTCA
GAGATGCCTTTAGAACACTGGGACTGTAGGCGGCCGCTCGATTGGTTTGGTGTGGCTCTAA

rAPOBEC1 (mammalian)(SEQ ID NO: 587)

CATCCTTGGTACCGAGCTCGGATCCAGCCACCATGAGCTCAGAGACTGGCCCAGTGGCTGTGGACC
CCACATTGAGACGGCGGATCGAGCCCCATGAGTTTGAGGTATTCTTCGATCCGAGAGAGCTCCGCA
AGGAGACCTGCCTGCTTTACGAAATTAATTGGGGGGGCCGGCACTCCATTTGGCGACATACATCACA
GAACACTAACAAGCACGTCGAAGTCAACTTCATCGAGAAGTTCACGACAGAAAGATATTTCTGTCCG
AACACAAGGTGCAGCATTACCTGGTTTCTCAGCTGGAGCCCATGCGGCGAATGTAGTAGGGCCATC
ACTGAATTCCTGTCAAGGTATCCCCACGTCACTCTGTTTATTTACATCGCAAGGCTGTACCACCACGC
TGACCCCCGCAATCGACAAGGCCTGCGGGATTTGATCTCTTCAGGTGTGACTATCCAAATTATGACT
GAGCAGGAGTCAGGATACTGCTGGAGAAACTTTGTGAATTATAGCCCGAGTAATGAAGCCCACTGG
CCTAGGTATCCCCATCTGTGGGTACGACTGTACGTTCTTGAACTGTACTGCATCATACTGGGCCTGC
CTCCTTGTCTCAACATTCTGAGAAGGAAGCAGCCACAGCTGACATTCTTTACCATCGCTCTTCAGTCT
TGTCATTACCAGCGACTGCCCCCACACATTCTCTGGGCCACCGGGTTGAAATGAGCGGCCGCTCGA
TTGGTTTGGTGTGGCTCTAA

pmCDA1 (SEQ ID NO: 588)

CATCCTTGGTACCGAGCTCGGATCCAGCCACCATGACAGACGCTGAATATGTTAGGATCCATGAAAA
ACTGGATATCTATACATTTAAGAAGCAGTTCTTCAATAACAAAAAGTCAGTATCTCACAGATGCTATGT
CCTGTTCGAACTCAAGAGAAGAGGAGAAAGGCGGGCCTGTTTCTGGGGGTACGCGGTTAATAAACC
CCAGTCCGGGACCGAGAGGGGGATTCACGCCGAGATCTTTTCAATTAGGAAGGTTGAAGAGTATCT
TCGCGACAATCCCGGTCAGTTCACAATTAACTGGTACAGCTCCTGGAGCCCTTGCGCTGATTGCGCC
GAGAAAATACTCGAATGGTACAACCAGGAGTTGAGAGGCAATGGCCACACTCTCAAGATTTGGGCTT
GCAAGCTTTACTACGAGAAGAACGCGAGAAATCAGATTGGCTTGTGGAACCTCAGGGACAACGGGG
TCGGGTTGAATGTTATGGTGTCCGAACATTACCAGTGCTGTAGAAAGATCTTCATTCAGTCCAGTCAC
AATCAGCTGAACGAGAACAGATGGCTGGAGAAAACACTGAAACGGGCAGAGAAAAGGCGCTCAGAG
CTGAGTATCATGATCCAGGTCAAAATCCTGCATACAACCAAAAGCCCGGCTGTATAAGCGGCCGCTC
GATTGGTTTGGTGTGGCTCTAA

haPOBEC3G (SEQ ID NO: 589)

CATCCTTGGTACCGAGCTCGGATCCAGCCACCATGGAGCTGAAGTATCACCCTGAGATGCGGTTTTT
CCACTGGTTTAGTAAGTGGCGCAAACTTCATCGGGATCAGGAGTATGAAGTGACCTGGTATATCTCT
TGGTCTCCCTGCACAAAATGTACACGCGACATGGCCACATTTCTGGCCGAGGATCCAAAGGTGACG
CTCACAATCTTTGTGGCCCGCCTGTATTATTTCTGGGACCCGGATTATCAGGAGGCACTTAGGTCAT
TGTGCCAAAAGCGCGACGGACCACGGGCGACTATGAAAATCATGAATTATGACGAATTCCAGCATTG
CTGGAGTAAGTTTGTGTACAGCCAGCGGGAGCTGTTCGAGCCCTGGAACAATCTTCCCAAGTACTAC
ATACTGCTTCACATTATGTTGGGGGAGATCCTTCGGCACTCTATGGATCCTCCTACCTTTACGTTTAA
CTTTAATAATGAGCCTTGGGTTCGCGGGCGCCATGAAACCTATTTGTGCTACGAGGTCGAGCGGATG
CATAATGATACGTGGGTCCTGCTGAATCAGAGGAGGGGGTTTCTGTGTAACCAGGCTCCACATAAAC
ATGGATTTCTCGAGGGGCGGCACGCCGAACTGTGTTTCCTTGATGTGATACCTTTCTGGAAGCTCGA
CCTTGATCAAGATTACAGGGTGACGTGTTTCACCTCCTGGTCACCCTGCTTCAGTTGCGCCCAAGAG
ATGGCTAAATTTATCAGTAAGAACAAGCATGTGTCCCTCTGTATTTTTACAGCCAGAATTTATGATGAC
CAGGGCCGGTGCCAGGAGGGGCTGCGGACACTCGCTGAGGCGGGCGCGAAGATCAGCATAATGA
CATACTCCGAATTCAAACACTGTTGGGACACTTTTGTGGACCACCAGGGCTGCCCATTTCAGCCGTG
GGATGGGCTCGACGAACATAGTCAGGATCTCTCAGGCCGGCTGCGAGCCATATTGCAGAACCAGGA
GAATTAGGCGGCCGCTCGATTGGTTTGGTGTGGCTCTAA

rAPOBEC1(E. Coli) (SEQ ID NO: 590)

GGCCGGGGATTCTAGAAATAATTTTGTTTAACTTTAAGAAGGAGATATACCATGGATGTCTTCTGAAA
CCGGTCCGGTTGCGGTTGACCCGACCCTGCGTCGTCGTATCGAACCGCACGAATTCGAAGTTTTCT
TCGACCCGCGTGAACTGCGTAAAGAAACCTGCCTGCTGTACGAAATCAACTGGGGTGGTCGTCACT
CTATCTGGCGTCACACCTCTCAGAACACCAACAAACACGTTGAAGTTAACTTCATCGAAAAATTCACC
ACCGAACGTTACTTCTGCCCGAACACCCGTTGCTCTATCACCTGGTTCCTGTCTTGGTCTCCGTGCG
GTGAATGCTCTCGTGCGATCACCGAATTCCTGTCTCGTTACCCGCACGTTACCCTGTTCATCTACATC
GCGCGTCTGTACCACCACGCGGACCCGCGTAACCGTCAGGGTCTGCGTGACCTGATCTCTTCTGGT
GTTACCATCCAGATCATGACCGAACAGGAATCTGGTTACTGCTGGCGTAACTTCGTTAACTACTCTCC
GTCTAACGAAGCGCACTGGCCGCGTTACCCGCACCTGTGGGTTCGTCTGTACGTTCTGGAACTGTA
CTGCATCATCCTGGGTCTGCCGCCGTGCCTGAACATCCTGCGTCGTAAACAGCCGCAGCTGACCTT

CTTCACCATCGCGCTGCAGTCTTGCCACTACCAGCGTCTGCCGCCGCACATCCTGTGGGCGACCGG
TCTGAAAGGTGGTAGTGGAGGGAGCGGCGGTTCAATGGATAAGAAATAC

[00343] Amino Acid Sequences of NBE1, NBE2, and NBE3.

NBE1 for *E. Coli* expression (His_6-rAPOBEC1-XTEN-dCas9) (SEQ ID NO: 591)

MGSSHHHHHHMSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTN
KHVEVNFIEKFTTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGL
RDLISSGVTIQIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLELYCIILGLPPCLNILRRKQPQ
LTFFTIALQSCHYQRLPPHILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSK
KFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRL
EESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDL
NPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSL
GLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSA
SMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELL
VKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRF
AWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTE
GMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDK
DFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQS
GKTILDFLKSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDEL
VKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQ
NGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQL
LNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLK
SKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIG
KATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQT
GGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSS
FEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKL
KGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNL
GAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGSPKKKRKV

NBE1 for Mammalian expression (rAPOBEC1-XTEN-dCas9-NLS) (SEQ ID NO: 592)

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKHVEVNFIEKF
TTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGLRDLISSGVTIQ
IMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSC
HYQRLPPHILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDR
HSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKK
HERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLF
IQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDL
AEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQ
DLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQ
RTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITP
WNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQ
KKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILE
DIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFA
NRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVI
EMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDI
NRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNL
TKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQF
YKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIM
NFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRN
SDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKG
YKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQL
FVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTI
DRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGSPKKKRKV

Alternative NBE1 for Mammalian expression with human APOBEC1 (hAPOBEC1-XTEN-dCas9-NLS) (SEQ ID NO: 5737)

MTSEKGPSTGDPTLRRRIEPWEFDVFYDPRELRKEACLLYEIKWGMSRKIWRSSGKNTTN
HVEVNFIKKFTSERDFHPSMSCSITWFLSWSPCWECSQAIREFLSRHPGVTLVIYVARLFW
HMDQQNRQGLRDLVNSGVTIQIMRASEYYHCWRNFVNYPPGDEAHWPQYPPLWMMLY
ALELHCIILSLPPCLKISRRWQNHLTFFRLHLQNCHYQTIPPHILLATGLIHPSVAWRGSETP
GTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALL
FDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKK
HERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGD
LNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKK
NGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAA
KNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQ
SKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIH
LGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITPW
NFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMR
KPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYH
DLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRY
TGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQ
GDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQK
NSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSD
YDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQ
RKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVK
VITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDY
KVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVW
DKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGG
FDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKD
LIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNE
QKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTL
TNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGSPKKK
RKV

NBE2 (rAPOBEC1-XTEN-dCas9-UGI-NLS) (SEQ ID NO: 593)

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKHVEVNFIEKF
TTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGLRDLISSGVTIQ
IMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSC
HYQRLPPHILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDR
HSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKK
HERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLF
IQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDL
AEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQ
DLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQ
RTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITP
WNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQ
KKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILE
DIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFA
NRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVI
EMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDI
NRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNL
TKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQF
YKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIM
NFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRN
SDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKG
YKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQL
FVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTI
DRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIG
NKPESDILVHTAYDESTDENVMLLTSDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV

NBE3 (rAPOBEC1-XTEN-Cas9n-UGI-NLS) (SEQ ID NO: 594)

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKHVEVNFIEKF
TTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGLRDLISSGVTIQ
IMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQSC
HYQRLPPHILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDR
HSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKK
HERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLF
IQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDL
AEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQ
DLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQ
RTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITP
WNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQ
KKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILE
DIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFA
NRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVI
EMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDI
NRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNL
TKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQF
YKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIM
NFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRN
SDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKG
YKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQL
FVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTI
DRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGDSGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIG
NKPESDILVHTAYDESTDENVMLLTSDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV

pmCDA1-XTEN-dCas9-UGI (bacteria) (SEQ ID NO: 5742)
MTDAEYVRIHEKLDIYTFKKQFFNNKKSVSHRCYVLFELKRRGERRACFWGYAVNKPQS
GTERGIHAEIFSIRKVEEYLRDNPGQFTINWYSSWSPCADCAEKILEWYNQELRGNGHTL
KIWACKLYYEKNARNQIGLWNLRDNGVGLNVMVSEHYQCCRKIFIQSSHNQLNENRWL

EKTLKRAEKRRSELSIMIQVKILHTTKSPAVSGSETPGTSESATPESDKKYSIGLAIGTNSV
GWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRK
NRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYH
LRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFE
ENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLA
EDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSAS
MIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILE
KMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEK
ILTFRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLP
NEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVK
QLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLT
LFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLK
SDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVV
DELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENT
QLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKN
RGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQL
VETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNY
HHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYS
NIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQ
TGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSV
KELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQ
KGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVIL
ADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEV
LDATLIHQSITGLYETRIDLSQLGGDSGGSMTNLSDIIEKETGKQLVIQESILMLPEEVEEVI
GNKPESDILVHTAYDESTDENVMLLTSDAPEYKPWALVIQDSNGENKIKML

pmCDA1-XTEN-nCas9-UGI-NLS (mammalian construct) (SEQ ID NO: 5743)

MTDAEYVRIHEKLDIYTFKKQFFNNKKSVSHRCYVLFELKRRGERRACFWGYAVNKPQS
GTERGIHAEIFSIRKVEEYLRDNPGQFTINWYSSWSPCADCAEKILEWYNQELRGNGHTL
KIWACKLYYEKNARNQIGLWNLRDNGVGLNVMVSEHYQCCRKIFIQSSHNQLNENRWL
EKTLKRAEKRRSELSIMIQVKILHTTKSPAVSGSETPGTSESATPESDKKYSIGLAIGTNSV
GWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRK
NRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYH
LRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFE
ENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLA
EDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSAS
MIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILE
KMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEK
ILTFRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLP
NEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVK
QLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLT
LFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLK
SDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVV
DELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENT
QLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKN
RGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQL

VETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNY
HHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYS
NIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQ
TGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSV
KELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQ
KGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVIL
ADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEV
LDATLIHQSITGLYETRIDLSQLGGDSGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIG
NKPESDILVHTAYDESTDENVMLLTSDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKR
KV

huAPOBEC3G-XTEN-dCas9-UGI (bacteria) (SEQ ID NO: 5744)
MDPPTFTFNFNNEPWVRGRHETYLCYEVERMHNDTWVLLNQRRGFLCNQAPHKHGFLE
GRHAELCFLDVIPFWKLDLDQDYRVTCFTSWSPCFSCAQEMAKFISKNKHVSLCIFTARIY
DDQGRCQEGLRTLAEAGAKISIMTYSEFKHCWDTFVDHQGCPFQPWDGLDEHSQDLSGR
LRAILQSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDR
HSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRL
EESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMI
KFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLE
NLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIG
DQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQL
PEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRT
FDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMT
RKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTK
VKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDR
FNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKV
MKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMQLIHDDSLTFKED
IQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMAREN
QTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQ
ELDINRLSDYDVDAIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQ
LLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDE
NDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLE
SEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETN
GETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKD
WDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAK
GYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKL
KGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQA
ENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD
SGGSMTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLL
TSDAPEYKPWALVIQDSNGENKIKML

huAPOBEC3G-XTEN-nCas9-UGI-NLS (mammalian construct) (SEQ ID NO: 5745)

MDPPTFTFNFNNEPWVRGRHETYLCYEVERMHNDTWVLLNQRRGFLCNQAPHKHGFLE
GRHAELCFLDVIPFWKLDLDQDYRVTCFTSWSPCFSCAQEMAKFISKNKHVSLCIFTARIY
DDQGRCQEGLRTLAEAGAKISIMTYSEFKHCWDTFVDHQGCPFQPWDGLDEHSQDLSGR
LRAILQSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDR

HSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRL
EESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMI
KFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLE
NLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIG
DQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQL
PEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRT
FDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMT
RKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTK
VKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDR
FNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKV
MKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMQLIHDDSLTFKED
IQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMAREN
QTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQ
ELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQ
LLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDE
NDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLE
SEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETN
GETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKD
WDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAK
GYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKL
KGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQA
ENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD
SGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLLT
SDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV

huAPOBEC3G (D316R_D317R)-XTEN-nCas9-UGI-NLS (mammalian construct) (SEQ ID NO: 5746)

MDPPTFTFNFNNEPWVRGRHETYLCYEVERMHNDTWVLLNQRRGFLCNQAPHKHGFLE
GRHAELCFLDVIPFWKLDLDQDYRVTCFTSWSPCFSCAQEMAKFISKNKHVSLCIFTARIY
RRQGRCQEGLRTLAEAGAKISIMTYSEFKHCWDTFVDHQGCPFQPWDGLDEHSQDLSGR
LRAILQSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDR
HSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRL
EESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMI
KFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLE
NLIAQLPGEKKNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIG
DQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQL
PEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRT
FDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMT
RKSEETITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTK
VKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDR
FNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKV
MKQLKRRRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMQLIHDDSLTFKED
IQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMAREN
QTTQKGQKNSRERMKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQ
ELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQ
LLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDE

NDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLE
SEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETN
GETGEIVWDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKD
WDPKKYGGFDSPTVAYSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAK
GYKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKL
KGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQA
ENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD
SGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDESTDENVMLLT
SDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV

[00344] Base Calling Matlab Script

WTnuc='GCGGACATGGAGGACGTGCGCGGCCGCCTGGTGCAGTACCGCGGCGAGGTGCAGGCCATGCTCGGC
CAGA
GCACCGAGGAGCTGCGGGTGCGCCTCGCCTCCCACCTGCGCAAGCTGCGTAAGCGGCTCCTCCGCGATGCCG
ATGAC
CTGCAGAAGCGCCTGGCAGTGTACCAGGCCGGGGCCCGCGAGGGCGCCGAGCGCGGCCTCAGCGCCATCCGC
GAGCG CCTGGGGCCCCTGGTGGAACAG'(SEQ ID NO: 595);

```
%cycle through fastq files for different samples  files=dir('*.fastq');
for d=1:20
filename=files(d).name;
%read fastq file
[header,seqs,qscore]=fastqread(filename);
seqsLength=length(seqs);                                    % number of sequences  seqsFile=
strrep(filename,'.fastq','');                               % trims off .fastq
%create a directory with the same name as fastq file  ifexist(seqsFile,'dir');
        error('Directory already exists. Please rename or move it before  moving on.');
end
mkdir(seqsFile);                                            % make directory
wtLength=length(WTnuc);                                     % length of wildtype sequence
%% aligning back to the wildtype nucleotide sequence
%
% AlN is a matrix of the nucleotide alignment  window=1:wtLength;
sBLength=length(seqs);                                      % number of sequences
% counts number of skips  nSkips = 0;
ALN=repmat('',[sBLength wtLength]);
% iterate through each sequencing read  for i = 1:sBLength
%If you only have forward read fastq files leave as is
%If you have R1 foward and R2 is reverse fastq files uncomment the
%next four lines of code and the subsequent end statement
%               if mod(d,2)==0;
%                       reverse=seqrcomplement(seqs{i});
%                       [score,alignment,start]=
swalign(reverse,WTnuc,'Alphabet','NT');
%               else

[score,alignment,start]=swalign(seqs{i},WTnuc,'Alphabet','NT');
%               end

% length of the sequencing read  len=
length(alignment(3,:));
% if there is a gap in the alignment , skip = 1 and we will
```

```
        % throw away the entire read  skip = 0;
        for j = 1:len
if (alignment(3,j) == '-' || alignment(1,j) == '-')   skip = 1;
                                        break;
        end
        %in addition if the qscore for any given base in the read is
                                %below 31 the nucleotide is turned into an N (fastq qscores that   are not letters)
        ifisletter(qscore{i}(start(1)+j-1)) else
        alignment(1,j) = 'N';
                end

        end
        if skip == 0 && len>10
        ALN(i,    start(2):(start(2)+length(alignment)-1))=alignment(1,:);
                        end
        end
```

```
% with the alignment matrices we can simply tally up the occurrences of
% each nucleotide at each column in the alignment these
% tallies ignore bases annotated as N
% due to low qscores
TallyNTD=zeros(5,wtLength); fori=1:wtLength

TallyNTD(:,i)=[sum(ALN(:,i)=='A'),sum(ALN(:,i)=='C'),sum(ALN(:,i)=='G'),sum(A
LN(:,i)=='T'),sum(ALN(:,i)=='N')];
end
% we then save these tally matrices in the respective folder for
% further processing

save(strcat(seqsFile,'/TallyNTD'),'TallyNTD'); dlmwrite(strcat(seqsFile,'/TallyNTD.txt'),TallyNTD,'precision',
'%.3f','newline','pc'); end
```

[00345] INDEL Detection Matlab Script

```
WTnuc='GCGGACATGGAGGACGTGCGCGGCCGCCTGGTGCAGTACCGCGGCGAGGTGCAGGCCATGCTCGGC
CAGA
GCACCGAGGAGCTGCGGGTGCGCCTCGCCTCCCACCTGCGCAAGCTGCGTAAGCGGCTCCTCCGCGATGCCG
ATGAC
CTGCAGAAGCGCCTGGCAGTGTACCAGGCCGGGGCCCGCGAGGGCGCCGAGCGCGGCCTCAGCGCCATCCGC
GAGCG CCTGGGGCCCCTGGTGGAACAG'(SEQ ID NO: 595);

%cycle through fastq files for different samples files=dir('*.fastq');
%specify start and width of indel window as well as length of each flank indelstart=154;
width=30; flank=10;

for d=1:3
filename=files(d).name;
%read fastq file
[header,seqs,qscore]=fastqread(filename);
seqsLength=length(seqs);                                    % number of sequences seqsFile
=strcat(strrep(filename,'.fastq',''),'_INDELS');
%create a directory with the same name as fastq file+_INDELS ifexist(seqsFile,'dir');
        error('Directory already exists. Please rename or move it before moving on.');
end
mkdir(seqsFile);                                            % make directory
wtLength=length(WTnuc);                                     % length of wildtype sequence sBLength=
length(seqs);                                               % number of sequences

% initialize counters and cell arrays
nSkips = 0; notINDEL=0;
ins={};
dels={}; NumIns=0;
NumDels=0;
% iterate through each sequencing read for i = 1:sBLength
    %search for 10BP sequences that should flank both sides of the "INDEL WINDOW"
  windowstart=strfind(seqs{i},WTnuc(indelstart-flank:indelstart));

                windowend=strfind(seqs{i},WTnuc(indelstart+width:indelstart+width+flank
));
%if the flanks are found proceed
iflength(windowstart)==1&& length(windowend)==1
%if the sequence length matches the INDEL window length save as
```

```
        %not INDEL
if   windowend-windowstart==width+flank notINDEL=notINDEL+1;
        %if the sequence is two or more bases longer than the INDEL
        %window length save as an Insertion
elseif   windowend-windowstart>=width+flank+2  NumIns=NumIns+1;
        ins{NumIns}=seqs{i};
        %if the sequence is two or more bases shorter than the INDEL
        %window length save as a Deletion
elseif   windowend-windowstart<=width+flank-2   NumDels=NumDels+1;
        dels{NumDels}=seqs{i};
        %keep track of skipped sequences that are either one base
        %shorter or longer than the INDEL window width  else
        nSkips=nSkips+1;
        end
        %keep track of skipped sequences that do not possess matching flank
        %sequences else
        nSkips=nSkips+1;

                        end
        end
```

```
fid=fopen(strcat(seqsFile,'/summary.txt'),'wt');
fprintf(fid, 'Skipped reads %i\n not INDEL %i\n Insertions %i\n Deletions
%i\n', [nSkips, notINDEL, NumIns, NumDels]); fclose(fid);
save(strcat(seqsFile,'/nSkips'),'nSkips'); save(strcat(seqsFile,'/notINDEL'),'notINDEL');
save(strcat(seqsFile,'/NumIns'),'NumIns'); save(strcat(seqsFile,'/NumDels'),'NumDels');
save(strcat(seqsFile,'/dels'),'dels');
C = dels;
fid = fopen(strcat(seqsFile, '/dels.txt'), 'wt'); fprintf(fid, '"%s"\n', C{:});
fclose(fid);
save(strcat(seqsFile,'/ins'),'ins'); C = ins;
fid = fopen(strcat(seqsFile, '/ins.txt'), 'wt'); fprintf(fid, '"%s"\n', C{:});
fclose(fid);

end
```

EXAMPLE 5: Cas9 variant sequences

[00346] The disclosure provides Cas9 variants, for example Cas9 proteins from one or more organisms, which may comprise one or more mutations (*e.g.*, to generate dCas9 or Cas9 nickase). In some embodiments, one or more of the amino acid residues, identified below by an asterek, of a Cas9 protein may be mutated. In some embodiments, the D10 and/or H840 residues of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, are mutated. In some embodiments, the D10 residue of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, is mutated to any amino acid residue, except for D. In some embodiments, the D10 residue of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, is mutated to an A. In some embodiments, the H840 residue of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding residue in any of the amino acid sequences provided in SEQ ID NOs: 11-260, is an H. In some embodiments, the H840 residue of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, is mutated to any amino acid residue, except for H. In some embodiments, the H840 residue of the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260, is mutated to an A. In some embodiments, the D10 residue of the amino acid sequence provided in SEQ ID NO: 10, or a

corresponding residue in any of the amino acid sequences provided in SEQ ID NOs: 11-260, is a D.

[00347] A number of Cas9 sequences from various species were aligned to determine whether corresponding homologous amino acid residues of D10 and H840 of SEQ ID NO: 10 or SEQ ID NO: 11 can be identified in other Cas9 proteins, allowing the generation of Cas9 variants with corresponding mutations of the homologous amino acid residues. The alignment was carried out using the NCBI Constraint-based Multiple Alignment Tool (COBALT(accessible at st-va.ncbi.nlm.nih.gov/tools/cobalt), with the following parameters. Alignment parameters: Gap penalties -11,-1; End-Gap penalties -5,-1. CDD Parameters: Use RPS BLAST on; Blast E-value 0.003; Find Conserved columns and Recompute on. Query Clustering Parameters: Use query clusters on; Word Size 4; Max cluster distance 0.8; Alphabet Regular.

[00348] An exemplary alignment of four Cas9 sequences is provided below. The Cas9 sequences in the alignment are: Sequence 1 (S1): SEQ ID NO: 11 | WP_010922251| gi 499224711 | type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogenes]; Sequence 2 (S2): SEQ ID NO: 12 | WP_039695303 | gi 746743737 | type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus gallolyticus]; Sequence 3 (S3): SEQ ID NO: 13 | WP_045635197 | gi 782887988 | type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus mitis]; Sequence 4 (S4): SEQ ID NO: 14 | 5AXW_A | gi 924443546 | *Staphylococcus aureus* Cas9. The HNH domain (bold and underlined) and the RuvC domain (boxed) are identified for each of the four sequences. Amino acid residues 10 and 840 in S1 and the homologous amino acids in the aligned sequences are identified with an asterisk following the respective amino acid residue.

```
S1 1    --MDKK-YSIGLD*IGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLI--GALLFDSG--ETAEATRLKRTARRRYT  73
S2 1    --MTKKNYSIGLD*IGTNSVGWAVITDDYKVPAKKMKVLGNTDKKYIKKNLL--GALLFDSG--ETAEATRLKRTARRRYT  74
S3 1    --M-KKGYSIGLD*IGTNSVGFAVITDDYKVPSKKMKVLGNTDKRFIKKNLI--GALLFDEG--TTAEARRLKRTARRRYT  73
S4 1    GSHMKRNYILGLD*IGITSVGYGII--DYET----------------RDVIDAGVRLFKEANVENNEGRRSKRGARRLKR  61

S1 74   RRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRL 153
S2 75   RRKNRLRYLQEIFANEIAKVDESFFQRLDESFLTDDDKTFDSHPIFGNKAEEDAYHQKFPTIYHLRKHLADSSEKADLRL 154
S3 74   RRKNRLRYLQEIFSEEMSKVDSSFFHRLDDSFLIPEDKRESKYPIFATLTEEKEYHKQFPTIYHLRKQLADSKEKTDLRL 153
S4 62   RRRHRIQRVKKLL--------------FDYNLLTD--------------------HSELSGINPYEARVKGLSQKLSEEE 107

S1 154  IYLALAHMIKFRGHFLIEGDLNPDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEK  233
S2 155  VYLALAHMIKFRGHFLIEGELNAENTDVQKIFADFVGVYNRTFDDSHLSEITVDVASILTEKISKSRRLENLIKYYPTEK  234
```

```
S3 154  IYLALAHMIKYRGHFLYEEAFDIKNNDIQKIFNEFISIYDNTFEGSSLSGQNAQVEAIFTDKISKSAKRERVLKLFPDEK  233
S4 108  FSAALLHLAKRRG----------------------VHNVNEVEEDT----------------------------------  131

S1 234  KNGLFGNLIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEIT 313
S2 235  KNTLFGNLIALALGLQPNFKTNFKLSEDAKLQFSKDTYEEDLEELLGKIGDDYADLFTSAKNLYDAILLSGILTVDDNST 314
S3 234  STGLFSEFLKLIVGNQADFKKHFDLEDKAPLQFSKDTYDEDLENLLGQIGDDFTDLFVSAKKLYDAILLSGILTVTDPST 313
S4 132  -----GNELS------------------TKEQISRN---------------------------------------------- 144

S1 314  KAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKM--DGTEELLV  391
S2 315  KAPLSASMIKRYVEHHEDLEKLKEFIKANKSELYHDIFKDKNKNGYAGYIENGVKQDEFYKYLKNILSKIKIDGSDYFLD  394
S3 314  KAPLSASMIERYENHQNDLAALKQFIKNNLPEKYDEVFSDQSKDGYAGYIDGKTTQETFYKYIKNLLSKF--EGTDYFLD  391
S4 145  ----SKALEEKYVAELQ-----------------------------------------------LERLKKDG------  165

S1 392  KLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEE  471
S2 395  KIEREDFLRKQRTFDNGSIPHQIHLQEMHAILRRQGDYYPFLKEKQDRIEKILTFRIPYYVGPLVRKDSRFAWAEYRSDE  474
S3 392  KIEREDFLRKQRTFDNGSIPHQIHLQEMNAILRRQGEYYPFLKDNKEKIEKILTFRIPYYVGPLARGNRDFAWLTRNSDE  471
S4 166  --EVRGSINRFKTSD--------YVKEAKQLLKVQKAYHQLDQSFIDTYIDLLETRRTYYEGP--GEGSPFGW------K  227

S1 472  TITPWNFEEVVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFLSGEQKKAIVDL 551
S2 475  KITPWNFDKVIDKEKSAEKFITRMTLNDLYLPEEKVLPKHSHVYETYAVYNELTKIKYVNEQGKE-SFFDSNMKQEIFDH 553
S3 472  AIRPWNFEEIVDKASSAEDFINKMTNYDLYLPEEKVLPKHSLLYETFAVYNELTKVKFIAEGLRDYQFLDSGQKKQIVNQ 551
S4 228  DIKEW---------------YEMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITRDENEK---LEYYEKFQIIEN 289

S1 552  LFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDR---FNASLGTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFED  628
S2 554  VFKENRKVTKEKLLNYLNKEFPEYRIKDLIGLDKENKSFNASLGTYHDLKKIL-DKAFLDDKVNEEVIEDIIKTLTLFED  632
S3 552  LFKENRKVTEKDIIHYLHN-VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDKEFMDDAKNEAILENIVHTLTIFED  627
S4 290  VFKQKKKPTLKQIAKEILVNEEDIKGYRVTSTGKPEF---TNLKVYHDIKDITARKEII---ENAELLDQIAKILTIYQS  363

S1 629  REMIEERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMQLIHDDSLTFKED 707
S2 633  KDMIHERLQKYSDIFTANQLKKLER-RHYTGWGRLSYKLINGIRNKENNKTILDYLIDDGSANRNFMQLINDDTLPFKQI 711
S3 628  REMIKQRLAQYDSLFDEKVIKALTR-RHYTGWGKLSAKLINGICDKQTGNTILDYLIDDGKINRNFMQLINDDGLSFKEI 706
S4 364  SEDIQEELTNLNSELTQEEIEQISNLKGYTGTHNLSLKAINLILDE------LWHTNDNQIAIFNRLKLVP--------- 428

S1 708  IQKAQVSGQGDSLHEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTT------QKGQKNSRERM 781
S2 712  IQKSQVVGDVDDIEAVVHDLPGSPAIKKGILQSVKIVDELVKVMG-GNPDNIVIEMARENQTT------NRGRSQSQQRL 784
S3 707  IQKAQVIGKTDDVKQVVQELSGSPAIKKGILQSIKIVDELVKVMG-HAPESIVIEMARENQTT------ARGKKNSQQRY 779
S4 429  -KKVDLSQQKEIPTTLVDDFILSPVVKRSFIQSIKVINAIIKKYG--LPNDIIIELAREKNSKDAQKMINEMQKRNRQTN 505

S1 782  KRIEEGIKELGSQIL-------KEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSD----YDVDH*IVPQSFLKDD 850
S2 785  KKLQNSLKELGSNILNEEKPSYIEDKVENSHLQNDQLFLYYIQNGKDMYTGDELDIDHLSD----YDIDH*IIPQAFIKDD 860
S3 780  KRIEDSLKILASGL---DSNILKENPTDNNQLQNDRLFLYYLQNGKDMYTGEALDINQLSS----YDIDH*IIPQAFIKDD 852
S4 506  ERIEEIIRTTGK---------------ENAKYLIEKIKLHDMQEGKCLYSLEAIPLEDLLNNPFNYEVDH*IIPRSVSFDN 570

S1 851  SIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDN-LTKAERGGL-SELD------KAGFIKRQLV 922
S2 861  SIDNRVLTSSAKNRGKSDDVPSLDIVRARKAEWVRLYKSGLISKRKFDN-LTKAERGGL-TEAD------KAGFIKRQLV 932
S3 853  SLDNRVLTSSKDNRGKSDNVPSIEVVQKRKAFWQQLLDSKLISERKFNN-LTKAERGGL-DERD------KVGFIKRQLV 924
S4 571  SFNNKVLVKQEEASKKGNRTPFQYLSSSDSKISYETFKKHILNLAKGKGRISKTKKEYLLEERDINRFSVQKDFINRNLV 650
```

```
S1 923  ETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYP 1002
S2 933  ETRQITKHVAQILDARFNTEHDENDKVIRDVKVITLKSNLVSQFRKDFEFYKVREINDYHHAHDAYLNAVVGTALLKKYP 1012
S3 925  ETRQITKHVAQILDARYNTEVNEKDKKNRTVKIITLKSNLVSNFRKEFRLYKVREINDYHHAHDAYLNAVVAKAILKKYP 1004
S4 651  DTRYATRGLMNLLRSYFRVN-------NLDVKVKSINGGFTSFLRRKWKFKKERNKGYKHHAEDALIIA---------- 712

S1 1003 KLESEFVYGDYKVYDVRKMIAKSEQ--EIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKG--- 1077
S2 1013 KLASEFVYGEYKKYDIRKFITNSSD-----KATAKYFFYSNLMNFFKTKVKYADGTVFERPIIETNAD-GEIAWNKQ--- 1083
S3 1005 KLEPEFVYGEYQKYDLKRYISRSKDPKEVEKATEKYFFYSNLLNFFKEEVHYADGTIVKRENIEYSKDTGEIAWNKE--- 1081
S4 713  --NADFIFKEWKKLDKAKKVMENQM-------------------------FEEKQAESMPEIETEQEYKEIFITPHQIK 764

S1 1078 -----RDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKD---WDPKKYGGFDSPTVAYSVLVVAKV 1149
S2 1084 -----IDFEKVRKVLSYPQVNIVKKVETQTGGFSKESILPKGDSDKLIPRKTKKVYWDTKKYGGFDSPTVAYSVFVVADV 1158
S3 1082 -----KDFAIIKKVLSLPQVNIVKKREVQTGGFSKESILPKGNSDKLIPRKTKDILLDTTKYGGFDSPVIAYSILLIADI 1156
S4 765  HIKDFKDYKYSHRVDKKPNRELINDTLYSTRKDDKGNTLIVNNLNGLYDKDNDKL----KKLIN-KSP----EKLLMYHH 835

S1 1150 EKGKSKKLKSVKELLGITIMERSSFEKNPI-DFLEAKG-----YKEVKKDLIIKLPKYSLFELENGRKRMLASAGELQKG 1223
S2 1159 EKGKAKKLKTVKELVGISIMERSFFEENPV-EFLENKG-----YHNIREDKLIKLPKYSLFEFEGGRRRLLASASELQKG 1232
S3 1157 EKGKAKKLKTVKTLVGITIMEKAAFEENPI-TFLENKG-----YHNVRKENILCLPKYSLFELENGRRRLLASAKELQKG 1230
S4 836  DPQTYQKLK--------LIMEQYGDEKNPLYKYYEETGNYLTKYSKKDNGPVIKKIKYYGNKLNAHLDITDDYPNSRNKV 907

S1 1224 NELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKH------ 1297
S2 1233 NEMVLPGYLVELLYHAHRADNF-----NSTEYLNYVSEHKKEFEKVLSCVEDFANLYVDVEKNLSKIRAVADSM------ 1301
S3 1231 NEIVLPVYLTTLLYHSKNVHKL-----DEPGHLEYIQKHRNEFKDLLNLVSEFSQKYVLADANLEKIKSLYADN------ 1299
S4 908  VKLSLKPYRFD-VYLDNGVYKFV-----TVKNLDVIK--KENYYEVNSKAYEEAKKLKKISNQAEFIASFYNNDLIKING 979

S1 1298 RDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIHQSIT--------GLYETRI----DLSQL 1365
S2 1302 DNFSIEEISNSFINLLTLTALGAPADFNFLGEKIPRKRYTSTKECLNATLIHQSIT--------GLYETRI----DLSKL 1369
S3 1300 EQADIEILANSFINLLTFTALGAPAAFKFFGKDIDRKRYTTVSEILNATLIHQSIT--------GLYETWI----DLSKL 1367
S4 980  ELYRVIGVNNDLLNRIEVNMIDITYR-EYLENMNDKRPPRIIKTIASKT---QSIKKYSTDILGNLYEVKSKKHPQIIKK 1055

S1 1366 GGD  1368
S2 1370 GEE  1372
S3 1368 GED  1370
S4 1056 G--  1056
```

[00349] The alignment demonstrates that amino acid sequences and amino acid residues that are homologous to a reference Cas9 amino acid sequence or amino acid residue can be identified across Cas9 sequence variants, including, but not limited to Cas9 sequences from different species, by identifying the amino acid sequence or residue that aligns with the reference sequence or the reference residue using alignment programs and algorithms known in the art. This disclosure provides Cas9 variants in which one or more of the amino acid residues identified by an asterisk in SEQ ID NOs: 11-14 (*e.g.,* S1, S2, S3, and S4, respectively) are

mutated as described herein. The residues D10 and H840 in Cas9 of SEQ ID NO: 10 that correspond to the residues identified in SEQ ID NOs: 11-14 by an asterisk are referred to herein as "homologous" or "corresponding" residues. Such homologous residues can be identified by sequence alignment, *e.g.*, as described above, and by identifying the sequence or residue that aligns with the reference sequence or residue. Similarly, mutations in Cas9 sequences that correspond to mutations identified in SEQ ID NO: 10 herein, *e.g.*, mutations of residues 10, and 840 in SEQ ID NO: 10, are referred to herein as "homologous" or "corresponding" mutations. For example, the mutations corresponding to the D10A mutation in SEQ ID NO: 10 or S1 (SEQ ID NO: 11) for the four aligned sequences above are D11A for S2, D10A for S3, and D13A for S4; the corresponding mutations for H840A in SEQ ID NO: 10 or S1 (SEQ ID NO: 11) are H850A for S2, H842A for S3, and H560A for S4.

[00350] A total of 250 Cas9 sequences (SEQ ID NOs: 11-260) from different species were aligned using the same algorithm and alignment parameters outlined above. Amino acid residues homologous to residues 10, and 840 of SEQ ID NO: 10 were identified in the same manner as outlined above. The alignments are provided below. The HNH domain (bold and underlined) and the RuvC domain (boxed) are identified for each of the four sequences. Single residues corresponding to amino acid residues 10, and 840 in SEQ ID NO: 10 are boxed in SEQ ID NO: 11 in the alignments, allowing for the identification of the corresponding amino acid residues in the aligned sequences.

WP_010922251.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_039695303.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus gallol
WP_045635197.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus mitis]
5AXW_A	Cas9, Chain A, Crystal Structure [Staphylococcus Aureus] SEQ ID
WP_009880683.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_010922251.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_011054416.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_011284745.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_011285506.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_011527619.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_012560673.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_014407541.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_020905136.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_023080005.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
WP_023610282.1	type II CRISPR RNA-guided endonuclease Cas9 [Streptococcus pyogen
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WP_003727705  1        MKNP-YTIGLDIGTNSVGWAVLTDQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
WP_003730785  1        MKNP-YTIGLDIGTNSVGWAVLTDQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
WP_003733029  1        MKKP-YTIGLDIGTNSVGWAVLTDQYDLVKRKMKISGDSEKKQIKKNFWGVRLFEKGET
WP_003739838  1        MKNP-YTIGLDIGTNSVGWAVLTDQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDEGET
WP_014601172  1        MKNP-YTIGLDIGTNSVGWAVLTNQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
WP_023548323  1        MKNP-YTIGLDIGTNSVGWAVLTNQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
WP_031665337  1        MKNP-YTIGLDIGTNSVGWAVLTDQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
WP_031669209  1        MKKP-YTIGLDIGTNSVGWAVLTDQYDLVKRKMKISGDSEKKQIKKNFWGVRLFEKGET
WP_033920898  1        MKNP-YTIGLDIGTNSVGWAVLTNQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
AKI42028      1        MKNP-YTIGLDIGTNSVGWAVLTNQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
AKI50529      1        MKNP-YTIGLDIGTNSVGWAVLTNQYDLVKRKMKVAGNSDKKQIKKNFWGVRLFDDGQT
EFR83390               ------------------------------------------------------------
WP_046323366  1        MKKP-YTIGLDIGTNSVGWAALTDQYDLVKRKMKVAGNSEKKQIKKNLWGVRLVDEGKT
AKE81011      1        ADKK-YSIGLDIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGET
CUO82355      1        I-VD-YCIGLDLGTGSVGWAVVDMNHRLMKRN-----------GKHLWGSRLFSNAET
WP_033162887  1        KDIR-YSIGLDIGTNSVGWAVMDEHYELLKKG-----------NHHMWGSRLFDAAEP
AGZ01981      1        ADKK-YSIGLDIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGET
AKA60242      1        MDKK-YSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGET
AKS40380      1        MDKK-YSIGLDIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGET
4UN5_B        1        MDKK-YSIGLAIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGET
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WP_010922251  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_039695303  75    RRKNRLRYLQEIFANEIAKVDESFFQRLDE-SFLT--DDDKT---F   DSHPIFGNKA-EED
WP_045635197  74    RRKNRLRYLQEIFSEEMSKVDSSFFHRLDD-SFLI--PEDKR---E   SKYPIFATLT-EEK
5AXW_A        62    RRRHRIQRVKKLLFD---------YNLLTDhSELS----------G   --NPYEARVK----
WP_009880683          ----------------------------------------------   --------------
WP_010922251  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_011054416  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_011284745  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_011285506  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_011527619  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_012560673  74    RRKNRICYLQEIFSNEIAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_014407541  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_020905136  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_023080005  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_023610282  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_030125963  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_030126706  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_031488318  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_032460140  74    RRKNRICYLQEIFSNEIAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_032461047  74    RRKNRICYLQEIFSNEIAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_032462016  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_032462936  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_032464890  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_033888930          ----------------------------------------------   --------------
WP_038431314  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_038432938  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_038434062  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
BAQ51233      1     ----------------MAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
KGE60162              ----------------------------------------------   --------------
KGE60856              ----------------------------------------------   --------------
WP_002989955  74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_003030002  74    RRRNRLRYLQEIFAEEMNKVDENFFQRLDD-SFLV--DEDKR---G   ERHPIFGNIA-AEV
WP_003065552  75    RRKNRLRYLQEIFAEEMTKVDESFFQRLDE-SFLRwdDDNKK---L   GRYPIFGNKA-DVV
WP_001040076  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G   SKYPIFATLQ-EEK
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WP_001040078  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFPTIYHLRKELA  143
WP_001040080  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040081  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040083  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040085  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040087  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040088  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040089  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040090  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040091  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040092  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_001040094  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040095  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040096  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040097  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040098  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040099  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040100  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_001040104  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYXIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040105  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_001040106  74  CRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_001040107  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_001040108  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_001040109  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_001040110  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_015058523  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_017643650  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_017647151  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_017648376  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_017649527  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFPTIYHLRKELA  143
WP_017771611  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_017771984  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
CFQ25032      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
CFV16040      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
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KLJ37842          74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
KLJ72361          74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
KLL20707          74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
KLL42645          74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_047207273      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_047209694      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKYYHEKFPTIYHLRKELA  143
WP_050198062      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_050201642      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_050204027      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EDDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
WP_050881965      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
WP_050886065      74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
AHN30376          74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATMQ-EEKDYHEKFPTIYHLRKELA  143
EAO78426          74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFSTIYHLRKELA  143
CCW42055          74  RRRNRILYLQEIFAEKMSKVDDSFFHRLED-SFLV--EEDKR---G  SKYPIFATLQ-EEKDYHEKFPTIYHLRKELA  143
WP_003041502      74  RRRNRLRYLQEIFAEEMMQVDESFFQRLDD-SFLV--DEDKR---G  ERHPIFGNIA-AEVKYHDEFPTIYHLRKHLA  143
WP_037593752      75  RRKNRLRYLQEIFTEEMNKVDENFFQRLDD-SFLV--EEDKQ---G  SKYPIFGTLK-EEKEYHKKFKTIYHLREELA  144
WP_049516684      75  RRRNRLRYLQEIFAEEMMQVDESFFQRLDD-SFLV--EEDKR---G  SRYPIFGNIA-AEVKYHDDFPTIYHLRKHLV  144
GAD46167          74  RRKNRLRYLQEITTEEMNKVDENFFQRLDD-SFLV--EEDKQ---G  SKYPIFGTLK-EEKEYHKKFKTIYHLREELA  143
WP_018363470      75  RRKNRLRYLQDITTEEMAKVDDSFFQRLDE-SFLT--DNDKN---F  DSHPIFGNKA-EEDAYHQKFPTIYHLRKHLA  144
WP_003043819      74  RRKNRIRYLQEIFANEMAKLDDSFFQRLEE-SFLV--EEDKK---N  ERHPIFGNLA-DEVAYHRNYPTIYHLRKKLA  143
WP_006269658      74  RRKNRLRYLQEIFTGEMNKVDENFFQRLDD-SFLV--DEDKR---G  EHHPIFGNIA-AEVKYHDDFPTIYHLRRHLA  143
WP_048800889      74  RRKNRLRYLQEIFIEEMNKVDENFFQRLDD-SFLV--TEDKR---G  SKYPIFGTLK-EEKEYYKEFETIYHLRKRLA  143
WP_012767106      74  RRKNRIRYLQEIFSSEMSKVDDSFFHRLEE-SFLV--EEDKK---H  ERHPIFGNIV-DEVAYHEKYPTIYHLRKKLA  143
WP_014612333      74  RRKNRIRYLQEIFSSEMSKVDDSFFHRLEE-SFLV--EEDKK---H  ERHPIFGNIV-DEVAYHEKYPTIYHLRKKLA  143
WP_015017095      74  RRKNRIRYLQEIFSSEMSKVDDSFFHRLEE-SFLV--EEDKK---H  ERHPIFGNIV-DEVAYHEKYPTIYHLRKKLA  143
WP_015057649      74  RRKNRIRYLQEIFSSEMSKVDDSFFHRLEE-SFLV--EEDKK---H  ERHPIFGNIV-DEVAYHEKYPTIYHLRKKLA  143
WP_048327215      74  RRKNRIRYLQEIFSSEMSKVDDSFFHRLEE-SFLV--EEDKK---H  ERHPIFGNIV-DEVAYHEKYPTIYHLRKKLA  143
WP_049519324      74  RRKNRIRYLQEIFSSEMSKVDDSFFHRLEE-SFLV--EEDKK---H  ERHPIFGNIV-DEVAYHEKYPTIYHLRKKLA  143
WP_012515931      74  RRKNRLRYLKEIFTEEMAKVDDGFFQRLED-SFYV--LEDKE---G  NKHPIFANLA-DEVAYHKKYPTIYHLRKELV  143
WP_021320964      74  RRKNRLRYLKEIFTEEMAKVDDGFFQRLED-SFYV--LEDKE---G  NKHPIFANLA-DEVAYHKKYPTIYHLRKELV  143
WP_037581760      74  RRKNRLRFLKEIFTEEMAKVDDGFFQRLED-SFYV--LEDKE---G  NKHPIFANLA-DEVAYHKKYPTIYHLRKELV  143
WP_004232481      74  RRKNRLRYLQEIFAKEMAKVDESFFQRLEE-SFLT--DDDKT---F  DSHPIFGNKA-EEDTYHQEFPTIYHLRKHLA  143
WP_009854540      75  RRKNRLRYLQEIFAEEMTKVDESFFYRLDE-SFLT--TDEKD---F  ERHPIFGNKA-EEDAYHQKFPTIYHLRNYLA  144
WP_012962174      75  RRKNRLRYLQEIFAEEMAKVDESFFYRLDE-SFLT--TDDKD---F  ERHPIFGNKA-DEIKYHQEFPTIYHLRKHLA  144
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WP_039695303  75    RRKNRLRYLQEIFANEIAKVDESFFQRLDE-SFLT--DDDKT---F   DSHPIFGNKA-EED
WP_014334983  74    RRKNRLRYLQEIFAKEMTKVDESFFQRLEE-SFLT--DDDKT---F   DSHPIFGNKA-EED
WP_003099269  74    RRKYRIKELQKIFSSEMNELDIAFFPRLSE-SFLV--SDDKE---F   ENHPIFGNLK-DEI
AHY15608      74    RRKYRIKELQKIFSSEMNELDIAFFPRLSE-SFLV--SDDKE---F   ENHPIFGNLK-DEI
AHY17476      74    RRKYRIKELQKIFSSEMNELDIAFFPRLSE-SFLV--SDDKE---F   ENHPIFGNLK-DEI
ESR09100            ----------------------------------------------   --------------
AGM98575      74    RRKYRIKELQKIFSSEMNELDIAFFPRLSE-SFLV--SDDKE---F   ENHPIFGNLK-DEI
ALF27331      74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_018372492  74    RRRNRIRYLQHIFAEEMNRADENFFHRLKE-SFFV--EEDKT---Y   SKYPIFGTLE-EEK
WP_045618028  75    RRKNRLRYLQEIFTEEMSKVDISFFHRLDD-SFLV--PEDKR---G   SKYPIFATLE-EEK
WP_045635197  74    RRKNRLRYLQEIFSEEMSKVDSSFFHRLDD-SFLI--PEDKR---E   SKYPIFATLT-EEK
WP_002263549  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002263887  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002264920  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLDE-SFLT--DDDKN---F   DSYPIFGNKA-EED
WP_002269043  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002269448  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002271977  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002272766  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002273241  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002275430  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002276448  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002277050  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLDE-SFLT--DDDKN---F   DSHPIFGNKA-EED
WP_002277364  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002279025  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-FFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002279859  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLDE-SFLT--DDDKN---F   DSHPIFGNKA-EED
WP_002280230  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002281696  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002282247  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLDE-SFLT--DDDKN---F   DSHPIFGNKA-EED
WP_002282906  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002283846  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002287255  74    RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002288990  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002289641  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
WP_002290427  74    RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G   ERHPIFGNLE-EEV
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WP_002295753  74  RRRNRILYLQEIFSEEMGKVNDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_002296423  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_002304487  74  RRRNRILYLQEIFAEEMMQVDESFFQRLDD-SFLV--EEDKR---G  SRYPIFGTLK-EEKKYHKEFKTIYHLREKLA  143
WP_002305844  74  RRRNRILYLQEIFSEEMDKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_002307203  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_002310390  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_002352408  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_012997688  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_014677909  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_019312892  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_019313659  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_019314093  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_019315370  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ECHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_019803776  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_019805234  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_024783594  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_024784288  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLDE-SFLT--DDDKN---F  DSHPIFGNKA-EEDAYHQKFPTIYHLRKHLA  143
WP_024784666  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_024784894  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
WP_024786433  74  RRRNRILYLQEIFAEEMNKVDDSFFHRLDE-SFLT--DDDKN---F  DSHPIFGNKA-EEDAYHQKFPTIYHLRKHLA  143
WP_049473442  74  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYYENFPTIYHLRQYLA  143
WP_049474547  74  RRRNRILYLQEIFSEEMGKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  143
EMC03581      67  RRRNRILYLQEIFAEEMSKVDDSFFHRLED-SFLV--TEDKR---G  ERHPIFGNLE-EEVKYHENFPTIYHLRQYLA  136
WP_000428612  75  RRKNRLRYLQEIFAEEMSKVDSSFFHRLDD-SFLI--PEDKK---G  SKYPIFATLI-EEKEYHKQFPTIYHLRKQLA  144
WP_000428613  75  RRKNRLRYLQEIFAEEMSKVDSSFFHRLDD-SFLI--PEDKR---G  SKYPIFATLA-EEKEYHKQFPTIYHLRKQLA  144
WP_049523028  74  RRRNRILYLQEIFAAEMNKVDESFFHRLDD-SFLV--PEDKR---G  SKYPIFGTLE-EEKEYHKQFPTIYYLRKILA  143
WP_003107102  43  RRINRIKYLQSIFDDEMSKIDSAFFQRIKD-SFLV--PDDKN---D  DRHPIFGNIK-DEVDYHKNYPTIYHLRKKLA  112
WP_054279288  76  RRKNRLCYLRDIFESEMHTIDKHFFLRLED-SFLH--KSDKR---Y  EAHPIFGTLQ-EEKAYHDNYPTIYHLRKALA  145
WP_049531101  75  RRKNRLRYLQEIFSEEISKVDNSFFHRLDD-SFLV--PEDKR---G  SKYPIFATLT-EEKEYYKQFPTIYHLRKQLA  144
WP_049538452  75  RRKNRLRYLQEIFAEEMNKVDSSFFHRLDD-SFLV--PEDKR---G  SKYPIFATLA-EEKEYHKNFPTIYHLRKQLA  144
WP_049549711  75  RRKNRLRYLQEIFSGEMSKVDSSFFHRLDD-SFLV--PEDKR---G  SKYPIFATLV-EEKEYHKQFPTIYHLRKQLA  144
WP_007896501  76  RRRYRLCQLQNIFATEMVKVDDTFFQRLSE-SFFY--YQDKA---F  DKHPIFGNSK-EERAYHKTYPTIYHLRKDLA  145
EFR44625      28  RRRYRLCQLQNIFATEMVKVDDTFFQRLSE-SFFY--YQDKA---F  DKHPIFGNSK-EERAYHKTYPTIYHLRKDLA  97
WP_002897477  74  RRRNRILYLQEIFTESMNEIDESFFHRLDD-SFLV--PEDKR---G  SKYPIFATLQ-EEKEYHKQFPTIYHLRKQLA  143
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WP_002906454  74    RRKNRLRYLQEIFSEEISKLDSSFFHRLDD-SFLV--PEDKR---G   SKYPIFATLE-EEK
WP_009729476  75    RRKNRLRYLQEIFSEEIGKVDSSFFHRLDD-SFLI--PEDKR---G   SKYPIFATLA-EEK
CQR24647      74    RRRNRILYLQDIFSPELNQVDESFLHRLDD-SFLVa--EDKR---G   ERHVIFGNIA-DEV
WP_000066813  75    RRKNRLRYLQEIFSQEISKVDSSFFHRLDD-FFLV--PEDKR---G   SKYPIFATLV-EEK
WP_009754323  75    RRKNRLRYLQEIFAEEMSKVDSSFFHRLDD-SFLV--PEDKS---G   SKYPIFATLA-EEK
WP_044674937  74    RRRNRILYLQEIFAEEINKIDDSFFQRLDD-SFLIv--EDKQ---G   SKHPIFGTLQ-EEK
WP_044676715  74    RRRNRILYLQEIFAEEINKIDDSFFQRLDD-SFLIv--EDKQ---G   SKHPIFGTLQ-EEK
WP_044680361  74    RRRNRILYLQEIFAEEINKIDDSFFQRLDD-SFLIv--EDKQ---G   SKHPIFGTLQ-EEK
WP_044681799  74    RRRNRILYLQEIFAEEINKIDDSFFQRLDD-SFLIv--EDKQ---G   SKHPIFGTLQ-EEK
WP_049533112  74    RRRNRLRYLQEIFAEEMNKVDENFFQRLDD-SFLV--DEDKR---G   ERHPIFGNIA-AEV
WP_029090905  28    HRKFRLRLLEDMFEKEILSKDPSFFIRLKE-AFLSpkDEQKQ---F   ----LFNDKDyTDA
WP_006506696  61    KRRERIRLLRAILQDMVLEKDPTFFIRLEHtSFLD--EEDKAkylG   DNYNLFIDEDfNDY
AIT42264      74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_034440723  73    RRRFRIRELQKIFDKSMGEVDSNFFHRLDE-SFLV--EEDKE---Y   SKYPIFSNEK-EDK
AKQ21048      74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_004636532  74    RRRNRLIYLQDIFQQPMLAIDENFFHRLDD-SFFV--PDDKS---Y   DRHPIFGSLE-EEV
WP_002364836  74    RRRNRLRYLQAFFEEAMTDLDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_016631044  25    RRRNRLRYLQAFFEEAMTDLDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
EMS75795            ---------------------------------------------   --------------
WP_002373311  74    RRRNRLRYLQAFFEEAMTDLDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_002378009  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_002407324  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_002413717  74    RRRNRLRYLQAFFEEAMTDLDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_010775580  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_010818269  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_010824395  74    RRRNRLRYLQAFFEEAMTDLDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_016622645  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_033624816  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_033625576  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_033789179  74    RRRNRLRYLQAFFEEAMTALDENFFARLQE-SFLV--PEDKK---W   HRHPIFAKLE-DEV
WP_002310644  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
WP_002312694  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--PDEKK---Q   SRHPVFATIK-QEK
WP_002314015  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
WP_002320716  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
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WP_002330729  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
WP_002335161  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
WP_002345439  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
WP_034867970  74    RRKYRLSKLQDLFAEELCKQDDCFFVRLEE-SFLV--PEEKQ---Y   KPASIFPTLE-EEK
WP_047937432  74    RRRQRILELQKIFAPEILKIDEHFFARLNE-SFLV--LDEKK---Q   SRHPVFATIK-QEK
WP_010720994  74    RRKYRLSKLQDLFAEELCKQDDCFFVRLEE-SFLV--PEEKQ---Y   KPASIFPTLE-EEK
WP_010737004  74    RRKYRLSKLQDLFAEELCKQDDCFFVRLEE-SFLV--PEEKQ---Y   KPASIFPTLE-EEK
WP_034700478  74    RRKYRLSKLQDLFAEELCKQDDCFFVRLEE-SFLV--PEEKQ---Y   KPASIFPTLE-EEK
WP_007209003  74    RRKNRICYLQEIFQPEMNHLDNNFFYRLNE-SFLVa--DDAK---Y   DKHPIFGTLD-EEI
WP_023519017  74    RRRQRVLALQDIFAEEIHKKDPNFFARLEE-GDRV--EADKR---F   AKFPVFATLS-EEK
WP_010770040  74    RRRNRICRLQDLFTEEMNQVDANFFHRLQE-SFLV--PDEKE---F   ERHAIFGKME-EEV
WP_048604708  74    RRRQRISYLQTFFQEEMNRIDPNFFNRLDE-SFLI--EEDKL---S   ERHPIFGTIE-EEV
WP_010750235  74    RRKYRILELQKIFSEEILKKDSHFFARLDE-SFLI--PEDKQ---Y   ARFPIFPTLL-EEK
AII16583      113   RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
WP_029073316  66    KRRERIRLLRGIMEDMVLDVDPTFFIRLANvSFLD--QEDKKdylK   SNYNLFIDKDfNDK
WP_031589969  66    KRRERIRLLREIMEDMVLDVDPTFFIRLANvSFLD--QEDKKdylK   SNYNLFIDKDfNDK
KDA45870      75    RRKNRLRYLQEIFAPALAKVDPNFFYRLEE-SSLVa--EDKK---Y   DVYPIFGKRE-EEL
WP_039099354  62    RRKWRLGLLREIFEPYITPVDDTFFLRKKQ-SNLS--PKDQR---K   -QTSLFNDRT--DR
AKP02966      65    RRKNRINWLNEIFSEELANTDPSFLIRLQN-SWVSkkDPDRK---R   DKYNLFIDNPyTDK
WP_010991369  74    RRRNRISYLQGIFAEEMSKTDANFFCRLSD-SFYV--DNEKR---N   SRHPFFATIE-EEV
WP_033838504  74    RRRNRISYLQGIFAEEMSKTDANFFCRLSD-SFYV--DNEKR---N   SRHPFFATIE-EEV
EHN60060      77    RRRNRISYLQGIFAEEMSKTDANFFCRLSD-SFYV--DNEKR---N   SRHPFFATIE-EEV
EFR89594            ---------------------------------------------   --------------
WP_038409211  74    RRRNRIAYLQEIFAAEMAEVDANFFYRLED-SFYI--ESEKR---H   SRHPFFATIE-EEV
EFR95520            ---------------------------------------------   --------------
WP_003723650  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_003727705  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_003730785  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_003733029  74    RRRNRISYLQEIFAIQMNEVDDNFFNRLKE-SFYA--ESDKK---Y   NRHPFFGTVE-EEV
WP_003739838  74    RRRNRISYLQEIFALEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_014601172  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_023548323  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_031665337  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
WP_031669209  74    RRRNRISYLQEIFAIQMNEVDDNFFNRLKE-SFYA--ESDKK---Y   NRHPFFGTVE-EEV
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WP_033920898  74    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
AKI42028      77    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
AKI50529      77    RRRNRISYLQEIFAVEMANIDANFFCRLND-SFYV--DSEKR---N   SRHPFFATIE-EEV
EFR83390            ------------------------------------------   --------------
WP_046323366  74    RRRNRISYLQEIFTAEMFEVDANFFYRLED-SFYI--ESEKR---Q   SRHPFFATIE-EEV
AKE81011      90    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
CUO82355      65    KRRERIRLLRAILQDMVLEKDPTFFIRLEHtSFLD--EEDKAkylG   DNYNLFIDEDfNDY
WP_033162887  66    KRRERIRLLRDLLGDMVMEVDPTFFIRLLNvSFLD--EEDKQknlG   DNYNLFIEKDfNDK
AGZ01981      107   RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
AKA60242      74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
AKS40380      74    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
4UN5_B        78    RRKNRICYLQEIFSNEMAKVDDSFFHRLEE-SFLV--EEDKK---H   ERHPIFGNIV-DEV
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WP_010922251  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_039695303  145   DSSEKADLRLVYLALAHMIKFRGHFLIEGE-LNAENTDVQKI--FADFVGVYNRT--FDDS-H
WP_045635197  144   DSKEKTDLRLIYLALAHMIKYRGHFLYEEA-FDIKNNDIQKI--FNEFISIYDNT--FEGS-S
5AXW_A        105   EEEFSA-------ALLHLAKRRG---VHNV------NEVE------------EDT----GN--
WP_009880683        ------------------------------------------------------------
WP_010922251  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_011054416  144   DSTDKVDLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_011284745  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_011285506  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_011527619  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_012560673  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_014407541  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQIYNQL--FEEN--
WP_020905136  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_023080005  144   DSTDKVDLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_023610282  144   DSTDKVDLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_030125963  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_030126706  144   DSTDKVDLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_031488318  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_032460140  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_032461047  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGG-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_032462016  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_032462936  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_032464890  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_033888930  1     ---------------------------------PDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_038431314  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_038432938  144   DSTDKVDLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_038434062  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
BAQ51233      55    DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
KGE60162            ------------------------------------------------------------
KGE60856            ------------------------------------------------------------
WP_002989955  144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_003030002  144   DISQKADLRLVYLALAHMIKFRGHFLIEGQ-LKAENTNVQAL--FKDFVEVYDKT--VEES-H
WP_003065552  147   DSSEKADLRLVYLALAHMIKFRGHFLIEGE-LNAENTDVQKI--FADFVGVYDRT--FDDS-H
WP_001040076  144   DKKEKADLRLVYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D
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WP_001040078	144	DKQEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-H	LLSQNVDVE---AI	212
WP_001040080	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNIT--FENN-D	LLSQNVDVE---AI	212
WP_001040081	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTI--FENN-D	LLSQNVDVE---AI	212
WP_001040083	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040085	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040087	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040088	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040089	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040090	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040091	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040092	144	DKKEKADLRLVYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTS--FENN-H	LLSQNVDVE---AI	212
WP_001040094	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040095	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040096	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040097	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040098	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040099	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040100	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_001040104	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040105	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_001040106	144	DKKEKANLRLVYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_001040107	144	DKKEKADLRLVYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_001040108	144	DKKEKADLRLVYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_001040109	144	DKKEKANLRLVYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_001040110	144	DKKEKANLRLVYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_015058523	144	DKKEKADLRLVYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTS--FENN-H	LLSQNVDVE---AI	212
WP_017643650	144	DKKEKADLRLIYLALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D	LLSQNVDVE---AI	212
WP_017647151	144	DKKEKADLRLFYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDIE---GI	212
WP_017648376	144	DKKEKADLRLFYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_017649527	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
WP_017771611	144	DKKEKADLRLVYLALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H	LLSQNIDVE---GI	212
WP_017771984	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
CFQ25032	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212
CFV16040	144	DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D	LLSQNVDVE---AI	212

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KLJ37842          144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
KLJ72361          144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
KLL20707          144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
KLL42645          144  DKKEKANLRLVYILALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H  LLSQNIDVE---GI  212
WP_047207273      144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
WP_047209694      144  DKKEKADLRLIYLALAHIIKFRGHFLIEDDsFDVRNTDIQKQ--YQAFLEIFDTT--FENN-D  LLSQNVDVE---AI  212
WP_050198062      144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
WP_050201642      144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
WP_050204027      144  DKKEKANLRLVYILALAHIIKFRGHFLIEDDsFDVRNTDIQRQ--YQAFLEIFDTT--FENN-H  LLSQNIDVE---GI  212
WP_050881965      144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
WP_050886065      144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
AHN30376          144  DKKEKADLRLVYILALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTS--FENN-H  LLSQNVDVE---AI  212
EAO78426          144  DKKEKADLRLIYIALAHIIKFRGHFLIEDDsFDVRNTDISKQ--YQDFLEIFNTT--FENN-D  LLSQNVDVE---AI  212
CCW42055          144  DKKEKADLRLVYILALAHIIKFRGHFLIEDDrFDVRNTDIQKQ--YQAFLEIFDTT--FENN-H  LLSQNVDVE---AI  212
WP_003041502      144  DISQKADLRLVYILALAHMIKFRGHFLIEGQ-LKAENTNVQAL--FKDFVEVYDKT--VEES-H  LSEITVDAL---SI  211
WP_037593752      145  NSKEKADLRLVYILALAHMIKFRGHFLYEGD-LKAENTDVQAL--FKDFVEEYDKT--IEES-H  LSEITVDAL---SI  212
WP_049516684      145  DISQKADLRLVYILALAHMIKFRGHFLIEGQ-LKAENTNVQAL--FKDFVEVYDKT--VEES-H  LSEMTVDAL---SI  212
GAD46167          144  NSKEKADLRLVYILALAHMIKFRGHFLYEGD-LKAENTDVQAL--FKDFVEEYDKT--IEES-H  LSEITVDAL---SI  211
WP_018363470      145  DSTEKADLRLVYILALAHMIKFRGHFLIEGE-LNAENTDVQKL--FTDFVGVYDRT--FDDS-H  LSEITVDAA---SI  212
WP_003043819      144  DSPEKADLRLIYILALAHIIKFRGHFLIEGK-LNAENSDVAKL--FYQLIQTYNQL--FEES--  LDEIEVDAK---GI  211
WP_006269658      144  DTSKKADLRLVYILALAHMIKFRGHFLYEGD-LKAENTDVQAL--FKDFVEEYDKT--IEES-H  LSEITVDAL---SI  211
WP_048800889      144  DSTGKVDLRLVYILALAHMIKFRGHFLIEGQ-LKAENTDVQTL--FNDFVEVYDKT--IEES-H  LAEITVDAL---SI  211
WP_012767106      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDMDKL--FIQLVQTYNQL--FEEN--  INASRVDAK---AI  211
WP_014612333      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEK--  INASGVDAK---AI  211
WP_015017095      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDMDKL--FIQLVQTYNQL--FEEN--  INASRVDAK---AI  211
WP_015057649      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDMDKL--FIQLVQTYNQL--FEEN--  INASRVDAK---AI  211
WP_048327215      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDMDKL--FIQLVQTYNQL--FEEN--  INASRVDAK---AI  211
WP_049519324      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--  INASRVDAK---AI  211
WP_012515931      144  DNPQKADLRLIYLAVAHIIKFRGHFLIEGT-LSSKNNNLQKS--FDHLVDTYNLL--FEEQ--  LLTEGINAK---EL  211
WP_021320964      144  DNPQKADLRLIYLAVAHIIKFRGHFLIEGT-LSSKNNNLQKS--FDHLVDTYNLL--FEEQ--  LLTEGINAK---EL  211
WP_037581760      144  DNPQKADLRLIYLAVAHIIKFRGHFLIEGT-LSSKNNNLQKS--FDHLVDTYNLL--FEEQ--  LLTEGINAK---EL  211
WP_004232481      144  DSPEKVDLRLVYILALAHMIKFRGHFLIEGQ-LNAENTDVQKI--FADFVGVYDRT--FDDS-H  LSEITVDAA---SI  211
WP_009854540      145  DSSEKADLRLVYILALAHMIKYRGHFLIEGK-LNAENTDVQKL--FTDFVGVYDRT--FDDS-H  LSEITVDVA---ST  212
WP_012962174      145  DSHEKADLRLIYLALAHMIKFRGHFLIEGE-LNAENTDVQKL--FEAFVEVYDRT--FDDS-N  LSEITVDAS---SI  212
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WP_039695303 145 DSSEKADLRLVYLALAHMIKFRGHFLIEGE-LNAENTDVQKI--FADFVGVYNRT--FDDS-H LSEITVDVA---SI 212
WP_014334983 144 DSQEKADLRLVYLALAHMIKYRGHFLIEGE-LNAENTDVQKL--FNVFVETYDKI--VDES-H LSEIEVDAS---SI 211
WP_003099269 144 DSDQKADLRLIYLALAHIIKFRGHFLIEGN-LDSENTDVHVL--FLNLVNIYNNL--FEED-- VETASIDAE---KI 211
AHY15608     144 DSDQKADLRLIYLALAHIIKFRGHFLIEGN-LDSENTDVHVL--FLNLVNIYNNL--FEED-- VETASIDAE---KI 211
AHY17476     144 DSDQKADLRLIYLALAHIIKFRGHFLIEGN-LDSENTDVHVL--FLNLVNIYNNL--FEED-- VETASIDAE---KI 211
ESR09100     ---------------------------------------------------------------- ---------- 
AGM98575     144 DSDQKADLRLIYLALAHIIKFRGHFLIEGN-LDSENTDVHVL--FLNLVNIYNNL--FEED-- VETASIDAE---KI 211
ALF27331     144 DNPEKTDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_018372492 144 DTPDKMDIRLIYLALAHIIKYRGHFLIEGD-LDIENIGIQDS--FKSFIEEYNTQ--FGTK-- -LDSTTKVE---AI 209
WP_045618028 145 DSKEKADFRLIYLALAHIIKYRGHFLYEES-FDIKNNDIQKI--FNEFISIYDNT--FEGS-S LNGQNAQVE---AI 212
WP_045635197 144 DSKEKTDLRLIYLALAHMIKYRGHFLYEEA-FDIKNNDIQKI--FNEFISIYDNT--FEGS-S LSGQNAQVE---AI 211
WP_002263549 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002263887 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002264920 144 DSTEKADLRLVYLALAHMIKFRGHFLIEGE-LNAENTDVQKL--FADFVGVYDRT--FDDS-H LSEITVDAS---SI 211
WP_002269043 144 DNPEKTDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002269448 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002271977 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002272766 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002273241 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002275430 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002276448 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002277050 144 DSTEKADLRLVYLALAHMIKFRGHFLIEGE-LNAENTDVQKL--FADFVGVYDRT--FDDS-H LSEITVDAS---SI 211
WP_002277364 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002279025 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
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WP_002280230 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002281696 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
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WP_002282906 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002283846 144 DNPEKTDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002287255 144 DNPEKVDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002288990 144 DNPEKTDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002289641 144 DNPEKTDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
WP_002290427 144 DNPEKTDLRLVYLALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S LQEQNVQVE---EI 211
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WP_002296423  144  DNPEKTDLRLVYILALAHIIKFGGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_002304487  144  NSTEKADLRLVYLSLAHMIKFRGHFLIEGQ-LKAENTNVQAL--FKDFVEVYDKT--VEES-H  LSEMTVDAL---SI  211
WP_002305844  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_002307203  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQKL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_002310390  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_002352408  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_012997688  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_014677909  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_019312892  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_019313659  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_019314093  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQKL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_019315370  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_019803776  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_019805234  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_024783594  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_024784288  144  DSTEKADLRLVYILALAHMIKFRGHFLIEGE-LNAENTDVQKL--FADFVGVYDRT--FDDS-H  LSEITVDAS---SI  211
WP_024784666  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_024784894  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_024786433  144  DSTEKADLRLVYILALAHMIKFRGHFLIEGE-LNAENTDVQKL--FADFVGVYDRT--FDDS-H  LSEITVDAS---SI  211
WP_049473442  144  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
WP_049474547  144  DNPEKTDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  211
EMC03581      137  DNPEKVDLRLVYILALAHIIKFRGHFLIEGK-FDTRNNDVQRL--FQEFLAVYDNT--FENS-S  LQEQNVQVE---EI  204
WP_000428612  145  DSKEKTDLRLIYILALAHMIKYRGHFLYEDT-FDIKNNDIQKI--FNEFISIYNNT--FEGN-S  LSGQNVQVE---AI  212
WP_000428613  145  DSKEKTDLRLIYILALAHMIKYRGHFLYEDT-FDIKNNDIQKI--FSEFISIYDNT--FEGS-S  LSGQNAQVE---AI  212
WP_049523028  144  DSKEKVDLRLIYILALAHIIKYRGHFLYEDS-FDIKNNDIQKI--FNEFTILYDNT--FEES-S  LSKGNAQVE---EI  211
WP_003107102  113  DSDEKADLRLIYILALAHIIKFRGHFLIEGD-LDSQNTDVNAL--FLKLVDTYNLM--FEDD--  IDTQTIDAT---VI  180
WP_054279288  146  DNTEKADLRLIYILALAHIIKFRGHFLIEGA-LSANNTDVQQL--VHALVDAYNIM--FEED--  LDIEAIDVK---AI  213
WP_049531101  145  DSKEKADLRLIYLTLAHMIKYRGHFLYEES-FDIKNNDIQKI--FNEFISIYDNT--FEGS-S  LSGQNAQVE---AI  212
WP_049538452  145  DSKEKADLRLIYILALAHMIKYRGHFLYEEA-FDIKNNDIQKI--FNEFINIYDNT--FEGS-S  LSGQNEQVE---AI  212
WP_049549711  145  DSKEKADLRLIYLVLAHMIKYRGHFLYEEA-FDIKNNDIQKI--FNEFISIYDNT--FEGS-S  LSGQNAQVE---TI  212
WP_007896501  146  DRDQKADLRLIYALSHIIKFRGHFLIEGK-LNSENTDVQKL--FIALVTVYNLL--FEEE--  IAGETCDAK---AL  213
EFR44625       98  DRDQKADLRLIYILALSHIIKFRGHFLIEGK-LNSENTDVQKL--FIALVTVYNLL--FEEE--  IAGETCDAK---AL  165
WP_002897477  144  DSKEKSDVRLIYILALAHMIKYRGHFLYEET-FDIKNNDIQKI--FNEFINIYDNT--FEGS-S  LSGQNAQVE---AI  211
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WP_009729476  145  DSKEKTDLRLIYLALAHMIKYRGHFLYEEA-FDIKNNDIQKI--FNEFISIYNNT--FEGN-S  LSGQNVQVE---AI  212
CQR24647      144  DSSEKADLRLVYLALAHIIKYRGHFLIDEP-IDIRNMNSQNL--FKEFLLAFDGI--QVDC-Y  LASKHTDIS---GI  211
WP_000066813  145  DSKEKTDLRLIYLALAHMIKYRGHFLYEES-FDIKNNDIQKI--FSEFISIYDNT--FEGK-S  LSGQNAQVE---AI  212
WP_009754323  145  DSKEKADLRLIYLALAHITKYRGHFLYEEA-FDIKNNDIQKI--FNEFINIYDNT--FEGS-S  LSGQNAQVE---AI  212
WP_044674937  144  DSSQKADIRLIYLALAHIIKYRGHFLFEGD-LKSENKDVQHL--FNDFVEMFDKT--VEGS-Y  LSENLPNVA---DV  211
WP_044676715  144  DSSQKADIRLIYLALAHIIKYRGHFLFEGD-LKSENKDVQHL--FNDFVEMFDKT--VEGS-Y  LSENLPNVA---DV  211
WP_044680361  144  DSSQKADIRLIYLALAHIIKYRGHFLFEGD-LKSENKDVQHL--FNDFVEMFDKT--VEGS-Y  LSENLPNVA---DV  211
WP_044681799  144  DSSQKADIRLIYLALAHIIKYRGHFLFEGD-LKSENKDVQHL--FNDFVEMFDKT--VEGS-Y  LSENLPNVA---DV  211
WP_049533112  144  DISQKADLRLVYLALAHMIKFRGHFLIEGQ-LKAENTNVQAL--FKDFVEVYDKT--VEES-H  LSEMTVDAL---SI  211
WP_029090905  101  SQHRQFDIREVYLAIHHLIKYRGHFIYEDQtFTTDGNQLQHH--IKAIITMINST1---NR--  IIPETIDINvfeKI  171
WP_006506696  140  ESTEKADPRLIYLALHHIVKYRGNFLYEGQkFNMDASNIEDK--LSDIFTQFTSFnnIPYEdD  --KKNLEIL---EI  210
AIT42264      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--  INASGVDAK---AI  211
WP_034440723  143  DSNQKADLRLIYLALAHMIKYRGHFLIEGD-LKMDGISISES--FQEFIDSYNEVcaLEDE-N  NDELLTQIE---NI  217
AKQ21048      144  DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--  INASGVDAK---AI  211
WP_004636532  144  DNPEKADLRLVYTALAHIVKYRGHFLIEGE-LNTENTSISET--FEQFLDTYSDI--FKEQ--  LVGDISKVE---EI  210
WP_002364836  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_016631044  95   DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENTSVKDQ--FQQFMVIYNQT--FVNGeS  PLPESVLIE---EE  168
EMS75795      ---  ----------------------------------------------------------  ----------  ---
WP_002373311  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENTSVKEQ--FQQFMVIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_002378009  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_002407324  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_002413717  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_010775580  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_010818269  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_010824395  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENTSVKDQ--FQQFMVIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_016622645  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEK--FQQFMIIYNQT--FVNGeG  PLPESVLIE---EE  217
WP_033624816  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKDQ--FQQFMVIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_033625576  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_033789179  144  DSSEQADLRLIYLALAHIVKYRGHFLIEGK-LSTENISVKEQ--FQQFMIIYNQT--FVNGeS  PLPESVLIE---EE  217
WP_002310644  144  DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D  KLDEAVDCS---FV  216
WP_002312694  144  DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-G  KLDEAVDCS---FV  216
WP_002314015  144  DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D  KLDEAVDCS---FV  216
WP_002320716  144  DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D  KLDEAVDCS---FV  216
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WP_002330729  144   DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D
WP_002335161  144   DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D
WP_002345439  144   DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D
WP_034867970  144   DSTEKEDLRLVYLALAHLLKYRGHFLFEGD-LDTENTSIEES--FRVFLEQYSKQ--SDQP--
WP_047937432  144   DSSEKADIRLVYLAMAHLLKYRGHFLIEGE-LNTENSSVTET--FRQFLSTYNQQ--FSEA-D
WP_010720994  144   DSTEKGDLRLVYLALAHLLKYRGHFLFEGD-LDTENTSIEES--FRVFLEQYGKQ--SDQP--
WP_010737004  144   DSTEKEDLRLVYLALAHLLKYRGHFLFEGD-LDTENTSIEES--FRVFLEQYSKQ--SDQP--
WP_034700478  144   DSTEKEDLRLVYLALAHLLKYRGHFLFEGD-LDTENTSIEES--FRVFLEQYGKQ--SDQP--
WP_007209003  144   DGDEKADLRLVYLAIAHIIKFRGNFLIEGE-LNTENNSVIELs--KVFVQLYNQTl-SELE--
WP_023519017  144   NSKEQADIRLVYLAIAHCLKYRGHFLFEGE-LDTENTSVTEN--YQQFLQAYQQF--FPEP--
WP_010770040  144   DTSEQADLRLVYLALAHIVKYRGHFLIEGE-LNTENSSVSET--FRTFIQVYNQI--FRENe-
WP_048604708  144   DAEEKADLRLVYLALAHIIKYRGHFLIEGR-LSTENTSTEET--FKTFLQKYNQT--FN----
WP_010750235  144   DSTEKADIRLVYLALAHMIKYRGHFLFEGE-LDTENTSVEET--FKEFIDIYNEQ--FEEG--
AII16583      183   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
WP_029073316  145   ESKEKEDPRLIYLALHHIVKYRGNFLYEGQkFSMDVSNIEDK--MIDVLRQFNEInlFEYVeD
WP_031589969  145   ESKEKEDPRLIYLALHHIVKYRGNFLYEGQkFSMDVSNIEDK--MIDVLRQFNEInlFEYVeD
KDA45870      145   NNDRPADLRLVYLALAHIIKYRGNFLLEGE-IDLRTTDINKV--FAEFSETLNEN--SDENlG
WP_039099354  133   TEKRQFDIREIYLAMHHIVKYRGHFLNEAPvSSFKSSEINLVahFDRLNTIFADL--FSESgF
AKP02966      138   INKNKADIRLVYLALHNILKYRGNFTYEHQkFNISTLNSNLS---KELIELNQQLikYDIS--
WP_010991369  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTQNTSVDGI--YKQFIQTYNQV--FASGiE
WP_033838504  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTQNTSVDGI--YKQFIQTYNQV--FASGiE
EHN60060      147   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTQNTSVDGI--YKQFIQTYNQV--FASGiE
EFR89594            ------------------------------------------------------------
WP_038409211  144   NSSDKADLRLVYLALAHIIKYRGNFLIEGM-LDTKNTSVDEV--FKQFIQTYNQI--FASDiE
EFR95520            ------------------------------------------------------------
WP_003723650  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDEV--YKQFIETYNQV--FMSNiE
WP_003727705  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDEV--YKQFIQTYNQV--FMSNiE
WP_003730785  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDEV--YKQFIQTYNQV--FMSNiE
WP_003733029  144   DSQKKADLRLVYLALAHIIKYRGHFLIEGA-LDTKNTSIDEM--FKQFLQIYNQV--FANDiE
WP_003739838  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YKQFIQTYNQV--FISNiE
WP_014601172  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YEQFIQTYNQV--FMSNiE
WP_023548323  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YEQFILTYNQV--FMSNiE
WP_031665337  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YEQFIQTYNQV--FMSNiE
WP_031669209  144   DSQKKADLRLVYLALAHIIKYRGHFLIEGA-LDTKNTSIDEM--FKQFLQIYNQV--FANDiE
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WP_033920898  144   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YEQFIQTYNQV--FMSNiE
AKI42028      147   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YEQFIQTYNQV--FMSNiE
AKI50529      147   NSSEKADLRLVYLALAHIIKYRGNFLIEGA-LDTKNTSVDGV--YEQFIQTYNQV--FMSNiE
EFR83390            ------------------------------------------------------------
WP_046323366  144   NSSDKADLRLVYLALAHIIKYRGNFLIEGK-LDTKNTSVDEV--FKQFIKTYNQV--FASDiE
AKE81011      160   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
CUO82355      144   ESTEKADPRLIYLALHHIVKYRGNFLYEGQkFNMDASNIEDK--LSDVFTQFADFnnIPYEdD
WP_033162887  145   ENKEKADPRLIYLALHHIVKYRGNFLYEGQsFTMDNSDIEER--LNSAIEKFMSIneFDNRiV
AGZ01981      177   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
AKA60242      144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
AKS40380      144   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
4UN5_B        148   DSTDKADLRLIYLALAHMIKFRGHFLIEGD-LNPDNSDVDKL--FIQLVQTYNQL--FEEN--
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LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
ISKSRRLENLIKY-Y-PT   EKKNTLFGNLIALALGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE  278
ISKSAKRERVLKL-F-PD   EKSTGLFSEFLKLIVGNQADFKKHF--DLEDK-A---PLQ--FSKDTYDEDLEN  277
-------EQISRN-S--K   ---------------------------LEEKyVa--ELQ--------------  157
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LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
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LSKSRRLENLIAQ-L-PG   EKRNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
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LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-T---KLQ--LSKDTYDDDLDN  277
LSKSRRLENLIAQ-L-PG   EKRNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  188
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VSKSRRLENLIAH-Y-PA   EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEG  277
ISKSRRLENLIKY-Y-PT   EKKNTLFGNLIALALGLQPNFKMNF--KLSED-A---KLQ--FSKDSYEEDLGE  280
ISKSAKKDRILAQ-Y-PN   QKSTGIFAEFLKLIVGNQADFKKHF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
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WP_001040081  213  LTDK-ISKSAKKDRILAQ-Y-PN  QKSTGIFAEFLKLIVGNQADFKKYF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
WP_001040083  213  LTDK-ISKSAKKDRILAQ-Y-PN  QKSTGIFAEFLKLIVGNQADFKKYF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
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WP_001040088  213  LTDK-ISKSAKKDRILAQ-Y-PN  QKSTGIFAEFLKLIVGNQADFKKYF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
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WP_001040098  213  LTDK-ISKSAKKDRILAR-Y-PN  QKSTGIFAEFLKLIVGNQADFKKHF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
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CFQ25032      213  LTDK-ISKSAKKDRILAQ-Y-PN  QKSTGIFAEFLKLIVGNQADFKKYF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
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KLL20707          213  LTDK-ISKSAKKDRILAQ-Y-PN  QKSTGIFAEFLKLIVGNQADFKKYF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
KLL42645          213  LTDK-ISKSAKKDRILAQ-Y-PN  QKSTGIFAEFLKLIVGNQADFKKHF--NLEDK-T---PLQ--FAKDSYDEDLEN  278
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CCW42055          213  LTEK-VSKSRRLENLIAH-Y-PA  EKKNTLFGNLIALFLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEG  277
WP_003041502      212  LTEK-VSKSRRLENLIAH-Y-PT  EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEE  278
WP_037593752      213  LTEK-VSKSSRLENLIAH-Y-PT  EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEE  278
WP_049516684      213  LTEK-VSKSRRLENLVEC-Y-PT  EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEG  278
GAD46167          212  LTEK-VSKSSRLENLIAH-Y-PT  EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEE  277
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WP_003043819      212  LSAR-LSKSKRLEKLIAV-F-PN  EKKNGLFGNIIALALGLTPNFKSNF--DLTED-A---KLQ--LSKDTYDDDLDE  277
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WP_004232481      212  LTEK-ISKSRRLENLIKQ-Y-PT  EKKNTLFGNLVALALGLQPNFKTNF--KLSED-A---KLQ--FSKDTYDEDLEE  277
WP_009854540      213  LTEK-ISKSRRLENLIKY-Y-PT  EKKNTLFGNLIALALGLQPNFKMNF--KLSED-A---KLQ--FSKDTYEEDLEE  278
WP_012962174      213  LTEK-FSKSRRLENLIKH-Y-PT  EKKNTLFGNLVALALGLQPNFKTSF--KLSED-A---KLQ--FSKDTYEEDLEE  278
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WP_014334983 212 LTEK-VSKSRRLENLIKQ-Y-PT EKKNTLFGNLIALALGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE 277
WP_003099269 212 LTSK-TSKSRRLENLIAE-I-PN QKRNMLFGNLVSLALGLTPNFKTNF--ELLED-A---KLQ--ISKDSYEEDLDN 277
AHY15608     212 LTSK-TSKSRRLENLIAE-I-PN QKRNMLFGNLVSLALGLTPNFKTNF--ELLED-A---KLQ--ISKDSYEEDLDN 277
AHY17476     212 LTSK-TSKSRRLENLIAE-I-PN QKRNMLFGNLVSLALGLTPNFKTNF--ELLED-A---KLQ--ISKDSYEEDLDN 277
ESR09100     ---------------------- ---------------------------------------------------------
AGM98575     212 LTSK-TSKSRRLENLIAE-I-PN QKRNMLFGNLVSLALGLTPNFKTNF--ELLED-A---KLQ--ISKDSYEEDLDN 277
ALF27331     212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEELEV 277
WP_018372492 210 FTEN-SSKAKRVETILGL-F-PD ETAAGNLDKFLKLMLGNQADFKKVF--DLEEK-----ITLQ--FSKDSYEEDLEL 275
WP_045618028 213 FTDK-ISKSAKRERVLKL-F-PD EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYDEDLEN 278
WP_045635197 212 FTDK-ISKSAKRERVLKL-F-PD EKSTGLFSEFLKLIVGNQADFKKHF--DLEDK-A---PLQ--FSKDTYDEDLEN 277
WP_002263549 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002263887 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002264920 212 LTEK-ISKSRRLEKLINN-Y-PK EKKNTLFRNLVALSLGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE 277
WP_002269043 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV 277
WP_002269448 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002271977 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEDLEE 277
WP_002272766 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEDLEE 277
WP_002273241 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002275430 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002276448 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002277050 212 LTEK-ISKSRRLEKLINN-Y-PK EKKNTLFGNLIALSLGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE 277
WP_002277364 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002279025 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002279859 212 LTEK-ISKSRRLEKLINN-Y-PK EKKNTLFGNLIALSLGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEELEV 277
WP_002280230 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002281696 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002282247 212 LTEK-ISKSRRLEKLINN-Y-PK EKKNTLFGNLIALSLGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE 277
WP_002282906 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002283846 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV 277
WP_002287255 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002288990 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV 277
WP_002289641 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGCFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV 277
WP_002290427 212 LTDK-ISKSAKKDRVLKL-F-PN EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV 277
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WP_002295753  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_002296423  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV  277
WP_002304487  212  LTEK-VSKSRRLENLVEC-Y-PT  EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEG  277
WP_002305844  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_002307203  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--LSKDTYEEELEV  277
WP_002310390  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV  277
WP_002352408  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV  277
WP_012997688  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_014677909  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV  277
WP_019312892  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_019313659  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_019314093  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--LSKDTYEEELEV  277
WP_019315370  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_019803776  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-V---PLQ--FSKDTYEEELEV  277
WP_019805234  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDIYEEELEV  277
WP_024783594  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_024784288  212  LTEK-ISKSRRLEKLINN-Y-PK  EKKNTLFGNLIALSLGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE  277
WP_024784666  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_024784894  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
WP_024786433  212  LTEK-ISKSRRLEKLINN-Y-PK  EKKNTLFGNLIALSLGLQPNFKTNF--KLSED-A---KLQ--FSKDTYEEDLEE  277
WP_049473442  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEDLEE  277
WP_049474547  212  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  277
EMC03581      205  LTDK-ISKSAKKDRVLKL-F-PN  EKSNGRFAEFLKLIVGNQADFKKHF--ELEEK-A---PLQ--FSKDTYEEELEV  270
WP_000428612  213  FTDK-ISKSAKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSRDTYDEDLEN  278
WP_000428613  213  FTDK-ISKSAKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLGEK-A---PLQ--FSKDTYDEDLEN  278
WP_049523028  212  FTDK-ISKSAKRDRVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYEEDLES  277
WP_003107102  181  LTEK-MSKSRRLENLIAK-I-PN  QKKNTLFGNLISLSLGLTPNFKANF--ELSED-A---KLQ--ISKESFEEDLDN  246
WP_054279288  214  LTEK-ISKTRRLENLISN-I-PG  QKKNGLFGNLIALSLGLTPNFKSHF--NLPED-A---KLQ--LAKDTYDEELNN  279
WP_049531101  213  FTDK-ISKSTKRERVLKL-F-PD  QKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYDEDLEN  278
WP_049538452  213  FSDK-ISKSAKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYDEDLEN  278
WP_049549711  213  FTDK-ISKSAKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLGEK-A---PLQ--FSKDTYDEDLEN  278
WP_007896501  214  LTAK-TSKSKRLESLISE-F-PG  QKKNGLFGNLLALALGLRPNFKSNF--GLSED-A---KLQ--ITKDTYEEELDN  279
EFR44625      166  LTAK-TSKSKRLESLISE-F-PG  QKKNGLFGNLLALALGLRPNFKSNF--GLSED-A---KLQ--ITKDTYEEELDN  231
WP_002897477  212  FTDK-ISKSAKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYDEELEN  277
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WP_002906454  212  FTDK-ISKSTKRERVLKL-F-SD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYDEDLEN  277
WP_009729476  213  FTDK-ISKSAKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSRDTYDEDLEN  278
CQR24647      212  ITAK-ISKSRKVEAVLEQ-F-PD  QKKNSFFGNMVSLVFGLMPNFKSNF--ELDED-A---KLQ--FSRDSYDEDLEN  277
WP_000066813  213  FTDK-ISKSTKRERVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---PLQ--FSKDTYDEDLEN  278
WP_009754323  213  FTGK-ISKSVKREHVLKL-F-PD  EKSTGLFSEFLKLIVGNQADFKKHF--DLEEK-A---SLQ--FSKDTYDEDLEN  278
WP_044674937  212  LVEK-VSKSRRLENILHY-F-PN  EKKNGLFGNFLALALGLQPNFKTNF--ELAED-A---KIQ--FSKETYEEDLEE  277
WP_044676715  212  LVEK-VSKSRRLENILHY-F-PN  EKKNGLFGNFLTLALGLQPNFKTNF--ELAED-A---KIQ--FSKETYEEDLEE  277
WP_044680361  212  LVEK-VSKSRRLENILHY-F-PN  EKKNGLFGNFLALALGLQPNFKTNF--ELAED-A---KIQ--FSKETYEEDLEE  277
WP_044681799  212  LVEK-VSKSRRLENILHY-F-PN  EKKNGLFGNFLALALGLQPNFKTNF--ELAED-A---KIQ--FSKETYEEDLEE  277
WP_049533112  212  LTEK-VSKSRRLENLIAH-Y-PA  EKKNTLFGNLIALSLGLQPNFKTNF--QLSED-A---KLQ--FSKDTYEEDLEG  277
WP_029090905  172  LLDRmMNRSSKVKFLIEL--TG  KQDKPLLKELFNLIVGLKAKPASIFe---QENlAtivETM-nMSTEQVQLDLLT  243
WP_006506696  211  LKKP-LSKKAKVDEVMTL-IaPE  KDYKSAFKELVTGIAGNKMNVTKMIlcEPIKQ-Gds-EIKlkFSDSNYDDQFSE  283
AIT42264      212  LSAR-LSKSRRLENLIAQ-L-PG  EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
WP_034440723  218  FKQD-ISRSKKLDQAIAL-F-QG  -KRQSLFGIFLTLIVGNKANFQKIF--NLEDD----iKLD--LKEEDYDENLEE  283
AKQ21048      212  LSAR-LSKSRRLENLIAQ-L-PG  EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---KLQ--LSKDTYDDDLDN  277
WP_004636532  211  LSSK-QSRSRKHEQIMAL-F-PN  ENKLGNFGRFMMLIVGNTSNFKPVF--DLDDE-Y---KLK--LSDETYEEDLDT  276
WP_002364836  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_016631044  169  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  234
EMS75795      1    ----------------------  ----------------------MDEE-A---KIQ--LSKESYEEELES  20
WP_002373311  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_002378009  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_002407324  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_002413717  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_010775580  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KIKitYASESYEEDLEG  285
WP_010818269  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_010824395  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_016622645  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_033624816  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_033625576  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_033789179  218  LTEK-ASRTKKSEKVLQQ-F-PQ  EKANGLFGQFLKLMVGNKADFKKVF--GLEEE-A---KI--tYASESYEEDLEG  283
WP_002310644  217  FTEK-MSKTKKAETLLKY-F-PH  EKSNGYLSQFTKLMVGNQGNFKNVF--GL-EE-A---KLQ--FSKETYEEDLEE  281
WP_002312694  217  FTEK-MSKTKKAETLLKY-F-PH  EKSNGYLSQFTKLMVGNQGNFKNVF--GL-EEeA---KLQ--FSKETYEEDLEE  282
WP_002314015  217  FTEK-MSKTKKAETLLKY-F-PH  EKSNGYLSQFTKLMVGNQGNFKNVF--GL-EEeA---KLQ--FSKETYEEDLEE  282
WP_002320716  217  FTEK-MSKTKKAETLLKY-F-PH  EKSNGYLSQFTKLMVGNQGNFKNVF--GL-EEeA---KLQ--FSKETYEEDLEE  282
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WP_002330729  217   FTEK-MSKTKKAETLLKY-F-PH   EKSNGYLSQFIKLMVGNQGNFKNVF--GL-EE-A---
WP_002335161  217   FTEK-MSKTKKAETLLKY-F-PH   EKSNGYLSQFIKLMVGNQGNFKNVF--GL-EEeA---
WP_002345439  217   FTEK-MSKTKKAETLLKY-F-PH   EKSNGYLSQFIKLMVGNQGNFKNVF--GL-EEeA---
WP_034867970  210   LTDK-LSKTKKVEEILKY-Y-PT   EKINSFFAQCLKLIVGNQANFKRIF--DLEAE-V---
WP_047937432  217   FTEK-MSKTKKAETLLKY-F-PH   EKSNGYLSQFIKLMVGNQGNFKNVF--GL-EEeA---
WP_010720994  210   LTDK-LSKTKKVEEILKY-Y-PT   EKINSFFAQCLKLIVGNQANFKRIF--DLEAE-V---
WP_010737004  210   LTDK-LSKTKKVEEILKY-Y-PT   EKINSFFAQCLKLIVGNQANFKRIF--DLEAE-V---
WP_034700478  210   LTDK-LSKTKKVEEILKY-Y-PT   EKINSFFAQCLKLIVGNQANFKRIF--DLEAE-V---
WP_007209003  215   LTQQ-LSKSERADNVLKL-F-PD   EKGTGIFAQFIKLIVGNQGNFKKVF--QLEED----q
WP_023519017  210   LTER-LSKAKRVEKVLAY-Y-PS   EKSTGNFAQFLKLMVGNQANFKKTF--DLEEE-M---
WP_010770040  213   FSEK-VSRARKVEAILSV-Y-SE   EKSTGTLAQFLKLMVGNQGRFKKTF--DLEED-G---
WP_048604708  209   FADK-VSRAKKAEGVLAL-F-PD   EKRNGTFDQFLKMIVGNQGNFKKTF--ELEED-A---
WP_010750235  210   LTDK-LSKSKKVEKILQY-Y-PK   EKTTGCLAQFLKLIVGNQGNFKQAF--HLDEE-V---
AII16583      251   LSAR-LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---
WP_029073316  216   LKEP-LSKKHKADKAFAL-FdTT   KDNKAAYKELCAALAGNKFNVTKMLkeAELHD-Edek
WP_031589969  216   LKEP-LSKKHKAEKAFAL-FdTT   KDNKAAYKELCAALAGNKFNVTKMLkeAELHD-Edek
KDA45870      210   FKDNtFSKTKKSEELLKL---SG   -KKNQLAHQLFKMMVGNMGSFKKVL--GTDEE----h
WP_039099354  207   LLDNhQSASNRQRQALLLiYtPS   KQNKAIATELLKAILGLKAKFNVLT--GIEAEdVktw
AKP02966      209   LIGR-GNATQKSSNILNN-F--T   KETKKLLKEVINLILGNVAHLNTIFktSLTKDeE---
WP_010991369  218   LVEK-VTRKEKLERILKL-Y-PG   EKSAGMFAQFISLIVGSKGNFQKPF--DLIEK-S---
WP_033838504  218   LVEK-VTRKEKLERILKL-Y-PG   EKSAGMFAQFISLIVGSKGNFQKPF--DLIEK-S---
EHN60060      221   LVEK-VTRKEKLERILKL-Y-PG   EKSAGMFAQFISLIVGSKGNFQKPF--DLIEK-S---
EFR89594      1     ---------------LKL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKPF--DLIEK-S---
WP_038409211  218   LSEK-LTRREKLDKILKL-Y-TG   EKSTGMFARFINLIIGSKGDFKKVF--DLDEK-A---
EFR95520            ----------------------   ------------------------------------
WP_003723650  218   LAGK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
WP_003727705  218   LAGK-FTRREKFERILRL-Y-PG   EKSTGMFAQFISLIVGNKGNFQKVF--NLVEK-T---
WP_003730785  218   LAGK-FTRREKFERILRL-Y-PG   EKSTGMFAQFISLIVGNKGNFQKVF--NLVEK-T---
WP_003733029  218   LAEK-FTRKDKLDKILSL-Y-PG   EKTTGVFAQFVNIIVGSTGKFKKHF--NLHEK-K---
WP_003739838  218   LAGK-FTRKEKLERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLVEK-T---
WP_014601172  218   LAGK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
WP_023548323  218   LAGK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
WP_031665337  218   LAGK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
WP_031669209  218   LAEK-FTRKDKLDKILSL-Y-PG   EKTTGVFAQFVNIIVGSTGKFKKHF--NLHEK-K---
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WP_033920898  218   LARK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
AKI42028      221   LAGK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
AKI50529      221   LARK-FTRREKFERILQL-Y-PG   EKSTGMFAQFISLIVGSKGNFQKVF--DLIEK-T---
EFR83390            ----------------------   ------------------------------------
WP_046323366  218   FSEK-LTKREKLDKILNL-Y-PN   EKSTDLFAQFISLIIGSKGNFKKFF--NLTEK-T---
AKE81011      228   LSAR-LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---
CUO82355      215   LKKP-LSKKAKVDEVMAL-IsPE   KEFKSAYKELVTGIAGNKMNVTKMIlcESIKQ-Gds-
WP_033162887  216   LSKI-YQRSKKADDLLKI-MnPT   KEEKAAYKEFTKALVGLKFNISKMIlaQEVKK-Gdt-
AGZ01981      245   LSAR-LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---
AKA60242      212   LSAR-LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---
AKS40380      212   LSAR-LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---
4UN5_B        216   LSAR-LSKSRRLENLIAQ-L-PG   EKKNGLFGNLIALSLGLTPNFKSNF--DLAED-A---
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WP_002906454  278   LLGQIGDGFADLFLVAKKLYDAILLSGILTVTDPSTKAPLSASMIERYENHQEDLAALKQFIK
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CQR24647      278   LLGIIGDEYADVFVAAKKVYDSILLSGILTTNNHSTKAPLSASMIDRYDEHNSDKKLLRDFIR
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WP_044681799  278   LLGKIGDDYADLFIATKSLYDGILLAGILSTTDSTTKAPLSSSMVNRYEEHKKDLALLKNFIH
WP_049533112  278   LLGEIGDEYADLFASAKNLYDAILLSGILTVDDNSTKAPLSASMVKRYEEHQKDLKKLKDFIK
WP_029090905  244   LADVLADEEYDLLLTAQKIYSAIILDESMDGYEYFA-----EAKKESYRKHQEELVLVKKMLK
WP_006506696  284   VEKDLGE-YVEFVDALHNVYSWVELQTIMGATHTD-NASISEAMVSRYNKHHDDLKLLKDCIK
AIT42264      278   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
WP_034440723  284   LLSNIDEGYRDVFLQAKNVYNAIELSKILKTDGKETKAPLSAQMVELYNQHREDLKKYKDYIK
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WP_004636532  277   LLGMTDDVFLDVFMAAKNVYDAVEMSAIISTDTGNSKAVLSNQMINFYDEHKVDLAQLKQFFK
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WP_016631044  235   ILAKVGDEYSDVFLAAKNVYDAVELSTILADSDKKSHAKLSSSMIVRFTEHQEDLKKFKRFIR
EMS75795      21    LLEKSGEEFRDVFLQAKKVYDAILLSDILSTKKQNSKAKLSLGMIERYDSHKKDLEELKQFVK
WP_002373311  284   ILAKVGDEYSDVFLAAKNVYDAVELSTILADSDKKSHAKLSSSMIVRFTEHQEDLKKFKRFIR
WP_002378009  284   ILAKVGDEYSDVFLAAKNVYDAVELSTILADSDKKSHAKLSSSMIVRFTEHQEDLKNFKRFIR
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WP_010824395  284   ILAKVGDEYSDVFLAAKNVYDAVELSTILADSDKKSHAKLSSSMIVRFTEHQEDLKNFKRFIR
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WP_002314015  283   LLEKIGDDYIDLFVQAKNVYDAVLLSEILSDSTKNTRAKLSAGMIRRYDAHKEDLVLLKRFVK
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WP_002345439  283   LLEKIGDDYIDLFVQAKNVYDAVLLSEILSDSTKNTRAKLSAGMIRRYDAHKEDLVLLKRFVK
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WP_007209003  281   LLAIIGDEYGDIFVAAQNLYQAILLAGILTSTEK-TRAKLSASMIQRYEEHAKDLKLLKRFVK
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WP_010770040  279   LLAIIGDEYAEIFSATKSVYDAVALSGILSVTDGDTKAKLSASMVERYEAHQKDLVQFKQFIR
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WP_010750235  276   LLRKSNEEMIDVFLQVKKVYDAILLSDILSTKMKDTKAKLSAGMIERYQNHKKDLEELKQFVR
AII16583      317   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
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KDA45870      276   LLAEAGDQYLDIFVAAKKVYDAAILASILDVKDTQTKTVFSQAMIERYEEHQKDLIELKRVFK
WP_039099354  286   LESSLDDNAHQIIESLQELYSGVLLAGIVPENQSLS-----QAMITKYDDHQKHLKMLKAVRE
AKP02966      279   LDSILDDDQFTVLDTANRIYSTITLNEIL-----NGESYFSMAKVNQYENHAIDLCKLRDMWH
WP_010991369  284   LLALIGDEYAELFVAAKNAYSAVVLSSIITVAETETNAKLSASMIERFDTHEEDLGELKAFIK
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EHN60060      287   LLALIGDEYAELFVAAKNAYSAVVLSSIITVAETETNAKLSASMIERFDTHEEDLGELKAFIK
EFR89594      53    LLALIGDEYAELFVAAKNAYSAVVLSSIITVAETETNAKLSASMIERFDTHEEDLGELKAFIK
WP_038409211  284   LLAKIGDEYAEIFVAAKSTYNAVVLSNIITVTDTETKAKLSASMIERFDKHAKDLKRLKAFFK
EFR95520              ------------------------------------------------------------
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WP_003730785  284   LLAIIGDEYAELFVAAKNTYNAVVLSSIITVTATETNAKLSASMIERFDAHEKELGELKAFIK
WP_003733029  284   LLAIIGDEFAEVFVAAKNAYNAVVLSNIITVTDSTTRAKLSASLIERFENHKEDLKKMKRFVR
WP_003739838  284   LLAIIGDEYAELFVAAKNTYNAVVLSSIITVTDTETNAKLSASMIERFDAHEKDLSELKAFIK
WP_014601172  284   LLAIIGDEYAELFVAAKNTYNAVVLSSIITVTATETNAKLSASMIERFDAHEKDLGELKAFIK
WP_023548323  284   LLAIIGDEYAELFVAAKNTYNAVVLSSIITVTDTETNAKLSASMIERFDAHEKDLVELKAFIK
WP_031665337  284   LLAIIGDEYAELFVAAKNTYNAVVLSSIITVNDTETNAKLSASMIERFDAHEKDLVELKAFIK
WP_031669209  284   LLAIIGDEFAEVFVAAKNAYNAVVLSNIITVTDSTTRAKLSASLIERFENHKEDLKKMKRFVR
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WP_046323366  284   LLARVGDEYAEIFVAAKNAYNAVVLSSIITVSNTETKAKLSASMIERFDKHDKDLKRMKAFFK
AKE81011      294   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
CUO82355      288   VENDLGE-YVEFIDSLHNIYSWVELQTIMGATHTD-NASISEAMVSRYNKHHEDLQLLKKCIK
WP_033162887  289   LQSELGE-YIEFIEMLHNIYSWVELQAILGATHTD-NPSISAAMVERYEEHKKDLRVLKKVIR
AGZ01981      311   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
AKA60242      278   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
AKS40380      278   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
4UN5_B        282   LLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVR
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WP_039695303  358   --NGYAG   YIEN   G    VKQDEFYKYLKNILSK-IkiDGSDYFL--DKIEREDFLRKQ
WP_045635197  357   --DGYAG   YIDG   K    TTQETFYKYIKNLLSK-F--EGTDYFL--DKIEREDFLRKQ
5AXW_A        169   ------G   SINR   -    ---------------K------TSDYVk-------------
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BAQ51233      268   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
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KGE60856            -------   ----   -    ----------------------------------------
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WP_003065552  360   --NGYAG   YIEN   G    VKQDEFYKYLKNTLSK-Ia--GSDYFL--DKIEREDFLRKQ
WP_001040076  358   --DGYAG   YIEG   K    TNQEAFYKYLSKLLTK-Q--EGSEYFL--EKIKNEDFLRKQ
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CCW42055          358  --DGYAG  YIEG  K  TNQEAFYKYLSKLLTK-Q--EGSEYLL--EKIKNEDFLRKQRTFDNGSIPHQVHLTEL  420
WP_003041502      357  --KGYAG  YIES  G  VKQDEFYKYLKGILLQ-I--NGSGDFL--DKIDREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_037593752      358  --KGYAG  YIES  G  VEQDEFYKYLKGILLK-I--NGSGDFL--DKIDCEDFLRKQRTFDNGSIPHQIHLQEM  420
WP_049516684      358  --NGYAG  YIEN  G  VKQDEFYKYLKNTLSK-I--DGSDYFL--DKIDREDFLRKQRTFDNGSIPHQIHLQEM  420
GAD46167          357  --KGYAG  YIES  G  VEQDEFYKYLKGILLK-I--NGSGDFL--DKIDCEDFLRKQRTFDNGSIPHQIHLQEM  419
WP_018363470      358  --NGYAG  YIEN  G  VKQDEFYKYLKGILTK-I--NGSDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_003043819      357  --NGYAG  YVGI  G  ATQEEFYKFIKPILEK-M--DGAEELLa--KLNRDDLLRKQRTFDNGSIPHQIHLKEL  429
WP_006269658      357  --KGYAS  YIES  G  VKQDEFYKYLKGILLK-I--NGSGDFL--DKIDREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_048800889      357  --NGYAG  YIEN  G  VKQDEFYKYLKNTLSK-I--DGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_012767106      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLa--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_014612333      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLa--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_015017095      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLa--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_015057649      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLa--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_048327215      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLa--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_049519324      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLa--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_012515931      357  --NGYAG  YIEG  Q  VSQEEFYKYLKPILAR-L--DGSEPLLl--KIDREDFLRKQRTFDNGSIPHQIHLEEL  419
WP_021320964      357  --NGYAG  YIEG  Q  VSQEEFYKYLKPILAR-L--DGSEPLLl--KIDREDFLRKQRTFDNGSIPHQIHLEEL  419
WP_037581760      357  --NGYAG  YIEG  Q  VSQEEFYKYLKPILAR-L--DGSEPLLl--KIDREDFLRKQRTFDNGSIPHQIHLEEL  419
WP_004232481      357  --NGYAG  YIEN  G  VKQDIFYKHLKSIISE-K--NGGQYFL--DKIEREDFLRKQRTFDNGSIPYQIHLQEM  419
WP_009854540      358  --NGYAG  YIEN  G  VKQDEFYKYLKNTLSK-I--DGSDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_012962174      358  --NGYAG  YIEN  G  VKQDEFYKYLKTTLSK-I--DGSDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
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WP_039695303 358 --NGYAG YIEN G VKQDEFYKYLKNILSK-IkiDGSDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 422
WP_014334983 357 --NGYAG YIDN G VKQDEFYKYLKTILTK-I--DDSDYFL--DKIERDDFLRKQRTFDNGSIPHQIHLQEM 419
WP_003099269 357 --NGYAG YIDG K TSQEEFYKYIKPILLK-L--DGTEKLIs--KLEREDFLRKQRTFDNGSIPHQIHLNEL 419
AHY15608     357 --NGYAG YIDG K TSQEEFYKYIKPILLK-L--DGTEKLIs--KLEREDFLRKQRTFDNGSIPHQIHLNEL 419
AHY17476     357 --NGYAG YIDG K TSQEEFYKYIKPILLK-L--DGTEKLIs--KLEREDFLRKQRTFDNGSIPHQIHLNEL 419
ESR09100     ------- ---- - ---------------------------------------------------------
AGM98575     357 --NGYAG YIDG K TSQEEFYKYIKPILLK-L--DGTEKLIs--KLEREDFLRKQRTFDNGSIPHQIHLNEL 419
ALF27331     357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_018372492 355 --PSYAA YVSG A VTEDDFYKFSKGLLID-V--EGAEYFL--EKIEREDFLRKQRTFDNGAIPNQVHVKEL 432
WP_045618028 358 --DGYAG YIDG K TTQEAFYKYIKNLLSK-L--EGADYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 420
WP_045635197 357 --DGYAG YIDG K TTQETFYKYIKNLLSK-F--EGTDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002263549 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002263887 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002264920 357 --NGYAG YIEN G VKQDEFYKYLKNTLSK-I--AGSDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002269043 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
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WP_002272766 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
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WP_002276448 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
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WP_002281696 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002282247 357 --NGYAG YIEN G VKQDEFYKYLKNTLSK-I--TGSDYFL--DQIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002282906 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
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WP_002289641 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
WP_002290427 357 --DGYAG YIDG K TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM 419
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WP_002296423  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_002304487  357  --NGYAG  YVGA  D  ATEEEFYKYVKGILNK-V--EGADVWL--DKIDREDFLRKQRTFDNGSIPHQIHLQEM  429
WP_002305844  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_002307203  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
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WP_002352408  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
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WP_019312892  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_019313659  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_019314093  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_019315370  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGNGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_019803776  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_019805234  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_024783594  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_024784288  357  --NGYAG  YIEN  G  VKQDEFYKYLKNTLSK-I--TGSDYFL--DQIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_024784666  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_024784894  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_024786433  357  --NGYAG  YIEN  G  VKQDEFYKYLKNTLSK-I--AGSDYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_049473442  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_049474547  357  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
EMC03581      350  --DGYAG  YIDG  K  TNQEAFYKYLKGLLNK-I--EGSGYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  412
WP_000428612  358  --DGYAG  YIDG  K  TTQESFYKYIKNLLSK-F--EGADYFL--EKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_000428613  358  --DGYAG  YIDG  K  TTQEAFYKYIKNLLSK-F--EGTDYFL--EKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_049523028  357  --DGYAG  YIDG  K  TTQEGFYKYIKNLISK-I--EGAEYFL--EKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
WP_003107102  326  --NGYAG  YING  K  TSQEDFYKYIKPILSK-L--KGAESLIs--KLEREDFLRKQRTFDNGSIPHQIHLNEL  388
WP_054279288  359  --DGYAG  YISG  K  TSQEAFYKYIKPILET-L--DGAEDFLt--KINREDFLRKQRTFDNGSIPHQIHLGEL  421
WP_049531101  358  --EGYAG  YIDS  K  TTQEAFYKYIKNLLSK-I--DGADYLL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_049538452  358  --DGYAG  YVDG  K  TTQEAFYKYIKNLLSK-F--EGADYFL--EKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_049549711  358  --DGYAG  YIDG  K  TTQEAFYKYIKNLLSK-F--EGTDYFL--EKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
WP_007896501  359  --NGYAG  YIEG  K  VSQEDFYRYIKPILSR-L--KGGDEFLa--KIDRDDFLRKQRTFDNGSIPHQIHLKEL  421
EFR44625      311  --NGYAG  YIEG  K  VSQEDFYRYIKPILSR-L--KGGDEFLa--KIDRDDFLRKQRTFDNGSIPHQIHLKEL  373
WP_002897477  357  --DGYAG  YIDG  K  TTQEAFYKYIKNLLSK-F--EGADYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  419
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WP_009729476  358  --DGYAG  YIDG  K  TTQETFYKYIKNLLSK-F--EGADYFL--DKIEREDFLRKQRTFDNGSIPHQIHLQEM  420
CQR24647      358  --NGYAG  YIDG  K  TNQEDFYKYLKNLLQK-V--DGGDYFI--EKIEREDFLRKQRTFDNGSIPHQVHLDEM  420
WP_000066813  358  --DGYAG  YIDG  K  TTQEAFYKYIKNLLSK-F--EGADYFL--DKIEREDFLKKQRTFDNGSIPHQIHLQEM  420
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WP_006506696  361  ksKGYYN  YINR  K  APVDEFYKYVKKCIEK-VdtPEAKQILn--DIELENFLLKQNSRTNGSVPYQMQLDEM  429
AIT42264      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_034440723  363  --NGYAG  YIDG  K  TSQEDFYKFVKAQLKG---eENGEYFL--EAIENENFLRKQRSFYNGVIPYQIHLQEL  425
AKQ21048      357  --NGYAG  YIDG  G  ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQRTFDNGSIPHQIHLGEL  419
WP_004636532  356  --NGYAG  YIDG  K  TNQEDFYKYIEKVMKT-IksDKKDYFL--DKIDREVFLRKQRSFYNSVIPHQIHLQEM  420
WP_002364836  363  --DGYAG  YIAH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
WP_016631044  314  --DGYAG  YIAH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  378
EMS75795      100  --NGYAG  YIDG  K  TTQEDFYKFLKKELNG-I--AGSERFM--EKVDQENFLLKQRTTANGVIPHQVHLTEL  162
WP_002373311  363  --DGYAG  YIAH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
WP_002378009  363  --DGYAG  YITH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
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WP_010824395  363  --DGYAG  YITH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
WP_016622645  363  --DGYAG  YIAH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
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WP_033625576  363  --DGYAG  YIAH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
WP_033789179  363  --DGYAG  YIAH  A  VSQLKFYQYVKKIIQD-I--AGAEYFL--EKIAQENFLRKQRTFDNGVIPHQIHLAEL  427
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WP_002314015  362  --NGYAG  YIEG  H  ATQEDFYKFVKKELTG-I--RGSEVFL--TKIEQENFLRKQRTFDNGVIPHQIHLTEL  424
WP_002320716  362  --NGYAG  YIEG  H  ATQEDFYKFVKKELTG-I--RGSEVFL--TKIEQENFLRKQRTFDNGVIPHQIHLTEL  424
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WP_002345439  362   --NGYAG   YIEG   H    ATQEDFYKFVKKELTG-I--RGSEVFL--TKIEQENFLRKQ
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WP_047937432  362   --NGYAG   YIEG   H    ATQEDFYKFVKKELTG-I--RGSEVFL--TKIEQENFLRKQ
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AII16583      396   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
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WP_031589969  367   kkNNYCN   YINH   K    TPVDEFYKYIKKLIEK-IddPDVKTILn--KIELESFMLKQ
KDA45870      354   -iSGYAG   YIDG   K    VSEEDFYKYTKKTLKG-I--PETEEILq--KIDANNYLRKQ
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EHN60060      366   --HGYAG   YIDG   -    TKQADFYKYMKMTLEN-I--EGADYFI--AKIEKENFLRKQ
EFR89594      132   --HGYAG   YIDG   -    TKQADFYKYMKTTLEN-I--EGADYFI--AKIEKENFLRKQ
WP_038409211  363   --DGYAG   YIDG   -    TTQEKFYKYMKKMLAN-I--DGADYFI--DQIEEENFLRKQ
EFR95520      1     -------   ----   -    ---------MKKMLAN-I--DGADYFI--DQIEEENFLRKQ
WP_003723650  363   --DGYAG   YIDG   -    TKQVDFYKYLKTILEN-I--EGSDYFI--AKIEEENFLRKQ
WP_003727705  363   --DGYAG   YIDG   -    TKQVDFYKYLKTTLEN-V--EGADYFI--TKIEEENFLRKQ
WP_003730785  363   --DGYAG   YIDG   -    TKQVDFYKYLKTTLEN-V--EGADYFI--TKIEEENFLRKQ
WP_003733029  363   --HGYAG   YISG   -    TKQADFYKYMKATLEK-I--EGADYFI--AKIEEENFLRKQ
WP_003739838  363   --DGYAG   YIDG   -    TKQVDFYKYLKTLLEN-I--EGADYFI--AKIEEENFLRKQ
WP_014601172  363   --DGYAG   YIDG   -    TKQVDFYKYLKTILEN-I--EGADYFI--AKIEEENFLRKQ
WP_023548323  363   --DGYAG   YIDG   -    TKQVDFYKYLKTTLEN-V--EGADYFI--TKIEEENFLRKQ
WP_031665337  363   --DGYAG   YIDG   -    TKQVDFYKYLKTILEN-I--EGSDYFI--AKIEEENFLRKQ
WP_031669209  363   --HGYAG   YISG   -    TKQADFYKYMKATLEK-I--EGADYFI--AKIEEENFLRKQ
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WP_033920898  363   --DGYAG   YIDG   -    TKQVDFYKYLKTILEN-I--EGADYFI--AKIEEENFLRKQ
AKI42028      366   --DGYAG   YIDG   -    TKQVDFYKYLKTILEN-I--EGADYFI--AKIEEENFLRKQ
AKI50529      366   --DGYAG   YIDG   -    TKQVDFYKYLKTILEN-I--EGADYFI--AKIEEENFLRKQ
EFR83390            -------   ----   -    ----------------------------------------
WP_046323366  363   --DGYAG   YIEG   -    TKQEAFYKYMKKMLEH-V--EGADYFI--NQIEEENFLRKQ
AKE81011      373   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
CUO82355      365   kvKGYYN   YINR   K    APVDEFYKFVKKCIEK-VdtPEAKQILh--DIELENFLLKQ
WP_033162887  366   klHNYLG   YIKY   D    TPVEEFYKYIKGLLAK-VdtDEAREILe--RIDLEKFMLKQ
AGZ01981      390   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
AKA60242      357   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
AKS40380      357   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
4UN5_B        361   --NGYAG   YIDG   G    ASQEEFYKFIKPILEK-M--DGTEELLv--KLNREDLLRKQ
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WP_010922251  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_039695303  423   HAILRRQGDYYPFLKE--KQD   RIEKILTFRIPYYVGPL   VRKD--SRFAWAEY---RS
WP_045635197  420   NAILRRQGEYYPFLKD--NKE   KIEKILTFRIPYYVGPL   ARGN--RDFAWLTR---NS
5AXW_A        184   KQLLKVQKAYHQLDQSfi--D   TYIDLLETRRTYYEGPG   ---Eg-SPFGWKDI-----
WP_009880683  104   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_010922251  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_011054416  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_011284745  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_011285506  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_011527619  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_012560673  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_014407541  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_020905136  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_023080005  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_023610282  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_030125963  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_030126706  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_031488318  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_032460140  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_032461047  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_032462016  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_032462936  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_032464890  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_033888930  245   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_038431314  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_038432938  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_038434062  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
BAQ51233      331   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
KGE60162            --------------------   ----------------   ------------------
KGE60856            --------------------   ----------------   ------------------
WP_002989955  420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
WP_003030002  420   HAILRRQEEHYPFLKE--NQD   KIEKILTFRIPYYVGPL   ARKG--SRFAWAEY---KA
WP_003065552  423   HAILRRQGDYYPFLKE--NQD   RIEKILTFRIPYYVGPL   ARKD--SRFSWAEY---HS
WP_001040076  421   RAIIRRQSEYYPFLKE--NLD   RIEKILTFRIPYYVGPL   AREK--SDFAWMTR---KT
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WP_001040078  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040080  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040081  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040083  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040085  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040087  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040088  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040089  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040090  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040091  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040092  421  KAIIRRQSEYYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARGN--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040094  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040095  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040096  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040097  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040098  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040099  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040100  421  RAIIRRQSEYYPLLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040104  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040105  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_001040106  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_001040107  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_001040108  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_001040109  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_001040110  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_015058523  421  KAIIRRQSEYYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARGN--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_017643650  421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_017647151  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_017648376  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_017649527  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_017771611  421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_017771984  421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
CFQ25032      421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
CFV16040      421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
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KLJ37842          421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
KLJ72361          421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
KLL20707          421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
KLL42645          421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_047207273      421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_047209694      421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_050198062      421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_050201642      421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_050204027      421  KAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_050881965      421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
WP_050886065      421  KDIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
AHN30376          421  KAIIRRQSEYYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARGN--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
EAO78426          421  KAIIRRQSEYYPFLKE--NQD  RIEKILTFRIPYYIGPL  AREK--SDFAWMTR---KTDDSIRPWNFEDLVDKEK  487
CCW42055          421  RAIIRRQSEYYPFLKE--NLD  RIEKILTFRIPYYVGPL  AREK--SDFAWMTR---KTDDSIRPWNFEELVDKEA  487
WP_003041502      420  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARKG--SRFAWAEY---KADEKITPWNFDDILDKEK  486
WP_037593752      421  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARKG--SRFAWAEY---KADEKITPWNFDDILDKEK  487
WP_049516684      421  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARKG--SRFAWAEY---KADEKITPWNFDDILDKEK  487
GAD46167          420  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARKG--SRFAWAEY---KADEKITPWNFDDILDKEK  486
WP_018363470      421  HAILRRQGDYYPFLKE--NQE  EIEKILTFRIPYYVGPL  ARKD--SRFAWAEY---RSDEKITPWNFDKVIDKEK  487
WP_003043819      430  HAILRRQEEFYPFLKE--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWLTR---KSEEAITPWNFEEVVDKGA  496
WP_006269658      420  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARKG--SRFAWAEY---KADEKITPWNFDDILDKEK  486
WP_048800889      420  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  VRKG--SRFAWAEY---KADEKITPWNFDDILDKEK  486
WP_012767106      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_014612333      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_015017095      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_015057649      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_048327215      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_049519324      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_012515931      420  HAILRRQEVFYPFLKD--NRK  KIESLLTFRIPYYVGPL  ARG-h-SRFAWVKR---KFDGAIRPWNFEEIVDEEA  486
WP_021320964      420  HAILRRQEVFYPFLKD--NRK  KIESLLTFRIPYYVGPL  ARG-h-SRFAWVKR---KFDGAIRPWNFEEIVDEEA  486
WP_037581760      420  HAILRRQEVFYPFLKD--NRK  KIESLLTFRIPYYVGPL  ARG-h-SRFAWVKR---KFDGAIRPWNFEEIVDEEA  486
WP_004232481      420  RTILRRQGEYYPFLKE--NQA  KIEKILTFRIPYYVGPL  ARKN--SRFAWAKY---HSDEPITPWNFDEVVDKEK  486
WP_009854540      421  HAILRRQGDYYPFLKE--KQD  RIEKILTFRIPYYVGPL  VRKD--SRFAWAEY---RSDEKITPWNFDKVIDKEK  487
WP_012962174      421  HAILRRQGEHYAFLKE--NQA  KIEKILTFRIPYYVGPL  ARKN--SRFAWAEY---HSDEKITPWNFDEIIDKEK  487
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WP_039695303  423  HAILRRQGDYYPFLKE--KQD  RIEKILTFRIPYYVGPL  VRKD--SRFAWAEY---RSDEKITPWNFDKVIDKEK  489
WP_014334983  420  HSILRRQGDYYPFLKE--NQA  KIEKILTFRIPYYVGPL  ARKD--SRFAWANY---HSDEPITPWNFDEVVDKEK  486
WP_003099269  420  KAIIRRQEKFYPFLKE--NQK  KIEKLFTFKIPYYVGPL  ANG-q-SSFAWLKR---QSNESITPWNFEEVVDQEA  486
AHY15608      420  KAIIRRQEKFYPFLKE--NQK  KIEKLFTFKIPYYVGPL  ANG-q-SSFAWLKR---QSNESITPWNFEEVVDQEA  486
AHY17476      420  KAIIRRQEKFYPFLKE--NQK  KIEKLFTFKIPYYVGPL  ANG-q-SSFAWLKR---QSNESITPWNFEEVVDQEA  486
ESR09100      ---  ------------------  ----------------  ----------------------------------  ---
AGM98575      420  KAIIRRQEKFYPFLKE--NQK  KIEKLFTFKIPYYVGPL  ANG-q-SSFAWLKR---QSNESITPWNFEEVVDQEA  486
ALF27331      420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYIGPL  ARGK--SDFSWLSR---KSADKITPWNFDEIVDKES  486
WP_018372492  433  QAIILNQSKYYPFLAE--NKE  KIEKILTFRIPYYVGPL  ARGN--SSFAWLQR---KSDEAIRPWNFEQVVDMET  499
WP_045618028  421  NAIIRRQGEHYPFLQE--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAR  487
WP_045635197  420  NAILRRQGEYYPFLKD--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDEAIRPWNFEEIVDKAS  486
WP_002263549  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002263887  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002264920  420  HAILRRQGDYYPFLKE--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002269043  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002269448  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002271977  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002272766  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002273241  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002275430  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002276448  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002277050  420  HAILRRQGDYYPFLKE--NQD  RIEKILTFRIPYYVGPL  ARKN--SRFAWAEY---HSDEAVMPWNFDQVIDKES  486
WP_002277364  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002279025  420  RAIIRRQSEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002279859  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002280230  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002281696  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002282247  420  HAILRRQGDYYPFLKE--NQD  RIEKILTFRIPYYVGPL  ARKN--SRFAWAEY---HSDEAVTPWNFDQVIDKES  486
WP_002282906  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002283846  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002287255  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002288990  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002289641  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ASGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002290427  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
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WP_002295753  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002296423  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002304487  430  HAILRRQGEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  VRKG--SRFAWAEY---KADEKITPWNFDDILDKEK  496
WP_002305844  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002307203  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002310390  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_002352408  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_012997688  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_014677909  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_019312892  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_019313659  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_019314093  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_019315370  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_019803776  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_019805234  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_024783594  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_024784288  420  HAILRRQGDYYPFLKE--NQD  RIEKILTFRIPYYVGPL  ARKN--SRFAWAEY---HSDEAVTPWNFDQVIDKES  486
WP_024784666  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_024784894  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ASGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_024786433  420  HAILRRQGDYYPFLKE--NQD  RIEKILTFRIPYYVGPL  ARKN--SRFAWAEY---HSDEAVTPWNFDQVIDKES  486
WP_049473442  420  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
WP_049474547  420  RAIIRRQAEFYPFLAD--NQD  RIEKLLTFRIPYYVGPL  ASGK--SDFAWLSR---KSADKITPWNFDEIVDKES  486
EMC03581      413  RAIIRRQAEFYPFLAD--NQD  RIEKILTFRIPYYVGPL  ARGK--SDFAWLSR---KSADKITPWNFDEIVDKES  479
WP_000428612  421  NAILRRQGEHYPFLKE--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAS  487
WP_000428613  421  NAILRRQGEHYPFLKD--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDEAIRPWNFEEIVDKAS  487
WP_049523028  420  NAILRHQGEYYPFLKE--NKD  KIEQILTFRIPYYVGPL  ARGN--SDFAWLSR---NSDEAIRPWNFEEMVDKSS  486
WP_003107102  389  KSIIRRQEKYYPFLKD--KQV  RIEKIFTFRIPYFVGPL  ANG-n-SSFAWVKR---RSNESITPWNFEEVVEQEA  455
WP_054279288  422  QAILERQQAYYPFLKD--NQE  KIEKILTFRIPYYIGPL  ARG-n-SRFAWLTR---TSDQKITPWNFDEMVDQEA  488
WP_049531101  421  NAILRRQGEHYPFLKE--NRE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAS  487
WP_049538452  421  NAILRRQGEHYPFLKE--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAS  487
WP_049549711  421  NAILRRQGEHYPFLKE--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAS  487
WP_007896501  422  HAILRRQEKYYPFLAE--QKE  KIEQLLCFRIPYYVGPL  AKGGn-SSFAWLKR---RSDEPITPWNFKDVVDEEA  489
EFR44625      374  HAILRRQEKYYPFLAE--QKE  KIEQLLCFRIPYYVGPL  AKGGn-SSFAWLKR---RSDEPITPWNFKDVVDEEA  441
WP_002897477  420  NAILRRQGEHYPFLKE--NRE  KIEKILTFRIPYYVGPL  ARDN--RDFSWLTR---NSDEPIRPWNFEEVVDKAR  486
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WP_002906454  420  NAILRRQGEHYLFLKE--NRE  KIEKILAFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEVVDKAS  486
WP_009729476  421  NAILRRQGEHYPFLKE--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAS  487
CQR24647      421  KAILRRQGEFYPFLKE--NAE  KIQQILTFKIPYYVGPL  ARGN--SRFAWASY---NSNEKMTPWNFDNVIDKTS  487
WP_000066813  421  NAIIRRQGEHYPFLQE--NKE  KIEKILTFRIPYYVGPL  ARGN--GDFAWLTR---NSDQAIRPWNFEEIVDQAS  487
WP_009754323  421  NAILRRQGEHYPLLKE--NKE  KIEKILTFRIPYYVGPL  ARGN--RDFAWLTR---NSDQAIRPWNFEEIVDKAS  487
WP_044674937  420  HAIIRRQAEFYPFLVE--NQD  KIEKILTFRIPYYVGPL  ARGK--SEFAWLNR---KSDEKIRPWNFDEMVDKET  486
WP_044676715  420  HAIIRRQAEFYPFLVE--NQD  KIEKILTFRIPYYVGPL  ARGK--SEFAWLNR---KSDEKIRPWNFDEMVDKET  486
WP_044680361  420  HAIIRRQAEFYPFLVE--NQD  KIEKILTFRIPYYVGPL  ARGK--SEFAWLNR---KSDEKIRPWNFDEMVDKET  486
WP_044681799  420  HAIIRRQAEFYPFLVE--NQD  KIEKILTFRIPYYVGPL  ARGK--SEFAWLNR---KSDEKIRPWNFDEMVDKET  486
WP_049533112  420  HAILRRQEEHYPFLKE--NQD  KIEKILTFRIPYYVGPL  ARKG--SRFAWAEY---KADEKITPWNFDDILDKEK  486
WP_029090905  404  VAILENQATYYPFLLE--QKD  NIHKLLTFRIPYYVGPL  ADQKd-SEFAWMVR---KQAGKITPFNFEEMVDIDA  471
WP_006506696  430  IKIIDNQAEYYPILKE--KRE  QLLSILTFRIPYYFGPL  ETSEh---AWIKRlegKENQRILPWNYQDIVDVDA  498
AIT42264      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_034440723  426  TAVLDQQEKHYSFLKE--NRD  KIISLLTFRIPYYVGPL  AKGE--SRFAWLER--sNSEEKIKPWNFDKIVDIDK  493
AKQ21048      420  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  486
WP_004636532  421  QAILDRQSQYYPFLAE--NRD  KIESLVTFRIPYYVGPL  TVSDq-SEFAWMER---QSDEPIRPWNFDEIVNKER  488
WP_002364836  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_016631044  379  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  446
EMS75795      163  KAIIERQKPYYPSLEE--ARD  KMIRLLTFRIPYYVGPL  AQGEetSSFAWLER---KTPEKVTPWNATEVIDYSA  231
WP_002373311  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_002378009  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-NTFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_002407324  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_002413717  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_010775580  430  QAIIHRQAAYYPFLKE--NQK  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  497
WP_010818269  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_010824395  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_016622645  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_033624816  428  QAIIHRQAAYYPFLKE--NQK  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_033625576  428  QAIIHRQAAYYPFLKE--NQE  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QNEKPIRPWNLQETVDLDQ  495
WP_033789179  428  QAIIHRQAAYYPFLKE--NQK  KIEQLVTFRIPYYVGPL  SKGDa-STFAWLKR---QSEEPIRPWNLQETVDLDQ  495
WP_002310644  424  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  492
WP_002312694  425  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  493
WP_002314015  425  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  493
WP_002320716  425  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  493
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WP_002330729  424  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  492
WP_002335161  425  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  493
WP_002345439  425  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  493
WP_034867970  418  KAIIDQQKQHYPFLEE--AGP  KIIALFKFRIPYYVGPL  AKEQeaSSFAWIER---KTAEKINPWNFSEVVDIEK  486
WP_047937432  425  RAIIANQKKHYPFLKE--EQE  KLESLLTFKIPYYVGPL  AKKQenSPFAWLIR---KSEEKIKPWNLPEIVDMEG  493
WP_010720994  418  KAIIDQQKQHYPFLEE--AGP  KIIALFKFRIPYYVGPL  AKEQeaSSFAWIER---KTAEKINPWNFSEVVDIEK  486
WP_010737004  418  KAIIDQQKQHYPFLEE--AGP  KIIALFKFRIPYYVGPL  AKEQeaSSFAWIER---KTAEKINPWNFSEVVDIEK  486
WP_034700478  418  KAIIDQQKQHYPFLEE--AGP  KIIALFKFRIPYYVGPL  AKEQeaSSFAWIER---KTAEKINPWNFSEVVDIEK  486
WP_007209003  422  RAILRKQEKYYSFLKE--NHE  KIEQIFKVRIPYYVGPL  AKHNeqSRFAWNIR---KSDEPIRPWNMNDVVDENA  490
WP_023519017  418  REIMDRQKRFYPFLKG--AQG  KIEKLLTFRIPYYVGPL  AQEGq-SPFAWIKR---KSPSQITPWNFAEVVDKEN  485
WP_010770040  423  EAIIQKQATYYPFLAD--NKE  EMKQLVTFRIPYYVGPL  ADGN--SPFAWLER---ISSEPIRPGNLAEVVDIKK  489
WP_048604708  419  KAILHHQAMYYPFLQE--KFS  NFVDLLTFRIPYYVGPL  ANGN--SRFSWLSR---KSDEPIRPWNLAEVVDLSK  485
WP_010750235  418  KAIIDQQKQYYPFLEK--SKE  KMIQLLTFRIPYYVGPL  AQDKetSSFAWLER---KTTEKIKPWNAKDVIDYGA  486
AII16583      459  HAILRRQEDFYPFLKD--NRE  KIEKILTFRIPYYVGPL  ARG-n-SRFAWMTR---KSEETITPWNFEEVVDKGA  525
WP_029073316  436  NKILENQSVYYSDLKD--NED  KIRSILTFRIPYYFGPL  ITKDr--QFDWIIKkegKENERILPWNANEIVDVDK  506
WP_031589969  436  NKILENQSVYYSDLKD--NED  KIRSILTFRIPYYFGPL  ITKDr--QFDWIIKkegKENERILPWNANEIVDVDK  506
KDA45870      418  VAIVENQGKYYPFLRE--NKD  KFEKILNFRIPYYVGPL  ARGN--SKFAWLTR--a-GEGKITPYNFDEMIDKET  484
WP_039099354  426  DRIIENQQQYYPWLAE-lNPN  KLDELVAFRVPYYVGPL  QQQSsdAKFAWMIR---KAEGQITPWNFDDKVDRQA  509
AKP02966      416  EKIIDNQSQYYPFLKE--NKE  KLLSILSFRIPYYVGPL  -QSSekNPFAWMER---KSNGHARPWNFDEIVDREK  483
WP_010991369  426  EAILHQQAKYYPFLKE--NYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_033838504  426  EAILHQQAKYYPFLKE--NYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
EHN60060      429  EAILHQQAKYYPFLKE--NYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  495
EFR89594      195  EAILHQQAKYYPFLKE--NYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  261
WP_038409211  426  EAILHQQAKYYPFLRK--DYE  KIRSLVTFRIPYFIGPL  ANGQ--SDFAWLTR---KADGEIRPWNIEEKVDFGK  492
EFR95520      45   EAILHQQAKYYPFLRK--DYE  KIRSLVTFRIPYFIGPL  ANGQ--SDFAWLTR---KADGEIRPWNIEEKVDFGK  111
WP_003723650  426  EAIIHQQAKYYPFLKE--DYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_003727705  426  EAILHQQAKYYPFLRE--GYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KDDGEIRPWNIEEKVDFGK  492
WP_003730785  426  EAILHQQAKYYPFLRE--GYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KDDGEIRPWNIEEKVDFGK  492
WP_003733029  426  EAILHQQAKYYPFLRE--DYE  KIKSLVTFRIPYFVGPL  AKGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_003739838  426  EAILHQQAKYYPFLKE--AYD  KIKSLVTFRIPYFVGPL  ANGQ--SDFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_014601172  426  EAIIHQQAKYYPFLRE--DYE  KIKSLVTFRIPYFVGPL  AKGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_023548323  426  EAILHQQAKYYPFLRE--DYE  KIKSLVTFRIPYFVGPL  AKGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_031665337  426  EAIIHQQAKYYTFLKE--DYD  KIKSLVTFRIPYFVGPL  ANGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
WP_031669209  426  EAILHQQAKYYPFLRE--DYE  KIKSLVTFRIPYFVGPL  AKGQ--SEFAWLTR---KADGEIRPWNIEEKVDFGK  492
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WP_033920898  426   EAIIHQQAKYYPFLRE--DYE   KIKSLVTFRIPYFVGPL   AKGQ--SEFAWLTR---KA
AKI42028      429   EAIIHQQAKYYPFLRE--DYE   KIKSLVTFRIPYFVGPL   AKGQ--SEFAWLTR---KA
AKI50529      429   EAIIHQQAKYYPFLRE--DYE   KIKSLVTFRIPYFVGPL   AKGQ--SEFAWLTR---KA
EFR83390            --------------------   -----------------   ------------------
WP_046323366  426   EAILHQQAKYYPFLKV--DYE   KIKSLVTFRIPYFVGPL   ANGQ--SEFSWLTR---KA
AKE81011      436   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
CUO82355      434   IKIIDNQAKYYPVLKE--KRE   QLLSILTFRIPYYFGPL   ETSEh----AWIKRlegKE
WP_033162887  435   IQIIDNQSVYYPQLKE--NRD   KLISILEFRIPYYFGPL   AHSE----FAWIKKfedKQ
AGZ01981      453   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
AKA60242      420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
AKS40380      420   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
4UN5_B        424   HAILRRQEDFYPFLKD--NRE   KIEKILTFRIPYYVGPL   ARG-n-SRFAWMTR---KS
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WP_010922251  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_039695303  490   SAEKFITRMTLNDLYLPEEKVLPKHSHVYETYAVYNELTKIKYVN--EQGKES-FFDSNMKQE
WP_045635197  487   SAEDFINKMTNYDLYLPEEKVLPKHSLLYETFAVYNELTKVKFIA--EGLRDYqFLDSGQKKQ
5AXW_A        230   --KEWYEMLMGHCTYFPEELRSVKYAYNADLYNALNDLNNLVITR--DENEKLeYYE---KFQ
WP_009880683  171   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_010922251  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_011054416  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_011284745  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_011285506  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_011527619  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_012560673  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_014407541  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_020905136  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_023080005  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_023610282  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_030125963  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_030126706  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_031488318  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPeFLSGEQKKA
WP_032460140  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_032461047  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_032462016  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_032462936  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_032464890  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_033888930  312   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_038431314  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_038432938  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_038434062  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
BAQ51233      398   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
KGE60162            ------------------------------------------------------------
KGE60856            ------------------------------------------------------------
WP_002989955  487   SAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVT--EGMRKPaFLSGEQKKA
WP_003030002  487   SAEKFITRMTLNDLYLPEEKVLPKHSLLYETFTVYNELTKVKYVN--EQGEAK-FFDANMKQE
WP_003065552  490   SAEKFITRMTLNDLYLPEEKVLPKHSHVYETYAVYNELTKIKYVN--EQGKDS-FFDSNMKQE
WP_001040076  488   SAEAFIHRMTNNDLYLPEEKVLPKHSLIYEKFTVYNELTKVRFLA--EGFKDFqFLNRKQKET
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WP_001040078  488  SAEAFIHRMTNNDFYLPEEKVLPKHSLIYEKFTVYNELTKVRYKN--EQGETY-FFDSNIKQEIFDGVFK--EHR-KVSK  561
WP_001040080  488  SAEAFIHRMTNNDFYLPEEKVLPKHSLIYEKFTVYNELTKVRYKN--EQGETY-FFDSNIKQEIFDGVFK--EHR-KVSK  561
WP_001040081  488  SAEAFIHRMTNNDFYLPEEKVLPKHSLIYEKFTVYNELTKVRYKN--EQGETY-FFDSNIKQEIFDGVFK--EHR-KVSK  561
WP_001040083  488  SAEAFIHRMTNNDFYLPEEKVLPKHSLIYEKFTVYNELTKVRYKN--EQGETY-FFDSNIKQEIFDGVFK--EHR-KVSK  561
WP_001040085  488  SAEAFIHRMTNNDFYLPEEKVLPKHSLIYEKFTVYNELTKVRYKN--EQGETY-FFDSNIKQEIFDGVFK--EHR-KVSK  561
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WP_001040088  488  SAEAFIHRMTNNDFYLPEEKVLPKHSLIYEKFTVYNELTKVRYKN--EQGETY-FFDSNIKQEIFDGVFK--EHR-KVSK  561
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WP_002264920  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002269043  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002269448  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002271977  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002272766  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002273241  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002275430  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002276448  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002277050  561   KKLRTFLDKN--FDEFRIVDIQGLDKEteTFNASYATYQDLLKVIKDK   VFMDNPENAEIL
WP_002277364  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002279025  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002279859  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002280230  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002281696  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002282247  561   KKLRTFLDKN--FDEFRIVDIQGLDKEteTFNASYATYQDLLKVIKDK   VFMDNPENAEIL
WP_002282906  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002283846  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002287255  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002288990  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002289641  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002290427  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
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WP_002295753  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002296423  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002304487  571   DKLLNYLNKE--FEEFRIVNLTGLDKEnkVFNSSLGTYHDLRKIL-NK   SFLDNKENEQII
WP_002305844  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_002307203  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002310390  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_002352408  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_012997688  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_014677909  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_019312892  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_019313659  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_019314093  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_019315370  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_019803776  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_019805234  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_024783594  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_024784288  561   KKLRTFLDKN--FDEFRIVDIQGLDKEteTFNASYATYQDLLKVIKDK   VFMDNPENAEIL
WP_024784666  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_024784894  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
WP_024786433  561   KKLRTFLDKN--FDEFRIVDIQGLDKEteTFNASYATYQDLLKVIKDK   VFMDNPENAEIL
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WP_049474547  561   DKLMDFLEKE--FDEFRIVDLTGLDKEnkVFNASYGTYHDLCKIL-DK   DFLDNSKNEKIL
EMC03581      554   DKLMDFLEKE--FDEFRIVDLTGLDKEnkAFNASYGTYHDLRKIL-DK   DFLDNSKNEKIL
WP_000428612  565   KDIIQYLHT---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   EFMDDPNNEEIL
WP_000428613  563   KDIIQFLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   EFMDDSKNEEIL
WP_049523028  562   KDIIHYLHN---VDGYDGIELKGIEKH---FNSSLSTYHDLLKIIKDK   EFMDDPKNEEIF
WP_003107102  531   KQLKENYFNK--IRCLDSITISGVEDK---FNASLGTYHDLLNIIKNQ   KILDDEQNQDSL
WP_054279288  564   KQLKEDFFSK--IECFDTVDISGVEDK---FNASLGTYHDLLKIIKDK   AFLDNSENENII
WP_049531101  563   KDLIHYLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   RFMDEPKNQEIL
WP_049538452  563   KDIIQYLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   EFMDDSKNEEIL
WP_049549711  563   KDIIHYLHT---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   EFMDDSKNEAIL
WP_007896501  565   KDLKEKYFSQ--IEGLENVDVTGVEGA---FNASLGTYNDLLKIIKDK   AFLDDEANAEIL
EFR44625      517   KDLKEKYFSQ--IEGLENVDVTGVEGA---FNASLGTYNDLLKIIKDK   AFLDDEANAEIL
WP_002897477  562   KDIIHYLHN---VDGYDGIELKGIEKQ---FNANLSTYHDLLKITKDK   EFMDDPKNEEIL
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WP_002906454  562   KDIIHYLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   EFMDNPKNGEIL
WP_009729476  563   KDIIQFLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   AFMDDAKNEAIL
CQR24647      562   KKILNFLDKN--FDEFRITDIQGLDNEtgNFNASYGTYHDLLKIIGDK   EFMDSSDNVDVL
WP_000066813  565   KDIIHYLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   AFMDDSKNEEIL
WP_009754323  563   KDIIHYLHN---VDGYDGIELKGIEKQ---FNASLSTYHDLLKIIKDK   EFMDNHKNQEIL
WP_044674937  562   KDIKAYL-EN--SNGYAGVELKGLEEQ---FNASLPTYHDLLKILRDK   AFIDAEENQEIL
WP_044676715  561   EKLMDFLGKE--FDEFRIVDLLGLDKDnkSFNASLGTYHDLKKIV-SK   DLLDNPENEDIL
WP_044680361  561   EKLMDFLGKE--FDEFRIVDLLGLDKDnkSFNASLGTYHDLKKIV-SK   DLLDNPENEDIL
WP_044681799  562   KDIKAYL-EN--SNGYAGVELKGLEEQ---FNASLPTYHDLLKILRDK   AFIDAEENQEIL
WP_049533112  561   DKLLNYLGKE--FDEFRIVDLTGLDKEnkVFNSSLGTYHDLRKIL-DK   SFLDNKENEQII
WP_029090905  541   TSLKKWLAEH---EHMTVSVVQGTQKEt-EFATSLQAFHRFVKIF-DR   ETVSNPANEEMF
WP_006506696  568   KKLKNWLVNNqcCS--KDAEIKGFQKEn-QFSTSLTPWIDFTNIFGKI   ----DQSNFDLI
AIT42264      562   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
WP_034440723  568   NQLVKYIENK---EQIIAPEIKGIEDS---FNSNYSTYIDLSKIPDMK   --LLEKDEDEIL
AKQ21048      562   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
WP_004636532  563   KDIANYLEQ---YGYVDGTDIKGVEDK---FNASLSTYNDLAKIDGAK   AYLDDPEYADVW
WP_002364836  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_016631044  521   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
EMS75795      306   KKLQQFLSAN--YN-IEDAEILGVDKA---FNSSYATYHDFLDLAKPN   ELLEQPEMNAMF
WP_002373311  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_002378009  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
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WP_002413717  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
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WP_010824395  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_016622645  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_033624816  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_033625576  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_033789179  570   KDIIQFYRNE--YN-TEIVTLSGLEED--QFNASFSTYQDLLKCGLTR   AELDHPDNAEKL
WP_002310644  567   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_002312694  568   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_002314015  568   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_002320716  568   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
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WP_002330729  567   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_002335161  568   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_002345439  568   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_034867970  562   KKLQNFLYTH--YH-IENAQIFGIEKA---FNASYSTYHDFMKLAKTN   EWLEQPEMEPIF
WP_047937432  568   KDLQEFLYLK--YD-IKHAELSGIEKA---FNASYTTYHDFLTMSENK   QWLEDPELASMF
WP_010720994  562   KKLQNFLYTH--YH-IENAQIFGIEKA---FNASYSTYHDFMKLAKTN   EWLEQPEMEPIF
WP_010737004  562   KKLQNFLYTH--YH-IENAQIFGIEKA---FNASYSTYHDFMKLAKTN   EWLEQPEMEPIF
WP_034700478  562   KKLQNFLYTH--YH-IENAQIFGIEKA---FNASYSTYHDFMKLAKTN   EWLEQPEMEPIF
WP_007209003  564   KKLENYLRIEl---SISSPSVKGIEEQ---FNANFGTYLDLKKFDELH   PYLDDEKYQDTL
WP_023519017  560   KQLRKFLELN--EQ-IDSTEIKGIETS---FNASYSTYHDLLKLS---   TLLDDPDMTTMF
WP_010770040  564   KLLEKFLSNE--FG-LVDVAIKGIE-T--SFNAGYGTYHDFLKIGITR   EQLDKEENSETL
WP_048604708  560   KDLSNFLRNE--YN-LDDVIIDGIE-N--KFNASFNTYHDFLKLKIDP   KVLDDPANEPMF
WP_010750235  561   KKLQHFLSAN--YN-IEDAEILGVDKV---FNSSYATYHDFLELAKPY   ELLEQPEMEEMF
AII16583      601   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
WP_029073316  576   NAMKKWLVKNqyFSNTDDIKIEGFQKEn-ACSTSLTPWIDFTKIFGEI   ----NNSNYELI
WP_031589969  576   NAMKKWLVKNqyFSNTDDIKIEGFQKEn-ACSTSLTPWIDFTKIFGKI   ----NESNYDFI
KDA45870      558   KMVIKHLKVV--MPAIRIQALKGLDNGk--FNASYGTYKDLVDMGVAP   ELLNDEVNSEKW
WP_039099354  579   KNIQDYLVSEk--RYASRPAITGLSDEnk-FNSRLSTYHDLKTIVGDA   --VDDVDKQADL
AKP02966      561   KKLTKWLIAQg---YYKNPILIGLSQKd-EFNSTLTTYLDMKKIFGSS   -FMENNKNYNQI
WP_010991369  567   KDLELFLRNM--SH-VESPTIEGLE-D--SFNSSYSTYHDLLKVGIKQ   EILDNPVNTEML
WP_033838504  567   KDLELFLRNM--SH-VESPTIEGLE-D--SFNSSYSTYHDLLKVGIKQ   EILDNPVNTEML
EHN60060      570   KDLELFLRNM--SH-VESPTIEGLE-D--SFNSSYSTYHDLLKVGIKQ   EILDNPVNTEML
EFR89594      336   KDLELFLRNM--SH-VESPTIEGLE-D--SFNSSYSTYHDLLKVGIKQ   EILDNPVNTEML
WP_038409211  567   KDLERFLYTI--NH-IESPTIEGVE-D--AFNSSFATYHDLQKGGVTQ   EILDNPLNADML
EFR95520      186   KDLERFLYTI--NH-IESPTIEGVE-D--AFNSSFATYHDLQKGGVTQ   EILDNPLNADML
WP_003723650  567   KDLELFLRNI--NH-IESPTIEGLE-D--SFNASYATYHDLLKVGMKQ   EILDNPLNTEML
WP_003727705  567   KDLELFLRNI--NH-IESPTIEGLE-D--SFNASYATYHDLLKVGLKQ   EILDNPLNTEIL
WP_003730785  567   KDLELFLRNI--NH-IESPTIEGLE-D--SFNASYATYHDLLKVGLKQ   EILDNPLNTEIL
WP_003733029  567   KDLELFLRNI--NQ-IESPTIEGLE-D--SFNASYATYHDLLKVGMKQ   EILDNPLNTEML
WP_003739838  567   KDLEQFLRNM--SH-IESPTIEGLE-D--SFNSSYATYHDLLKVGIKQ   EVLENPLNTEML
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WP_023548323  567   KDLELFLRNI--NH-VESPTIEGLE-D--SFNASYATYHDLMKVGIKQ   EILDNPLNTEML
WP_031665337  567   KDLELFLRNI--NQ-IESPTIEGLE-D--SFNASYATYHDLLKVGMKQ   EILDNPLNTEML
WP_031669209  567   KDLELFLRNI--NQ-IESPTIEGLE-D--SFNASYATYHDLLKVGMKQ   EILDNPLNTEML
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WP_033920898  567   KDLELFLRNI--NH-VESPTIEGLE-D--SFNASYATYHDLMKVGIKQ   EILDNPLNTEML
AKI42028      570   KDLELFLRNI--NH-IESPTIEGLE-D--SFNASYATYHDLLKVGMKQ   EILDNPLNTEML
AKI50529      570   KDLELFLRNI--NH-VESPTIEGLE-D--SFNASYATYHDLMKVGIKQ   EILDNPLNTEML
EFR83390      15    KDLELFLRNI--NQ-IESPTIEGLE-D--SFNASYATYHDLLKVGMKQ   EILDNPLNTEML
WP_046323366  567   KDLELFLYNM--NH-VESPTVEGVE-D--AFNSSFTTYHDLQKVGVPQ   EILDDPLNTEML
AKE81011      578   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
CUO82355      572   KKLKNWLVNNqcCR--KDAEIKGFQKEn-QFSTSLTPWIDFTNIFGKI   ----DQSNFDLI
WP_033162887  573   KKLKDWLVTHqyYDINEELKIEGYQKDl-QFSTSLAPWIDFTKIFGEI   ----NASNYQLI
AGZ01981      595   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
AKA60242      562   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
AKS40380      562   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
4UN5_B        566   KQLKEDYFKK--IECFDSVEISGVEDR---FNASLGTYHDLLKIIKDK   DFLDNEENEDIL
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WP_010922251  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_039695303  638   ERLQKYSDIFTANQLKKLER-RHYTGWGRLSYKLINGIRNK   ENNKTILDYLI    DDG-
WP_045635197  633   QRLAQYDSLFDEKVIKALTR-RHYTGWGKLSAKLINGICDK   QTGNTILDYLI    DDG-
5AXW_A        369   EELTNLNSELTQEEIEQISNlKGYTGTHNLSLKAINLILDE   ----------LW    ----
WP_009880683  318   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_010922251  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_011054416  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_011284745  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_011285506  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_011527619  634   ERLKTYAHLFDDKVMKQLKR-RRYTVWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_012560673  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_014407541  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_020905136  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_023080005  634   ERLKKYANLFDDKVMKQLKR-RHYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_023610282  634   ERLKKYANLFDDKVMKQLKR-RHYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_030125963  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_030126706  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
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WP_032460140  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
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WP_032462016  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_032462936  634   ERLKKYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_032464890  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_033888930  459   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_038431314  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_038432938  634   ERLKKYANLFDDKVMKQLKR-RHYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_038434062  634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
BAQ51233      545   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
KGE60162            ----------------------------------------   -----------   ----
KGE60856            ----------------------------------------   -----------   ----
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WP_003030002  635   QRLQKYSDIFTKAQLKKLER-RHYTGWGRLSYKLINGIRNK   ENKKTILDYLI    DDG-
WP_003065552  638   ERLQKYSDIFTADQLKKLER-RHYTGWGRLSYKLINGIRNK   ENNKTILDYLI    DDG-
WP_001040076  633   KRLDIYKDFFTESQLKKLYR-RHYTGWGRLSAKLINGIRNK   ENQKTILDYLI    DDG-
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WP_001040078  636  KRLENYKDLFTESQLKKLYR-RHYTGWGRLSAKLINGIRDK  ESQKTILDYLI  DDG---RSNRNFMQLINDDGL  704
WP_001040080  636  KRLENYKDLFTESQLKKLYR-RHYTGWGRLSAKLINGIRDK  ESQKTILDYLI  DDG---RSNRNFMQLINDDGL  704
WP_001040081  636  KRLENYKDLFTESQLKKLYR-RHYTGWGRLSAKLINGIRDK  ESQKTILDYLI  DDG---RSNRNFMQLINDDGL  704
WP_001040083  636  KRLENYKDLFTESQLKKLYR-RHYTGWGRLSAKLINGIRDK  ESQKTILDYLI  DDG---RSNRNFMQLINDDGL  704
WP_001040085  636  KRLENYKDLFTESQLKKLYR-RHYTGWGRLSAKLINGIRDK  ESQKTILDYLI  DDG---RSNRNFMQLINDDGL  704
WP_001040087  636  KRLENYKDLFTESQLKKLYR-RHYTGWGRLSAKLINGIRDK  ESQKTILDYLI  DDG---RSNRNFMQLINDDGL  704
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WP_034440723  638   RQLMKFKDKLSEKAINQLSK-KHYTGWGQLSEKLINGIRDE   QSNKTILDYLI    DNGc
AKQ21048      634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_004636532  634   KQLQTYSDTLSPEILKKLER-KHYTGWGRFSKKLINGLRDE   GSNKTILDYLK    DEGs
WP_002364836  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILGYLI    DDGv
WP_016631044  593   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLV    DDGv
EMS75795      382   TQLKKYQSVLGDGFFKKLVK-KHYTGWGRLSERLINGIRDK   KTNKTILDYLI    DDDf
WP_002373311  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLV    DDGv
WP_002378009  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLI    DDGv
WP_002407324  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLV    DDGv
WP_002413717  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLV    DDGv
WP_010775580  644   TQLSTFKGQFSEEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLI    DDGv
WP_010818269  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILGYLI    DDGv
WP_010824395  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLI    DDGv
WP_016622645  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLV    DDGv
WP_033624816  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLI    DDGv
WP_033625576  642   TQLSTFKGQFSAEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLI    DDGv
WP_033789179  642   TQLSTFKGQFSEEVLKKLER-KHYTGWGRLSKKLINGIYDK   ESGKTILDYLI    DDGv
WP_002310644  642   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
WP_002312694  643   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
WP_002314015  643   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
WP_002320716  643   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
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WP_002330729  642   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
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WP_002345439  643   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
WP_034867970  638   HQLSKYQEVFGEKLLKEFAR-KHYTGWGRFSAKLIHGIRDR   KTNKTILDYLI    DDDv
WP_047937432  643   TRLSHHEATLGKHIIKKLTK-KHYTGWGRLSKELIQGIRDK   QSNKTILDYLI    DDDf
WP_010720994  638   HQLSKYQEVFGEKLLKEFAR-KHYTGWGRFSAKLIHGIRDR   KTNKTILDYLI    DDDv
WP_010737004  638   HQLSKYQEVFGEKLLKEFAR-KHYTGWGRFSAKLIHGIRDR   KTNKTILDYLI    DDDv
WP_034700478  638   HQLSKYQEVFGEKLLKEFAR-KHYTGWGRFSAKLIHGIRDR   KTNKTILDYLI    DDDv
WP_007209003  635   NQLEQLPLNLSTKTIKALSR-RKYTGWGRLSARLIDGIHDK   NSGKTILDYLI    DESd
WP_023519017  632   EQLKPYETVLGLPAIKKLAK-KHYTGWGRLSEKMIQGMREK   QSRKTILDYLI    DDDf
WP_010770040  635   EQLKKYTYLFDEEVLKKLER-RHYTGWGRLSAKLLIGIKEK   RTHKTILDYLI    DDGg
WP_048604708  631   EQLSKFSDRLSEKTIKDLER-RHYTGWGRLSAKLINGIHDK   QSNKTILDYLI    DDAp
WP_010750235  637   TQLKKYQRILGEEIFKKLVK-KKYTGWGRLSKRLINGIRDQ   KTNKTILDYLI    DDDf
AII16583      673   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
WP_029073316  648   RRLKKEYDLDEEKIKKILKL--KYSGWSRLSKKLLSGIKTK   RTPETVLEVME    ----
WP_031589969  648   RRLKKEYDLDEEKIKKILKL--KYSGWSRLSKKLLSGIKTK   RTPETVLEVME    ----
KDA45870      631   RRLENYRDFLGEDILRKLSR-KKYTGWGRLSAKLLDGIYDK   KTHKTILDCLM    EDYs
WP_039099354  651   AKLNEIDWLTDQQRVQLAAK--RYRGWGRLSAKLLTQIVN-   ANGQRIMDLLW    ----
AKP02966      633   EKLHSSNYSYTSDQIKKISN-MRYKGWGRLSKKILTCITTE   TNTPKSLQLSN    -DLm
WP_010991369  638   EQLQQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLMGIRDK   QSHLTILDYLM    DDG-
WP_033838504  638   EQLQQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLMGIRDK   QSHLTILDYLM    DDG-
EHN60060      641   EQLQQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLMGIRDK   QSHLTILDYLM    DDG-
EFR89594      407   EQLQQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLMGIRDK   QSHLTILDYLM    DDG-
WP_038409211  638   EQLQSFSDVLDGTILKKLER-RHYTGWGRLSAKLLTGIRDK   HSHLTILDYLM    DDG-
EFR95520      257   EQLQSFSDVLDGTILKKLER-RHYTGWGRLSAKLLTGIRDK   HSHLTILDYLM    DDG-
WP_003723650  638   EQLQQFSDVLDGGVLKKLER-RHYTGWGRLSAKLLVGIREK   QSHLTILDYLM    DDG-
WP_003727705  638   EQLEQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILDYLM    DDG-
WP_003730785  638   EQLEQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILDYLM    DDG-
WP_003733029  638   EQLQQFSDVLDGTVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILDYLM    DDG-
WP_003739838  638   EQLQQFSDVLDGAVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILDYLM    DDG-
WP_014601172  638   EQLQQFSDVLDGGVLKKLER-RHYTGWGRLSAKLLVGIREK   QSHLTILDYLM    DDG-
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WP_031665337  638   EQLQQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILEYLM    DDG-
WP_031669209  638   EQLQQFSDVLDGTVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILDYLM    DDG-
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AKI42028      641   EQLQQFSDVLDGGVLKKLER-RHYTGWGRLSAKLLVGIREK   QSHLTILDYLM    DDG-
AKI50529      641   EQLQQFSDVLDGTVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILDYLM    DDG-
EFR83390      86    EQLQQFSDVLDGVVLKKLER-RHYTGWGRLSAKLLVGIRDK   QSHLTILEYLM    DDG-
WP_046323366  638   ERLQEFSNVLDEAVLKKLER-RHYTGWGRLSAKLLIGIRDK   ESHLTILDYLM    DDK-
AKE81011      650   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
CUO82355      642   RRLKKKYALPDDKIKQILKL--KYKDWSRLSKKLLDGIVAD   SV--TVLDVLE    ----
WP_033162887  645   RRLKKVYQLDDLLVDKILKL--NYTGWSRLSEKLLTGMTAD   KA--TVLFVLE    ----
AGZ01981      667   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
AKA60242      634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
AKS40380      634   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
4UN5_B        638   ERLKTYAHLFDDKVMKQLKR-RRYTGWGRLSRKLINGIRDK   QSGKTILDFLK    -DGf
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WP_039695303  707   PFKQIIQKSQVVG-DVDD-IEAVVHDLPGSPAIKKGILQSVKIVDELVKVMG-GNPDNIVIEM
WP_045635197  702   SFKEIIQKAQVIG-KTDD-VKQVVQELSGSPAIKKGILQSIKIVDELVKVMG-HAPESIVIEM
5AXW_A        427   VPKKVDLSQQKEI---PT---TLVDDFILSPVVKRSFIQSIKVINAIIKKYG--LPNDIIIEL
WP_009880683  387   TFKEDLQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKVVDELVKVMGrHKPENIVIEM
WP_010922251  703   TFKEDIQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKVVDELVKVMGrHKPENIVIEM
WP_011054416  703   TFKEDIQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKVVDELVKVMGrHKPENIVIEM
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WP_032464890  703   TFKEDIQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKVVDELVKVMGrHKPENIVIEM
WP_033888930  528   TFKEDIQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKVVDELVKVMGrHKPENIVIEM
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WP_038434062  703   TFKEDIQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKIVDELVKVMGrHKPENIVIEM
BAQ51233      614   TFKEDIQKAQVSG-QGDS-LHEHIANLAGSPAIKKGILQTVKVVDELVKVMGrHKPENIVIEM
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KGE60856            ------------------------------------------------------------
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WP_003065552  707   PFKQIIQKSQVVG-DVDD-IEAVVHDLPGSPAIKKGILQSVKIVDELVKVMG-DNPDNIVIEM
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WP_001040080	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040081	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040083	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040085	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040087	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040088	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
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WP_001040091	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040092	705	SFKSIISKAQSGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
WP_001040094	702	SFKPIIDKARTGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	775	TTAKGLSRS	775
WP_001040095	702	SFKPIIDKARTGS-HLDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	775	TTAKGLSRS	775
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WP_001040097	702	SFKPIIDKARTGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	775	TTAKGLSRL	775
WP_001040098	702	SFKPIIDKARTGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	775	TTAKGLSRS	775
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WP_001040104	705	SFKSIISKAQAGS-HSDN-LKEVVGELAGSPAIKKGILQSLKIVDELVKVMG-YEPEQIVVEMARENQ	778	TTNQGRRNS	778
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WP_045635197	776	QQRYKRIEDSLK	ILAS	NILKENP--TD---NNQLQNDRLFLYYLQNGKDMYTGEAL--D--INQLSSYDIDHI	843
5AXW_A	488	KDAQKMINEMQK	QTNE	EIIRTTGk--E---NAKYLIEKIKLHDMQEGKCLYSLEAIplEdlNNPFNYEVDHI	561
WP_009880683	462	RERMKRIEEGIK	ELGS	DILKEYP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	525
WP_010922251	778	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_011054416	778	RERMKRIEEGIK	ELGS	DILKEYP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_011284745	778	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_011285506	778	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_011527619	778	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_012560673	778	RERMKRIEEGIK	ELGS	DILKEYP--VE---TTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_014407541	777	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	840
WP_020905136	778	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
WP_023080005	777	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	840
WP_023610282	777	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	840
WP_030125963	778	RERMKRIEEGIK	ELGS	DILKEYP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	841
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WP_033888930	603	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	666
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BAQ51233	689	RERMKRIEEGIK	ELGS	QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI	752
KGE60162	1	------------	-----	----------------------------QEL--D--INRLSGYDVDHI	16
KGE60856		------------	-----	--	
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WP_003065552	781	QQRLKKLQNSLK	PSYI	E----DK--VE---NSHLQNDQLFLYYIQNGKDMYTGDEL--D--IDHLSDYDIDHI	851
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KLL20707          793  RQRYKLLDDGVK  NLAS  NILKEYP--TD---NQALQNERLFLYYLQNGRDMYTGEAL--D--IDNLSQYDIDHI  860
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ESR09100      ---  ----------  ----  ----------  ---  ----------  ----------  ----------  ---
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WP_019313659 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_019314093 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_019315370 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_019803776 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_019805234 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_024783594 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_024784288 779 QQRYKRLKEAIK DLNH KILKEHP--TD---NQALQNNRLFLYYLQNGRDMYTGESL--D--INRLSDYDIDHV 846
WP_024784666 778 QQRLKGLTDSIK EFGS QILKEHP--VE---HSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_024784894 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_024786433 779 QQRYKRLKEAIK DLNH KILKEHP--TD---NQALQNNRLFLYYLQNGRDMYTGESL--D--INRLSDYDIDHV 846
WP_049473442 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
WP_049474547 778 QQRLKGLTDSIK EFGS QILKEHP--VE---NSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 841
EMC03581     771 QQRLKGLTDSIK EFGS QILKEHP--VE---HSQLQNDRLFLYYLQNGRDMYTGEEL--D--IDYLSQYDIDHI 834
WP_000428612 779 QQRYKRIEDSLK ILAS KILKEHP--TD---NIQLQNDRLFLYYLQNGRDMYTGKPL--D--INQLSSYDIDHI 846
WP_000428613 777 QQRYKRIEDALK NLAS NILKEHP--TN---NIQLQNDRLFLYYLQNGRDMYTGKPL--D--INQLSSYDIDHI 844
WP_049523028 776 QQRLKTLSDAIS ELG- NILKEHP--TD---NIQLQNDRLFLYYLQNGKDMYTGEAL--D--INQLSNYDIDHI 839
WP_003107102 747 RERLRKLEEVHK NIGS KILKEHE--IS---NAQLQSDRVYLYLLQDGKDMYTGKDL--D--FDRLSQYDIDHI 810
WP_054279288 779 RERMKRVQEVLK KLGS QLLKEHP--VE---NFQLQNERLYLYYLQNGKDMYTGEEL--S--ISNLSHYDIDHI 842
WP_049531101 777 QQRYKRIEDSLK ILAS NILKEHP--TD---NIQLQNDRLFLYYLQNGKDMYTGNPL--D--INHLSSYDIDHI 844
WP_049538452 777 QQRYKRIENSLK ILAS KILKEHP--TD---NNQLQNDRLFLYYLQNGKDMYTGEAL--D--INQLSSCDIDHI 844
WP_049549711 777 QQRYKRIEDSLK ILAS NILKENP--TD---NNQLQNDRLFLYYLQNGKDMYTGEAL--D--INQLSSYDIDHI 844
WP_007896501 781 RQRLKKIKEVHK KTGS RILEDNSerIT---NLTLQDNRLYLYLLQDGKDMYTGQDL--D--INNLSQYDIDHI 846
EFR44625     733 RQRLKKIKEVHK KTGS RILEDNSerIT---NLTLQDNRLYLYLLQDGKDMYTGQDL--D--INNLSQYDIDHI 798
WP_002897477 776 QQRYKRIEDALK NLAP NILKENP--TD---NIQLKNDRLFLYYLQNGKDMYTGKPL--D--INQLSSYDIDHI 843
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WP_002906454  776  QQRYKRIEDALK  NLAP  NILKENP--TD---NIQLQNDRLFLYYLQNGKDMYTGKAI--D--INQLSNYDIDHI  843
WP_009729476  777  QQRYKRIEDSLK  ILAS  KILKEHP--TD---NIQLQNDRLFLYYLQNGKDMYTGEAL--D--INQLSSCDIDHI  844
CQR24647      780  QQRLGSLTKAIQ  DFGS  DILKRYP--VE---NNQLQNDQLYLYYLQNGKDMYTGDTL--D--IHNLSQYDIDHI  843
WP_000066813  779  QQRYKRIEDSLK  NLAS  NILKENP--TD---NIQLQNDRLFLYYLQNGRDMYTGKPL--E--INQLSNYDIDHI  846
WP_009754323  777  QQRYKRIEDALK  NLAP  TISKENP--TD---NIQLQNDRLFLYYLQNGKDMYTGEAL--D--INQLSSYDIDHI  844
WP_044674937  776  QQRYKKIENAIK  NLNS  KILKEYP--TN---NQALQNDRLFLYYLQNGKDMYTDEEL--D--IDQLSQYDIDHI  843
WP_044676715  778  QQRYKKIENAIK  NLNS  KILKEYP--TN---NQALQNDRLFLYYLQNGKDMYTDEEL--D--IDQLSQYDIDHI  845
WP_044680361  778  QQRYKKIENAIK  NLNS  KILKEYP--TN---NQALQNDRLFLYYLQNGKDMYTDEEL--D--IDQLSQYDIDHI  845
WP_044681799  776  QQRYKKIENAIK  NLNS  KILKEYP--TN---NQALQNDRLFLYYLQNGKDMYTDEEL--D--IDQLSQYDIDHI  843
WP_049533112  778  QQRLKLLQDSLK  PVNI  K-----N--VE---NQQLQNDRLFLYYIQNGKDMYTGETL--D--INNLSQYDIDHI  840
WP_029090905  753  TPRDKFIEKAYA  ETDT  EHLKELK---Qr--SKQLSSQRLFLYFIQNGKCMYSGEHL--D--IERLDSYEVDHI  823
WP_006506696  777  ESKIKKLENVYK  DEQT  SVLEELKg-FDn--TKKISSDSLFLYFTQLGKCMYSGKKL--D--IDSLDKYQIDHI  849
AIT42264      778  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI  841
WP_034440723  785  KARLKKIQEGLE  NLDS  HVEKQAL---D---EEMLKSPKYYLYCLQNGKDIYTGKDL--D--IGQLQTYDIDHI  848
AKQ21048      778  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI  841
WP_004636532  781  KERLEKLTEAIK  EFDG  --VKVKD--LK---NENLRNDRLYLYYLQNGRDMYTNEPL--D--INNLSKYDIDHI  845
WP_002364836  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_016631044  740  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  803
EMS75795      525  KPRLKALEEALK  SFDS  PLLKEQP--VD---NQALQKDRLYLYYLQNGKDMYTGEAL--D--IDRLSEYDIDHI  588
WP_002373311  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_002378009  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_002407324  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_002413717  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_010775580  791  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  854
WP_010818269  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_010824395  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_016622645  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_033624816  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_033625576  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_033789179  789  IQRLKIVEKAMA  EIGS  NLLKEQP--TT---NEQLRDTRLFLYYMQNGKDMYTGDEL--S--LHRLSHYDIDHI  852
WP_002310644  789  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  852
WP_002312694  790  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  853
WP_002314015  790  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  853
WP_002320716  790  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  853
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WP_002330729  789  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  852
WP_002335161  790  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  853
WP_002345439  790  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  853
WP_034867970  781  SPRLKALENGLK  QIGS  TLLKEQP--TD---NKALQKERLYLYYLQNGRDMYTGEPL--E--IENLHQYEVDHI  844
WP_047937432  790  RPRLKALEESLK  DFGS  QLLKEYP--TD---NSSLQKDRLYLYYLQNGRDMYTGAPL--D--IHRLSDYDIDHI  853
WP_010720994  781  KPRLKALENGLK  QIGS  TLLKEQP--TD---NKALQKERLYLYYLQNGRDMYTGEPL--E--IENLHQYEVDHI  844
WP_010737004  781  SPRLKALENGLK  QIGS  TLLKEQP--TD---NKALQKERLYLYYLQNGRDMYTGEPL--E--IENLHQYEVDHI  844
WP_034700478  781  KPRLKALENGLK  QIGS  TLLKEQP--TD---NKALQKERLYLYYLQNGRDMYTGEPL--E--IENLHQYEVDHI  844
WP_007209003  782  KPRLKGIENGLK  EFSD  SVLKGSS--ID---NKQLQNDRLYLYYLQNGKDMYTGHEL--D--IDHLSTYDIDHI  845
WP_023519017  775  RPRLKALEEALK  NIDS  PLLKDYP--TD---NQALQKDRLYLYYLQNGKDMYTGEPL--E--IHRLSEYDIDHI  838
WP_010770040  782  NPRMKALEEAMR  NLRS  NLLKEYP--TD---NQALQNDRLYLYYLQNGKDMYTGLDL--S--LHNLSSYDIDHI  845
WP_048604708  779  RPRLKNLEKAID  DLDS  EILKKHP--VD---NKALQKDRLYLYYLQNGKDMYTNEEL--D--IHKLSTYDIDHI  842
WP_010750235  784  KPRLKSLEEALK  NFDS  QLLKERP--VD---NQSLQKDRLYLYYLQNGKDMYTGESL--D--IDRLSEYDIDHI  847
AII16583      817  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI  880
WP_029073316  789  DSFVNQMLKLYK  DFED  EANKHLKg-EDa--KSKIRSERLKLYYTQMGKCMYTGKSL--D--IDRLDTYQVDHI  860
WP_031589969  789  DSFVNQMLKLYK  DFED  EANKHLKg-EDa--KSKIRSERLKLYYTQMGKCMYTGKSL--D--IDRLDTYQVDHI  860
KDA45870      772  EDRVQQIVKNLK  ELPK  ------P---S---NAELSDERKYLYCLQNGRDMYTGAPL--D--YDHLQFYDVDHI  833
WP_039099354  789  KRQVEQVYQNIS  EL--  EIRNELK---Dl-sNSALSNTRLFLYFMQGGRDMYTGDSL--N--IDRLSTYDIDHI  856
AKP02966      786  RLQSKLLNKANG  -LVP  EELKKHKn--D-----LSSERIMLYFLQNGKSLYSEESL--N--INKLSDYQVDHI  858
WP_010991369  781  RPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELRNNRLYLYYLQNGKDMYTGQDL--D--IHNLSNYDIDHI  844
WP_033838504  781  RPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELRNNRLYLYYLQNGKDMYTGQDL--D--IHNLSNYDIDHI  844
EHN60060      784  RPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELRNNRLYLYYLQNGKDMYTGQDL--D--IHNLSNYDIDHI  847
EFR89594      550  RPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQDL--D--IHNLSNYDIDHI  613
WP_038409211  781  KPRFISLEKAIK  EFGS  QILKEHP--TD---NQCLKNDRLYLYYLQNGKDMYTGKEL--D--IHNLSNYDIDHI  844
EFR95520      400  KPRFISLEKAIK  EFGS  QILKEHP--TD---NQCLKNDRLYLYYLQNGKDMYTGKEL--D--IHNLSNYDIDHI  463
WP_003723650  781  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
WP_003727705  781  KPRYKSLEKAIK  DFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDIYTGQEL--D--IHNLSNYDIDHI  844
WP_003730785  781  KPRYKSLEKAIK  DFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDIYTGQEL--D--IHNLSNYDIDHI  844
WP_003733029  781  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
WP_003739838  781  RPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELRNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
WP_014601172  781  KPRYKSLEKAIK  EFGS  KILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
WP_023548323  781  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
WP_031665337  781  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
WP_031669209  781  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
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WP_033920898  781  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  844
AKI42028      784  KPRYKSLEKAIK  EFGS  KILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  847
AKI50529      784  KPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHI  847
EFR83390      229  RPRYKSLEKAIK  EFGS  QILKEHP--TD---NQELKNNRLYLYYLQNGKDIYTGQEL--D--IHNLSNYDIDHI  292
WP_046323366  781  KPRFTSLEKAIK  ELGS  QILKEHP--TD---NQGLKNDRLYLYYLQNGKDMYTGQEL--D--IHNLSNYDIDHV  844
AKE81011      794  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI  857
CUO82355      781  ESKIKKLENVYK  DEQT  SVLEELKg-FDn--TKKISSDSLFLYFTQLGKCMYSGKKL--D--IDSLDKYQIDHI  853
WP_033162887  784  ESKIAKLQKIYE  NLQT  QVYESLKk-EDa--KKRMETDALYLYYLQMGKSMYSGKPL--D--IDKLSTYQIDHI  855
AGZ01981      811  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI  874
AKA60242      778  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDAI  841
AKS40380      778  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDHI  841
4UN5_B        782  RERMKRIEEGIK  ELGS  QILKEHP--VE---NTQLQNEKLYLYYLQNGRDMYVDQEL--D--INRLSDYDVDAI  845
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WP_010922251  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_039695303  852  IPQAFIKDDSIDNRVLTSSAKNRG-KSDD--VP  S--LDIVRARKA-EWVRLYKSGLISKRKFDNLTKA--ERGGLTE  920
WP_045635197  844  IPQAFIKDDSLDNRVLTSSKDNRG-KSDN--VP  S--IEVVQKRKA-FWQQLLDSKLISERKFNNLTKA--ERGGLDE  912
5AXW_A        562  IPRSVSFDNSFNNKVLVKQEEASK-KGNR--TP  Fqy-LSSSDSKI-SYETFKKHILNLAKGKGRISKTk-KEYLLEE  632
WP_009880683  526  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWKQLLNAKLITQRKFDNLTKA--ERGGLSE  594
WP_010922251  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_011054416  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_011284745  842  VPQSFIKDDSIDNKVLTRSDKNRG-KSNN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_011285506  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_011527619  842  VPQSFIKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_012560673  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWKQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_014407541  841  VPQSFIKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  909
WP_020905136  842  VPQSFIKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
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WP_023610282  841  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  909
WP_030125963  842  VPQSFIKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
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WP_033888930  667  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  735
WP_038431314  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
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WP_038434062  842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWKQLLNAKLITQRKFDNLTKA--ERGGLSE  910
BAQ51233      753  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  821
KGE60162       17  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE   85
KGE60856           ----------------------------------  ----------------------------------------
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WP_003030002  841  IPQAFIKDNSLDNRVLTRSDKNRG-KSDD--VP  S--IEVVHEMKS-FWSKLLSVKLITQRKFDNLTKA--ERGGLTE  909
WP_003065552  852  IPQAFIKDDSIDNRVLTSSAKNRG-KSDD--VP  S--LDIVRARKA-EWVRLYKSGLISKRKFDNLTKA--ERGGLTE  920
WP_001040076  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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WP_001040080  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040081  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040083  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040085  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040087  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040088  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040089  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040090  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040091  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040092  847  VPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--IDIVKARKA-FWKKLLDAKLISQRKYDNLTKA--ERGGLTP  915
WP_001040094  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--VEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
WP_001040095  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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WP_001040105  847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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CFQ25032      847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
CFV16040      847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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KLJ72361          847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
KLL20707          861  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  929
KLL42645          847  VPQAFIKDDSIDNRVLVSSAKNRG-KSDN--VP  S--IDIVKARKA-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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WP_050198062      847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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EAO78426          847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
CCW42055          847  IPQAFIKDDSIDNRVLVSSAKNRG-KSDD--VP  S--LEIVKDCKV-FWKKLLDAKLMSQRKYDNLTKA--ERGGLTS  915
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GAD46167          841  IPQAFIKDNSLDNRVLTRSDKNRG-KSDD--VP  S--IEVVHEMKS-FWSKLLSVKLITQRKFDNLTKA--ERGGLTE  909
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WP_009854540      850  IPQAFIKDDSIDNRVLTSSAKNRG-KSDD--VP  S--LDIVRARKA-EWVRLYKSGLISKRKFDNLTKA--ERGGLTE  918
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WP_014334983  849  IPQAFIKDNSIDNKVLTSSAKNRG-KSDD--VP  S--IETVRNRRS-YWYKLYKSGLISKRKFDNLTKA--ERGGLTE  917
WP_003099269  842  IPQSFIKDNSIDNTVLTTQASNRG-KSDN--VP  N--IETVNKMKS-FWYKQLKSGAISQRKFDHLTKA--ERGALSD  910
AHY15608      842  IPQSFIKDNSIDNTVLTTQASNRG-KSDN--VP  N--IETVNKMKS-FWYKQLKSGAISQRKFDHLTKA--ERGALSD  910
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ESR09100           ----------------------------  ----------------------------------------
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ALF27331      842  IPQAFIKDNSIDNRVLTSSKENRG-KSDD--VP  S--KDVVRKMKS-YWSKLLSAKLITQRKFDNLTKA--ERGGLTD  910
WP_018372492  858  VPRSYIKNDSFDNKVLTTSKGNRK-KLDD--VP  A--KEVVEKMEN-TWRRLHAAGLISDIKLSYILMKGe-----LTE  923
WP_045618028  845  IPQAFIKDDSLDNRVLTSSKDNRG-KSDN--VP  S--LEIVQKRKA-FWQQLLDSKLISERKFNNLTKA--ERGGLDE  913
WP_045635197  844  IPQAFIKDDSLDNRVLTSSKDNRG-KSDN--VP  S--IEVVQKRKA-FWQQLLDSKLISERKFNNLTKA--ERGGLDE  912
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WP_054279288  843  IPRSFIKDDSIDNKVLTRSEHNRG-KTDN--VP  S--IEVVKRMKP-YWQKLLDTKVISQRKFDNLTKA--ERGGLQE  911
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WP_049549711  845  IPQAFIKDDSLDNRVLTSSKDNRG-KSDN--VP  S--LEVVQKRKA-FWQQLLDSKLISERKFNNLTKAerERDGLNE  915
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WP_044674937  844  IPQAFIKDDSLDNKVLTKSAKNRG-KSDD--VP  S--LEIVHKKKN-FWKQLLDSQLISQRKFDNLTKA--ERGGLTN  912
WP_044676715  846  IPQAFIKDDSLDNKVLTKSAKNRG-KSDD--VP  S--LEIVHKKKN-FWKQLLDSQLISQRKFDNLTKA--ERGGLTN  914
WP_044680361  846  IPQAFIKDDSLDNKVLTKSAKNRG-KSDD--VP  S--LEIVHKKKN-FWKQLLDSQLISQRKFDNLTKA--ERGGLTN  914
WP_044681799  844  IPQAFIKDDSLDNKVLTKSAKNRG-KSDD--VP  S--LEIVHKKKN-FWKQLLDSQLISQRKFDNLTKA--ERGGLTN  912
WP_049533112  841  IPQAFIKDDSFDNRVLTSSSENRG-KSDN--VP  S--IEVVRARKA-DWMRLRKAGLISQRKFDNLTKA--ERGGLTE  909
WP_029090905  824  LPQSYIKDNSIENLALVKKVENQR-KKDSllLN  S---SIINQNYS-RWEQLKNAGLIGEKKFRNLTRTk-----ITD  890
WP_006506696  850  VPQSLVKDDSFDNRVLVVPSENQR-KLDDlvVP  ---FDIRDKMYR-FWKLLFDHELISPKKFYSLIKTe-----YTE  916
AIT42264      842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_034440723  849  IPRSFTTDNSFDNLVLTSSTVNRG-KLDN--VP  Sp--DIVRQQKG-FWKQLLRAGLMSQRKFNNLTKGk-----LTD  914
AKQ21048      842  VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP  S--EEVVKKMKN-YWRQLLNAKLITQRKFDNLTKA--ERGGLSE  910
WP_004636532  846  IPQSFTTDNSIDNKVLVSRTKNQGnKSDD--VP  S--INIVHKMKP-FWRQLHKAGLISDRKFKNLTKA--EHGGLTE  915
WP_002364836  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_016631044  804  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  872
EMS75795      589  IPRSFIVDNSIDNKVLVSSKENRL-KMDD--VP  D--QKVVIRMRR-YWEKLLRANLISERKFAYLTKLe-----LTP  654
WP_002373311  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KKVVKKMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_002378009  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_002407324  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_002413717  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_010775580  855  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKKMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  923
WP_010818269  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_010824395  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_016622645  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKDMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_033624816  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKKMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_033625576  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKKMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_033789179  853  IPQSFMKDDSLDNLVLVGSTENRG-KSDD--VP  S--KEVVKKMKA-YWEKLYAAGLISQRKFQRLTKG--EQGGLTL  921
WP_002310644  853  IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP  S--EKVVKKMRS-FWYDLYSSKLISKRKLDNLTKIk-----LTE  918
WP_002312694  854  IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP  S--EKVVKKMRS-FWYDLYSSKLISKRKLDNLTKIk-----LTE  919
WP_002314015  854  IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP  S--EKVVKKMRS-FWYDLYSSKLISKRKLDNLTKIk-----LTE  919
WP_002320716  854  IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP  S--EKVVKKMRS-FWYDLYSSKLISKRKLDNLTKIk-----LTE  919
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WP_002330729  853   IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP   S--EKVVKKMRS-FWYDLYSSKLISKR
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WP_002345439  854   IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP   S--EKVVKKMRS-FWYDLYSSKLISKR
WP_034867970  845   IPRSFIVDNSIDDKVLVASKQNQK-KRDD--VP   K--KQIVNEQRI-FWNQLKEAKLISTK
WP_047937432  854   IPRSFTTDNSIDNKVLVSSKENRL-KKDD--VP   S--EKVVKKMRS-FWYDLYSSKLISKR
WP_010720994  845   IPRSFIVDNSIDNKVLVASKQNQK-KRDD--VP   K--KQIVNEQRI-FWNQLKEAKLISPK
WP_010737004  845   IPRSFIVDNSIDNKVLVASKQNQK-KRDD--VP   K--KQIVNEQRI-FWNQLKEAKLISPK
WP_034700478  845   IPRSFIVDNSIDNKVLVASKQNQK-KRDD--VP   N--KQIVNEQRI-FWNQLKEAKLISPK
WP_007209003  846   IPQSFLTDNSIDNRVLTTSKSNRG-KSDN--VP   S--EEVVRKMDR-FWRKLLNAKLISER
WP_023519017  839   IPRSFIVDNSLDNKVLVSSKVNRG-KLDN--AP   D--PLVVKRMRS-HWEKLHQAKLISDK
WP_010770040  846   VPQSFTTDNSLDNRVLVSSKENRG-KKDD--VP   S--KEVVQKNIT-LWETLKNSNLISQK
WP_048604708  843   IPQSFIVDNSLDNRVLVSSSKNRG-KLDD--VP   S--KEVVKKMRA-FWESLYRSGLISKK
WP_010750235  848   IPRSFIVDHSLDNKVLVSSKENRL-KKDD--VP   D--SKVVKRMKA-YWEKLLRANLISER
AII16583      881   VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP   S--EEVVKKMKN-YWRQLLNAKLITQR
WP_029073316  861   VPQSLLKDDSIDNKVLVLSSENQR-KLDDlvIP   ---EMIRNKMFG-FWNKLYENKIISPK
WP_031589969  861   VPQSLLKDDSIDNKVLVLSSENQR-KLDDlvIP   ---SSIRNKMYG-FWEKLFNNKIISPK
KDA45870      834   IPQSFLKDDSIENKVLTIKKENVR-KTNG--LP   S--EAVIQKMGS-FWKKLLDAGAMTNK
WP_039099354  857   LPQSFIKDNSLDNRVLVSQRMNRS-KADQ--VP   S--VELGQKMQI-QWEQMLRAGLITKK
AKP02966      859   LPRTYIPDDSLENKALVLAKENQR-KADDllLN   S---NVIDKNLE-RWTYMLNNNMMGLK
WP_010991369  845   VPQSFITDNSIDNLVLTSSAGNRE-KGDD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_033838504  845   VPQSFITDNSIDNLVLTSSAGNRE-KGDD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
EHN60060      848   VPQSFITDNSIDNLVLTSSAGNRE-KGDD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
EFR89594      614   VPQSFITDNSIDNLVLTSSAGNRE-KGND--VP   P--LEIVQKRKV-FWEKLYQGNLMSKR
WP_038409211  845   IPQSFITDNSIDNRVLVSSTANRE-KGDN--VP   L--LEVVRKRKA-FWEKLYQAKLMSKR
EFR95520      464   IPQSFITDNSIDNRVLVSSTANRE-KGDN--VP   L--LEVVRKRKA-FWEKLYQAKLMSKR
WP_003723650  845   VPQSFITDNSIDNLVLTSSAGNRE-KGGD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_003727705  845   VPQSFITDNSIDNLVLTSSAGNRE-KGGD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_003730785  845   VPQSFITDNSIDNLVLTSSAGNRE-KGGD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_003733029  845   VPQSFITDNSVDNLVLTSSAGNRE-KGDN--VP   P--LEIVQKRKI-FWEKLYQGNLMSKR
WP_003739838  845   VPQSFITDNSIDNLVLTSSAGNRE-KGDD--VP   P--LEIVRKRKV-FWEKLFQGNLMSKR
WP_014601172  845   VPQSFITDNSIDNLVLTSSAGNRE-KGGD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_023548323  845   VPQSFITDNSIDNLVLTSSAGNRE-KGDN--VP   P--LEIVQKRKI-FWEKLYQGNLMSKR
WP_031665337  845   VPQSFITDNSIDNLVLTSSAGNRE-KGGD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_031669209  845   VPQSFITDNSVDNLVLTSSAGNRE-KGDN--VP   P--LEIVQKRKI-FWEKLYQGNLMSKR
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WP_033920898  845   VPQSFITDNSIDNLVLTSSAGNRE-KGDN--VP   P--LEIVQKRKI-FWEKLYQGNLMSKR
AKI42028      848   VPQSFITDNSIDNLVLTSSAGNRE-KGGD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
AKI50529      848   VPQSFITDNSIDNLVLTSSAGNRE-KGDN--VP   P--LEIVQKRKI-FWEKLYQGNLMSKR
EFR83390      293   VPQSFITDNSIDNLVLTSSAGNRE-KGDD--VP   P--LEIVRKRKV-FWEKLYQGNLMSKR
WP_046323366  845   VPQSFITDNSIDNRVLASSAANRE-KGDN--VP   S--LEVVRKRKV-YWEKLYQAKLMSKR
AKE81011      858   VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP   S--EEVVKKMKN-YWRQLLNAKLITQR
CUO82355      854   VPQSLVKDDSFDNRVLVLPSENQR-KLDDlvVP   ---FDIRDKMYR-FWKLLFDHELISPK
WP_033162887  856   LPQSLIKDDSFDNRVLVLPEENQW-KLDSetVP   ---FEIRNKMIG-FWQMLHENGLMSNK
AGZ01981      875   VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP   S--EEVVKKMKN-YWRQLLNAKLITQR
AKA60242      842   VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP   S--EEVVKKMKN-YWRQLLNAKLITQR
AKS40380      842   VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP   S--EEVVKKMKN-YWRQLLNAKLITQR
4UN5_B        846   VPQSFLKDDSIDNKVLTRSDKNRG-KSDN--VP   S--EEVVKKMKN-YWRQLLNAKLITQR
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WP_010922251  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_039695303  921   AD   KAGFIKRQLVETRQITKHVAQILDARFNTEHDENDKVIR--DVKVITLKSNLVSQFRK
WP_045635197  913   RD   KVGFIKRQLVETRQITKHVAQILDARYNTEVNEKDKKNR--TVKIITLKSNLVSNFRK
5AXW_A        633   RD   QKDFINRNLVDTRYATRGLMNLLRSYFR---------VNnlDVKVKSINGGFTSFLRR
WP_009880683  595   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVRVITLKSKLVSDFRK
WP_010922251  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_011054416  911   LD   KVGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVRVITLKSKLVSDFRK
WP_011284745  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_011285506  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_011527619  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_012560673  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVRVITLKSKLVSDFRK
WP_014407541  910   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_020905136  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_023080005  910   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_023610282  910   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_030125963  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_030126706  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_031488318  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
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WP_032462936  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVRVITLKSKLVSDFRK
WP_032464890  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
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WP_038431314  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_038432938  910   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_038434062  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
BAQ51233      822   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
KGE60162      86    LD   KVGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVRVITLKSKLVSDFRK
KGE60856            --   ----------------------------------------------------------
WP_002989955  911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_003030002  910   ED   KAGFIKRQLVETRQITKHVAQILDERFNTEFDGNKRRIR--NVKIITLKSNLVSNFRK
WP_003065552  921   AD   KAGFIKRQLVETRQITKHVAQILDARFNTESDENDKVIR--DVKVITLKSNLVSQFRK
WP_001040076  916   DD   KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRK
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CFQ25032      916   DD   KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRK
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KLJ37842          916  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTVKSNLVSNFRKEF  GFYKIREVNNY  986
KLJ72361          916  DD  KARFIQRQLVETRQITKHVARILDELFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNNY  986
KLL20707          930  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNNY  1000
KLL42645          916  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNDY  986
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WP_047209694      916  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNNY  986
WP_050198062      916  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNNY  986
WP_050201642      916  DD  KARFIQRQLVETRQITKHVASILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNNY  986
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EAO78426          916  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNNY  986
CCW42055          916  DD  KARFIQRQLVETRQITKHVARILDERFNNELDSKGRRIR--KVKIVTLKSNLVSNFRKEF  GFYKIREVNDY  986
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GAD46167          910  ED  KAGFIKRQLVETRQITKHVAQILDERFNTEFDGAQRRIR--NVKIITLKSNLVSNFRKEF  ELYKVREINDY  980
WP_018363470      919  AD  KAGFIKRQLVETRQITKHVAQILDARFNTERDENDKVIR--DVKVITLKSNLVSQFRKEF  KFYKVREINDY  989
WP_003043819      920  AD  KAGFIKRQLVETRQITKHVARILDSRMNTKRDKNDKPIR--EVKVITLKSKLVSDFRKDF  QLYKVRDINNY  990
WP_006269658      910  ED  KAGFIKRQLVETRQITKHVAQILDERFNTEFDGNKRRIR--NVKIITLKSNLVSNFRKEF  ELYKVREINDY  980
WP_048800889      910  ND  KAGFIHRQLVETRQITKHVAQILDARFNPKRDDNKKVIR--DVKIITLKSNLVSQFRRDF  KLYKVREINDY  980
WP_012767106      910  LD  KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRKDF  QFYKVREINNY  980
WP_014612333      910  LD  KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRKDF  QFYKVREINNY  980
WP_015017095      910  LD  KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRKDF  QFYKVREINNY  980
WP_015057649      910  LD  KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRKDF  QFYKVREINNY  980
WP_048327215      910  LD  KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRKDF  QFYKVREINNY  980
WP_049519324      910  LD  KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRKDF  QFYKVREINNY  980
WP_012515931      910  VD  KAGFIQRQLVETRQITKHVAQILDSRFNTEFDDHNKRIR--KVHIITLKSKLVSDFRKEF  GLYKIRDINHY  980
WP_021320964      910  VD  KAGFIQRQLVETRQITKHVAQILDSRFNTEFDDHNKRIR--KVHIITLKSKLVSDFRKEF  GLYKIRDINHY  980
WP_037581760      910  VD  KAGFIQLQLVETRQITKHVAQILDSRFNTEFDDHNKRIR--KVHIITLKSKLVSDFRKEF  GLYKIRDINHY  980
WP_004232481      918  TD  KAGFIKRQLVETRQITKHVAQILDARFNTKCDENDKVIR--DVKVITLKSSLVSQFRKEF  KFYKVREINDY  988
WP_009854540      919  AD  KAGFIKRQLVETRQITKHVAQILDARFNTEHDENDKVIR--DVKVITLKSNLVSQFRKDF  EFYKVREINDY  989
WP_012962174      919  ND  KAGFIKRQLVETRQITKHVAQILDSRFNTERDENDKVIR--NVKVITLKSNLVSQFRKDF  KFYKVREINDY  989
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WP_039695303  921   AD   KAGFIKRQLVETRQITKHVAQILDARFNTEHDENDKVIR--DVKVITLKSNLVSQFRK
WP_014334983  918   AD   KAGFIKRQLVETRQITKHVAQILDARFNTKRDENDKVIR--DVKVITLKSNLVSQFRK
WP_003099269  911   FD   KAGFIKRQLVETRQITKHVAQILDSRFNSNLTEDSKSNR--NVKIITLKSKMVSDFRK
AHY15608      911   FD   KAGFIKRQLVETRQITKHVAQILDSRFNSNLTEDSKSNR--NVKIITLKSKMVSDFRK
AHY17476      911   FD   KAGFIKRQLVETRQITKHVAQILDSRFNSNLTEDSKSNR--NVKIITLKSKMVSDFRK
ESR09100            --   ----------------------------------------------------------
AGM98575      911   FD   KAGFIKRQLVETRQITKHVAQILDSRFNSNLTEDSKSNR--NVKIITLKSKMVSDFRK
ALF27331      911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_018372492  924   ED   KAGFIRRQLVETRQITKHVARLLDEKLNRKKNENGEKLR--TTKIITLKSVFASRFRA
WP_045618028  914   RD   KVGFIKRQLVETRQITKHVAQILDARFNTEVTEKDKKDR--SVKIITLKSNLVSNFRK
WP_045635197  913   RD   KVGFIKRQLVETRQITKHVAQILDARYNTEVNEKDKKNR--TVKIITLKSNLVSNFRK
WP_002263549  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002263887  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002264920  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_002271977  911   DD   KAGFIKRQLVETRQITKHVARILDERFYTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002272766  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002273241  911   DD   KAGFIKRQLVETRQITKHVARILDERFYTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002275430  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_002277050  913   DD   KAGFIKRQLVETRQITKHVARMLDERFNKEFDDNNKRIR--RVKIVTLKSNLVSSFRK
WP_002277364  911   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_002290427  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_002295753  911   DD   KAGFIKRQLVETRQITKHVARILDERFYTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002296423  911   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002304487  925   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_002307203  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002310390  911   DD   KAGFIKHQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_002352408  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_012997688  911   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_019312892  911   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_019313659  911   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_019803776  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
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WP_024784288  913   DD   KAGFIKRQLVETRQITKHVARMLDERFNKEFDDNNKRIR--RVKIVTLKSNLVSSFRK
WP_024784666  911   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_024784894  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_024786433  913   DD   KAGFIKRQLVETRQITKHVARMLDERFNKEFDDNNKRIR--RVKIVTLKSNLVSSFRK
WP_049473442  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_049474547  911   DD   KAGFIKRQLVETRQITKHVARILDERFNTETDENNKKIR--QVKIVTLKSNLVSNFRK
EMC03581      904   DD   KAGFIKRQLVETRQITKHVARILDERFHTETDENNKKIR--QVKIVTLKSNLVSNFRK
WP_000428612  916   RD   KVGFIKRQLVETRQITKHVAQILDARYNTEVNEKDKKNR--TVKIITLKSNLVSNFRK
WP_000428613  914   RD   KVGFIKRQLVETRQITKHVAQILDARFNKEVNEKDKKNR--TVKIITLKSNLVSNFRK
WP_049523028  909   RD   KVGFIKRQLVETRQITKHVAQILDDRFNAEVNEKNQKLR--SVKIITLKSNLVSNFRK
WP_003107102  880   YD   KAGFIKRQLVETRQITKHVAQILNNRFNNNVDDSSKNKR--PVKIITLKSKMVSDFRK
WP_054279288  912   SD   KANFIQRQLVETRQITKHVAQILDSRFNTERDEKDRPIR--RVKVITLKSKFVSDFRQ
WP_049531101  914   RD   KVGFIKRQLVETRQITKHVAQILDSRFNTKVNEKNQKIR--TVKIITLKSNLVSNFRK
WP_049538452  914   RD   KVGFIRRQLVETRQITKHVAQILDSRFNTEVTEKDKKNR--NVKIITLKSNLVSNFRK
WP_049549711  916   LD   KVGFIKRQLVETRQITKHVAQILDARFNKEVTEKDKKNR--NVKIITLKSNLVSNFRK
WP_007896501  917   SD   KARFLRRQLVETRQITKHVAQLLDSRFNSKSNQNKKLAR--NVKIITLKSKIVSDFRK
EFR44625      869   SD   KARFLRRQLVETRQITKHVAQLLDSRFNSKSNQNKKLAR--NVKIITLKSKIVSDFRK
WP_002897477  913   RD   KVGFIRRQLVETQQITKNVAQILDARFNTEVKEKNQKIR--TVKIITLKSNLVSNFRK
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WP_002906454  913   RD   KVGFIKRQLVETRQITKHVAQLLDTRFNTEVNEENQKIR--TVKIITLKSNLVSNFRK
WP_009729476  914   LD   KVGFIKRQLVETRQITKHVAQILDARFNKEVTEKDKKNR--TVKIITLKSNLVSNFRK
CQR24647      913   ED   KAGFIKRQLVETRQITKHVARILDERFNRDFDKNDKRIR--NVKIVTLKSNLVSNFRK
WP_000066813  918   LD   KVGFIKRQLVETRQITKHVAQFLDARFNKEVTEKDKKNR--NVKIITLKSNLVSNFRK
WP_009754323  914   RD   KVGFIKRQLVETRQITKHVARILDARFNTEVSEKNQKIR--SVKIITLKSNLVSNFRK
WP_044674937  913   ED   KARFIQRQLVETRQITKHVARILDTRFNTKLDEAGNRIRdpKVNIITLKSNLVSQFRK
WP_044676715  915   ED   KARFIQRQLVETRQITKHVARILDTRFNTKLDEAGNRIRdpKVNIITLKSNLVSQFRK
WP_044680361  915   ED   KARFIQRQLVETRQITKHVARILDTRFNTKLDEAGNRIRdpKVNIITLKSNLVSQFRK
WP_044681799  913   ED   KARFIQRQLVETRQITKHVARILDTRFNTKLDEAGNRIRdpKVNIITLKSNLVSQFRK
WP_049533112  910   ND   KAGFIKRQLVETRQITKHVAQVLDARFNAKHDENKKVIR--DVKIITLKSNLVSQFRK
WP_029090905  891   RD   KEGFIARQLVETRQITKHVTQLLQQEY-----------K-dTTKVFAIKATLVSGLRR
WP_006506696  917   RD   EERFINRQLVETRQITKNVTQIIEDHYST-------------TKVAAIRANLSHEFRV
AIT42264      911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_034440723  915   RD   RQQFINRQLVETRQITKHVANLLSHHLNEK-----KEVG--EINIVLLKSALTSQFRK
AKQ21048      911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_004636532  916   AD   RAHFLNRQLVETRQITKHVANLLDSQYNTAEEQ-----R---INIVLLKSSMTSRFRK
WP_002364836  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNANSKE-----K--KVQIITLKASLTSQFRS
WP_016631044  873   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
EMS75795      655   ED   KARFIQRQLVETRQITKHVAAILDQYFN-QPEE-SK-NK--GIRIITLKSSLVSQFRK
WP_002373311  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_002378009  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_002407324  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_002413717  922   ED   KAHFIQRQLVETRQITKNVAGILNQRYNANSKE-----K--KVQIITLKASLTSQFRS
WP_010775580  924   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_010818269  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_010824395  922   ED   KAHFIQRQLVETRQITKNVAGILDQLYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_016622645  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_033624816  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_033625576  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_033789179  922   ED   KAHFIQRQLVETRQITKNVAGILDQRYNAKSKE-----K--KVQIITLKASLTSQFRS
WP_002310644  919   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
WP_002312694  920   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNDPIR--KVRIITLKSALVSQFRN
WP_002314015  920   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
WP_002320716  920   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
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WP_002330729  919   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
WP_002335161  920   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
WP_002345439  920   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
WP_034867970  911   ED   KARFIQRQLVETRQITKHVANILHQSFN-QEEEGTD-CD--GVQIITLKATLTSQFRQ
WP_047937432  920   ED   KAGFIKRQLVETRQITKHVAGILHHRFN-KAEDTNEPIR--KVRIITLKSALVSQFRN
WP_010720994  911   ED   KARFIQRQLVETRQITKHVANILHQSFN-QEEEGTD-CD--GVQIITLKATLTSQFRQ
WP_010737004  911   ED   KARFIQRQLVETRQITKHVANILHQSFN-QEEEGTD-CD--GVQIITLKATLTSQFRQ
WP_034700478  911   ED   KARFIQRQLVETRQITKHVANILHQSFN-QEEEGTD-CD--GVQIITLKATLTSQFRQ
WP_007209003  912   SD   KAGFLKRQLVETRQITKHVATILDSKFNE--DSNNRDVQ-----IITLKSALVSEFRK
WP_023519017  905   AD   KARFIQRQLVETRQITKHVANLLHQHFN-LPEEVSA-TE--KTSIITLKSTLTSQFRQ
WP_010770040  915   DD   RAHFIKRQLVETRQITKHVARILDQRFNSQKDEEGKTIR--AVRVVTLKSSLTSQFRK
WP_048604708  912   DD   KAGFIHRQLVETRQITKNVARILHQRFNSEKDEEGNLIR--KVRIITLKSALTSQFRK
WP_010750235  914   DD   KARFIQRQLVETRQITKHVAAILHQYFN-QTQELEK-EK--DIRIITLKSSLVSQFRQ
AII16583      950   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
WP_029073316  928   KD   KERFINRQIVETRQITKHVAQIISNHYET-------------TKVVTVRADLSHAFRE
WP_031589969  928   KD   QERFINRQIVETRQITKHVAQIIDNHYEN-------------TKVVTVRADLSHQFRE
KDA45870      903   KL   KERFIERQLVETRQITKYVAQLLDQRLN--YDGNGVELD-eKIAIVTLKAQLASQFRS
WP_039099354  924   -D   MKGFINRQLVETRQVIKLATNLLMEQYGED-----------NIELITVKSGLTHQMRT
AKP02966      926   KD   KLGFIHRQLVQTSQMVKGVANILNSMYK---NQGTTCIQ--------ARANLSTAFRK
WP_010991369  914   AD   KARFIHRQLVETRQITKNVANILHQRFNYEKDDHGNTMK--QVRIVTLKSALVSQFRK
WP_033838504  914   AD   KARFIHRQLVETRQITKNVANILHQRFNYEKDDHGNTMK--QVRIVTLKSALVSQFRK
EHN60060      917   AD   KARFIHRQLVETRQITKNVANILHQRFNYEKDDHGNTMK--QVRIVTLKSALVSQFRK
EFR89594      683   AD   KARFIHRQLVETRQITKNVANILHQRFNYGKDDHGNTMK--QVRIVTLKSALVSQFRK
WP_038409211  914   AD   KANFIQRQLVETRQITKNVANILYQRFNCKQDENGNEVE--QVRIVTLKSTLVSQFRK
EFR95520      533   AD   KANFIQRQLVETRQITKNVANILYQRFNCKQDENGNEVE--QVRIVTLKSTLVSQFRK
WP_003723650  914   AD   KARFIHRQLVETRQITKNVANILYQRFNKETDNHGNTME--QVRIVTLKSALVSQFRK
WP_003727705  914   AD   KARFIHRQLVETRQITKNVANILHQRFNKETDNHGNTME--QVRIVTLKSALVSQFRK
WP_003730785  914   AD   KARFIHRQLVETRQITKNVANILHQRFNKETDNHGNTME--QVRIVTLKSALVSQFRK
WP_003733029  914   AD   KARFIHRQLVETRQITKNVANILHQRFNYKTDGNKDTME--TVRIVTLKSALVSQFRK
WP_003739838  914   AD   KATFIHRQLVETRQITKNVANILHQRFNNETDNHGNNME--QVRIVMLKSALVSQFRK
WP_014601172  914   AD   KARFIHRQLVETRQITKNVANILHQRFNNETDNHGNTME--QVRIVTLKSALVSQFRK
WP_023548323  914   AD   KARFIHRQLVETRQITKNVANILHQRFNYKTDDNEDTME--PVRIVTLKSALVSQFRK
WP_031665337  914   AD   KARFIHRQLVETRQITKNVANILHQRFNKETDNHGNTME--QVRIVTLKSALVSQFRK
WP_031669209  914   AD   KARFIHRQLVETRQITKNVANILHQRFNYKTDGNKDTME--TVRIVTLKSALVSQFRK
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WP_033920898  914   AD   KARFIHRQLVETRQITKNVANILHQRFNYKTDDNEDTME--PVRIVTLKSALVSQFRK
AKI42028      917   AD   KARFIHRQLVETRQITKNVANILHQRFNNETDNHGNTME--QVRIVTLKSALVSQFRK
AKI50529      917   AD   KARFIHRQLVETRQITKNVANILHQRFNYKTDDNEDTME--PVRIVTLKSALVSQFRK
EFR83390      362   AD   KARFIHRQLVETRQITKNVANILHQRFNNETDNHGNTME--QVRIVTLKSALVSQFRK
WP_046323366  914   AD   KARFIHRQLVETRQITKNVANILHQRFNCKKDESGNVIE--QVRIVTLKAALVSQFRK
AKE81011      927   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
CUO82355      921   RD   EERFINRQLVETRQITKNVTQIIEDHYST-------------TKVAAIRANLSHEFRV
WP_033162887  923   KD   KERFINRQLVETRQIIKNVAVIINDHYTN-------------TNIVTVRAELSHQFRE
AGZ01981      944   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
AKA60242      911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
AKS40380      911   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
4UN5_B        915   LD   KAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIR--EVKVITLKSKLVSDFRK
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WP_010922251  982   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDV   S---EQEi--GK   ATAKY--
WP_039695303  992   HHAHDAYLNAVVGTALLKKYPKL-ASEFVYGEYKKYDI   S---SD------   KATAK--
WP_045635197  984   HHAHDAYLNAVVAKAILKKYPKL-EPEFVYGEYQKYDL   SkdpKEV---EK   ATEKY--
5AXW_A        703   HHAEDALI--------------IaNADFIFKEWKKLDK   Nq-mFE----EK   ETEQEyk
WP_009880683  666   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDI   S---EQEi--GK   ATAKY--
WP_010922251  982   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDV   S---EQEi--GK   ATAKY--
WP_011054416  982   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDV   S---EQEi--GK   ATAKY--
WP_011284745  982   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDV   S---EQEi--GK   ATAKY--
WP_011285506  982   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDV   S---EQEi--GK   ATAKY--
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WP_012560673  982   HHAHDAYLNAVVGTALIKKYPKL-ESEFVYGDYKVYDI   S---EQEi--GK   ATAKY--
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KLL20707       1001  HHAHDAYLNAVVAKAILTKYPQL-EPEFVYGDYPKYN-   S---YKT---RK   ATEKL--
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WP_045635197	1056	YAD-GTIVKRENIE	Y-SKDtGE-IAWNKEKDFAIIKKVLS-LPQVNIVKKREVQT	GGFSK	ESIL-PKG-	1118
5AXW_A	772	YKYsHRVDKKPNRE	VNNLN-GL---YDKDND--KLKKLINksPEKLMYHHDPQT	--YQK	KLIMeQYGd	852
WP_009880683	736	LAN-GEIRKRPLIE	TNGET-GE-IVWDKGRDFATVRKVLS-MPQVNIVKKTEVQT	GGFSK	ESIL-PKR-	798
WP_010922251	1052	LAN-GEIRKRPLIE	TNGET-GE-IVWDKGRDFATVRKVLS-MPQVNIVKKTEVQT	GGFSK	ESIL-PKR-	1114
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WP_002310644 1049 --T---------P VCDEN-GE-IFWDKSKSIAQVKKVIN-HHHMNIVKKTEIQK GGFSK ETVE-PKK- 1100
WP_002312694 1050 --T---------P VCDEN-GE-IFWDKSKSIAQVKKVIN-HHHMNIVKKTEIQK GGFSK ETVE-PKK- 1101
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WP_002345439 1050 --T----------P VCDEN-GE-IFWDKSKSIAQVKKVIN-HHHMNIVKKTEIQK GGFSK ETVE-PKK- 1101
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CUO82355      1043  --D-----------    ------GK-LIWNP-DLINEIKKCFY-YKDCYCTTKLDQKS   G
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4UN5_B        1056  LAN-GEIRKRPLIE    TNGET-GE-IVWDKGRDFATVRKVLS-MPQVNIVKKTEVQT   G
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WP_021320964    1095  --NSD  KLIP----RKNNW-DTRKYGGFDSPTVAYSV-LVIAK--ME--KGKAKVLKPVKEMVGITIME  RIAFEE  1156
WP_037581760    1095  --NSD  KLIP----RKNNW-DTRKYGGFDSPTVAYSV-LVIAK--ME--KGKAKVLKPVKEMVGITIME  RIAFEE  1156
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WP_009854540    1119  --DSD  KLIPRKTkKV-YW-DTKKYGGFDSPTVAYSV-FVVAD--VE--KGKAKKLKTVKELVGISIME  RSFFEE  1183
WP_012962174    1120  --NSD  KLIPRKTkKF-RW-DTPKYGGFDSPNIAYSV-FVIAD--VE--KGKAKKLKTVKELVGISIME  RSSFEE  1184
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WP_014334983 1124 --DSD  KLIPRKTkKV-YW-NTKKYGGFDSPTVAYSV-LVVAD--IE--KGKAKKLKTVKELVGISIME 1188 RSFEEE 1188
WP_003099269 1115 --DSD  KLIA----RKKSW-DPKKYGGFDSPIIAYSV-LVVAK--IA--KGKTQKLKTIKELVGIKIME 1176 QDEFEK 1176
AHY15608     1115 --DSD  KLIA----RKKSW-DPKKYGGFDSPIIAYSV-LVVAK--IA--KGKTQKLKTIKELVGIKIME 1176 QDEFEK 1176
AHY17476     1115 --DSD  KLIA----RKKSW-DPKKYGGFDSPIIAYSV-LVVAK--IA--KGKTQKLKTIKELVGIKIME 1176 QDEFEK 1176
ESR09100        1 -----  ------------------------------------------------------------ME 8    QDEFEK 8
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ALF27331     1094 --NSD  KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAN--IE--KGKSKKLKLVKDLVGITIME 1158 RTIFEK 1158
WP_018372492 1109 ---NN  KLIP----RKKDW-SVDKYGGFIEPAESYSLaIFYTD--IN-----GKKPKKKSTIIAISRME 1167 KKDYEK 1167
WP_045618028 1125 vvDAS  KLTPIKS-G---L-SPEKYGGYARPTIAYSV-LVIAD--IE--KGKAKKLKRIKEMVGITVQD 1188 KKKFEA 1188
WP_045635197 1119 --NSD  KLIPRKT-KDILL-DTTKYGGFDSPVIAYSI-LLIAD--IE--KGKAKKLKTVKTLVGITIME 1183 KAAFEE 1183
WP_002263549 1094 --NSD  KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158 KMTFER 1158
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WP_002272766 1094 --NSD  KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158 KMTFER 1158
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WP_002288990 1094 --NSY  KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158 KMTFER 1158
WP_002289641 1094 --NSD  KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158 KMTFER 1158
WP_002290427 1094 --DSD  KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158 KMTFER 1158
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WP_002305844 1094 --NSD KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158
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WP_024784666 1094 --NSD KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158
WP_024784894 1094 --NSD KLIPRKT-KKFYW-DTKKYGGFDSPIVAYSI-LVIAD--IE--KGKSKKLKTVKALVGVTIME 1158
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EFR44625     1073 --NSD KLIE----RKKGW-DPKKYGGFDSPNTAYSI-FVVAK--VA--KRKAQKLKTVKEIVGITIME 1134
WP_002897477 1119 --NSD KLIPRKT-KDILW-DTTKYGGFDSPVIAYSI-LLIAD--IE--KGKAKKLKTVKTLVGITIME 1183

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WP_031589969 1100 nsDKD ATVP---vNKYRS-NVNKYGGFSG--VNSFI----VA--IKgkKKKGKKVIEVNKLTGIPLMY KNADEE 1164
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WP_003733029 1096 --KSN KLIP----RKKDW-DPIKYGGFDGSKMAYAI-II--E--YE--KQK-RKVRIEKKLIQINIME REAFEK 1154
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WP_014601172 1096 --NSS KLIP----RKENW-DPMKYGGLDSPNMAYAV-II--E--HA--KGK-KKLIFEKKIIRITIME RKMFEK 1154
WP_023548323 1096 --DSS KLIP----KKTNL-NPIKYGGFEGSNMAYAI-II--E--HE--KRK-KKVTIEKKLIQINIME RKAFEK 1154
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WP_031669209 1096 --KSN KLIP----RKKDW-DPIKYGGFDGSKMAYAI-II--E--YE--KQK-RKVRIEKKLIQINIME REAFEK 1154
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EFR83390      544   --NSS  KLIP----RKENW-DPMKYGGLDSPNMAYAV-II--E--HA--KGK-KKIVIEKKLIQINIME  RKMFEK  602
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AKE81011      1131  --NSD  KLIA----RKKDW-DPKKYGGFDSPTVAYSV-LVVAK--VE--KGKSKKLKSVKELLGITIME  RSSFEK  1192
CUO82355      1089  hsAKG  AVIP---vNKNRK-DVNKYGGFSG--LQYVI----AA--IEgtKKKGKKLVKVRKLSGIPLYL  KQADIK  1153
WP_033162887  1090  hsEKG  ATVP---lNKYRA-DVHKYGGFGN--VQSII----VA--IEgkKKKGKKLIDVRKLTSIPLHL  KNAPVE  1154
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AKA60242      1115  --NSD  KLIA----RKKDW-DPKKYGGFDSPTVAYSV-LVVAK--VE--KGKSKKLKSVKELLGITIME  RSSFEK  1176
AKS40380      1115  --NSD  KLIA----RKKDW-DPKKYGGFDSPTVAYSV-LVVAK--VE--KGKSKKLKSVKELLGITIME  RSSFEK  1176
4UN5_B        1119  --NSD  KLIA----RKKDW-DPKKYGGFDSPTVAYSV-LVVAK--VE--KGKSKKLKSVKELLGITIME  RSSFEK  1180
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WP_045635197  1184  NPI---TFLE---NKGYHN--V-RKENILC--LPKYSLFE---LENGRRRLLAS  AKELQKGNEIVLPVYLTTLLYHS  1246
5AXW_A         913   KPYrfdVYLD---NGVYKFvtV-KNLDVIK----KENYYE---VNSKAYEEAKK  -KKISNQAEFIASFYNNDLIKIN  978
WP_009880683   861   DPV---DFLE---AKGYKE--V-RKDLIIK--LPKYSLFE---LENGRKRMLAS  -GELQKGNELALPSKYVNFLYLA  923
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KGE60856       115  DPI---DFLE---AKGYKE--V-RKDLIIK--LPKYSLFE---LENGRKRMLAS  -GELQKGNELALPSKYVNFLYLA  177
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KLL20707          1192  NPS---AFLE---SKGYLN--I-RADKLII--LPKYSLFE---LENGRRRLLAS  AGELQKGNELALPTQFMKFLYLA  1254
KLL42645          1178  NPS---AFLE---SKGYLN--I-RDDKLMI--LPKYSLFE---LENGRRRLLAS  AGELQKGNELALPTQYMKFLYLA  1240
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GAD46167          1159  NPV---DFLE---QRGYRN--V-RLEKIIK--LPKYSLFE---LENKRRRLLAS  ARELQKGNELVIPQRFTTLIYHS  1221
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WP_009854540      1184  NPV---EFLE---NKGYHN--I-REDKLIK--LPKYSLFE---FEGGRRRLLAS  ASELQKGNEMVLPGYLVELLYHA  1246
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EMC03581      1152  DPV---AFLE---RKGYRN--V-QEENIIK--LPKYSLFK---LENGRKRLLAS  ARELQKGNEIVLPNHLGTLLYHA  1214
WP_000428612  1187  SPI---AFLE---NKGYHN--V-RKENILC--LPKYSLFE---LKNGRRRMLAS  AKELQKGNEIVLPVHLTTLLYHA  1249
WP_000428613  1185  NPI---TFLE---NKGYHN--V-RKENILC--LPKYSLFE---LENGRRRLLAS  AKELQKGNEIVLPVYLTTLLYHS  1247
WP_049523028  1180  NPV---AFLE---GKGYQN--V-VEENIIR--LPKYSLFE---LENGRRRMLAS  AKELQKGNEMVLPSYLIALLYHA  1242
WP_003107102  1146  DRI---TFLE---KKGYQD--I-QESLIIK--LPKFSLFE---LENGRKRLLAS  --ELQKGNELSLPNKYIQFLYLA  1208
WP_054279288  1179  NPI---AFLE---SKGYHD--I-QEHLMIT--LPKYSLFE---LENGRRRLLAS  --ELQKGNEMVLPQHLVTFLYRV  1241
WP_049531101  1189  NPT---AYLE---EYGYKN--I-NPNLIIK--LPKYSLFK---FNDGQRRLLAS  SIELQKGNELILPYHFTTLLYHA  1251
WP_049538452  1189  NPI---AYLE---ECGYKN--I-NPNLIIK--LPKYSLFE---FNGGQRRLLAS  SIELQKGNELILPYHFTALLYHT  1251
WP_049549711  1191  NPI---AYLE---ECGYKN--I-NPNLIIK--LPKYSLFE---FNGGQRRLLAS  SIELQKGNELILPYHFTALLYHA  1253
WP_007896501  1183  DNI---AFLE---KKGYQD--I-QEKLLIK--LPKYSLFE---LENGRRRLLAS  --EFQKGNELALSGKYMKFLYLA  1245
EFR44625      1135  DNI---AFLE---KKGYQD--I-QEKLLIK--LPKYSLFE---LENGRRRLLAS  --EFQKGNELALSGKYMKFLYLA  1197
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WP_002906454 1190 NPV---TYLE---ECGYKN--I-NSNLIIK--LPKYSLFE---FNDGQRRLLAS SIELQKGNELILPYHLTALLYHA 1252
WP_009729476 1185 NPI---AFLE---NKGYHN--V-CKENILC--LPKYSLFE---LENGRRRLLAS AKELQKCNEIVLPVYLTTLLYHS 1247
CQR24647     1175 NSV---TFLE---EKGYRN--I-RENTIIK--FPKYSLFE---LENGRRRLLAS AIELQKGNEMFLPQQFVNLLYHA 1237
WP_000066813 1189 NPI---TFLE---NKGYHN--V-RKENILC--LPKYSLFE---LESGRRRMLAS AKELQKGNEIVLPVYLTTLLYHS 1251
WP_009754323 1185 NPI---TFLE---NKGYHN--V-RKENILC--LPKYSLFE---LENGRRRLLAS AKELQKGNEIVLPVYLTTLLYHS 1247
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WP_044676715 1180 NPI---EFLE---HKGYKN--I-LEKNIIK--LPKYSLFE---LENGRRRLLAS AKELQKGNEMILPPHLVTLLYHS 1242
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WP_044681799 1178 NPI---EFLE---HKGYKN--I-LEKNIIK--LPKYSLFE---LENGRRRLLAS AKELQKGNEMILPPHLVTLLYHS 1240
WP_049533112 1186 DKR---AFLL---GKGYKD--I-K--KIIE--LPKYSLFE---LKDGSRRMLAS RGEIHKGNELFVPQKFTTLLYHA 1253
WP_029090905 1118 SETslqAYIA---EQIKSE--VeIIN----grILKYQLIS----NNGNRLYIAG -SERHNARQLIVSDEAAKVIWLI 1181
WP_006506696 1150 EKI---NYIE--eKEGLSD--VrIIK---Dn-IPVNQMIEm----DGGEYLLTS --EYVNARQLVLNEKQCALIADI 1211
AIT42264     1177 NPI---DFLE---AKGYKE--V-KKDLIIK--LPKYSLFE---LENGRKRMLAS -GELQKGNELALPSKYVNFLYLA 1239
WP_034440723 1150 QTT---EYLG---KIGFNK--AsIIN---S--FKNYTLFE---LENGSRRMIVG KGELQKGNQMYLPQNLLEFVYHL 1217
AKQ21048     1177 NPI---DFLE---AKGYKE--V-KKDLIIK--LPKYSLFE---LENGRKRMLAS -GELQKGNELALPSKYVNFLYLA 1239
WP_004636532 1151 HPT---AYLE---EAGYNN--P-TV--LHE--LFKYQLFE---LEDGSRRMIAS AKEFQKGNQMVLPLELVELLYHA 1211
WP_002364836 1157 NPI---LFLE---EKGFLR--P-RV--LMK--LPKYTLYE---FPEGRRRLLAS AKEAQKGNQMVLPEHLLTLLYHA 1217
WP_016631044 1108 NPI---LFLE---EKGFLR--P-RV--LMK--LPKYTLYE---FPEGRRRLLAS AKEAQKGNQMVLPEHLLTLLYHA 1168
EMS75795     893  DPI---GFLS---NKGYSN--V-TKF--IK--LSKYTLYE---LENGRRRMVAS -KEAQKANSFTLPEKLVTLLYHA 953
WP_002373311 1157 NPI---LFLE---EKGFLR--P-RV--LMK--LPKYTLYE---FPEGRRRLLAS AKEAQKGNQMVLPEHLLTLLYHA 1217
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WP_002312694 1160 SPV---LFLK---NKGYEQ--A-EIE--MK--LPKYALFE---LENGRKRMVAS -KEAQKANSFLLPEHLVTLLYHA 1220
WP_002314015 1160 SPV---LFLK---NKGYEQ--A-EIE--MK--LPKYALFE---LENGRKRMVAS -KEAQKANSFLLPEHLVTLLYHA 1220
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WP_002330729 1159 SPV---LFLK---NKGYEQ--A-EIE--MK--LPKYALFE---LENGRKRMVAS -KEAQKANSFLLPEHLVTLLYHA 1219
WP_002335161 1160 SPV---LFLK---NKGYEQ--A-EIE--MK--LPKYALFE---LENGRKRMVAS -KEAQKANSFLLPEHLVTLLYHA 1220
WP_002345439 1160 SPV---LFLK---NKGYEQ--A-EIE--MK--LPKYALFE---LENGRKRMVAS -KEAQKANSFLLPEHLVTLLYHA 1220
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WP_007209003 1148 NPIl---YLS---KKDYHN---pKVEAI----LPKYSLFE---FENGRRRMVAS -SETQKGNQLIIPGHLMELLYHS 1208
WP_023519017 1144 DPR---EFLK---TKGYEG--V-KQW--LI--LPKYILFE---AQGGYRRMIAS -QETQKANSLILPENLVTLLYHA 1204
WP_010770040 1155 DPV---SLLE---EKGYAN--P-EV--LIH--LPKYTLYE---LENGRRLLAS ANEAQKGNQLVLPASLVTLLYHA 1215
WP_048604708 1152 NER---EFLK---NKGYQN--P-QI--CMK--LPKYSLYE---FDDGRRRLLAS AKEAQKGNQMVLPAHLVTFLYHA 1212
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AII16583     1216 NPI---DFLE---AKGYKE--V-KKDLIIK--LPKYSLFE---LENGRKRMLAS -GELQKGNELALPSKYVNFLYLA 1278
WP_029073316 1165 TKI---NYIK--eSEGLEE--VkIIK---E--ILKNQLIEi----NGGLFYVTS --EIVNARQLILDFNCTRIIDGI 1225
WP_031589969 1165 IKI---NYLK--qAEDLEE--VqIGK---E--ILKNQLIEk----DGGLYYIVA --EIINAKQLILNESQTKLVCEI 1225
KDA45870     1139 DPT---AYLA---SRGYTNvtNsFIL-------PKYSLLEd---PEGRRRYLAS -KEFQKANELILPQHLVELLYWV 1199
WP_039099354 1171 QKI-spQFTKv---KKQKGTiV-KVVEDFEv-IAPHILINqrfFDNGQELTLGS ----HNEQELILDKTAVKLLNGA 1241
AKP02966     1173 KTL--qNWLE---ENVKHKksIqIIK---Nn-VPIGQIIY-------SKKVGLLS -RETANRQQLILPPEHSALLRIL 1237
WP_010991369 1155 DEK---AFLE---EQGYRQ--P-KV--LAK--LPKYTLYE---CEEGRRRMLAS ANEAQKGNQQVLPNHLVTLLHHA 1215
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WP_003727705 1155 DEE---AFLE---EKGYHQ--P-KV--LTK--LPKYTLYE---CEKGRRRMLSS ANEAQKGNQLVLSNHLVSLLYHA 1215
WP_003730785 1155 DEE---AFLE---EKGYHQ--P-KV--LTK--LPKYTLYE---CEKGRRRMLSS ANEAQKGNQLVLSNHLVSLLYHA 1215
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WP_003739838 1155 DEK---SFLE---KQGYRQ--P-KV--LTK--LPKYTLYE---CENGRRRMLAS ANEAQKGNQQVLKGQLITLLHHA 1215
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EFR83390      603   DEE---AFLE---EKGYRH--P-KV--LTK--LPKYTLYE---CEKGRRRMLAS   ANEAQK
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WP_033162887  1155  EQL---SYIAspeHEDLID--VrIVK---E--ILKNQLIEi----DGGLYYVTS   --EYVT
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AKA60242      1177  NPI---DFLE---AKGYKE--V-KKDLIIK--LPKYSLFE---LENGRKRMLAS   -GELQK
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4UN5_B        1181  NPI---DFLE---AKGYKE--V-KKDLIIK--LPKYSLFE---LENGRKRMLAS   -GELQK
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WP_039695303  1249  HRAD-----NFNS-TEYLN--YVSEHKKEFEKVLSCVEDFANLYVDVE--KNLSKIR-A  VAD-SM---DNFSIEE--  1308
WP_045635197  1247  KNVH-----KLDE-PGHLE--YIQKHRNEFKDLLNLVSEFSQKYVLAD--ANLEKIK-S  LYA-DN---EQADIEI--  1306
5AXW_A        979   GELYRVIgVNNDlLNRIE---VNMIDITYREYLENMNDKRPPRIIKTiaSKTQSIK-K  LYEvKSk--KHPQIIKkg  1056
WP_009880683  924   SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  989
WP_010922251  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_011054416  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
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WP_012560673  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_014407541  1239  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1304
WP_020905136  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
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WP_023610282  1239  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1304
WP_030125963  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_030126706  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_031488318  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_032460140  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_032461047  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_032462016  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_032462936  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_032464890  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_033888930  1065  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1130
WP_038431314  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_038432938  1239  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1304
WP_038434062  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
BAQ51233      1151  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1216
KGE60162      415   SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  480
KGE60856      178   SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  243
WP_002989955  1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq-  1305
WP_003030002  1222  YQIE-----KNYE-PEHRE--YVEKHKDEFKELLEYISVFSRKYVLAD--NNLTKIE-M  LFS-KN---KDAEVSS--  1281
WP_003065552  1250  QRIN-----SFNS-TKYLD--YVSAHKKEFEKVLSCVEDFANLYVDVE--KNLSKIR-A  VAD-SM---DNFSIEE--  1309
WP_001040076  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDIFQIINDFSKRVILAD--ANLEKIN-R  LYQ-DNk--ENIPVDE--  1306
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WP_001040078  1249  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040080  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040081  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040083  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040085  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040087  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040088  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040089  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040090  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040091  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040092  1241  SRYNESKgKPEEiEKKQE--FVNQHISYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040094  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040095  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040096  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040097  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040098  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040099  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040100  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040104  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040105  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_001040106  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040107  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040108  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040109  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_001040110  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_015058523  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_017643650  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_017647151  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_017648376  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_017649527  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_017771611  1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_017771984  1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
CFQ25032      1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
CFV16040      1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
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KLJ37842       1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
KLJ72361       1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
KLL20707       1255  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
KLL42645       1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_047207273   1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_047209694   1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_050198062   1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_050201642   1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_050204027   1241  SRYNELKgKPEEiEQKQE--FVVQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_050881965   1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
WP_050886065   1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
AHN30376       1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
EAO78426       1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQLINDFSKRVILAD--ANLEKIN-K    L
CCW42055       1241  SRYNESKgKPEEiEKKQE--FVNQHVSYFDDILQIINDFSNRVILAD--ANLEKIN-K    L
WP_003041502   1254  KRIN----NPIN-KDHIE--YVKKHRDDFKELLNYVLEFNEKYVGAT--KNGERLK-E    A
WP_037593752   1223  YQIE----KNYE-PEHRE--YVEKHKDEFKELLEYISVFSRKYVLAD--NNLTKIE-M    L
WP_049516684   1223  YRIE----KDYE-PEHRE--YVEKHKDEFKELLEYISVFSRKYVLAD--NNLTKIE-M    L
GAD46167       1222  YQIE----KNYE-PEHRE--YVEKHKDEFKELLEYISVFSRKYVLAD--NNLTKIE-M    L
WP_018363470   1254  HRIG----NFNS-AEHLK--YVSEHKKEFEEVLSCVENFANVYVDVE--KNLSKIR-A    A
WP_003043819   1249  QNISATTgSNNLg-------YIEQHREEFKEIFEKIIDFSEKYILKN--KVNSNLK-S    S
WP_006269658   1222  YRIE----KDYE-PEHRE--YVEKHKDEFKELLEYISVFSRKYVLAD--NNLTKIE-M    L
WP_048800889   1242  HRID----NSDN-SEHLK--YITEHKEEFGKLLSYIENFAKSYVDVD--KNLEKIQ-L    A
WP_012767106   1246  -HAHKIEsSKE--LEHEA--YILDHYNDLYQLLSYIERFASLYVDVE--KNISKVK-E    L
WP_014612333   1246  -HAHKIEsSKE--LEHEA--YILDHYNDLYQLLSYIERFASLYVDVE--KNISKVK-E    L
WP_015017095   1246  -HAHKIEsSKE--LEHEA--YILDHYNDLYQLLSYIERFASLYVDVE--KNISKVK-E    L
WP_015057649   1246  -HAHKIEsSKE--LEHEA--YILDHYNDLYQLLSYIERFASLYVDVE--KNISKVK-E    L
WP_048327215   1246  -HAHKIEsSKE--LEHEA--YILDHYNDLYQLLSYIERFASLYVDVE--KNISKVK-E    L
WP_049519324   1246  -HAHKIEsSKE--LEHEA--YILDHYNDLYQLLSYIERFASLYVDVE--KNISKVK-E    L
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WP_021320964   1220  AHYHELTgSSEDvLRKKY--FVERHLHYFDDIIQMINDFAERHILAS--SNLEKIN-H    T
WP_037581760   1220  AHYHELTgSSEDvLRKKY--FVERHLHYFDDIIQMINDFAERHILAS--SNLEKIN-H    T
WP_004232481   1252  QHVN----NSHK-PEHLN--YVKQHKDEFKDIFNLIISIARINILKP--KVVDNL---    -
WP_009854540   1247  HRAD----NFNS-TEYLN--YVSEHKKEFEKVLSCVEDFANLYVDVE--KNLSKIR-A    V
WP_012962174   1248  HRVN----SFNN-SEHLK--YVSEHKKEFGEVLSCVENFAKSYVDVE--KNLGKIR-A    V
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WP_039695303  1249  HRAD----NFNS-TEYLN--YVSEHKKEFEKVLSCVEDFANLYVDVE--KNLSKIR-A    V
WP_014334983  1252  HRID----SFNS-TEHLK--YVSEHKKEFEKVLSCVENFSNLYVDVE--KNLSKVR-A    A
WP_003099269  1240  SHYTKFTgKEEDrEKKRS--YVESHLYYFDEIMQIIVEYSNRYILAD--SNLIKIQ-N    L
AHY15608      1240  SHYTKFTgKEEDrEKKRS--YVESHLYYFXEVKSSF----------------------    -
AHY17476      1240  SHYTKFTgKEEDrEKKRS--YVESHLYXFX---------------------------    -
ESR09100      72    SHYTKFTgKEEDrEKKRS--YVESHLYYFDEIMQIIVEYSNRYILAD--SNLIKIQ-N    L
AGM98575      1240  SHYTKFTgKEEDrEKKRS--YVESHLYYFDVRLSQVFRVTNVEF--------------    -
ALF27331      1222  KNIH----KVDE-PKHLD--YVKKHKDEFKELLDVVSNFSKKNILAE--SNLEKIE-E    L
WP_018372492  1229  KKVDVLVkSKDD---DYD---LEEHRAEFAELLDCIKKFNDMYILAS--SNMSKIE-E    I
WP_045618028  1252  QRIN----KISE-PIHKQ--YVETHQSEFKELLTAIISLSKKYI-QK--PNVESL---    L
WP_045635197  1247  KNVH----KLDE-PGHLE--YIQKHRNEFKDLLNLVSEFSQKYVLAD--ANLEKIK-S    L
WP_002263549  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002263887  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002264920  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
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WP_002269448  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002271977  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002272766  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002273241  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002275430  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002276448  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002277050  1230  HHL-------DN-DYSNE--YVKNHYQQFDILFNEITSFSKKCKLGK--EHIQKIE-E    A
WP_002277364  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
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WP_002280230  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002281696  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002282247  1230  HHL-------DN-DYSNE--YVKNHYQQFDILFNEITSFSKKCKLGK--EHIQKIE-E    A
WP_002282906  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002283846  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002287255  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002288990  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002289641  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
WP_002290427  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E    L
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WP_002295753  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_002296423  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_002304487  1236  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1295
WP_002305844  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_002307203  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
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WP_019803776  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_019805234  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_024783594  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_024784288  1230  HHL-------DN-DYSNE--YVKNHYQQFDILFNEITSFSKKCKLGK--EHIQKIE-E  AYSkER---DFASIEE--  1287
WP_024784666  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_024784894  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_024786433  1230  HHL-------DN-DYSNE--YVKNHYQQFDILFNEITSFSKKCKLGK--EHIQKIE-E  AYSkER---DSASIEE--  1287
WP_049473442  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
WP_049474547  1222  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1281
EMC03581      1215  KNIH----KVDE-PKHLD--YVDKHKDEFKELLDVVSNFSKKYTLAE--GNLEKIK-E  LYA-QN---NGEDLKE--  1274
WP_000428612  1250  KNIH----RLDE-PEHLE--YIQKHRNEFKGLLNLVSEFSQKYVLAD--ANLEKIK-N  LYA-DN---EQADIEI--  1309
WP_000428613  1248  KNVH----KLDE-PEHLE--YIQKHRNEFKDLLNLVSEFSQKYVLAD--ANLEKIQ-N  LYA-DN---EQADIEI--  1307
WP_049523028  1243  KRIQ----KKDE-PEHLE--YIKQHHSEFNDLLNFVSEFSQKYVLAE--SNLEKIK-N  LYI-DN---EQTNMEE--  1302
WP_003107102  1209  SRYTSFSgKEEDrEKHRH--FVESHLHYFDEIKDIIADFSRRYILAD--ANLEKIL-T  LYN-EKn---QFSIEEq-  1274
WP_054279288  1242  SKRDK--gTQSEnME-----YISNHKEKFIEIFHYIIRYAEKNVIKP--KVIERLN-D  TFNqKF---NDSDLTEl-  1303
WP_049531101  1252  QRIN----KISE-PIHKQ--YVETHQSEFEELLTTIISLSKKYI-QK--PIVESL---  LQQ-AF---EQADKDIyq  1310
WP_049538452  1252  QRIN----KISE-PIHKQ--YVEAHQNEFKELLTTIISLSKKYI-QK--PNVESL---  LQQ-AF---EQADKDIyq  1310
WP_049549711  1254  QRIN----KFSE-PIHKQ--YVEAHQNEFKELLTIIISLSKKYI-QK--PNVESL---  LHQ-AF---EQADNDIyq  1312
WP_007896501  1246  SRYDKLSsKIESeQQKKL--FVEQHLHYFDEILDIVVKHATCYIKAE--NNLKKII-S  LYK-KK---EAYSINEq-  1311
EFR44625      1198  SRYDKLSsKIESeQQKKL--FVEQHLHYFDEILDIVVKHATCYIKAE--NNLKKII-S  LYK-KK---EAYSINEq-  1263
WP_002897477  1247  KNLH----KLDE-PEHLE--YIQKHRNEFKDLLNLVSEFSQKYILAE--ANLEKIK-D  LYA-DN---EQADIEI--  1306
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WP_002906454 1253 QRIN----KISE-PIHKQ--YVEAHQNEFKELLTTIISLSKKYI-QK--PNVELL---  LQQ-AF---DQADKDIyq 1311
WP_009729476 1248 KNVH----KLDE-PGHLE--YIQKHRNEFKDLLNLVSEFSQKYVLAD--ANLEKIK-N  LYA-DN---EQADIEI-- 1307
CQR24647     1238 QHAN----KEDS----VI--YLEKHRHELSELFHHIIGVSEKTILKP--KVEMTLN-E  AFE-KHf--EFDEVSE-- 1295
WP_000066813 1252 KNVH----KLDE-PEHLE--YIQKHRYEFKDLLNLVSEFSQKYVLAD--ANLEKIK-N  LYA-DN---EQADIEI-- 1311
WP_009754323 1248 KNVH----KLDE-PEHLE--YIQKHRYEFKDLLNLVSEFSQKYVLAE--ANLEKIK-S  LYV-DN---EQADIEI-- 1307
WP_044674937 1241 SNIH----KITE-PIHLN--YVNKNKHEFKELLRHISDFSTRYILAQ--DRLSKIE-E  LYD-KN---DGDDISD-- 1300
WP_044676715 1243 SNIH----KITE-PIHLN--YVNKNKHEFKELLRHISDFSTRYILAQ--DRLSKIE-E  LYD-KN---DGDDISD-- 1302
WP_044680361 1243 SNIH----KITE-PIHLN--YVNKNKHEFKELLRHISDFSTRYILAQ--DRLSKIE-E  LYD-KN---DGDDISD-- 1302
WP_044681799 1241 SNIH----KITE-PIHLN--YVNKNKHEFKELLRHISDFSTRYILAQ--DRLSKIE-E  LYD-KN---DGDDISD-- 1300
WP_049533112 1254 KRIN----NPIN-KDHIE--YVKKHRDDFKELLNYVLEFNEKYVGAT--KNGERLK-E  AVA-DF---DSKSNEE-- 1313
WP_029090905 1182 STKQA----DE-AMFLKyYRLEHLEAVFEEL---IRKQAADYQIFE--KLIKKIEvN  FYS----c----TYNEk- 1240
WP_006506696 1212 YNAIYKQ-DYDNlDDILMi----------QLYIELTNKMKVLYPAY-rGLAEKFE-S  YVV----i----SKEEk- 1268
AIT42264     1240 SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq- 1305
WP_034440723 1218 KHYNE----DE--TSHK--FIVEHKAYFDELLNYIVEFANKYLELE--NSIEKIK-D  LYH-----gKGPDVEEke 1276
AKQ21048     1240 SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S  AYN-KH---RDKPIREq- 1305
WP_004636532 1212 NRYDKVK-----fPDSIE--YVHDNLAKFDDLLEYVIDFSNKYINAD--KNVQKIQ-K  IYK-EH---GTEDVEL-- 1271
WP_002364836 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_016631044 1169 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1228
EMS75795     954  QHYDEIAhKESF----D--YVNDHLSEFREILDQVIDFSNRYTIAA--KNTEKIA-E  LFE-QN---QESTVQS-- 1013
WP_002373311 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_002378009 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_002407324 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_002413717 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_010775580 1220 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1279
WP_010818269 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_010824395 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_016622645 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_033624816 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_033625576 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-TN---QTADVKE-- 1277
WP_033789179 1218 KQCLL----PNQ-SESLA--YVEQHQPEFQEILERVVDFAEVHTLAK--SKVQQIV-K  LFE-AN---QTADVKE-- 1277
WP_002310644 1220 KQYDEIShKESF----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K  IYK-EN---QTDDLAK-- 1279
WP_002312694 1221 KQYDEIShKESF----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K  IYK-EN---QTDDLAK-- 1280
WP_002314015 1221 KQYDEIShKESF----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K  IYK-EN---QTDDLAK-- 1280
WP_002320716 1221 KQYDEIShKESF----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K  IYK-EN---QTDDLAK-- 1280
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WP_002330729  1220  KQYDEIShKESF-----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K    I
WP_002335161  1221  KQYDEIShKESF-----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K    I
WP_002345439  1221  KQYDEIShKESF-----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K    I
WP_034867970  1211  QHYDKITyQESF-----D--YVNTHLSDFSAILTEVLAFAEKYTLAD--KNIERIQ-E    L
WP_047937432  1221  KQYDEIShKESF-----D--YVNEHHKEFSEVFARVLEFAGKYTLAE--KNIEKLE-K    I
WP_010720994  1211  QHYDKITyQESF-----D--YVNTHLSDFSAILTEVLAFAEKYTLAD--KNIERIQ-E    L
WP_010737004  1211  QHYDKITyQESF-----D--YVNTHLSDFSAILTEVLAFAEKYTLAD--KNIERIQ-E    L
WP_034700478  1211  QHYDKITyQESF-----D--YVNTHLSDFSAILTEVLAFAEKYTLAD--KNIERIQ-E    L
WP_007209003  1209  KKIIN--gKNSD---SVS--YIQNNKEKFREIFEYIVDFSSKYISAD--ANLNKIE-K    I
WP_023519017  1205  RHYDEINhKVSF-----D--YVNAHKEGFNDIFDFISDFGVRYILAP--QHLEKIK-V    A
WP_010770040  1216  KQVDE-----DS-GKSEE--YVREHRAEFAEILNYVQAFSETKILAN--KNLQTIL-K    L
WP_048604708  1213  KHCNE-----KP-D-SLK--YVTEHQSGFSEIMAHVKDFAEKYTLVD--KNLEKIL-S    L
WP_010750235  1214  NHYDEIAyKDSY-----D--YVNEHFSNFQDILDKVIIFAEKYTSAP--QKLNQII-A    T
AII16583      1279  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S    A
WP_029073316  1226  YKAMKYK-NYSElSQEEIm-----------NVYDIFVEKLKLYYPTY-kNIATNFE-N    F
WP_031589969  1226  YKAMKYK-NYDNlDSEKIi-----------DLYRLLINKMELYYPEYrkQLVKKFE-D    L
KDA45870      1200  NAKDG--------EQKLE-----DHKAEFKELFDKIMEFADKYVVAP--KNSEKIR-R    L
WP_039099354  1242  LPLTQ-----SEeLAEQV----------YDEILDQVMHYFPLYDTNQfrAKLSAGKaA    D
AKP02966      1238  QIPDE------DpDQILAf----YDKNILVEILQELITKMKKFYPFY--KNEQEFLaS    F
WP_010991369  1216  ANCEV-----SD-GKSLD--YIESNREMFAELLAHVSEFAKRYTLAE--ANLNKIN-Q    L
WP_033838504  1216  ANCEV-----SD-GKSLD--YIESNREMFAELLAHVSEFAKRYTLAE--ANLNKIN-Q    L
EHN60060      1219  ANCEV-----SD-GKSLD--YIESNREMFAELLAHVSEFAKRYTLAE--ANLNKIN-Q    L
EFR89594      985   ANCEV-----SD-GKSLD--YIESNREMFAELLAHVSEFAKRYTLAE--ANLNKIN-Q    L
WP_038409211  1216  KNCEA-----ND-GESLA--YIEMHREMFAELLAYISEFAKRYTLAN--DRLEKIN-M    F
EFR95520      835   KNCEA-----ND-GESLA--YIEMHREMFAELLAYISEFAKRYTLAN--DRLEKIN-M    F
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WP_003730785  1216  KNCEA-----SD-GKSLK--YIEAHRETFSELLAQVSEFATKYTLAD--ANLSKIN-N    L
WP_003733029  1216  EKYEA-----ID-GESLA--YIEVHRALFDELLAYISEFARKYTLSN--DRLDEIN-M    L
WP_003739838  1216  KNCEA-----SD-GKSLD--YIESNREMFGELLAHVSEFAKRYTLAD--ANLSKIN-Q    L
WP_014601172  1216  KNCEA-----SD-GKSLK--YIEAHRETFSELLAQVSEFATRYTLAD--ANLSKIN-N    L
WP_023548323  1216  EKREA-----ID-GESLA--YIEAHKAVFGELLAHISEFARKYTLAN--DKLDEIN-M    L
WP_031665337  1216  KNCEA-----SD-GKSLK--YIEAHRETFSELLAQVSEFATRYTLAD--ANLSKIN-N    L
WP_031669209  1216  EKYEA-----ID-GESLA--YIEVHRALFDELLAYISEFARKYTLSN--DRLDEIN-M    L
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WP_033920898  1216  EKREA-----ID-GESLA--YIEAHKAVFGELLAHISEFARKYTLAN--DKLDEIN-M    L
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AKI50529      1219  EKREA-----ID-GESLA--YIEAHKAVFGELLAHISEFARKYTLAN--DKLDEIN-M    L
EFR83390      664   KNCEA-----SD-GKSLK--YTEAHRETFSELLAQVSEFATRYTLAD--ANLSKIN-N    L
WP_046323366  1216  KNCEA-----SD-GKSLA--YIESHREMFAELLDSISEFASRYTLAD--ANLEKIN-T    I
AKE81011      1256  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S    A
CUO82355      1216  YNAIYKQ-DFDGlDNMLMi-----------QLYLQLIDKLKTLYPIY-mGIVEKFE-K    F
WP_033162887  1218  YAAMLKK-RYEYlDEEEIf-----------DLYLQLLQKMDTLYPAY-kGIAKRFF-D    F
AGZ01981      1273  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S    A
AKA60242      1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S    A
AKS40380      1240  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S    A
4UN5_B        1244  SHYEKLKgSPEDnEQKQL--FVEQHKHYLDEIIEQISEFSKRVILAD--ANLDKVL-S    A
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WP_010922251 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1365
WP_039695303 1309 ISN---SFI NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL NATLIHQSITGLYETRIDLSKL-- 1369
WP_045635197 1307 LAN---SFI NLLTFTALGAP-AAFKFFG--KDI--DRK--R-YTTVSEIL NATLIHQSITGLYETWIDLSKL-- 1367
5AXW_A       ------------------------------------------------------ ------------------------ 
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WP_011054416 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1365
WP_011284745 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATFIHQSITGLYETRIDLSQL-- 1365
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WP_014407541 1305 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1364
WP_020905136 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1365
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WP_023610282 1305 -AK---NII HLFTLTNLGAP-AAFKYFD--TTI--ERN--R-YKSIKEVL DATLIHQSITGLYEIRIDLSQL-- 1364
WP_030125963 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--GRN--R-YKSIKEVL DATLIHQSITGLYETRIDLSQL-- 1365
WP_030126706 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1365
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WP_032460140 1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--GRN--R-YKSIKEVL DATLIHQSITGLYETRIDLSQL-- 1365
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KGE60856     244  -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 303
WP_002989955 1306 -AE---NII HLFTLTNLGAP-TAFKYFD--TTI--DRK--R-YTSTKEVL DATFIHQSITGLYETRIDLSQL-- 1365
WP_003030002 1282 LAK---SFI SLLTFTAFGAP-AAFNFFG--ENI--DRK--R-YTSVTECL NATLIHQSITGLYETRIDLSKL-- 1342
WP_003065552 1310 ISN---SFI NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL NATLIHQSITGLYETRIDLSKI-- 1370
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KLL20707        1321  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1381
KLL42645        1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_047207273    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_047209694    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHKSITGLYETRIDLGKL--  1367
WP_050198062    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_050201642    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_050204027    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_050881965    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KII--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_050886065    1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
AHN30376        1307  LAK---NII  NLFTFTSLGAP-AAFKFFD--KSV--DRK--R-YTSTKEVL  DSTLIHQSITGLYETRIDLGKL--  1367
EAO78426        1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
CCW42055        1307  LAN---NII  NLFTFTSLGAP-AAFKFFD--KIV--DRK--R-YTSTKEVL  NSTLIHQSITGLYETRIDLGKL--  1367
WP_003041502    1314  ICT---SFL  GLFELTSLGSA-SDFEFLG--VKI--PRY--RdYTPSSLLK  DSTLIHQSITGLYETRIDLSKL--  1383
WP_037593752    1283  LAK---SFI  SLLTFTAFGAP-AAFNFFG--ENI--DRK--R-YTSVTECL  NATLIHQSITGLYETRIDLSKL--  1343
WP_049516684    1283  LAK---SFI  SLLTFTAFGAP-AAFNFFG--ENI--DRK--R-YTSVTECL  NATLIHQSITGLYETRIDLSKL--  1343
GAD46167        1282  LAK---SFI  SLLTFTAFGAP-AAFNFFG--ENI--DRK--R-YTSVTECL  NATLIHQSITGLYETRIDLSKL--  1342
WP_018363470    1314  ISD---SFI  NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YNSTKECL  NATLIHQSITGLYETRIDLSKL--  1374
WP_003043819    1311  -SN---SFV  SLLKYTSFGAS-GGFTFLD--LDVkqGRL--R-YQTVTEVL  DATLIYQSITGLYETRTDLSQL--  1372
WP_006269658    1282  LAK---SFI  SLLTFTAFGAP-AAFNFFG--ENI--DRK--R-YTSVTECL  NATLIHQSITGLYETRIDLSKL--  1342
WP_048800889    1302  ISN---SFI  HLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL  NATLIHQSITGLYETQTDLSKL--  1362
WP_012767106    1309  -CS---SVI  NLLTLTASGAP-ADFKFLG--TTI--PRK--R-YGSPQSIL  SSTLIHQSITGLYETRIDLSQL--  1368
WP_014612333    1309  -CS---SVI  NLLTLTASGAP-ADFKFLG--TTI--PRK--R-YGSPQSIL  SSTLIHQSITGLYETRIDLSQL--  1368
WP_015017095    1309  -CS---SVI  NLLTLTASGAP-ADFKFLG--TTI--PRK--R-YGSPQSIL  SSTLIHQSITGLYETRIDLSQL--  1368
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WP_048327215    1309  -CS---SVI  NLLTLTASGAP-ADFKFLG--TTI--PRK--R-YGSPQSIL  SSTLIHQSITGLYETRIDLSQL--  1368
WP_049519324    1309  -CS---SVI  NLLTLTASGAP-ADFKFLG--TTI--PRK--R-YGSPQSIL  SSTLIHQSITGLYETRIDLSQL--  1368
WP_012515931    1286  -AE---NII  NVFTFVALGAP-AAFKFFD--ATI--DRK--R-YTSTKEVL  NATLIHQSVTGLYETRIDLSQL--  1345
WP_021320964    1286  -AE---NII  NVFTFVALGAP-AAFKFFD--ATI--DRK--R-YTSTKEVL  NATLIHQSVTGLYETRIDLSQL--  1345
WP_037581760    1286  -AE---NII  NVFTFVALGAP-AAFKFFD--ATI--DRK--R-YTSTKEVL  NATLIHQSVTGLYETRIDLSQL--  1345
WP_004232481    1309  ISSlseSFI  NLLKFISFGAP-GAFKFLK--LDV--KQSnlR-YKSTTEAL  SATLIHQSVTGLYETRIDLSKL--  1374
WP_009854540    1307  ISN---SFI  NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL  TATLIHQSITGLYETRIDLSKL--  1367
WP_012962174    1308  ISI---SFV  NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL  NATLIHQSITGLYETRIDLSKL--  1368
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WP_039695303  1309  ISN---SFI   NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL   NATLIHQ
WP_014334983  1312  ISA---SFI   NLLTLTALGAP-ADFNFLG--EKI--PRK--R-YTSTKECL   SATLIHQ
WP_003099269  1306  -AI---NML   NLFTFTDLGAP-SAFKFFN--GDI--DRK--R-YSSTNEII   NSTLIYQ
AHY15608              ---------   ----------------------------------------   -------
AHY17476              ---------   ----------------------------------------   -------
ESR09100      138   -AI---NML   NLFTFTDLGAP-SAFKFFNg--DI--DRK--R-YSSTNEII   NSTLIYQ
AGM98575              ---------   ----------------------------------------   -------
ALF27331      1282  LAS---SFI   NLLTFTAIGAP-AAFKFFD--NNI--DRK--R-YTSTTEIL   NATLIHQ
WP_018372492  1290  VAR---SFV   -LLNFTMMGAA-TDFKFFG--QII--PRK--R-YPSTTECL   KSTLIHQ
WP_045618028  1311  LSE---SFI   SLLKLISFGAP-GTFKFLG--VEI--SQSnvR-YQSVSSCF   NATLIHQ
WP_045635197  1307  LAN---SFI   NLLTFTALGAP-AAFKFFG--KDI--DRK--R-YTTVSEIL   NATLIHQ
WP_002263549  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002263887  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002264920  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002269043  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002269448  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002271977  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002272766  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002273241  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002275430  1282  LSS---SFI   NLLTFTAIGAP-AAFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002276448  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002277050  1288  LAD---GFI   KLLGFTQLGAT-SPFSFLG--IKL--NQK--Q-YTGKKDYL   EATLIHQ
WP_002277364  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002279025  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002279859  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002280230  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002281696  1282  LSS---SFI   NLLTFTAIGAP-AAFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002282247  1288  LAD---GFI   KLLGFTQLGAT-SPFSFLG--IKL--NQK--Q-YTGKKDYL   EATLIHQ
WP_002282906  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002283846  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002287255  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002288990  1282  LAS---SFI   NLLTFTAIGAP-AAFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002289641  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
WP_002290427  1282  LAS---SFI   NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL   NATLIHQ
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WP_002295753 1282 LAS---SFI NLLTFTAIGAP-AAFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
WP_002296423 1282 LAS---SFI NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
WP_002304487 1296 LAS---SFI NLLTFTAIGAP-AAFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLNKL-- 1356
WP_002305844 1282 LAS---SFI NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
WP_002307203 1282 LAS---SFI NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
WP_002310390 1282 LAS---SFI NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
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WP_012997688 1282 LAS---SFI NLLTFTAIGAP-ATFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
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WP_019314093 1282 LAS---SFI NLLTFTAIGAP-AAFKFFD--KNI--DRK--R-YTSTTEIL NATLIHQSITGLYETRIDLSKL-- 1342
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WP_000428612 1310 LAN---SFI NLLTFTALGAP-AAFKFFG--KDV--DRK--R-YTTVSEIL NATLIHQSITGLYETRIDLSKL-- 1370
WP_000428613 1308 LAN---SFI NLLTFTALGAP-AAFKFFG--KDI--DRK--R-YTTVSEIL NATLIHQSITGLYETRIDLSKL-- 1368
WP_049523028 1303 IAN---SFI NLLTFTAFGAP-AVFKFFG--KDI--ERK--R-YSTVTEIL KATLIHQSLTGLYETRIDLSKL-- 1363
WP_003107102 1275 -AT---NML NLFTFTGLGAP-ATLKFFN--VDI--DRK--R-YTSSTEIL NSTLIRQSITGLYETRIDLSKI-- 1334
WP_054279288 1304 -SI---SFL NLFKFTSFGAP-EKFTFLN--SEIkqDDV--R-YRSTKECL NSTLIHQSVTGLYETRIDLSQF-- 1365
WP_049531101 1311 LSE---SFI SLLKLTSFGAP-GAFRFLG--VEI--SQSnvR-YQSVSSCF NATLIHQSITGLYETRIDLSKL-- 1373
WP_049538452 1311 LSE---SFI SLLKLTSFGAP-GAFKFLG--VEI--SQSsvR-YKPNSQFL DATLIHQSITGLYETRIDLSKL-- 1373
WP_049549711 1313 LSE---SFI SLLKLTSFGAP-GAFKFLG--AEI--SQSsvR-YKPNSQFL DTTLIHQSITGLYETRIDLSKL-- 1375
WP_007896501 1312 -AL---NML NLFIFTSLGAP-STFVFFD--ETI--DRK--R-YTTSSDVL NGILIQQSITGLYETRIDLSRF-- 1371
EFR44625     1264 -AL---NML NLFIFTSLGAP-STFVFFD--ETI--DRK--R-YTTSSDVL NGILIQQSITGLYETRIDLSRF-- 1323
WP_002897477 1307 LAN---SFI NLLTFTALGAP-AAFKFFG--KDV--DRK--R-YTTVSEIL NATLIHQSITGLYETRIDLSKL-- 1367
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WP_009729476 1308 LAN---SFI NLLTFTALGAP-AAFKFFG--KDV--DRK--R-YTTVSEIL NATLIHQSITGLYETRIDLSKL-- 1368
CQR24647     1296 LAQ---SFI SLLKFTAFGAP-GGFKFLD--ADI--KQSnlR-YQTVTEVL SSTLIHQSVTGLYETRIDLSKL-- 1358
WP_000066813 1312 LAN---SFI NLLTFTALGAP-AAFKFLG--KDV--DRK--R-YTTVSEIL NATLIHQSITGLYETRIDLSKL-- 1372
WP_009754323 1308 LAN---SFI NLLTFTALGAP-AAFKFFG--KDV--DRK--R-YTTVSEIL NATLIHQSITGLYETRIDLSKL-- 1368
WP_044674937 1301 LTS---SFV NLLTFTAIGAP-AAFKFLG--SVI--DRK--R-YTSIAEIL EATLIHQSVTGLYETRIDLSKL-- 1361
WP_044676715 1303 LTS---SFV NLLTFTAIGAP-AAFKFLG--SVI--DRK--R-YTSIAEIL EATLIHQSVTGLYETRIDLSKL-- 1363
WP_044680361 1303 LTS---SFV NLLTFTAIGAP-AAFKFLG--SVI--DRK--R-YTSIAEIL EATLIHQSVTGLYETRIDLSKL-- 1363
WP_044681799 1301 LTS---SFV NLLTFTAIGAP-AAFKFLG--SVI--DRK--R-YTSIAEIL EATLIHQSVTGLYETRIDLSKL-- 1361
WP_049533112 1314 ICT---SFL GLFELTSLGSA-SDFEFLG--VKI--PRY--RdYTPSSLLK DSTLIHQSITGLYETRIDLSKL-- 1383
WP_029090905 1241 -VK---VI ELLKITQANATnGDLKLLK---M-sNREg-R-LGSVSVAL DFKIINQSVTGLYQSIEDYNN--- 1300
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AIT42264     1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1365
WP_034440723 1277 LVE---SFI NLLAITKCGPA-ADITFLG--EKI--SRK--R-YRSTNCLW GSEVIFQSPTGLYETRLRLE---- 1335
AKQ21048     1306 -AE---NII HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL DATLIHQSITGLYETRIDLSQL-- 1365
WP_004636532 1272 TVE---SFV NLMTFTAMGAP-ATFKFYG--ESI--TRS--R-YTSITEFR GSTLIFQSITGLYETRYKL----- 1329
WP_002364836 1278 IAA---SFI QLMQFNAMGAP-STFKFFQ--KDI--ERA--R-YTSIKEIF DATIIYQSTTGLYETRRKV----- 1335
WP_016631044 1229 IAA---SFI QLMQFNAMGAP-STFKFFQ--KDI--ERA--R-YTSIKEIF DATIIYQSPTGLYETRRKV----- 1286
EMS75795     1014 LSQ---SFI NLMQLNAMGAP-ADFKFFD--VTI--PRK--R-YPSLTEIW ESTIIYQSITGLRETRTRMATLwd 1076
WP_002373311 1278 IAA---SFI QLMQFNAMGAP-STFKFFQ--KDI--ERA--R-YTSIKEIF DATIIYQSTTGLYETRRKV----- 1335
WP_002378009 1278 IAA---SFI QLMQFNAMGAP-STFKFFQ--KDI--ERA--R-YTSIKEIF DATIIYQSTTGLYETRRKV----- 1335
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WP_033789179 1278 IAA---SFI QLMQFNAMGAP-STFKFFQ--KDI--ERA--R-YTSIKEIF DATIIYQSTTGLYETRRKV----- 1335
WP_002310644 1280 LAS---SFV NLMQFNAMGAP-ADFKFFD--VTI--PRK--R-YTSLTEIW QSTIIHQSITGLYETRIRMGK--- 1339
WP_002312694 1281 LAS---SFV NLMQFNAMGAP-ADFKFFD--VTI--PRK--R-YTSLTEIW QSTIIHQSITGLYETRIRMGK--- 1340
WP_002314015 1281 LAS---SFV NLMQFNAMGAP-ADFKFFD--VTI--PRK--R-YTSLTEIW QSTIIHQSITGLYETRIRMGK--- 1340
WP_002320716 1281 LAS---SFV NLMQFNAMGAP-ADFKFFD--VTI--PRK--R-YTSLTEIW QSTIIHQSITGLYETRIRMGK--- 1340
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WP_002335161	1281	LAS---SFV	NLMQFNAMGAP-ADFKEFD--VTI--PRK--R-YTSLTEIW	QSTIIHQSITGLYETRIRMGK---	1340
WP_002345439	1281	LAS---SFV	NLMQFNAMGAP-ADFKEFD--VTI--PRK--R-YTSLTEIW	QSTIIHQSITGLYETRIRMGK---	1340
WP_034867970	1271	IAQ---SFL	QLLQFNAIGAP-ADFKEFG--VTI--PRK--R-YTSLTEIW	DATIIYQSVTGLYETRIRMGDLwa	1333
WP_047937432	1281	LAS---SFV	NLMQFNAMGAP-ADFKEFD--VTI--PRK--R-YTSLTEIW	QSTIIHQSITGLYETRIRMGK---	1340
WP_010720994	1271	IAQ---SFL	QLLQFNAIGAP-ADFKEFG--VTI--PRK--R-YTSLTEIW	DATIIYQSVTGLYETRIRMGDLwa	1333
WP_010737004	1271	IAQ---SFL	QLLQFNAIGAP-ADFKEFG--VTI--PRK--R-YTSLTEIW	DATIIYQSVTGLYETRIRMGDLwa	1333
WP_034700478	1271	IAQ---SFL	QLLQFNAIGAP-ADFKEFG--VTI--PRK--R-YTSLTEIW	DATIIYQSVTGLYETRIRMGDLwa	1333
WP_007209003	1270	IAK---SFI	NLLTFTAMGAP-ADFEEFG--EKI--PRK--R-YVSISEII	DAVFIHQSITGLYETRVRLTEV--	1330
WP_023519017	1265	MID---AIL	SLLKFTLFGAS-VEFKFFD--IKI---LK--R-YKSLTDIW	EATIIYQSVTGLYERRVEVRKLwd	1326
WP_010770040	1275	IAE---SFV	NLMKFSAYGAP-MDFKFFG--KTI--PRS--R-YTSVGELL	SATIINQSITGLYETRRKL-----	1332
WP_048604708	1271	IAQ---SFV	DLMQLNAFGAP-ADFKFFG--ETI--PRK--R-YTSVNELL	EATIINQSITGLYETRRRL-----	1328
WP_010750235	1274	MAH---SFV	NLMQFNALGAP-ADFKEFD--TTI--TRK--R-YTSLTEIW	QSTIIYQSVTGLYETRRRMADLwd	1336
AII16583	1345	-AE---NII	HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL	DATLIHQSITGLYETRIDLSQL--	1404
WP_029073316	1283	-CE---VI	QMLVVMHAGPQnGNITFDDf--KL-sNRLg-R-LNCKTISL	TTVFIADSPTGMYSKKYKL-----	1343
WP_031589969	1284	-CN---II	QILATLHCNSSiGKIMYSDf--KI-sTTIg-R-LNGRTISL	DISFIAESPTGMYSKKYKL-----	1344
KDA45870	1254	LGK---NFV	ELLRYTADGAA-SDFKFFG--ENI--PRK--R-YNSAGSLL	NGTLIYQSKTGLYETRIDLGKL--	1314
WP_039099354	1307	ILDr----V	-LIGLHANAAV-SDLGVLKisTPL--GKM--Q---QPSGIS	DTQIIYQSPTGLFERRVALRDL--	1368
AKP02966	1297	INSl-eELI	TLLHANSTSAH-LIFNNIE-kKAF--GRK---------THGLT	DTDFIYQSVTGLYETRIHIE----	1356
WP_010991369	1275	IAQ---SFV	DLMAFNAMGAP-ASFKEFE--TTI--ERK--R-YNNLKELL	NSTIIYQSITGLYESRKRL-----	1332
WP_033838504	1275	IAQ---SFV	DLMAFNAMGAP-ASFKEFE--TTI--ERK--R-YNNLKELL	NSTIIYQSITGLYESRKRL-----	1332
EHN60060	1278	IAQ---SFV	DLMAFNAMGAP-ASFKFFE--TTI--ERK--R-YNNLKELL	NSTIIYQSITGLYESRKRL-----	1335
EFR89594	1044	IAQ---SFV	DLMAFNAMGAP-ASFKFFE--TTI--ERK--R-YNNLKELL	NSTIIYQSITGLYESRKRL-----	1101
WP_038409211	1275	IAK---SFD	KLKVFNAFGAP-RDFEFFE--TTI--KRK--R-YYNIKELL	NATIIYQSITGLYEARKRL-----	1332
EFR95520	894	IAK---SFD	KLKVFNAFGAP-RDFEFFE--TTI--KRK--R-YYNIKELL	NATIIYQSITGLYEARKRL-----	951
WP_003723650	1275	IAQ---SFV	DLMAFNAMGAP-ASFKFFE--ATI--DRK--R-YTNLKELL	SSTIIYQSITGLYESRKRL-----	1332
WP_003727705	1275	IAQ---SFV	DLMAFNAMGAP-ASFKFFE--ATI--DRK--R-YTNLKELL	SSTIIYQSITGLYESRKRL-----	1332
WP_003730785	1275	IAQ---SFV	DLMAFNAMGAP-ASFKFFE--ATI--DRK--R-YTNLKELL	SSTIIYQSITGLYESRKRL-----	1332
WP_003733029	1275	IAE---SFV	SLKKFNAFGVH-QDFSFFG--TKI--ERK--R-DRKLNELL	NSTIIYQSITGLYESRKRL-----	1332
WP_003739838	1275	IAQ---SFV	NLMAFNAMGAP-ASFKFFE--ATI--ERK--R-YTNLKELL	SATIIYQSITGLYEARKRL-----	1332
WP_014601172	1275	IAQ---SFV	DLMAFNAMGAP-ASFKEFE--ATI--DRK--R-YTNLKELL	SSTIIYQSITGLYESRKRL-----	1332
WP_023548323	1275	IAE---SFV	SLKKFNAFGVH-KDFNEFG--TTI--KRK--R-DRKLKELL	NSTIIYQSITGLYESRKRL-----	1332
WP_031665337	1275	IAQ---SFV	DLMAFNAMGAP-ASFKFFE--ATI--DRK--R-YTNLKELL	SSTIIYQSITGLYESRKRL-----	1332
WP_031669209	1275	IAE---SFV	SLKKFNAFGVH-QDFSFFG--TKI--ERK--R-DRKLNELL	NSTIIYQSITGLYESRKRL-----	1332

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WP_033920898  1275  IAE---SFV   SLKKFNAFGVH-KDFNFFG--TTI--KRK--R-DRKLKELL   NSTIIYQ
AKI42028      1278  IAQ---SFV   DLMAFNAMGAP-ASFKFFE--ATI--DRK--R-YTNLKELL   SSTIIYQ
AKI50529      1278  IAE---SFV   SLKKFNAFGVH-KDFNFFG--TTI--KRK--R-DRKLKELL   NSTIIYQ
EFR83390      723   IAQ---SFV   DLMVFNAMGAP-ASFKFFE--ATI--DRK--R-YTNLKELL   SSTIIYQ
WP_046323366  1275  IAQ---SFV   NLLEFNAMGAP-ASFKYFE--TNI--ERK--R-YNNLKELL   NATIIYQ
AKE81011      1322  -AE---NII   HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL   DATLIHQ
CUO82355      1273  -AN----VI   QMLIIMHKGPQnGNIIYDDf--NV-gKRIg-R-LNGRTFYL   NIEFISQ
WP_033162887  1275  -CD----VI   QILIIMHAGPMnGNIMYDDf--KF-tNRIg-R-FTHKNIDL   KTTFIST
AGZ01981      1339  -AE---NII   HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL   DATLIHQ
AKA60242      1306  -AE---NII   HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL   DATLIHQ
AKS40380      1306  -AE---NII   HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL   DATLIHQ
4UN5_B        1310  -AE---NII   HLFTLTNLGAP-AAFKYFD--TTI--DRK--R-YTSTKEVL   DATLIHQ
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WP_045635197  1368  GED   1370
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WP_033888930  1191  GGD   1193
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KGE60162       541  GGD    543
KGE60856       304  GGD    306
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WP_001040076  1368  GED   1370
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CUO82355           ---
WP_033162887        ---
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AKS40380      1366  GGD    1376
4UN5_B        1370  GGD    1372
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Table 2. T to C changes with NGG PAM. Table 2 shows a list of T to C mutations that may be corrected using any of the base editors provided herein. GRNAs and gRNAall indicate the protospacer and PAM sequence, where the PAM sequence is the last 3 nucleotides of each of the sequences in GRNAs and gRNAall.

Name	Gene ID	Gene Symbol	SEQ ID NO:	Flanks	SEQ ID NO:	GRNAs	SEQ ID NO:	gRNAall
NM_000071.2(CBS):c.833T>C (p.Ile278Thr)	875	CBS	2540	['CTGAAGCCGCGCCCTCTGCAGATCAYTGGGGTGGATCCCGAAGGGTCCATC']	2703-2704	['ATCAYTGGGGTGGATCCCGAAGG', 'TCAYTGGGGTGGATCCCGAAGGG']	2907-2908	['ATCAYTGGGGTGGATCCCGAAGG', 'TCAYTGGGGTGGATCCCGAAGGG']
NM_001385.2(DPYS):c.1078T>C (p.Trp360Arg)	1807	DPYS	2541	['TGTTGAAGATCGGATGTCCGTAATAYGGGAAAAAGGCGTGGTGGGTTTCAC']	2705-2707	['CGTAATAYGGGAAAAAGGCGTGG', 'AATAYGGGAAAAAGGCGTGGTGG', 'ATAYGGGAAAAAGGCGTGGTGGG']	2909-2911	['CGTAATAYGGGAAAAAGGCGTGG', 'AATAYGGGAAAAAGGCGTGGTGG', 'ATAYGGGAAAAAGGCGTGGTGGG']
NM_000027.3(AGA):c.916T>C (p.Cys306Arg)	175	AGA	2542	['TCCAGAATTCTTTGGGGCTGTTATAYGTGCCAATGTGACTGGAAGTTACGG']	2708	['GTTATAYGTGCCAATGTGACTGG']	2912	['GTTATAYGTGCCAATGTGACTGG']
NM_000035.3(ALDOB):c.442T>C (p.Trp148Arg)	229	ALDOB	2543	['GAAAGATGGTGTTGACTTTGGGAAGYGGCGTGCTGTGCTGAGGATTGCCGA']	2709	['GGAAGYGGCGTGCTGTGCTGAGG']	2913	['GGAAGYGGCGTGCTGTGCTGAGG']
NM_173560.3(RFX6):c.380+2T>C	222546	RFX6	2544	['GCAGACACAGCTCACGCTGCAGTGGYGAGACTCGCCCGCAGGGTACACTGA']	2710-2711	['CAGTGGYGAGACTCGCCCGCAGG', 'AGTGGYGAGACTCGCCCGCAGG']	2914-2915	['CAGTGGYGAGACTCGCCCGCAGG', 'AGTGGYGAGACTCGCCCGCAGGG']
NM_153704.5(TMEM67):c.1843T>C (p.Cys615Arg)	91147	TMEM67	2545	['AGAACGTTTTGTCACTTATGTTGGAHGTGCCTTTGCTCTGAAGGTAAGTTT']	2712	['TGGAHGTGCCTTTGCTCTGAAGG']	2916	['TGGAHGTGCCTTTGCTCTGAAGG']

NM_000124.3(ERCC6):c.2960T>C (p.Leu987Pro)	2074	ERCC6	2546	['AAGCAGTTTTTGACAAATAGAGTGCYAAAAGACCCAAAACAAAGGCGGTTT']	2713	['TGCYAAAAGACCCAAAACAAAGG']	2917	['TGCYAAAAGACCCAAAACAAAGG']
NM_020435.3(GJC2):c.857T>C (p.Met286Thr)	57165	GJC2	2547	['TGCCTGCTGCTCAACCTCTGTGAGAYGGCCCACCTGGGCTTGGGCAGCGCG']	2714	['TGAGAYGGCCCACCTGGGCTTGG']	2918-2919	['TGAGAYGGCCCACCTGGGCTTGG', 'GAGAYGGCCCACCTGGGCTTGGG']
NM_000920.3(PC):c.434T>C (p.Val145Ala)	5091	PC	2548	['CGGTTTATTGGGCCAAGCCCAGAAGBGGTCCGCAAGATGGGAGACAAGGTG']	2715	['CCAGAAGBGGTCCGCAAGATGGG']	2920	['CCAGAAGBGGTCCGCAAGATGGG']
NM_000026.2(ADSL):c.674T>C (p.Met225Thr)	158	ADSL	2549	['TCCAAGGTAGAGCAGCTTGACAAGAYGGTGACAGAAAAGGCAGGATTTAAG']	2716	['AAGAYGGTGACAGAAAAGGCAGG']	2921	['AAGAYGGTGACAGAAAAGGCAGG']
NM_000391.3(TPP1):c.1093T>C (p.Cys365Arg)	1200	TPP1	2550	['TCTCTCAGGTGACAGTGGGGCCGGGYGTTGGTCTGTCTCTGGAAGACACCA']	2717	['GCCGGGYGTTGGTCTGTCTCTGG']	2922	['GCCGGGYGTTGGTCTGTCTCTGG']
NM_004183.3(BEST1):c.704T>C (p.Val235Ala)	7439	BEST1	2551	['TACGACTGGATTAGTATCCCACTGGYGTATACACAGGTGAGGACTAGGCTG']	2718	['CACTGGYGTATACACAGGTGAGG']	2923	['CACTGGYGTATACACAGGTGAGG']
NM_000019.3(ACAT1):c.935T>C (p.Ile312Thr)	38	ACAT1	2552	['CTCAATGTTACACCACTGGCAAGAAYAGTAGGTAAGGCCAGGCGAGGTGGC']	2719	['CAAGAAYAGTAGGTAAGGCCAGG']	2924	['CAAGAAYAGTAGGTAAGGCCAGG']
NM_000543.4(SMPD1):c.911T>C (p.Leu304Pro)	6609	SMPD1	2553	['CGGGCCCTGACCACCGTCACAGCACYTGTGAGGAAGTTCCTGGGGCCAGTG']	2720	['CACYTGTGAGGAAGTTCCTGGGG']	2925-2927	['AGCACYTGTGAGGAAGTTCCTGG', 'GCACYTGTGAGGAAGTTCCTGGG', 'CACYTGTGAGGAAGTTCCTGGGG']
NM_000527.4(	3949	LDLR	2554	['ACAAATCTGA	2721	['CGGYATGGGCGG	2928	['ACTGCGGYATGG

LDLR):c.694+2T>C				CGAGGAAAACTGCGGYATGGGCGGGGCCAGGGTGGGGGCGG']		GGCCAGGGTGG']	-2930	GCGGGGCCAGG', 'CTGCGGYATGGGCGGGGCCAGGG', 'CGGYATGGGCGGGGCCAGGGTGG']
NM_012464.4(TLL1):c.713T>C (p.Val238Ala)	7092	TLL1	2555	['AAGAACTGTGATAAATTTGGGATTGYTGTTCATGAATTGGGTCATGTGATA']	2722	['GGGATTGYTGTTCATGAATTGGG']	2931	['GGGATTGYTGTTCATGAATTGGG']
NM_000112.3(SLC26A2):c.-26+2T>C	1836	SLC26A2	2556	['CCTGCAGCGGCCCGGACCCGAGAGGYGAGAAGAGGGAAGCGGACCAGGGAA']	2723	['GAGAGGYGAGAAGAGGGAAGCGG']	2932	['GAGAGGYGAGAAGAGGGAAGCGG']
NM_001005741.2(GBA):c.751T>C (p.Tyr251His)	2629	GBA	2557	['CATCTACCACCAGACCTGGGCCAGAYACTTTGTGAAGTAAGGGATCAGCAA']	2724	['GCCAGAYACTTTGTGAAGTAAGG']	2933-2934	['GCCAGAYACTTTGTGAAGTAAGG', 'CCAGAYACTTTGTGAAGTAAGGG']
NM_020365.4(EIF2B3):c.1037T>C (p.Ile346Thr)	8891	EIF2B3	2558	['CCACCAGTCCATTCGTCAGCCCAGAYGTCAGCAAACACCTGGTAAGTGCT']	2725	['CCAGAYGTCAGCAAACACCTGG']	2935	['CCAGAYGTCAGCAAACACCTGG']
NM_022041.3(GAN):c.1268T>C (p.Ile423Thr)	8139	GAN	2559	['TGCTATGCAGCTATGAAAAAGAAAAYCTACGCCATGGGTGGAGGCTCCTAC']	2726	['AAGAAAAYCTACGCCATGGGTGG']	2936-2937	['AAGAAAAYCTACGCCATGGGTGG', 'AAAAYCTACGCCATGGGTGGAGG']
NM_054027.4(ANKH):c.143T>C (p.Met48Thr)	56172	ANKH	2560	['GCTGTCAAGGAGGATGCAGTCGAGAYGCTGGCCAGCTACGGGCTGGCGTAC']	2727-2728	['GTCGAGAYGCTGGCCAGCTACGG', 'TCGAGAYGCTGGCCAGCTACGGG']	2938-2939	['GTCGAGAYGCTGGCCAGCTACGG', 'TCGAGAYGCTGGCCAGCTACGGG']
NM_006329.3(FBLN5):c.506T>C (p.Ile169Thr)	10516	FBLN5	2561	['TTGCTTGCATTTCTGTTTCCAGACAYTGATGAATGTCGCTATGGTTACTGC']	2729	['GACAYTGATGAATGTCGCTATGG']	2940	['GACAYTGATGAATGTCGCTATGG']
NM_004086.2(	-1	-	2562	['GCACCTCTGG	2730	['AGATAYGGCTTC	2941	['AGATAYGGCTTC

COCH):c.1535T>C (p.Met512Thr)				ATGACCTGAAAGATAYGGCTTCTAAACCGAAGGAGTCTCAT']		TAAACCGAAGG']		TAAACCGAAGG']
NM_002942.4(ROBO2):c.2834T>C (p.Ile945Thr)	6092	ROBO2	2563	['AATAGCAACAGTGGCCCAAATGAGAYTGGAAATTTTGGCCGTGGGAGGTAAG']	2731	['GAGAYTGGAAATTTGGCCGTGG']	2942	['GAGAYTGGAAATTTGGCCGTGG']
NM_001300.5(KLF6):c.190T>C (p.Trp64Arg)	1316	KLF6	2564	['CAAATTTGACAGCCAGGAAGATCTGYGGACCAAAATCATTCTGGCTCGGGA']	2732	['TCTGYGGACCAAAATCATTCTGG']	2943	['TCTGYGGACCAAAATCATTCTGG']
NM_030653.3(DDX11):c.2271+2T>C	1663	DDX11	2565	['CTGGCATATTCCAGGTGCATCCAGGYGCGGGCGTCATGCTGGGCTTGGGTC']	2733	['TCCAGGYGCGGGCGTCATGCTGG']	2944-2945	['TCCAGGYGCGGGCGTCATGCTGG', 'CCAGGYGCGGGCGTCATGCTGGG']
NM_001451.2(FOXF1):c.1138T>C (p.Ter380Arg)	2294	FOXF1	2566	['CCAAGACATCAAGCCTTGCGTGATGYGAGGCTGCCGCCGCAGGCCCTCCTG']	2734	['TGATGYGAGGCTGCCGCCGCAGG']	2946	['TGATGYGAGGCTGCCGCCGCAGG']
NM_000435.2(NOTCH3):c.1363T>C (p.Cys455Arg)	4854	NOTCH3	2567	['CCTCGACCGCATAGGCCAGTTCACCYGTATCTGTATGGCAGGTGGGTGGTG']	2735	['ACCYGTATCTGTATGGCAGGTGG']	2947-2948	['TTCACCYGTATCTGTATGGCAGG', 'ACCYGTATCTGTATGGCAGGTGG']
NM_002427.3(MMP13):c.272T>C (p.Met91Thr)	4322	MMP13	2568	['CTTGACGATAACACCTTAGATGTCAYGAAAAAGCCAAGATGCGGGGTTCCT']	2736-2737	['GTCAYGAAAAAGCCAAGATGCGG', 'TCAYGAAAAAGCCAAGATGCGGG']	2949-2950	['GTCAYGAAAAAGCCAAGATGCGG', 'TCAYGAAAAAGCCAAGATGCGGG']
NM_000211.4(ITGB2):c.446T>C (p.Leu149Pro)	3689	ITGB2	2569	['GATGACCTCAGGAATGTCAAGAAGCYAGGTGGCGACCTGCTCCGGGCCCTC']	2738	['AGCYAGGTGGCGACCTGCTCCGG']	2951	['AGCYAGGTGGCGACCTGCTCCGG']
NM_005502.3(ABCA1):c.4429T>C (p.Cys1477Arg)	19	ABCA1	2570	['CAAAATCAAGAAGATGCTGCCTGTGYGTCCCCCAGGGGCAGGGGGGCTGCC']	2739-2740	['CCTGTGYGTCCCCCAGGGGCAGG', 'CTGTGYGTCCCCCAGGGGCAGGG']	2952-2955	['CCTGTGYGTCCCCCAGGGGCAGG', 'CTGTGYGTCCCCCAGGGGCAGGG', 'TGTGYGTCCCCCA

								GGGGCAGGGG', 'GTGYGTCCCCCA GGGGCAGGGGG']
m.12297T>C	4568	MT-TL2	2571	['AAAGGATAAC AGCTATCCATT GGTCYTAGGCC CCAAAAATTTT GGTGCAAC']	2741	['GTCYTAGGCCCC AAAAATTTTGG']	2956	['GTCYTAGGCCCC AAAAATTTTGG']
m.4290T>C	4565	MT-TI	2572	['AAATATGTCT GATAAAAGAG TTACTYTGATA GAGTAAATAA TAGGAGCTTA']	2742	['ACTYTGATAGAG TAAATAATAGG']	2957	['ACTYTGATAGAG TAAATAATAGG']
m.4291T>C	4565	MT-TI	2573	['AATATGTCTG ATAAAAGAGT TACTTYGATAG AGTAAATAAT AGGAGCTTAA']	2743	['ACTTYGATAGAG TAAATAATAGG']	2958	['ACTTYGATAGAG TAAATAATAGG']
m.3394T>C	4535	MT-ND1	2574	['GCTTACCGAA CGAAAAATTCT AGGCYATATA CAACTACGCA AAGGCCCCAA']	2744	['GGCYATATACAA CTACGCAAAGG']	2959	['GGCYATATACAA CTACGCAAAGG']
NM_002764.3(PRPS1):c.344T>C (p.Met115Thr)	5631	PRPS1	2575	['ATCTCAGCCA AGCTTGTTGCA AATAYGCTATC TGTAGCAGGTG CAGATCAT']	2745	['GCAAATAYGCTA TCTGTAGCAGG']	2960	['GCAAATAYGCTA TCTGTAGCAGG']
NM_000132.3(F8):c.5372T>C (p.Met1791Thr)	2157	F8	2576	['AGAGCAGAA GTTGAAGATA ATATCAYGGTG AGTTAAGGAC AGTGGAATTAC ']	2746	['TCAYGGTGAGTT AAGGACAGTGG']	2961	['TCAYGGTGAGTT AAGGACAGTGG']
NM_000132.3(F8):c.1754T>C (p.Ile585Thr)	2157	F8	2577	['CCTTTCAATA TATGTAATTAA CAGAYAATGT CAGACAAGAG GAATGTCATC']	2747	['AACAGAYAATGT CAGACAAGAGG']	2962	['AACAGAYAATGT CAGACAAGAGG']
NM_000133.3(F9):c.1328T>C (p.Ile443Thr)	2158	F9	2578	['TGTGCAATGA AAGGCAAATA TGGAAYATAT ACCAAGGTATC CCGGTATGTC']	2748	['GAAYATATACCA AGGTATCCCGG']	2963	['GAAYATATACCA AGGTATCCCGG']

NM_000169.2(GLA):c.806T>C (p.Val269Ala)	-1	-	2579	['TTATTTCATTCTTTTTCTCAGTTAGYGATTGGCAACTTTGGCCTCAGCTGG']	2749	['CAGTTAGYGATTGGCAACTTTGG']	2964	['CAGTTAGYGATTGGCAACTTTGG']
NM_000116.4(TAZ):c.352T>C (p.Cys118Arg)	6901	TAZ	2580	['CTCCCACTTCTTCAGCTTGGGCAAGYGTGTGCCTGTGTGCCGAGGTGAGCT']	2750	['AAGYGTGTGCCTGTGTGCCGAGG']	2965	['AAGYGTGTGCCTGTGTGCCGAGG']
NM_000061.2(BTK):c.2T>C (p.Met1Thr)	695	BTK	2581	['GGTGAACTCCAGAAAGAAGAAGCTAYGGCCGCAGTGATTCTGGAGAGCATC']	2751	['AGCTAYGGCCGCAGTGATTCTGG']	2966	['AGCTAYGGCCGCAGTGATTCTGG']
NM_000061.2(BTK):c.1223T>C (p.Leu408Pro)	695	BTK	2582	['AAGGACCTGACCTTCTTGAAGGAGCYGGGGACTGGACAATTTGGGGGTAGTG']	2752	['AGCYGGGGACTGGACAATTTGGG']	2967 - 2968	['GAGCYGGGGACTGGACAATTTGG', 'AGCYGGGGACTGGACAATTTGGG']
NM_000061.2(BTK):c.1741T>C (p.Trp581Arg)	695	BTK	2583	['CAAGTTCAGCAGCAAATCTGACATTYGGGCTTTTGGTAAGTGGATAAGATT']	2753	['ACATTYGGGCTTTTGGTAAGTGG']	2969	['ACATTYGGGCTTTTGGTAAGTGG']
NM_014009.3(FOXP3):c.970T>C (p.Phe324Leu)	50943	FOXP3	2584	['GATTCATCCCCACCCTCTGACAGAGYTCCTCCACAACATGGACTACTTCAA']	2754	['GACAGAGYTCCTCCACAACATGG']	2970	['GACAGAGYTCCTCCACAACATGG']
NM_003688.3(CASK):c.2740T>C (p.Trp914Arg)	8573	CASK	2585	['TGAGCTCGTGTGCACAGCCCCACAGYGGGTCCCTGTCTCCTGGGTCTATTA']	2755 - 2756	['CACAGYGGGTCCCTGTCTCCTGG', 'ACAGYGGGTCCCTGTCTCCTGGG']	2971 - 2972	['CACAGYGGGTCCCTGTCTCCTGG', 'ACAGYGGGTCCCTGTCTCCTGGG']
NM_004992.3(MECP2):c.464T>C (p.Phe155Ser)	4204	MECP2	2586	['GACACATCCCTGGACCCTAATGATTBTGACTTCACGGTAACTGGGGAGAGGG']	2757	['GATTBTGACTTCACGGTAACTGG']	2973 - 2974	['GATTBTGACTTCACGGTAACTGG', 'ATTBTGACTTCACGGTAACTGGG']
NM_000431.3(MVK):c.803T>C (p.Ile268Thr)	4598	MVK	2587	['ATCGTGGCCCCCCTCCTGACCTCAAYAGATGCCATCTCCCTG	2758	['CTCAAYAGATGCCATCTCCCTGG']	2975	['CTCAAYAGATGCCATCTCCCTGG']

				GAGTGTGAG']				
NM_021961.5(TEAD1):c.1261T>C (p.Tyr?His)	7003	TEAD1	2588	['TGAACACGGAGCACAACATCATATTYACAGGCTTGTAAAGGACTGAACATG']	2759	['TCATATTYACAGGCTTGTAAAGG']	2976	['TCATATTYACAGGCTTGTAAAGG']
NM_005633.3(SOS1):c.1294T>C (p.Trp432Arg)	6654	SOS1	2589	['CGAGATTCAGAAGAATATTGATGGTYGGGAGGGAAAAGACATTGGACAGTG']	2760	['GGTYGGGAGGGAAAAGACATTGG']	2977	['GGTYGGGAGGGAAAAGACATTGG']
NM_006920.4(SCN1A):c.3577T>C (p.Trp1193Arg)	-1	-	2590	['TGTGGAAGAAGGCAGAGGAAAACAAYGGTGGAACCTGAGAAGGACGTGTTT']	2761	['AACAAYGGTGGAACCTGAGAAGG']	2978	['AACAAYGGTGGAACCTGAGAAGG']
NM_000141.4(FGFR2):c.1018T>C (p.Tyr340His)	2263	FGFR2	2591	['TGTAACTTTTGAGGACGCTGGGGAAYATACGTGCTTGGCGGGTAATTCTAT']	2762 - 2763	['TGGGGAAYATACGTGCTTGGCGG', 'GGGGAAYATACGTGCTTGGCGGG']	2979 - 2980	['TGGGGAAYATACGTGCTTGGCGG', 'GGGGAAYATACGTGCTTGGCGGG']
NM_000174.4(GP9):c.70T>C (p.Cys24Arg)	2815	GP9	2592	['GGCCACCAAGGACTGCCCCAGCCCAYGTACCTGCCGCGCCCTGGAAACCAT']	2764	['CCCAYGTACCTGCCGCGCCCTGG']	2981	['CCCAYGTACCTGCCGCGCCCTGG']
NM_000175.3(GPI):c.1574T>C (p.Ile525Thr)	2821	GPI	2593	['CTGGGAAAGCAGCTGGCTAAGAAAABAGAGCCTGAGCTTGATGGCAGTGCT']	2765	['AAAABAGAGCCTGAGCTTGATGG']	2982	['AAAABAGAGCCTGAGCTTGATGG']
NM_000315.2(PTH):c.52T>C (p.Cys18Arg)	5741	PTH	2594	['AGTTATGATTGTCATGTTGGCAATTYGTTTTCTTACAAAATCGGATGGGAA']	2766	['AATTYGTTTTCTTACAAAATCGG']	2983	['AATTYGTTTTCTTACAAAATCGG']
NM_000222.2(KIT):c.1676T>C (p.Val559Ala)	3815	KIT	2595	['CCCATGTATGAAGTACAGTGGAAGGNTGTTGAGGAGATAAATGGAAACAAT']	2767	['AAGGNTGTTGAGGAGATAAATGG']	2984	['AAGGNTGTTGAGGAGATAAATGG']

NM_016835.4(MAPT):c.1839T>C (p.Asn613=)	4137	MAPT	2596	['AGTCCAAGTGTGGCTCAAAGGATAAYATCAAACACGTCCCGGGAGGCGGCA']	2768 - 2769	['GGATAAYATCAAACACGTCCCGG', 'GATAAYATCAAACACGTCCCGGG']	2985 - 2986	['GGATAAYATCAAACACGTCCCGG', 'GATAAYATCAAACACGTCCCGGG']
NM_170707.3(LMNA):c.1139T>C (p.Leu380Ser)	4000	LMNA	2597	['GAGATCCACGCCTACCGCAAGCTCTYGGAGGGCGAGGAGGAGAGGTGGGCT']	2770	['TCTYGGAGGGCGAGGAGGAGAGG']	2987	['TCTYGGAGGGCGAGGAGGAGAGG']
NM_000424.3(KRT5):c.20T>C (p.Val7Ala)	3852	KRT5	2598	['GCCACCATGTCTCGCCAGTCAAGTGYGTCCTTCCGGAGCGGGGGCAGTCGT']	2771 -- 2773	['TCAAGTGYGTCCTTCCGGAGCGG', 'CAAGTGYGTCCTTCCGGAGCGGG', 'AAGTGYGTCCTTCCGGAGCGGGG']	2988 - 2991	['TCAAGTGYGTCCTTCCGGAGCGG', 'CAAGTGYGTCCTTCCGGAGCGGG', 'AAGTGYGTCCTTCCGGAGCGGGG', 'AGTGYGTCCTTCCGGAGCGGGGG']
NM_000184.2(HBG2):c.125T>C (p.Phe42Ser)	3048	HBG2	2599	['GTTGTCTACCCATGGACCCAGAGGTYCTTTGACAGCTTTGGCAACCTGTCC']	2774	['CAGAGGTYCTTTGACAGCTTTGG']	2992	['CAGAGGTYCTTTGACAGCTTTGG']
NM_000515.4(GH1):c.291+6T>C	2688	GH1	2600	['AGGAAACACAACAGAAATCCGTGAGYGGATGCCTTCTCCCCAGGCGGGGAT']	2775	['TGAGYGGATGCCTTCTCCCCAGG']	2993	['TGAGYGGATGCCTTCTCCCCAGG']
NM_002087.3(GRN):c.2T>C (p.Met1Thr)	2896	GRN	2601	['TCCTTGGTACTTTGCAGGCAGACCAYGTGGACCCTGGTGAGCTGGGTGGCC']	2776	['CCAYGTGGACCCTGGTGAGCTGG']	2994	['CCAYGTGGACCCTGGTGAGCTGG']
NM_001083112.2(GPD2):c.1904T>C (p.Phe635Ser)	2820	GPD2	2602	['AGGTATAAGAAGAGATTTCATAAGTYTGATGCAGACCAGAAAGGCTTTATT']	2777	['AAGTYTGATGCAGACCAGAAAGG']	2995	['AAGTYTGATGCAGACCAGAAAGG']
NM_001018077.1(NR3C1):c.1712T>C (p.Val571Ala)	2908	NR3C1	2603	['CTCAACATGTTAGGAGGGCGGCAAGYGATTGCAGCAGTGA']	2778	['AAGYGATTGCAGCAGTGAAATGG']	2996	['AAGYGATTGCAGCAGTGAAATGG']

				AATGGGCAAAG']				
NM_000138.4(FBN1):c.3793T>C (p.Cys1265Arg)	2200	FBN1	2604	['TATCCCTGGAGAGTACAGGTGCTTGYGTTATGATGGATTCATGGCATCTGA']	2779	['CTTGYGTTATGATGGATTCATGG']	2997	['CTTGYGTTATGATGGATTCATGG']
NM_000129.3(F13A1):c.728T>C (p.Met243Thr)	2162	F13A1	2605	['ATCCTGGACACTTGCCTGTATGTGAYGGACAGAGCACAAATGGACCTCTCT']	2780	['TGTGAYGGACAGAGCACAAATGG']	2998	['TGTGAYGGACAGAGCACAAATGG']
NM_031226.2(CYP19A1):c.743+2T>C	-1	-	2606	['ATACAAAAAGTATGAGAAGTCTGTGYAAGTAATACAACTTTGGAAGATTTA']	2781	['CTGTGYAAGTAATACAACTTTGG']	2999	['CTGTGYAAGTAATACAACTTTGG']
NM_000416.2(IFNGR1):c.260T>C (p.Ile87Thr)	3459	IFNGR1	2607	['AATATTTCTCATCATTATTGTAATAYTTCTGATCATGTTGGTGATCCATCA']	2782	['TGTAATAYTTCTGATCATGTTGG']	3000	['TGTAATAYTTCTGATCATGTTGG']
NM_000018.3(ACADVL):c.848T>C (p.Val283Ala)	37	ACADVL	2608	['GTGAAGGAGAAGATCACAGCTTTTGYGGTGGAGAGGGGCTTCGGGGGCATT']	2783 - 2784	['TTTGYGGTGGAGAGGGGCTTCGG', 'TTGYGGTGGAGAGGGGCTTCGGG']	3001 - 3002	['TTTGYGGTGGAGAGGGGCTTCGG', 'TTGYGGTGGAGAGGGGCTTCGGG']
NM_000195.4(HPS1):c.716T>C (p.Leu239Pro)	3257	HPS1	2609	['GCTGGGGTAGAGGTCCTGAACCAGGRGGATGAGGGCAAGCAGGTCGGCCGG']	2785	['CCAGGRGGATGAGGGCAAGCAGG']	3003	['CCAGGRGGATGAGGGCAAGCAGG']
NM_000352.4(ABCC8):c.257T>C (p.Val86Ala)	6833	ABCC8	2610	['ACCTTCATGCTGCTCTTCGTCCTGGBGTGTGAGATTGCAGAGGGCATCCTG']	2786 - 2787	['CCTGGBGTGTGAGATTGCAGAGG', 'CTGGBGTGTGAGATTGCAGAGGG']	3004 - 3005	['CCTGGBGTGTGAGATTGCAGAGG', 'CTGGBGTGTGAGATTGCAGAGGG']
NM_000528.3(MAN2B1):c.2426T>C (p.Leu809Pro)	4125	MAN2B1	2611	['GGCAGCAGCCTGAGAGATGGCTCGCYGGAGCTCATGGTGAGTGGGTCAGAG']	2788	['CGCYGGAGCTCATGGTGAGTGGG']	3006 - 3007	['TCGCYGGAGCTCATGGTGAGTGG', 'CGCYGGAGCTCATGGTGAGTGGG']
NM_002863.4(	5836	PYGL	2612	['CTCCAGTGAC	2789	['AAGAAYATGCCC	3008	['AAGAAYATGCCC

PYGL):c.2461T>C (p.Tyr821His)				CGAACAATTAAAGAAYATGCCCAAAACATCTGGAACGTGGA']		AAAACATCTGG']		AAAACATCTGG']
NM_000495.4(COL4A5):c.4699T>C (p.Cys1567Arg)	1287	COL4A5	2613	['CTTTTCCTTTACCAGATGTGCAGTAYGTGAAGCTCCAGCTGTGGTGATCGC']	2790	['AGTAYGTGAAGCTCCAGCTGTGG']	3009	['AGTAYGTGAAGCTCCAGCTGTGG']
NM_000155.3(GALT):c.460T>C (p.Trp154Arg)	2592	GALT	2614	['TGAGATCCGGGCTGTTGTTGATGCABGGGCCTCAGTCACAGAGGAGCTGGG']	2791	['TGCABGGGCCTCAGTCACAGAGG']	3010	['TGCABGGGCCTCAGTCACAGAGG']
NM_000155.3(GALT):c.482T>C (p.Leu161Pro)	2592	GALT	2615	['GCATGGGCCTCAGTCACAGAGGAGCYGGGTGCCCAGTACCCTTGGGTGCAG']	2792	['AGCYGGGTGCCCAGTACCCTTGG']	3011	['AGCYGGGTGCCCAGTACCCTTGG']
NM_000155.3(GALT):c.499T>C (p.Trp167Arg)	2592	GALT	2616	['AGAGGAGCTGGGTGCCCAGTACCCTYGGGTGCAGGTTTGTGAGGTCGCCCC']	2793	['CCCTYGGGTGCAGGTTTGTGAGG']	3012	['CCCTYGGGTGCAGGTTTGTGAGG']
NM_000155.3(GALT):c.509T>C (p.Ile170Thr)	2592	GALT	2617	['GAGCTCCGTATCCCTATCTGATAGAHCTTTGAAAACAAAGGTGCCATGATG']	2794	['TGATAGAHCTTTGAAAACAAAGG']	3013	['TGATAGAHCTTTGAAAACAAAGG']
NM_004523.3(KIF11):c.2547+2T>C	3832	KIF11	2618	['CAGGAACTTCACAACTTATTGGAGGYAATAACTTTGTAAGTGGAACTTACT']	2795	['GGAGGYAATAACTTTGTAAGTGG']	3014	['GGAGGYAATAACTTTGTAAGTGG']
NM_005211.3(CSF1R):c.2381T>C (p.Ile794Thr)	1436	CSF1R	2619	['TTGACCAATGGTCATGTGGCCAAGAYTGGGGACTTCGGGCTGGCTAGGGAC']	2796	['CAAGAYTGGGGACTTCGGGCTGG']	3015	['CAAGAYTGGGGACTTCGGGCTGG']
NM_005188.3(CBL):c.1150T>C (p.Cys384Arg)	867	CBL	2620	['CTCCACATTCCAACTATGTAAAATABGTGCTGAAAATGATAAGGATGTAAA']	2797	['AATABGTGCTGAAAATGATAAGG']	3016	['AATABGTGCTGAAAATGATAAGG']

NM_006306.3(SMC1A):c.2351T>C (p.Ile784Thr)	8243	SMC1A	2621	['GTGTTTGAAGAGTTTTGTCGGGAGAYTGGTGTGCGCAACATCCGGGAGTTT']	2798	['AGAYTGGTGTGCGCAACATCCGG']	3017	['AGAYTGGTGTGCGCAACATCCGG']
NM_002242.4(KCNJ13):c.722T>C (p.Leu241Pro)	-1	-	2622	['TGGTGTAATGGAGTGATAGTACGTTDGTGGAAAGATGAAGAATGGACATTC']	2799	['GTTDGTGGAAAGATGAAGAATGG']	3018	['GTTDGTGGAAAGATGAAGAATGG']
NM_000199.3(SGSH):c.892T>C (p.Ser298Pro)	6448	SGSH	2623	['CCCCAGCGTTTTGGGTGCTCCGGGGRTGACACCAGTAAGGGTTCAGCAGTG']	2800	['TCCGGGGRTGACACCAGTAAGGG']	3019	['TCCGGGGRTGACACCAGTAAGGG']
NM_020191.2(MRPS22):c.644T>C (p.Leu215Pro)	56945	MRPS22	2624	['CCAATAATTTTCAAGGAAGAAAATCYTAGGGTAAGGTGACTTAGGTTTTAT']	2801	['ATCYTAGGGTAAGGTGACTTAGG']	3020	['ATCYTAGGGTAAGGTGACTTAGG']
NM_017882.2(CLN6):c.200T>C (p.Leu67Pro)	54982	CLN6	2625	['CCCATTCTTCCATTTGCTCCGCAGCYGGTATTCCCTCTCGAGTGGTTTCCA']	2802	['AGCYGGTATTCCCTCTCGAGTGG']	3021	['AGCYGGTATTCCCTCTCGAGTGG']
NM_014874.3(MFN2):c.1392+2T>C	9927	MFN2	2626	['GTAGTCCTCAAGGTTTATAAGAATGWGAGTCATGGAGCAACAGGTCCTCTT']	2803	['AATGWGAGTCATGGAGCAACAGG']	3022	['AATGWGAGTCATGGAGCAACAGG']
NM_024599.5(RHBDF2):c.557T>C (p.Ile186Thr)	79651	RHBDF2	2627	['GCTTACCGCCCCCCTCCCTTCCAGAYTGTGGATCCGCTGGCCCGGGGCCGG']	2804	['AGAYTGTGGATCCGCTGGCCCGG']	3023	['AGAYTGTGGATCCGCTGGCCCGG']
NM_020894.2(UVSSA):c.94T>C (p.Cys32Arg)	57654	UVSSA	2628	['GAAAATGAAGGAACTGAAGAAAATTYGCAAGTATGTCTTAGGGTTCAGTAA']	2805	['AAAATTYGCAAGTATGTCTTAGG']	3024-3025	['AAAATTYGCAAGTATGTCTTAGG', 'AAATTYGCAAGTATGTCTTAGGG']
NM_001161581.1(POC1A):c.398T>C	25886	POC1A	2629	['GCCAGTGATGACAAGACTGTTAAGCYGTGGG	2806	['AGCYGTGGGACAAGAGCAGCCGG']	3026	['AGCYGTGGGACAAGAGCAGCCGG']

(p.Leu133Pro)				ACAAGAGCAG CCGGGAATGT']				
NM_005340.6(HINT1):c.250T >C (p.Cys84Arg)	3094	HINT1	2630	['ACACTTAATG ATTGTTGGCAA GAAAYGTGCT GCTGATCTGGG CCTGAATAA']	2807 - 2808	['CAAGAAAYGTGC TGCTGATCTGG', 'AAGAAAYGTGCTG CTGATCTGGG']	3027 - 3028	['CAAGAAAYGTGC TGCTGATCTGG', 'AAGAAAYGTGCT GCTGATCTGGG']
NM_000495.4(COL4A5):c.43 8+2T>C	1287	COL4 A5	2631	['TTTCCTGGTTT ACAGGGTCCTC CAGYAAGTTAT AAAATTTGGG ATTATGAT']	2809	['TCCAGYAAGTTA TAAAATTTGGG']	3029 - 3030	['CTCCAGYAAGTT ATAAAATTTGG', 'TCCAGYAAGTTA TAAAATTTGGG']
NM_000344.3(SMN1):c.388T >C (p.Tyr130His)	6606	SMN1	2632	['AACCTGTGTT GTGGTTTACAC TGGAYATGGA AATAGAGAGG AGCAAAATCT']	2810	['CACTGGAYATGG AAATAGAGAGG']	3031	['CACTGGAYATGG AAATAGAGAGG']
NM_005334.2(HCFC1):c.- 970T>C	3054	HCFC1	2633	['TTAGTTGTTA CTTCTTCACAC AAGAYGGCGG CTCCCAGGGA GGAGGCATGA']	2811	['CAAGAYGGCGGC TCCCAGGGAGG']	3032	['CAAGAYGGCGG CTCCCAGGGAGG']
NM_000431.3(MVK):c.1039+ 2T>C	4598	MVK	2634	['GTGGCATCAC ACTCCTCAAGC CAGGYATCCC GGGGGTAGGT GGGCCAGGCT']	2812	['CCAGGYATCCCG GGGGTAGGTGG']	3033 - 3034	['CCAGGYATCCCG GGGGTAGGTGG', 'CAGGYATCCCGG GGGTAGGTGGG']
NM_018344.5(SLC29A3):c.60 7T>C (p.Ser203Pro)	5531 5	SLC29 A3	2635	['TATGAGGAAC TCCCAGGCACT GATAYCAGGT GAGAGCCAGG GTCCGGGCAG']	2813	['ACTGATAYCAGG TGAGAGCCAGG']	3035 - 3036	['ACTGATAYCAGG TGAGAGCCAGG', 'CTGATAYCAGGT GAGAGCCAGGG']
NM_000108.4(DLD):c.140T> C (p.Ile47Thr)	1738	DLD	2636	['GTAGTTGATG CTGATGTAACA GTTAYAGGTTC TGGTCCTGGAG GATATGTT']	2814 - 2815	['ACAGTTAYAGGT TCTGGTCCTGG', 'GTTAYAGGTTCTG GTCCTGGAGG']	3037 - 3038	['ACAGTTAYAGGT TCTGGTCCTGG', 'GTTAYAGGTTCTG GTCCTGGAGG']
NM_004333.4(BRAF):c.1403 T>C (p.Phe468Ser)	673	BRAF	2637	['GGACAAAGA ATTGGATCTGG ATCATYTGGAA CAGTCTACAAG GGAAAGTGG']	2816 - 2817	['ATCATYTGGAAC AGTCTACAAGG', 'TCATYTGGAACAG TCTACAAGGG']	3039 - 3040	['ATCATYTGGAAC AGTCTACAAGG', 'TCATYTGGAACA GTCTACAAGGG']
NM_000540.2(RYR1):c.1205T	6261	RYR1	2638	['CAGGAGGAGT CCCAGGCCGCC	2818	['CGCAYGATCCAC AGCACCAATGG']	3041	['CGCAYGATCCAC AGCACCAATGG']

>C (p.Met402Thr)				CGCAYGATCC ACAGCACCAA TGGCCTATAC']				
NM_000256.3(MYBPC3):c.1351+2T>C	4607	MYBPC3	2639	['GTAGCACGGA GCTCTTTGTGA AAGGYGGGCC TGGGACCTGA GGATGTGGGA']	2819	['AAAGGYGGGCCT GGGACCTGAGG']	3042	['AAAGGYGGGCCT GGGACCTGAGG']
NM_000256.3(MYBPC3):c.821+2T>C	4607	MYBPC3	2640	['CCTCCTATCA GCCTTCCGCCG CACGYGAGTG GCCATCCTCAG GGCCTGGGG']	2820	['CACGYGAGTGGC CATCCTCAGGG']	3043 - 3044	['GCACGYGAGTGG CCATCCTCAGG', 'CACGYGAGTGGC CATCCTCAGGG']
NM_000257.3(MYH7):c.2546T>C (p.Met849Thr)	4625	MYH7	2641	['AAGAGTGCAG AAAGAGAGAA GGAGAYGGCC TCCATGAAGG AGGAGTTCAC A']	2821	['GGAGAYGGCCTC CATGAAGGAGG']	3045	['GGAGAYGGCCTC CATGAAGGAGG']
NM_206933.2(USH2A):c.1606T>C (p.Cys536Arg)	7399	USH2A	2642	['CGACACAACA AGCCAGCCAT ATAGAYGCCTC TGCTCCCAGGA GAGCTTCAC']	2822	['ATATAGAYGCCT CTGCTCCCAGG']	3046	['ATATAGAYGCCT CTGCTCCCAGG']
NM_000059.3(BRCA2):c.316+2T>C	675	BRCA2	2643	['TAGATAAATT CAAATTAGACT TAGGYAAGTA ATGCAATATGG TAGACTGGG']	2823	['CTTAGGYAAGTA ATGCAATATGG']	3047	['CTTAGGYAAGTA ATGCAATATGG']
NM_007294.3(BRCA1):c.5291T>C (p.Leu1764Pro)	672	BRCA1	2644	['CTCTTCTTCC AGATCTTCAGG GGGCYAGAAA TCTGTTGCTAT GGGCCCTTC']	2824	['GGCYAGAAATCT GTTGCTATGGG']	3048 - 3049	['GGGCYAGAAATC TGTTGCTATGG', 'GGCYAGAAATCT GTTGCTATGGG']
NM_001130089.1(KARS):c.517T>C (p.Tyr173His)	3735	KARS	2645	['AGCTTCTGGG GGAAAGCTCA TCTTCYATGAT CTTCGAGGAG AGGGGGTGAA']	2825	['TTCYATGATCTTC GAGGAGAGGG']	3050 - 3051	['CTTCYATGATCT TCGAGGAGAGG', 'TTCYATGATCTTC GAGGAGAGGG']
NM_001283009.1(RTEL1):c.3730T>C (p.Cys1244Arg	-1	-	2646	['CGGGCCCCTC TCAGCAGGCTG TGTGYGCCAG GGCTGTGGGG	2826	['CTGTGTGYGCCA GGGCTGTGGGG']	3052	['CTGTGTGYGCCA GGGCTGTGGGG']

)				CAGAGGACGT']				
NM_005554.3(KRT6A):c.1406T>C (p.Leu469Pro)	3853	KRT6A	2647	['GAGATCGCCACCTACCGCAAGCTGCBGGAGGGTGAGGAGTGCAGGTGGGTA']	2827	['TGCBGGAGGGTGAGGAGTGCAGG']	3053	['TGCBGGAGGGTGAGGAGTGCAGG']
NM_000218.2(KCNQ1):c.550T>C (p.Tyr184His)	3784	KCNQ1	2648	['CTGGTCCGCCGGCTGCCGCAGCAAGBACGTGGGCCTCTGGGGGCGGCTGCG']	2828	['AGCAAGBACGTGGGCCTCTGGGG']	3054 - 3056	['CAGCAAGBACGTGGGCCTCTGGG', 'AGCAAGBACGTGGGCCTCTGGGG', 'GCAAGBACGTGGGCCTCTGGGGG']
NM_198056.2(SCN5A):c.5624T>C (p.Met1875Thr)	6331	SCN5A	2649	['GAGATGGACGCCCTGAAGATCCAGAHGGAGGAGAAGTTCATGGCAGCCAAC']	2829	['CCAGAHGGAGGAGAAGTTCATGG']	3057	['CCAGAHGGAGGAGAAGTTCATGG']
NM_006920.4(SCN1A):c.269T>C (p.Phe90Ser)	6323	SCN1A	2650	['TGTTGTGTTCCTGTCTTACAGACTTYTATAGTATTGAATAAAGGGGAAGGCC']	2830 - 2831	['ACTTYTATAGTATTGAATAAAGG', 'CTTYTATAGTATTGAATAAAGGG']	3058 - 3059	['ACTTYTATAGTATTGAATAAAGG', 'CTTYTATAGTATTGAATAAAGGG']
NM_006920.4(SCN1A):c.272T>C (p.Ile91Thr)	6323	SCN1A	2651	['TGTGTTCCTGTCTTACAGACTTTAYAGTATTGAATAAAGGGAAGGCCATC']	2832 - 2833	['ACTTTTAYAGTATTGAATAAAGG', 'CTTTTAYAGTATTGAATAAAGGG']	3060 - 3061	['ACTTTTAYAGTATTGAATAAAGG', 'CTTTTAYAGTATTGAATAAAGGG']
NM_006514.3(SCN10A):c.1661T>C (p.Leu554Pro)	6336	SCN10A	2652	['GGAGTCAGGGTTGCTGGGTTGAGGARGAGGGCTTCTAGGGAGGGGGCCTTG']	2834 - 2836	['GAGGARGAGGGCTTCTAGGGAGG', 'AGGARGAGGGCTTCTAGGGAGGG', 'GGARGAGGGCTTCTAGGGAGGG']	3062 - 3064	['GAGGARGAGGGCTTCTAGGGAGG', 'AGGARGAGGGCTTCTAGGGAGGG', 'GGARGAGGGCTTCTAGGGAGGG']
NM_000251.2(MSH2):c.2005+2T>C	4436	MSH2	2653	['AACAGATGTTCCACATCATTACTGGYAAAAAACCTGGTTTTTGGGCTTTGT']	2837 - 2838	['CTGGYAAAAAACCTGGTTTTTGG', 'TGGYAAAAAACCTGGTTTTTGGG']	3065 - 3066	['CTGGYAAAAAACCTGGTTTTTGG', 'TGGYAAAAAACCTGGTTTTTGGG']
NM_000251.2(MSH2):c.595T>C (p.Cys199Arg)	4436	MSH2	2654	['CCTCATCCAGATTGGACCAAAGGAAYGTGTTTTACCCGGAGGAGAGACTGC']	2839	['AAGGAAYGTGTTTTACCCGGAGG']	3067	['AAGGAAYGTGTTTTACCCGGAGG']
NM_00100574	2629	GBA	2655	['TTCACCGCTC	2840	['GCCRAGTGGGTG	3068	['GAGCCRAGTGGG

1.2(GBA):c.667T>C (p.Trp223Arg)				CATTGGTCTTGAGCCRAGTGGGTGATGTCCAGGGGCTGGCA']		ATGTCCAGGGG']	-3070	TGATGTCCAGG', 'AGCCRAGTGGGTGATGTCCAGGG', 'GCCRAGTGGGTGATGTCCAGGGG']
NM_003494.3(DYSF):c.1284+2T>C	8291	DYSF	2656	['GAGGTCAGCTTTGCGGGGAAAATGGYAAGGAGCAAGGGAGCAGGAGGGTTC']	2841	['ATGGYAAGGAGCAAGGGAGCAGG']	3071	['ATGGYAAGGAGCAAGGGAGCAGG']
NM_012463.3(ATP6V0A2):c.825+2T>C	23545	ATP6V0A2	2657	['ACCCGCATCCAGGATCTCTACACTGYGAGTAAGCTGGAAGTGGATTGCCTC']	2842	['CACTGYGAGTAAGCTGGAAGTGG']	3072	['CACTGYGAGTAAGCTGGAAGTGG']
NM_016725.2(FOLR1):c.493+2T>C	2348	FOLR1	2658	['ACAAGGGCTGGAACTGGACTTCAGGYGAGGGCTGGGGTGGGCAGGAATGGA']	2843	['AGGYGAGGGCTGGGGTGGGCAGG']	3073-3074	['CTTCAGGYGAGGGCTGGGGTGGG', 'AGGYGAGGGCTGGGGTGGGCAGG']
NM_003764.3(STX11):c.173T>C (p.Leu58Pro)	8676	STX11	2659	['GACATTCAGGATGAAAACCAGCTGCYGGTGGCCGACGTGAAGCGGCTGGGA']	2844	['TGCYGGTGGCCGACGTGAAGCGG']	3075	['TGCYGGTGGCCGACGTGAAGCGG']
NM_014714.3(IFT140):c.4078T>C (p.Cys1360Arg)	9742	IFT140	2660	['GGACCCCAAGGAGTCCATCAAGCAGYGTGAGCTGCTCCTGGAGGAACCAGA']	2845	['GCAGYGTGAGCTGCTCCTGGAGG']	3076-3077	['CAAGCAGYGTGAGCTGCTCCTGG', 'GCAGYGTGAGCTGCTCCTGGAGG']
NM_000531.5(OTC):c.1005+2T>C	5009	OTC	2661	['GAAAACAGAAAGTGGACAATCATGGYAAGCAAGAAACAAGGAATGGAGGAT']	2846	['ATCATGGYAAGCAAGAAACAAGG']	3078	['ATCATGGYAAGCAAGAAACAAGG']
NM_000531.5(OTC):c.158T>C (p.Ile53Thr)	5009	OTC	2662	['CTAAAAAACTTTACCGGAGAAGAAABTAAATATATGCTATGGCTATCAGCA']	2847	['AAGAAABTAAATATATGCTATGG']	3079	['AAGAAABTAAATATATGCTATGG']
NM_000531.5(	5009	OTC	2663	['GAGAAAAGA	2848	['ACAAGATYGTCT	3080	['ACAAGATYGTCT

OTC):c.284T>C (p.Leu95Ser)				AGTACTCGAACAAGATYGTCTACAGAAACAGGTAAGTCCACT']		ACAGAAACAGG']		ACAGAAACAGG']
NM_000531.5(OTC):c.2T>C (p.Met1Thr)	5009	OTC	2664	['CGTCCTTTACACAATTAAAAGAAGAYGCTGTTTAATCTGAGGATCCTGTTA']	2849	['AGAAGAYGCTGTTTAATCTGAGG']	3081	['AGAAGAYGCTGTTTAATCTGAGG']
NM_000531.5(OTC):c.526T>C (p.Tyr176His)	5009	OTC	2665	['CCATCCTATCCAGATCCTGGCTGATYACCTCACGCTCCAGGTTGGTTTATT']	2850	['GGCTGATYACCTCACGCTCCAGG']	3082 - 3083	['GGCTGATYACCTCACGCTCCAGG', 'GATYACCTCACGCTCCAGGTTGG']
NM_000531.5(OTC):c.779T>C (p.Leu260Ser)	5009	OTC	2666	['GAAGCAGCGCATGGAGGCAATGTATYAATTACAGACACTTGGATAAGCATG']	2851	['ATGTATYAATTACAGACACTTGG']	3084	['ATGTATYAATTACAGACACTTGG']
NM_000322.4(PRPH2):c.736T>C (p.Trp246Arg)	5961	PRPH2	2667	['CCACCAGACGGAGGAGCTCAACCTGYGGGTGCGTGGCTGCAGGGCTGCCCT']	2852 - 2853	['ACCTGYGGGTGCGTGGCTGCAGG', 'CCTGYGGGTGCGTGGCTGCAGGG']	3085 - 3086	['ACCTGYGGGTGCGTGGCTGCAGG', 'CCTGYGGGTGCGTGGCTGCAGGG']
NM_000211.4(ITGB2):c.1877+2T>C	3689	ITGB2	2668	['CCCCTCACCCTGTGGCAAGTACATGYGAGTGCAGGCGGAGCAGGCAGGGCG']	2854	['CATGYGAGTGCAGGCGGAGCAGG']	3087	['CATGYGAGTGCAGGCGGAGCAGG']
NM_015474.3(SAMHD1):c.1106T>C (p.Leu369Ser)	25939	SAMHD1	2669	['TTTGTGTTGATAAGCTCTACGGTGTRAAGAGTTGCGAGTGTGGAACATGTC']	2855	['GGTGTRAAGAGTTGCGAGTGTGG']	3088	['GGTGTRAAGAGTTGCGAGTGTGG']
NM_001101.3(ACTB):c.356T>C (p.Met119Thr)	60	ACTB	2670	['AACCCCAAGGCCAACCGCGAGAAGAYGACCCAGGTGAGTGGCCCGCTACCT']	2856	['GAGAAGAYGACCAGGTGAGTGG']	3089	['GAGAAGAYGACCCAGGTGAGTGG']
NM_015713.4(RRM2B):c.368T>C	50484	RRM2B	2671	['CTCGATGAGAATTTGAAAGCCATAGRAACAG	2857	['CCATAGRAACAGCGAGCCTCTGG']	3090	['CCATAGRAACAGCGAGCCTCTGG']

(p.Phe123Ser)				CGAGCCTCTGG AACCTGCAC']				
NM_015599.2(PGM3):c.248T >C (p.Leu83Ser)	5238	PGM3	2672	['TTGGTTGATC CTTTGGGTGAA ATGTYGGCACC ATCCTGGGAG GAACATGCC']	2858	['AATGTYGGCACC ATCCTGGGAGG']	3091	['AATGTYGGCACC ATCCTGGGAGG']
NM_002136.2(HNRNPA1):c.8 17T>C (p.Phe273Leu)	3178	HNRN PA1	2673	['GAATTACAAC AATCAGTCTTC AAATBTTGGAC CCATGAAGGG AGGAAATTT']	2859 - 2861	['TTCAAATBTTGGA CCCATGAAGG', 'TCAAATBTTGGAC CCATGAAGGG', 'AATBTTGGACCCA TGAAGGGAGG']	3092 - 3094	['TTCAAATBTTGG ACCCATGAAGG', 'TCAAATBTTGGAC CCATGAAGGG', 'AATBTTGGACCC ATGAAGGGAGG']
NM_002136.2(HNRNPA1):c.8 41T>C (p.Phe281Leu)	3178	HNRN PA1	2674	['TTTTGGACCC ATGAAGGGAG GAAATYTTGG AGGCAGAAGC TCTGGCCCCTA']	2862	['AATYTTGGAGGC AGAAGCTCTGG']	3095	['AATYTTGGAGGC AGAAGCTCTGG']
NM_022552.4(DNMT3A):c.27 05T>C (p.Phe902Ser)	1788	DNMT 3A	2675	['CGCAAAATAC TCCTTCAGCGG AGCGRAGAGG TGGCGGATGA CTGGCACGCT']	2863	['GCGRAGAGGTGG CGGATGACTGG']	3096	['GCGRAGAGGTGG CGGATGACTGG']
NM_000076.2(CDKN1C):c.*5 +2T>C	1028	CDKN 1C	2676	['GCGCAAGAG GCTGCGGTGA GCCAAGYGAG TACAGCGCACC TGGGGGGGCG C']	2864 - 2866	['CCAAGYGAGTAC AGCGCACCTGG', 'CAAGYGAGTACA GCGCACCTGGG', 'AAGYGAGTACAG CGCACCTGGGG']	3097 - 3099	['CCAAGYGAGTAC AGCGCACCTGG', 'CAAGYGAGTACA GCGCACCTGGG', 'AAGYGAGTACAG CGCACCTGGGG']
NC_012920.1: m.9478T>C	4514	MT-CO3	2677	['ATAATCCTAT TTATTACCTCA GAAGYTTTTTT CTTCGCAGGAT TTTTCTGA']	2867	['TCAGAAGYTTTTT TCTTCGCAGG']	3100	['TCAGAAGYTTTT TTCTTCGCAGG']
NM_002049.3(GATA1):c.2T> C (p.Met1Thr)	2623	GATA 1	2678	['CGCAGGTTAA TCCCCAGAGGC TCCAYGGAGTT CCCTGGCCTGG GGTCCCTG']	2868 - 2869	['TCCAYGGAGTTC CCTGGCCTGGG', 'CCAYGGAGTTCCC TGGCCTGGGG']	3101 - 3103	['CTCCAYGGAGTT CCCTGGCCTGG', 'TCCAYGGAGTTC CCTGGCCTGGG', 'CCAYGGAGTTCC CTGGCCTGGGG']
NM_005740.2(DNAL4):c.153 +2T>C	1012 6	DNAL 4	2679	['GAGAAATTCT CCAACAACAA CGAGGYATTG	2870	['CGAGGYATTGCC AGCAGTGCAGG']	3104	['CGAGGYATTGCC AGCAGTGCAGG']

				CCAGCAGTGC AGGCGGCCCCT ']				
NM_00128722 3.1(SCN11A):c .1142T>C (p.Ile381Thr)	1128 0	SCN11 A	2680	['GGGCTCTACT CAGTCTTCTTC TTCAYTGTGGT CATTTTCCTGG GCTCCTTC']	2871	['TTCAYTGTGGTCA TTTTCCTGGG']	3105 - 3106	['CTTCAYTGTGGT CATTTTCCTGG', 'TTCAYTGTGGTCA TTTTCCTGGG']
NM_00130294 6.1(TRNT1):c.4 97T>C (p.Leu166Ser)	5109 5	TRNT1	2681	['TAATGAATAG GTTTTGATGGC ACTTYATTTGA CTACTTTAATG GTTATGAA']	2872	['ACTTYATTTGACT ACTTTAATGG']	3107	['ACTTYATTTGAC TACTTTAATGG']
NM_178151.2(DCX):c.2T>C (p.Met1Thr)	1641	DCX	2682	['AGGTCTCTGA GGTTCCACCAA AATAYGGAAC TTGATTTTGGA CACTTTGAC']	2873	['CAAAATAYGGAA CTTGATTTTGG']	3108	['CAAAATAYGGA ACTTGATTTTGG']
NM_000169.2(GLA):c.758T> C (p.Ile253Thr)	-1	-	2683	['TGGACATCTT TTAACCAGGA GAGAAYTGTT GATGTTGCTGG ACCAGGGGGT']	2874	['GAGAGAAYTGTT GATGTTGCTGG']	3109	['GAGAGAAYTGTT GATGTTGCTGG']
NM_170707.3(LMNA):c.710T >C (p.Phe237Ser)	4000	LMNA	2684	['ATTGACAATG GGAAGCAGCG TGAGTYTGAG AGCCGGCTGG CGGATGCGCTG ']	2875	['TGAGTYTGAGAG CCGGCTGGCGG']	3110	['TGAGTYTGAGAG CCGGCTGGCGG']
NM_000256.3(MYBPC3):c.33 30+2T>C	4607	MYBP C3	2685	['CAGAAAGCCG ACAAGAAGAC CATGGBGAGC CCAGGGTCTGG GGTCCCCACG']	2876 - 2878	['ACCATGGBGAGC CCAGGGTCTGG', 'CCATGGBGAGCCC AGGGTCTGGG', 'CATGGBGAGCCCA GGGTCTGGGG']	3111 - 3113	['ACCATGGBGAGC CCAGGGTCTGG', 'CCATGGBGAGCC CAGGGTCTGGG', 'CATGGBGAGCCC AGGGTCTGGGG']
NM_005957.4(MTHFR):c.153 0+2T>C	4524	MTHF R	2686	['AGCGGGGGCT ATGTCTTCCAG AAGGYGTGGT AGGGAGGCAC GGGGTGCCCC']	2879 - 2881	['GAAGGYGTGGTA GGGAGGCACGG', 'AAGGYGTGGTAG GGAGGCACGGG', 'AGGYGTGGTAGG GAGGCACGGGG']	3114 - 3116	['GAAGGYGTGGTA GGGAGGCACGG', 'AAGGYGTGGTAG GGAGGCACGGG', 'AGGYGTGGTAGG GAGGCACGGGG']
NM_000264.3(PTCH1):c.3168 +2T>C	5727	PTCH1	2687	['AACCCCTGGA CGGCCGGGAT CATTGYGAGTG	2882 - 2884	['ATCATTGYGAGT GTATTATAAGG', 'TCATTGYGAGTGT	3117 - 3119	['ATCATTGYGAGT GTATTATAAGG', 'TCATTGYGAGTGT

				TATTATAAGGG GCTTTGTGG']		ATTATAAGGG', 'CATTGYGAGTGTA TTATAAGGGG']		ATTATAAGGG', 'CATTGYGAGTGT ATTATAAGGGG']
NM_000030.2(AGXT):c.322T>C (p.Trp108Arg)	189	AGXT	2688	['CTTCCTGGTT GGGGCCAATG GCATTYGGGG GCAGCGAGCC GTGGACATCG G']	2885	['CATTYGGGGGCA GCGAGCCGTGG']	3120	['CATTYGGGGGCA GCGAGCCGTGG']
NM_000023.2(SGCA):c.371T>C (p.Ile124Thr)	6442	SGCA	2689	['ACTCGGCAGA GGCTGGTGCTG GAGAYTGGGG ACCCAGAAGG TACCTCTAGC']	2886	['CTGGAGAYTGGG GACCCAGAAGG']	3121	['CTGGAGAYTGGG GACCCAGAAGG']
NM_001103.3(ACTN2):c.683T>C (p.Met228Thr)	88	ACTN 2	2690	['GAGAAGCACC TGGATATTCCT AAAAYGTTGG ATGCTGAAGGT GAGATGAAA']	2887	['CCTAAAAYGTTG GATGCTGAAGG']	3122	['CCTAAAAYGTTG GATGCTGAAGG']
NM_001165963.1(SCN1A):c.4055T>C (p.Leu1352Pro)	-1	-	2691	['ATTCCATCCA TCATGAATGTG CTTCYGGTTTG TCTTATATTCT GGCTAATT']	2888	['TTCYGGTTTGTCT TATATTCTGG']	3123	['TTCYGGTTTGTC TTATATTCTGG']
NM_001165963.1(SCN1A):c.1265T>C (p.Val422Ala)	6323	SCN1A	2692	['CTAATAAATT TGATCCTGGCT GTGGHGGCCA TGGCCTACGAG GAACAGAAT']	2889	['TGTGGHGGCCAT GGCCTACGAGG']	3124	['TGTGGHGGCCAT GGCCTACGAGG']
NM_000426.3(LAMA2):c.8282T>C (p.Ile2761Thr)	3908	LAMA 2	2693	['GCAGAATCAG AACCAGCTCTT TTGAYAGGGA GCAAGCAGTTC GGGCTTTCA']	2890 - 2891	['TTGAYAGGGAGC AAGCAGTTCGG', 'TGAYAGGGAGCA AGCAGTTCGGG']	3125 - 3126	['TTGAYAGGGAGC AAGCAGTTCGG', 'TGAYAGGGAGCA AGCAGTTCGGG']
NM_000257.3(MYH7):c.5117T>C (p.Leu1706Pro)	-1	-	2694	['TCCCGGAAGC TGGCGGAGCA GGAGCYGATT GAGACTAGTG AGCGGGTGCA G']	2892	['AGCYGATTGAGA CTAGTGAGCGG']	3127	['AGCYGATTGAGA CTAGTGAGCGG']
NM_001399.4(EDA):c.396+2T>C	1896	EDA	2695	['TCTGACTCCC AGGACGGGCA CCAGGKGAGT CACCTAGTAGG	2893 - 2894	['ACCAGGKGAGTC ACCTAGTAGGG', 'CCAGGKGAGTCAC CTAGTAGGGG']	3128 - 3130	['CACCAGGKGAGT CACCTAGTAGG', 'ACCAGGKGAGTC ACCTAGTAGGG',

				GGCGGCGGCG']				'CCAGGKGAGTCA CCTAGTAGGGG']
NM_001848.2(COL6A1):c.957+2T>C	1291	COL6A1	2696	['TCCAGGGGAC CCAAGGGCTA CAAGGYGAGC GTGGGCTGCTG GGAGGGGGGA']	2895 - 2896	['ACAAGGYGAGCG TGGGCTGCTGG', 'CAAGGYGAGCGT GGGCTGCTGGG']	3131 - 3132	['ACAAGGYGAGC GTGGGCTGCTGG', 'CAAGGYGAGCGT GGGCTGCTGGG']
NM_000238.3(KCNH2):c.1945+6T>C	3757	KCNH2	2697	['CTGCGTCATG CTCATTGGCTG TGAGYGTGCCC AGGGGCGGGC GGCGGGGAG']	2897 - 2898	['CTGTGAGYGTGC CCAGGGGCGGG', 'TGAGYGTGCCCAG GGGCGGGCGG']	3133 - 3134	['CTGTGAGYGTGC CCAGGGGCGGG', 'TGAGYGTGCCCA GGGGCGGGCGG']
NM_021007.2(SCN2A):c.1271T>C (p.Val424Ala)	6326	SCN2A	2698	['CTAATAAATT TGATCTTGGCT GTGGYGGCCA TGGCCTATGAG GAACAGAAT']	2899	['TGTGGYGGCCAT GGCCTATGAGG']	3135	['TGTGGYGGCCAT GGCCTATGAGG']
NM_021007.2(SCN2A):c.4308+2T>C	6326	SCN2A	2699	['TATGCAGCTG TTGATTCACGA AATGYAAGTCT AGTTAGAGGG AAATTGTTT']	2900 - 2901	['CGAAATGYAAGT CTAGTTAGAGG', 'GAAATGYAAGTCT AGTTAGAGGG']	3136 - 3137	['CGAAATGYAAGT CTAGTTAGAGG', 'GAAATGYAAGTC TAGTTAGAGGG']
NM_000083.2(CLCN1):c.1283T>C (p.Phe428Ser)	1180	CLCN1	2700	['CCCCGCGAAG CCATCAGTACT TTGTYTGACAA CAATACATGG GTGAAACAC']	2902 - 2903	['CTTTGTYTGACAA CAATACATGG', 'TTTGTYTGACAAC AATACATGGG']	3138 - 3139	['CTTTGTYTGACA ACAATACATGG', 'TTTGTYTGACAAC AATACATGGG']
NM_004550.4(NDUFS2):c.875T>C (p.Met292Thr)	4720	NDUFS2	2701	['CATTATGCTC TCCACAGTGGA GTGAYGCTTCG GGGCTCAGGC ATCCAGTGG']	2904	['GGAGTGAYGCTT CGGGGCTCAGG']	3140	['GGAGTGAYGCTT CGGGGCTCAGG']
NM_000546.5(TP53):c.584T>C (p.Ile195Thr)	7157	TP53	2702	['CACACGCAAA TTTCCTTCCAC TCGGRTAAGAT GCTGAGGAGG GGCCAGACC']	2905 - 2906	['CTCGGRTAAGAT GCTGAGGAGGG', 'TCGGRTAAGATGC TGAGGAGGGG']	3141 - 3143	['ACTCGGRTAAGA TGCTGAGGAGG', 'CTCGGRTAAGAT GCTGAGGAGGG', 'TCGGRTAAGATG CTGAGGAGGGG']

Table 3. A to G with NGG PAM. Table 2 shows a list of A to G mutations that may be corrected using any of the base editors provided herein. GRNAs and gRNAall indicate the protospacer and PAM sequence, where the PAM sequence is the last 3 nucleotides of each of the sequences in GRNAs and gRNAall.

Name	Gene ID	Gene Symbol	SEQ ID NO:	Flanks	SEQ ID NO:	gRNAs	SEQ ID NO:	gRNAall
NM_017547.3(FOXRED1):c.1289A>G (p.Asn430Ser)	55572	FOXRED1	5084	['GTGGGCCCCCACCCGCTAGTTGTCAVCATGTACTTTGCTACTGGCTTCAGT']	5261	['CCACCCGCTAGTTGTCAVCATGT']	5464-5466	['CCCACCCGCTAGTTGTCAVCATG', 'CCACCCGCTAGTTGTCAVCATGT', 'CCCGCTAGTTGTCAVCATGTACT']
NM_000071.2(CBS):c.1150A>G (p.Lys384Glu)	875	CBS	5085	['GGTGACTCCCCCATCCCGCAGGACCRAGTTCCTGAGCGACAGGTGGATGCT']	5262	['CCCCATCCCGCAGGACCRAGTTC']	5467-5470	['CCCCCATCCCGCAGGACCRAGTT', 'CCCCATCCCGCAGGACCRAGTTC', 'CCCATCCCGCAGGACCRAGTTCC', 'CCATCCCGCAGGACCRAGTTCCT']
NM_000552.3(VWF):c.2384A>G (p.Tyr795Cys)	7450	VWF	5086	['GAGTGTACCAAAACGTGCCAGAACTRTGACCTGGAGTGCATGAGCATGGGC']	5263	['CCAAAACGTGCCAGAACTRTGAC']	5471	['CCAAAACGTGCCAGAACTRTGAC']
NM_000552.3(VWF):c.1583A>G (p.Asn528Ser)	7450	VWF	5087	['ACCTGCGGCCTGTGTGGGAATTACARTGGCAACCAGGGCGACGACTTCCTT']	5264	['CCTGTGTGGGAATTACARTGGCA']	5472	['CCTGTGTGGGAATTACARTGGCA']
NM_000308.2(CTSA):c.1238A>G (p.Tyr413Cys)	5476	CTSA	5088	['CTTTAGAAATACCAGATCCTATTATRTAATGGAGATGTAGACATGGCCTGC']	5265	['CCAGATCCTATTATRTAATGGAG']	5473	['CCAGATCCTATTATRTAATGGAG']
NM_000277.1(PAH):c.916A>G (p.Ile306Val)	5053	PAH	5089	['TTCTATTTTCCCCCAATTACAGGAARTTGGCCTTGCCTCTCTGGGTGCACC']	5266	['CCCCCAATTACAGGAARTTGGCC']	5474-5476	['CCCCCAATTACAGGAARTTGGCC', 'CCCCAATTACAGGAARTTGGCCT', 'CCCAATTACAGGAARTTGGCCTT']
NM_000512.4(GALNS):c.1460A>G (p.Asn487Ser)	2588	GALNS	5090	['TTGGTCCCCGCGCAGCCCCAGCTCARCGTGTGCAACTGGGCGGTCATGGTA']	5267	['CCGCGCAGCCCCAGCTCARCGTG']	5477	['CCGCGCAGCCCCAGCTCARCGTG']
NM_013319.2(UBIAD1):c.305A>G (p.Asn102Ser)	29914	UBIAD1	5091	['GTGCACGGGGCCGGTAATTTGGTCARCACTTACTATGACTTTTCCAAGGGC']	5268	['CCGGTAATTTGGTCARCACTTAC']	5478	['CCGGTAATTTGGTCARCACTTAC']
NM_013319.2(U	2991	UBIAD	5092	['AGCACCGAGGCCA	5269	['CCATTCTCCATT	5479	['CCATTCTCCATT

BIAD1):c.695A>G (p.Asn232Ser)	4	1		TTCTCCATTCCARCAACACCAGGGACATGGAGTCCGAC']		CCARCAACACC']		CCARCAACACC']
NM_000275.2(OCA2):c.1465A>G (p.Asn489Asp)	4948	OCA2	5093	['TGCCACTGCCATCGGGGACCCTCCARATGTCATTATTGTTTCCAACCAAGA']	5270	['CCATCGGGGACCCTCCARATGTC']	5480	['CCATCGGGGACCCTCCARATGTC']
NM_001127255.1(NLRP7):c.2738A>G (p.Asn913Ser)	-1	-	5094	['CTCACAAACCTGGACTTGAGTATCARCCAGATAGCTCGTGGATTGTGGATT']	5271	['CCTGGACTTGAGTATCARCCAGA']	5481	['CCTGGACTTGAGTATCARCCAGA']
NM_152783.4(D2HGDH):c.1315A>G (p.Asn439Asp)	728294	D2HGDH	5095	['TGCCCTTGTCCCTCCAGGAGATGGTRACCTGCACCTCAATGTGACGGCGGA']	5272	['CCTCCAGGAGATGGTRACCTGCA']	5482-5483	['CCCTCCAGGAGATGGTRACCTGC', 'CCTCCAGGAGATGGTRACCTGCA']
NM_022132.4(MCCC2):c.1309A>G (p.Ile437Val)	640887	MCCC2	5096	['TGTGGCCTGTGCCCAAGTGCCTAAGDTAACCCTCATCATTGGGGGCTCCTA']	5273	['CCCAAGTGCCTAAGDTAACCCTC']	5484	['CCCAAGTGCCTAAGDTAACCCTC']
NM_000022.2(ADA):c.219-2A>G	100	ADA	5097	['TTCCCAACCCCTTTCTTCCCTTCCCRGGGGCTGCCGGGAGGCTATCAAAAG']	5274	['CCCCTTTCTTCCCTTCCCRGGGG']	5485-5487	['CCCCTTTCTTCCCTTCCCRGGGG', 'CCCTTTCTTCCCTTCCCRGGGGC', 'CCTTTCTTCCCTTCCCRGGGGCT']
NM_017780.3(CHD7):c.3082A>G (p.Ile1028Val)	556376	CHD7	5098	['TTTAGTAATTGCCCCATTGTCCACARTCCCCAACTGGGAAAGGGAATTCCG']	5275	['CCCCATTGTCCACARTCCCCAAC']	5488	['CCCCATTGTCCACARTCCCCAAC']
NM_000483.4(APOC2):c.1A>G (p.Met1Val)	-1	-	5099	['TCAATGTTCCAGGTCTCTGGACACTRTGGGCACACGACTCCTCCCAGCTCT']	5276	['CCAGGTCTCTGGACACTRTGGGC']	5489	['CCAGGTCTCTGGACACTRTGGGC']
NM_000391.3(TPP1):c.887-10A>G	1200	TPP1	5100	['TGTCCCTCATGCCGGCCTGGATTTTYTTTTTTTTTTTTTTGAGGGATGGG']	5277	['CCGGCCTGGATTTTYTTTTTTTT']	5490	['CCGGCCTGGATTTTYTTTTTTTT']
NM_017890.4(VPS13B):c.8978A>G (p.Asn2993Ser)	157680	VPS13B	5101	['CTTCTGCCCTGGGCCCTGCTTATCARTGAATCCAAATGGGACCTCTGGCTA']	5278	['CCTGGGCCCTGCTTATCARTGAA']	5491	['CCTGGGCCCTGCTTATCARTGAA']
NM_000226.3(KRT9):c.482A>G (p.Asn161Ser)	3857	KRT9	5102	['GAGAAGAGCACCATGCAGGAACTCADTTCTCGGCTGGCCTCT	5279	['CCATGCAGGAACTCADTTCTCGG']	5492	['CCATGCAGGAACTCADTTCTCGG']

				TACTTGGAT']				
NM_000529.2(MC2R):c.761A>G (p.Tyr254Cys)	4158	MC2R	5103	['CCAAGTAACCCCTACTGCGCCTGCTRCATGTCTCTCTTCCAGGTGAACGGC']	5280-5281	['CCCTACTGCGCCTGCTRCATGTC', 'CCTACTGCGCCTGCTRCATGTCT']	5493-5495	['CCCCTACTGCGCCTGCTRCATGT', 'CCCTACTGCGCCTGCTRCATGTC', 'CCTACTGCGCCTGCTRCATGTCT']
NM_005957.4(MTHFR):c.971A>G (p.Asn324Ser)	4524	MTHFR	5104	['CCAGGCCTCCACTTCTACACCCTCARCCGCGAGATGGCTACCACAGAGGTG']	5282	['CCACTTCTACACCCTCARCCGCG']	5496	['CCACTTCTACACCCTCARCCGCG']
NM_000403.3(GALE):c.101A>G (p.Asn34Ser)	2582	GALE	5105	['GGCTACTTGCCTGTGGTCATCGATARCTTCCATAATGCCTTCCGTGGTGAG']	5283	['CCTGTGGTCATCGATARCTTCCA']	5497	['CCTGTGGTCATCGATARCTTCCA']
NM_000356.3(TCOF1):c.149A>G (p.Tyr50Cys)	6949	TCOF1	5106	['CAGCCCGTAACCCTTCTGGACATCTRTACACACTGGCAACAGTAAGTGGTG']	5284-5285	['CCCTTCTGGACATCTRTACACAC', 'CCTTCTGGACATCTRTACACACT']	5498-5499	['CCCTTCTGGACATCTRTACACAC', 'CCTTCTGGACATCTRTACACACT']
NM_012464.4(TLL1):c.1885A>G (p.Ile629Val)	7092	TLL1	5107	['ACTTCTTACCAAACTTAACGGCACCRTAACCACCCCTGGCTGGCCCAAGGA']	5286	['CCAAACTTAACGGCACCRTAACC']	5500	['CCAAACTTAACGGCACCRTAACC']
NM_000112.3(SLC26A2):c.1273A>G (p.Asn425Asp)	1836	SLC26A2	5108	['GGAAATGTATGCCATTGGCTTTTGTRATATCATCCCTTCCTTCTTCCACTG']	5287	['CCATTGGCTTTTGTRATATCATC']	5501	['CCATTGGCTTTTGTRATATCATC']
NM_000157.3(GBA):c.680A>G (p.Asn227Ser)	2629	GBA	5109	['ACATCACCCACTTGGCTCAAGACCARTGGAGCGGTGAATGGGAAGGGGTCA']	5288	['CCACTTGGCTCAAGACCARTGGA']	5502	['CCACTTGGCTCAAGACCARTGGA']
NM_175073.2(APTX):c.602A>G (p.His201Arg)	54840	APTX	5110	['GATAAATACCCAAAGGCCCGTTACCRTTGGCTGGTCTTACCGTGGACCTCC']	5289-5290	['CCCAAAGGCCCGTTACCRTTGGC', 'CCAAAGGCCCGTTACCRTTGGCT']	5503-5504	['CCCAAAGGCCCGTTACCRTTGGC', 'CCAAAGGCCCGTTACCRTTGGCT']
NM_020638.2(FGF23):c.211A>G (p.Ser71Gly)	8074	FGF23	5111	['TGGCGCACCCCATCAGACCATCTACRGTGAGTAGGGCTTCAGGCTGGGAAG']	5291	['CCCCATCAGACCATCTACRGTGA']	5505-5507	['CCCCATCAGACCATCTACRGTGA', 'CCCATCAGACCATCTACRGTGAG', 'CCATCAGACCATCTACRGTGAGT']
NM_021102.3(SPINT2):c.488A>G (p.Tyr163Cys)	10653	SPINT2	5112	['AGGAACTCCTGCAATAACTTCATCTRTGGAGGCTGCCGGGGC	5292	['CCTGCAATAACTTCATCTRTGGA']	5508	['CCTGCAATAACTTCATCTRTGGA']

				AATAAGAAC']				
NM_004795.3(KL):c.578A>G (p.His193Arg)	9365	KL	5113	['GTGCAGCCCGTGGTCACCCTGTACCRCTGGGACCTGCCCCAGCGCCTGCAG']	5293	['CCGTGGTCACCCTGTACCRCTGG']	5509	['CCGTGGTCACCCTGTACCRCTGG']
NM_012193.3(FZD4):c.766A>G (p.Ile256Val)	-1	-	5114	['GTTTTCCTACCCTGAGCGCCCCATCRTATTTCTCAGTATGTGCTATAATAT']	5294 - 5295	['CCCTGAGCGCCCCATCRTATTTC', 'CCTGAGCGCCCCATCRTATTTCT']	5510-5511	['CCCTGAGCGCCCCATCRTATTTC', 'CCTGAGCGCCCCATCRTATTTCT']
NM_001099274.1(TINF2):c.838A>G (p.Lys280Glu)	26277	TINF2	5115	['ATGGGCCTCCACTAGGGGAGGCCATDAGGAGCGCCCCACAGTCATGCTGTT']	5296	['CCACTAGGGGAGGCCATDAGGAG']	5512	['CCACTAGGGGAGGCCATDAGGAG']
NM_005682.6(ADGRG1):c.263A>G (p.Tyr88Cys)	9289	ADGRG1	5116	['TCCTTCCCTGACCCCAGGGGCCTCTRCCACTTCTGCCTCTACTGGAACCGA']	5297	['CCCCAGGGGCCTCTRCCACTTCT']	5513	['CCCCAGGGGCCTCTRCCACTTCT']
NM_000369.2(TSHR):c.1856A>G (p.Asp619Gly)	7253	TSHR	5117	['CCGCAGTACAACCCAGGGGACAAAGRTACCAAAATTGCCAAGAGGATGGCT']	5298	['CCCAGGGGACAAAGRTACCAAAA']	5514	['CCCAGGGGACAAAGRTACCAAAA']
NM_024009.2(GJB3):c.497A>G (p.Asn166Ser)	2707	GJB3	5118	['ATGCCGCGCCTGGTGCAGTGTGCCADCGTGGCCCCCTGCCCCAACATCGTG']	5299	['CCTGGTGCAGTGTGCCADCGTGG']	5515	['CCTGGTGCAGTGTGCCADCGTGG']
NM_003722.4(TP63):c.697A>G (p.Lys233Glu)	8626	TP63	5119	['TATCCGCGCCATGCCTGTCTACAAARAAGCTGAGCACGTCACGGAGGTGGT']	5300	['CCATGCCTGTCTACAAARAAGCT']	5516	['CCATGCCTGTCTACAAARAAGCT']
NM_003494.3(DYSF):c.3443-33A>G	8291	DYSF	5120	['CAGCTCTTAACCACTCCAGCCACTCRCTCTGGCACCTCTGTTTTTCCCTT']	5301	['CCACTCCAGCCACTCRCTCTGGC']	5517	['CCACTCCAGCCACTCRCTCTGGC']
NM_003494.3(DYSF):c.1285-2A>G	8291	DYSF	5121	['AACTTGTCCCCTCCCTGTGTCTTCTRGCTGTGCAGCAAGATCTTGGAGAAG']	5302	['CCCCTCCCTGTGTCTTCTRGCTG']	5518-5520	['CCCCTCCCTGTGTCTTCTRGCTG', 'CCCTCCCTGTGTCTTCTRGCTGT', 'CCTCCCTGTGTCTTCTRGCTGTG']
NM_002408.3(MGAT2):c.785A>G (p.His262Arg)	4247	MGAT2	5122	['CTTATACTTTTCCTAGAAGAGGATCRCTACTTAGCCCCAGACTTTTACCAT']	5303	['CCTAGAAGAGGATCRCTACTTAG']	5521	['CCTAGAAGAGGATCRCTACTTAG']
NM_000492.3(CF	1080	CFTR	5123	['GTGATTATCACCA	5304	['CCAGCACCAGTT	5522	['CCAGCACCAGTT

TR):c.2738A>G (p.Tyr913Cys)				GCACCAGTTCGTRT TATGTGTTTTACATT TACGTGGGA']		CGTRTTATGTG']		CGTRTTATGTG']
NM_001814.4(CTSC):c.857A>G (p.Gln286Arg)	1075	CTSC	5124	['TCTCAGACCCCAAT CCTAAGCCCTCRGG AGGTTGTGTCTTGTA GCCAGTAT']	5305	['CCCCAATCCTAA GCCCTCRGGAG']	5523-5525	['CCCCAATCCTAA GCCCTCRGGAG', 'CCCAATCCTAAG CCCTCRGGAGG', 'CCAATCCTAAGC CCTCRGGAGGT']
NM_005144.4(HR):c.-218A>G	5580 6	HR	5125	['TCCGACCCCTCCAA CCTGCGGCCCTRGA GCGCCCCCGCCGCC CCGGGGGAA']	5306	['CCTCCAACCTGC GGCCCTRGAGC']	5526-5527	['CCTCCAACCTGC GGCCCTRGAGC', 'CCAACCTGCGGC CCTRGAGCGCC']
NM_018488.2(TBX4):c.1592A>G (p.Gln531Arg)	9496	TBX4	5126	['TCCTTGTCCCGAGA ATCTTCCTTACRGTA CCATTCAGGAATGG GGACTGTG']	5307	['CCCGAGAATCTT CCTTACRGTAC']	5528-5529	['CCCGAGAATCTT CCTTACRGTAC', 'CCGAGAATCTTCC TTACRGTACC']
NM_001089.2(ABCA3):c.1702A>G (p.Asn568Asp)	21	ABCA3	5127	['ACAGATCACCGTC CTGCTGGGCCACRA CGGTGCCGGGAAGA CCACCACCCT']	5308	['CCGTCCTGCTGG GCCACRACGGT']	5530	['CCGTCCTGCTGG GCCACRACGGT']
NM_000525.3(KCNJ11):c.776A>G (p.His259Arg)	3767	KCNJ11	5128	['CTGGTGGCCCCGCT GATCATCTACCRTG TCATTGATGCCAAC AGCCCACTC']	5309 - 5310	['CCCCGCTGATCA TCTACCRTGTC', 'CCCGCTGATCATC TACCRTGTCA']	5531-5533	['CCCCGCTGATCA TCTACCRTGTC', 'CCCGCTGATCATC TACCRTGTCA', 'CCGCTGATCATCT ACCRTGTCAT']
NM_005587.2(MEF2A):c.788A>G (p.Asn263Ser)	4205	MEF2A	5129	['TCTCCCCCTCCACC AGGTGGTGGTARTC TTGGAATGAACAGT AGGAAACCA']	5311	['CCACCAGGTGGT GGTARTCTTGG']	5534	['CCACCAGGTGGT GGTARTCTTGG']
NM_000098.2(CPT2):c.359A>G (p.Tyr120Cys)	1376	CPT2	5130	['TTTTTAGGACCCTG GTTTGATATGTRCCT ATCTGCTCGAGACT CCGTTGTT']	5312	['CCTGGTTTGATA TGTRCCTATCT']	5535-5536	['CCCTGGTTTGAT ATGTRCCTATC', 'CCTGGTTTGATAT GTRCCTATCT']
NM_178138.4(LHX3):c.332A>G (p.Tyr111Cys)	8022	LHX3	5131	['GTGCGCCGCGCCC AGGACTTCGTGTRC CACCTGCACTGCTTT GCCTGCGTC']	5313 - 5314	['CCCAGGACTTCG TGTRCCACCTG', 'CCAGGACTTCGT GTRCCACCTGC']	5537-5538	['CCCAGGACTTCG TGTRCCACCTG', 'CCAGGACTTCGT GTRCCACCTGC']
NM_005502.3(ABCA1):c.2804A>G (p.Asn935Ser)	19	ABCA1	5132	['CAGATCACCTCCTT CCTGGGCCACARTG GAGCGGGGAAGAC GACCACCATG']	5315	['CCTCCTTCCTGG GCCACARTGGA']	5539-5540	['CCTCCTTCCTGG GCCACARTGGA', 'CCTTCCTGGGCCA CARTGGAGCG']
m.3260A>G	4567	MT-	5133	['GATGGCAGAGCCC	5316	['CCCGGTAATCGC	5541-	['CCCGGTAATCGC

		TL1		GGTAATCGCATARA ACTTAAAACTTTAC AGTCAGAGGT']	- 5317	ATARAACTTAA', 'CCGGTAATCGCA TARAACTTAAA']	5542	ATARAACTTAA', 'CCGGTAATCGCA TARAACTTAAA']
m.4269A>G	4565	MT-TI	5134	['GCATTCCCCCTCAA ACCTAAGAAATRTG TCTGATAAAAGAGT TACTTTGAT']	5318 - 5319	['CCCTCAAACCTA AGAAATRTGTC', 'CCTCAAACCTAA GAAATRTGTCT']	5543-5544	['CCCTCAAACCTA AGAAATRTGTC', 'CCTCAAACCTAA GAAATRTGTCT']
m.14495A>G	4541	MT-ND6	5135	['TCCAAAGACAACC ATCATTCCCCCTRA ATAAATTAAAAAAA CTATTAAACC']	5320	['CCATCATTCCCC CTRAATAAATT']	5545	['CCATCATTCCCC CTRAATAAATT']
NM_002764.3(PR PS1):c.341A>G (p.Asn114Ser)	5631	PRPS1	5136	['CCAATCTCAGCCA AGCTTGTTGCAART ATGCTATCTGTAGC AGGTGCAGAT']	5321	['CCAAGCTTGTTG CAARTATGCTA']	5546	['CCAAGCTTGTTG CAARTATGCTA']
NM_000054.4(A VPR2):c.614A>G (p.Tyr205Cys)	554	AVPR2	5137	['GCGGAGCCCTGGG GCCGTCGCACCTRT GTCACCTGGATTGC CCTGATGGTG']	5322	['CCTGGGGCCGTC GCACCTRTGTC']	5547	['CCTGGGGCCGTC GCACCTRTGTC']
NM_000033.3(A BCD1):c.443A>G (p.Asn148Ser)	215	ABCD1	5138	['ATCGCCCTCCCTGC TACCTTCGTCARCA GTGCCATCCGTTAC CTGGAGGGC']	5323 - 5324	['CCCTGCTACCTT CGTCARCAGTG', 'CCTGCTACCTTCG TCARCAGTGC']	5548-5549	['CCCTGCTACCTT CGTCARCAGTG', 'CCTGCTACCTTCG TCARCAGTGC']
NM_000061.2(BT K):c.1082A>G (p.Tyr361Cys)	695	BTK	5139	['AGCACCATCCCTG AGCTCATTAACTRC CATCAGCACAACTC TGCAGGTGAG']	5325 - 5326	['CCCTGAGCTCAT TAACTRCCATC', 'CCTGAGCTCATTA ACTRCCATCA']	5550-5551	['CCCTGAGCTCAT TAACTRCCATC', 'CCTGAGCTCATTA ACTRCCATCA']
NM_003413.3(ZI C3):c.1213A>G (p.Lys405Glu)	7547	ZIC3	5140	['CTACACGCACCCG AGCTCCCTGCGCRA ACACATGAAGGTAA TTACCTCTTT']	5327	['CCGAGCTCCCTG CGCRAACACAT']	5552-5553	['CCCGAGCTCCCT GCGCRAACACA', 'CCGAGCTCCCTGC GCRAACACAT']
NM_005448.2(B MP15):c.704A>G (p.Tyr235Cys)	9210	BMP15	5141	['TTGGACATTGCCTT CTTGTTACTCTRTTT CAATGATACTCATA AAAGCATT']	5328	['CCTTCTTGTTACT CTRTTTCAAT']	5554	['CCTTCTTGTTACT CTRTTTCAAT']
NM_001363.4(D KC1):c.1069A>G (p.Thr357Ala)	1736	DKC1	5142	['ATTAATGACCACA GCGGTCATCTCTRC CTGCGACCATGGTA TAGTAGCCAA']	5329	['CCACAGCGGTCA TCTCTRCCTGC']	5555	['CCACAGCGGTCA TCTCTRCCTGC']
NM_000481.3(A MT):c.125A>G (p.His42Arg)	275	AMT	5143	['CGCAGGACACCGC TCTATGACTTCCRCC TGGCCCACGGCGGG AAAATGGTG']	5330	['CCGCTCTATGAC TTCCRCCTGGC']	5556	['CCGCTCTATGAC TTCCRCCTGGC']
NM_003361.3(U	7369	UMOD	5144	['TGCCACGCCCTGG	5331	['CCTGGCCACATG	5557-	['CCCTGGCCACAT

MOD):c.383A>G (p.Asn128Ser)				CCACATGTGTCART GTGGTGGGCAGCTA CTTGTGCGTA']		TGTCARTGTGG']	5558	GTGTCARTGTG', 'CCTGGCCACATGT GTCARTGTGG']
NM_001382.3(DP AGT1):c.509A>G (p.Tyr170Cys)	1798	DPAG T1	5145	['TCTCTCCCCGCAGG AATCCTGTACTRTGT CTACATGGGGCTGC TGGCAGTG']	5332	['CCGCAGGAATCC TGTACTRTGTC']	5559	['CCGCAGGAATCC TGTACTRTGTC']
NM_001128177.1 (THRB):c.1324A >G (p.Met442Val)	7068	THRB	5146	['CTGCCATGCCAGC CGCTTCCTGCACRT GAAGGTGGAATGCC CCACAGAACT']	5333	['CCAGCCGCTTCC TGCACRTGAAG']	5560	['CCAGCCGCTTCC TGCACRTGAAG']
NM_000141.4(FG FR2):c.874A>G (p.Lys292Glu)	2263	FGFR2	5147	['TGCCCAGCCCCAC ATCCAGTGGATCRA GCACGTGGAAAAGA ACGGCAGTAA']	5334	['CCCACATCCAGT GGATCRAGCAC']	5561-5563	['CCCCACATCCAG TGGATCRAGCA', 'CCCACATCCAGT GGATCRAGCAC', 'CCACATCCAGTG GATCRAGCACG']
NM_000371.3(TT R):c.401A>G (p.Tyr134Cys)	7276	TTR	5148	['ACCATTGCCGCCCT GCTGAGCCCCTRCT CCTATTCCACCACG GCTGTCGTC']	5335 - 5337	['CCGCCCTGCTGA GCCCCTRCTCC', 'CCCTGCTGAGCCC CTRCTCCTAT', 'CCTGCTGAGCCCC TRCTCCTATT']	5564-5566	['CCGCCCTGCTGA GCCCCTRCTCC', 'CCCTGCTGAGCCC CTRCTCCTAT', 'CCTGCTGAGCCCC TRCTCCTATT']
NM_000371.3(TT R):c.379A>G (p.Ile127Val)	7276	TTR	5149	['CGACTCCGGCCCC CGCCGCTACACCRT TGCCGCCCTGCTGA GCCCCTACTC']	5338	['CCCCCGCCGCTA CACCRTTGCCG']	5567-5569	['CCCCCGCCGCTA CACCRTTGCCG', 'CCCCGCCGCTAC ACCRTTGCCGC', 'CCCGCCGCTACA CCRTTGCCGCC']
NM_000174.4(GP 9):c.182A>G (p.Asn61Ser)	2815	GP9	5150	['ACCCGCCACCTTCT GCTGGCCAACARCA GCCTTCAGTCCGTG CCCCCGGGA']	5339	['CCTTCTGCTGGC CAACARCAGCC']	5570	['CCTTCTGCTGGC CAACARCAGCC']
NM_000222.2(KI T):c.1924A>G (p.Lys642Glu)	3815	KIT	5151	['ACGGGAAGCCCTC ATGTCTGAACTCRA AGTCCTGAGTTACC TTGGTAATCA']	5340 - 5341	['CCCTCATGTCTG AACTCRAAGTC', 'CCTCATGTCTGAA CTCRAAGTCC']	5571-5572	['CCCTCATGTCTG AACTCRAAGTC', 'CCTCATGTCTGAA CTCRAAGTCC']
NM_000530.6(M PZ):c.242A>G (p.His81Arg)	4359	MPZ	5152	['TCCCCTCATTCCTC ATAGATCTTCCRCT ATGCCAAGGGACAA CCCTACATT']	5342	['CCTCATAGATCT TCCRCTATGCC']	5573	['CCTCATAGATCT TCCRCTATGCC']
NM_000233.3(L HCGR):c.1733A> G (p.Asp578Gly)	-1	-	5153	['AAAATGGCAATCC TCATCTTCACCGRTT TCACCTGCATGGCA	5343	['CCTCATCTTCAC CGRTTTCACCT']	5574	['CCTCATCTTCAC CGRTTTCACCT']

				CCTATCTCT']				
NM_000421.3(KRT10):c.1374-2A>G	-1	-	5154	['CCGCCGCGTCCGCCGCCTCCGGAACYAAACGGGGTGAGGTCACATTCGGTT']	5344	['CCGCCGCCTCCGGAACYAAACGG']	5575	['CCGCCGCCTCCGGAACYAAACGG']
NM_000422.2(KRT17):c.274A>G (p.Asn92Asp)	3872	KRT17	5155	['TGAGAAGGCCACCATGCAGAACCTCVATGACCGCCTGGCCTCCTACCTGGA']	5345	['CCACCATGCAGAACCTCVATGAC']	5576-5577	['CCACCATGCAGAACCTCVATGAC', 'CCATGCAGAACCTCVATGACCGC']
NM_000422.2(KRT17):c.275A>G (p.Asn92Ser)	3872	KRT17	5156	['GAGAAGGCCACCATGCAGAACCTCARTGACCGCCTGGCCTCCTACCTGGAC']	5346	['CCACCATGCAGAACCTCARTGAC']	5578-5579	['CCACCATGCAGAACCTCARTGAC', 'CCATGCAGAACCTCARTGACCGC']
NM_000823.3(GHRHR):c.985A>G (p.Lys329Glu)	2692	GHRHR	5157	['TTGTCTTTCCTGCAGGCGTCTCTCCRAGTCGACACTTTTCCTGATCCCACT']	5347	['CCTGCAGGCGTCTCTCCRAGTCG']	5580	['CCTGCAGGCGTCTCTCCRAGTCG']
NM_000407.4(GP1BB):c.338A>G (p.Tyr113Cys)	-1	-	5158	['GCCGGCCGCCCCGAGCGTGCGCCCTDCCGCGACCTGCGTTGCGTGGCGCCC']	5348-5349	['CCCCGAGCGTGCGCCCTDCCGCG', 'CCCGAGCGTGCGCCCTDCCGCGA']	5581-5583	['CCCCGAGCGTGCGCCCTDCCGCG', 'CCCGAGCGTGCGCCCTDCCGCGA', 'CCGAGCGTGCGCCCTDCCGCGAC']
NM_001146040.1(GLRA1):c.920A>G (p.Tyr307Cys)	2741	GLRA1	5159	['CCTCCACCCCCACTCTAGGTGTCCTVTGTGAAAGCCATTGACATTTGGATG']	5350-5351	['CCCCACTCTAGGTGTCCTVTGTG', 'CCCACTCTAGGTGTCCTVTGTGA']	5584-5586	['CCCCACTCTAGGTGTCCTVTGTG', 'CCCACTCTAGGTGTCCTVTGTGA', 'CCACTCTAGGTGTCCTVTGTGAA']
NM_182925.4(FLT4):c.3104A>G (p.His1035Arg)	2324	FLT4	5160	['CGCCTCCCCGCACCCCAGTGCATCCRCAGAGACCTGGCTGCTCGGAACATT']	5352	['CCGCACCCCAGTGCATCCRCAGA']	5587	['CCGCACCCCAGTGCATCCRCAGA']
NM_212482.1(FN1):c.2918A>G (p.Tyr973Cys)	2335	FN1	5161	['ACCGGGCTGTCCCCTGGGGTCACCTRTTACTTCAAAGTCTTTGCAGTGAGC']	5353-5354	['CCCCTGGGGTCACCTRTTACTTC', 'CCCTGGGGTCACCTRTTACTTCA']	5588-5589	['CCCCTGGGGTCACCTRTTACTTC', 'CCCTGGGGTCACCTRTTACTTCA']
NM_000121.3(EPOR):c.1460A>G (p.Asn487Ser)	2057	EPOR	5162	['GGCTTATCCGATGGCCCCTACTCCARCCCTTATGAGAACAGCCTTATCCCA']	5355	['CCGATGGCCCCTACTCCARCCCT']	5590	['CCGATGGCCCCTACTCCARCCCT']
NM_001735.2(C5):c.1115A>G (p.Lys372Arg)	727	C5	5163	['CGTCTACCCCCTCACCCAATCTACCYTGATGGGATATGGAAT	5356-5357	['CCCCTCACCCAATCTACCYTGAT', 'CCCTCACCCAATC	5591-5593	['CCCCTCACCCAATCTACCYTGAT', 'CCCTCACCCAATC

				CCCAGGCTT']		TACCYTGATG']		TACCYTGATG', 'CCTCACCCAATCTACCYTGATGG']
NM_001844.4(COL2A1):c.4172A>G (p.Tyr1391Cys)	1280	COL2A1	5164	['ACGGAAGGCTCCCAGAACATCACCTRCCACTGCAAGAACAGCATTGCCTAT']	5358-5359	['CCCAGAACATCACCTRCCACTGC', 'CCAGAACATCACCTRCCACTGCA']	5594-5595	['CCCAGAACATCACCTRCCACTGC', 'CCAGAACATCACCTRCCACTGCA']
NM_001904.3(CTNNB1):c.121A>G (p.Thr41Ala)	1499	CTNNB1	5165	['CTCTGGAATCCATTCTGGTGCCACTNCCACAGCTCCTTCTCTGAGTGGTAA']	5360	['CCATTCTGGTGCCACTNCCACAG']	5596	['CCATTCTGGTGCCACTNCCACAG']
NM_000040.1(APOC3):c.280A>G (p.Thr94Ala)	345	APOC3	5166	['GGATTTGGACCCTGAGGTCAGACCARCTTCAGCCGTGGCTGCCTGAGACCT']	5361-5362	['CCCTGAGGTCAGACCARCTTCAG', 'CCTGAGGTCAGACCARCTTCAGC']	5597-5598	['CCCTGAGGTCAGACCARCTTCAG', 'CCTGAGGTCAGACCARCTTCAGC']
NM_000488.3(SERPINC1):c.655A>G (p.Asn219Asp)	462	SERPINC1	5167	['TGCAGAGCAATCCAGAGCGGCCATCRACAAATGGGTGTCCAATAAGACCGA']	5363	['CCAGAGCGGCCATCRACAAATGG']	5599	['CCAGAGCGGCCATCRACAAATGG']
NM_001085.4(SERPINA3):c.1240A>G (p.Met414Val)	12	SERPINA3	5168	['TACAGACACCCAGAACATCTTCTTCRTGAGCAAAGTCACCAATCCCAAGCA']	5364	['CCCAGAACATCTTCTTCRTGAGC']	5600-5601	['CCCAGAACATCTTCTTCRTGAGC', 'CCAGAACATCTTCTTCRTGAGCA']
NM_001145.4(ANG):c.121A>G (p.Lys41Glu)	-1	-	5169	['CTTCCTGACCCAGCACTATGATGCCRAACCACAGGGCCGGGATGACAGATA']	5365	['CCAGCACTATGATGCCRAACCAC']	5602-5603	['CCCAGCACTATGATGCCRAACCA', 'CCAGCACTATGATGCCRAACCAC']
NM_001100.3(ACTA1):c.350A>G (p.Asn117Ser)	58	ACTA1	5170	['GAGGCCCCCCTCAATCCCAAGGCCARCCGCGAGAAGATGACCCAGATCATG']	5366	['CCTCAATCCCAAGGCCARCCGCG']	5604-5605	['CCCTCAATCCCAAGGCCARCCGC', 'CCTCAATCCCAAGGCCARCCGCG']
NM_014053.3(FLVCR1):c.361A>G (p.Asn121Asp)	28982	FLVCR1	5171	['GATCTTCAGCCTGTACTCGCTGGTCRACGCCTTTCAGTGGATCCAGTACAG']	5367	['CCTGTACTCGCTGGTCRACGCCT']	5606	['CCTGTACTCGCTGGTCRACGCCT']
NM_000334.4(SCN4A):c.4078A>G (p.Met1360Val)	6329	SCN4A	5172	['GAAGCAGGCCTTCGACATCACCATCRTGATCCTCATCTGCCTCAACATGGT']	5368	['CCTTCGACATCACCATCRTGATC']	5607	['CCTTCGACATCACCATCRTGATC']
NM_004519.3(KCNQ3):c.1403A>G (p.Asn468Ser)	3786	KCNQ3	5173	['GAACCAAAGCCTGTTGGCTTAAACARTAAAGAGCGTTTCCGCACGGCCTTC']	5369	['CCTGTTGGCTTAAACARTAAAGA']	5608	['CCTGTTGGCTTAAACARTAAAGA']
NM_007375.3(T	2343	TARD	5174	['AATGCCGAACCTA	5370	['CCTAAGCACAAT	5609	['CCTAAGCACAAT

ARDBP):c.800A>G (p.Asn267Ser)	5	BP		AGCACAATAGCARTAGACAGTTAGAAAGAAGTGGAAGA']		AGCARTAGACA']		AGCARTAGACA']
NM_032520.4(GNPTG):c.610-2A>G	84572	GNPTG	5175	['TGCTGCCCCTGCATCCTCCACCTTCRGGGCCATGAGAAGTTGCTGAGGACA']	5371	['CCTGCATCCTCCACCTTCRGGGC']	5610	['CCTGCATCCTCCACCTTCRGGGC']
NM_000495.4(COL4A5):c.466-2A>G	1287	COL4A5	5176	['AGAACTTCCATTGATGGCTTCTTTTRGGGTGAACCAGGTAGTATAATTATG']	5372	['CCATTGATGGCTTCTTTTRGGGT']	5611	['CCATTGATGGCTTCTTTTRGGGT']
NM_000495.4(COL4A5):c.1340-2A>G	1287	COL4A5	5177	['TTGCTATCCTTTCTTTATCTTACTCRGGTGATGAGATATGTGAACCAGGCC']	5373	['CCTTTCTTTATCTTACTCRGGTG']	5612	['CCTTTCTTTATCTTACTCRGGTG']
NM_000060.3(BTD):c.278A>G (p.Tyr93Cys)	686	BTD	5178	['CTCATGAACCAGAACCTTGACATCTRTGAACAGCAAGTGATGACTGCAGCC']	5374	['CCAGAACCTTGACATCTRTGAAC']	5613	['CCAGAACCTTGACATCTRTGAAC']
NM_000060.3(BTD):c.641A>G (p.Asn214Ser)	686	BTD	5179	['CTTGTTGACCGCTACCGTAAACACARCCTCTACTTTGAGGCAGCATTCGAT']	5375	['CCGCTACCGTAAACACARCCTCT']	5614	['CCGCTACCGTAAACACARCCTCT']
NM_000094.3(COL7A1):c.425A>G (p.Lys142Arg)	1294	COL7A1	5180	['CAGCTGGCCCGACCTGGTGTCCCCARGGTGATCCCTACCCCTACCATGCCT']	5376	['CCCGACCTGGTGTCCCCARGGTG']	5615-5616	['CCCGACCTGGTGTCCCCARGGTG', 'CCGACCTGGTGTCCCCARGGTGA']
NM_005247.2(FGF3):c.146A>G (p.Tyr49Cys)	2248	FGF3	5181	['GGGGCGCCCCGGCGCCGCAAGCTCTRCTGCGCCACGAAGTACCACCTCCAG']	5377-5378	['CCCGGCGCCGCAAGCTCTRCTGC', 'CCGGCGCCGCAAGCTCTRCTGCG']	5617-5618	['CCCGGCGCCGCAAGCTCTRCTGC', 'CCGGCGCCGCAAGCTCTRCTGCG']
NM_000313.3(PROS1):c.701A>G (p.Tyr234Cys)	5627	PROS1	5182	['TGTGAATGCCCCGAAGGCTACAGATRTAATCTCAAATCAAAGTCTTGTGAA']	5379-5380	['CCCGAAGGCTACAGATRTAATCT', 'CCGAAGGCTACAGATRTAATCTC']	5619-5621	['CCCCGAAGGCTACAGATRTAATC', 'CCCGAAGGCTACAGATRTAATCT', 'CCGAAGGCTACAGATRTAATCTC']
NM_004612.3(TGFBR1):c.134A>G (p.Asn45Ser)	7046	TGFBR1	5183	['TTCTGCCACCTCTGTACAAAAGACARTTTTACTTGTGTGACAGATGGGCTC']	5381	['CCTCTGTACAAAAGACARTTTTA']	5622	['CCTCTGTACAAAAGACARTTTTA']
m.608A>G	4558	MT-TF	5184	['GTAGCTTACCTCCTCAAAGCAATACRCTGAAAATGTTTAGAC	5382	['CCTCCTCAAAGCAATACRCTGAA']	5623-5624	['CCTCCTCAAAGCAATACRCTGAA', 'CCTCAAAGCAAT

				GGGCTCACA']				ACRCTGAAAAT']
NM_001376.4(DYNC1H1):c.2909A>G (p.Tyr970Cys)	1778	DYNC1H1	5185	['CTAAGAATAACCAATCAGGTAATCTRCTTGAATCCACCAATTGAAGAGTGC']	5383	['CCAATCAGGTAATCTRCTTGAAT']	5625	['CCAATCAGGTAATCTRCTTGAAT']
NM_000459.4(TEK):c.2690A>G (p.Tyr897Cys)	7010	TEK	5186	['ATGCTCTCTTCCTTCCCTCCAGGCTVCTTGTACCTGGCCATTGAGTACGCG']	5384	['CCTTCCCTCCAGGCTVCTTGTAC']	5626	['CCTTCCCTCCAGGCTVCTTGTAC']
NM_014191.3(SCN8A):c.5302A>G (p.Asn1768Asp)	6334	SCN8A	5187	['CATGTACATTGCCATCATCCTGGAGRACTTCAGTGTAGCCACAGAGGAAAG']	5385	['CCATCATCCTGGAGRACTTCAGT']	5627	['CCATCATCCTGGAGRACTTCAGT']
NM_002552.4(ORC4):c.521A>G (p.Tyr174Cys)	5000	ORC4	5188	['CATCATAAAAACCAAACACTTCTCTRTAATCTTTTTGACATTTCTCAGTCT']	5386	['CCAAACACTTCTCTRTAATCTTT']	5628	['CCAAACACTTCTCTRTAATCTTT']
NM_004813.2(PEX16):c.992A>G (p.Tyr331Cys)	9409	PEX16	5189	['TACTTGCCCACCTGGCAGAAAATCTRCTTCTACAGTTGGGGCTGACAGACC']	5387 - 5388	['CCACCTGGCAGAAAATCTRCTTC', 'CCTGGCAGAAAATCTRCTTCTAC']	5629-5630	['CCACCTGGCAGAAAATCTRCTTC', 'CCTGGCAGAAAATCTRCTTCTAC']
NM_016952.4(CDON):c.2368A>G (p.Thr790Ala)	50937	CDON	5190	['GTTTTTGTTTTCCCTCAAAGGTTCARCATACAAATTTAGGGTCATTGCCAT']	5389	['CCCTCAAAGGTTCARCATACAAA']	5631	['CCCTCAAAGGTTCARCATACAAA']
NM_016464.4(TMEM138):c.287A>G (p.His96Arg)	51524	TMEM138	5191	['TACTTTGCCCTCAGCATCTCCCTTCRTGTCTGGGTCATGGTAAGAGTGGCA']	5390 - 5391	['CCCTCAGCATCTCCCTTCRTGTC', 'CCTCAGCATCTCCCTTCRTGTCT']	5632-5633	['CCCTCAGCATCTCCCTTCRTGTC', 'CCTCAGCATCTCCCTTCRTGTCT']
NM_005022.3(PFN1):c.350A>G (p.Glu117Gly)	5216	PFN1	5192	['GTTGATCAAACCACCGTGGACACCTYCTTTGCCCATCAGCAGGACTAGCGC']	5392	['CCACCGTGGACACCTYCTTTGCC']	5634	['CCACCGTGGACACCTYCTTTGCC']
NM_022787.3(NMNAT1):c.817A>G (p.Asn273Asp)	64802	NMNAT1	5193	['GGTCATCCTGGCCCCTTTGCAGAGARACACTGCAGAAGCTAAGACATAGGA']	5393	['CCCCTTTGCAGAGARACACTGCA']	5635	['CCCCTTTGCAGAGARACACTGCA']
NM_005340.6(HINT1):c.152A>G (p.His51Arg)	3094	HINT1	5194	['GACATTTCCCCTCAAGCACCAACACRTTTTCTGGTGATACCCAAGAAACAT']	5394 - 5396	['CCCCTCAAGCACCAACACRTTTT', 'CCCTCAAGCACCAACACRTTTTC', 'CCTCAAGCACCAACACRTTTTCT']	5636-5638	['CCCCTCAAGCACCAACACRTTTT', 'CCCTCAAGCACCAACACRTTTTC', 'CCTCAAGCACCAACACRTTTTCT']
NM_005211.3(CS	1436	CSF1R	5195	['GACTAACCCTGCA	5397	['CCTGCAGTGCTT	5639	['CCTGCAGTGCTT

F1R):c.2320-2A>G				GTGCTTTCCCTCRGTGCATCCACCGGGACGTGGCAGCG']		TCCCTCRGTGC']		TCCCTCRGTGC']
NM_001039958.1(MESP2):c.271A>G (p.Lys91Glu)	145873	MESP2	5196	['GCGGCAGAGCGCCAGCGAGCGGGAGRAACTGCGCATGCGCACGCTGGCCCG']	5398	['CCAGCGAGCGGGAGRAACTGCGC']	5640	['CCAGCGAGCGGGAGRAACTGCGC']
NM_001099274.1(TINF2):c.850A>G (p.Thr284Ala)	26277	TINF2	5197	['TAGGGGAGGCCATAAGGAGCGCCCCRCAGTCATGCTGTTTCCCTTTAGGAA']	5399	['CCATAAGGAGCGCCCCRCAGTCA']	5641	['CCATAAGGAGCGCCCCRCAGTCA']
NM_003863.3(DPM2):c.68A>G (p.Tyr23Cys)	8818	DPM2	5198	['GCCGTTAGCCTGATCATCTTCACCTRCTACACCGCCTGGGTGATTCTCTTG']	5400	['CCTGATCATCTTCACCTRCTACA']	5642	['CCTGATCATCTTCACCTRCTACA']
NM_000530.6(MPZ):c.347A>G (p.Asn116Ser)	4359	MPZ	5199	['AAGGATGGCTCCATTGTCATACACARCCTAGACTACAGTGACAATGGCACG']	5401	['CCATTGTCATACACARCCTAGAC']	5643	['CCATTGTCATACACARCCTAGAC']
NM_000138.4(FBN1):c.3058A>G (p.Thr1020Ala)	2200	FBN1	5200	['ACCCGGATTTGCCACAAAAGAAATTRCAAATGGAAAGCCTTTCTTCAAAGG']	5402	['CCACAAAAGAAATTRCAAATGGA']	5644	['CCACAAAAGAAATTRCAAATGGA']
NM_000169.2(GLA):c.1153A>G (p.Thr385Ala)	-1	-	5201	['GGCCTGTAATCCTGCCTGCTTCATCRCACAGCTCCTCCCTGTGAAAAGGAA']	5403	['CCTGCCTGCTTCATCRCACAGCT']	5645	['CCTGCCTGCTTCATCRCACAGCT']
NM_000257.3(MYH7):c.2206A>G (p.Ile736Val)	4625	MYH7	5202	['AGCGGCCATCCCTGAGGGACAGTTCRTTGATAGCAGGAAGGGGGCAGAGAA']	5404	['CCCTGAGGGACAGTTCRTTGATA']	5646-5647	['CCCTGAGGGACAGTTCRTTGATA', 'CCTGAGGGACAGTTCRTTGATAG']
NM_018972.2(GDAP1):c.368A>G (p.His123Arg)	54332	GDAP1	5203	['AGCATGTATTACCCACGGGTACAACRTTACCGAGAGCTGCTTGACTCCTTG']	5405	['CCCACGGGTACAACRTTACCGAG']	5648	['CCCACGGGTACAACRTTACCGAG']
NM_001946.3(DUSP6):c.566A>G (p.Asn189Ser)	1848	DUSP6	5204	['ACTACCATCCGAGTCTGTTGCACTAYTGGGGTCTCGGTCAAGGTCAGACTC']	5406	['CCGAGTCTGTTGCACTAYTGGGG']	5649	['CCGAGTCTGTTGCACTAYTGGGG']
NM_003867.3(FGF17):c.560A>G (p.Asn187Ser)	8822	FGF17	5205	['TACCAAGGCCAGCTGCCCTTCCCCARCCACGCCGAGAAGCAGAAGCAGTTC']	5407	['CCAGCTGCCCTTCCCCARCCACG']	5650	['CCAGCTGCCCTTCCCCARCCACG']
NM_015560.2(OP	4976	OPA1	5206	['TTTTTATTTTTCCT	5408	['CCTGAGTAGACC	5651	['CCTGAGTAGACC

A1):c.1146A>G (p.Ile382Met)				GAGTAGACCATRTC CTTAAATGTAAAAG GCCCTGGAC']		ATRTCCTTAAA']		ATRTCCTTAAA']
NM_002972.3(SBF1):c.1249A>G (p.Met417Val)	6305	SBF1	5207	['AAGGCCATGCCCT CCAGCACCTTCAYC AGGAAATCGTCCTC TACCAGCCCA']	5409	['CCCTCCAGCACC TTCAYCAGGAA']	5652-5653	['CCCTCCAGCACC TTCAYCAGGAA', 'CCTCCAGCACCTT CAYCAGGAAA']
NM_006876.2(B4GAT1):c.1168A>G (p.Asn390Asp)	11041	B4GAT1	5208	['GTTCCATCCCCAAA AGGAGGCTGAARAT CAGCACAATAAGAT CCTATATCG']	5410	['CCAAAAGGAGG CTGAARATCAGC']	5654-5656	['CCCCAAAAGGAG GCTGAARATCA', 'CCCAAAAGGAGG CTGAARATCAG', 'CCAAAAGGAGGC TGAARATCAGC']
NM_000218.2(KCNQ1):c.332A>G (p.Tyr111Cys)	3784	KCNQ1	5209	['CGCACCCACGTCC AGGGCCGCGTCTRC AACTTCCTCGAGCG TCCCACCGGC']	5411	['CCAGGGCCGCGT CTRCAACTTCC']	5657	['CCAGGGCCGCGT CTRCAACTTCC']
NM_000492.3(CFTR):c.1A>G (p.Met1Val)	1080	CFTR	5210	['CAGGGACCCCAGC GCCCGAGAGACCRT GCAGAGGTCGCCTC TGGAAAAGGC']	5412	['CCAGCGCCCGAG AGACCRTGCAG']	5658-5659	['CCCAGCGCCCGA GAGACCRTGCA', 'CCAGCGCCCGAG AGACCRTGCAG']
NM_007294.3(BRCA1):c.122A>G (p.His41Arg)	672	BRCA1	5211	['GAACCTGTCTCCAC AAAGTGTGACCRCA TATTTTGCAAGTAA GTTTGAATG']	5413	['CCACAAAGTGTG ACCRCATATTT']	5660	['CCACAAAGTGTG ACCRCATATTT']
NM_007294.3(BRCA1):c.4485-2A>G	672	BRCA1	5212	['GTTTTCTCCTTCCA TTTATCTTTCTRGGT CATCCCCTTCTAAAT GCCCATC']	5414	['CCTTCCATTTATC TTTCTRGGTC']	5661-5662	['CCTTCCATTTATC TTTCTRGGTC', 'CCATTTATCTTTC TRGGTCATCC']
NM_014795.3(ZEB2):c.3134A>G (p.His1045Arg)	9839	ZEB2	5213	['AAACACAAGCACC ACCTTATCGAGCRC TCAAGGCTTCACTC GGGCGAGAAG']	5415	['CCACCTTATCGA GCRCTCAAGGC']	5663	['CCACCTTATCGA GCRCTCAAGGC']
NM_001287.5(CLCN7):c.296A>G (p.Tyr99Cys)	1186	CLCN7	5214	['TGTCCCGGCCTGCA GAGCTTGGACTRTG ACAACAGTGAGAAC CAGCTGTTC']	5416	['CCTGCAGAGCTT GGACTRTGACA']	5664	['CCTGCAGAGCTT GGACTRTGACA']
NM_080605.3(B3GALT6):c.1A>G (p.Met1Val)	126792	B3GALT6	5215	['CGCCACGCCCGCC GCAGCAGCTTCAYG GCGCCCGCGCCGGG CCGGCGGCCC']	5417	['CCCGCCGCAGCA GCTTCAYGGCG']	5665-5667	['CCCGCCGCAGCA GCTTCAYGGCG', 'CCGCCGCAGCAG CTTCAYGGCGC', 'CCGCAGCAGCTT CAYGGCGCCCG']
NM_000207.2(IN	-1	-	5216	['TCCTGCACCGAGA	5418	['CCGAGAGAGATG	5668	['CCGAGAGAGATG

S):c.*59A>G				GAGATGGAATAARG CCCTTGAACCAGCC CTGCTGTGCC']		GAATAARGCCC']		GAATAARGCCC']
NM_000784.3(C YP27A1):c.1061 A>G (p.Asp354Gly)	1593	CYP27 A1	5217	['TGGGCCCTGTACC ACCTCTCAAAGGRC CCTGAGATCCAGGA GGCCTTGCAC']	5419	['CCACCTCTCAAA GGRCCCTGAGA']	5669	['CCACCTCTCAAA GGRCCCTGAGA']
NM_000540.2(R YR1):c.14572A> G (p.Asn4858Asp)	6261	RYR1	5218	['CTACCTGTACACCG TGGTGGCCTTCRAC TTCTTCCGCAAGTTC TACAACAA']	5420	['CCGTGGTGGCCT TCRACTTCTTC']	5670	['CCGTGGTGGCCT TCRACTTCTTC']
NM_000238.3(K CNH2):c.1478A> G (p.Tyr493Cys)	3757	KCNH 2	5219	['CACCCCGGCCGCA TCGCCGTCCACTNC TTCAAGGGCTGGTT CCTCATCGAC']	5421	['CCGCATCGCCGT CCACTNCTTCA']	5671	['CCGCATCGCCGT CCACTNCTTCA']
NM_000335.4(SC N5A):c.688A>G (p.Ile230Val)	6331	SCN5A	5220	['CCGAGTCCTCCGG GCCCTGAAAACTRT ATCAGTCATTTCAG GTGAAAATCA']	5422	['CCGGGCCCTGAA AACTRTATCAG']	5672	['CCGGGCCCTGAA AACTRTATCAG']
NM_000169.2(G LA):c.548-2A>G	-1	-	5221	['TATTTTACCCATTG TTTTCTCATACRGGT TATAAGCACATGTC CTTGGCCC']	5423	['CCCATTGTTTTCT CATACRGGTT']	5673-5674	['CCCATTGTTTTCT CATACRGGTT', 'CCATTGTTTTCTC ATACRGGTTA']
NM_000146.3(FT L):c.1A>G (p.Met1Val)	2512	FTL	5222	['GTTAGCTCCTTCTT GCCAACCAACCRTG AGCTCCCAGATTCG TCAGAATTA']	5424	['CCTTCTTGCCAA CCAACCRTGAG']	5675	['CCTTCTTGCCAA CCAACCRTGAG']
NM_000531.5(O TC):c.1034A>G (p.Tyr345Cys)	5009	OTC	5223	['GTCATGGTGTCCCT GCTGACAGATTRCT CACCTCAGCTCCAG AAGCCTAAA']	5425-5426	['CCCTGCTGACAG ATTRCTCACCT', 'CCTGCTGACAGA TTRCTCACCTC']	5676-5677	['CCCTGCTGACAG ATTRCTCACCT', 'CCTGCTGACAGA TTRCTCACCTC']
NM_000531.5(O TC):c.350A>G (p.His117Arg)	5009	OTC	5224	['TGTTTTCTTACCAC ACAAGATATTCDTT TGGGTGTGAATGAA AGTCTCACG']	5427	['CCACACAAGATA TTCDTTTGGGT']	5678	['CCACACAAGATA TTCDTTTGGGT']
NM_000531.5(O TC):c.524A>G (p.Asp175Gly)	5009	OTC	5225	['TACCATCCTATCCA GATCCTGGCTGDTT ACCTCACGCTCCAG GTTGGTTTA']	5428	['CCAGATCCTGGC TGDTTACCTCA']	5679	['CCAGATCCTGGC TGDTTACCTCA']
NM_000531.5(O TC):c.527A>G (p.Tyr176Cys)	5009	OTC	5226	['CATCCTATCCAGAT CCTGGCTGATTRCCT CACGCTCCAGGTTG GTTTATTT']	5429	['CCAGATCCTGGC TGATTRCCTCA']	5680	['CCAGATCCTGGC TGATTRCCTCA']
NM_000531.5(O	5009	OTC	5227	['TCTCCTTCATCCCG	5430	['CCGTGCCTTTTA	5681-	['CCCGTGCCTTTT

TC):c.542A>G (p.Glu181Gly)				TGCCTTTTAGGRAC ACTATAGCTCTCTG AAAGGTCTT']		GGRACACTATA']	5682	AGGRACACTAT', 'CCGTGCCTTTTAG GRACACTATA']
NM_024301.4(FK RP):c.1A>G (p.Met1Val)	7914 7	FKRP	5228	['CCAGCTAGCCCCA GACTTCGGCCCCRT GCGGCTCACCCGCT GCCAGGCTGC']	5431	['CCCCAGACTTCG GCCCCRTGCGG']	5683-5685	['CCCCAGACTTCG GCCCCRTGCGG', 'CCCAGACTTCGG CCCCRTGCGGC', 'CCAGACTTCGGC CCCRTGCGGCT']
NM_000321.2(R B1):c.1927A>G (p.Lys643Glu)	5925	RB1	5229	['AGCCTTCCAGACC CAGAAGCCATTGRA ATCTACCTCTCTTTC ACTGTTTTA']	5432	['CCCAGAAGCCAT TGRAATCTACC']	5686	['CCCAGAAGCCAT TGRAATCTACC']
NM_015713.4(R RM2B):c.581A> G (p.Glu194Gly)	5048 4	RRM2 B	5230	['AAAAGATCCTGAG AAGAAAACTCCTYC TACAGCAGCAAAGG CCACCACTCT']	5433	['CCTGAGAAGAAA ACTCCTYCTAC']	5687	['CCTGAGAAGAAA ACTCCTYCTAC']
NM_000219.5(K CNE1):c.242A>G (p.Tyr81Cys)	3753	KCNE1	5231	['CACTCGAACGACC CATTCAACGTCTDC ATCGAGTCCGATGC CTGGCAAGAG']	5434	['CCCATTCAACGT CTDCATCGAGT']	5688	['CCCATTCAACGT CTDCATCGAGT']
NM_003108.3(SO X11):c.347A>G (p.Tyr116Cys)	6664	SOX11	5232	['AAGCACATGGCCG ACTACCCCGACTRC AAGTACCGGCCCCG GAAAAAGCCC']	5435	['CCGACTACCCCG ACTRCAAGTAC']	5689	['CCGACTACCCCG ACTRCAAGTAC']
NM_002764.3(PR PS1):c.343A>G (p.Met115Val)	5631	PRPS1	5233	['AATCTCAGCCAAG CTTGTTGCAAATRTG CTATCTGTAGCAGG TGCAGATCA']	5436	['CCAAGCTTGTTG CAAATRTGCTA']	5690	['CCAAGCTTGTTG CAAATRTGCTA']
NM_000546.5(TP 53):c.1101-2A>G	7157	TP53	5234	['TCTCCTCCCTGCTT CTGTCTCCTACRGCC ACCTGAAGTCCAAA AAGGGTCA']	5437	['CCTGCTTCTGTCT CCTACRGCCA']	5691	['CCTGCTTCTGTCT CCTACRGCCA']
NM_000166.5(GJ B1):c.580A>G (p.Met194Val)	2705	GJB1	5235	['CGAGAAAACCGTC TTCACCGTCTTCRTG CTAGCTGCCTCTGG CATCTGCAT']	5438	['CCGTCTTCACCG TCTTCRTGCTA']	5692	['CCGTCTTCACCG TCTTCRTGCTA']
NM_003159.2(C DKL5):c.449A>G (p.Lys150Arg)	6792	CDKL5	5236	['TTAATCAGCCACA ATGATGTCCTAARA CTGTGTGACTTTGGT AAGTTAAAA']	5439	['CCACAATGATGT CCTAARACTGT']	5693	['CCACAATGATGT CCTAARACTGT']
NM_000053.3(A TP7B):c.122A>G (p.Asn41Ser)	540	ATP7B	5237	['ATCCAGACCACCTT CATAGCCAACAYTG TCAAAAGCAAAACT	5440	['CCACCTTCATAG CCAACAYTGTC']	5694-5695	['CCACCTTCATAG CCAACAYTGTC', 'CCTTCATAGCCAA

				CTTCTTCAT']				CAYTGTCAAA']
NM_006306.3(SMC1A):c.3254A>G (p.Tyr1085Cys)	8243	SMC1A	5238	['GTGGCTACCAACATTGATGAGATCTRTAAGGCCCTGTCCCGCAATAGCAGT']	5441	['CCAACATTGATGAGATCTRTAAG']	5696	['CCAACATTGATGAGATCTRTAAG']
NM_005154.4(USP8):c.2150A>G (p.Tyr717Cys)	9101	USP8	5239	['GAACCTTCCAAACTGAAGCGCTCCTDCTCCTCCCCAGATATAACCCAGGCT']	5442	['CCAAACTGAAGCGCTCCTDCTCC']	5697	['CCAAACTGAAGCGCTCCTDCTCC']
NM_000117.2(EMD):c.266-2A>G	2010	EMD	5240	['TCTGCTACCGCTGCCCCCTTCCCARGGCTACAATGACGACTACTATGAAG']	5443	['CCGCTGCCCCCCTTCCCARGGCT']	5698	['CCGCTGCCCCCCTTCCCARGGCT']
NM_207352.3(CYP4V2):c.1393A>G (p.Arg465Gly)	285440	CYP4V2	5241	['CTACGTGCCCTTCTCTGCTGGCCCCRGGAACTGTATAGGTTTGTATCCATC']	5444	['CCCTTCTCTGCTGGCCCCRGGAA']	5699-5700	['CCCTTCTCTGCTGGCCCCRGGAA', 'CCTTCTCTGCTGGCCCCRGGAAC']
NM_000546.5(TP53):c.709A>G (p.Met237Val)	7157	TP53	5242	['CTGTACCACCATCCACTACAACTACRTGTGTAACAGTTCCTGCATGGGCGG']	5445	['CCATCCACTACAACTACRTGTGT']	5701	['CCATCCACTACAACTACRTGTGT']
NM_016069.9(PAM16):c.226A>G (p.Asn76Asp)	-1	-	5243	['CTCACCCGTCCCCTCTCCTCTGCAGRACTATGAACACTTATTTAAGGTGAA']	5446	['CCTCTCCTCTGCAGRACTATGAA']	5702-5704	['CCCCTCTCCTCTGCAGRACTATG', 'CCCTCTCCTCTGCAGRACTATGA', 'CCTCTCCTCTGCAGRACTATGAA']
NM_006785.3(MALT1):c.1019-2A>G	10892	MALT1	5244	['AACACCCCCTTTCTTTTTTTTCAARGCGAAGGACAAGGTTGCCCTTTTGA']	5447	['CCTTTCTTTTTTTTTCAARGCGA']	5705	['CCTTTCTTTTTTTTTCAARGCGA']
NM_004771.3(MMP20):c.611A>G (p.His204Arg)	9313	MMP20	5245	['GGAGAAGGCCTGGGAGGAGATACACRTTTCGACAATGCTGAGAAGTGGACT']	5448	['CCTGGGAGGAGATACACRTTTCG']	5706	['CCTGGGAGGAGATACACRTTTCG']
NM_003159.2(CDKL5):c.458A>G (p.Asp153Gly)	6792	CDKL5	5246	['CACAATGATGTCCTAAAACTGTGTGRCTTTGGTAAGTTAAAAAGAAATTAA']	5449	['CCTAAAACTGTGTGRCTTTGGTA']	5707	['CCTAAAACTGTGTGRCTTTGGTA']
NM_001204830.1(LIPT1):c.535A>G (p.Thr179Ala)	-1	-	5247	['CCGGACTACTGCCTATCACCATTGCRCTTTATTATGTAGTACTGATGGGAC']	5450	['CCTATCACCATTGCRCTTTATTA']	5708	['CCTATCACCATTGCRCTTTATTA']
NM_000921.4(PD	5139	PDE3A	5248	['AGTTTCTTCCACTT	5451	['CCACTTGGACCA	5709	['CCACTTGGACCA

E3A):c.1333A>G (p.Thr445Ala)				GGACCACCACCRCC TCGGCCACAGGTCT ACCCACCTT']		CCACCRCCTCG']		CCACCRCCTCG']
NM_000182.4(H ADHA):c.919-2A>G	3030	HADH A	5249	['TTGCTCAATTCCAG TCTTTACCACCYAA AAAACATATAAAGC ACTTGCTCA']	5452	['CCAGTCTTTACC ACCYAAAAAAC']	5710	['CCAGTCTTTACC ACCYAAAAAAC']
NM_000169.2(G LA):c.620A>G (p.Tyr207Cys)	-1	-	5250	['GTGTACTCCTGTGA GTGGCCTCTTTRTAT GTGGCCCTTTCAAA AGGTGAGA']	5453	['CCTGTGAGTGGC CTCTTTRTATG']	5711	['CCTGTGAGTGGC CTCTTTRTATG']
NM_000238.3(K CNH2):c.2582A>G (p.Asn861Ser)	3757	KCNH 2	5251	['TGGTCCAGCCTGG AGATCACCTTCANC CTGCGAGATGTGAG TTGGCTGCCC']	5454	['CCTGGAGATCAC CTTCANCCTGC']	5712	['CCTGGAGATCAC CTTCANCCTGC']
NM_000218.2(K CNQ1):c.605A>G (p.Asp202Gly)	3784	KCNQ 1	5252	['GCTCCCCCTCTCCT GCACTCCACAGRCC TCATCGTGGTCGTG GCCTCCATG']	5455	['CCTGCACTCCAC AGRCCTCATCG']	5713	['CCTGCACTCCAC AGRCCTCATCG']
NM_012203.1(G RHPR):c.934A>G (p.Asn312Asp)	9380	GRHP R	5253	['CACCATGTCCTTGT TGGCAGCTAACRAC TTGCTGGCTGGCCT GAGAGGGGA']	5456	['CCTTGTTGGCAG CTAACRACTTG']	5714	['CCTTGTTGGCAG CTAACRACTTG']
NM_021007.2(SC N2A):c.387-2A>G	6326	SCN2A	5254	['ACTTTGTCTTCCTT GACGATATTCTRCTT TATTCAATATGCTCA TTATGTG']	5457	['CCTTGACGATAT TCTRCTTTATT']	5715	['CCTTGACGATAT TCTRCTTTATT']
NM_002693.2(PO LG):c.2840A>G (p.Lys947Arg)	5428	POLG	5255	['GTGGGCATCAGCC GTGAGCATGCCARA ATCTTCAACTACGG CCGCATCTAT']	5458	['CCGTGAGCATGC CARAATCTTCA']	5716	['CCGTGAGCATGC CARAATCTTCA']
NM_020533.2(M COLN1):c.1406A>G (p.Asn469Ser)	5719 2	MCOL N1	5256	['TCTGAGTGCCTGTT CTCGCTCATCARTG GGGACGACATGTTT GTGACGTTC']	5459	['CCTGTTCTCGCT CATCARTGGGG']	5717	['CCTGTTCTCGCT CATCARTGGGG']
NM_000069.2(C ACNA1S):c.3526-2A>G	779	CACN A1S	5257	['TCGCTTTCCCATCC TTTTCCTTCCCRGGG CTACTTTGGAGACC CCTGGAAT']	5460	['CCCATCCTTTTCC TTCCCRGGGC']	5718-5719	['CCCATCCTTTTCC TTCCCRGGGC', 'CCATCCTTTTCCT TCCCRGGGCT']
NM_017662.4(TR PM6):c.3173A>G (p.Tyr1058Cys)	1408 03	TRPM6	5258	['CAAGCTGTCTACCT CTTCGTGCAATRTAT CATCATGGTGAACC TGTTGATT']	5461	['CCTCTTCGTGCA ATRTATCATCA']	5720	['CCTCTTCGTGCA ATRTATCATCA']
NM_006642.3(SD	1080	SDCC	5259	['AATAAACCCTCTG	5462	['CCTCTGCTTTTGC	5721	['CCTCTGCTTTTGC

CCAG8):c.221-2A>G	6	AG8		CTTTTGCTCTATRGTTAATCAGCTCAAAGATTTGTTGC']		TCTATRGTTA']		TCTATRGTTA']
NM_003560.2(PLA2G6):c.1349-2A>G	8398	PLA2G6	5260	['CAGCATGCCCTGCTCTGTGCCTCACRGAACTACAGGATCTCATGCACATCT']	5463	['CCCTGCTCTGTGCCTCACRGAAC']	5722-5723	['CCCTGCTCTGTGCCTCACRGAAC', 'CCTGCTCTGTGCCTCACRGAACT']

EXAMPLE 6: Next Generation C to T Editors

[00351] Other familes of cytidine deaminases as alterantives to base etitor 3 (BE3) constructs were examined. The different C to T editors were developed to have a narrow or different editing window, alternate sequence specificity to expand targetable substrates, and to have higher activity.

[00352] Using the methods described in Example 4, the pmCDA1 (cytidine deaminase 1 from *Petromyzon marinus*) activity at the HeK-3 site is evaluated (Figure 42). The pmCDA1-nCas9-UGI-NLS (nCas9 indicates the Cas9 nickase described herein) construct is active on some sites (e.g., the C bases on the complementary strand at position 9, 5, 4, and 3) that are not accessible with rAPOBEC1 (BE3).

[00353] The pmCDA1 activity at the HeK-2 site is given in Figure 43. The pmCDA1-XTEN-nCas9-UGI-NLS construct is active on sites adjacent to "G," while rAPOBEC1 analog (BE3 construct) has low activity on "C"s that are adjacent to "G"s, e.g., the C base at position 11 on the complementary strand.

[00354] The percent of total sequencing reads with target C converted to T (Figure 44), C converted to A (Figure 45), and C converted to G (Figure 46) are shown for CDA and APOBEC1 (the BE3 construct).

[00355] The huAPOBEC3G activity at the HeK-2 site is shown in Figure 47. Two constructs were used: huAPOBEC3G-XTEN-nCas9-UGI-NLS and huAPOBEC3G*(D316R_D317R)-XTEN-nCas9-UGI-NLS. The huAPOBEC3G-XTEN-nCas9-UGI-NLS construct has different sequence specificity than rAPOBEC1 (BE3), as shown in Figure 47, the editing window appears narrow, as indicated by APOBEC3G's descreased activity at position 4 compared to APOBEC1. Mutations made in huAPOBEC3G (D316R and D317R) increased ssDNA binding and resulted in an observable effect on expanding the sites which were edited (compare APOBEC3G with APOBEC3G_RR in Figure 47). Mutations were chosen based on APOBEC3G crystal structure, *see*: Holden *et al.,* Crystal structure of the anti-viral APOBEC3G catalytic domain and functional

implication. *Nature.* (2008); 121-4, the entire contents of which are incorporated herein by reference.

EXAMPLE 7: pmCDA1/huAPOBEC3G/rAPOBEC1 work in E. coli

[00356] LacZ selection optimization for the A to I conversion was performed using a bacterial strain with lacZ encoded on the F plasmid. A critical glutamic acid residue was mutated (e.g., GAG to GGG, Glu to Gly mutation) so that G to A by a cytidine deaminase would restore lacZ activity (Figure 48). Strain CC102 was selected for the selection assay. APOBEC1 and CDA constructs were used in a selection assay to optimize G to A conversion.

[00357] To evaluate the the effect of copy number of the plasmids encoding the deaminase constructs on lacZ reversion frequency, the CDA and APOBEC1 deaminases were cloned into 4 plasmids with different replication origins (hence different copy numbers), SC101, CloDF3, RSF1030, and PUC (copy number: PUC>RSF1030>CloDF3>SC101) and placed under an inducible promoter. The plasmids were individually transformed into E. coli cells harboring F plasmid containing the mutated LacZ gene. The expression of the deaminases were induced and LacZ activity was detected for each construct (Figure 49). As shown in Figure 49, CDA exhibited significantly higher activity than APOBEC1 in all instances, regardless of the plasmid copy number the deaminases were cloned in. Further, In terms of the copy number, the deaminase activity was positively correlated with the copy number of the plasmid they are cloned in, i.e., PUC> CloDF3>SC101.

[00358] LacZ reversions were confirmed by sequencing of the genomic DNA at the lacZ locus. To obtain the genomic DNA containing the corrected LacZ gene, cells were grown media containg X-gal, where cells having LacZ activity form blue colonies. Blue colonies were selected and grown in minimial media containing lactose. The cells were spun down, washed, and re-plated on minimal media plates (lactose). The blue colony at the highest dilution was then selected, and its genomic DNA was sequenced at the lacZ locus (Figure 50).

[00359] A chloramphenicol reversion assay was designed to test the activity of different cytidine deaminases (e.g., CDA, and APOBEC1). A plasmid harboring a mutant CAT1 gene which confers chloramphenicol resistance to bacteria is constructed with RSF1030 as the replication origin. The mutant CAT1 gene encodings a CAT1 protein that has a H195R (CAC to CGC) mutation, rendering the protein inactive (Figure 51). Deamination of the C base-paired to the G base in the CGC codon would convert the codon back to a CAC codon,

restoring the activity of the protein. As shown in Figure 52, CDA outperforms rAPOBEC in *E. coli* in restoring the acitivyt of the chloramphenicol reisitance gene. The minimum inhibitory concentration (MIC) of chlor in S1030 with the selection plasmid (pNMG_ch_5) was approximately 1 μg/mL. Both rAPOBEC-XTEN-dCas9-UGI and CDA-XTEN-dCas9-UGI induced DNA correction on the selection plasmid (Figure 53).

[00360] Next, the huAPOBEC3G-XTEN-dCas9-UGI protein was tested in the same assay. Interestingly, huAPOBEC3G-XTEN-dCas9-UGI exhibited different sequence specificity than the rAPOBEC1-XTEN-dCas9-UGI fusion protein. Only position 8 was edited with APOBEC3G-XTEN-dCas9-UGI fusion, as compared to the rAPOBEC11-XTEN-dCas9-UGIfusion (in which positions 3, 6, and 8 were edited) (Figure 54).

EXAMPLE 8: C to T Base Editors with Less Off Target Editing

[00361] Current base editing technologies allow for the sequence-specific conversion of a C:G base pair into a T:A base pair in genomic DNA. This is done via the direct catalytic conversion of cytosine to uracil by a cytidine deaminase enzyme and thus, unlike traditional genome editing technologies, does not introduce double-stranded DNA breaks (DSBs) into the DNA as a first step. See, Komor, A.C., Kim, Y.B., Packer, M.S., Zuris, J.A., and Liu, D.R. (2016), "Programmable editing of a target base in genomic DNA without double-stranded DNA cleavage." Nature *533*, 420-424; the entire contents of which are incorporated by reference herein. Instead, catalytically dead SpCas9 (dCas9) or a SpCas9 nickase (dCas9(A840H)) is tethered to a cytidine deaminase enzyme such as rAPOBEC1, pmCDA1, or hAPOBEC3G. The genomic locus of interest is encoded by an sgRNA, and DNA binding and local denaturation is facilitated by the dCas9 portion of the fusion. However, just as wt dCas9 and wt Cas9 exhibit off-target DNA binding and cleavage, current base editors also exhibit C to T editing at Cas9 off-target loci, which limits their therapeutic usefulness.

[00362] It has been reported that the introduction of just three to four mutations into SpCas9 that neutralize nonspecific electrostatic interactions between the protein and the sugar-phosphate backbone of its target DNA, increases the DNA binding specificity of SpCas9. See, Kleinstiver, B.P., Pattanayak, V., Prew, M.S., Tsai, S.Q., Nguyen, N.T., Zheng, Z., and Joung, J.K. (2016) "High-fidelity CRISPR–Cas9 nucleases with no detectable genome-wide off-target effects." Nature *529*, 490-495; and Slaymaker, I.M., Gao, L., Zetsche, B., Scott, D.A., Yan, W.X., and Zhang, F. (2015) "Rationally engineered Cas9 nucleases with improved specificity. Science *351*, 84-88; the entire contents of each are hereby incorporated by reference herein. Four reported neutralizing mutations were therefore

incorporated into the initially reported base editor BE3 (SEQ ID NO: 285), and found that off-target C to T editing of this enzyme is also drastically reduced (Figure 55), with no decrease in on-target editing (Figure 56).

[00363] As shown in Figure 55, HEK293T cells were transfected with plasmids expressing BE3 or HF-BE3 and a sgRNA matching the EMX1 sequence using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target locus, plus the top ten known Cas9 off-target loci for the EMX1 sgRNA, as previously determined by Joung and coworkers using the GUIDE-seq method. See Tsai, S.Q., Zheng, Z., Nguyen, N.T., Liebers, M., Topkar, V.V., Thapar, V., Wyvekens, N., Khayter, C., Iafrate, A.J., Le, L.P., *et al.* (2015) "GUIDE-seq enables genome-wide profiling of off-target cleavage by CRISPR-Cas nucleases." *Nat Biotech 33*, 187-197; the entire contents of which are incorporated by reference herein. EMX1 off-target 5 locus did not amplify and is not shown. Sequences of the on-target and off-target protospacers and protospacer adjacent motifs (PAMs) are displayed (Figure 55). Cellular C to T conversion percentages, defined as the percentage of total DNA sequencing reads with T at each position of an original C within the protospacer, are shown for BE3 and HF-BE3.

[00364] In Figure 56, HEK293T cells were transfected with plasmids expressing BE3 or HF-BE3 and sgRNAs matching the genomic loci indicated using Lipofectamine 2000. Three days after transfection, genomic DNA was extracted, amplified by PCR, and analyzed by high-throughput DNA sequencing at the on-target loci. The percentage of total DNA sequencing reads with all four bases at the target Cs within each protospacer are shown for treatment with BE3 or HF-BE3 (Figure 56). Frequencies of indel formation are shown as well.

[00365] Primary Protein Sequence of HF-BE3 (SEQ ID NO: 285):

MSSETGPVAVDPTLRRRIEPHEFEVFFDPRELRKETCLLYEINWGGRHSIWRHTSQNTNKHVEVNFIEKF
TTERYFCPNTRCSITWFLSWSPCGECSRAITEFLSRYPHVTLFIYIARLYHHADPRNRQGLRDLISSGVTI
QIMTEQESGYCWRNFVNYSPSNEAHWPRYPHLWVRLYVLELYCIILGLPPCLNILRRKQPQLTFFTIALQ
SCHYQRLPPHILWATGLKSGSETPGTSESATPESDKKYSIGLAIGTNSVGWAVITDEYKVPSKKFKVLG
NTDRHSIKKNLIGALLFDSGETAEATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFL
VEEDKKHERHPIFGNIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLN
PDNSDVDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGNLIALS
LGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAILLSDILRVNTEITK
APLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYAGYIDGGASQEEFYKFIKPILEKM
DGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELHAILRRQEDFYPFLKDNREKIEKILTFRIPYYVGP
LARGNSRFAWMTRKSEETITPWNFEEVVDKGASAQSFIERMTAFDKNLPNEKVLPKHSLLYEYFTVYN
ELTKVKYVTEGMRKPAFLSGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASL
GTYHDLLKIIKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWG
ALSRKLINGIRDKQSGKTILDFLKSDGFANRNFMALIHDDSLTFKEDIQKAQVSGQGDSLHEHIANLAGS
PAIKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRERMKRIEEGIKELGSQILKEH

PVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDHIVPQSFLKDDSIDNKVLTRSDKNRGKS
DNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNLTKAERGGLSELDKAGFIKRQLVETRAITKHVAQIL
DSRMNTKYDENDKLIREVKVITLKSKLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPK
LESEFVYGDYKVYDVRKMIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIV
WDKGRDFATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVA
YSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPKYSLFELENGR
KRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVEQHKHYLDEIIEQISEFSKR
VILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGAPAAFKYFDTTIDRKRYTSTKEVLDATLIH
QSITGLYETRIDLSQLGGDSGGSTNLSDIIEKETGKQLVIQESILMLPEEVEEVIGNKPESDILVHTAYDES
TDENVMLLTSDAPEYKPWALVIQDSNGENKIKMLSGGSPKKKRKV

EXAMPLE 9: Development of Base Editors that Use Cas9 Variants and Modulation of the Base Editor Processivity to Increase the Target Range and Precision of the Base Editing Technology

[00366] Unlike traditional genome editing platforms, base editing technology allows precise single nucleotide changes in the DNA without inducing double-stranded breaks(DSBs). See, Komor, A. C. *et al. Nature* **533**, 420-424 (2016). The current generation of base editor uses the NGG PAM exclusively. This limits its ability to edit desired bases within the genome, as the base editor needs to be placed at a precise location where the target base is placed within a 4-base region (the 'deamination window'), approximately 15 bases upstream of the PAM . See, Komor, A. C. *et al. Nature* **533**, 420-424 (2016). Moreover, due to the high processivity of cytidine deaminase, the base editor may convert all cytidines within its deamination window into thymidines, which could induce amino acid changes other than the one desired by the researcher. See, Komor, A. C. *et al. Nature* **533**, 420-424 (2016).

Expanding the scope of base editing through the development of base editors with Cas9 variants

[00367] Cas9 homologs and other RNA-guided DNA binders that have different PAM specificities were incorporated into the base editor architecture. See, Kleinstiver, B. P. *et al. Nature* **523**, 481–485 (2015); Kleinstiver, B. P. *et al. Nature Biotechnology* **33**, 1293–1298 (2015); and Zetsche, B. *et al. Cell* **163**, 759-771 (2015); the entire contents of each are incorporated by reference herein. Furthermore, innovations that have broadened the PAM specificities of various Cas9 proteins were also incorporated to expand the target reach of the base editor even more. See, Kleinstiver, B. P. *et al. Nature* **523**, 481–485 (2015); and Kleinstiver, B. P. *et al. Nature Biotechnology* **33**, 1293–1298 (2015). The current palette of base editors is summarized in Table 4.

Table 4. New base editors made from Cas9 Variants

Species	PAM	Base Editor Name	Reference for Cas9 variant
S. pyogenes	...NGG	BE3	Wild-type
	...NGA	VQR BE3 or EQR BE3	Kleinstiver, B. P. *et al.*
	...NGCG	VRER BE3	Kleinstiver, B. P. *et al.*
S. aureus	...NNGRRT	SaBE3	Wild-type
	...NNNRRT	SaKKH BE3	Kleinstiver, B. P. *et al.*
L. bacterium	TTTN...	dCpf1 BE2	Zetsche, B. *et al.*

Modulating base editor's processivity through site-directed mutagenesis of rAPOBEC1

[00368] It was reasoned that the processivity of the base editor could be modulated by making point mutations in the deaminase enzyme. The incorporatation of mutations that slightly reduce the catalytic activity of deaminase in which the base editor could still catalyze on average one round of cytidine deamination but was unlikely to access and catalyze another deamination within the relevant timescale were pursued. In effect, the resulting base editor would have a narrower deamination window.

[00369] rAPOBEC1 mutations probed in this work are listed in Table 5. Some of the mutations resulted in slight apparent impairment of rAPOBEC1 catalysis, which manifested as preferential editing of one cytidine over another when multiple cytidines are found within the deamination window. Combining some of these mutations had an additive effect, allowing the base editor to discriminate substrate cytidines with higher stringency. Some of the double mutants and the triple mutant allowed selective editing of one cytidine among multiple cytidines that are right next to one another (Figure 57).

Table 5. rAPOBEC1 Point Mutations Investigated

rAPOBEC1 mutation studied in this work	Corresponding mutation in APOBEC3G	Reference
H121R / H122R	D315R / D316R	Holden, L. G. *et al.*
R126A	R320A	Chen, K-M. *et al.*
R126E	R320E	Chen, K-M. *et al.*
R118A	R313A	Chen, K-M. *et al.*
W90A	W285A	Chen, K-M. *et al.*
W90Y	W285Y	
R132E	R326E	

Base Editor PAM Expansion and Processivity Modulation

[00370] The next generation of base editors were designed to expand editable cytidines in the genome by using other RNA-guided DNA binders (Figure 58). Using a NGG PAM only

allows for a single target within the "window" whereas the use of multiple different PAMs allows for Cas9 to be positioned anywhere to effect selective deamination. A variety of new base editors have been created from Cas9 variants (Figure 59 and Table 4). Different PAM sites (NGA, Figure 60; NGCG, Figure 61; NNGRRT, Figure 62; and NNHRRT, Figure 63) were explored. Selective deamination was successfully achieved through kinetic modulation of cytidine deaminase point mutagenesis (Figure 65 and Table 5).

[00371] The effect of various mutations on the deamination window was then investigated in cell culture using spacers with multiple cytidines (Figures 66 and 67).

[00372] Further, the effect of various mutations on different genomic sites with limited numbers of cytidines was examined (Figures 68 to 71). It was found that approximately one cytidine will be edited within the deamination windown in the spacer, while the rest of the cytidines will be left intact. Overall, the preference for editing is as follows: $C_6 > C_5 >> C_7 \approx C_4$.

Base Editing Using Cpf1

[00373] Cpf1, a Cas9 homolog, can be obtained as AsCpf1, LbCpf1, or from any other species. Schematics of fusion constructs, including BE2 and BE3 equivalents, are shown in Figure 73. The BE2 equivalent uses catalytically inactive Cpf2 enzyme (dCpf1) instead of Cas9, while the BE3 equivalent includes the Cpf1 mutant, which nicks the target strand. The bottom schematic depicts different fusion architectures to combine the two innovations illustrated above it (Figure 73). The base editing results of HEK293T cell TTTN PAM sites using Cpf1 BE2 were examined with different spacers (Figures 64A to 64C). In some embodiments, Cpf1 may be used in place of a Cas9 domain in any of the base editors provided herein. In some embodiments, the Cpf1 is a protein that is at lesst 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 98%, 99%, or 99.5% identical to SEQ ID NO 313.

Full Protein Sequence of Cpf1 (SEQ ID NO: 313):

MSIYQEFVNKYSLSKTLRFELIPQGKTLENIKARGLILDDEKRAKDYKKAKQIIDKYHQFFIEEILSSVCIS
EDLLQNYSDVYFKLKKSDDDNLQKDFKSAKDTIKKQISEYIKDSEKFKNLFNQNLIDAKKGQESDLILW
LKQSKDNGIELFKANSDITDIDEALEIIKSFKGWTTYFKGFHENRKNVYSSNDIPTSIIYRIVDDNLPKFLE
NKAKYESLKDKAPEAINYEQIKKDLAEELTFDIDYKTSEVNQRVFSLDEVFEIANFNNYLNQSGITKFNT
IIGGKFVNGENTKRKGINEYINLYSQQINDKTLKKYKMSVLFKQILSDTESKSFVIDKLEDDSDVVTTMQ
SFYEQIAAFKTVEEKSIKETLSLLFDDLKAQKLDLSKIYFKNDKSLTDLSQQVFDDYSVIGTAVLEYITQ
QIAPKNLDNPSKKEQELIAKKTEKAKYLSLETIKLALEEFNKHRDIDKQCRFEEILANFAAIPMIFDEIAQ
NKDNLAQISIKYQNQGKKDLLQASAEDDVKAIKDLLDQTNNLLHKLKIFHISQSEDKANILDKDEHFYL
VFEECYFELANIVPLYNKIRNYITQKPYSDEKFKLNFENSTLANGWDKNKEPDNTAILFIKDDKYYLGV
MNKKNNKIFDDKAIKENKGEGYKKIVYKLLPGANKMLPKVFFSAKSIKFYNPSEDILRIRNHSTHTKNG
SPQKGYEKFEFNIEDCRKFIDFYKQSISKHPEWKDFGFRFSDTQRYNSIDEFYREVENQGYKLTFENISES
YIDSVVNQGKLYLFQIYNKDFSAYSKGRPNLHTLYWKALFDERNLQDVVYKLNGEAELFYRKQSIPKK
ITHPAKEAIANKNKDNPKKESVFEYDLIKDKRFTEDKFFFHCPITINFKSSGANKFNDEINLLLKEKAND

VHILSIDRGERHLAYYTLVDGKGNIIKQDTFNIIGNDRMKTNYHDKLAAIEKDRDSARKDWKKINNIKE
MKEGYLSQVVHEIAKLVIEYNAIVVFEDLNFGFKRGRFKVEKQVYQKLEKMLIEKLNYLVFKDNEFDK
TGGVLRAYQLTAPFETFKKMGKQTGIIYYVPAGFTSKICPVTGFVNQLYPKYESVSKSQEFFSKFDKICY
NLDKGYFEFSFDYKNFGDKAAKGKWTIASFGSRLINFRNSDKNHNWDTREVYPTKELEKLLKDYSIEY
GHGECIKAAICGESDKKFFAKLTSVLNTILQMRNSKTGTELDYLISPVADVNGNFFDSRQAPKNMPQDA
DANGAYHIGLKGLMLLGRIKNNQEGKKLNLVIKNEEYFEFVQNRNN

EXAMPLE 10: Increased Fidelity of Base Editing

[00374] Examining the difference between plasmid delivery of BE3 and HF-BE3, it was found that the two edit on-target loci with comparable efficiency (Figures 74 and 75). However, HF-BE3 edited off-target loci much less than BE3, meaning that HF-BE3 has a much higher DNA specificity than BE3 (Figure 76). Deaminase protein lipofection to HEK cells demonstrated that protein delivery of BE3 results in comparable on-target activity, but much better specificity, than plasmid DNA delivery of BE3. Using improved transfection procedures and better plasmids (n=2), the experiment used the following conditions: protein delivery was 125 nM Cas9:sgRNA complex, plasmid delivery was 750ng BE3/HF-BE3 plasmid + 250ng sgRNA plasmid, and lipofection was with 1.5μL of Lipofectamine 2000 per well. EMX-1 off target site 2 and FANCF off-target site 1 showed the most off-target editing with BE3, compared to all of the off-targets assayed (Figures 77 and 78), while HEK-3 showed no significant editing at off-targets for any of the delivery methods (Figure 79). HEK-4 shows some C-to-G editing on at the on-target site, while its off-target sites 1, 3, and 4 showed the most off-target editing of all the assayed sites (Figure 80).

Delivery of BE3 Protein via Micro-injection to Zebrafish

[00375] TYR guide RNAs were tested in an *in vitro* assay for sgRNA activity (Figures 81 and 82). The % HTS reads shows how many C residues were converted to T residues during a 2h incubation with purified BE3 protein and PCR of the resulting product. Experiments used an 80-mer synthetic DNA substate with the target deamination site in 60bp of its genomic context. This is not the same as % edited DNA strands because only one strand was nicked, so the product is not amplified by PCR. The proportion of HTS reads edited is equal to x/(2-x), where x is the actual proportion of THS reads edited. For 60% editing, the actual proportion of bases edited is 75%. "Off target" is represents BE3 incubated with the same DNA substrate, while bound to an off-target sgRNA. It was found sgRNAs sgRH_13, sgHR_17, and possibly sgHR_16 appeared to be promising targets for *in vivo* injection experiments.

[00376] The delivery of BE3 protein in was tested *in vivo* in zebrafish. Zebrafish embryos (n=16-24) were injected with either scramled sgRNA, sgHR_13, sgHR_16, or sgHR_17 and purified BE3. Three embryos from each condition were analyzed independently (single embryo) and for each condition, all of the injected embryos were pooled and sequenced as a pool. The results are shown in Figures 83 to 85.

EXAMPLE 11: Uses of Base Editors to Treat Disease

[00377] Base editors or complexes provided herein (*e.g.*, BE3) may be used to modify nucleic acids. For example, base editors may be used to change a cytosine to a thymine in a nucleic acid (*e.g.,* DNA). Such changes may be made to, *inter alia*, alter the amino acid sequence of a protein, to destroy or create a start codon, to create a stop codon, to distupt splicing donors, to disrupt splicing acceptors or edit regulatory sequences. Examples of possible nucleotide changes are shown in Figure 86.

[00378] Base editors or complexes provided herein (*e.g.,* BE3) may be used to edit an isoform of Apolipoprotein E in a subject. For example, an Apolipoprotein E isoform may be edited to yield an isoform associated with a lower risk of developing Alzheimer's disease. Apolipoprotein E has four isoforms that differ at amino acids 112 and 158. APOE4 is the largest and most common genetic risk factor for late-onset Alzheimer's disease. Arginine residue 158 of APOE4, encoded by the nucleic acid sequence CGC, may be changed to a cysteine by using a base editor (*e.g.*, BE3) to change the CGC nucleic acid sequence to TGC, which encodes cysteine at residue 158. This change yields an APOE3r isoform, which is associated with lower Alzheimer's disease risk. See Figure 87.

[00379] It was tested whether base editor BE3 could be used to edit APOE4 to APOE3r in mouse astrocytes (Figure 88). APOE 4 mouse astrocytes were nucleofected with Cas9 + template or BE3, targeting the nucleic acid encoding Arginine 158 of APOE4. The Cas9 + template yielded only 0.3% editing with 26% indels, while BE3 yielded 75% editing with 5% indels. Two additional base-edited cytosines are silent and do not yield changes to the amino acid sequence (Figure 88).

[00380] Base editors or complexes provided herein may be used to treat prion protein diseases such as Creutzfeldt-Jakob disease and fatal familial insomnia, for example, by introducing mutations into a PRNP gene. Reverting PRNP mutations may not yield therapeutic results, and intels in PRNP may be pathogenic. Accordingly, it was tested whether PRNP could be mutated using base editors (*e.g.,* BE3) to introduce a premature stop codon in the PRNP gene. BE3, associated with its guide RNA,was introduced into HEK cells

or glioblastoma cells and was capable of editing the PRNP gene to change the encoded arginine at residue 37 to a stop codon. BE3 yielded 41% editing (Figure 89).

[00381] Additional genes that may be edited include the following: *APOE* editing of Arg 112 and Arg 158 to treat increased Alzheimer's risk; *APP* editing of Ala 673 to decrease Alzheimer's risk; *PRNP* editing of Arg 37 to treat fatal familial insomnia and other prion protein diseases; *DMD* editing of the exons 23 and 51 splice sites to treat Duchenne muscular dystrophy; *FTO* editing of intron 1 to treat obesity risk; *PDS* editing of exon 8 to treat Pendred syndrome (genetic deafness); *TMC1* editing of exon 8 to treat congenital hearing loss; *CYBB* editing of various patient-relevant mutations to treat chronic granulomatous disease. Additional diseases that may be treated using the base editors provided herein are shown in Table 6, below.

[00382] UGI also plays a key role. Knocking out UDG (which UGI inhibits) was shown to dramatically improve the cleanliness and efficiency of C to T base editing (Figure 90). Furthermore, base editors with nickase and without UGI were shown to produce a mixture of outcomes, with very high indel rates (Figure 91).

EXAMPLE 12: Expanding the Targeting Scope of Base Editing

[00383] Base editing is a new approach to genome editing that uses a fusion protein containing a catalytically defective *Streptococcus pyogenes* Cas9, a cytidine deaminase, and an inhibitor of base excision repair to induce programmable, single-nucleotide C→T (or G→A) changes in DNA without generating double-strand DNA breaks, without requiring a donor DNA template, and without inducing an excess of stochastic insertions and deletions[1]. The development of five new C→T (or G→A) base editors that use natural and engineered Cas9 variants with different protospacer-adjacent motif (PAM) specificities to expand the number of sites that can be targeted by base editing by 2.5-fold are described herein. Additionally, new base editors containing mutated cytidine deaminase domains that narrow the width of the apparent editing window from approximately 5 nucleotides to 1 or 2 nucleotides were engineered, enabling the discrimination of neighboring C nucleotides that would previously be edited with comparable efficiency. Together, these developments substantially increase the targeting scope of base editing.

[00384] CRISPR-Cas9 nucleases have been widely used to mediate targeted genome editing[2]. In most genome editing applications, Cas9 forms a complex with a single guide RNA (sgRNA) and induces a double-stranded DNA break (DSB) at the target site specified by the sgRNA sequence. Cells primarily respond to this DSB through the non-homologuous

end-joining (NHEJ) repair pathway, which results in stochastic insertions or deletions (indels) that can cause frameshift mutations that disrupt the gene. In the presence of a donor DNA template with a high degree of homology to the sequences flanking the DSB, gene correction can be achieved through an alternative pathway known as homology directed repair (HDR).[3,4] Unfortunately, under most non-perturbative conditions HDR is inefficient, dependent on cell state and cell type, and dominated by a larger frequency of indels.[3,4] As most of the known genetic variations associated with human disease are point mutations[5], methods that can more efficiently and cleanly make precise point mutations are needed.

[00385] Base editing, which enables targeted replacement of a C:G base pair with a T:A base pair in a programmable manner without inducing DSBs[1], has been recently described. Base editing uses a fusion protein between a catalytically inactivated (dCas9) or nickase form of *Streptococcus pyogenes* Cas9 (SpCas9), a cytidine deaminase such as APOBEC1, and an inhibitor of base excision repair such as uracil glycosylase inhibitor (UGI) to convert cytidines into uridines within a five-nucleotide window specified by the sgRNA.[1] The third-generation base editor, BE3, converts C:G base pairs to T:A base pairs, including disease-relevant point mutations, in a variety of cell lines with higher efficiency and lower indel frequency than what can be achieved using other genome editing methods[1]. Subsequent studies have validated the deaminase-dCas9 fusion approach in a variety of settings[6,7].

[00386] Efficient editing by BE3 requires the presence of an NGG PAM that places the target C within a five-nucleotide window near the PAM-distal end of the protospacer (positions 4-8, counting the PAM as positions 21-23)[1]. This PAM requirement substantially limits the number of sites in the human genome that can be efficiently targeted by BE3, as many sites of interest lack an NGG 13- to 17- nucleotides downstream of the target C. Moreover, the high activity and processivity of BE3 results in conversion of all Cs within the editing window to Ts, which can potentially introduce undesired changes to the target locus. Herein, new C:G to T:A base editors that address both of these limitations are described.

[00387] It was thought that any Cas9 homolog that binds DNA and forms an "R-loop" complex[8] containing a single-stranded DNA bubble could in principle be converted into a base editor. These new base editors would expand the number of targetable loci by allowing non-NGG PAM sites to be edited. The Cas9 homolog from *Staphylococcus aureus* (SaCas9) is considerably smaller than SpCas9 (1053 vs. 1368 residues), can mediate efficient genome editing in mammalian cells, and requires an NNGRRT PAM[9]. SpCas9 was replaced with SaCas9 in BE3 to generate SaBE3 and transfected HEK293T cells with plasmids encoding SaBE3 and sgRNAs targeting six human genomic loci (Figures 92A and 92B). After 3 d, the

genomic loci were subjected to high-throughput DNA sequencing (HTS) to quantify base editing efficiency. SaBE3 enabled C to T base editing of target Cs at a variety of genomic sites in human cells, with very high conversion efficiencies (approximately 50-75% of total DNA sequences converted from C to T, without enrichment for transfected cells) arising from targeting Cs at positions 6-11. The efficiency of SaBE3 on NNGRRT-containing target sites in general exceeded that of BE3 on NGG-containing target sites[1]. Perhaps due to its higher average efficiency, SaBE3 can also result in detectable base editing at target Cs at positions outside of the canonical BE3 activity window (Figure 92C). In comparison, BE3 showed significantly reduced editing under the same conditions (0-11%), in accordance with the known SpCas9 PAM preference (Figure 106A)[10]. These data show that SaBE3 can facilitate very efficient base editing at sites not accessible to BE3.

[00388] The targeting range of base editors was further expanded by applying recently engineered Cas9 variants that expand or alter PAM specificities. Joung and coworkers recently reported three SpCas9 mutants that accept NGA (VQR-Cas9), NGAG (EQR-Cas9), or NGCG(VRER-Cas9) PAM sequences[11]. In addition, Joung and coworkers engineered a SaCas9 variant containing three mutations (SaKKH-Cas9) that relax its PAM requirement to NNNRRT[12]. The SpCas9 portion of BE3 was replaced with these four Cas9 variants to produce VQR-BE3, EQR-BE3, VRER-BE3, and SaKKH-BE3, which target NNNRRT,NGA, NGAG, and NGCG PAMs respectively. HEK293T cells were transfected with plasmids encoding these constructs and sgRNAs targeting six genomic loci for each new base editor, and measured C to T base conversions using HTS.

[00389] SaKKH-BE3 edited sites with NNNRRT PAMs with efficiencies up to 62% of treated, non-enriched cells (Figure 92D). As expected, SaBE3 was unable to efficiently edit targets containing PAMs that were NNNHRRT (where H = A, C, or T) (Figure 92D). VQR-BE3, EQR-BE3, and VRER-BE3 exhibited more modest, but still substantial base editing efficiencies of up to 50% of treated, non-enriched cells at genomic loci with the expected PAM requirements with an editing window similar to that of BE3 (Figures 92E and 92F). Base editing efficiencies of VQR-BE3, EQR-BE3, and VRER-BE3 in general closely paralleled the reported PAM requirements of the corresponding Cas9 nucleases; for example, EQR-BE3 was unable to efficiently edit targets containing NGAH PAM sequences (Figure 92F). In contrast, BE3 was unable to edit sites with NGA or NGCG PAMs efficiently (0-3%), likely due to its PAM restrictions (Figure 106B).

[00390] Collectively, the properties of SaBE3, SaKKH-BE3, VQR-BE3, EQR-BE3, and VRER-BE3 establish that base editors exhibit a modularity that facilitates their ability to exploit Cas9 homologs and engineered variants.

[00391] Next, base editors with altered activity window widths were developed. All Cs within the activity window of BE3 can be efficiently converted to Ts[1]. The ability to modulate the width of this window would be useful in cases in which it is important to edit only a subset of Cs present in the BE3 activity window.

[00392] The length of the linker between APOBEC1 and dCas9 was previously observed to modulate the number of bases that are accessible by APOBEC1 *in vitro*[1]. In HEK293T cells, however, varying the linker length did not significantly modulate the width of the editing window, suggesting that in the complex cellular milieu, the relative orientation and flexibility of dCas9 and the cytidine deaminase are not strongly determined by linker length (Figure 96). Next, it was thought that truncating the 5' end of the sgRNA might narrow the base editing window by reducing the length of single-stranded DNA accessible to the deaminase upon formation of the RNA-DNA heteroduplex. HEK293T cells were co-transfected with plasmids encoding BE3 and sgRNAs of different spacer lengths targeting a locus with multiple Cs in the editing window. No consistent changes in the width of base editing when using truncated sgRNAs with 17- to 19-base spacers were observed (Figures 95A to 95C). Truncating the sgRNA spacer to fewer than 17 bases resulted in large losses in activity (Figure 95A).

[00393] As an alternative approach, it was thought that mutations to the deaminase domain might narrow the width of the editing window through multiple possible mechanisms. First, some mutations may alter substrate binding, the conformation of bound DNA, or substrate accessibility to the active site in ways that reduce tolerance for non-optimal presentation of a C to the deaminase active site. Second, because the high activity of APOBEC1 likely contributes to the deamination of multiple Cs per DNA binding event,[1,13,14] mutations that reduce the catalytic efficiency of the deaminase domain of a base editor might prevent it from catalyzing successive rounds of deamination before dissociating from the DNA. Once any C:G to T:A editing event has taken place, the sgRNA no longer perfectly matches the target DNA sequence and re-binding of the base editor to the target locus should be less favorable. Both strategies were tested in an effort to discover new base editors that distinguish among multiple cytidines within the original editing window.

[00394] Given the absence of an available APOBEC1 structure, several mutations previously reported to modulate the catalytic activity of APOBEC3G, a cytidine deaminase from the same family that shares 42% sequence similarity of its active site-containing domain

to that of APOBEC1, were identified[15]. Corresponding APOBEC1 mutations were incorporated into BE3 and evaluated their effect on base editing efficiency and editing window width in HEK293T cells at two C-rich genomic sites containing Cs at positions 3, 4, 5, 6, 8, 9, 10, 12, 13, and 14 (site A); or containing Cs at positions 5 ,6, 7, 8, 9, 10, 11, and 13 (site B).

[00395] The APOBEC1 mutations R118A and W90A each led to dramatic loss of base editing efficiency (Figure 97C). R132E led to a general decrease in editing efficiency but did not change the substantially narrow the shape of the editing window (Figure 97C). In contrast, several mutations that narrowed the width of the editing window while maintaining substantial editing efficiency were found (Figures 93A and 97C). The "editing window width" was defined to represent the artificially calculated window width within which editing efficiency exceeds the half-maximal value for that target. The editing window width of BE3 for the two C-rich genomic sites tested was 5.0 (site A) and 6.1 (site B) nucleotides.

[00396] R126 in APOBEC1 is predicted to interact with the phosphate backbone of ssDNA[13]. Previous studies have shown that introducing the corresponding mutation into APOBEC3G decreased catalysis by at least 5-fold[14]. Interestingly, when introduced into APOBEC1 in BE3, R126A and R126E increased or maintained activity relative to BE3 at the most strongly edited positions (C5, C6, and C7), while decreasing editing activity at other positions (Figures 93A and 97C). Each of these two mutations therefore narrowed the width of the editing window at site A and site B to 4.4 and 3.4 nucleotides (R126A), or to 4.2 and 3.1 nucleotides (R126E), respectively (Figures 93A and97C).

[00397] W90 in APOBEC1 (corresponding to W285 in APOBEC3G) is predicted to form a hydrophobic pocket in the APOBEC3G active site and assist in substrate binding[13]. Mutating this residue to Ala abrogated APOBEC3G's catalytic activity[13]. In BE3, W90A almost completely abrogated base editing efficiency (Figure 97C). In contrast, it was found that W90Y only modestly decreased base editing activity while narrowing the editing window width at site A and site B to 3.8 and 4.9 nucleotides, respectively (Figure 93A). These results demonstrate that mutations to the cytidine deaminase domain can narrow the activity window width of the corresponding base editors.

[00398] W90Y, R126E, and R132E, the three mutations that narrowed the editing window without drastically reducing base editing activity, were combined into doubly and triply mutated base editors. The double mutant W90Y+R126E resulted in a base editor (YE1-BE3) with BE3-like maximal editing efficiencies, but substantially narrowed editing window width (width at site A and site B = 2.9 and 3.0 nucleotides, respectively (Figure 93A). The

W90Y+R132E base editor (YE2-BE3) exhibited modestly lower editing efficiencies (averaging 1.4-fold lower maximal editing yields across the five sites tested compared with BE3), and also substantially narrowed editing window width (width at site A and site B = 2.7 and 2.8 nucleotides, respectively) (Figure 97C). The R126E+R132E double mutant (EE-BE3) showed similar maximal editing efficiencies and editing window width as YE2-BE3 (Figure 97C). The triple mutant W90Y+R126E+R132E (YEE-BE3) exhibited 2.0-fold lower average maximal editing yields but very little editing beyond the C6 position and an editing window width of 2.1 and 1.4 nucleotides for site A and site B, respectively (Figure 97C). These data taken together indicate that mutations in the cytidine deaminase domain can strongly affect editing window widths, in some cases with minimal or only modest effects on editing efficiency.

[00399] The base editing outcomes of BE3, YE1-BE3, YE2-BE3, EE-BE3, and YEE-BE3 were further compared in HEK293T cells targeting four well-studied human genomic sites that contain multiple Cs within the BE3 activity window[1]. These target loci contained target Cs at positions 4 and 5 (HEK site 3), positions 4 and 6 (HEK site 2), positions 5 and 6 (EMX1), or positions 6, 7, 8, and 11 (FANCF). BE3 exhibited little (< 1.2-fold) preference for editing any Cs within the position 4-8 activity window. In contrast, YE1-BE3, exhibited a 1.3-fold preference for editing C5 over C4 (HEK site 3), 2.6-fold preference for C6 over C4 (HEK site 2), 2.0-fold preference for C5 over C6 (EMX1), and 1.5-fold preference for C6 over C7 (FANCF) (Figure 93B). YE2-BE3 and EE-BE3 exhibited somewhat greater positional specificity (narrower activity window) than YE1-BE3, averaging 2.4-fold preference for editing C5 over C4 (HEK site 3), 9.5-fold preference for C6 over C4 (HEK site 2), 2.9-fold preference for C5 over C6 (EMX1), and 2.6-fold preference for C7 over C6 (FANCF) (Figure 93B). YEE-BE3 showed the greatest positional selectivity, with a 2.9-fold preference for editing C5 over C4 (HEK site 3), 29.7-fold preference for C6 over C4 (HEK site 2), 7.9-fold preference for C5 over C6 (EMX1), and 7.9-fold preference for C7 over C6 (FANCF) (Figure 93B). The findings establish that mutant base editors can discriminate between adjacent Cs, even when both nucleotides are within the BE3 editing window.

[00400] The product distributions of these four mutants and BE3 were further analyzed by HTS to evaluate their apparent processivity. BE3 generated predominantly T4-T5 (HEK site 3), T4-T6 (HEK site 2), and T5-T6 (EMX1) products in treated HEK293T cells, resulting in, on average, 7.4-fold more products containing two Ts, than products containing a single T. In contrast, YE1-BE3, YE2-BE3, EE-BE3, and YEE-BE3 showed substantially higher preferences for singly edited C4-T5, C4-T6, and T5-C6 products (Figure 93C). YE1-BE3

yielded products with an average single-T to double-T product ratio of 1.4. YE2-BE3 and EE-BE3 yielded products with an average single-T to double-T product ratio of 4.3 and 5.1, respectively (Figure 93C). Consistent with the above results, the YEE-BE3 triple mutant favored single-T products by an average of 14.3-fold across the three genomic loci. (Figure 93C). For the target site in which only one C is within the target window (HEK site 4, at position C5), all four mutants exhibited comparable editing efficiencies as BE3 (Figure 98). These findings indicate that these BE3 mutants have decreased apparent processivity and can favor the conversion of only a single C at target sites containing multiple Cs within the BE3 editing window. These data also suggest a positional preference of C5 > C6 > C7 ≈ C4 for these mutant base editors, although this preference could differ depending on the target sequence.

[00401] The window-modulating mutations in APOBEC1 were applied to VQR-BE3, allowing selective base editing of substrates at sites targeted by NGA PAM (Figure 107A). However, when these mutations were applied to SaKKH-BE3, a linear decrease in base editing efficiency was observed without the improvement in substrate selectivity, suggesting a different kinetic equilibrium and substrate accessibility of this base editor than those of BE3 and its variants (Figure 107B).

[00402] The five base editors with altered PAM specificities described in this study together increase the number of disease-associated mutations in the ClinVar database that can in principle be corrected by base editing by 2.5-fold (Figures 94A and 94B). Similarly, the development of base editors with narrowed editing windows approximately doubles the fraction of ClinVar entries with a properly positioned NGG PAM that can be corrected by base editing without comparable modification of a non-target C (from 31% for BE3 to 59% for YEE-BE3) (Figures 94A and 94B).

[00403] In summary, the targeting scope of base editing was substantially expanded by developing base editors that use Cas9 variants with different PAM specificities, and by developing a collection of deaminase mutants with varying editing window widths. In theory, base editing should be possible using other programmable DNA-binding proteins (such as Cpf1[16]) that create a bubble of single-stranded DNA that can serve as a substrate for a single-strand-specific nucleotide deaminase enzyme.

Materials and Methods

[00404] *Cloning.* PCR was performed using Q5 Hot Start High-Fidelity DNA Polymerase (New England Biolabs). Plasmids for BE and sgRNA were constructed using USER cloning

(New England Biolabs), obtained from previously reported plasmids[1]. DNA vector amplification was carried out using NEB 10beta competent cells (New England Biolabs).

[00405] *Cell culture.* HEK293T (ATCC CRL-3216) were cultured in Dulbecco's Modified Eagle's Medium plus GlutaMax (ThermoFisher) supplemented with 10% (v/v) fetal bovine serum (FBS), at 37 °C with 5% CO_2. Immortalized rat astrocytes containing the ApoE4 isoform of the *APOE* gene (Taconic Biosciences) were maintained in Dulbecco's Modified Eagle's Medium plus GlutaMax (ThermoFisher Scientific) supplemented with 10% (v/v) fetal bovine serum (FBS) and 200 µg/mL Geneticin (ThermoFisher Scientific).

[00406] *Transfections.* HEK293T cells were seeded on 48-well collagen-coated BioCoat plates (Corning) and transfected at approximately 85% confluency. 750 ng of BE and 250 ng of sgRNA expression plasmids were transfected using 1.5 µl of Lipofectamine 2000 (ThermoFisher Scientific) per well according to the manufacturer's protocol.

[00407] *High-throughput DNA sequencing of genomic DNA samples.* Transfected cells were harvested after 3 d and the genomic DNA was isolated using the Agencourt DNAdvance Genomic DNA Isolation Kit (Beckman Coulter) according to the manufacturer's instructions. Genomic regions of interest were amplified by PCR with flanking HTS primer pairs listed in the Supplementary Sequences. PCR amplification was carried out with Phusion hot-start II DNA polymerase (ThermoFisher) according to the manufacturer's instructions. PCR products were purified using RapidTips (Diffinity Genomics). Secondary PCR was performed to attach sequencing adaptors. The products were gel-purified and quantified using the KAPA Library Quantification Kit-Illumina (KAPA Biosystems). Samples were sequenced on an Illumina MiSeq as previously described[1].

[00408] *Data analysis.* Nucleotide frequencies were assessed using a previously described MATLAB script [1]. Briefly, the reads were aligned to the reference sequence via the Smith-Waterman algorithm. Base calls with Q-scores below 30 were replaced with a placeholder nucleotide (N). This quality threshold results in nucleotide frequencies with an expected theoretical error rate of 1 in 1000.

[00409] Analyses of base editing processivity were performed using a custom python script. This program trims sequencing reads to the 20 nucleotide protospacer sequence as determined by a perfect match for the 7 nucleotide sequences that should flank the target site. These targets were then consolidated and sorted by abundance to assess the frequency of base editing products.

[00410] Bioinformatic analysis of the ClinVar database of human disease-associated mutations was performed in a manner similar to that previously described but with small

adjustments[1]. These adjustments enable the identification of targets with PAMs of customizable length and sequence. In addition, this improved script includes a priority ranking of target C positions (C5 > C6 > C7 >C8 ≈ C4), thus enabling the identification of target sites in which the on-target C is either the only cytosine within the window or is placed at a position with higher predicted editing efficiency than any off-target C within the editing window.

References for Example 12

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EXAMPLE 13:

[00411] Using improved transfection procedures and better plasmids, biological replicates (n=3) were used to install the four HF mutations into the Cas9 portion of BE3. The muations do not significantly effect on-targeting editing with plasmid delivery (Figure 99). At the tested concentration, BE3 protein delivery works; however, the on-target editing is lower than for plasmid delivery (Figure 100). Protein delivery of BE3 with the HF mutations installed reduces on-targeting ediing efficiency but still yields some edited cells (Figure 101).

[00412] Both lipofection and installing HF mutations were shown to decrease off-target deamination events. For the four sites shown in Figure 102, the off-target sitest (OT) with the highest GUIDE-Seq reads and deamination events were assayed (Komor *et al., Nature,* 2016). The specificity ratio was calculated by dividing the off-target editing by the on-target editing at the closest corresponding C. In cases where off-target editing was not detectable, the ratio was set to 100. Thus, a higher specificity ratio indicates a more specific construct. BE3 plasmid delivery showed much higher off-target/on-target editing than protein delivery of BE3, plasmid delivery of HF-BE3, or protein delivery of HF-BE3 (Figures 102 and 105).

[00413] Purified proteins HF-BE3 and BE3 were analyzed *in vitro* for their capabilities to convert C to T residues at different positions in the spacer with the most permissive motif. Both BE3 and HF-BE3 proteins were found to have the same "window" for base editing (Figures 103 and 104).

[00414] A list of the disease targets is given in Table 9. The base to be edited in Table 9 is indicated in bold and underlined.

Table 9. Base Editor Disease Targets

GENE	DISEASE	SPACER	PAM	EDITOR	DEFECT	CELL
RB1	RETINOBLASTOMA	AAT**C**TAGTAAATAAATTGATGT	AAAAGT	SAKKH-BE3	SPLICING IMPAIRMENT	J82
PTEN	CANCER	GACCAA**C**GGCTAAGTGAAGA	TGA	VQR-BE3	W111R	MC116
PIK3CA	CANCER	TC**C**TTTCTTCACGGTTGCCT	ACTGGT	SAKKH-BE3	K111R	CRL-5853
PIK3CA	CANCER	CTC**C**TGCTCAGTGATTTCAG	AGA	VQR-BE3	Q546R	CRL-2505
TP53	CANCER	TGT**C**ACACATGTAGTTGTAG	TGG	YEE-BE3	N239D	SNU475
HRAS	CANCER	CCTCC**C**GGCCGGCGGTATCC	AGG	YEE-BE3	Q61R	MC/CAR

Table 6. Exemplary diseases that may be treated using base editors. The protospacer and PAM sequences are shown in the sgRNA (PAM) column. The PAM sequence is shown in parentheses and with the base to be edited indicated by underlining.

Disease target	gene symbol	Base changed	sgRNA (PAM)	Base editor
Prion disease	PRNP	R37*	GGCAGCCGATACCCGGGGCA(GGG)	BE3
			GGGCAGCCGATACCCGGGGC(AGG)	
Pendred syndrome	Slc26a4	c.919-2A>G	TTATTGTCCGAAATAAAAGA(AGA)	BE3 (VQR SaCas9)
			ATTGTCCGAAATAAAAGAAG(AGG)	
			TTGTCCGAAATAAAAGAAGA(GGA)	
			GTCCGAAATAAAAGAAGAGGAAAA(AAT)	
			GTCCGAAATAAAAGAAGAGGAAAAA(ATT)	
Congenital deafness	Tmc1	c.545A>G	CAGGAAGCACGAGGCCACTG(AGG)	BE3 YE-BE3 YEE-BE3
			AACAGGAAGCACGAGGCCAC(TGA)	
			AGGAAGCACGAGGCCACTGA(GGA)	
Acquired deafness	SNHL	S33F	TTGGATTCTGGAATCCATTC(TGG)	BE3
Alzheimer's Disease	APP	A673T	TCTGCATCCATCTTCACTTC(AGA)	BE3 VQR
Niemann-Pick Disease Type C	NPC1	I1061T	CTTACAGCCAGTAATGTCAC(CGA)	BE3 VQR

[00415] Additional exemplary genes in the human genome that may be targeted by the base editors or complexes of this disclosure are provided herein in Tables 7 and 8. Table 7 includes gene mutations that may be correcteded by changing a cytosine (C) to a thymine (T), for example, using a BE3 nucleobase editor. Table 8 includes gene mutations that may be corrected by changing a guanine (G) to an adenine (A), for example, using a BE3 nucleobase editor.

Table 7. Human gene mutations that may be corrected by changing a cytosine (C) to a thymine (T). The gene name, gene symbol,and dbSNP database reference number (RS#) are indicated. Also indicated are exemplary protospacers with their PAM sequences (gRNAs and gRNAall) and the base to be edited, *e.g.*, a C, indicated by a "Y". The "gRNAs" sequences, from top to bottom, correspond to SEQ ID NOs: 1914-2091. The "gRNA all" sequences, from top to bottom, correspond to SEQ ID NOs: 2192-2540, 3144-3433.

Name	**RS# (dbSNP)**	**GeneSymbol**	**gRNAs**	**gRNAall**	**Phenotypes**
NM_000138.4(FBN1): c.3220T>C (p.Cys1074Arg)	137854465	FBN1	[]	[]	[]
NM_001927.3(DES):c. 1154T>C (p.Leu385Pro)	57955682	DES	[]	[]	['Myofibrillar myopathy 1', 'not provided']
NM_025152.2(NUBP L):c.815-27T>C	118161496	NUBPL	[]	['TGGTTCYAATGG ATGTCTGCTGG', 'GGTTCYAATGGA TGTCTGCTGGG']	['Mitochondrial complex I deficiency', 'not provided']
NM_003000.2(SDHB) :c.574T>C (p.Cys192Arg)	786202732	SDHB	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_004586.2(RPS6K A3):c.803T>C (p.Phe268Ser)	122454131	RPS6KA3	[]	[]	['Coffin-Lowry syndrome']

NM_005609.2(PYGM):c.425_528del	764313717	PYGM	[]	['TGGCTGYCAGGGACCCAGCAAGG', 'CTGYCAGGGACCCAGCAAGGAGG']	[]
NM_000124.3(ERCC6):c.2830-2A>G	373227647	ERCC6	[]	[]	['Cockayne syndrome, type B']
NM_000059.3(BRCA2):c.316+2T>C	81002805	BRCA2	['CTTAGGYAAGTAATGCAATATGG']	['CTTAGGYAAGTAATGCAATATGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2', 'Hereditary cancer-predisposing syndrome']
NM_003242.5(TGFBR2):c.923T>C (p.Leu308Pro)	28934568	TGFBR2	[]	['AGTTCCYGACGGCTGAGGAGCGG']	['Loeys-Dietz syndrome 2']
NM_000410.3(HFE):c.314T>C (p.Ile105Thr)	28934596	HFE	[]	[]	['Hemochromatosis type 1']
NM_000308.2(CTSA):c.247T>C (p.Trp83Arg)	28934603	CTSA	[]	[]	['Combined deficiency of sialidase AND beta galactosidase']
NM_033290.3(MID1):c.1877T>C (p.Leu626Pro)	28934611	MID1	[]	[]	['Opitz-Frias syndrome']
NM_000329.2(RPE65):c.1102T>C (p.Tyr368His)	62653011	RPE65	[]	[]	['Leber congenital amaurosis 2', 'Retinitis pigmentosa 20', 'not provided']
NM_007313.2(ABL1):c.814T>C (p.Tyr272His)	121913461	ABL1	[]	['CCAGYACGGGGAGGTGTACGAGG', 'CAGYACGGGGAGGTGTACGAGGG']	[]
NM_000546.5(TP53):c.398T>C (p.Met133Thr)	28934873	TP53	[]	[]	['Li-Fraumeni syndrome 1']
NM_000490.4(AVP):c.200T>C (p.Val67Ala)	28934878	AVP	[]	[]	['Neurohypophyseal diabetes insipidus']
NM_021961.5(TEAD1):c.1261T>C (p.Tyr?His)	11567847	TEAD1	['TCATATTYACAGGCTTGTAAAGG']	['TCATATTYACAGGCTTGTAAAGG']	[]
NM_002609.3(PDGFRB):c.1973T>C (p.Leu658Pro)	397509381	PDGFRB	[]	[]	['Basal ganglia calcification, idiopathic, 4']
NM_005236.2(ERCC4):c.689T>C (p.Leu230Pro)	397509402	ERCC4	[]	[]	['Fanconi anemia, complementation group Q']
NM_005236.2(ERCC4):c.706T>C (p.Cys236Arg)	397509403	ERCC4	[]	[]	['XERODERMA PIGMENTOSUM, TYPE F/COCKAYNE SYNDROME']
NM_173551.4(ANKS6):c.1322A>G (p.Gln441Arg)	377750405	ANKS6	[]	['AGGGCYGTCGGACCTTCGAGTGG', 'GGGCYGTCGGACCTTCGAGTGGG',	['Nephronophthisis 16']

				'GGCYGTCGGACC TTCGAGTGGGG']	
NM_000142.4(FGFR3):c.1612A>G (p.Ile538Val)	80053154	FGFR3	[]	[]	['Hypochondroplasia ']
NM_000441.1(SLC26 A4):c.707T>C (p.Leu236Pro)	80338848	SLC26A4	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_000518.4(HBB):c .337T>C (p.Cys113Arg)	35849199	HBB	[]	[]	[]
NM_000104.3(CYP1B 1):c.2T>C (p.Met1Thr)	72549389	CYP1B1	[]	[]	['Irido-corneo-trabecular dysgenesis']
NM_000169.2(GLA):c .484T>C (p.Trp162Arg)	28935196	-	[]	[]	['Fabry disease']
NM_001927.3(DES):c. 1034T>C (p.Leu345Pro)	57639980	DES	[]	['ATTCCCYGATGA GGCAGATGCGG', 'TTCCCYGATGAG GCAGATGCGGG']	['Myofibrillar myopathy 1', 'not provided']
NM_006517.4(SLC16 A2):c.1190T>C (p.Leu397Pro)	122455132	SLC16A2	[]	[]	['Allan-Herndon-Dudley syndrome']
NM_020320.3(RARS2):c.35A>G (p.Gln12Arg)	147391618	RARS2	[]	['ATACCYGGCAA GCAATAGCGCGG']	['Pontocerebellar hypoplasia type 6']
NM_000239.2(LYZ):c .221T>C (p.Ile74Thr)	121913547	LYZ	[]	[]	['Familial visceral amyloidosis, Ostertag type']
NM_002977.3(SCN9A):c.2215A>G (p.Ile739Val)	182650126	-	[]	['GTAAYTGCAAG ATCTACAAAAGG']	['Small fiber neuropathy', 'not provided']
NM_004700.3(KCNQ 4):c.842T>C (p.Leu281Ser)	80358278	KCNQ4	[]	['ACATYGACAAC CATCGGCTATGG']	['DFNA 2 Nonsyndromic Hearing Loss']
NM_000169.2(GLA):c .806T>C (p.Val269Ala)	28935488	-	['CAGTT AGYGA TTGGC AACTT TGG']	['CAGTTAGYGATT GGCAACTTTGG']	['Fabry disease']
NM_000228.2(LAMB 3):c.565-2A>G	370148688	LAMB3	[]	[]	['Junctional epidermolysis bullosa gravis of Herlitz']
NM_052867.2(NALC N):c.1526T>C (p.Leu509Ser)	786203987	NALCN	[]	[]	['CONGENITAL CONTRACTURES OF THE LIMBS AND FACE, HYPOTONIA, AND DEVELOPMENTA L DELAY']
NM_001031.4(RPS28) :c.1A>G (p.Met1Val)	786203997	RPS28	['CCAY GATGG CGGCG CGGCG GCGG']	['TGTCCAYGATGG CGGCGCGGCGG', 'CCAYGATGGCGG CGCGGCGGCGG']	['Diamond-Blackfan anemia with microtia and cleft palate']
NM_005957.4(MTHF	786204012	MTHFR	[]	['GACCYGCTGCCG	['Homocysteinemia

R):c.388T>C (p.Cys130Arg)				TCAGCGCCTGG']	due to MTHFR deficiency']
NM_005957.4(MTHFR):c.1530+2T>C	786204027	MTHFR	['GAAGGYGTGGTAGGGAGGCACGG', 'AAGGYGTGGTAGGGAGGCACGGG', 'AGGYGTGGTAGGGAGGCACGGGG']	['GAAGGYGTGGTAGGGAGGCACGG', 'AAGGYGTGGTAGGGAGGCACGGG', 'AGGYGTGGTAGGGAGGCACGGGG']	['Homocysteinemia due to MTHFR deficiency']
NM_005957.4(MTHFR):c.1793T>C (p.Leu598Pro)	786204034	MTHFR	[]	[]	['Homocysteinemia due to MTHFR deficiency']
NM_005957.4(MTHFR):c.1883T>C (p.Leu628Pro)	786204037	MTHFR	[]	['TCCCACYGGACAACTGCCTCTGG']	['Homocysteinemia due to MTHFR deficiency']
NM_000264.3(PTCH1):c.3168+2T>C	786204056	PTCH1	['ATCATTGYGAGTGTATTATAAGG', 'TCATTGYGAGTGTATTATAAGGG', 'CATTGYGAGTGTATTATAAGGGG']	['ATCATTGYGAGTGTATTATAAGG', 'TCATTGYGAGTGTATTATAAGGG', 'CATTGYGAGTGTATTATAAGGGG']	['Gorlin syndrome']
NM_000182.4(HADHA):c.919-2A>G	200017313	HADHA	[]	[]	['Mitochondrial trifunctional protein deficiency', 'Long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency', 'not provided']
NM_000030.2(AGXT):c.806T>C (p.Leu269Pro)	180177271	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_006121.3(KRT1):c.1436T>C (p.Ile479Thr)	57837128	KRT1	[]	[]	['Ichthyosis, cyclic, with epidermolytic hyperkeratosis', 'not provided']
NM_000521.3(HEXB):c.185C>T (p.Ser62Leu)	820878	HEXB	[]	[]	['Sandhoff disease, infantile type']
NM_000140.3(FECH):c.1137+3A>G	202147607	FECH	[]	['GTAGAYACCTTAGAGAACAATGG']	['Erythropoietic protoporphyria']
NM_015046.5(SETX):c.1166T>C	29001584	SETX	[]	[]	['Amyotrophic lateral sclerosis type

(p.Leu389Ser)					4']
NM_020365.4(EIF2B3):c.1037T>C (p.Ile346Thr)	119474039	EIF2B3	['CCAGAYTGTCAGCAAACACCTGG']	['CCAGAYTGTCAGCAAACACCTGG']	['Leukoencephalopathy with vanishing white matter']
NM_139058.2(ARX):c.98T>C (p.Leu33Pro)	28936077	ARX	[]	[]	['Mental retardation, with or without seizures, ARX-related, X-linked']
NM_005183.3(CACNA1F):c.2267T>C (p.Ile756Thr)	122456136	CACNA1F	[]	['TGCCAYTGCTGTGGACAACCTGG']	[]
NM_007374.2(SIX6):c.110T>C (p.Leu37Pro)	786204851	SIX6	[]	['GTCGCYGCCCGTGGCCCCTGCGG']	['Cataract, microphthalmia and nystagmus']
NM_000339.2(SLC12A3):c.1261T>C (p.Cys421Arg)	28936387	SLC12A3	[]	[]	['Familial hypokalemia-hypomagnesemia']
NM_003865.2(HESX1):c.77T>C (p.Ile26Thr)	28936416	HESX1	[]	[]	['Pituitary hormone deficiency, combined 5']
NM_022114.3(PRDM16):c.2660T>C (p.Leu887Pro)	202115331	PRDM16	[]	[]	['Dilated cardiomyopathy 1LL']
NM_001159287.1(TPI1):c.832T>C (p.Phe278Leu)	121964847	TPI1	[]	[]	['Triosephosphate isomerase deficiency']
NM_001692.3(ATP6V1B1):c.242T>C (p.Leu81Pro)	121964880	ATP6V1B1	[]	[]	[]
NM_000490.4(AVP):c.61T>C (p.Tyr21His)	121964893	AVP	[]	[]	['Neurohypophyseal diabetes insipidus']
NM_000027.3(AGA):c.916T>C (p.Cys306Arg)	121964906	AGA	['GTTATAYGTGCCAATGTGACTGG']	['GTTATAYGTGCCAATGTGACTGG']	['Aspartylglycosaminuria']
NM_000138.4(FBN1):c.1468+2T>C	794728167	FBN1	[]	['ATTGGYACGTGATCCATCCTAGG']	['Thoracic aortic aneurysms and aortic dissections']
NM_000027.3(AGA):c.214T>C (p.Ser72Pro)	121964909	AGA	[]	['GACGGCYCTGTAGGCTTTGGAGG']	['Aspartylglycosaminuria']
NM_004453.3(ETFDH):c.1001T>C (p.Leu334Pro)	377686388	ETFDH	[]	[]	['Glutaric aciduria, type 2']
NM_001385.2(DPYS):c.1078T>C (p.Trp360Arg)	121964924	DPYS	['CGTAATAYGGGAAAAAGGCGTGG', 'AATAYGGGAAAAAGGCGTGGTGG', 'ATAYGGGAAAAAGGCGTGGTGGG']	['CGTAATAYGGGAAAAAGGCGTGG', 'AATAYGGGAAAAAGGCGTGGTGG', 'ATAYGGGAAAAAGGCGTGGTGGG']	['Dihydropyrimidinase deficiency']

			GTGGTGGG']		
NM_004453.3(ETFDH):c.2T>C (p.Met1Thr)	121964953	ETFDH	[]	[]	['Glutaric acidemia IIC']
NM_000071.2(CBS):c.1616T>C (p.Leu539Ser)	121964968	CBS	[]	[]	['Homocystinuria, pyridoxine-responsive']
NM_000170.2(GLDC):c.2T>C (p.Met1Thr)	121964978	GLDC	[]	['CGGCCAYGCAGTCCTGTGCCAGG', 'GGCCAYGCAGTCCTGTGCCAGGG']	['Non-ketotic hyperglycinemia']
NM_000108.4(DLD):c.1178T>C (p.Ile393Thr)	121964991	DLD	[]	[]	['Maple syrup urine disease, type 3']
NM_014425.3(INVS):c.1478T>C (p.Leu493Ser)	121964995	INVS	[]	[]	['Infantile nephronophthisis']
NM_000398.6(CYB5R3):c.382T>C (p.Ser128Pro)	121965006	CYB5R3	[]	[]	['Methemoglobinemia type 2']
NM_000398.6(CYB5R3):c.446T>C (p.Leu149Pro)	121965008	CYB5R3	[]	['CTGCYGGTCTACCAGGGCAAAGG']	['METHEMOGLOBINEMIA, TYPE I']
NM_000398.6(CYB5R3):c.610T>C (p.Cys204Arg)	121965011	CYB5R3	[]	[]	['Methemoglobinemia type 2']
NM_000398.6(CYB5R3):c.218T>C (p.Leu73Pro)	121965013	CYB5R3	[]	[]	['METHEMOGLOBINEMIA, TYPE I']
NM_001103.3(ACTN2):c.683T>C (p.Met228Thr)	786205144	ACTN2	['CCTAAAAYGTTGGATGCTGAAGG']	['CCTAAAAYGTTGGATGCTGAAGG']	['Dilated cardiomyopathy 1AA']
NM_000548.3(TSC2):c.3106T>C (p.Ser1036Pro)	45517281	TSC2	[]	[]	['Tuberous sclerosis syndrome', 'Tuberous sclerosis 2']
NM_000203.4(IDUA):c.1469T>C (p.Leu490Pro)	121965027	IDUA	[]	[]	['Mucopolysaccharidosis, MPS-I-H/S', 'Hurler syndrome', 'not provided']
NM_001122764.1(PPOX):c.35T>C (p.Ile12Thr)	28936677	PPOX	[]	[]	['Variegate porphyria']
NM_000525.3(KCNJ11):c.440T>C (p.Leu147Pro)	28936678	KCNJ11	[]	[]	['Islet cell hyperplasia']
NM_001025107.2(ADAR):c.1883T>C (p.Leu628Pro)	28936680	ADAR	[]	[]	['Symmetrical dyschromatosis of extremities']
NM_001025107.2(ADAR):c.2609T>C (p.Phe870Ser)	28936681	ADAR	[]	[]	['Symmetrical dyschromatosis of extremities']
NM_000557.4(GDF5):c.1322T>C (p.Leu441Pro)	28936683	-	[]	[]	['Brachydactyly type A2', 'Fibular hypoplasia and complex brachydactyly']

NM_000274.3(OAT):c.163T>C (p.Tyr55His)	121965037	OAT	[]	[]	['Ornithine aminotransferase deficiency']
NM_000274.3(OAT):c.1205T>C (p.Leu402Pro)	121965043	OAT	[]	[]	['Ornithine aminotransferase deficiency']
NM_000223.3(KRT12):c.386T>C (p.Met129Thr)	28936695	KRT12	[]	[]	['Meesman corneal dystrophy', 'not provided']
NM_000128.3(F11):c.901T>C (p.Phe301Leu)	121965064	F11	[]	['TGATYTCTTGGGAGAAGAACTGG']	['Hereditary factor XI deficiency disease']
NM_000128.3(F11):c.166T>C (p.Cys56Arg)	121965069	F11	[]	[]	['Hereditary factor XI deficiency disease']
NM_000235.3(LIPA):c.599T>C (p.Leu200Pro)	121965086	LIPA	[]	[]	['Lysosomal acid lipase deficiency']
NM_001199.3(BMP1):c.*241T>C	786205217	BMP1	[]	[]	['Osteogenesis imperfecta type 13']
NM_004974.3(KCNA2):c.788T>C (p.Ile263Thr)	786205231	KCNA2	[]	[]	['EPILEPTIC ENCEPHALOPATHY, EARLY INFANTILE, 32']
NM_000548.3(TSC2):c.5150T>C (p.Leu1717Pro)	45517398	TSC2	[]	['GCCCYGCACGCAAATGTGAGTGG', 'CCCYGCACGCAAATGTGAGTGGG']	['Tuberous sclerosis syndrome', 'not provided']
NM_000212.2(ITGB3):c.176T>C (p.Leu59Pro)	5918	ITGB3	[]	[]	['Myocardial infarction', 'Posttransfusion purpura', 'Thrombocytopenia, neonatal alloimmune', 'Fracture, hip, susceptibility to']
m.9191T>C	386829069	MT-ATP6	[]	[]	['Leigh disease']
NM_000419.3(ITGA2B):c.1787T>C (p.Ile596Thr)	76811038	ITGA2B	[]	[]	['Glanzmann thrombasthenia']
NM_002294.2(LAMP2):c.864+2T>C	730880485	LAMP2	[]	[]	['Cardiomyopathy', 'Danon disease']
NM_000138.4(FBN1):c.7111T>C (p.Trp2371Arg)	794728264	FBN1	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_000531.5(OTC):c.143T>C (p.Phe48Ser)	72554315	OTC	[]	[]	['not provided']
NM_178454.4(DRAM2):c.79T>C (p.Tyr27His)	786205662	DRAM2	[]	[]	['Retinal dystrophy']
NM_000138.4(FBN1):c.7531T>C (p.Cys2511Arg)	794728272	FBN1	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_016218.2(POLK):c.609T>C (p.Asn203=)	786205684	POLK	[]	[]	['Malignant tumor of prostate']
NM_016218.2(POLK):c.*66T>C	786205688	POLK	[]	[]	['Malignant tumor of prostate']
NM_000354.5(SERPI	28937312	SERPINA7	[]	[]	[]

NA7):c.740T>C (p.Leu247Pro)					
NM_000531.5(OTC):c.284T>C (p.Leu95Ser)	72554346	OTC	['ACAAGATYGTCTACAGAAACAGG']	['ACAAGATYGTCTACAGAAACAGG']	['not provided']
NM_015662.2(IFT172):c.770T>C (p.Leu257Pro)	786205857	IFT172	[]	['TTGTGCYAGGAAGTTATGACAGG']	['RETINITIS PIGMENTOSA 71']
NM_000531.5(OTC):c.386+2T>C	72554359	OTC	[]	[]	['not provided']
NM_001135669.1(XPR1):c.434T>C (p.Leu145Pro)	786205901	XPR1	[]	[]	['BASAL GANGLIA CALCIFICATION, IDIOPATHIC, 6']
NM_001135669.1(XPR1):c.419T>C (p.Leu140Pro)	786205903	XPR1	[]	[]	['BASAL GANGLIA CALCIFICATION, IDIOPATHIC, 6']
NM_001135669.1(XPR1):c.653T>C (p.Leu218Ser)	786205904	XPR1	[]	['GCGTTYACGTGTCCCCCCTTTGG', 'CGTTYACGTGTCCCCCTTTGGG']	['BASAL GANGLIA CALCIFICATION, IDIOPATHIC, 6']
NM_181457.3(PAX3):c.268T>C (p.Tyr90His)	104893654	PAX3	[]	[]	['Klein-Waardenberg syndrome']
NM_001987.4(ETV6):c.1046T>C (p.Leu349Pro)	786205155	ETV6	[]	[]	['Thrombocytopenia', 'LEUKEMIA, ACUTE LYMPHOBLASTIC; ALL']
NM_000055.2(BCHE):c.1004T>C (p.Leu335Pro)	104893684	BCHE	[]	[]	['Deficiency of butyrylcholine esterase']
NM_000388.3(CASR):c.382T>C (p.Phe128Leu)	104893696	CASR	[]	[]	['Hypocalcemia, autosomal dominant 1']
NM_000388.3(CASR):c.1835T>C (p.Phe612Ser)	104893698	CASR	[]	[]	['Hypocalcemia, autosomal dominant 1']
NM_000388.3(CASR):c.2641T>C (p.Phe881Leu)	104893704	CASR	[]	['ACGCTYTCAAGGTGGCTGCCCGG', 'CGCTYTCAAGGTGGCTGCCCGGG']	['Hypercalciuric hypercalcemia']
NM_000388.3(CASR):c.374T>C (p.Leu125Pro)	104893708	CASR	[]	[]	['Hypocalcemia, autosomal dominant 1', 'Hypocalcemia, autosomal dominant 1, with bartter syndrome']
NM_000388.3(CASR):c.2362T>C (p.Phe788Leu)	104893711	CASR	[]	[]	['Hypocalcemia, autosomal dominant 1']
NM_000388.3(CASR):c.38T>C (p.Leu13Pro)	104893717	CASR	[]	[]	['Hypocalciuric hypercalcemia, familial, type 1']
NM_006580.3(CLDN16):c.500T>C	104893725	CLDN16	[]	[]	['Primary hypomagnesemia']

(p.Leu167Pro)					
NM_006580.3(CLDN16):c.434T>C (p.Leu145Pro)	104893731	CLDN16	[]	[]	['Primary hypomagnesemia']
NM_000041.3(APOE):c.388T>C (p.Cys130Arg)	429358	APOE	[]	[]	['Familial type 3 hyperlipoproteinemia']
NM_198159.2(MITF):c.1051T>C (p.Ser351Pro)	104893744	MITF	[]	[]	['Waardenburg syndrome type 2A']
NM_198159.2(MITF):c.1195T>C (p.Ser399Pro)	104893747	MITF	[]	['ACTTYCCCTTATTCCATCCACGG', 'CTTYCCCTTATTCCATCCACGGG']	['Waardenburg syndrome type 2A']
NM_001122757.2(POU1F1):c.655T>C (p.Trp219Arg)	104893758	POU1F1	[]	[]	['Pituitary hormone deficiency, combined 1']
NM_000539.3(RHO):c.133T>C (p.Phe45Leu)	104893770	RHO	[]	['CATGYTTCTGCTGATCGTGCTGG', 'ATGYTTCTGCTGATCGTGCTGGG']	['Retinitis pigmentosa 4']
NM_003106.3(SOX2):c.290T>C (p.Leu97Pro)	104893802	-	[]	[]	['Microphthalmia syndromic 3']
NM_024009.2(GJB3):c.101T>C (p.Leu34Pro)	28937583	GJB3	[]	[]	['Erythrokeratodermia variabilis']
NM_003907.2(EIF2B5):c.1882T>C (p.Trp628Arg)	28937596	EIF2B5	[]	['AGGCCYGGAGCCCTGTTTTTAGG']	['Leukoencephalopathy with vanishing white matter']
NM_000551.3(VHL):c.188T>C (p.Leu63Pro)	104893827	VHL	[]	[]	['Pheochromocytoma']
NM_000320.2(QDPR):c.106T>C (p.Trp36Arg)	104893865	QDPR	[]	[]	['Dihydropteridine reductase deficiency']
NM_001151.3(SLC25A4):c.293T>C (p.Leu98Pro)	104893876	SLC25A4	[]	['GCAGCYCTTCTTAGGGGGTGTGG']	['Autosomal dominant progressive external ophthalmoplegia with mitochondrial DNA deletions 2']
NM_006005.3(WFS1):c.2486T>C (p.Leu829Pro)	104893883	WFS1	[]	['ACCATCCYGGAGGGCCGCCTGGG']	['WFS1-Related Disorders']
NM_001018077.1(NR3C1):c.1712T>C (p.Val571Ala)	104893911	NR3C1	['AAGYGATTGCAGCAGTGAAATGG']	['AAGYGATTGCAGCAGTGAAATGG']	['Pseudohermaphroditism, female, with hypokalemia, due to glucocorticoid resistance']
NM_001018077.1(NR3C1):c.2318T>C (p.Leu773Pro)	104893912	NR3C1	[]	[]	['Glucocorticoid resistance, generalized']
NM_003122.4(SPINK1):c.2T>C (p.Met1Thr)	104893938	SPINK1	[]	[]	['Hereditary pancreatitis']
NM_000165.4(GJA1):c.52T>C (p.Ser18Pro)	104893962	GJA1	[]	['CTACYCAACTGCTGGAGGGAAGG']	['Oculodentodigital dysplasia']
NM_000416.2(IFNGR1):c.260T>C (p.Ile87Thr)	104893973	IFNGR1	['TGTAATAYTTCTGA	['TGTAATAYTTCTGATCATGTTGG']	['Disseminated atypical mycobacterial

			TCATGTTGG']		infection', 'Mycobacterium tuberculosis, susceptibility to']
NM_000434.3(NEU1):c.718T>C (p.Trp240Arg)	104893978	NEU1	[]	['GCCTCCYGGCGCTACGGAAGTGG', 'CCTCCYGGCGCTACGGAAGTGGG', 'CTCCYGGCGCTACGGAAGTGGGG']	['Sialidosis, type II']
NM_153704.5(TMEM67):c.755T>C (p.Met252Thr)	202149403	TMEM67	[]	[]	['Joubert syndrome 6']
NM_000162.3(GCK):c.391T>C (p.Ser131Pro)	104894010	GCK	[]	[]	['Maturity-onset diabetes of the young, type 2']
NM_004577.3(PSPH):c.155T>C (p.Met52Thr)	104894036	PSPH	[]	[]	['Deficiency of phosphoserine phosphatase']
NM_000193.3(SHH):c.995T>C (p.Val332Ala)	104894052	SHH	[]	[]	['Single upper central incisor']
NM_000282.3(PCCA):c.491T>C (p.Ile164Thr)	202247815	PCCA	[]	[]	['Propionic acidemia']
NM_002546.3(TNFRSF11B):c.349T>C (p.Phe117Leu)	104894092	TNFRSF11B	[]	['TAGAGYTCTGCTTGAAACATAGG']	['Hyperphosphatasemia with bone disease']
NM_000532.4(PCCB):c.1556T>C (p.Leu519Pro)	202247822	PCCB	[]	[]	['Propionic acidemia']
NM_006412.3(AGPAT2):c.683T>C (p.Leu228Pro)	104894100	AGPAT2	[]	[]	['Congenital generalized lipodystrophy type 1']
NM_000238.3(KCNH2):c.2366T>C (p.Ile789Thr)	794728388	KCNH2	[]	[]	['Cardiac arrhythmia']
NM_001243133.1(NLRP3):c.1058T>C (p.Leu353Pro)	28937896	NLRP3	[]	[]	['Familial cold urticaria']
NM_021020.3(LZTS1):c.85T>C (p.Ser29Pro)	28937897	LZTS1	[]	[]	[]
NM_000102.3(CYP17A1):c.316T>C (p.Ser106Pro)	104894135	CYP17A1	[]	['CATCGCGYCCAACAACCGTAAGG', 'ATCGCGYCCAACAACCGTAAGGG']	['Complete combined 17-alpha-hydroxylase/17,20-lyase deficiency']
NM_000102.3(CYP17A1):c.1216T>C (p.Trp406Arg)	104894143	CYP17A1	[]	[]	['Complete combined 17-alpha-hydroxylase/17,20-lyase deficiency']
NM_000102.3(CYP17A1):c.1358T>C (p.Phe453Ser)	104894151	CYP17A1	[]	['AGCTCTYCCTCATCATGGCCTGG']	['Combined partial 17-alpha-hydroxylase/17,20-lyase deficiency']
NM_005097.3(LGI1):c.136T>C (p.Cys46Arg)	104894166	LGI1	[]	[]	['Epilepsy, lateral temporal lobe, autosomal dominant']

NM_005097.3(LGI1):c.695T>C (p.Leu232Pro)	104894167	LGI1	[]	[]	['Epilepsy, lateral temporal lobe, autosomal dominant']
NM_000281.3(PCBD1):c.244T>C (p.Cys82Arg)	104894177	PCBD1	[]	[]	['Hyperphenylalaninemia, BH4-deficient, D']
NM_003476.4(CSRP3):c.131T>C (p.Leu44Pro)	104894205	CSRP3	[]	[]	['Familial hypertrophic cardiomyopathy 12', 'not specified']
NM_000315.2(PTH):c.67T>C (p.Ser23Pro)	104894272	PTH	[]	[]	['Hypoparathyroidism familial isolated']
NM_005055.4(RAPSN):c.848T>C (p.Leu283Pro)	104894293	RAPSN	[]	[]	['Myasthenic syndrome, congenital, associated with acetylcholine receptor deficiency', 'MYASTHENIC SYNDROME, CONGENITAL, 11, ASSOCIATED WITH ACETYLCHOLINE RECEPTOR DEFICIENCY']
NM_000518.4(HBB):c.344T>C (p.Leu115Pro)	36015961	HBB	[]	['TGTGTGCYGGCCCATCACTTTGG']	['Beta thalassemia intermedia']
NM_005055.4(RAPSN):c.41T>C (p.Leu14Pro)	104894300	RAPSN	[]	[]	['MYASTHENIC SYNDROME, CONGENITAL, 11, ASSOCIATED WITH ACETYLCHOLINE RECEPTOR DEFICIENCY']
NM_000531.5(OTC):c.2T>C (p.Met1Thr)	72552295	OTC	['AGAAGAYGCTGTTTAATCTGAGG']	['AGAAGAYGCTGTTTAATCTGAGG']	['not provided']
NM_020661.2(AICDA):c.238T>C (p.Trp80Arg)	104894320	AICDA	[]	[]	['Immunodeficiency with hyper IgM type 2']
NM_020661.2(AICDA):c.317T>C (p.Leu106Pro)	104894321	AICDA	[]	[]	['Immunodeficiency with hyper IgM type 2']
NM_020661.2(AICDA):c.452T>C (p.Phe151Ser)	104894327	AICDA	[]	[]	['Immunodeficiency with hyper IgM type 2']
NM_000486.5(AQP2):c.646T>C (p.Ser216Pro)	104894329	-	[]	[]	[]
NM_020638.2(FGF23):c.287T>C (p.Met96Thr)	104894343	FGF23	[]	[]	['Tumoral calcinosis, familial, hyperphosphatemic']
NM_021044.2(DHH):c.2T>C (p.Met1Thr)	104894346	DHH	[]	[]	['46,XY gonadal dysgenesis, partial,

					with minifascicular neuropathy']
NM_000217.2(KCNA1):c.1223T>C (p.Val408Ala)	104894352	KCNA1	[]	[]	['Episodic ataxia type 1']
NM_000432.3(MYL2):c.52T>C (p.Phe18Leu)	104894370	MYL2	[]	[]	['Familial hypertrophic cardiomyopathy 10']
NM_080911.2(UNG):c.752T>C (p.Phe251Ser)	104894380	UNG	[]	[]	['Immunodeficiency with hyper IgM type 5']
NM_000192.3(TBX5):c.161T>C (p.Ile54Thr)	104894384	TBX5	[]	[]	['Holt-Oram syndrome']
NM_175929.2(FGF14):c.449T>C (p.Phe150Ser)	104894393	FGF14	[]	[]	['Spinocerebellar ataxia 27']
NM_007262.4(PARK7):c.497T>C (p.Leu166Pro)	28938172	PARK7	[]	[]	['Parkinson disease 7']
NM_004004.5(GJB2):c.229T>C (p.Trp77Arg)	104894397	GJB2	[]	[]	['Deafness, autosomal recessive 1A', 'not provided']
NM_001130089.1(KARS):c.517T>C (p.Tyr173His)	397514745	KARS	['TTCYATGATCTTCGAGGAGAGGG']	['CTTCYATGATCTTCGAGGAGAGGG', 'TTCYATGATCTTCGAGGAGAGGG']	['Deafness, autosomal recessive 89']
NM_000161.2(GCH1):c.662T>C (p.Met221Thr)	104894434	GCH1	[]	[]	['Dystonia, dopa-responsive, with or without hyperphenylalaninemia, autosomal recessive']
NM_032409.2(PINK1):c.1040T>C (p.Leu347Pro)	28940285	-	[]	[]	['Parkinson disease 6, autosomal recessive early-onset']
NM_006177.3(NRL):c.479T>C (p.Leu160Pro)	104894463	NRL	[]	[]	['Retinal degeneration, autosomal recessive, clumped pigment type']
NM_152443.2(RDH12):c.523T>C (p.Ser175Pro)	104894472	RDH12	[]	['TCCYCGGTGGCTCACCACATTGG']	['Leber congenital amaurosis 13']
NM_002435.2(MPI):c.413T>C (p.Met138Thr)	104894495	MPI	[]	[]	['Congenital disorder of glycosylation type 1B']
NM_001159702.2(FHL1):c.457T>C (p.Cys153Arg)	122458144	FHL1	[]	[]	['Myopathy, reducing body, X-linked, childhood-onset']
NM_183235.2(RAB27A):c.389T>C (p.Leu130Pro)	104894498	RAB27A	[]	[]	['Griscelli syndrome type 2']
NM_001018005.1(TPM1):c.284T>C (p.Val95Ala)	104894504	TPM1	[]	[]	['Familial hypertrophic cardiomyopathy 3', 'Cardiomyopathy']

NM_000485.2(APRT):c.329T>C (p.Leu110Pro)	104894508	APRT	[]	[]	['Adenine phosphoribosyltransferase deficiency']
NM_000303.2(PMM2):c.131T>C (p.Val44Ala)	104894534	PMM2	[]	[]	['Carbohydrate-deficient glycoprotein syndrome type I']
NM_024006.5(VKORC1):c.134T>C (p.Val45Ala)	104894540	VKORC1	[]	[]	['Warfarin response']
NM_001614.3(ACTG1):c.1109T>C (p.Val370Ala)	104894547	ACTG1	[]	[]	['Deafness, autosomal dominant 20']
NM_001128085.1(ASPA):c.454T>C (p.Cys152Arg)	104894548	-	[]	[]	['Spongy degeneration of central nervous system']
NM_004870.3(MPDU1):c.356T>C (p.Leu119Pro)	104894587	MPDU1	[]	['TTCCYGGTCATGCACTACAGAGG']	['Congenital disorder of glycosylation type 1F']
NM_004870.3(MPDU1):c.2T>C (p.Met1Thr)	104894588	MPDU1	[]	['AATAYGGCGGCCGAGGCGGACGG']	['Congenital disorder of glycosylation type 1F']
NM_004870.3(MPDU1):c.221T>C (p.Leu74Ser)	104894589	MPDU1	[]	[]	['Congenital disorder of glycosylation type 1F']
NM_153006.2(NAGS):c.1289T>C (p.Leu430Pro)	104894605	-	[]	[]	['Hyperammonemia, type III']
NM_000304.3(PMP22):c.47T>C (p.Leu16Pro)	104894617	PMP22	[]	[]	['Charcot-Marie-Tooth disease, type IA']
NM_000304.3(PMP22):c.82T>C (p.Trp28Arg)	104894626	PMP22	[]	['TAGCAAYGGATCGTGGGCAATGG']	['Charcot-Marie-Tooth disease, type IE']
NM_018129.3(PNPO):c.784T>C (p.Ter262Gln)	104894631	PNPO	[]	['ACCTYAACTCTGGGACCTGCTGG']	["Pyridoxal 5'-phosphate-dependent epilepsy"]
NM_173477.4(USH1G):c.143T>C (p.Leu48Pro)	104894651	USH1G	[]	[]	['Usher syndrome, type 1G']
NM_000371.3(TTR):c.191T>C (p.Phe64Ser)	104894665	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis', 'AMYLOIDOSIS, LEPTOMENINGEAL, TRANSTHYRETIN-RELATED']
NM_024301.4(FKRP):c.899T>C (p.Val300Ala)	104894691	FKRP	[]	[]	['Limb-girdle muscular dystrophy-dystroglycanopathy, type C5']
NM_032551.4(KISS1R):c.305T>C (p.Leu102Pro)	104894703	KISS1R	[]	['GCCCTGCYGTACCCGCTGCCCGG', 'TGCYGTACCCGCTGCCCGGCTGG']	[]
NM_000660.5(TGFB1):c.673T>C	104894719	TGFB1	[]	[]	['Diaphyseal dysplasia']

(p.Cys225Arg)					
NM_000229.1(LCAT):c.524-22T>C	794726664	LCAT	[]	[]	['Fish-eye disease']
NM_003332.3(TYROBP):c.2T>C (p.Met1Thr)	104894732	TYROBP	[]	[]	['Polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy']
NM_000074.2(CD40LG):c.464T>C (p.Leu155Pro)	104894769	CD40LG	[]	[]	['Immunodeficiency with hyper IgM type 1']
NM_000495.4(COL4A5):c.438+2T>C	281874738	COL4A5	['TCCAGYAAGTTATAAAATTGGG']	['CTCCAGYAAGTTATAAAATTTGG', 'TCCAGYAAGTTATAAAATTTGGG']	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.4690T>C (p.Cys1564Arg)	281874745	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_178152.2(DCX):c.373T>C (p.Tyr125His)	104894781	DCX	[]	[]	['Lissencephaly, X-linked', 'Subcortical laminar heterotopia, X-linked']
NM_006579.2(EBP):c.53T>C (p.Leu18Pro)	104894795	EBP	[]	[]	['MEND SYNDROME']
NM_001097642.2(GJB1):c.397T>C (p.Trp133Arg)	104894813	GJB1	[]	[]	['X-linked hereditary motor and sensory neuropathy']
NM_001165963.1(SCN1A):c.2690T>C (p.Leu897Ser)	794726761	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000166.5(GJB1):c.407T>C (p.Val136Ala)	104894826	GJB1	[]	['ATGYCATCAGCGTGGTGTTCCGG']	['Dejerine-Sottas disease', 'X-linked hereditary motor and sensory neuropathy']
NM_001165963.1(SCN1A):c.769T>C (p.Cys257Arg)	794726771	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy', 'not provided']
NM_001165963.1(SCN1A):c.1033T>C (p.Cys345Arg)	794726782	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_001122606.1(LAMP2):c.961T>C (p.Trp321Arg)	104894859	LAMP2	[]	['CAGCTACYGGGATGCCCCCCTGG', 'AGCTACYGGGATGCCCCCCTGGG']	['Danon disease']
m.10237T>C	193302927	MT-ND3	[]	[]	['Leber optic atrophy']
NM_033290.3(MID1):c.884T>C (p.Leu295Pro)	104894866	MID1	[]	[]	['Opitz-Frias syndrome']
m.10663T>C	193302933	MT-ND4L	[]	[]	['Leber optic atrophy']
NM_001165963.1(SCN1A):c.4055T>C (p.Leu1352Pro)	794726821	-	['TTCYGGTTTGTCTTATATTCTGG']	['TTCYGGTTTGTCTTATATTCTGG']	['Severe myoclonic epilepsy in infancy']
NM_000475.4(NR0B1	104894907	NR0B1	[]	[]	['Congenital adrenal

):c.890T>C (p.Leu297Pro)					hypoplasia, X-linked']
NM_022567.2(NYX):c.302T>C (p.Ile101Thr)	104894911	NYX	[]	[]	['Congenital stationary night blindness, type 1A']
NM_000513.2(OPN1MW):c.607T>C (p.Cys203Arg)	104894914	OPN1MW	[]	[]	['Colorblindness, partial, deutan series', 'Cone monochromatism']
NM_006517.4(SLC16A2):c.1313T>C (p.Leu438Pro)	104894931	SLC16A2	[]	['TGAGCYGGTGGGCCCAATGCAGG']	['Allan-Herndon-Dudley syndrome']
NM_000330.3(RS1):c.38T>C (p.Leu13Pro)	104894935	RS1	[]	['TTACTTCYCTTTGGCTATGAAGG']	['Juvenile retinoschisis', 'not provided']
NM_000116.4(TAZ):c.352T>C (p.Cys118Arg)	104894937	TAZ	['AAGYGTGTGCCTGTGTGCCGAGG']	['AAGYGTGTGCCTGTGTGCCGAGG']	['3-Methylglutaconic aciduria type 2']
NM_006517.4(SLC16A2):c.1481T>C (p.Leu494Pro)	104894938	SLC16A2	[]	[]	['Allan-Herndon-Dudley syndrome']
NM_001109878.1(TBX22):c.641T>C (p.Leu214Pro)	104894946	TBX22	[]	[]	['Cleft palate with ankyloglossia']
NM_001011658.3(TRAPPC2):c.248T>C (p.Phe83Ser)	104894948	-	[]	[]	['Spondyloepiphysea l dysplasia tarda']
NM_003140.2(SRY):c.326T>C (p.Phe109Ser)	104894956	SRY	[]	[]	['46,XY sex reversal, type 1']
NM_003140.2(SRY):c.203T>C (p.Ile68Thr)	104894968	SRY	[]	[]	['46,XY sex reversal, type 1']
NM_201269.2(ZNF644):c.1759A>G (p.Ile587Val)	146936371	ZNF644	[]	[]	['Myopia 21, autosomal dominant']
NM_001004434.2(SLC30A2):c.161A>G (p.His54Arg)	587776926	SLC30A2	[]	[]	['Reduced zinc in breast milk']
NM_000492.3(CFTR):c.3469-20T>C	373002889	CFTR	[]	[]	['Cystic fibrosis']
NM_001848.2(COL6A1):c.957+2T>C	794727060	COL6A1	['ACAAGGYGAGCGTGGGCTGCTGG', 'CAAGGYGAGCGTGGGCTGCTGGG']	['ACAAGGYGAGCGTGGGCTGCTGG', 'CAAGGYGAGCGTGGGCTGCTGGG']	['Ullrich congenital muscular dystrophy', 'Bethlem myopathy']
m.4336T>C	41456348	MT-TQ	[]	[]	[]
NM_001065.3(TNFRSF1A):c.175T>C (p.Cys59Arg)	104895217	TNFRSF1A	[]	['TGCYGTACCAAGTGCCACAAAGG']	['TNF receptor-associated periodic fever syndrome (TRAPS)']
NM_003072.3(SMARCA4):c.3032T>C	281875229	SMARCA4	[]	[]	['Mental retardation, autosomal dominant

(p.Met1011Thr)					16', 'not provided']
NM_019885.3(CYP26B1):c.436T>C (p.Ser146Pro)	281875232	CYP26B1	[]	[]	['Radiohumeral fusions with other skeletal and craniofacial anomalies', 'not provided']
NM_000182.4(HADHA):c.180+3A>G	781222705	HADHA	[]	[]	['Mitochondrial trifunctional protein deficiency', 'Long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency', 'not provided']
NM_000208.2(INSR):c.1124-2A>G	587776819	INSR	[]	[]	['Pineal hyperplasia AND diabetes mellitus syndrome']
NM_006329.3(FBLN5):c.506T>C (p.Ile169Thr)	28939072	FBLN5	['GACAYTGATGAATGTCGCTATGG']	['GACAYTGATGAATGTCGCTATGG']	['Age-related macular degeneration 3']
NM_000431.3(MVK):c.803T>C (p.Ile268Thr)	104895304	MVK	['CTCAAYAGATGCCATCTCCCTGG']	['CTCAAYAGATGCCATCTCCCTGG']	['Hyperimmunoglobulin D with periodic fever', 'Mevalonic aciduria']
NM_024960.4(PANK2):c.437T>C (p.Met146Thr)	28939088	PANK2	[]	[]	['Hypoprebetalipoproteinemia, acanthocytosis, retinitis pigmentosa, and pallidal degeneration']
NM_005359.5(SMAD4):c.1499T>C (p.Ile500Thr)	281875321	SMAD4	[]	[]	['Myhre syndrome', 'not provided']
NM_003793.3(CTSF):c.692A>G (p.Tyr231Cys)	143889283	CTSF	[]	['CTCCAYACTGAGCTGTGCCACGG']	['Ceroid lipofuscinosis, neuronal, 13']
NM_001159702.2(FHL1):c.310T>C (p.Cys104Arg)	122459147	FHL1	[]	['GGGGYGCTTCAAGGCCATTGTGG']	['Myopathy, reducing body, X-linked, childhood-onset']
NM_001159702.2(FHL1):c.625T>C (p.Cys209Arg)	122459149	FHL1	[]	[]	['Emery-dreifuss muscular dystrophy 6']
NM_006214.3(PHYH):c.135-2A>G	201578674	PHYH	[]	[]	['Refsum disease, adult, 1']
NM_006329.3(FBLN5):c.679T>C (p.Ser227Pro)	28939370	FBLN5	[]	[]	['Autosomal recessive cutis laxa type IA']
NM_004329.2(BMPR1A):c.1409T>C (p.Met470Thr)	199476089	BMPR1A	[]	[]	['Juvenile polyposis syndrome']
NM_005154.4(USP8):c.2152T>C (p.Ser718Pro)	672601307	USP8	[]	[]	['Pituitary dependent hypercortisolism']
NM_020184.3(CNNM	74552543	CNNM4	[]	['AAGCTCCYGGA	['Cone-rod

4):c.971T>C (p.Leu324Pro)				CTTTTTTCTGGG']	dystrophy amelogenesis imperfecta']
NM_000734.3(CD247) :c.2T>C (p.Met1Thr)	672601318	CD247	[]	[]	['Immunodeficiency due to defect in cd3-zeta']
NM_016042.3(EXOS C3):c.712T>C (p.Trp238Arg)	672601332	EXOSC3	[]	[]	['Pontocerebellar hypoplasia, type 1b']
NC_012920.1:m.1448 4T>C	199476104	MT-ND6	[]	[]	['Leber optic atrophy', 'Leigh disease']
m.10158T>C	199476117	MT-ND3	[]	['AAAYCCACCCCT TACGAGTGCGG']	['Leigh disease', 'Leigh syndrome due to mitochondrial complex I deficiency', 'Mitochondrial complex I deficiency']
NM_020451.2(SEPN1):c.872+2T>C	794727808	SEPN1	[]	['TTCCGGYGAGTG GGCCACACTGG']	['Congenital myopathy with fiber type disproportion', 'Eichsfeld type congenital muscular dystrophy']
NM_005022.3(PFN1): c.350A>G (p.Glu117Gly)	140547520	PFN1	[]	['CACCTYCTTTGC CCATCAGCAGG']	['Amyotrophic lateral sclerosis 18']
NM_032551.4(KISS1 R):c.443T>C (p.Leu148Ser)	28939719	KISS1R	[]	[]	[]
NM_000084.4(CLCN5):c.674T>C (p.Leu225Pro)	273585645	CLCN5	[]	[]	['Dent disease 1']
NM_000030.2(AGXT) :c.605T>A (p.Ile202Asn)	536352238	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_000060.3(BTD):c .212T>C (p.Leu71Pro)	397514333	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c .248T>C (p.Leu83Ser)	397514347	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c .445T>C (p.Phe149Leu)	397514359	BTD	[]	['TCACCGCYTCAA TGACACAGAGG']	['Biotinidase deficiency']
NM_000060.3(BTD):c .743T>C (p.Ile248Thr)	397514382	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c .764T>C (p.Ile255Thr)	397514384	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c .833T>C (p.Leu278Pro)	397514389	BTD	[]	[]	['Biotinidase deficiency']
NM_000061.2(BTK):c .2T>C (p.Met1Thr)	128620186	BTK	['AGCT AYGGC CGCAG TGATT CTGG']	['AGCTAYGGCCG CAGTGATTCTGG']	['X-linked agammaglobulinemi a']
m.15572T>C	207459996	MT-CYB	[]	[]	['Familial colorectal cancer']

NM_000060.3(BTD):c.1096T>C (p.Ser366Pro)	397514399	BTD	[]	[]	['Biotinidase deficiency']
m.15197T>C	207460001	MT-CYB	[]	['CTAYCCGCCATCCCATACATTGG']	['Exercise intolerance']
m.14849T>C	207460004	MT-CYB	[]	[]	[]
NM_000060.3(BTD):c.1214T>C (p.Leu405Pro)	397514406	BTD	[]	['TTCACCCYGGTCCCTGTCTGGGG']	['Biotinidase deficiency']
NM_000060.3(BTD):c.1252T>C (p.Cys418Arg)	397514408	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c.1267T>C (p.Cys423Arg)	397514412	BTD	[]	[]	['Biotinidase deficiency']
NM_001128177.1(THRB):c.1336T>C (p.Cys446Arg)	121918703	THRB	[]	[]	['Thyroid hormone resistance, generalized, autosomal dominant']
NM_000060.3(BTD):c.1459T>C (p.Trp487Arg)	397514422	BTD	[]	[]	['Biotinidase deficiency']
NM_198056.2(SCN5A):c.3963+2T>C	397514447	SCN5A	[]	[]	['Progressive familial heart block type 1A']
NM_020461.3(TUBGCP6):c.2546A>G (p.Glu849Gly)	368449236	TUBGCP6	[]	[]	['Microcephaly with chorioretinopathy, autosomal recessive']
NM_006225.3(PLCD1):c.562T>C (p.Cys188Arg)	397514471	PLCD1	[]	[]	['Leukonychia totalis']
NM_001161581.1(POC1A):c.398T>C (p.Leu133Pro)	397514488	POC1A	['AGCYGTGGGACAAGAGCAGCCGG']	['AGCYGTGGGACAAGAGCAGCCGG']	['Short stature, onychodysplasia, facial dysmorphism, and hypotrichosis']
NM_005340.6(HINT1):c.250T>C (p.Cys84Arg)	397514489	HINT1	['CAAGAAAYGTGCTGCTGATCTGG', 'AAGAAAYGTGCTGCTGATCTGGG']	['CAAGAAAYGTGCTGCTGATCTGG', 'AAGAAAYGTGCTGCTGATCTGGG']	['Gamstorp-Wohlfart syndrome']
NM_000051.3(ATM):c.2T>C (p.Met1Thr)	786203606	ATM	[]	[]	['Ataxia-telangiectasia syndrome', 'Hereditary cancer-predisposing syndrome']
NM_004281.3(BAG3):c.1385T>C (p.Leu462Pro)	397514507	BAG3	[]	[]	['Dilated cardiomyopathy 1HH']
NM_183075.2(CYP2U1):c.784T>C	397514515	CYP2U1	[]	[]	['Spastic paraplegia 56, autosomal

(p.Cys262Arg)					recessive']
NM_006177.3(NRL):c.287T>C (p.Met96Thr)	397514516	NRL	[]	['GAGGCCAYGGAGCTGCTGCAGGG']	['Retinitis pigmentosa 27']
NM_000344.3(SMN1):c.388T>C (p.Tyr130His)	397514518	SMN1	['CACTGGAYATGGAAATAGAGAGG']	['CACTGGAYATGGAAATAGAGAGG']	['Kugelberg-Welander disease']
NM_152692.4(C1GALT1C1):c.577T>C (p.Ser193Pro)	397514537	C1GALT1C1	[]	[]	['Polyagglutinable erythrocyte syndrome']
NM_024531.4(SLC52A2):c.368T>C (p.Leu123Pro)	397514538	SLC52A2	[]	[]	['Brown-Vialetto-Van Laere syndrome 2']
NM_000138.4(FBN1):c.5746T>C (p.Cys1916Arg)	794728238	FBN1	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_000138.4(FBN1):c.6274T>C (p.Trp2092Arg)	794728246	FBN1	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_017802.3(DNAAF5):c.2384T>C (p.Leu795Pro)	397514561	DNAAF5	[]	[]	['Ciliary dyskinesia, primary, 18']
NM_206933.2(USH2A):c.12295-2A>G	151148854	USH2A	[]	[]	['Usher syndrome, type 2A']
NM_000531.5(OTC):c.134T>C (p.Leu45Pro)	72554312	OTC	[]	['CTCACTCYAAAAAACTTTACCGG']	['Ornithine carbamoyltransferase deficiency', 'not provided']
NM_178012.4(TUBB2B):c.350T>C (p.Leu117Pro)	397514569	TUBB2B	[]	['GGTCCYGGATGTGGTGAGGAAGG']	['Polymicrogyria, asymmetric']
NM_000431.3(MVK):c.764T>C (p.Leu255Pro)	397514570	MVK	[]	[]	['Porokeratosis, disseminated superficial actinic 1']
NM_000431.3(MVK):c.122T>C (p.Leu41Pro)	397514571	MVK	[]	['CGGCYTCAACCCCACAGCAATGG', 'GGCYTCAACCCCACAGCAATGGG']	['Porokeratosis, disseminated superficial actinic 1']
NM_000531.5(OTC):c.167T>C (p.Met56Thr)	72554320	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.188T>C (p.Leu63Pro)	72554324	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.227T>C (p.Leu76Ser)	72554328	OTC	[]	[]	['not provided']
NM_004055.4(CAPN5):c.731T>C (p.Leu244Pro)	397514602	CAPN5	[]	[]	['Vitreoretinopathy, neovascular inflammatory']
NM_133497.3(KCNV2):c.491T>C (p.Phe164Ser)	397514604	KCNV2	[]	[]	['Retinal cone dystrophy 3B']
NM_006567.3(FARS2):c.986T>C (p.Ile329Thr)	397514611	FARS2	[]	[]	['Combined oxidative phosphorylation deficiency 14']
NM_018344.5(SLC29A3):c.607T>C (p.Ser203Pro)	397514626	SLC29A3	['ACTGATAYCAGGTGAGAGC	['ACTGATAYCAGGTGAGAGCCAGG', 'CTGATAYCAGGT	['Histiocytosis-lymphadenopathy plus syndrome']

			CAGG']	GAGAGCCAGGG']	
NM_000108.4(DLD):c.140T>C (p.Ile47Thr)	397514651	DLD	['ACAGTTAYAGGTTCTGGTCCTGG', 'GTTAYAGGTTCTGGTCCTGGAGG']	['ACAGTTAYAGGTTCTGGTCCTGG', 'GTTAYAGGTTCTGGTCCTGGAGG']	['Maple syrup urine disease, type 3']
NM_020632.2(ATP6V0A4):c.1739T>C (p.Met580Thr)	3807153	ATP6V0A4	[]	[]	['Renal tubular acidosis, distal, autosomal recessive']
NM_000238.3(KCNH2):c.1945+6T>C	794728380	KCNH2	['CTGTGAGYGTGCCCAGGGGCGGG', 'TGAGYGTGCCCAGGGGCGGG']	['CTGTGAGYGTGCCCAGGGGCGGG', 'TGAGYGTGCCCAGGGGCGGGCGG']	['Cardiac arrhythmia']
NM_001033053.2(NLRP1):c.230T>C (p.Met77Thr)	397514692	NLRP1	[]	[]	['Corneal intraepithelial dyskeratosis and ectodermal dysplasia']
NM_000238.3(KCNH2):c.2396T>C (p.Leu799Pro)	794728390	KCNH2	[]	['GCCATCCYGGGTATGGGGTGGGG', 'CCATCCYGGGTATGGGGTGGGGG', 'CATCCYGGGTATGGGGTGGGGG']	['Cardiac arrhythmia']
NM_014845.5(FIG4):c.524T>C (p.Leu175Pro)	397514707	FIG4	[]	[]	['Yunis Varon syndrome']
NM_001199107.1(TBC1D24):c.686T>C (p.Phe229Ser)	397514713	TBC1D24	[]	['GGTCTYTGACGTCTTCCTGGTGG']	['Early infantile epileptic encephalopathy 16']
NM_080605.3(B3GALT6):c.193A>G (p.Ser65Gly)	397514719	B3GALT6	[]	['CGCYGGCCACCAGCACTGCCAGG']	['Spondyloepimetaphyseal dysplasia with joint laxity']
NM_004183.3(BEST1):c.253T>C (p.Tyr85His)	28940274	BEST1	[]	[]	['Vitelliform dystrophy', 'not provided']
NM_005689.2(ABCB6):c.1067T>C (p.Leu356Pro)	397514756	ABCB6	[]	[]	['Dyschromatosis universalis hereditaria 3']
NM_000551.3(VHL):c.488T>C (p.Leu163Pro)	28940297	VHL	[]	[]	[]
NM_000218.2(KCNQ1):c.1025T>C (p.Leu342Pro)	794728522	KCNQ1	[]	[]	['Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.1251+2T>C	794728528	KCNQ1	[]	[]	['Cardiac arrhythmia']

NM_000498.3(CYP11B2):c.1382T>C (p.Leu461Pro)	72554627	-	[]	[]	['Corticosterone methyloxidase type 1 deficiency']
NM_130799.2(MEN1):c.547T>C (p.Trp183Arg)	794728649	MEN1	[]	[]	['not provided']
NM_213653.3(HFE2):c.238T>C (p.Cys80Arg)	28940586	HFE2	[]	[]	['Hemochromatosis type 2A']
NM_198056.2(SCN5A):c.4299+6T>C	794728934	SCN5A	[]	[]	['not provided']
NM_000548.3(TSC2):c.1946+2T>C	397515247	TSC2	[]	[]	['Tuberous sclerosis syndrome']
NM_000256.3(MYBPC3):c.3796T>C (p.Cys1266Arg)	730880608	MYBPC3	[]	['GAGYGCCGCCTGGAGGTGCGAGG']	['Cardiomyopathy']
NM_016381.5(TREX1):c.530T>C (p.Val177Ala)	79993407	TREX1	[]	[]	['Aicardi Goutieres syndrome 1']
NM_001382.3(DPAGT1):c.503T>C (p.Leu168Pro)	397515329	DPAGT1	[]	['AATCCYGTACTATGTCTACATGG', 'ATCCYGTACTATGTCTACATGGG', 'TCCYGTACTATGTCTACATGGGG']	['Congenital disorder of glycosylation type 1J']
NM_000372.4(TYR):c.265T>C (p.Cys89Arg)	28940877	TYR	[]	[]	['Tyrosinase-negative oculocutaneous albinism', 'not provided']
NM_000375.2(UROS):c.-26-177T>C	397515348	UROS	[]	[]	['Congenital erythropoietic porphyria']
NM_015102.4(NPHP4):c.2972T>C (p.Phe991Ser)	28940891	NPHP4	[]	[]	['Nephronophthisis 4']
NM_020822.2(KCNT1):c.2386T>C (p.Tyr796His)	397515406	KCNT1	[]	[]	['Epilepsy, nocturnal frontal lobe, 5']
NM_000061.2(BTK):c.1516T>C (p.Cys506Arg)	128621200	BTK	[]	[]	['X-linked agammaglobulinemia']
NM_006383.3(CIB2):c.272T>C (p.Phe91Ser)	397515411	CIB2	[]	[]	['Deafness, autosomal recessive 48']
NM_000061.2(BTK):c.1741T>C (p.Trp581Arg)	128621205	BTK	['ACATTYGGGCTTTTGGTAAGTGG']	['ACATTYGGGCTTTTGGTAAGTGG']	['X-linked agammaglobulinemia']
NM_018127.6(ELAC2):c.460T>C (p.Phe154Leu)	397515465	ELAC2	[]	['ATAYTTTCTGGTCCATTGAAAGG']	['Combined oxidative phosphorylation deficiency 17']
NM_199355.2(ADAMTS18):c.605T>C (p.Leu202Pro)	397515468	ADAMTS18	[]	[]	['Microcornea, myopic chorioretinal atrophy, and telecanthus']
NM_023110.2(FGFR1	397515481	FGFR1	[]	[]	['Hartsfield

):c.494T>C (p.Leu165Ser)					syndrome']
NM_001059.2(TACR3):c.766T>C (p.Tyr256His)	397515483	TACR3	[]	[]	['Hypogonadotropic hypogonadism 11 with or without anosmia']
m.14325T>C	397515505	MT-ND6	[]	[]	['Leber optic atrophy']
NM_004333.4(BRAF):c.1783T>C (p.Phe595Leu)	794729219	BRAF	[]	[]	['Cardiofaciocutaneous syndrome']
NM_000370.3(TTPA):c.548T>C (p.Leu183Pro)	397515525	TTPA	[]	[]	['Ataxia with vitamin E deficiency']
NM_000375.2(UROS):c.139T>C (p.Ser47Pro)	397515527	UROS	[]	[]	['Congenital erythropoietic porphyria']
NM_001006657.1(WDR35):c.1592T>C (p.Leu531Pro)	397515533	WDR35	[]	[]	['Cranioectodermal dysplasia 2']
NM_004595.4(SMS):c.449T>C (p.Ile150Thr)	397515552	SMS	[]	[]	['Snyder Robinson syndrome']
NM_005211.3(CSF1R):c.2483T>C (p.Phe828Ser)	397515557	CSF1R	[]	['CATCTYTGACTGTGTCTACACGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_000026.2(ADSL):c.1339T>C (p.Ser447Pro)	777821034	ADSL	[]	[]	['not provided']
NM_194248.2(OTOF):c.3413T>C (p.Leu1138Pro)	397515599	OTOF	[]	['AGGTGCYGTTCTGGGGCCTACGG', 'GGTGCYGTTCTGGGGCCTACGGG']	['Deafness, autosomal recessive 9']
NM_002608.2(PDGFB):c.356T>C (p.Leu119Pro)	397515632	PDGFB	[]	[]	['Idiopathic basal ganglia calcification 5']
NM_000404.2(GLB1):c.152T>C (p.Ile51Thr)	72555390	GLB1	[]	[]	['Gangliosidosis GM1 type 3']
NM_000116.4(TAZ):c.310T>C (p.Phe104Leu)	397515741	TAZ	[]	[]	['3-Methylglutaconic aciduria type 2']
NM_000138.4(FBN1):c.2341T>C (p.Cys781Arg)	397515766	FBN1	[]	['GGACAAYGTAGAAATACTCCTGG']	['Marfan syndrome']
NM_000138.4(FBN1):c.4222T>C (p.Cys1408Arg)	397515802	FBN1	[]	[]	['Marfan syndrome']
NM_000112.3(SLC26A2):c.-26+2T>C	386833492	SLC26A2	['GAGAGGYGAGAAGAGGGAAGCGG']	['GAGAGGYGAGAAGAGGGAAGCGG']	['Diastrophic dysplasia']
NM_000256.3(MYBPC3):c.1351+2T>C	397515897	MYBPC3	['AAAGGYGGGCCTGGGACCTGAGG']	['AAAGGYGGGCCTGGGACCTGAGG']	['Familial hypertrophic cardiomyopathy 4', 'Cardiomyopathy']
NM_000045.3(ARG1):c.32T>C (p.Ile11Thr)	28941474	ARG1	[]	[]	['Arginase deficiency']
NM_004820.3(CYP7B	587777222	CYP7B1	[]	[]	['Spastic paraplegia',

1):c.889A>G (p.Thr297Ala)					'Spastic paraplegia 5A']
NM_017909.3(RMND 1):c.713A>G (p.Asn238Ser)	144972972	RMND1	[]	[]	['Combined oxidative phosphorylation deficiency 11']
NM_000314.6(PTEN): c.545T>C (p.Leu182Ser)	794729664	PTEN	[]	[]	['Macrocephaly/autism syndrome']
NM_000256.3(MYBP C3):c.821+2T>C	397516076	MYBPC3	['CACG YGAGT GGCCA TCCTC AGGG']	['GCACGYGAGTG GCCATCCTCAGG', 'CACGYGAGTGGC CATCCTCAGGG']	['Familial hypertrophic cardiomyopathy 4', 'not specified']
NM_000257.3(MYH7) :c.1370T>C (p.Ile457Thr)	397516103	MYH7	[]	[]	['Cardiomyopathy', 'not specified']
NM_000257.3(MYH7) :c.2093T>C (p.Val698Ala)	397516130	MYH7	[]	[]	['Familial hypertrophic cardiomyopathy 1', 'not specified']
NM_000257.3(MYH7) :c.2546T>C (p.Met849Thr)	397516156	MYH7	['GGAG AYGGC CTCCA TGAAG GAGG']	['GGAGAYGGCCT CCATGAAGGAGG']	['Primary familial hypertrophic cardiomyopathy', 'Cardiomyopathy']
NM_000271.4(NPC1): c.1133T>C (p.Val378Ala)	120074134	NPC1	[]	[]	['Niemann-Pick disease type C1']
NM_000520.4(HEXA) :c.538T>C (p.Tyr180His)	28941771	HEXA	[]	[]	[]
NM_024426.4(WT1):c .1351T>C (p.Phe451Leu)	28941777	WT1	[]	[]	['Diffuse mesangial sclerosis']
NM_024426.4(WT1):c .1378T>C (p.Phe460Leu)	28941779	WT1	[]	[]	['Frasier syndrome']
NM_000257.3(MYH7) :c.602T>C (p.Ile201Thr)	397516258	MYH7	[]	[]	['Dilated cardiomyopathy 1S', 'Cardiomyopathy']
NM_000257.3(MYH7) :c.788T>C (p.Ile263Thr)	397516269	MYH7	[]	[]	['Primary familial hypertrophic cardiomyopathy', 'Familial hypertrophic cardiomyopathy 1', 'Cardiomyopathy']
NM_001429.3(EP300) :c.3573T>A (p.Tyr1191Ter)	565779970	EP300	[]	['CTTAYTACAGTT ACCAGAACAGG']	['Rubinstein-Taybi syndrome 2']
NM_080605.3(B3GAL T6):c.1A>G (p.Met1Val)	786200938	B3GALT6	[]	['AGCTTCAYGGCG CCCGCGCCGGG', 'TCAYGGCGCCCG CGCCGGGCCGG']	['Spondyloepimetaphyseal dysplasia with joint laxity']
NM_032551.4(KISS1 R):c.937T>C (p.Tyr313His)	587777844	KISS1R	[]	[]	[]
NM_000257.3(MYH7)	369437262	MYH7	[]	[]	['Familial

:c.5326A>G (p.Ser1776Gly)					hypertrophic cardiomyopathy 1', 'Cardiomyopathy', 'not specified']
NM_000441.1(SLC26A4):c.164+2T>C	397516420	SLC26A4	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_000441.1(SLC26A4):c.765+2T>C	397516432	SLC26A4	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_000551.3(VHL):c.497T>C (p.Val166Ala)	397516445	VHL	[]	[]	['Von Hippel-Lindau syndrome', 'Hereditary cancer-predisposing syndrome']
NM_000256.3(MYBPC3):c.709T>C (p.Tyr237His)	730880624	MYBPC3	[]	[]	['Cardiomyopathy']
NM_000531.5(OTC):c.392T>C (p.Leu131Ser)	72556252	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.394T>C (p.Ser132Pro)	72556253	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.416T>C (p.Leu139Ser)	72556259	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.476T>C (p.Ile159Thr)	72556269	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.490T>C (p.Ser164Pro)	72556273	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.526T>C (p.Tyr176His)	72556282	OTC	['GGCTGATYACCTCACGCTCCAGG']	['GGCTGATYACCTCACGCTCCAGG', 'GATYACCTCACGCTCCAGGTTGG']	['not provided']
NM_000531.5(OTC):c.536T>C (p.Leu179Pro)	72556286	OTC	[]	[]	['not provided']
NM_000229.1(LCAT):c.698T>C (p.Leu233Pro)	28942087	LCAT	[]	['ATCTCTCYTGGGGCTCCCTGGGG', 'TCTCYTGGGGCTCCCTGGGGTGG']	['Norum disease']
NM_174936.3(PCSK9):c.646T>C (p.Phe216Leu)	28942112	PCSK9	[]	[]	['Hypercholesterolemia, autosomal dominant, 3']
NM_004572.3(PKP2):c.2386T>C (p.Cys796Arg)	794729098	PKP2	[]	[]	['not provided']
NM_000061.2(BTK):c.1223T>C (p.Leu408Pro)	128621198	BTK	['AGCYGGGGACTGGACAATTTGGG']	['GAGCYGGGGACTGGACAATTTGGG', 'AGCYGGGGACTGGACAATTTGGG']	['X-linked agammaglobulinemia']
NM_000061.2(BTK):c.1625T>C (p.Leu542Pro)	128621203	BTK	[]	['TCGGCCYGTCCAGGTGAGTGTGG']	['X-linked agammaglobulinemia with growth hormone

					deficiency']
NM_006383.3(CIB2):c.368T>C (p.Ile123Thr)	397515412	CIB2	[]	['CTTCAYCTGCAAGGAGGACCTGG']	['Deafness, autosomal recessive 48']
NM_001943.3(DSG2):c.523+2T>C	397516709	DSG2	[]	[]	['Arrhythmogenic right ventricular cardiomyopathy, type 10', 'Cardiomyopathy']
NM_032575.2(GLIS2):c.523T>C (p.Cys175Arg)	587777353	GLIS2	[]	[]	['Nephronophthisis 7']
NM_000492.3(CFTR):c.3230T>C (p.Leu1077Pro)	139304906	CFTR	[]	[]	['Cystic fibrosis']
NM_000492.3(CFTR):c.1853T>C (p.Ile618Thr)	139468767	CFTR	[]	[]	['Cystic fibrosis', 'not provided']
NM_002755.3(MAP2K1):c.388T>C (p.Tyr130His)	397516793	MAP2K1	[]	[]	['Cardiofaciocutaneous syndrome 3']
NM_000525.3(KCNJ11):c.755T>C (p.Val252Ala)	193929352	KCNJ11	[]	[]	['Permanent neonatal diabetes mellitus']
NM_000352.4(ABCC8):c.404T>C (p.Leu135Pro)	193929364	ABCC8	[]	['AAGCYGCTAATTGGTAGGTGAGG']	['Permanent neonatal diabetes mellitus']
NM_000071.2(CBS):c.833T>C (p.Ile278Thr)	5742905	CBS	['ATCAYTGGGGTGGATCCCGAAGG', 'TCAYTGGGGTGGATCCCGAAGGG']	['ATCAYTGGGGTGGATCCCGAAGG', 'TCAYTGGGGTGGATCCCGAAGGG']	['Homocystinuria due to CBS deficiency', 'Homocystinuria, pyridoxine-responsive', 'not provided']
NM_001038.5(SCNN1A):c.1477T>C (p.Trp493Arg)	5742912	SCNN1A	[]	[]	['Bronchiectasis with or without elevated sweat chloride 2', 'not specified']
NM_000030.2(AGXT):c.2T>C (p.Met1Thr)	138584408	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_005633.3(SOS1):c.1649T>C (p.Leu550Pro)	397517153	SOS1	[]	[]	['Noonan syndrome 4', 'Rasopathy']
NM_014714.3(IFT140):c.4078T>C (p.Cys1360Arg)	431905520	IFT140	['GCAGYGTGAGCTGCTCCTGGGAGG']	['CAAGCAGYGTGAGCTGCTCCTGG', 'GCAGYGTGAGCTGCTCCTGGAGG']	['Renal dysplasia, retinal pigmentary dystrophy, cerebellar ataxia and skeletal dysplasia']
NM_022168.3(IFIH1):c.1009A>G (p.Arg337Gly)	587777447	IFIH1	[]	[]	['Aicardi-goutieres syndrome 7']
NG_012123.1:g.2493A>G	1024611	CCL2	[]	[]	['Coronary artery disease, modifier of', 'Coronary artery disease,

					development of, in hiv', 'Mycobacterium tuberculosis, susceptibility to']
m.3394T>C	41460449	MT-ND1	['GGCYATATACAACTACGCAAAGG']	['GGCYATATACAACTACGCAAAGG']	['Leber optic atrophy']
NM_001127328.2(ACADM):c.997A>G (p.Lys333Glu)	77931234	ACADM	[]	[]	['Medium-chain acyl-coenzyme A dehydrogenase deficiency', 'not provided']
NM_005859.4(PURA):c.299T>C (p.Leu100Pro)	587782995	PURA	[]	[]	['Neonatal hypotonia', 'Intellectual disability', 'Seizures', 'Delayed speech and language development', 'Global developmental delay', 'Mental retardation, autosomal dominant 31']
NM_000368.4(TSC1):c.539T>C (p.Leu180Pro)	118203396	TSC1	[]	[]	['Tuberous sclerosis syndrome', 'Tuberous sclerosis 1']
NM_000256.3(MYBPC3):c.1696T>C (p.Cys566Arg)	730880695	MYBPC3	[]	[]	['Cardiomyopathy']
m.7275T>C	267606884	MT-CO1	[]	[]	['Familial colorectal cancer']
NM_000257.3(MYH7):c.1400T>C (p.Ile467Thr)	730880872	MYH7	[]	['TCGAGAYCTTCGATGTGAGTTGG', 'CGAGAYCTTCGATGTGAGTTGGG']	['Cardiomyopathy']
NM_002977.3(SCN9A):c.647T>C (p.Phe216Ser)	80356469	SCN9A	[]	[]	['Primary erythromelalgia']
NM_002977.3(SCN9A):c.2543T>C (p.Ile848Thr)	80356474	-	[]	['AAGATCAYTGGTAACTCAGTAGG', 'AGATCAYTGGTAACTCAGTAGGG', 'GATCAYTGGTAACTCAGTAGGGG']	['Primary erythromelalgia']
NM_001164277.1(SLC37A4):c.352T>C (p.Trp118Arg)	80356489	SLC37A4	[]	['GGGCYGGCCCCCATGTGGGAAGG']	['Glucose-6-phosphate transport defect', 'not provided']
NM_001457.3(FLNB):c.4804T>C (p.Ser1602Pro)	80356501	FLNB	[]	[]	[]
NM_152296.4(ATP1A3):c.2338T>C (p.Phe780Leu)	80356536	ATP1A3	[]	['GCCCYTCCTGCTGTTCATCATGG']	['Dystonia 12']

NM_206933.2(USH2A):c.5857+2T>C	397518022	-	[]	[]	['Usher syndrome, type 2A']
NM_194248.2(OTOF):c.1544T>C (p.Ile515Thr)	80356586	OTOF	[]	[]	['Deafness, autosomal recessive 9', 'Auditory neuropathy, autosomal recessive, 1']
NM_000335.4(SCN5A):c.3745T>C (p.Phe1249Leu)	45589741	SCN5A	[]	[]	['Acquired long QT syndrome']
NM_194248.2(OTOF):c.3032T>C (p.Leu1011Pro)	80356596	OTOF	[]	['GATGCYGGTGTTCGACAACCTGG']	['Deafness, autosomal recessive 9', 'Auditory neuropathy, autosomal recessive, 1']
NM_000525.3(KCNJ11):c.124T>C (p.Cys42Arg)	80356610	KCNJ11	[]	[]	['Permanent neonatal diabetes mellitus', 'Transient neonatal diabetes mellitus 3', 'MATURITY-ONSET DIABETES OF THE YOUNG, TYPE 13']
NM_000257.3(MYH7):c.2723T>C (p.Leu908Pro)	730880900	MYH7	[]	[]	['Cardiomyopathy']
NM_152296.4(ATP1A3):c.1112T>C (p.Leu371Pro)	606231433	ATP1A3	[]	[]	['Alternating hemiplegia of childhood 2']
NM_000083.2(CLCN1):c.857T>C (p.Val286Ala)	80356689	CLCN1	[]	['AGGAGYGCTATTTAGCATCGAGG']	['Myotonia congenita']
NM_000083.2(CLCN1):c.920T>C (p.Phe307Ser)	80356701	CLCN1	[]	[]	['Myotonia congenita']
NM_007375.3(TARDBP):c.*83T>C	80356744	TARDBP	[]	[]	['Amyotrophic lateral sclerosis type 10']
NM_152296.4(ATP1A3):c.1250T>C (p.Leu417Pro)	606231449	ATP1A3	[]	[]	['Dystonia 12']
NM_001876.3(CPT1A):c.1451T>C (p.Leu484Pro)	80356793	CPT1A	[]	[]	['Carnitine palmitoyltransferase I deficiency']
NM_000088.3(COL1A1):c.4391T>C (p.Leu1464Pro)	72656353	COL1A1	[]	[]	['Osteogenesis imperfecta type III']
NM_000089.3(COL1A2):c.279+2T>C	72656357	COL1A2	[]	[]	['Ehlers-Danlos syndrome, type 7B']
NM_015046.5(SETX):c.1807A>G (p.Asn603Asp)	116205032	SETX	[]	[]	['Spinocerebellar ataxia autosomal recessive 1']
m.4409T>C	118203884	MT-TM	[]	['AGGYCAGCTAAATAAGCTATCGG']	['Mitochondrial myopathy']
m.5874T>C	118203891	MT-TY	[]	[]	[]
NM_000130.4(F5):c.1160T>C (p.Ile387Thr)	118203911	F5	[]	[]	['Thrombophilia due to activated protein

					C resistance']
NM_173596.2(SLC39A5):c.911T>C (p.Met304Thr)	587777625	SLC39A5	[]	['AGAACAYGCTGGGGCTTTTGCGG']	['Myopia 24, autosomal dominant']
NM_024120.4(NDUFAF5):c.686T>C (p.Leu229Pro)	118203929	NDUFAF5	[]	[]	['Mitochondrial complex I deficiency']
NM_003159.2(CDKL5):c.602T>C (p.Leu201Pro)	587783087	CDKL5	[]	['ATTCYTGGGGAGCTTAGCGATGG']	['not provided']
NM_000046.3(ARSB):c.349T>C (p.Cys117Arg)	118203939	ARSB	[]	[]	['MUCOPOLYSACCHARIDOSIS, TYPE VI, SEVERE']
NM_000046.3(ARSB):c.707T>C (p.Leu236Pro)	118203940	ARSB	[]	[]	['MUCOPOLYSACCHARIDOSIS, TYPE VI, MILD']
NM_013319.2(UBIAD1):c.511T>C (p.Ser171Pro)	118203951	UBIAD1	[]	['TCTGGCYCCTTTCTCTACACAGG', 'GGCYCCTTTCTCTACACAGGAGG']	['Schnyder crystalline corneal dystrophy']
NM_138387.3(G6PC3):c.554T>C (p.Leu185Pro)	118203969	G6PC3	[]	[]	['Severe congenital neutropenia 4, autosomal recessive']
NM_006364.2(SEC23A):c.1144T>C (p.Phe382Leu)	118204000	SEC23A	[]	[]	['Craniolenticulosutural dysplasia']
NM_000429.2(MAT1A):c.914T>C (p.Leu305Pro)	118204004	MAT1A	[]	[]	['Methionine adenosyltransferase deficiency, autosomal recessive']
NM_000018.3(ACADVL):c.1372T>C (p.Phe458Leu)	118204017	ACADVL	[]	['TCGCATCYTCCGGATCTTTGAGG', 'CGCATCYTCCGGATCTTTGAGGG', 'GCATCYTCCGGATCTTTGAGGGG']	['Very long chain acyl-CoA dehydrogenase deficiency']
NM_000833.4(GRIN2A):c.2T>C (p.Met1Thr)	397518466	GRIN2A	[]	['CTAYGGGCAGAGTGGGCTATTGG']	['Focal epilepsy with speech disorder with or without mental retardation']
NM_015702.2(MMADHC):c.776T>C (p.Leu259Pro)	118204044	MMADHC	[]	[]	['Homocystinuria, cblD type, variant 1']
NM_018077.2(RBM28):c.1052T>C (p.Leu351Pro)	118204055	RBM28	[]	[]	['Alopecia, neurologic defects, and endocrinopathy syndrome']
NM_000237.2(LPL):c.662T>C (p.Ile221Thr)	118204061	LPL	[]	[]	['Hyperlipoproteinemia, type I']
NM_000237.2(LPL):c.337T>C (p.Trp113Arg)	118204069	LPL	[]	['GGACYGGCTGTCACGGGCTCAGG']	['Hyperlipoproteinemia, type I']
NM_000237.2(LPL):c.755T>C (p.Ile252Thr)	118204080	LPL	[]	['GTGAYTGCAGAGAGAGGACTTGG']	['Hyperlipoproteinemia, type I']
NM_000155.3(GALT)	111033726	GALT	[]	[]	['Deficiency of

:c.580T>C (p.Phe194Leu)					UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000190.3(HMBS) :c.739T>C (p.Cys247Arg)	118204111	HMBS	[]	['GCTTCGCYGCATCGCTGAAAGGG']	['Acute intermittent porphyria']
NM_000190.3(HMBS) :c.242T>C (p.Leu81Pro)	118204119	HMBS	[]	[]	['Acute intermittent porphyria']
NM_001363.4(DKC1): c.1193T>C (p.Leu398Pro)	199422253	DKC1	[]	[]	['Dyskeratosis congenita X-linked']
NM_004278.3(PIGL): c.500T>C (p.Leu167Pro)	145303331	PIGL	[]	[]	['Zunich neuroectodermal syndrome']
NM_000531.5(OTC):c .602T>C (p.Leu201Pro)	72558407	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c .779T>C (p.Leu260Ser)	72558441	OTC	['ATGTATYAATTACAGACACTTGG']	['ATGTATYAATTACAGACACTTGG']	['not provided']
NM_000531.5(OTC):c .803T>C (p.Met268Thr)	72558449	OTC	[]	[]	['not provided']
NM_000256.3(MYBPC3):c.1814A>G (p.Asp605Gly)	372371774	MYBPC3	[]	[]	['Primary dilated cardiomyopathy', 'Cardiomyopathy', 'not specified']
NM_000531.5(OTC):c .947T>C (p.Phe316Ser)	72558471	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c .1005+2T>C	72558484	OTC	['ATCATGGYAAGCAAGAAACAAGG']	['ATCATGGYAAGCAAGAAACAAGG']	['not provided']
NM_000531.5(OTC):c .1018T>C (p.Ser340Pro)	72558489	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c .1022T>C (p.Leu341Pro)	72558490	OTC	[]	[]	['not provided']
NM_007294.3(BRCA1):c.5291T>C (p.Leu1764Pro)	80357281	BRCA1	['GGCYAGAAATCTGTTGCTATGGG']	['GGGCYAGAAATCTGTTGCTATGGG', 'GGCYAGAAATCTGTTGCTATGGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1']
NM_000035.3(ALDOB):c.442T>C (p.Trp148Arg)	118204430	ALDOB	['GGAAGYGGCGTGCTGTGCTGAGG']	['GGAAGYGGCGTGCTGTGCTGAGG']	['Hereditary fructosuria']
NM_000512.4(GALNS):c.413T>C (p.Val138Ala)	118204436	GALNS	[]	[]	['Mucopolysaccharidosis, MPS-IV-A']
NM_024782.2(NHEJ1):c.367T>C	118204452	NHEJ1	[]	[]	['Severe combined immunodeficiency

(p.Cys123Arg)					with microcephaly, growth retardation, and sensitivity to ionizing radiation']
NM_007294.3(BRCA1):c.65T>C (p.Leu22Ser)	80357438	BRCA1	[]	['AAATCTYAGAGTGTCCCATCTGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
m.12297T>C	121434464	MT-TL2	['GTCYTAGGCCCCAAAAATTTGG']	['GTCYTAGGCCCCAAAAATTTTGG']	['Cardiomyopathy, mitochondrial']
NM_001040431.2(COA3):c.215A>G (p.Tyr72Cys)	139877390	COA3	[]	['CCAYCTGGGGAGGTAGGTTCAGG']	[]
m.10010T>C	121434476	MT-TG	[]	[]	['Exercise intolerance']
NM_000860.5(HPGD):c.577T>C (p.Ser193Pro)	121434481	HPGD	[]	[]	['Digital clubbing, isolated congenital']
NM_024915.3(GRHL2):c.1192T>C (p.Tyr398His)	587777737	GRHL2	[]	[]	['Ectodermal dysplasia/short stature syndrome']
NM_032374.4(APOPT1):c.353T>C (p.Phe118Ser)	587777786	APOPT1	[]	[]	['Cytochrome-c oxidase deficiency']
NM_001605.2(AARS):c.2251A>G (p.Arg751Gly)	143370729	AARS	[]	[]	['EPILEPTIC ENCEPHALOPATHY, EARLY INFANTILE, 29']
NM_000257.3(MYH7):c.2479T>C (p.Trp827Arg)	730880744	MYH7	[]	[]	['Cardiomyopathy']
NM_017415.2(KLHL3):c.1160T>C (p.Leu387Pro)	199469630	KLHL3			['Pseudohypoaldosteronism, type 2']
NM_017415.2(KLHL3):c.1280T>C (p.Met427Thr)	199469642	KLHL3			['Pseudohypoaldosteronism, type 2']
NM_005859.4(PURA):c.563T>C (p.Ile188Thr)	793888527	PURA	[]	['GACCAYTGCGCTGCCCGCGCAGG', 'ACCAYTGCGCTGCCCGCGCAGGG', 'CCAYTGCGCTGCCCGCGCAGGGG']	['not provided', 'Mental retardation, autosomal dominant 31']
NM_007294.3(BRCA1):c.212+2T>C	80358026	BRCA1	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1']
NM_002878.3(RAD51D):c.1A>G (p.Met1Val)	561425038	-	[]	['CGCCCAYGTTCCCCGCAGGCCGG']	['Hereditary cancer-predisposing syndrome']
NM_018960.4(GNMT):c.149T>C (p.Leu50Pro)	121907888	GNMT	[]	[]	['Glycine N-methyltransferase deficiency']

NM_007294.3(BRCA1):c.5074+2T>C	80358089	BRCA1	[]	[]	['Breast-ovarian cancer, familial 1']
NC_012920.1:m.11984T>C	200911567	MT-ND4	[]	[]	['Leigh disease']
NM_000280.4(PAX6):c.773T>C (p.Phe258Ser)	121907925	PAX6	[]	[]	['Congenital ocular coloboma', 'Coloboma of optic disc']
NM_020117.9(LARS):c.1118A>G (p.Tyr373Cys)	201861847	LARS	[]	[]	['Infantile liver failure syndrome 1']
NM_024105.3(ALG12):c.473T>C (p.Leu158Pro)	121907934	ALG12	[]	['TCCYGCTGGCCCTCGCGGCCTGG']	['Congenital disorder of glycosylation type 1G']
NM_000152.3(GAA):c.953T>C (p.Met318Thr)	121907936	GAA	[]	[]	['Glycogen storage disease type II, infantile']
NM_000520.4(HEXA):c.1453T>C (p.Trp485Arg)	121907968	HEXA	[]	[]	['Tay-Sachs disease']
NM_000520.4(HEXA):c.632T>C (p.Phe211Ser)	121907974	HEXA	[]	[]	['Tay-Sachs disease']
NM_000053.3(ATP7B):c.2123T>C (p.Leu708Pro)	121908000	ATP7B	[]	[]	['Wilson disease']
NM_000375.2(UROS):c.217T>C (p.Cys73Arg)	121908012	UROS	[]	[]	['Congenital erythropoietic porphyria']
NM_153212.2(GJB4):c.409T>C (p.Phe137Leu)	80358207	GJB4	[]	['CCTCATCYTCAAGGCCGCCGTGG']	['Erythrokeratodermia variabilis']
NM_000403.3(GALE):c.548T>C (p.Leu183Pro)	121908045	GALE	[]	[]	['UDPglucose-4-epimerase deficiency']
NM_002353.2(TACSTD2):c.557T>C (p.Leu186Pro)	80358228	TACSTD2	[]	['TCGGCYGCACCCCAAGTTCGTGG']	['Lattice corneal dystrophy Type III']
NM_001563.3(IMPG1):c.461T>C (p.Leu154Pro)	713993047	IMPG1	[]	[]	['Macular dystrophy, vitelliform, 4']
NM_138691.2(TMC1):c.1543T>C (p.Cys515Arg)	121908076	TMC1	[]	['AGGACCTYGCTGGGAAACAATGG', 'ACCTYGCTGGGAAACAATGGTGG', 'CCTYGCTGGGAAACAATGGTGGG']	['Deafness, autosomal recessive 7']
NM_000271.4(NPC1):c.3182T>C (p.Ile1061Thr)	80358259	NPC1	[]	[]	['Niemann-Pick disease type C1']
NM_006432.3(NPC2):c.295T>C (p.Cys99Arg)	80358264	NPC2	[]	[]	['Niemann-Pick disease type C2']
NM_017838.3(NHP2):c.415T>C (p.Tyr139His)	121908089	NHP2	[]	['GGAGGCTYACGATGAGTGCCTGG', 'GGCTYACGATGAGTGCCTGGAGG']	['Dyskeratosis congenita autosomal recessive 1', 'Dyskeratosis congenita, autosomal recessive

					2']
NM_005857.4(ZMPSTE24):c.1018T>C (p.Trp340Arg)	121908093	ZMPSTE24	[]	[]	['Mandibuloacral dysplasia with type B lipodystrophy', 'not provided']
NM_001195794.1(CLRN1):c.488T>C (p.Leu163Pro)	121908142	CLRN1	[]	[]	['Usher syndrome, type 3']
NM_057176.2(BSND):c.35T>C (p.Ile12Thr)	121908144	BSND	[]	[]	['Sensorineural deafness with mild renal dysfunction']
NM_001243133.1(NLRP3):c.1718T>C (p.Phe573Ser)	121908152	NLRP3	[]	[]	['Familial cold urticaria', 'Chronic infantile neurological, cutaneous and articular syndrome']
NM_001243133.1(NLRP3):c.926T>C (p.Phe309Ser)	121908154	NLRP3	[]	['GGTGCCTYTGACGAGCACATAGG']	['Familial cold urticaria', 'Chronic infantile neurological, cutaneous and articular syndrome']
NM_153741.1(DPM3):c.254T>C (p.Leu85Ser)	121908155	DPM3	[]	[]	['Congenital disorder of glycosylation type 1O']
NM_001033855.2(DCLRE1C):c.2T>C (p.Met1Thr)	121908158	DCLRE1C	[]	['GGCGCTAYGAGTTCTTTCGAGGG', 'GCGCTAYGAGTTCTTTCGAGGGG']	['Histiocytic medullary reticulosis']
NM_017696.2(MCM9):c.1732+2T>C	587777871	MCM9	[]	[]	['Premature ovarian failure 1', 'Ovarian dysgenesis 4']
NM_031433.3(MFRP):c.545T>C (p.Ile182Thr)	121908190	-	[]	[]	['Nanophthalmos 2']
NM_001127221.1(CACNA1A):c.2141T>C (p.Val714Ala)	121908213	CACNA1A	[]	[]	['Familial hemiplegic migraine type 1']
NM_001127221.1(CACNA1A):c.4469T>C (p.Phe1490Ser)	121908233	CACNA1A	[]	[]	['Episodic ataxia type 2']
NM_133459.3(CCBE1):c.520T>C (p.Cys174Arg)	121908254	CCBE1	[]	[]	['Hennekam lymphangiectasia-lymphedema syndrome']
NM_018129.3(PNPO):c.2T>C (p.Met1Thr)	796052870	PNPO	[]	['CCCCCAYGACGTGCTGGCTGCGG', 'CCCCAYGACGTGCTGGCTGCGGG', 'CCCAYGACGTGCTGGCTGCGGGG']	['not provided']
NM_014845.5(FIG4):c.122T>C (p.Ile41Thr)	121908287	FIG4	[]	[]	['Charcot-Marie-Tooth disease, type 4J', 'not provided']
NM_001005741.2(GBA):c.751T>C (p.Tyr251His)	121908300	GBA	['GCCAGAYACTTTGTGAAGT	['GCCAGAYACTTTGTGAAGTAAGG', 'CCAGAYACTTTGTGAAGTAAGGG']	['Gaucher disease, type 1']

			AAGG']		
NM_020427.2(SLURP1):c.43T>C (p.Trp15Arg)	121908318	SLURP1	[]	['GCAGCCYGGAGCATGGGCTGTGG']	['Acroerythrokeratoderma']
NM_020427.2(SLURP1):c.229T>C (p.Cys77Arg)	121908319	SLURP1	[]	[]	['Acroerythrokeratoderma']
NM_000787.3(DBH):c.339+2T>C	74853476	DBH	[]	[]	['Dopamine beta hydroxylase deficiency']
NM_017882.2(CLN6):c.200T>C (p.Leu67Pro)	154774633	CLN6	['AGCYGGTATTCCCTCTCGAGTGG']	['AGCYGGTATTCCCTCTCGAGTGG']	['Adult neuronal ceroid lipofuscinosis', 'not provided']
NM_022124.5(CDH23):c.5663T>C (p.Phe1888Ser)	121908352	CDH23	[]	['CTCACCTYCAACATCACTGCGGG']	['Deafness, autosomal recessive 12']
NM_054027.4(ANKH):c.143T>C (p.Met48Thr)	121908407	ANKH	['GTCGAGAYGCTGGCCAGCTACGG', 'TCGAGAYGCTGGCCAGCTACGGG']	['GTCGAGAYGCTGGCCAGCTACGG', 'TCGAGAYGCTGGCCAGCTACGGG']	['Chondrocalcinosis 2']
NM_004924.4(ACTN4):c.784T>C (p.Ser262Pro)	121908417	ACTN4	[]	[]	['Focal segmental glomerulosclerosis 1']
NM_014384.2(ACAD8):c.455T>C (p.Met152Thr)	121908418	ACAD8	[]	[]	['Deficiency of isobutyryl-CoA dehydrogenase']
NM_153717.2(EVC):c.919T>C (p.Ser307Pro)	121908426	EVC	[]	[]	['Chondroectodermal dysplasia', 'Curry-Hall syndrome']
NM_001040108.1(MLH3):c.3826T>C (p.Trp1276Arg)	121908439	MLH3	[]	[]	['Hereditary nonpolyposis colorectal cancer type 7']
NM_013339.3(ALG6):c.1432T>C (p.Ser478Pro)	121908444	ALG6	[]	[]	['Congenital disorder of glycosylation type 1C']
NM_003835.3(RGS9):c.895T>C (p.Trp299Arg)	121908449	RGS9	[]	[]	['Prolonged electroretinal response suppression']
NM_022336.3(EDAR):c.259T>C (p.Cys87Arg)	121908451	EDAR	[]	[]	['Autosomal recessive hypohidrotic ectodermal dysplasia syndrome']
NM_014270.4(SLC7A9):c.131T>C (p.Ile44Thr)	121908485	SLC7A9	[]	[]	['Cystinuria']
NM_000030.2(AGXT):c.613T>C (p.Ser205Pro)	121908520	AGXT	[]	['CCTGTACYCGGGCTCCCAGAAGG']	['Primary hyperoxaluria, type I']

NM_000030.2(AGXT):c.731T>C (p.Ile244Thr)	121908525	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_000026.2(ADSL):c.1312T>C (p.Ser438Pro)	119450940	ADSL	[]	[]	['Adenylosuccinate lyase deficiency']
NM_000026.2(ADSL):c.674T>C (p.Met225Thr)	119450945	ADSL	['AAGAYGGTGACAGAAAAGGCAGG']	['AAGAYGGTGACAGAAAAGGCAGG']	['Adenylosuccinate lyase deficiency']
NM_014985.3(CEP152):c.2000A>G (p.Lys667Arg)	200879436	CEP152	[]	[]	['Seckel syndrome 5', 'not specified']
NM_002755.3(MAP2K1):c.158T>C (p.Phe53Ser)	121908594	MAP2K1	[]	[]	['Cardiofaciocutaneous syndrome 3', 'Rasopathy']
NM_004820.3(CYP7B1):c.647T>C (p.Phe216Ser)	121908612	CYP7B1	[]	[]	['Spastic paraplegia 5A']
NM_004273.4(CHST3):c.776T>C (p.Leu259Pro)	121908616	CHST3	[]	[]	['Spondyloepiphyseal dysplasia with congenital joint dislocations']
NM_004273.4(CHST3):c.920T>C (p.Leu307Pro)	121908618	CHST3	[]	['CGTGCYGGCCTCGCGCATGGTGG']	['Spondyloepiphyseal dysplasia with congenital joint dislocations']
NM_004273.4(CHST3):c.857T>C (p.Leu286Pro)	121908620	CHST3	[]	[]	['Spondyloepiphyseal dysplasia with congenital joint dislocations']
NM_000050.4(ASS1):c.535T>C (p.Trp179Arg)	121908646	ASS1	[]	[]	[]
NM_030761.4(WNT4):c.35T>C (p.Leu12Pro)	121908653	WNT4	[]	[]	['Mullerian aplasia and hyperandrogenism']
NM_006432.3(NPC2):c.199T>C (p.Ser67Pro)	11694	NPC2	[]	['TATTCAGYCTAAAAGCAGCAAGG']	['Niemann-Pick disease type C2']
NM_003839.3(TNFRSF11A):c.523T>C (p.Cys175Arg)	121908656	TNFRSF11A	[]	[]	['Osteopetrosis autosomal recessive 7']
NM_000022.2(ADA):c.320T>C (p.Leu107Pro)	121908739	ADA	[]	['CCTGCYGGCCAACTCCAAAGTGG']	['Severe combined immunodeficiency due to ADA deficiency']
NM_000140.3(FECH):c.315-48T>C	2272783	FECH	[]	[]	['Erythropoietic protoporphyria']
NM_000059.3(BRCA2):c.7529T>C (p.Leu2510Pro)	80358979	BRCA2	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2', 'Fanconi anemia, complementation group D1']
NM_003722.4(TP63):c.1033T>C (p.Cys345Arg)	121908837	TP63	[]	[]	['Ectrodactyly, ectodermal dysplasia, and cleft

					lip/palate syndrome 3']
NM_003722.4(TP63):c.1646T>C (p.Ile549Thr)	121908845	TP63	[]	[]	['Hay-Wells syndrome of ectodermal dysplasia', 'Rapp-Hodgkin ectodermal dysplasia syndrome']
NM_000059.3(BRCA2):c.7958T>C (p.Leu2653Pro)	80359022	BRCA2	[]	['TGCYTCTTCAACTAAAATACAGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2']
NM_000369.2(TSHR):c.1891T>C (p.Phe631Leu)	121908861	TSHR	[]	[]	['Hyperthyroidism, nonautoimmune', 'Thyroid adenoma, hyperfunctioning']
NM_000369.2(TSHR):c.1358T>C (p.Met453Thr)	121908864	TSHR	[]	[]	['Hyperthyroidism, nonautoimmune']
NM_000369.2(TSHR):c.1526T>C (p.Val509Ala)	121908874	TSHR	[]	[]	['Hyperthyroidism, nonautoimmune']
NM_000369.2(TSHR):c.1798T>C (p.Cys600Arg)	121908884	TSHR	[]	[]	['Hypothyroidism, congenital, nongoitrous, 1']
NM_000369.2(TSHR):c.1400T>C (p.Leu467Pro)	121908885	TSHR	[]	[]	['Hypothyroidism, congenital, nongoitrous, 1']
NM_001457.3(FLNB):c.703T>C (p.Ser235Pro)	121908896	FLNB	[]	[]	['Boomerang dysplasia']
NM_003880.3(WISP3):c.232T>C (p.Cys78Arg)	121908902	WISP3	[]	['AAAATCYGTGCCAAGCAACCAGG', 'AAATCYGTGCCAAGCAACCAGGG', 'AATCYGTGCCAAGCAACCAGGGG']	['Progressive pseudorheumatoid dysplasia']
NM_003880.3(WISP3):c.1000T>C (p.Ser334Pro)	121908903	WISP3	[]	[]	['Progressive pseudorheumatoid dysplasia']
NM_002977.3(SCN9A):c.4382T>C (p.Ile1461Thr)	121908914	-	[]	[]	['Paroxysmal extreme pain disorder']
NM_004086.2(COCH):c.349T>C (p.Trp117Arg)	121908929	-	[]	[]	['Deafness, autosomal dominant 9']
NM_004086.2(COCH):c.1535T>C (p.Met512Thr)	121908934	-	['AGATAYGGCTTCTAAACCGAAGG']	['AGATAYGGCTTCTAAACCGAAGG']	['Deafness, autosomal dominant 9']
NM_006892.3(DNMT3B):c.808T>C (p.Ser270Pro)	121908947	DNMT3B	[]	['CAAGTTCYCCGAGGTGAGTCCGG', 'AAGTTCYCCGAGGTGAGTCCGGG', 'AGTTCYCCGAGGTGAGTCCGGGG']	['Centromeric instability of chromosomes 1,9 and 16 and immunodeficiency']
NM_000226.3(KRT9):c.503T>C	61157095	KRT9	[]	[]	['Epidermolytic palmoplantar

(p.Leu168Ser)					keratoderma', 'not provided']
NM_000051.3(ATM): c.8584+2T>C	730881326	-	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000492.3(CFTR): c.3857T>C (p.Phe1286Ser)	121909028	CFTR	[]	['AGCCTYTGGAGT GATACCACAGG']	['Cystic fibrosis']
NM_000492.3(CFTR): c.3763T>C (p.Ser1255Pro)	121909041	CFTR	[]	[]	['Cystic fibrosis']
NM_001040667.2(HSF4):c.341T>C (p.Leu114Pro)	121909048	HSF4	[]	[]	['Cataract, zonular']
NM_005025.4(SERPINI1):c.145T>C (p.Ser49Pro)	121909051	SERPINI1	[]	[]	['Familial encephalopathy with neuroserpin inclusion bodies']
NM_002700.2(POU4F3):c.668T>C (p.Leu223Pro)	121909057	POU4F3	[]	[]	['Deafness, autosomal dominant 15']
NM_003322.4(TULP1):c.1471T>C (p.Phe491Leu)	121909074	TULP1	[]	[]	['Retinitis pigmentosa 14']
NM_003322.4(TULP1):c.1145T>C (p.Phe382Ser)	121909076	TULP1	[]	[]	['Retinitis pigmentosa', 'Retinitis pigmentosa 14']
NM_003000.2(SDHB):c.487T>C (p.Ser163Pro)	33927012	SDHB	[]	[]	['Pheochromocytoma', 'Hereditary Paraganglioma-Pheochromocytoma Syndromes', 'Hereditary cancer-predisposing syndrome', 'Cowden syndrome 2', 'not specified', 'not provided']
NM_005603.4(ATP8B1):c.863T>C (p.Leu288Ser)	121909099	ATP8B1	[]	[]	['Progressive intrahepatic cholestasis']
NM_005603.4(ATP8B1):c.1982T>C (p.Ile661Thr)	121909100	ATP8B1	[]	[]	['Progressive intrahepatic cholestasis', 'Benign recurrent intrahepatic cholestasis 1']
NM_000483.4(APOC2):c.142T>C (p.Trp48Arg)	120074115	-	[]	[]	['Apolipoprotein C2 deficiency']
NM_000543.4(SMPD1):c.911T>C (p.Leu304Pro)	120074124	SMPD1	['CACYTGTGAGGAAGTTCCTGGGG']	['AGCACYTGTGAGGAAGTTCCTGG', 'GCACYTGTGAGGAAGTTCCTGGG', 'CACYTGTGAGGAAGTTCCTGGGG']	['Sphingomyelin/cholesterol lipidosis', 'Niemann-Pick disease, type A', 'Niemann-Pick disease, type B', 'not provided']
NM_000085.4(CLCNKB):c.1294T>C	121909135	CLCNKB	[]	['CTTTGTCYATGGTGAGTCTGGGG']	['Bartter syndrome type 3']

(p.Tyr432His)					
NM_001300.5(KLF6): c.346T>C (p.Ser116Pro)	121909139	KLF6	[]	[]	[]
m.12338T>C	201863060	MT-ND5	[]	[]	['Leber optic atrophy']
NM_001300.5(KLF6): c.190T>C (p.Trp64Arg)	121909142	KLF6	['TCTGYGGACCAAAATCATTCTGG']	['TCTGYGGACCAAAATCATTCTGG']	[]
NM_001300.5(KLF6): c.506T>C (p.Leu169Pro)	121909143	KLF6	[]	['GGAGCYGCCCTCGCCAGGGAAGG']	[]
NM_000271.4(NPC1): c.337T>C (p.Cys113Arg)	120074136	NPC1	[]	[]	['Niemann-Pick disease type C1']
NM_000019.3(ACAT1):c.935T>C (p.Ile312Thr)	120074146	ACAT1	['CAAGAAYAGTAGGTAAGGCCAGG']	['CAAGAAYAGTAGGTAAGGCCAGG']	['Deficiency of acetyl-CoA acetyltransferase']
NM_017890.4(VPS13B):c.8459T>C (p.Ile2820Thr)	120074155	VPS13B	[]	[]	['Cohen syndrome']
NM_017653.3(DYM): c.1624T>C (p.Cys542Arg)	120074165	DYM	[]	[]	['Smith McCort dysplasia']
NM_001089.2(ABCA3):c.302T>C (p.Leu101Pro)	121909182	ABCA3	[]	['GCACYTGTGATCAACATGCGAGG']	['Surfactant metabolism dysfunction, pulmonary, 3']
NM_001089.2(ABCA3):c.4658T>C (p.Leu1553Pro)	121909183	ABCA3	[]	[]	['Surfactant metabolism dysfunction, pulmonary, 3']
NM_001089.2(ABCA3):c.977T>C (p.Leu326Pro)	121909185	ABCA3	[]	[]	['Surfactant metabolism dysfunction, pulmonary, 3']
NM_000474.3(TWIST1):c.392T>C (p.Leu131Pro)	121909189	TWIST1	[]	[]	['Saethre-Chotzen syndrome']
NM_000503.5(EYA1): c.1459T>C (p.Ser487Pro)	121909200	EYA1	[]	['CACTCYCGCTCATTCACTCCCGG']	['Melnick-Fraser syndrome']
NM_000358.2(TGFBI):c.1619T>C (p.Phe540Ser)	121909214	TGFBI	[]	[]	['Lattice corneal dystrophy type 3A']
NM_000426.3(LAMA2):c.8282T>C (p.Ile2761Thr)	115650537	LAMA2	['TTGAYAGGGAGCAAGCAGTTCGG', 'TGAYAGGGAGCAAGCAGTTCGG']	['TTGAYAGGGAGCAAGCAGTTCGG', 'TGAYAGGGAGCAAGCAGTTCGGG']	['Merosin deficient congenital muscular dystrophy']

NM_000314.6(PTEN):c.209T>C (p.Leu70Pro)	121909226	PTEN	[]	[]	['Cowden syndrome 1', 'Hereditary cancer-predisposing syndrome']
NM_000314.6(PTEN):c.335T>C (p.Leu112Pro)	121909230	PTEN	[]	[]	['Lhermitte-Duclos disease']
NM_000314.6(PTEN):c.722T>C (p.Phe241Ser)	121909240	PTEN	[]	[]	['Macrocephaly/autism syndrome']
NM_004970.2(IGFALS):c.1618T>C (p.Cys540Arg)	121909247	IGFALS	[]	['GGACYGTGGCTGCCCTCTCAAGG']	['Acid-labile subunit deficiency']
NM_005570.3(LMAN1):c.2T>C (p.Met1Thr)	121909253	LMAN1	[]	['AGAYGGCGGGATCCAGGCAAAGG']	['Combined deficiency of factor V and factor VIII, 1']
NM_005055.4(RAPSN):c.416T>C (p.Phe139Ser)	121909256	RAPSN	[]	[]	['Pena-Shokeir syndrome type I']
NM_000391.3(TPP1):c.887-10A>G	755445790	TPP1	['TTTYTTTTTTTTTTTTTTGAGG']	['TTTYTTTTTTTTTTTTTTGAGG']	['Ceroid lipofuscinosis, neuronal, 2', 'not provided']
NM_006302.2(MOGS):c.1954T>C (p.Phe652Leu)	121909292	MOGS	[]	[]	['Congenital disorder of glycosylation type 2B']
NM_005379.3(MYO1A):c.2728T>C (p.Ser910Pro)	121909306	MYO1A	[]	[]	['Deafness, autosomal dominant 48']
NM_178151.2(DCX):c.128T>C (p.Leu43Ser)	587783521	DCX	[]	[]	['Heterotopia']
NM_001127221.1(CACNA1A):c.5126T>C (p.Ile1709Thr)	121909326	CACNA1A	[]	[]	['Spinocerebellar ataxia 6', 'Familial hemiplegic migraine type 1', 'Episodic ataxia type 2']
NM_001451.2(FOXF1):c.1138T>C (p.Ter380Arg)	121909337	FOXF1	['TGATGYGAGGCTGCCGCCGCAGG']	['TGATGYGAGGCTGCCGCCGCAGG']	['Alveolar capillary dysplasia with misalignment of pulmonary veins']
NM_000163.4(GHR):c.341T>C (p.Phe114Ser)	121909357	GHR	[]	[]	['Laron-type isolated somatotropin defect']
NM_000163.4(GHR):c.512T>C (p.Ile171Thr)	121909367	GHR	[]	[]	['Laron-type isolated somatotropin defect']
NM_000339.2(SLC12A3):c.1868T>C (p.Leu623Pro)	121909385	SLC12A3	[]	['CAACCYGGCCCTCAGCTACTCGG']	['Familial hypokalemia-hypomagnesemia']
NM_001174089.1(SLC4A11):c.2480T>C (p.Leu827Pro)	121909394	SLC4A11	[]	[]	['Corneal dystrophy and perceptive deafness']
NM_001174089.1(SLC4A11):c.589T>C (p.Ser197Pro)	121909395	SLC4A11	[]	[]	['Corneal dystrophy and perceptive deafness']
NM_000519.3(HBD):c	34975911	HBD	[]	[]	['delta Thalassemia']

.-127T>C					
NM_002427.3(MMP13):c.224T>C (p.Phe75Ser)	121909497	MMP13	[]	['TTCTYCGGCTTAGAGGTGACTGG']	['Spondyloepimetaphyseal dysplasia, Missouri type']
NM_002427.3(MMP13):c.221T>C (p.Phe74Ser)	121909498	MMP13	[]	[]	[]
NM_002427.3(MMP13):c.272T>C (p.Met91Thr)	121909499	MMP13	['GTCAYGAAAAAGCCAAGATGCGG', 'TCAYGAAAAAGCCAAGATGCGGG']	['GTCAYGAAAAAGCCAAGATGCGG', 'TCAYGAAAAAGCCAAGATGCGGG']	[]
NM_000751.2(CHRND):c.283T>C (p.Phe95Leu)	121909506	CHRND	[]	[]	['Lethal multiple pterygium syndrome']
NM_000751.2(CHRND):c.188T>C (p.Leu63Pro)	121909508	CHRND	[]	['AACCYCATCTCCCTGGTGAGAGG']	['MYASTHENIC SYNDROME, CONGENITAL, 3B, FAST-CHANNEL']
NM_001100.3(ACTA1):c.287T>C (p.Leu96Pro)	121909519	ACTA1	[]	['CGAGCYTCGCGTGGCTCCCGAGG']	['Nemaline myopathy 3']
NM_001100.3(ACTA1):c.668T>C (p.Leu223Pro)	121909530	ACTA1	[]	[]	['Congenital myopathy with fiber type disproportion']
NM_000488.3(SERPINC1):c.1141T>C (p.Ser381Pro)	121909565	SERPINC1	[]	[]	['Antithrombin III deficiency']
NM_000488.3(SERPINC1):c.442T>C (p.Ser148Pro)	121909569	SERPINC1	[]	[]	['Antithrombin III deficiency']
NM_000488.3(SERPINC1):c.667T>C (p.Ser223Pro)	121909572	SERPINC1	[]	['TGGGTGYCCAATAAGACCGAAGG']	['Antithrombin III deficiency']
NM_000488.3(SERPINC1):c.379T>C (p.Cys127Arg)	121909573	SERPINC1	[]	[]	['Antithrombin III deficiency']
NM_023110.2(FGFR1):c.899T>C (p.Ile300Thr)	121909633	FGFR1	[]	[]	['Interfrontal craniofaciosynostosis']
NM_023110.2(FGFR1):c.1141T>C (p.Cys381Arg)	121909634	FGFR1	[]	[]	['Osteoglophonic dysplasia']
NM_182925.4(FLT4):c.3131T>C (p.Leu1044Pro)	121909651	FLT4	[]	[]	['Hereditary lymphedema type I']
NM_182925.4(FLT4):c.3257T>C (p.Ile1086Thr)	121909655	FLT4	[]	[]	['Hereditary lymphedema type I']
NM_000145.3(FSHR):c.479T>C (p.Ile160Thr)	121909659	FSHR	[]	[]	['Ovarian dysgenesis 1']
NM_000145.3(FSHR):c.1634T>C	121909664	FSHR	[]	[]	['Ovarian hyperstimulation

(p.Ile545Thr)					syndrome']
NM_000821.6(GGCX):c.896T>C (p.Phe299Ser)	121909677	GGCX	[]	['TATGTYCTCCTACGTCATGCTGG']	['Pseudoxanthoma elasticum-like disorder with multiple coagulation factor deficiency']
NM_001018077.1(NR3C1):c.2209T>C (p.Phe737Leu)	121909727	NR3C1	[]	['CTATTGCYTCCAAACATTTTTGG']	['Glucocorticoid resistance, generalized']
NM_005271.3(GLUD1):c.1501T>C (p.Ser501Pro)	121909732	GLUD1	[]	[]	['Hyperinsulinism-hyperammonemia syndrome']
NM_004614.4(TK2):c.278A>G (p.Asn93Ser)	142291440	TK2	[]	[]	['Mitochondrial DNA depletion syndrome 2']
NM_032977.3(CASP10):c.440T>C (p.Met147Thr)	121909776	CASP10	[]	[]	['Neoplasm of stomach']
NM_000250.1(MPO):c.518A>G (p.Tyr173Cys)	78950939	MPO	['TGCGGYATTTGTCCTGCTCCGGG']	['GTGCGGYATTTGTCCTGCTCCGG', 'TGCGGYATTTGTCCTGCTCCGGG']	['Myeloperoxidase deficiency']
NM_001139.2(ALOX12B):c.1562A>G (p.Tyr521Cys)	199766569	ALOX12B	[]	[]	['Autosomal recessive congenital ichthyosis 2']
NM_022041.3(GAN):c.1268T>C (p.Ile423Thr)	119485091	GAN	['AAGAAAAYCTACGCCATGGGTGG']	['AAGAAAAYCTACGCCATGGGTGG', 'AAAAYCTACGCCATGGGTGGAGG']	['Giant axonal neuropathy']
NM_014009.3(FOXP3):c.970T>C (p.Phe324Leu)	122467173	FOXP3	['GACAGAGYTCCTCCACAACATGG']	['GACAGAGYTCCTCCACAACATGG']	['Insulin-dependent diabetes mellitus secretory diarrhea syndrome']
NM_014009.3(FOXP3):c.1099T>C (p.Phe367Leu)	122467175	FOXP3	[]	[]	['Insulin-dependent diabetes mellitus secretory diarrhea syndrome']
NM_004239.3(TRIP11):c.2102A>G (p.Asn701Ser)	139539448	TRIP11	[]	[]	['Achondrogenesis, type IA']
NM_001104.3(ACTN3):c.1729C>T (p.Arg577Ter)	1815739	ACTN3	[]	[]	['Sprinting performance', 'Actn3 deficiency']
NM_002693.2(POLG):c.2636A>G (p.Gln879Arg)	368587966	POLG	[]	[]	['not provided']
NM_000552.3(VWF):c.3178T>C (p.Cys1060Arg)	61748497	VWF	[]	[]	['von Willebrand disease type 2N', 'not provided']
NM_000552.3(VWF):c.3445T>C (p.Cys1149Arg)	61748511	VWF	[]	[]	['von Willebrand disease type 1', 'not provided']
NM_000184.2(HBG2):c.125T>C (p.Phe42Ser)	34878913	HBG2	['CAGAGGTYCTTTGACAGCT	['CAGAGGTYCTTTGACAGCTTTGG']	['Cyanosis, transient neonatal']

			TTGG']		
NM_000371.3(TTR):c.190T>C (p.Phe64Leu)	138065384	TTR	[]	[]	['Cardiomyopathy', 'not specified']
NM_000402.4(G6PD):c.1058T>C (p.Leu353Pro)	76723693	G6PD	[]	[]	['Glucose 6 phosphate dehydrogenase deficiency', 'Anemia, nonspherocytic hemolytic, due to G6PD deficiency', 'not provided']
NM_177405.2(CECR1):c.355A>G (p.Thr119Ala)	775440641	CECR1	[]	[]	['Idiopathic livedo reticularis with systemic involvement']
NM_000218.2(KCNQ1):c.401T>C (p.Leu134Pro)	199472685	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.625T>C (p.Ser209Pro)	199472705	KCNQ1			['Atrial fibrillation, familial, 3', 'Atrial fibrillation']
NM_000218.2(KCNQ1):c.752T>C (p.Leu251Pro)	199472716	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.824T>C (p.Phe275Ser)	199472729	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.832T>C (p.Tyr278His)	199472731	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.845T>C (p.Leu282Pro)	199472733	KCNQ1			['Congenital long QT syndrome']
NM_000257.3(MYH7):c.730T>C (p.Phe244Leu)	730880849	MYH7	[]	[]	['Cardiomyopathy']
NM_000218.2(KCNQ1):c.908T>C (p.Leu303Pro)	199472740	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.913T>C (p.Trp305Arg)	199472741	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.1045T>C (p.Ser349Pro)	199472764	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.1117T>C (p.Ser373Pro)	199472766	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.1165T>C (p.Ser389Pro)	199472772	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.1174T>C (p.Trp392Arg)	199472774	KCNQ1			['Congenital long QT syndrome']

NM_000218.2(KCNQ1):c.1541T>C (p.Ile514Thr)	199472786	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.1696T>C (p.Ser566Pro)	199472803	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.1805T>C (p.Leu602Pro)	199472818	-			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.608T>C (p.Leu203Pro)	199472823	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.65T>C (p.Phe22Ser)	199472826	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.86T>C (p.Phe29Ser)	199472831	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.89T>C (p.Ile30Thr)	199472832	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.160T>C (p.Tyr54His)	199472843	KCNH2			['Congenital long QT syndrome']
NM_001165963.1(SCN1A):c.662T>C (p.Leu221Pro)	796052961	SCN1A	[]	[]	['not provided']
NM_000238.3(KCNH2):c.287T>C (p.Ile96Thr)	199472853	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.322T>C (p.Cys108Arg)	199472859	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.872T>C (p.Met291Thr)	199472881	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1238T>C (p.Leu413Pro)	199472893	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1279T>C (p.Tyr427His)	199472898	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1387T>C (p.Phe463Leu)	199472904	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1655T>C (p.Leu552Ser)	199472918	KCNH2	[]	[]	['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000061.2(BTK):c.1955T>C (p.Leu652Pro)	128622212	BTK	[]	[]	['X-linked agammaglobulinemia']
NM_000238.3(KCNH2):c.1691T>C (p.Leu564Pro)	199472924	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1702T>C (p.Trp568Arg)	199472927	KCNH2			['Congenital long QT syndrome']
NM_000138.4(FBN1):	794728333	FBN1	[]	[]	['Thoracic aortic

c.5726T>C (p.Ile1909Thr)					aneurysms and aortic dissections']
NM_015884.3(MBTPS2):c.1424T>C (p.Phe475Ser)	122468179	MBTPS2	[]	[]	['IFAP syndrome with or without BRESHECK syndrome']
NM_000238.3(KCNH2):c.1985T>C (p.Ile662Thr)	199472980	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.2033T>C (p.Leu678Pro)	199472981	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.2078T>C (p.Leu693Pro)	199472983	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.2309T>C (p.Val770Ala)	199472994	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.3146T>C (p.Leu1049Pro)	199473026	KCNH2			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.278T>C (p.Phe93Ser)	199473052	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.544T>C (p.Cys182Arg)	199473066	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.689T>C (p.Ile230Thr)	199473073	SCN5A			['Cardiac conduction defect, nonspecific']
NM_000335.4(SCN5A):c.1187T>C (p.Val396Ala)	199473103	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.1190T>C (p.Ile397Thr)	199473105	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.2018T>C (p.Leu673Pro)	199473141	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.2516T>C (p.Leu839Pro)	199473164	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.2783T>C (p.Leu928Pro)	199473178	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.2804T>C (p.Leu935Pro)	199473179	SCN5A			['Brugada syndrome']
NM_198056.2(SCN5A):c.3010T>C (p.Cys1004Arg)	199473183	SCN5A	[]	[]	['Congenital long QT syndrome', 'not specified', 'not provided']
NM_000335.4(SCN5A):c.3679T>C (p.Tyr1227His)	199473205	SCN5A	[]	[]	['Brugada syndrome']
NM_000335.4(SCN5A):c.3713T>C (p.Leu1238Pro)	199473210	SCN5A			['Brugada syndrome']
NM_000492.3(CFTR):	139573311	CFTR	[]	['TTCACYTCTAAT	['Cystic fibrosis']

c.1400T>C (p.Leu467Pro)				GGTGATTATGG', 'TCACYTCTAATG GTGATTATGGG']	
NM_000335.4(SCN5A):c.3929T>C (p.Leu1310Pro)	199473219	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4027T>C (p.Phe1343Leu)	199473228	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4028T>C (p.Phe1343Ser)	199473229	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4034T>C (p.Leu1345Pro)	199473231	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4340T>C (p.Ile1447Thr)	199473251	SCN5A			['Brugada syndrome']
NM_198056.2(SCN5A):c.4493T>C (p.Met1498Thr)	199473263	SCN5A			['Congenital long QT syndrome', 'not provided']
NM_000335.4(SCN5A):c.4742T>C (p.Leu1581Pro)	199473275	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4778T>C (p.Phe1593Ser)	199473277	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.5179T>C (p.Cys1727Arg)	199473302	SCN5A			['Brugada syndrome']
NM_000219.5(KCNE1):c.158T>C (p.Phe53Ser)	199473355	KCNE1			['Congenital long QT syndrome']
NM_000219.5(KCNE1):c.259T>C (p.Trp87Arg)	199473361	KCNE1			['Congenital long QT syndrome']
NM_000891.2(KCNJ2):c.301T>C (p.Cys101Arg)	199473374	KCNJ2			['Ventricular tachycardia']
NM_000218.2(KCNQ 1):c.560T>C (p.Leu187Pro)	199473399	KCNQ1	[]	[]	['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ 1):c.572T>C (p.Leu191Pro)	199473401	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ 1):c.1052T>C (p.Phe351Ser)	199473402	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ 1):c.1058T>C (p.Leu353Pro)	199473403	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_000238.3(KCNH 2):c.202T>C	199473417	KCNH2			['Congenital long QT syndrome',

(p.Phe68Leu)					'Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.341T>C (p.Leu114Pro)	199473448	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.716T>C (p.Leu239Pro)	199473458	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.742T>C (p.Trp248Arg)	199473459	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.797T>C (p.Leu266Pro)	199473460	KCNQ1			['Long QT syndrome', 'Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.829T>C (p.Ser277Pro)	199473461	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.910T>C (p.Trp304Arg)	199473466	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.1550T>C (p.Ile517Thr)	199473478	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.1661T>C (p.Val554Ala)	199473481	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.2T>C (p.Met1Thr)	199473485	KCNQ1			['Congenital long QT syndrome', 'KCNQ1-related Jervell and Lange-Nielsen syndrome']
NM_000238.3(KCNH2):c.260T>C (p.Leu87Pro)	199473495	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1700T>C (p.Ile567Thr)	199473519	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1705T>C (p.Tyr569His)	199473520	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1816T>C (p.Ser606Pro)	199473523	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1889T>C (p.Val630Ala)	199473526	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1945T>C (p.Ser649Pro)	199473530	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.2452T>C (p.Ser818Pro)	199473537	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.2573T>C	199473539	KCNH2			['Congenital long QT syndrome']

(p.Ile858Thr)					
NM_000335.4(SCN5A):c.407T>C (p.Leu136Pro)	199473557	SCN5A			['Brugada syndrome']
NM_020166.4(MCCC1):c.1310T>C (p.Leu437Pro)	119103215	MCCC1	[]	[]	['3 Methylcrotonyl-CoA carboxylase 1 deficiency']
NM_198056.2(SCN5A):c.944T>C (p.Leu315Pro)	199473564	SCN5A			['Brugada syndrome', 'not provided']
NM_002972.3(SBF1):c.1249A>G (p.Met417Val)	587776986	SBF1	[]	[]	['Charcot-Marie-Tooth disease, type 4B3']
NM_000335.4(SCN5A):c.2551T>C (p.Phe851Leu)	199473586	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.2743T>C (p.Cys915Arg)	199473588	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4046T>C (p.Ile1349Thr)	199473607	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4453T>C (p.Phe1485Leu)	199473615	SCN5A			['Sudden infant death syndrome']
NM_000335.4(SCN5A):c.5111T>C (p.Phe1704Ser)	199473627	SCN5A			['Sudden infant death syndrome']
NM_000891.2(KCNJ2):c.650T>C (p.Leu217Pro)	199473656	KCNJ2			['Congenital long QT syndrome']
NM_000218.2(KCNQ1):c.550T>C (p.Tyr184His)	199473661	KCNQ1	['AGCAAGBACGTGGGCCTCTGGGG']	['CAGCAAGBACGTGGGCCTCTGGG', 'AGCAAGBACGTGGGCCTCTGGGG', 'GCAAGBACGTGGGCCTCTGGGGG']	['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.206T>C (p.Leu69Pro)	199473665	KCNH2			['Congenital long QT syndrome']
NM_001130823.1(DNMT1):c.1531T>C (p.Tyr511His)	199473692	DNMT1			['Hereditary sensory neuropathy type IE']
NM_031226.2(CYP19A1):c.743+2T>C	786205107	-	['CTGTGYAAGTAATACAACTTGG']	['CTGTGYAAGTAATACAACTTTGG']	['Aromatase deficiency']
NM_016373.3(WWOX):c.872T>C (p.Leu291Pro)	119487098	WWOX	[]	[]	[]
NM_001287223.1(SCN11A):c.1142T>C (p.Ile381Thr)	606231280	SCN11A	['TTCAYTGTGGTCATTTTCCTGG']	['CTTCAYTGTGGTCATTTTCCTGG', 'TTCAYTGTGGTCATTTTCCTGGG']	['Episodic pain syndrome, familial, 3']
NM_003640.3(IKBKAP):c.2204+6T>C	111033171	IKBKAP	[]	[]	['Familial dysautonomia', 'not provided']

NM_000238.3(KCNH2):c.1282T>C (p.Ser428Pro)	794728368	KCNH2	[]	[]	['Cardiac arrhythmia']
NM_000441.1(SLC26A4):c.1588T>C (p.Tyr530His)	111033254	SLC26A4	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_206933.2(USH2A):c.10561T>C (p.Trp3521Arg)	111033264	USH2A	[]	[]	['Usher syndrome, type 2A']
NM_206933.2(USH2A):c.1606T>C (p.Cys536Arg)	111033273	USH2A	['ATATAGAYGCCTCTGCTCCCAGG']	['ATATAGAYGCCTCTGCTCCCAGG']	['Usher syndrome, type 2A']
NM_001363.4(DKC1):c.1049T>C (p.Met350Thr)	121912300	DKC1	[]	[]	['Dyskeratosis congenita X-linked']
NM_000274.3(OAT):c.1180T>C (p.Cys394Arg)	121965054	OAT	[]	[]	['Ornithine aminotransferase deficiency']
NM_001302946.1(TRNT1):c.497T>C (p.Leu166Ser)	606231289	TRNT1	['ACTTYATTTGACTACTTTAATGG']	['ACTTYATTTGACTACTTTAATGG']	['Sideroblastic anemia with B-cell immunodeficiency, periodic fevers, and developmental delay']
NM_000454.4(SOD1):c.341T>C (p.Ile114Thr)	121912441	SOD1	[]	['CATCAYTGGCCGCACACTGGTGG']	['Amyotrophic lateral sclerosis type 1']
NM_000454.4(SOD1):c.434T>C (p.Leu145Ser)	121912446	SOD1	[]	['CGTTYGGCTTGTGGTGTAATTGG', 'GTTYGGCTTGTGGTGTAATTGGG']	['Amyotrophic lateral sclerosis type 1']
NM_000454.4(SOD1):c.455T>C (p.Ile152Thr)	121912449	SOD1	[]	[]	['Amyotrophic lateral sclerosis type 1']
NM_000213.3(ITGB4):c.467T>C (p.Leu156Pro)	121912461	ITGB4	[]	[]	['Epidermolysis bullosa with pyloric atresia']
NM_000213.3(ITGB4):c.1684T>C (p.Cys562Arg)	121912463	ITGB4	[]	['GGCCAGYGTGTGTGTGAGCCTGG']	['Epidermolysis bullosa with pyloric atresia']
NM_000213.3(ITGB4):c.112T>C (p.Cys38Arg)	121912465	ITGB4	[]	[]	['Epidermolysis bullosa with pyloric atresia']
NM_002198.2(IRF1):c.31T>C (p.Trp11Arg)	121912470	IRF1	[]	[]	['Non-small cell lung cancer']
NM_000424.3(KRT5):c.20T>C (p.Val7Ala)	121912474	KRT5	['TCAAGTGYGTCCTTCCGGAGCGG', 'CAAGTGYGTCCTTCCGGAGCGGG', 'AAGTGYGTCCTTCCGGAGCGGGGG', 'AGTGYGTCCTTCCGGAGCGGGGG']	['TCAAGTGYGTCCTTCCGGAGCGG', 'CAAGTGYGTCCTTCCGGAGCGGG', 'AAGTGYGTCCTTCCGGAGCGGGGG', 'AGTGYGTCCTTCCGGAGCGGGGG']	['Epidermolysis bullosa simplex, Koebner type']

			GAGCGGGG']		
NM_002292.3(LAMB2):c.961T>C (p.Cys321Arg)	121912492	LAMB2	[]	['CCTCAACYGCGAGCAGTGTCAGG']	['Nephrotic syndrome, type 5, with or without ocular abnormalities']
NM_170707.3(LMNA):c.1139T>C (p.Leu380Ser)	121912495	LMNA	['TCTYGGAGGGCGAGGAGGAGAGG']	['TCTYGGAGGGCGAGGAGGAGAGG']	['Congenital muscular dystrophy, LMNA-related', 'not provided']
NM_001399.4(EDA):c.2T>C (p.Met1Thr)	397516659	EDA	[]	['GGCCAYGGGCTACCCGGAGGTGG']	['Hypohidrotic X-linked ectodermal dysplasia']
NM_000493.3(COL10A1):c.1951T>C (p.Trp651Arg)	111033549	-	[]	[]	['Metaphyseal chondrodysplasia, Schmid type']
NM_000233.3(LHCGR):c.1193T>C (p.Met398Thr)	121912526	-	[]	[]	['Gonadotropin-independent familial sexual precocity']
NM_000233.3(LHCGR):c.391T>C (p.Cys131Arg)	121912527	-	[]	[]	['Leydig cell hypoplasia, partial']
NM_000493.3(COL10A1):c.1798T>C (p.Ser600Pro)	111033555	-	[]	[]	['Metaphyseal chondrodysplasia, Schmid type']
NM_000233.3(LHCGR):c.1103T>C (p.Leu368Pro)	121912533	-	[]	[]	['Gonadotropin-independent familial sexual precocity']
NM_000233.3(LHCGR):c.1627T>C (p.Cys543Arg)	121912537	-	[]	[]	['Leydig cell agenesis']
NM_000233.3(LHCGR):c.1505T>C (p.Leu502Pro)	121912538	-	[]	[]	['Leydig cell agenesis']
NM_000901.4(NR3C2):c.2771T>C (p.Leu924Pro)	121912563	NR3C2	[]	[]	['Pseudohypoaldosteronism type 1 autosomal dominant']
NM_021044.2(DHH):c.485T>C (p.Leu162Pro)	111033589	DHH	[]	['GTTGCYGGCGCGCCTCGCAGTGG']	['46,XY gonadal dysgenesis, complete, dhh-related']
NM_000901.4(NR3C2):c.2936T>C (p.Leu979Pro)	121912567	NR3C2	[]	[]	['Pseudohypoaldosteronism type 1 autosomal dominant']
NM_000517.4(HBA2):c.2T>C (p.Met1Thr)	111033603	HBA2	[]	[]	['alpha Thalassemia']
NM_000762.5(CYP2A6):c.670T>C (p.Ser224Pro)	111033610	-	[]	[]	['Tegafur response']
NM_000660.5(TGFB1):c.241T>C (p.Tyr81His)	111033611	TGFB1	[]	[]	['Diaphyseal dysplasia']
NM_001173464.1(KIF21A):c.1067T>C (p.Met356Thr)	121912588	KIF21A	[]	[]	['Fibrosis of extraocular muscles, congenital, 1']

NM_000206.2(IL2RG):c.343T>C (p.Cys115Arg)	111033622	IL2RG	[]	['TGGCYGTCAGTTGCAAAAAAAGG']	['X-linked severe combined immunodeficiency']
NM_001041.3(SI):c.1859T>C (p.Leu620Pro)	121912613	SI	[]	['ATGCYGGAGTTCAGTTTGTTTGG']	['Sucrase-isomaltase deficiency']
NM_016180.4(SLC45A2):c.1082T>C (p.Leu361Pro)	121912619	SLC45A2	[]	['GAGTTTCYCATCTACGAAAGAGG']	['Oculocutaneous albinism type 4', 'not provided']
NM_000552.3(VWF):c.4837T>C (p.Ser1613Pro)	61750581	VWF	[]	['CTGCCYCTGATGAGATCAAGAGG']	['von Willebrand disease, type 2a', 'not provided']
NM_000552.3(VWF):c.4883T>C (p.Ile1628Thr)	61750584	VWF	[]	[]	['von Willebrand disease, type 2a', 'not provided']
NM_000180.3(GUCY2D):c.1694T>C (p.Phe565Ser)	61749755	GUCY2D	[]	[]	['Leber congenital amaurosis 1', 'not provided']
NM_003235.4(TG):c.3733T>C (p.Cys1245Arg)	121912647	TG	[]	[]	['Iodotyrosyl coupling defect']
NM_000546.5(TP53):c.755T>C (p.Leu252Pro)	121912653	TP53	[]	['CATCCYCACCATCATCACACTGG']	['Li-Fraumeni syndrome 1']
NM_000155.3(GALT):c.350T>C (p.Phe117Ser)	111033679	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.374T>C (p.Val125Ala)	111033680	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.386T>C (p.Met129Thr)	111033683	GALT	[]	['AGGTCAYGTGCTTCCACCCCTGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000546.5(TP53):c.1031T>C (p.Leu344Pro)	121912662	TP53	[]	[]	['Li-Fraumeni syndrome 1']
NM_000155.3(GALT):c.416T>C (p.Leu139Pro)	111033687	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.452T>C (p.Val151Ala)	111033701	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.499T>C (p.Trp167Arg)	111033708	GALT	['CCCTYGGGTGCAGGTTTGTGAGG']	['CCCTYGGGTGCAGGTTTGTGAGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.507+2T>C	111033710	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.512T>C (p.Phe171Ser)	111033715	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']

NM_000341.3(SLC3A1):c.2033T>C (p.Leu678Pro)	121912693	-	[]	[]	['Cystinuria']
NM_000540.2(RYR1):c.9242T>C (p.Met3081Thr)	147012990	RYR1	[]	[]	['Minicore myopathy with external ophthalmoplegia']
NM_000155.3(GALT):c.584T>C (p.Leu195Pro)	111033728	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.650T>C (p.Leu217Pro)	111033741	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.687+2T>C	111033748	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000039.1(APOA1):c.220T>C (p.Trp74Arg)	121912726	-	[]	[]	['Familial visceral amyloidosis, Ostertag type']
NM_000155.3(GALT):c.677T>C (p.Leu226Pro)	111033752	GALT	[]	['CAGGAGCYACTCAGGAAGGTGGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000039.1(APOA1):c.593T>C (p.Leu198Ser)	121912729	APOA1	[]	['GCGCTYGGCCGCGCGCCTTGAGG']	['Familial visceral amyloidosis, Ostertag type']
NM_000155.3(GALT):c.745T>C (p.Trp249Arg)	111033757	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_001681.3(ATP2A2):c.1678T>C (p.Cys560Arg)	121912734	ATP2A2	[]	[]	['Keratosis follicularis']
NM_000041.3(APOE):c.137T>C (p.Leu46Pro)	769452	APOE	[]	['AACYGGCACTGGGTCGCTTTTGG']	[]
NM_000342.3(SLC4A1):c.2317T>C (p.Ser773Pro)	121912753	SLC4A1	[]	[]	['Renal tubular acidosis, distal, with normal red cell morphology']
NM_003002.3(SDHD):c.284T>C (p.Leu95Pro)	80338846	SDHD	[]	[]	['Hereditary Paraganglioma-Pheochromocytoma Syndromes']
NM_016124.4(RHD):c.329T>C (p.Leu110Pro)	121912762	RHD	[]	['ACACYGTTCAGGTATTGGGATGG']	[]
NM_003002.3(SDHD):c.416T>C (p.Leu139Pro)	80338847	SDHD	[]	[]	['Hereditary Paraganglioma-Pheochromocytoma Syndromes', 'Paragangliomas 1']
NM_001822.5(CHN1):c.427T>C (p.Tyr143His)	121912794	CHN1	[]	[]	['Duane syndrome type 2']
NM_000155.3(GALT):c.1138T>C	111033824	GALT	[]	['CGCCYGACCACGCCGACCACAGG'	['Deficiency of UDPglucose-

(p.Ter380Arg)				, 'GCCYGACCACGCCGACCACAGGG']	hexose-1-phosphate uridylyltransferase']
m.3271T>C	199474658	MT-TL1	[]	[]	['Juvenile myopathy, encephalopathy, lactic acidosis AND stroke']
NM_000155.3(GALT):c.980T>C (p.Leu327Pro)	111033832	GALT	[]	['TCCYGCGCTCTGCCACTGTCCGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
m.3290T>C	199474665	MT-TL1	[]	[]	['Sudden infant death syndrome']
NM_020549.4(CHAT):c.629T>C (p.Leu210Pro)	121912820	CHAT	[]	[]	['Familial infantile myasthenia']
NM_020549.4(CHAT):c.1007T>C (p.Ile336Thr)	121912823	CHAT	[]	[]	['Familial infantile myasthenia']
NM_000155.3(GALT):c.328+2T>C	111033849	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000267.3(NF1):c.1595T>C (p.Leu532Pro)	199474737	NF1	[]	[]	['Neurofibromatosis, type 1', 'not provided']
NM_000455.4(STK11):c.545T>C (p.Leu182Pro)	730881974	STK11	[]	['GGGAACCYGCTGCTCACCACCGG', 'AACCYGCTGCTCACCACCGGTGG']	['Hereditary cancer-predisposing syndrome']
NM_001042492.2(NF1):c.2288T>C (p.Leu763Pro)	199474762	NF1	[]	[]	['Hereditary cancer-predisposing syndrome', 'not provided']
NM_001042492.2(NF1):c.5858T>C (p.Leu1953Pro)	199474792	NF1	[]	[]	['Neurofibromatosis, type 1', 'not provided']
m.7512T>C	199474817	MT-TS1	[]	[]	['MERRF/MELAS overlap syndrome']
m.7510T>C	199474820	MT-TS1	[]	[]	['Deafness, nonsyndromic sensorineural, mitochondrial']
m.7511T>C	199474821	MT-TS1	[]	[]	['Deafness, nonsyndromic sensorineural, mitochondrial']
m.2991T>C	199474823	MT-RNR2	[]	[]	['Chloramphenicol resistance']
NM_201253.2(CRB1):c.3122T>C (p.Met1041Thr)	62635656	CRB1	[]	[]	['Retinitis pigmentosa 12', 'not provided']
m.7587T>C	199474825	MT-CO2	[]	[]	['Cytochrome-c oxidase deficiency']
NM_000400.3(ERCC2):c.1454T>C (p.Leu485Pro)	121913025	ERCC2	[]	[]	['Xeroderma pigmentosum, group D']
NM_000157.3(GBA):c.703T>C	1064644	GBA	[]	['GGGYCACTCAAGGGACAGCCCGG'	['Gaucher disease']

(p.Ser235Pro)				]	
NM_001113755.2(TYMP):c.854T>C (p.Leu285Pro)	121913042	TYMP	[]	[]	[]
NM_000122.1(ERCC3):c.296T>C (p.Phe99Ser)	121913045	ERCC3	[]	[]	['Xeroderma pigmentosum, complementation group b']
NM_000186.3(CFH):c.1606T>C (p.Cys536Arg)	121913052	CFH	[]	[]	['Factor H deficiency']
NM_138413.3(HOGA1):c.533T>C (p.Leu178Pro)	796052090	HOGA1	[]	['GGACCYGCCTGTGGATGCAGTGG']	['Primary hyperoxaluria, type III']
NM_004370.5(COL12A1):c.7001T>C (p.Ile2334Thr)	796052093	COL12A1	[]	[]	['BETHLEM MYOPATHY 2']
NM_000043.4(FAS):c.532T>C (p.Cys178Arg)	121913084	FAS	[]	[]	[]
NM_000208.2(INSR):c.779T>C (p.Leu260Pro)	121913141	INSR	[]	['CTACCYGGACGGCAGGTGTGTGG']	['Leprechaunism syndrome']
NM_000208.2(INSR):c.164T>C (p.Val55Ala)	121913152	INSR	[]	[]	['Leprechaunism syndrome']
NM_000026.2(ADSL):c.340T>C (p.Tyr114His)	374259530	ADSL	[]	[]	['Adenylosuccinate lyase deficiency', 'not provided']
NM_153490.2(KRT13):c.356T>C (p.Leu119Pro)	60440396	KRT13	[]	[]	['White sponge nevus 2', 'not provided']
NM_022455.4(NSD1):c.5885T>C (p.Ile1962Thr)	587784162	NSD1	[]	[]	['Sotos syndrome 1']
NM_006218.2(PIK3CA):c.1258T>C (p.Cys420Arg)	121913272	PIK3CA	[]	['GGAACACYGTCCATTGGCATGGG', 'GAACACYGTCCATTGGCATGGGG']	['Congenital lipomatous overgrowth, vascular malformations, and epidermal nevi', 'Neoplasm of ovary', 'PIK3CA Related Overgrowth Spectrum']
NM_000552.3(VWF):c.8317T>C (p.Cys2773Arg)	61751310	VWF	[]	['GCTCCYGCTGCTCTCCGACACGG']	['von Willebrand disease, type 2a', 'not provided']
NM_000335.4(SCN5A):c.5504T>C (p.Ile1835Thr)	45563942	SCN5A	[]	[]	['Primary dilated cardiomyopathy', 'Dilated cardiomyopathy 1E', 'not provided']
NM_183415.2(UBE3B):c.1741+2T>C	398123020	UBE3B	[]	[]	['Kaufman oculocerebrofacial syndrome']
NM_000021.3(PSEN1):c.1175T>C (p.Leu392Pro)	63750218	PSEN1	[]	[]	['Alzheimer disease, type 3', 'not provided']
NM_000350.2(ABCA	61751392	ABCA4	[]	[]	['Stargardt disease

4):c.1622T>C (p.Leu541Pro)					1', 'Cone-rod dystrophy 3', 'not provided']
NM_007313.2(ABL1): c.1109T>C (p.Met370Thr)	121913457	ABL1	[]	[]	[]
NM_024408.3(NOTCH2):c.1117T>C (p.Cys373Arg)	312262793	NOTCH2	[]	[]	['Alagille syndrome 2']
NM_024408.3(NOTCH2):c.1438T>C (p.Cys480Arg)	312262799	NOTCH2	[]	['TTCACAYGTCTGTGCATGCCAGG']	['Alagille syndrome 2']
NM_003611.2(OFD1): c.111+2T>C	312262809	OFD1	[]	[]	['Oral-facial-digital syndrome', 'not provided']
NM_003611.2(OFD1): c.274T>C (p.Ser92Pro)	312262819	OFD1	[]	[]	['Oral-facial-digital syndrome']
NM_020631.4(PLEKHG5):c.1940T>C (p.Phe647Ser)	63750315	PLEKHG5	[]	[]	['Distal spinal muscular atrophy, autosomal recessive 4']
NM_001288953.1(TTC7A):c.1912T>C (p.Ser638Pro)	149602485	TTC7A	[]	[]	['Multiple gastrointestinal atresias']
NM_000391.3(TPP1): c.1093T>C (p.Cys365Arg)	119455953	TPP1	['GCCGGGYGTTGGTCTGTCTCTGG']	['GCCGGGYGTTGGTCTGTCTCTGG']	['Ceroid lipofuscinosis, neuronal, 2', 'not provided']
NM_000426.3(LAMA2):c.7691T>C (p.Leu2564Pro)	121913570	LAMA2	[]	['ATCATTCYTTTGGGAAGTGGAGG', 'TCATTCYTTTGGGAAGTGGAGGG']	['Merosin deficient congenital muscular dystrophy']
NM_000426.3(LAMA2):c.2584T>C (p.Cys862Arg)	121913573	LAMA2	[]	[]	['Congenital muscular dystrophy due to partial LAMA2 deficiency']
NM_000530.6(MPZ):c.404T>C (p.Ile135Thr)	121913587	MPZ	[]	[]	['Charcot-Marie-Tooth disease type 1B']
NM_201253.2(CRB1): c.3541T>C (p.Cys1181Arg)	62636291	CRB1	[]	[]	['Retinitis pigmentosa 12', 'not provided']
NM_000257.3(MYH7):c.1046T>C (p.Met349Thr)	121913640	MYH7	[]	['AACTCCAYGTATAAGCTGACAGG']	['Familial hypertrophic cardiomyopathy 1', 'Cardiomyopathy']
NM_000257.3(MYH7):c.1594T>C (p.Ser532Pro)	121913642	MYH7	[]	['CATCATGYCCATCCTGGAAGAGG']	['Dilated cardiomyopathy 1S']
NM_000257.3(MYH7):c.5378T>C (p.Leu1793Pro)	121913654	MYH7	[]	[]	['Familial hypertrophic cardiomyopathy 1', 'Myosin storage myopathy', 'Left ventricular noncompaction 5', 'Cardiomyopathy']
NM_001127500.1(ME	121913668	MET	[]	[]	['Renal cell

T):c.3446T>C (p.Met1149Thr)					carcinoma, papillary, 1']
NM_002443.3(MSMB):c.-89T=	10993994	MSMB	[]	[]	['Prostate cancer, hereditary, 13']
NM_001079802.1(FK TN):c.527T>C (p.Phe176Ser)	119463996	FKTN	[]	['GTAGTCTYTCAT GAGAGGAGTGG']	['Limb-girdle muscular dystrophy-dystroglycanopathy, type C4']
NM_000169.2(GLA):c .865A>T (p.Ile289Phe)	140329381	-	[]	[]	['Fabry disease']
NM_000420.2(KEL):c .1790T>C (p.Leu597Pro)	8176038	KEL	[]	[]	[]
NM_003999.2(OSMR) :c.2072T>C (p.Ile691Thr)	63750567	OSMR	[]	[]	['Primary localized cutaneous amyloidosis 1']
NC_012920.1:m.9478 T>C	587776437	MT-CO3	['TCAG AAGYT TTTTTC TTCGC AGG']	['TCAGAAGYTTTT TTCTTCGCAGG']	['Leigh disease']
NM_002775.4(HTRA1):c.1091T>C (p.Leu364Pro)	587776447	HTRA1	[]	[]	['Cerebral autosomal recessive arteriopathy with subcortical infarcts and leukoencephalopath y']
NM_002049.3(GATA 1):c.2T>C (p.Met1Thr)	587776451	GATA1	['TCCA YGGAG TTCCCT GGCCT GGG', 'CCAYG GAGTT CCCTG GCCTG GGG']	['CTCCAYGGAGTT CCCTGGCCTGG', 'TCCAYGGAGTTC CCTGGCCTGGG', 'CCAYGGAGTTCC CTGGCCTGGGG']	['GATA-1-related thrombocytopenia with dyserythropoiesis']
NM_000021.3(PSEN1):c.254T>C (p.Leu85Pro)	63750599	PSEN1	[]	[]	['Alzheimer disease, familial, 3, with spastic paraparesis and apraxia', 'not provided']
NM_002049.3(GATA 1):c.1240T>C (p.Ter414Arg)	587776456	GATA1	[]	['GCTCAYGAGGG CACAGAGCATGG']	['GATA-1-related thrombocytopenia with dyserythropoiesis']
NM_006158.4(NEFL): c.281T>C (p.Leu94Pro)	62636505	NEFL	[]	[]	['Charcot-Marie-Tooth disease type 2E', 'not provided']
NM_000184.2(HBG2): c.-228T>C	63750654	HBG2	[]	['ATGCAAAYATCT GTCTGAAACGG']	['Fetal hemoglobin quantitative trait locus 1']
NM_000353.2(TAT):c .236-5A>G	587776512	TAT	[]	[]	['Tyrosinemia type 2']
NM_173560.3(RFX6): c.380+2T>C	587776514	RFX6	['CAGT GGYGA GACTC GCCCG CAGG',	['CAGTGGYGAGA CTCGCCCGCAGG', 'AGTGGYGAGACT CGCCCGCAGGG']	['Mitchell-Riley syndrome']

			'AGTGGYGAGACTCGCCCGCAGGG']		
NM_001999.3(FBN2):c.3725-15A>G	587776519	FBN2	[]	['AGCAYTGCAACCACATTGTCAGG']	['Congenital contractural arachnodactyly']
NM_000404.2(GLB1):c.1480-2A>G	587776526	GLB1	[]	[]	['GM1-GANGLIOSIDOSIS, TYPE I, WITH CARDIAC INVOLVEMENT']
NM_000402.4(G6PD):c.473T>C (p.Leu158Pro)	78365220	G6PD	[]	['TGCCCYCCACCTGGGGTCACAGG']	['Anemia, nonspherocytic hemolytic, due to G6PD deficiency', 'not provided']
NM_000179.2(MSH6):c.1346T>C (p.Leu449Pro)	63750741	MSH6	[]	['CTGGGGCYGGTATTCATGAAAGG']	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_145046.4(CALR3):c.245A>G (p.Lys82Arg)	142951029	CALR3	['CGGTYTGAAGCGTGCAGAGATGG']	['CGGTYTGAAGCGTGCAGAGATGG']	['Arrhythmogenic right ventricular cardiomyopathy', 'Familial hypertrophic cardiomyopathy 19', 'Hypertrophic cardiomyopathy']
NM_000260.3(MYO7A):c.5573T>C (p.Leu1858Pro)	368657015	MYO7A	[]	[]	['Usher syndrome, type 1']
NM_024753.4(TTC21B):c.2758-2A>G	766132877	TTC21B	[]	[]	['Nephronophthisis 12']
NM_001195129.1(PRSS56):c.1183T>C (p.Cys395Arg)	730882161	PRSS56	[]	[]	['Microphthalmia, isolated 6']
NM_001184.3(ATR):c.2022A>G (p.Gly674=)	587776690	ATR	[]	[]	['Seckel syndrome 1']
NM_000354.5(SERPINA7):c.623-2A>G	587776720	SERPINA7	[]	[]	[]
NM_000133.3(F9):c.277+2T>C	587776735	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_004006.2(DMD):c.9225-285A>G	587776747	DMD	[]	[]	['Becker muscular dystrophy']
NM_016835.4(MAPT):c.1839T>C (p.Asn613=)	63750912	MAPT	['GGATAAYATCAAACACGTCCCGG', 'GATAAYATCAAACACGTCCCGGG']	['GGATAAYATCAAACACGTCCCGG', 'GATAAYATCAAACACGTCCCGGG']	['Frontotemporal dementia', 'not provided']
NM_000321.2(RB1):c.1960+2T>C	587776780	RB1	[]	[]	['Retinoblastoma']

NM_006517.4(SLC16A2):c.1253T>C (p.Leu418Pro)	367543059	SLC16A2	[]	[]	['Allan-Herndon-Dudley syndrome']
NM_000421.3(KRT10):c.1374-2A>G	587776815	-	[]	[]	['Erythroderma, ichthyosiform, congenital reticular']
NM_000251.2(MSH2):c.2089T>C (p.Cys697Arg)	63750961	MSH2	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_002618.3(PEX13):c.977T>C (p.Ile326Thr)	61752115	PEX13	[]	[]	['Peroxisome biogenesis disorder 11B']
NM_001079867.1(PEX2):c.739T>C (p.Cys247Arg)	61752128	PEX2	[]	[]	['Peroxisome biogenesis disorder 5A']
NM_017929.5(PEX26):c.134T>C (p.Leu45Pro)	61752132	PEX26	[]	[]	['Peroxisome biogenesis disorder 7B']
NG_012088.1:g.2209A>G	587776843	IL10	['ACCYTATGATCCGCCCGCCTTGG']	['ACCYTATGATCCGCCCGCCTTGG']	[]
NM_001735.2(C5):c.1115A>G (p.Lys372Arg)	587776846	C5	[]	[]	['Leiner disease']
NM_002087.3(GRN):c.2T>C (p.Met1Thr)	63751006	GRN	['CCAYGTGGACCCTGGTGAGCTGG']	['CCAYGTGGACCCTGGTGAGCTGG']	['Frontotemporal dementia, ubiquitin-positive', 'not provided']
NM_004656.3(BAP1):c.2057-2A>G	587776878	BAP1	[]	[]	['Tumor predisposition syndrome']
NM_004656.3(BAP1):c.438-2A>G	587776879	BAP1	['GCCYGGGGAAAACAGAGTCAGG']	['GCCYGGGGAAAACAGAGTCAGG']	['Tumor predisposition syndrome']
NM_004329.2(BMPR1A):c.370T>C (p.Cys124Arg)	199476087	BMPR1A	[]	[]	['Juvenile polyposis syndrome', 'Hereditary cancer-predisposing syndrome']
NM_000510.2(FSHB):c.298T>C (p.Cys100Arg)	5030777	FSHB	[]	[]	['Follicle-stimulating hormone deficiency, isolated']
NM_001009944.2(PKD1):c.2534T>C (p.Leu845Ser)	199476100	PKD1	[]	[]	['Polycystic kidney disease, adult type']
NM_004963.3(GUCY2C):c.1160A>G (p.Asp387Gly)	587776905	GUCY2C	[]	[]	['Meconium ileus']
m.14487T>C	199476109	MT-ND6	[]	[]	['Leigh disease', 'Leigh syndrome due to mitochondrial complex I deficiency']

NM_017565.3(FAM20A):c.813-2A>G	587776912	-	[]	[]	['Enamel-renal syndrome']
NM_017565.3(FAM20A):c.590-2A>G	587776914	FAM20A	[]	['GTAATCYGCAAAGGAGGAGAAGG', 'TAATCYGCAAAGGAGGAGAAGGG']	['Enamel-renal syndrome']
NM_014165.3(NDUFAF4):c.194T>C (p.Leu65Pro)	63751061	NDUFAF4	[]	[]	['Mitochondrial complex I deficiency']
m.4160T>C	199476119	MT-ND1	[]	[]	['Leber optic atrophy']
NM_000551.3(VHL):c.292T>C (p.Tyr98His)	5030809	VHL	[]	['CCCYACCCAACGCTGCCGCCTGG']	['Von Hippel-Lindau syndrome', 'Hereditary cancer-predisposing syndrome']
m.3949T>C	199476124	MT-ND1	[]	[]	['Juvenile myopathy, encephalopathy, lactic acidosis AND stroke']
m.6742T>C	199476126	MT-CO1	[]	[]	[]
m.6721T>C	199476127	MT-CO1	[]	[]	[]
m.5692T>C	199476131	MT-TN	[]	[]	[]
m.5728T>C	199476132	MT-TN	[]	['CAATCYACTTCTCCCGCCGCCGG', 'AATCYACTTCTCCCGCCGCCGGG']	['Cytochrome-c oxidase deficiency', 'Mitochondrial complex I deficiency']
m.9101T>C	199476134	MT-ATP6	[]	[]	['Leber optic atrophy']
m.8851T>C	199476136	MT-ATP6	[]	[]	['Leigh disease', 'Striatonigral degeneration, infantile, mitochondrial']
m.9185T>C	199476138	MT-ATP6	[]	[]	['Leigh disease']
NM_000277.1(PAH):c.143T>C (p.Leu48Ser)	5030841	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_000277.1(PAH):c.691T>C (p.Ser231Pro)	5030845	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_000251.2(MSH2):c.595T>C (p.Cys199Arg)	63751110	MSH2	['AAGGAAYGTGTTTACCCGGAGG']	['AAGGAAYGTGTTTACCCGGAGG']	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_000039.1(APOA1):c.341T>C (p.Leu114Pro)	28931575	-	[]	[]	[]
NM_001288953.1(TTC7A):c.2366T>C (p.Leu789Pro)	587776972	TTC7A	[]	[]	['Multiple gastrointestinal atresias']
NM_014336.4(AIPL1):c.715T>C (p.Cys239Arg)	62637012	AIPL1	[]	['CTGCCAGYGCCTGCTGAAGAAGG', 'CCAGYGCCTGCTGAAGAAGGAGG']	['Leber congenital amaurosis 4', 'not provided']
NM_000155.3(GALT)	367543254	GALT	[]	[]	['Deficiency of

:c.336T>C (p.Ser112=)					UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_207352.3(CYP4V2):c.655T>C (p.Tyr219His)	199476191	CYP4V2	[]	[]	['Bietti crystalline corneoretinal dystrophy']
NM_207352.3(CYP4V2):c.1021T>C (p.Ser341Pro)	199476199	CYP4V2	[]	['AAACTGGYCCTTATACCTGTTGG', 'AACTGGYCCTTATACCTGTTGGG']	['Bietti crystalline corneoretinal dystrophy']
NM_001142519.1(FAM111A):c.1531T>C (p.Tyr511His)	587777012	FAM111A	[]	[]	['Kenny-Caffey syndrome type 2']
NM_000435.2(NOTCH3):c.4556T>C (p.Leu1519Pro)	367543285	NOTCH3	[]	[]	['Infantile myofibromatosis 1', 'Infantile myofibromatosis 2']
NM_000021.3(PSEN1):c.749T>C (p.Leu250Ser)	63751163	PSEN1	[]	[]	['Alzheimer disease, type 3', 'not provided']
NM_001283009.1(RTEL1):c.3730T>C (p.Cys1244Arg)	587777037	-	['CTGTGTGYGCCAGGGCTGTGGG']	['CTGTGTGYGCCAGGGCTGTGGGG']	['Dyskeratosis congenita, autosomal recessive, 5']
NM_001135021.1(ELMOD3):c.794T>C (p.Leu265Ser)	587777040	ELMOD3	[]	[]	['Deafness, autosomal recessive 88']
NM_001001557.2(GDF6):c.866T>C (p.Leu289Pro)	63751220	GDF6	[]	[]	['Klippel-Feil syndrome 1, autosomal dominant']
NM_014754.2(PTDSS1):c.794T>C (p.Leu265Pro)	587777090	PTDSS1	[]	[]	['Lenz-Majewski hyperostosis syndrome']
NM_052844.3(WDR34):c.1307A>G (p.Lys436Arg)	587777098	WDR34	[]	[]	['Short-rib thoracic dysplasia 11 with or without polydactyly']
NM_001290048.1(ATL3):c.521A>G (p.Tyr174Cys)	587777108	ATL3	[]	[]	['Hereditary sensory neuropathy type IF']
NM_001018005.1(TPM1):c.515T>C (p.Ile172Thr)	199476312	TPM1	[]	[]	['Primary familial hypertrophic cardiomyopathy', 'Cardiomyopathy', 'not provided']
NM_018849.2(ABCB4):c.523A>G (p.Thr175Ala)	58238559	ABCB4	[]	[]	['Cholecystitis']
NM_001018005.1(TPM1):c.842T>C (p.Met281Thr)	199476321	TPM1	[]	[]	['Cardiomyopathy', 'not specified', 'not provided']
NM_005763.3(AASS):c.874A>G (p.Ile292Val)	587777122	AASS	[]	[]	['Hyperlysinemia']
NM_194442.2(LBR):c.1639A>G (p.Asn547Asp)	587777171	LBR	[]	[]	['Greenberg dysplasia']

NM_006702.4(PNPLA6):c.3053T>C (p.Phe1018Ser)	587777183	PNPLA6	[]	['CCTYTAACCGCAGCATCCATCGG']	['Boucher Neuhauser syndrome']
NM_000487.5(ARSA):c.899T>C (p.Leu300Ser)	199476389	ARSA	[]	['GGTCTCTYGCGGTGTGGAAAGGG']	['Metachromatic leukodystrophy', 'not provided']
NM_016599.4(MYOZ2):c.142T>C (p.Ser48Pro)	199476398	MYOZ2	[]	['TTAYCCCATCTCAGTAACCGTGG']	['Familial hypertrophic cardiomyopathy 16', 'not provided']
NM_014740.3(EIF4A3):c.809A>G (p.Asp270Gly)	587777204	EIF4A3	[]	[]	['Richieri Costa Pereira syndrome']
NM_001040436.2(YARS2):c.1303A>G (p.Ser435Gly)	587777215	YARS2	[]	[]	['Myopathy, lactic acidosis, and sideroblastic anemia 2']
NM_001278503.1(STT3A):c.1877T>C (p.Val626Ala)	587777216	STT3A	[]	[]	['Congenital disorder of glycosylation type 1w']
NM_001037633.1(SIL1):c.1370T>C (p.Leu457Pro)	119456967	SIL1	[]	['TTGCYGAAGGAGCTGAGATGAGG']	['Marinesco-Sj\xc3\xb6gren syndrome']
NM_006888.4(CALM1):c.268T>C (p.Phe90Leu)	730882253	CALM1	[]	['GGCAYTCCGAGTCTTTGACAAGG']	['Long QT syndrome 14']
NM_001003811.1(TEX11):c.511A>G (p.Met171Val)	143246552	TEX11	['TCCAYGGTCAAGTCAGCCTCAGG']	['TCCAYGGTCAAGTCAGCCTCAGG', 'CCAYGGTCAAGTCAGCCTCAGGG']	['Spermatogenic failure, X-linked, 2']
NM_033419.4(PGAP3):c.914A>G (p.Asp305Gly)	587777252	PGAP3	[]	[]	['Hyperphosphatasia with mental retardation syndrome 4']
NM_000021.3(PSEN1):c.338T>C (p.Leu113Pro)	63751399	PSEN1	[]	[]	['Alzheimer disease, type 3', 'Frontotemporal dementia', 'not provided']
NM_000097.5(CPOX):c.980A>G (p.His327Arg)	587777271	CPOX	[]	[]	['Harderoporphyria']
NM_005654.5(NR2F1):c.755T>C (p.Leu252Pro)	587777276	NR2F1	[]	[]	['Bosch-boonstra-schaaf optic atrophy syndrome']
NM_012338.3(TSPAN12):c.413A>G (p.Tyr138Cys)	587777283	TSPAN12	[]	['TAATCCAYAATTTGTCATCCTGG']	['Exudative vitreoretinopathy 5']
NM_003181.3(T):c.512A>G (p.His171Arg)	587777303	T	[]	[]	['Sacral agenesis with vertebral anomalies']
NM_015884.3(MBTPS2):c.1391T>C (p.Phe464Ser)	587777306	MBTPS2	[]	['GCTYTGCTTTGGATGGACAATGG']	['Palmoplantar keratoderma, mutilating, with periorificial keratotic plaques, X-linked']
NM_020435.3(GJC2):	74315311	GJC2	['TGAG	['TGAGAYGGCCC	['Leukodystrophy,

c.857T>C (p.Met286Thr)			AYGGCCCACCTGGGCTTGG']	ACCTGGGCTTGG', 'GAGAYGGCCCACCTGGGCTTGGG']	hypomyelinating, 2']
NM_005356.4(LCK):c.1022T>C (p.Leu341Pro)	587777335	LCK	[]	[]	['Immunodeficiency 22']
NM_005861.3(STUB1):c.719T>C (p.Met240Thr)	587777345	-	[]	[]	['Spinocerebellar ataxia, autosomal recessive 16']
NM_000250.1(MPO):c.752T>C (p.Met251Thr)	56378716	MPO	[]	['TCACTCAYGTTCATGCAATGGGG']	['Myeloperoxidase deficiency']
NM_017890.4(VPS13B):c.11119+2T>C	587777382	VPS13B	[]	[]	['Cohen syndrome']
NM_005026.3(PIK3CD):c.1246T>C (p.Cys416Arg)	587777390	PIK3CD	[]	['GCAGGACYGCCCCATTGCCTGGG']	['Activated PI3K-delta syndrome']
NM_002633.2(PGM1):c.1547T>C (p.Leu516Pro)	587777401	PGM1	[]	[]	['Congenital disorder of glycosylation type 1t']
NM_000261.1(MYOC):c.1309T>C (p.Tyr437His)	74315328	MYOC	[]	[]	['Primary open angle glaucoma juvenile onset 1']
NM_001159287.1(TPI1):c.833T>C (p.Phe278Ser)	587777440	TPI1	[]	[]	['Triosephosphate isomerase deficiency']
NM_000414.3(HSD17B4):c.1547T>C (p.Ile516Thr)	587777443	HSD17B4	[]	[]	['Gonadal dysgenesis with auditory dysfunction, autosomal recessive inheritance']
NM_005359.5(SMAD4):c.970T>C (p.Cys324Arg)	377767339	SMAD4	[]	[]	['Juvenile polyposis syndrome']
NM_000211.4(ITGB2):c.1877+2T>C	483352818	ITGB2	['CATGYGAGTGCAGGCGGAGCAGG']	['CATGYGAGTGCAGGCGGAGCAGG']	['Leukocyte adhesion deficiency type 1']
NM_005359.5(SMAD4):c.1087T>C (p.Cys363Arg)	377767348	SMAD4	[]	[]	['Juvenile polyposis syndrome']
NM_001128159.2(VPS53):c.2084A>G (p.Gln695Arg)	587777465	VPS53	[]	[]	['Pontocerebellar hypoplasia, type 2e']
NM_000249.3(MLH1):c.1745T>C (p.Leu582Pro)	63751616	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_003108.3(SOX11):c.178T>C (p.Ser60Pro)	587777480	SOX11	[]	['TATGGYCCAAGATCGAACGCAGG']	['Mental retardation, autosomal dominant 27']
NM_020630.4(RET):c.1888T>C (p.Cys630Arg)	377767404	RET	[]	[]	[]
NM_017565.3(FAM20A):c.720-2A>G	587777530	-	[]	[]	['Enamel-renal syndrome']

NM_015599.2(PGM3):c.737A>G (p.Asn246Ser)	587777562	PGM3	['TAAATGAYTGAGTTTGCCCTTGG']	['TAAATGAYTGAGTTTGCCCTTGG']	['Immunodeficiency 23']
NM_015599.2(PGM3):c.1352A>G (p.Gln451Arg)	587777565	PGM3	[]	[]	['Immunodeficiency 23']
NM_000206.2(IL2RG):c.452T>C (p.Leu151Pro)	137852511	IL2RG	[]	[]	['X-linked severe combined immunodeficiency']
NM_198282.3(TMEM173):c.461A>G (p.Asn154Ser)	587777609	TMEM173	[]	[]	['Sting-associated vasculopathy, infantile-onset']
NM_000329.2(RPE65):c.1022T>C (p.Leu341Ser)	61752909	RPE65	[]	[]	['Retinitis pigmentosa 20', 'not provided']
NM_001127899.3(CLCN5):c.1768T>C (p.Ser590Pro)	151340623	CLCN5	[]	[]	['Dent disease 1']
NM_005027.3(PIK3R2):c.1202T>C (p.Leu401Pro)	587777624	PIK3R2	[]	[]	['Megalencephaly polymicrogyria-polydactyly hydrocephalus syndrome']
NM_007315.3(STAT1):c.854A>G (p.Gln285Arg)	587777629	STAT1	[]	[]	['Immunodeficiency 31C']
NM_139276.2(STAT3):c.1175A>G (p.Lys392Arg)	587777648	STAT3	[]	[]	['Autoimmune disease, multisystem, infantile-onset']
NM_001037811.2(HSD17B10):c.257A>G (p.Asp86Gly)	587777651	HSD17B10	[]	[]	['2-methyl-3-hydroxybutyric aciduria']
NM_001288767.1(ARMC5):c.1928T>C (p.Leu643Pro)	587777661	ARMC5	[]	[]	['Acth-independent macronodular adrenal hyperplasia 2']
NM_001288767.1(ARMC5):c.1379T>C (p.Leu460Pro)	587777663	ARMC5	[]	['GCCCGACYGCGGGATGCTGGTGG']	['Acth-independent macronodular adrenal hyperplasia 2']
NM_007315.3(STAT1):c.2018A>G (p.Lys673Arg)	587777704	STAT1	[]	[]	['Immunodeficiency 31a']
NM_007315.3(STAT1):c.1909A>G (p.Lys637Glu)	587777705	STAT1	[]	[]	['Immunodeficiency 31a']
NM_014845.5(FIG4):c.50T>C (p.Leu17Pro)	587777713	FIG4	[]	[]	['Charcot-Marie-Tooth disease, type 4J']
NM_000350.2(ABCA4):c.5819T>C (p.Leu1940Pro)	61753033	ABCA4	[]	['AAGGCYACATGAACTAACCAAGG']	['Stargardt disease', 'Stargardt disease 1', 'Cone-rod dystrophy 3', 'not provided']
NM_002972.3(SBF1):c.4768A>G (p.Thr1590Ala)	200488568	SBF1	[]	['CAGGCGYCCTCTTGCTCAGCCGG']	['Charcot-Marie-Tooth disease, type 4B3']

NM_000377.2(WAS): c.244T>C (p.Ser82Pro)	132630272	WAS	[]	[]	[]
NM_000377.2(WAS): c.809T>C (p.Leu270Pro)	132630274	WAS	[]	['CGGAGTCYGTTC TCCAGGGCAGG']	['Severe congenital neutropenia X-linked']
NM_001128834.2(PLP 1):c.487T>C (p.Trp163Arg)	132630279	PLP1	[]	[]	['Pelizaeus-Merzbacher disease', 'not provided']
NM_001128834.2(PLP 1):c.671T>C (p.Leu224Pro)	132630283	PLP1	[]	[]	['Pelizaeus-Merzbacher disease']
NM_001128834.2(PLP 1):c.560T>C (p.Ile187Thr)	132630288	PLP1	[]	[]	['Spastic paraplegia 2']
NM_001128834.2(PLP 1):c.710T>C (p.Phe237Ser)	132630291	PLP1	[]	[]	['Spastic paraplegia 2']
NM_001015877.1(PH F6):c.2T>C (p.Met1Thr)	132630300	PHF6	[]	[]	['Borjeson-Forssman-Lehmann syndrome']
NM_001399.4(EDA):c .181T>C (p.Tyr61His)	132630308	EDA	[]	['CTGCYACCTAGA GTTGCGCTCGG']	['Hypohidrotic X-linked ectodermal dysplasia']
NM_001205019.1(GK):c.1525T>C (p.Trp509Arg)	132630330	GK	[]	[]	['Deficiency of glycerol kinase']
NM_000076.2(CDKN 1C):c.*5+2T>C	587777866	CDKN1C	['CCAA GYGAG TACAG CGCAC CTGG', 'CAAGY GAGTA CAGCG CACCT GGG', 'AAGYG AGTAC AGCGC ACCTG GGG']	['CCAAGYGAGTA CAGCGCACCTGG', 'CAAGYGAGTACA GCGCACCTGGG', 'AAGYGAGTACAG CGCACCTGGGG']	['Beckwith-Wiedemann syndrome']
NM_000271.4(NPC1): c.2054T>C (p.Ile685Thr)	483352888	NPC1	[]	[]	['Niemann-Pick disease type C1']
NM_170707.3(LMNA):c.1589T>C (p.Leu530Pro)	60934003	LMNA	[]	['ACGGCTCYCATC AACTCCACTGG', 'CGGCTCYCATCA ACTCCACTGGG', 'GGCTCYCATCAA CTCCACTGGGG']	['Benign scapuloperoneal muscular dystrophy with cardiomyopathy', 'not provided']
NM_001165963.1(SC N1A):c.5536A>T (p.Lys1846Ter)	372098964	-	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_130838.1(UBE3A):c.2485T>C (p.Tyr829His)	587784526	UBE3A	[]	[]	['Angelman syndrome']
NM_014588.5(VSX1): c.50T>C (p.Leu17Pro)	74315436	VSX1	[]	[]	['Keratoconus 1']
NM_000404.2(GLB1):	398123354	GLB1	[]	[]	['Mucopolysaccharid

c.457+2T>C					osis, MPS-IV-B', 'Infantile GM1 gangliosidosis', 'Juvenile GM>1< gangliosidosis', 'Gangliosidosis GM1 type 3', 'not provided']
NM_003159.2(CDKL5):c.145+2T>C	267608430	CDKL5	[]	[]	['Atypical Rett syndrome', 'not provided']
NM_002764.3(PRPS1):c.869T>C (p.Ile290Thr)	180177153	PRPS1	[]	[]	['Deafness, high-frequency sensorineural, X-linked']
NM_000030.2(AGXT):c.1076T>C (p.Leu359Pro)	180177160	AGXT	[]	['GGTGCYGCGGATCGGCCTGCTGG', 'GTGCYGCGGATCGGCCTGCTGGG']	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT):c.1151T>C (p.Leu384Pro)	180177165	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT):c.449T>C (p.Leu150Pro)	180177222	AGXT	[]	['GTGCYGCTGTTCTTAACCCACGG', 'TGCYGCTGTTCTTAACCCACGGG']	['Primary hyperoxaluria, type I']
NM_000268.3(NF2):c.1079T>C (p.Leu360Pro)	74315492	NF2	[]	[]	['Neurofibromatosis, type 2']
NM_000030.2(AGXT):c.661T>C (p.Ser221Pro)	180177254	AGXT	[]	['GCTCATCYCCTTCAGTGACAAGG']	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT):c.757T>C (p.Cys253Arg)	180177264	AGXT	[]	['GGGGCYGTGACGACCAGCCCAGG']	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT):c.77T>C (p.Leu26Pro)	180177268	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT):c.851T>C (p.Leu284Pro)	180177287	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT):c.893T>C (p.Leu298Pro)	180177293	AGXT	[]	['GTATCYGCATGGGCGCCTGCAGG']	['Primary hyperoxaluria, type I']
NM_012203.1(GRHPR):c.203T>C (p.Leu68Pro)	180177305	GRHPR	[]	[]	['Primary hyperoxaluria, type II']
NM_000017.3(ACADS):c.1057T>C (p.Ser353Pro)	796051904	ACADS	[]	[]	['not provided']
NM_000406.2(GNRHR):c.392T>C (p.Met131Thr)	606231406	GNRHR	[]	[]	['Hypogonadotropic hypogonadism']
NM_000255.3(MUT):c.842T>C (p.Leu281Ser)	796052007	MUT	[]	[]	['not provided']
NM_000030.2(AGXT):c.947T>C (p.Leu316Pro)	796052063	AGXT	[]	[]	['Primary hyperoxaluria, type I']

NM_138413.3(HOGA1):c.875T>C (p.Met292Thr)	796052087	HOGA1	[]	[]	['Primary hyperoxaluria, type III']
NM_013382.5(POMT2):c.1997A>G (p.Tyr666Cys)	200198778	POMT2	['GGAAGYAGTGGTGGAAGTAGAGG']	['GGAAGYAGTGGTGGAAGTAGAGG']	['Congenital muscular dystrophy', 'Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies, type A2', 'Muscular dystrophy', 'Congenital muscular dystrophy-dystroglycanopathy with mental retardation, type B2', 'not provided']
NM_015909.3(NBAS):c.3164T>C (p.Leu1055Pro)	796052121	NBAS	[]	[]	['Infantile liver failure syndrome 2']
NM_000263.3(NAGLU):c.1208T>C (p.Ile403Thr)	796052122	NAGLU	[]	[]	['Charcot-Marie-Tooth disease, axonal type 2V']
NM_203290.2(POLR1C):c.436T>C (p.Cys146Arg)	796052125	POLR1C	[]	[]	['LEUKODYSTROPHY, HYPOMYELINATING, 11']
NM_018359.3(UFSP2):c.868T>C (p.Tyr290His)	796052130	-	[]	[]	['Hip dysplasia, beukes type']
NM_000053.3(ATP7B):c.122A>G (p.Asn41Ser)	201738967	ATP7B	[]	[]	['Wilson disease']
NM_001356.4(DDX3X):c.704T>C (p.Leu235Pro)	796052224	DDX3X	[]	[]	['not provided']
NM_001356.4(DDX3X):c.1541T>C (p.Ile514Thr)	796052226	DDX3X	[]	[]	['not provided']
NM_001356.4(DDX3X):c.1175T>C (p.Leu392Pro)	796052232	DDX3X	[]	[]	['not provided']
NM_000321.2(RB1):c.2663+2T>C	587778839	RB1	[]	[]	['Retinoblastoma']
NM_000321.2(RB1):c.1472T>C (p.Leu491Pro)	587778848	RB1	[]	[]	['Retinoblastoma']
NM_006894.5(FMO3):c.1079T>C (p.Leu360Pro)	28363581	FMO3	[]	[]	['Trimethylaminuria']
NM_172107.2(KCNQ2):c.583T>C (p.Ser195Pro)	796052620	KCNQ2	[]	[]	['not provided']
NM_001282227.1(CECR1):c.1232A>G (p.Tyr411Cys)	376785840	CECR1	[]	['GAAATCAYAGGACAAGCCTTTGG']	['Polyarteritis nodosa']
NM_170707.3(LMNA):c.644T>C	61295588	LMNA	[]	[]	['Dilated cardiomyopathy 1A',

(p.Leu215Pro)					'not provided']
NM_005249.4(FOXG1):c.673T>C (p.Trp225Arg)	796052482	FOXG1	[]	[]	['not provided']
NM_000806.5(GABRA1):c.788T>C (p.Met263Thr)	796052491	GABRA1	[]	[]	['not provided']
NM_000251.2(MSH2):c.1319T>C (p.Leu440Pro)	587779084	MSH2	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_003401.3(XRCC4):c.127T>C (p.Trp43Arg)	587779351	XRCC4	[]	[]	['Ateleiotic dwarfism']
NM_198056.2(SCN5A):c.1247A>G (p.Tyr416Cys)	372395294	SCN5A	['CTCAYAGGCCATTGCGACCACGG']	['CTCAYAGGCCATTGCGACCACGG']	['not provided']
NM_000257.3(MYH7):c.4835T>C (p.Leu1612Pro)	587779392	-	[]	[]	['Myopathy, distal, 1']
NM_000257.3(MYH7):c.4937T>C (p.Leu1646Pro)	587779393	-	[]	['GAGCCYCCAGAGCTTGTTGAAGG']	['Myopathy, distal, 1']
NM_000404.2(GLB1):c.922T>C (p.Phe308Leu)	587779404	GLB1	[]	[]	['Infantile GM1 gangliosidosis']
NM_012434.4(SLC17A5):c.500T>C (p.Leu167Pro)	587779410	SLC17A5	[]	['ATTGTACYCAGAGCACTAGAAGG']	['Sialic acid storage disease, severe infantile type']
NM_000257.3(MYH7):c.4442T>C (p.Leu1481Pro)	587779414	-	[]	[]	['Myopathy, distal, 1']
NM_000090.3(COL3A1):c.2022+2T>C (p.Gly660_Lys674del)	587779429	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_004453.3(ETFDH):c.1852T>C (p.Ter618Gln)	765742496	ETFDH	[]	[]	['not provided']
NM_000090.3(COL3A1):c.2337+2T>C (p.Gly762_Lys779del)	587779513	COL3A1	[]	['AGGYAACCCTTAATACTACCTGG']	['Ehlers-Danlos syndrome, type 4']
NM_000310.3(PPT1):c.2T>C (p.Met1Thr)	796052927	PPT1	[]	[]	['not provided']
NM_020376.3(PNPLA2):c.757+2T>C	777539013	PNPLA2	[]	['GAACGGYGCGCGGACCCGGGCGG', 'AACGGYGCGCGGACCCGGGCGGG']	['Neutral lipid storage disease with myopathy']
NM_000090.3(COL3A1):c.3039+6T>C (p.Asp1013_Gly1014insVSSSFYSTSQ)	587779532	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_012452.2(TNFRSF13B):c.310T>C (p.Cys104Arg)	34557412	TNFRSF13B	[]	['ACTTCYGTGAGAACAAGCTCAGG']	['Immunoglobulin A deficiency 2', 'Common variable immunodeficiency 2']

NM_000570.4(FCGR3B):c.244A=(p.Asn82=)	147574249	FCGR3B	[]	[]	[]
NM_001165963.1(SCN1A):c.1094T>C (p.Phe365Ser)	796052970	SCN1A	[]	['CAAGCTYTGATACCTTCAGTTGG', 'AAGCTYTGATACCTTCAGTTGGG']	['not provided']
NC_012920.1:m.7505T>C	724159989	MT-TS1	[]	['CCTCCAYGACTTTTTCAAAAAGG']	['Deafness, nonsyndromic sensorineural, mitochondrial']
NM_000663.4(ABAT):c.1433T>C (p.Leu478Pro)	724159991	ABAT	[]	[]	['Gamma-aminobutyric acid transaminase deficiency']
NM_153818.1(PEX10):c.890T>C (p.Leu297Pro)	724160000	PEX10	[]	[]	['Peroxisome biogenesis disorder 6B']
NM_153818.1(PEX10):c.2T>C (p.Met1Thr)	724160002	PEX10	[]	[]	['Peroxisome biogenesis disorder 6B']
NM_000090.3(COL3A1):c.4399T>C (p.Ter1467Gln)	587779618	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_002485.4(NBN):c.511A>G (p.Ile171Val)	61754966	NBN	[]	[]	['Microcephaly, normal intelligence and immunodeficiency', 'Aplastic anemia', 'Hereditary cancer-predisposing syndrome', 'Leukemia, acute lymphoblastic, susceptibility to', 'not specified', 'not provided']
NM_000090.3(COL3A1):c.2553+2T>C (p.Gly816_Ala851del)	587779684	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_021007.2(SCN2A):c.4308+2T>C	796053139	SCN2A	['CGAAATGYAAGTCTAGTTAGAGG', 'GAAATGYAAGTCTAGTTAGAGGG']	['CGAAATGYAAGTCTAGTTAGAGG', 'GAAATGYAAGTCTAGTTAGAGGG']	['not provided']
NM_021007.2(SCN2A):c.4718T>C (p.Leu1573Pro)	796053152	SCN2A	[]	[]	['not provided']
NM_001101.3(ACTB):c.356T>C (p.Met119Thr)	587779773	ACTB	['GAGAAGAYGACCCAGGTGAGTGG']	['GAGAAGAYGACCCAGGTGAGTGG']	['Baraitser-Winter syndrome 1']
NM_021007.2(SCN2A):c.2306T>C	796053191	SCN2A	[]	[]	['not provided']

(p.Ile769Thr)					
NM_014191.3(SCN8A):c.4889T>C (p.Leu1630Pro)	796053222	SCN8A	[]	['CGTCYGATCAAAGGCGCCAAAGG', 'GTCYGATCAAAGGCGCCAAAGGG']	['not provided']
NM_012415.3(RAD54B):c.1778A>G (p.Asn593Ser)	114216685	RAD54B	[]	[]	['Malignant lymphoma, non-Hodgkin']
NM_007215.3(POLG2):c.1105A>G (p.Arg369Gly)	201936720	POLG2	[]	[]	['Autosomal dominant progressive external ophthalmoplegia with mitochondrial DNA deletions 4', 'not specified']
NM_004006.2(DMD):c.6982A>T (p.Lys2328Ter)	754896795	DMD	['GCTTTTYTTCAAGCTGCCCAAGG']	['GCTTTTYTTCAAGCTGCCCAAGG']	['Duchenne muscular dystrophy', 'Becker muscular dystrophy', 'Dilated cardiomyopathy 3B']
NM_001061.4(TBXAS1):c.248T>C (p.Leu83Pro)	140005285	TBXAS1	[]	[]	['not provided']
NM_000540.2(RYR1):c.10817T>C (p.Leu3606Pro)	118192127	RYR1	[]	['TACTACCYGGACCAGGTGGGTGG', 'ACTACCYGGACCAGGTGGGTGGG', 'CTACCYGGACCAGGTGGGTGGGG']	['Central core disease', 'not provided']
NM_000138.4(FBN1):c.7754T>C (p.Ile2585Thr)	727503054	FBN1	[]	[]	['Thoracic aortic aneurysms and aortic dissections', 'Marfan syndrome']
NM_001128227.2(GNE):c.2228T>C (p.Met743Thr)	28937594	GNE	[]	[]	['Inclusion body myopathy 2', 'Nonaka myopathy']
NM_000540.2(RYR1):c.14693T>C (p.Ile4898Thr)	118192170	RYR1	[]	['AGGCAYTGGGGACGAGATCGAGG']	['Malignant hyperthermia susceptibility type 1', 'Central core disease', 'not provided']
NM_005247.2(FGF3):c.466T>C (p.Ser156Pro)	121917703	FGF3	[]	['GTACGTGYCTGTGAACGGCAAGG', 'TACGTGYCTGTGAACGGCAAGGG']	['Deafness with labyrinthine aplasia microtia and microdontia (LAMM)']
NM_005247.2(FGF3):c.17T>C (p.Leu6Pro)	121917706	FGF3	[]	[]	['Deafness with labyrinthine aplasia microtia and microdontia (LAMM)']
NM_005211.3(CSF1R):c.2450T>C (p.Leu817Pro)	690016549	CSF1R	[]	['CCGCCYGCCTGTGAAGTGGATGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_005211.3(CSF1R):c.2480T>C (p.Ile827Thr)	690016550	CSF1R	[]	[]	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_005211.3(CSF1R	690016552	CSF1R	[]	['GAATCCCYACCC	['Hereditary diffuse

):c.2566T>C (p.Tyr856His)				TGGCATCCTGG']	leukoencephalopathy with spheroids']
NM_001098668.2(SFTPA2):c.593T>C (p.Phe198Ser)	121917738	SFTPA2	[]	['GGAGACTYCCGCTACTCAGATGG', 'GAGACTYCCGCTACTCAGATGGG']	['Idiopathic fibrosing alveolitis, chronic form']
NM_005211.3(CSF1R):c.1957T>C (p.Cys653Arg)	690016559	CSF1R	[]	['AGCCYGTACCCATGGAGGTAAGG', 'GCCYGTACCCATGGAGGTAAGGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_005211.3(CSF1R):c.2717T>C (p.Ile906Thr)	690016560	CSF1R	[]	['GCAGAYCTGCTCCTTCCTTCAGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_000540.2(RYR1):c.14378T>C (p.Leu4793Pro)	118192179	RYR1	[]	[]	['Central core disease', 'not provided']
NM_199292.2(TH):c.707T>C (p.Leu236Pro)	121917763	TH	[]	[]	['Segawa syndrome, autosomal recessive']
NM_000256.3(MYBPC3):c.2994+2T>C	727503176	MYBPC3	[]	[]	['Familial hypertrophic cardiomyopathy 4']
NM_003361.3(UMOD):c.376T>C (p.Cys126Arg)	121917769	UMOD	[]	['GGCCACAYGTGTCAATGTGGTGG', 'GCCACAYGTGTCAATGTGGTGGG']	['Familial juvenile gout']
NM_003361.3(UMOD):c.943T>C (p.Cys315Arg)	121917773	UMOD	[]	['ATGGCACYGCCAGTGCAAACAGG']	['Glomerulocystic kidney disease with hyperuricemia and isosthenuria']
NM_024649.4(BBS1):c.1553T>C (p.Leu518Pro)	121917778	-	[]	[]	['Bardet-Biedl syndrome 1']
NM_172107.2(KCNQ2):c.2T>C (p.Met1Thr)	118192186	KCNQ2	[]	[]	['Benign familial neonatal seizures 1']
NM_014324.5(AMACR):c.154T>C (p.Ser52Pro)	121917814	-	[]	[]	['Alpha-methylacyl-CoA racemase deficiency', 'Bile acid synthesis defect, congenital, 4']
NM_014324.5(AMACR):c.320T>C (p.Leu107Pro)	121917816	-	[]	[]	['Bile acid synthesis defect, congenital, 4']
NM_007255.2(B4GALT7):c.617T>C (p.Leu206Pro)	121917818	B4GALT7	[]	['TGCYCTCCAAGCAGCACTACCGG']	['Ehlers-Danlos syndrome progeroid type']
NM_021615.4(CHST6):c.827T>C (p.Leu276Pro)	121917824	CHST6	[]	['GGACCYGGCGCGGGAGCCGCTGG']	['Macular corneal dystrophy Type I']
NM_006261.4(PROP1):c.263T>C (p.Phe88Ser)	121917841	PROP1	[]	[]	['Pituitary hormone deficiency, combined 2']
NM_000452.2(SLC10A2):c.728T>C (p.Leu243Pro)	121917848	SLC10A2	[]	['TTTCYTCTGGCTAGAATTGCTGG']	['Bile acid malabsorption, primary']
NM_000322.4(PRPH2):c.637T>C (p.Cys213Arg)	61755802	PRPH2	[]	[]	['Patterned dystrophy of retinal pigment epithelium',

					'not provided', 'Leber congenital amaurosis 18']
NM_000517.4(HBA2):c.89T>C (p.Leu30Pro)	41341344	HBA2	[]	[]	['Hemoglobin H disease, nondeletional']
NM_002181.3(IHH):c.569T>C (p.Val190Ala)	121917857	IHH	[]	[]	['Acrocapitofemoral dysplasia']
NM_000322.4(PRPH2):c.736T>C (p.Trp246Arg)	61755817	PRPH2	['ACCTGYGGGTGCGTGGCTGCAGG', 'CCTGYGGGTGCGTGGCTGCAGG']	['ACCTGYGGGTGCGTGGCTGCAGGG']	['Retinitis pigmentosa', 'not provided']
NM_005413.3(SIX3):c.749T>C (p.Val250Ala)	121917880	SIX3	[]	[]	['Holoprosencephaly 2']
NM_000124.3(ERCC6):c.2960T>C (p.Leu987Pro)	121917905	ERCC6	['TGCYAAAAGACCCAAAACAAAGG']	['TGCYAAAAGACCCAAAACAAAGG']	['Cerebro-oculo-facio-skeletal syndrome']
NM_006920.4(SCN1A):c.4729T>C (p.Cys1577Arg)	121917919	-	[]	[]	['Severe myoclonic epilepsy in infancy', 'not provided']
NM_006920.4(SCN1A):c.5113T>C (p.Cys1705Arg)	121917926	-	[]	[]	['Severe myoclonic epilepsy in infancy', 'not provided']
NM_006920.4(SCN1A):c.3577T>C (p.Trp1193Arg)	121917930	-	['AACAAYGGTGGAACCTGAGAAGG']	['AACAAYGGTGGAACCTGAGAAGG']	['Generalized epilepsy with febrile seizures plus, type 1', 'Generalized epilepsy with febrile seizures plus, type 2']
NM_006920.4(SCN1A):c.838T>C (p.Trp280Arg)	121917938	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000478.4(ALPL):c.1306T>C (p.Tyr436His)	121918006	ALPL	[]	['TGGACYATGGTGAGACCTCCAGG']	['Infantile hypophosphatasia']
NM_000478.4(ALPL):c.979T>C (p.Phe327Leu)	121918010	ALPL	[]	['CAAAGGCYTCTTCTTGCTGGTGG', 'GGCYTCTTCTTGCTGGTGGAAGG']	['Infantile hypophosphatasia']
NM_000301.3(PLG):c.1771T>C (p.Ser591Pro)	121918029	PLG	[]	[]	['Dysplasminogenemia']
NM_000174.4(GP9):c.20T>C (p.Leu7Pro)	121918038	GP9	[]	[]	['Bernard-Soulier syndrome type C']
NM_000371.3(TTR):c.224T>C (p.Leu75Pro)	121918079	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis']
NM_000371.3(TTR):c.88T>C (p.Cys30Arg)	121918083	TTR	[]	[]	['Amyloidogenic transthyretin

					amyloidosis', 'Cardiomyopathy']
NM_000371.3(TTR):c.272T>C (p.Val91Ala)	121918084	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis']
NM_000371.3(TTR):c.400T>C (p.Tyr134His)	121918088	TTR	[]	['CCCCYACTCCTATTCCACCACGG']	[]
NM_012275.2(IL36RN):c.115+6T>C	148755083	IL36RN	[]	[]	['Pustular psoriasis, generalized']
NM_000371.3(TTR):c.95T>C (p.Leu32Pro)	121918094	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis']
NM_000371.3(TTR):c.265T>C (p.Tyr89His)	121918100	TTR	[]	[]	['AMYLOIDOSIS, LEPTOMENINGEAL, TRANSTHYRETIN-RELATED']
NM_001042465.1(PSAP):c.1055T>C (p.Leu352Pro)	121918110	PSAP	[]	['GAAGCYGCCGAAGTCCCTGTCGG']	['Gaucher disease, atypical, due to saposin C deficiency']
NM_013251.3(TAC3):c.269T>C (p.Met90Thr)	121918123	TAC3	[]	[]	['not provided']
NM_199069.1(NDUFAF3):c.2T>C (p.Met1Thr)	121918136	NDUFAF3	[]	[]	['Mitochondrial complex I deficiency']
NM_003730.4(RNASET2):c.550T>C (p.Cys184Arg)	121918137	RNASET2	[]	['CCAGYGCCTTCCACCAAGCCAGG']	['Leukoencephalopathy, cystic, without megalencephaly']
NM_203395.2(IYD):c.347T>C (p.Ile116Thr)	121918139	IYD	[]	[]	['Iodotyrosine deiodination defect']
NM_001127628.1(FBP1):c.581T>C (p.Phe194Ser)	121918191	FBP1	[]	['GGAGTYCATTTTGGTGGACAAGG']	['Fructose-biphosphatase deficiency']
NM_015506.2(MMACHC):c.347T>C (p.Leu116Pro)	121918240	MMACHC	[]	[]	['Methylmalonic acidemia with homocystinuria']
NM_000255.3(MUT):c.313T>C (p.Trp105Arg)	121918249	MUT	[]	[]	['METHYLMALONIC ACIDURIA, mut(0) TYPE']
NM_022370.3(ROBO3):c.14T>C (p.Leu5Pro)	121918275	ROBO3	[]	[]	['Gaze palsy, familial horizontal, with progressive scoliosis']
NM_018400.3(SCN3B):c.29T>C (p.Leu10Pro)	121918282	SCN3B	[]	[]	['Brugada syndrome', 'Brugada syndrome 7', 'Cardiac arrhythmia', 'Atrial fibrillation, familial, 16']
NM_004183.3(BEST1):c.122T>C (p.Leu41Pro)	121918288	BEST1	[]	[]	['Bestrophinopathy, autosomal recessive', 'not provided']
NM_020166.4(MCCC1):c.640-2A>G	772395858	MCCC1	[]	[]	['3 Methylcrotonyl-CoA carboxylase 1 deficiency']
NM_004817.3(TJP2):c.143T>C (p.Val48Ala)	121918299	TJP2	[]	[]	['Hypercholanemia, familial']

NM_006946.2(SPTBN2):c.758T>C (p.Leu253Pro)	121918306	SPTBN2	[]	['ACCAAGCYGCTGGATCCCGAAGG', 'AAGCYGCTGGATCCCGAAGGTGG', 'AGCYGCTGGATCCCGAAGGTGGG']	['Spinocerebellar ataxia 5']
NM_000214.2(JAG1):c.110T>C (p.Leu37Ser)	121918352	JAG1	[]	[]	['Alagille syndrome 1']
NM_007194.3(CHEK2):c.470T>C (p.Ile157Thr)	17879961	CHEK2	[]	[]	['Familial cancer of breast', 'Hereditary cancer-predisposing syndrome', 'Li-Fraumeni syndrome 2', 'not specified']
NM_015384.4(NIPBL):c.7637T>C (p.Leu2546Pro)	727503772	NIPBL	[]	[]	['Cornelia de Lange syndrome 1']
NM_001083112.2(GPD2):c.1904T>C (p.Phe635Ser)	121918407	GPD2	['AAGTYTGATGCAGACCAGAAAGG']	['AAGTYTGATGCAGACCAGAAAGG']	['Diabetes mellitus type 2']
NM_021957.3(GYS2):c.1447T>C (p.Ser483Pro)	121918424	GYS2	[]	[]	['Hypoglycemia with deficiency of glycogen synthetase in the liver']
NM_018849.2(ABCB4):c.1207T>C (p.Tyr403His)	121918443	ABCB4	[]	[]	['Progressive familial intrahepatic cholestasis 3']
NM_000212.2(ITGB3):c.2332T>C (p.Ser778Pro)	121918447	-	[]	[]	['Glanzmann thrombasthenia']
NM_000506.3(F2):c.1139T>C (p.Met380Thr)	121918481	F2	[]	[]	['Hereditary factor II deficiency disease']
NM_000141.4(FGFR2):c.1018T>C (p.Tyr340His)	121918489	FGFR2	['TGGGGAAYATACGTGCTTGGCGG', 'GGGGAAYATACGTGCTTGGCGGG']	['TGGGGAAYATACGTGCTTGGCGG', 'GGGGAAYATACGTGCTTGGCGGG']	['Crouzon syndrome']
NM_000141.4(FGFR2):c.799T>C (p.Ser267Pro)	121918505	FGFR2	[]	['AATGCCYCCACAGTGGTCGGAGG']	['Pfeiffer syndrome', 'Neoplasm of stomach']
NM_002739.3(PRKCG):c.355T>C (p.Ser119Pro)	121918512	PRKCG	[]	[]	['Spinocerebellar ataxia 14']
NM_002739.3(PRKCG):c.1927T>C (p.Phe643Leu)	121918516	PRKCG	[]	[]	['Spinocerebellar ataxia 14']
NM_015107.2(PHF8):c.836T>C (p.Phe279Ser)	121918524	PHF8	[]	[]	['Siderius X-linked mental retardation syndrome']
NM_000311.3(PRNP):c.593T>C	74315405	PRNP	[]	[]	['Gerstmann-Straussler-Scheinker

(p.Phe198Ser)					syndrome', 'Genetic prion diseases']
NM_015665.5(AAAS):c.787T>C (p.Ser263Pro)	121918550	AAAS	[]	[]	['Glucocorticoid deficiency with achalasia']
NM_003018.3(SFTPC):c.581T>C (p.Leu194Pro)	121918560	SFTPC	[]	[]	['Surfactant metabolism dysfunction, pulmonary, 2']
NM_000322.4(PRPH2):c.554T>C (p.Leu185Pro)	121918563	PRPH2	[]	[]	['Patterned dystrophy of retinal pigment epithelium', 'Retinitis pigmentosa 7, digenic', 'not provided', 'Leber congenital amaurosis 18']
NM_000322.4(PRPH2):c.2T>C (p.Met1Thr)	121918565	PRPH2	[]	[]	['Macular dystrophy, vitelliform, adult-onset', 'not provided']
NM_001035.2(RYR2):c.1298T>C (p.Leu433Pro)	121918602	RYR2	[]	[]	['Arrhythmogenic right ventricular cardiomyopathy, type 2', 'Catecholaminergic polymorphic ventricular tachycardia', 'Long QT syndrome']
NM_001199138.1(NLRC4):c.1022T>C (p.Val341Ala)	587781260	NLRC4	[]	[]	['Syndrome of entercolitis and autoinflmmation caused by mutation of NLRC4 (SCAN4)', 'Autoinflammation with infantile enterocolitis']
NM_000702.3(ATP1A2):c.857T>C (p.Ile286Thr)	121918617	ATP1A2	[]	[]	['Familial hemiplegic migraine type 2']
NM_006920.4(SCN1A):c.4250T>C (p.Val1417Ala)	121918627	-	[]	[]	['Generalized epilepsy with febrile seizures plus, type 1', 'Generalized epilepsy with febrile seizures plus, type 2']
NM_006920.4(SCN1A):c.434T>C (p.Met145Thr)	121918631	SCN1A	[]	[]	['Generalized epilepsy with febrile seizures plus, type 2']
NM_006920.4(SCN1A):c.4462T>C (p.Phe1488Leu)	121918632	-	[]	[]	['Familial hemiplegic migraine type 3']
NM_003126.2(SPTA1):c.781T>C (p.Ser261Pro)	121918636	SPTA1	[]	[]	['Elliptocytosis 2']

NM_003126.2(SPTA1):c.620T>C (p.Leu207Pro)	121918643	SPTA1	[]	['GTGGAGCYGGTAGCTAAAGAAGG', 'TGGAGCYGGTAGCTAAAGAAGGG']	['Hereditary pyropoikilocytosis', 'Elliptocytosis 2']
NM_001024858.2(SPTB):c.604T>C (p.Trp202Arg)	121918646	SPTB	[]	['CTCCAGCYGGAAGGATGGCTTGG']	['Spherocytosis type 2']
NM_001024858.2(SPTB):c.6055T>C (p.Ser2019Pro)	121918648	SPTB	[]	['ATGCCYCTGTGGCTGAGGCGTGG']	[]
NM_001128177.1(THRB):c.929T>C (p.Met310Thr)	121918699	THRB	[]	[]	[]
NM_001128177.1(THRB):c.1373T>C (p.Val458Ala)	121918704	THRB	[]	[]	['Thyroid hormone resistance, generalized, autosomal recessive']
NM_000421.3(KRT10):c.482T>C (p.Leu161Ser)	60118264	-	[]	[]	['Bullous ichthyosiform erythroderma', 'not provided']
NM_006920.4(SCN1A):c.269T>C (p.Phe90Ser)	121918733	SCN1A	['ACTTYTATAGTATTGAATAAAGG', 'CTTYTATAGTATTGAATAAAGGG']	['ACTTYTATAGTATTGAATAAAGG', 'CTTYTATAGTATTGAATAAAGGG']	['Severe myoclonic epilepsy in infancy']
NM_006920.4(SCN1A):c.272T>C (p.Ile91Thr)	121918734	SCN1A	['ACTTTTAYAGTATTGAATAAAGG', 'CTTTTAYAGTATTGAATAAAGGG']	['ACTTTTAYAGTATTGAATAAAGG', 'CTTTTAYAGTATTGAATAAAGGG']	['Severe myoclonic epilepsy in infancy']
NM_006920.4(SCN1A):c.3827T>C (p.Leu1276Pro)	121918740	-	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000543.4(SMPD1):c.475T>C (p.Cys159Arg)	727504166	SMPD1	[]	['TGAGGCCYGTGGCCTGCTCCTGG', 'GAGGCCYGTGGCCTGCTCCTGGG']	['Niemann-Pick disease, type A', 'Niemann-Pick disease, type B']
NM_006920.4(SCN1A):c.568T>C (p.Trp190Arg)	121918773	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_006920.4(SCN1A):c.5522T>C (p.Met1841Thr)	121918783	-	[]	[]	['Severe myoclonic epilepsy in infancy', 'Generalized epilepsy with febrile seizures plus, type 1']

NM_002538.3(OCLN):c.656T>C (p.Phe219Ser)	267606926	OCLN	[]	[]	['Band-like calcification with simplified gyration and polymicrogyria']
NM_000434.3(NEU1):c.1088T>C (p.Leu363Pro)	193922915	NEU1	[]	['CAGCYATGGCCAGGCCCCAGTGG']	['Sialidosis, type II']
NM_198578.3(LRRK2):c.6059T>C (p.Ile2020Thr)	35870237	LRRK2	[]	[]	['Parkinson disease 8, autosomal dominant']
NM_000501.3(ELN):c.889+2T>C	727504419	ELN	[]	['CAGGYAACATCTGTCCCAGCAGG', 'AGGYAACATCTGTCCCAGCAGGG']	['Supravalvar aortic stenosis']
NM_001085.4(SERPINA3):c.233T>C (p.Leu78Pro)	1800463	SERPINA3	[]	[]	[]
NM_000238.3(KCNH2):c.1736T>C (p.Met579Thr)	199473425	KCNH2			['Congenital long QT syndrome']
NM_001211.5(BUB1B):c.3035T>C (p.Leu1012Pro)	28989185	-	[]	[]	['Mosaic variegated aneuploidy syndrome', 'Premature chromatid separation trait']
NM_000256.3(MYBPC3):c.26-2A>G	376395543	MYBPC3	[]	['GAGACYGAAGGGCCAGGTGGAGG']	['Primary familial hypertrophic cardiomyopathy', 'Familial hypertrophic cardiomyopathy 4', 'Cardiomyopathy']
NM_000051.3(ATM):c.4776+2T>C	587781927	ATM	[]	[]	['Ataxia-telangiectasia syndrome', 'Hereditary cancer-predisposing syndrome']
NM_006412.3(AGPAT2):c.589-2A>G	116807569	AGPAT2	[]	[]	['Congenital generalized lipodystrophy type 1']
NM_000545.6(HNF1A):c.1720G>A (p.Gly574Ser)	1169305	HNF1A	[]	['GATGCYGGCAGGGTCCTGGCTGG', 'ATGCYGGCAGGGTCCTGGCTGGG', 'TGCYGGCAGGGTCCTGGCTGGGG']	['Maturity-onset diabetes of the young, type 3']
NM_024514.4(CYP2R1):c.296T>C (p.Leu99Pro)	61495246	CYP2R1	[]	[]	['Vitamin d hydroxylation-deficient rickets, type 1b']
NM_012213.2(MLYCD):c.119T>C (p.Met40Thr)	28937908	MLYCD	[]	[]	['Deficiency of malonyl-CoA decarboxylase']
NM_001101.3(ACTB):c.224T>C (p.Ile75Thr)	587779771	ACTB	[]	[]	['Baraitser-Winter syndrome 1']

NM_021007.2(SCN2A):c.1271T>C (p.Val424Ala)	796053181	SCN2A	['TGTGGYGGCCATGGCCTATGAGG']	['TGTGGYGGCCATGGCCTATGAGG']	['not provided']
NM_002880.3(RAF1):c.769T>C (p.Ser257Pro)	727505017	RAF1	[]	[]	['Rasopathy', 'not specified']
NM_000527.4(LDLR):c.1468T>C (p.Trp490Arg)	730880130	LDLR	[]	['CTACYGGACCGACTCTGTCCTGG', 'TACYGGACCGACTCTGTCCTGGG']	['Familial hypercholesterolemia']
NM_170707.3(LMNA):c.710T>C (p.Phe237Ser)	730880132	LMNA	['TGAGTYTGAGAGCCGGCTGGCGG']	['TGAGTYTGAGAGCCGGCTGGCGG']	['Primary dilated cardiomyopathy']
NM_000080.3(CHRNE):c.223T>C (p.Trp75Arg)	193919341	-	[]	[]	['MYASTHENIC SYNDROME, CONGENITAL, 4B, FAST-CHANNEL']
NM_005211.3(CSF1R):c.2297T>C (p.Met766Thr)	281860270	CSF1R	[]	[]	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_005211.3(CSF1R):c.2381T>C (p.Ile794Thr)	281860274	CSF1R	['CAAGAYTGGGGACTTCGGGCTGG']	['CAAGAYTGGGGACTTCGGGCTGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_005211.3(CSF1R):c.2546T>C (p.Phe849Ser)	281860277	CSF1R	[]	[]	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_005211.3(CSF1R):c.2624T>C (p.Met875Thr)	281860279	CSF1R	[]	[]	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_018713.2(SLC30A10):c.266T>C (p.Leu89Pro)	281860284	SLC30A10	[]	[]	['Hypermanganesemia with dystonia, polycythemia and cirrhosis']
NM_018713.2(SLC30A10):c.500T>C (p.Phe167Ser)	281860286	SLC30A10	[]	['GGCGCTTYCGGGGGGCCTCAGGG']	['Hypermanganesemia with dystonia, polycythemia and cirrhosis']
NM_018713.2(SLC30A10):c.1046T>C (p.Leu349Pro)	281860291	SLC30A10	[]	[]	['Hypermanganesemia with dystonia, polycythemia and cirrhosis']
NM_016218.2(POLK):c.2287T>A (p.Tyr763Asn)	772307321	POLK	[]	[]	['Malignant tumor of prostate']
NM_000570.4(FCGR3B):c.194A>G (p.Asn65Ser)	448740	FCGR3B	[]	[]	[]
NM_030653.3(DDX11):c.2271+2T>C	730880279	DDX11	['TCCAGGYGCGGGCGTCATGCTGG']	['TCCAGGYGCGGGCGTCATGCTGG', 'CCAGGYGCGGGCGTCATGCTGGG']	['Warsaw breakage syndrome']
NM_145693.2(LPIN1)	730880306	LPIN1	[]	['AAGGYACCGCG	['Myoglobinuria,

:c.1441+2T>C				GGCCTCGCGCGG', 'AGGYACCGCGGG CCTCGCGCGGG']	acute recurrent, autosomal recessive']
NM_002546.3(TNFRSF11B):c.226A>C (p.Thr76Pro)	200071478	TNFRSF11B	[]	[]	['Hyperphosphatasemia with bone disease']
NM_000166.5(GJB1): c.145T>C (p.Ser49Pro)	116840817	GJB1	[]	[]	['X-linked hereditary motor and sensory neuropathy']
NM_005159.4(ACTC1):c.755T>C (p.Ile252Thr)	730880398	-	[]	[]	['Cardiomyopathy']
NM_020166.4(MCCC1):c.205A>T (p.Lys69Ter)	147741073	MCCC1	[]	[]	['3 Methylcrotonyl-CoA carboxylase 1 deficiency']
NM_000454.4(SOD1): c.338T>C (p.Ile113Thr)	74315452	SOD1	[]	['TTGCAYCATTGG CCGCACACTGG']	['Amyotrophic lateral sclerosis type 1']
NM_000169.2(GLA):c.41T>C (p.Leu14Pro)	730880455	-	[]	['CGCGCYTGCGCT TCGCTTCCTGG']	['not provided']
NM_152743.3(BRAT1):c.176T>C (p.Leu59Pro)	727505363	BRAT1	[]	[]	['Rigidity and multifocal seizure syndrome, lethal neonatal']
NM_005633.3(SOS1): c.2104T>C (p.Tyr702His)	727505381	SOS1	[]	[]	['Noonan syndrome', 'Rasopathy']
m.1095T>C	267606618	MT-RNR1	[]	[]	['Aminoglycoside-induced deafness', 'Auditory neuropathy', 'Deafness, nonsyndromic sensorineural, mitochondrial', 'not specified']
m.1291T>C	267606620	MT-RNR1	[]	[]	['Deafness, nonsyndromic sensorineural, mitochondrial']
NM_020247.4(ADCK3):c.1398+2T>C	606231138	ADCK3	[]	[]	['Coenzyme Q10 deficiency, primary, 4']
NM_000256.3(MYBPC3):c.467T>C (p.Leu156Pro)	730880616	MYBPC3	[]	[]	['Cardiomyopathy']
NM_022458.3(LMBR1):c.423+4842T>C	606231149	LMBR1	[]	[]	['Triphalangeal thumb', 'Preaxial polydactyly 2']
NM_022458.3(LMBR1):c.423+4808T>C	606231152	LMBR1	[]	[]	['Triphalangeal thumb', 'Preaxial polydactyly 2']
NM_021102.3(SPINT2):c.337+2T>C	606231155	SPINT2	[]	[]	['Diarrhea 3, secretory sodium, congenital, syndromic']
NM_001004127.2(ALG11):c.257T>C (p.Leu86Ser)	267606652	ALG11	[]	[]	['Congenital disorder of glycosylation type 1P']

NM_054027.4(ANKH):c.1015T>C (p.Cys339Arg)	267606656	ANKH	[]	['AGCTCYGTTTCGTGATGTTTTGG']	['Craniometaphyseal dysplasia, autosomal dominant']
NM_054027.4(ANKH):c.1172T>C (p.Leu391Pro)	267606658	-	[]	[]	['Craniometaphyseal dysplasia, autosomal dominant']
NM_175073.2(APTX):c.668T>C (p.Leu223Pro)	267606665	APTX	[]	[]	['Adult onset ataxia with oculomotor apraxia']
NM_004183.3(BEST1):c.704T>C (p.Val235Ala)	267606679	BEST1	['CACTGGYGTATACACAGGTGAGG']	['CACTGGYGTATACACAGGTGAGG']	['Vitreoretinochoroidopathy dominant']
NM_004183.3(BEST1):c.614T>C (p.Ile205Thr)	267606680	BEST1	[]	[]	['Retinitis pigmentosa 50']
NM_033409.3(SLC52A3):c.670T>C (p.Phe224Leu)	267606685	SLC52A3	[]	[]	['Brown-Vialetto-Van laere syndrome']
NM_033409.3(SLC52A3):c.1238T>C (p.Val413Ala)	267606687	SLC52A3	[]	['AGTTACGYCAAGGTGATGCTGGG']	['Brown-Vialetto-Van laere syndrome']
NM_004056.4(CA8):c.298T>C (p.Ser100Pro)	267606695	CA8	[]	[]	['Cerebellar ataxia, mental retardation, and dysequilibrium syndrome 3']
NM_000256.3(MYBPC3):c.3713T>C (p.Leu1238Pro)	730880702	MYBPC3	[]	[]	['Cardiomyopathy']
NM_001928.2(CFD):c.640T>C (p.Cys214Arg)	267606721	CFD	[]	['GGTGYGCGGGGGCGTGCTCGAGG', 'GTGYGCGGGGGCGTGCTCGAGGG']	['Complement factor d deficiency']
NM_005740.2(DNAL4):c.153+2T>C	606231254	DNAL4	['CGAGGYATTGCCAGCAGTGCAGG']	['CGAGGYATTGCCAGCAGTGCAGG']	['Mirror movements 3']
NM_020975.4(RET):c.2753T>C (p.Met918Thr)	74799832	RET	[]	[]	['Multiple endocrine neoplasia, type 2a', 'Multiple endocrine neoplasia, type 2b', 'Multiple endocrine neoplasia, type 2', 'Pheochromocytoma', 'not provided']
NM_001849.3(COL6A2):c.2329T>C (p.Cys777Arg)	267606747	COL6A2	[]	['CGCCYGCGACAAGCCACAGCAGG']	['Ullrich congenital muscular dystrophy']
NM_001006657.1(WDR35):c.781T>C (p.Trp261Arg)	431905505	WDR35	[]	[]	['Short rib polydactyly syndrome 5']
NM_003764.3(STX11):c.173T>C (p.Leu58Pro)	431905512	STX11	['TGCYGGTGGCCGACGTGAAGCGG']	['TGCYGGTGGCCGACGTGAAGCGG']	['Hemophagocytic lymphohistiocytosis, familial, 4']
NM_001044.4(SLC6A	431905515	SLC6A3	[]	['CTGCACCYCCAC	['Infantile

3):c.671T>C (p.Leu224Pro)				CAGAGCCATGG']	Parkinsonism-dystonia']
NM_000277.1(PAH):c.764T>C (p.Leu255Ser)	62642930	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_000277.1(PAH):c.932T>C (p.Leu311Pro)	62642936	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_000118.3(ENG):c.2T>C (p.Met1Thr)	267606783	ENG	[]	[]	['Osler hemorrhagic telangiectasia syndrome']
NM_000129.3(F13A1):c.728T>C (p.Met243Thr)	267606788	F13A1	['TGTGAYGGACAGAGCACAAATGG']	['TGTGAYGGACAGAGCACAAATGG']	['Factor xiii, a subunit, deficiency of']
NM_000257.3(MYH7):c.1952A>G (p.His651Arg)	606231328	MYH7	[]	[]	['Familial cardiomyopathy']
NM_014053.3(FLVCR1):c.574T>C (p.Cys192Arg)	267606821	FLVCR1	[]	[]	['Posterior column ataxia with retinitis pigmentosa']
NM_005249.4(FOXG1):c.643T>C (p.Phe215Leu)	267606828	FOXG1	[]	[]	['Rett syndrome, congenital variant']
NM_015474.3(SAMHD1):c.1153A>G (p.Met385Val)	515726140	SAMHD1	[]	[]	['Aicardi Goutieres syndrome 5']
NM_015474.3(SAMHD1):c.1411-2A>G	515726141	SAMHD1	[]	[]	['Aicardi Goutieres syndrome 5']
NM_000180.3(GUCY2D):c.2846T>C (p.Ile949Thr)	267606857	GUCY2D	[]	['AGAGAYCGCCAACATGTCACTGG']	['Cone-rod dystrophy 6']
NM_022489.3(INF2):c.556T>C (p.Ser186Pro)	267606877	INF2	[]	[]	['Focal segmental glomerulosclerosis 5']
NM_000257.3(MYH7):c.1048T>C (p.Tyr350His)	730880863	MYH7	[]	[]	['Cardiomyopathy']
NM_022489.3(INF2):c.125T>C (p.Leu42Pro)	267606880	INF2	[]	['GCTGCYCCAGATGCCCTCTGTGG']	['Focal segmental glomerulosclerosis 5']
m.4681T>C	267606889	MT-ND2	[]	[]	['Leigh disease', 'Leigh syndrome due to mitochondrial complex I deficiency']
m.10191T>C	267606890	MT-ND3	[]	[]	['Leigh disease', 'Mitochondrial complex I deficiency']
m.10563T>C	267606892	MT-ND4L	[]	[]	['Familial colorectal cancer']
m.12706T>C	267606893	MT-ND5	[]	[]	['Leigh disease', 'Leigh syndrome due to mitochondrial complex I deficiency']
NM_015713.4(RRM2	515726190	RRM2B	[]	[]	['RRM2B-related

B):c.556A>G (p.Arg186Gly)					mitochondrial disease']
NM_015713.4(RRM2B):c.581A>G (p.Glu194Gly)	515726191	RRM2B	[]	['AACTCCTYCTACAGCAGCAAAGG']	['RRM2B-related mitochondrial disease']
NM_000315.2(PTH):c.52T>C (p.Cys18Arg)	104894271	PTH	['AATTYGTTTTCTTACAAAATCGG']	['AATTYGTTTTCTTACAAAATCGG']	['Hypoparathyroidism familial isolated']
NM_001136271.2(NKX2-6):c.451T>C (p.Phe151Leu)	267606914	NKX2-6	[]	[]	['Persistent truncus arteriosus']
NM_004646.3(NPHS1):c.793T>C (p.Cys265Arg)	267606917	NPHS1	[]	['GCTGCCGYGCGTGGCCCGAGGGG', 'CTGCCGYGCGTGGCCCGAGGGGG']	['Finnish congenital nephrotic syndrome']
NM_000406.2(GNRHR):c.94A>G (p.Thr32Ala)	515726219	GNRHR	[]	[]	['Hypogonadotropic hypogonadism']
NM_152296.4(ATP1A3):c.2270T>C (p.Leu757Pro)	606231436	ATP1A3	[]	[]	['Alternating hemiplegia of childhood 2']
NM_000513.2(OPN1MW):c.529T>C (p.Trp177Arg)	267606927	OPN1MW	[]	[]	['Cone dystrophy 5, X-linked']
NM_152296.4(ATP1A3):c.1144T>C (p.Trp382Arg)	606231448	ATP1A3	[]	[]	['Dystonia 12']
NM_024411.4(PDYN):c.632T>C (p.Leu211Ser)	267606940	PDYN	[]	[]	['Spinocerebellar ataxia 23']
NM_000444.5(PHEX):c.755T>C (p.Phe252Ser)	267606945	PHEX	[]	[]	['Familial X-linked hypophosphatemic vitamin D refractory rickets']
NM_001543.4(NDST1):c.1918T>C (p.Phe640Leu)	606231458	NDST1	[]	[]	['Mental retardation, autosomal recessive 46']
NM_013382.5(POMT2):c.2242T>C (p.Trp748Arg)	267606964	POMT2	[]	[]	['Congenital muscular dystrophy-dystroglycanopathy with mental retardation, type B2']
NM_006121.3(KRT1):c.482T>C (p.Leu161Pro)	57695159	KRT1	[]	[]	['Bullous ichthyosiform erythroderma', 'not provided']
NM_016203.3(PRKAG2):c.1459T>C (p.Tyr487His)	267606976	PRKAG2	[]	[]	['Familial hypertrophic cardiomyopathy 6', 'not provided']
NM_016203.3(PRKAG2):c.1642T>C (p.Ser548Pro)	267606979	PRKAG2	[]	[]	['Familial hypertrophic cardiomyopathy 6', 'not provided']
NM_198965.1(PTHLH):c.179T>C	267606985	PTHLH	[]	[]	['Brachydactyly type E2']

(p.Leu60Pro)					
NM_198965.1(PTHLH):c.131T>C (p.Leu44Pro)	267606986	PTHLH	[]	[]	['Brachydactyly type E2']
NM_004990.3(MARS):c.1108T>C (p.Phe370Leu)	140467171	MARS	[]	[]	['Interstitial lung and liver disease']
NM_173560.3(RFX6):c.649T>C (p.Ser217Pro)	267607012	RFX6	[]	[]	['Mitchell-Riley syndrome']
NM_002942.4(ROBO2):c.2834T>C (p.Ile945Thr)	267607014	ROBO2	['GAGAYTGGAAATTTTGGCCGTGG']	['GAGAYTGGAAATTTTGGCCGTGG']	['Vesicoureteral reflux 2']
NM_178857.5(RP1L1):c.2878T>C (p.Trp960Arg)	267607018	RP1L1	[]	[]	['Occult macular dystrophy']
NM_002880.3(RAF1):c.1423T>C (p.Phe475Leu)	730881003	RAF1	[]	[]	['Rasopathy']
NM_015272.3(RPGRIP1L):c.1975T>C (p.Ser659Pro)	267607020	RPGRIP1L	[]	[]	['Joubert syndrome 7', 'COACH syndrome']
NM_015559.2(SETBP1):c.2612T>C (p.Ile871Thr)	267607038	SETBP1	[]	[]	['Schinzel-Giedion syndrome']
NM_000433.3(NCF2):c.481A>G (p.Lys161Glu)	137878529	NCF2	[]	[]	['Chronic granulomatous disease, autosomal recessive cytochrome b-positive, type 2']
NM_001041.3(SI):c.1022T>C (p.Leu341Pro)	267607049	SI	[]	[]	['Sucrase-isomaltase deficiency']
NM_005633.3(SOS1):c.1294T>C (p.Trp432Arg)	267607080	SOS1	['GGTYGGGAGGGAAAGACATTGG']	['GGTYGGGAGGGAAAAGACATTGG']	['Noonan syndrome 4', 'Rasopathy']
NM_018136.4(ASPM):c.2419+2T>C	587783225	ASPM	[]	[]	['Primary autosomal recessive microcephaly 5']
NM_001199107.1(TBC1D24):c.751T>C (p.Phe251Leu)	267607104	TBC1D24	[]	['CAAGTTCYTCCACAAGGTGAGGG', 'TTCYTCCACAAGGTGAGGGCCGG']	['Myoclonic epilepsy, familial infantile']
NM_153704.5(TMEM67):c.1769T>C (p.Phe590Ser)	267607115	TMEM67	[]	[]	['Joubert syndrome 6', 'COACH syndrome']
NM_153704.5(TMEM67):c.2498T>C (p.Ile833Thr)	267607119	TMEM67	[]	[]	['Joubert syndrome 6', 'COACH syndrome']
NM_133378.4(TTN):c.100163T>C (p.Leu33388Pro)	267607156	-	[]	[]	['Distal myopathy Markesbery-Griggs type']
m.12811T>C	199974018	MT-ND5	[]	[]	['Leber optic atrophy']
NM_144631.5(ZNF51	267607182	ZNF513	[]	['TGGGCGCYGCAT	['Retinitis

3):c.1015T>C (p.Cys339Arg)				GCGAGGAGAGG', 'CGCYGCATGCGAGGAGAGGCTGG']	pigmentosa 58']
NM_004737.4(LARGE):c.1483T>C (p.Trp495Arg)	267607209	LARGE	[]	[]	['Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies, type A6']
NM_000229.1(LCAT):c.508T>C (p.Trp170Arg)	267607211	LCAT	[]	['TATGACYGGCGGCTGGAGCCCGG']	['Norum disease']
NM_016269.4(LEF1):c.181T>C (p.Ser61Pro)	267607215	-	[]	['GAACGAGYCTGAAATCATCCCGG']	['Sebaceous tumors, somatic']
NM_139248.2(LIPH):c.322T>C (p.Trp108Arg)	267607219	LIPH	[]	[]	['Woolly hair, autosomal recessive 2, with or without hypotrichosis']
NM_004268.4(MED17):c.1112T>C (p.Leu371Pro)	267607232	MED17	[]	[]	['Microcephaly, postnatal progressive, with seizures and brain atrophy']
NM_000530.6(MPZ):c.341T>C (p.Ile114Thr)	267607241	MPZ	[]	[]	[]
NM_000489.4(ATRX):c.4840T>C (p.Cys1614Arg)	122445094	ATRX	[]	[]	['ATR-X syndrome']
NM_000489.4(ATRX):c.6250T>C (p.Tyr2084His)	122445097	ATRX	[]	[]	['ATR-X syndrome']
NM_000489.4(ATRX):c.1226T>C (p.Leu409Ser)	122445109	ATRX	[]	[]	[]
NM_000489.4(ATRX):c.6149T>C (p.Ile2050Thr)	122445110	ATRX	[]	[]	['Multiple congenital anomalies']
NM_178151.2(DCX):c.272T>C (p.Leu91Pro)	587783536	DCX	[]	[]	['Heterotopia']
NM_178151.2(DCX):c.2T>C (p.Met1Thr)	587783539	DCX	['CAAAATAYGGAACTTGATTTTGG']	['CAAAATAYGGAACTTGATTTTGG']	['Heterotopia']
NM_178151.2(DCX):c.412T>C (p.Tyr138His)	587783551	DCX	[]	[]	['Heterotopia']
NM_000212.2(ITGB3):c.2231T>C (p.Leu744Pro)	398122374	-	[]	[]	['Platelet-type bleeding disorder 16']
NM_178151.2(DCX):c.641T>C (p.Ile214Thr)	587783574	DCX	[]	[]	['Heterotopia']
NM_178151.2(DCX):c.683T>C (p.Leu228Pro)	587783580	DCX	[]	['AAAAAACYCTACACTCTGGATGG']	['Heterotopia']
NM_001005360.2(DNM2):c.1862T>C (p.Leu621Pro)	587783597	DNM2	[]	[]	['Myopathy, centronuclear']
NM_006579.2(EBP):c.310T>C (p.Tyr104His)	587783609	EBP	[]	[]	['Chondrodysplasia punctata 2 X-linked

					dominant']
NM_004004.5(GJB2):c.107T>C (p.Leu36Pro)	587783644	GJB2	[]	['GATCCYCGTTGTGGCTGCAAAGG']	['Hearing impairment']
NM_005682.6(ADGRG1):c.1460T>C (p.Leu487Pro)	587783653	ADGRG1	[]	['CCCTGCYCACCTGCCTTTCCTGG']	['Polymicrogyria, bilateral frontoparietal']
NM_000525.3(KCNJ11):c.988T>C (p.Tyr330His)	587783675	KCNJ11	[]	[]	['Diabetes mellitus']
NM_170707.3(LMNA):c.799T>C (p.Tyr267His)	267607593	LMNA	[]	[]	['Dilated cardiomyopathy 1A', 'not provided']
NM_000252.2(MTM1):c.1353+2T>C	587783780	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.1367T>C (p.Phe456Ser)	587783783	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.1433T>C (p.Phe478Ser)	587783794	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.1495T>C (p.Trp499Arg)	587783801	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.260T>C (p.Leu87Pro)	587783816	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.683T>C (p.Leu228Pro)	587783851	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.958T>C (p.Ser320Pro)	587783863	MTM1	[]	['GGAAYCTTTAAAAAAAGTGAAGG']	['Severe X-linked myotubular myopathy']
NM_000526.4(KRT14):c.1151T>C (p.Leu384Pro)	59629244	KRT14	[]	[]	['Epidermolysis bullosa simplex, Koebner type', 'not provided']
NM_000249.3(MLH1):c.453+2T>C	267607751	MLH1	[]	['ATCACGGYAAGAATGGTACATGG', 'TCACGGYAAGAATGGTACATGGG']	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_002764.3(PRPS1):c.455T>C (p.Leu152Pro)	80338676	PRPS1	[]	[]	['Arts syndrome', 'not provided']
NM_022132.4(MCCC2):c.499T>C (p.Cys167Arg)	119103222	MCCC2	[]	[]	['3-methylcrotonyl CoA carboxylase 2 deficiency']
NM_000411.6(HLCS):c.710T>C (p.Leu237Pro)	119103227	HLCS	[]	['CTATCYTTCTCAGGGAGGGAAGG']	['Holocarboxylase synthetase deficiency']
NM_005787.5(ALG3):c.211T>C (p.Trp71Arg)	119103237	ALG3	[]	['GATTGACYGGAAGGCCTACATGG']	['Congenital disorder of glycosylation type 1D']
NM_005609.2(PYGM):c.1187T>C (p.Leu396Pro)	119103254	PYGM	[]	[]	['Glycogen storage disease, type V']
m.3250T>C	199474664	MT-TL1	[]	[]	[]

NM_002764.3(PRPS1):c.344T>C (p.Met115Thr)	80338732	PRPS1	['GCAAATAYGCTATCTGTAGCAGG']	['GCAAATAYGCTATCTGTAGCAGG']	['Charcot-Marie-Tooth disease, X-linked recessive, type 5']
NM_003172.3(SURF1):c.679T>C (p.Trp227Arg)	398122806	SURF1	[]	['CCACYGGCATTATCGAGACCTGG']	['Congenital myasthenic syndrome, acetazolamide-responsive']
NM_004525.2(LRP2):c.7564T>C (p.Tyr2522His)	80338747	LRP2	[]	['GTACCTGYACTGGGCTGACTGGG']	['Donnai Barrow syndrome']
NM_006329.3(FBLN5):c.649T>C (p.Cys217Arg)	80338766	FBLN5	[]	[]	['Autosomal recessive cutis laxa type IA']
NM_001271723.1(FBXO38):c.616T>C (p.Cys206Arg)	398122838	FBXO38	[]	['TTCCTYGTATCCCAATGCTAAGG']	['Distal hereditary motor neuronopathy 2D']
NM_133433.3(NIPBL):c.7062+2T>C	587784032	NIPBL	[]	[]	['Cornelia de Lange syndrome 1']
NM_000334.4(SCN4A):c.4468T>C (p.Phe1490Leu)	80338790	SCN4A	[]	[]	['Hyperkalemic Periodic Paralysis Type 1']
NM_058216.2(RAD51C):c.404+2T>C	730881931	RAD51C	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_001111.4(ADAR):c.2615T>C (p.Ile872Thr)	398122897	ADAR	[]	[]	['Aicardi-goutieres syndrome 6']
NM_005334.2(HCFC1):c.-970T>C	398122908	HCFC1	['CAAGAYGGCGGCTCCCAGGGAGG']	['CAAGAYGGCGGCTCCCAGGGAGG']	['Mental retardation 3, X-linked']
NM_000431.3(MVK):c.1039+2T>C	398122910	MVK	['CCAGGYATCCCGGGGGTAGGTGG']	['CCAGGYATCCCGGGGGTAGGTGG', 'CAGGYATCCCGGGGGTAGGTGGG']	['Porokeratosis, disseminated superficial actinic 1']
NM_000431.3(MVK):c.1094T>C (p.Phe365Ser)	398122911	MVK	[]	[]	['Porokeratosis, disseminated superficial actinic 1']
NM_005050.3(ABCD4):c.956A>G (p.Tyr319Cys)	201777056	ABCD4	['GATGAGGYAGATGCACACAAAGG']	['GATGAGGYAGATGCACACAAAGG']	['METHYLMALONIC ACIDURIA AND HOMOCYSTINURIA, cblJ TYPE', 'not provided']
NM_000251.2(MSH2):c.2005+2T>C	267607987	MSH2	['CTGGYAAAAAACCTGGTTTTTGG', 'TGGYAAAAAACCTGGTTTTTGGG']	['CTGGYAAAAAACCTGGTTTTTGG', 'TGGYAAAAAACCTGGTTTTTGGG']	['Hereditary Nonpolyposis Colorectal Neoplasms']

NM_000518.4(HBB):c.257T>C (p.Phe86Ser)	35693898	HBB	[]	[]	['Hemoglobinopathy']
NM_001194998.1(CEP152):c.3149T>C (p.Leu1050Pro)	398122977	CEP152	[]	[]	['Primary autosomal recessive microcephaly 9']
NM_022455.4(NSD1):c.5989T>C (p.Tyr1997His)	587784171	NSD1	[]	[]	['Sotos syndrome 1']
NM_014495.3(ANGPTL3):c.883T>C (p.Phe295Leu)	398122989	-	[]	['ACAAAACYTCAATGAAACGTGGG']	['Hypobetalipoproteinemia, familial, 2']
NM_024577.3(SH3TC2):c.1982T>C (p.Leu661Pro)	80338927	SH3TC2	[]	[]	['Charcot-Marie-Tooth disease, type 4C']
NM_000551.3(VHL):c.227T>C (p.Phe76Ser)	730882033	VHL	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_004004.5(GJB2):c.269T>C (p.Leu90Pro)	80338945	GJB2	[]	['GCTCCYAGTGGCCATGCACGTGG']	['Deafness, autosomal recessive 1A', 'Hearing impairment', 'not provided']
NM_000334.4(SCN4A):c.2078T>C (p.Ile693Thr)	80338956	SCN4A	[]	['AAGATCAYTGGCAATTCAGTGGG', 'AGATCAYTGGCAATTCAGTGGGG', 'GATCAYTGGCAATTCAGTGGGGG']	['Hyperkalemic Periodic Paralysis Type 1', 'Paramyotonia congenita of von Eulenburg']
NM_004523.3(KIF11):c.2547+2T>C	730882063	KIF11	['GGAGGYAATAACTTTGTAAGTGG']	['GGAGGYAATAACTTTGTAAGTGG']	['Microcephaly with or without chorioretinopathy, lymphedema, or mental retardation']
NM_003060.3(SLC22A5):c.1051T>C (p.Trp351Arg)	68018207	SLC22A5	[]	[]	['Renal carnitine transport defect']
NM_001070.4(TUBG1):c.1160T>C (p.Leu387Pro)	398123045	TUBG1	[]	[]	['Cortical dysplasia, complex, with other brain malformations 4']
NM_000441.1(SLC26A4):c.-103T>C	60284988	-	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_000016.5(ACADM):c.233T>C (p.Ile78Thr)	398123074	ACADM	[]	[]	['Medium-chain acyl-coenzyme A dehydrogenase deficiency', 'not provided']
NM_000179.2(MSH6):c.4001+2T>C	267608131	MSH6	[]	['CGGYAACTAACTAACTATAATGG']	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_000019.3(ACAT1):c.730+2T>C	398123096	ACAT1	[]	[]	['Deficiency of acetyl-CoA acetyltransferase', 'not provided']
NM_000430.3(PAFAH1B1):c.841T>C (p.Cys281Arg)	587784288	PAFAH1B1	[]	[]	['Lissencephaly 1']

NM_000899.4(KITLG):c.98T>C (p.Val33Ala)	730882156	KITLG	[]	[]	['Familial progressive hyperpigmentation with or without hypopigmentation']
NM_015599.2(PGM3):c.248T>C (p.Leu83Ser)	267608260	PGM3	['AATGTYGGCACCATCCTGGGAGG']	['AATGTYGGCACCATCCTGGGAGG']	['Immunodeficiency 23']
NM_000169.2(GLA):c.899T>C (p.Leu300Pro)	398123223	-	[]	[]	['Fabry disease']
NM_001256047.1(C19orf12):c.391A>G (p.Lys131Glu)	146170087	C19orf12	[]	[]	['Neurodegeneration with brain iron accumulation 4']
NM_172337.2(OTX2):c.674A>G (p.Asn225Ser)	370761964	OTX2	[]	[]	['Pituitary hormone deficiency, combined 6']
NM_000202.6(IDS):c.587T>C (p.Leu196Ser)	398123250	IDS	[]	[]	['Mucopolysaccharidosis, MPS-II', 'not provided']
NM_000252.2(MTM1):c.688T>C (p.Trp230Arg)	398123274	MTM1	[]	[]	['Severe X-linked myotubular myopathy', 'not provided']
NM_022445.3(TPK1):c.656A>G (p.Asn219Ser)	371271054	TPK1	[]	[]	['THIAMINE METABOLISM DYSFUNCTION SYNDROME 5 (EPISODIC ENCEPHALOPATHY TYPE)']
NM_014139.2(SCN11A):c.2432T>C (p.Leu811Pro)	483352920	SCN11A	[]	[]	['NEUROPATHY, HEREDITARY SENSORY AND AUTONOMIC, TYPE VII']
NM_000733.3(CD3E):c.520+2T>C	483352928	CD3E	[]	[]	['Immunodeficiency 18']
NM_017653.3(DYM):c.621-2A>G	775414124	DYM	[]	[]	['Dyggve-Melchior-Clausen syndrome']
NM_001253816.1(SLC52A2):c.1016T>C (p.Leu339Pro)	148234606	SLC52A2	[]	[]	['Brown-Vialetto-Van Laere syndrome 2']
NM_004963.3(GUCY2C):c.2782T>C (p.Cys928Arg)	587784573	-	[]	['TCCCYGTGCTGCTGGAGTTGTGG', 'CCCYGTGCTGCTGGAGTTGTGGG']	['Meconium ileus']
NM_003159.2(CDKL5):c.659T>C (p.Leu220Pro)	267608511	CDKL5	[]	['CCAACYTTTTACTATTCAGAAGG']	['Early infantile epileptic encephalopathy 2']
NM_000528.3(MAN2B1):c.2436+2T>C	398123457	MAN2B1	[]	[]	['not provided']
NM_002136.2(HNRNPA1):c.841T>C (p.Phe281Leu)	483353031	HNRNPA1	['AATYTTGGAGGCAGAAGCTCTGG']	['AATYTTGGAGGCAGAAGCTCTGG']	['Chronic progressive multiple sclerosis']

NM_006920.4(SCN1A):c.4251+2T>C	398123595	-	[]	[]	['Severe myoclonic epilepsy in infancy', 'Generalized epilepsy with febrile seizures plus, type 2']
NM_002225.3(IVD):c.465+2T>C	398123683	IVD	[]	[]	['Isovaleryl-CoA dehydrogenase deficiency', 'not provided']
NM_000118.3(ENG):c.1273-2A>G	373842615	ENG	[]	['CCGCCYGCGGGGATAAAGCCAGG', 'CGCCYGCGGGGATAAAGCCAGGG']	['Haemorrhagic telangiectasia 1']
NM_005859.4(PURA):c.218T>C (p.Phe73Ser)	793888535	PURA	[]	[]	['not provided']
NM_003494.3(DYSF):c.1284+2T>C	398123765	DYSF	['ATGGYAAGGAGCAAGGGAGCAGG']	['ATGGYAAGGAGCAAGGGAGCAGG']	['Limb-girdle muscular dystrophy, type 2B', 'not provided']
NM_004006.2(DMD):c.2380+2T>C	398123885	DMD	[]	[]	['Dilated cardiomyopathy 3B']
NM_006364.2(SEC23A):c.2104A>G (p.Met702Val)	138568622	SEC23A	[]	[]	['Craniolenticulosutural dysplasia']
NM_000335.4(SCN5A):c.376A>G (p.Lys126Glu)	185492581	SCN5A	[]	['GAATCTYCACAGCCGCTCTCCGG']	['Brugada syndrome']
NM_015865.6(SLC14A1):c.871T>C (p.Ser291Pro)	78242949	SLC14A1	[]	[]	[]
NM_003995.3(NPR2):c.226T>C (p.Ser76Pro)	796065355	NPR2	[]	[]	['SHORT STATURE WITH NONSPECIFIC SKELETAL ABNORMALITIES']
NM_012463.3(ATP6V0A2):c.825+2T>C	398124257	ATP6V0A2	['CACTGYGAGTAAGCTGGAAGTGG']	['CACTGYGAGTAAGCTGGAAGTGG']	['Cutis laxa with osteodystrophy', 'not provided']
NM_014795.3(ZEB2):c.73+2T>C	398124282	ZEB2	[]	[]	['Mowat-Wilson syndrome']
NM_000424.3(KRT5):c.1388T>C (p.Leu463Pro)	57599352	KRT5	[]	[]	['Epidermolysis bullosa simplex, Koebner type', 'not provided']
NM_133499.2(SYN1):c.1699A>G (p.Thr567Ala)	200533370	SYN1	[]	['GATGYCTGACGGGTAGCCTGTGG', 'ATGYCTGACGGGTAGCCTGTGGG']	['Epilepsy, X-linked, with variable learning disabilities and behavior disorders', 'not specified']
NM_148960.2(CLDN19):c.269T>C (p.Leu90Pro)	118203981	CLDN19	[]	['GCTCCYGGGCTTCGTGGCCATGG']	['Hypomagnesemia 5, renal, with ocular involvement']

NM_018006.4(TRMU):c.229T>C (p.Tyr77His)	118203990	TRMU	[]	[]	['Liver failure acute infantile']
NM_000056.3(BCKDHB):c.752T>C (p.Val251Ala)	398124593	BCKDHB	[]	[]	['Maple syrup urine disease', 'not provided']
NM_182680.1(AMELX):c.2T>C (p.Met1Thr)	104894737	-	[]	[]	['Amelogenesis imperfecta, type 1E']
NM_018105.2(THAP1):c.241T>C (p.Phe81Leu)	118204013	THAP1	[]	[]	['Dystonia 6, torsion']
NM_001235.3(SERPINH1):c.233T>C (p.Leu78Pro)	137853892	SERPINH1	[]	['GTCGCYAGGGCTCGTGTCGCTGG', 'TCGCYAGGGCTCGTGTCGCTGGG']	['Osteogenesis imperfecta type 10']
NM_004482.3(GALNT3):c.516_688del	761396172	GALNT3	[]	[]	['Tumoral calcinosis, familial, hyperphosphatemic']
NM_000263.3(NAGLU):c.142T>C (p.Phe48Leu)	118204024	NAGLU	[]	['GGCCGACYTCTCCGTGTCGGTGG']	['Mucopolysaccharidosis, MPS-III-B']
NM_000559.2(HBG1):c.-251T>C	35710727	HBG1	[]	[]	['Fetal hemoglobin quantitative trait locus 1']
NM_000527.4(LDLR):c.694+2T>C	200238879	LDLR	['CGGYATGGGCGGGGCCAGGGTGG']	['ACTGCGGYATGGGCGGGGCCAGG', 'CTGCGGYATGGGCGGGGCCAGGG', 'CGGYATGGGCGGGGCCAGGGTGG']	['Familial hypercholesterolemia']
NM_001012515.2(FECH):c.1268T>C (p.Phe423Ser)	118204039	FECH	[]	[]	['Erythropoietic protoporphyria']
NM_005211.3(CSF1R):c.1745T>C (p.Leu582Pro)	690016563	CSF1R	[]	['CAACCYGCAGTTTGGTGAGATGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_000526.4(KRT14):c.1243T>C (p.Tyr415His)	58380626	KRT14	[]	['CGCCACCYACCGCCGCCTGCTGG', 'CACCYACCGCCGCCTGCTGGAGG', 'ACCYACCGCCGCCTGCTGGAGGG']	['Epidermolysis bullosa herpetiformis, Dowling-Meara', 'not provided']
NM_006493.2(CLN5):c.2T>C (p.Met1Thr)	201615354	CLN5	[]	[]	['not provided']
NM_002863.4(PYGL):c.2461T>C (p.Tyr821His)	113993988	PYGL	['AAGAAYATGCCCAAAACATCTGG']	['AAGAAYATGCCCAAAACATCTGG']	['Glycogen storage disease, type VI']
NM_016038.2(SBDS):c.258+2T>C	113993993	SBDS	[]	[]	['Shwachman syndrome', 'Aplastic anemia, susceptibility to']
NM_000110.3(DPYD):c.85T>C (p.Cys29Arg)	1801265	DPYD	[]	[]	['Dihydropyrimidine dehydrogenase deficiency']
NM_001034116.1(EIF	113994038	EIF2B4	[]	[]	['Ovarioleukodystro

2B4):c.1393T>C (p.Cys465Arg)					phy']
NM_001165963.1(SCN1A):c.323T>C (p.Leu108Pro)	794726793	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_001034116.1(EIF2B4):c.1465T>C (p.Tyr489His)	113994040	EIF2B4	[]	[]	['Ovarioleukodystrophy']
NM_004304.4(ALK):c.3749T>C (p.Ile1250Thr)	113994092	ALK	[]	[]	['Neuroblastoma 3']
NM_207346.2(TSEN54):c.277T>C (p.Ser93Pro)	113994151	TSEN54	[]	['TTGAAGYCTCCCGCGGTGAGCGG', 'AAGYCTCCCGCGGTGAGCGGCGG']	['Pontocerebellar hypoplasia type 4']
NM_000018.3(ACADVL):c.848T>C (p.Val283Ala)	113994167	ACADVL	['TTTGYGGTGGAGAGGGGCTTCGG', 'TTGYGGTGGAGAGGGGCTTCGGG']	['TTTGYGGTGGAGAGGGGCTTCGG', 'TTGYGGTGGAGAGGGGCTTCGGG']	['Very long chain acyl-CoA dehydrogenase deficiency', 'not provided']
NM_000430.3(PAFAH1B1):c.569-10T>C	113994202	PAFAH1B1	[]	[]	['Lissencephaly 1']
NM_004937.2(CTNS):c.473T>C (p.Leu158Pro)	113994206	CTNS	[]	['TGGTCYGAGCTTCGACTTCGTGG']	['Cystinosis']
NM_000546.5(TP53):c.488A>G (p.Tyr163Cys)	148924904	TP53	['GCTTGYAGATGGCCATGGCGCGG']	['GCTTGYAGATGGCCATGGCGCGG']	['Hereditary cancer-predisposing syndrome']
NM_004211.3(SLC6A5):c.1444T>C (p.Trp482Arg)	281864925	SLC6A5	[]	[]	['Hyperekplexia 3']
NM_024312.4(GNPTAB):c.1208T>C (p.Ile403Thr)	281864973	GNPTAB	[]	[]	['Pseudo-Hurler polydystrophy']
NM_024312.4(GNPTAB):c.3002T>C (p.Leu1001Pro)	281865006	GNPTAB	[]	[]	['I cell disease']
NM_000540.2(RYR1):c.7358T>C (p.Ile2453Thr)	118192123	RYR1	[]	[]	['Central core disease', 'not provided']
NM_000748.2(CHRNB2):c.923T>C (p.Val308Ala)	281865070	CHRNB2	[]	[]	['Epilepsy, nocturnal frontal lobe, type 3']
NM_000526.4(KRT14):c.356T>C (p.Met119Thr)	28928893	KRT14	[]	[]	['Epidermolysis bullosa herpetiformis, Dowling-Meara', 'not provided']
NM_000093.4(COL5A1):c.5137-11T>A	183495554	-	[]	[]	['Ehlers-Danlos syndrome, classic type']
NM_000277.1(PAH):c	62516109	PAH	[]	['CCACTTCYTGAA	['Phenylketonuria',

.638T>C (p.Leu213Pro)				AAGTACTGTGG']	'not provided']
NM_000531.5(OTC):c.663+2T>C	72558427	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.717+2T>C	72558431	OTC	[]	[]	['Ornithine carbamoyltransferase deficiency', 'not provided']
NM_000531.5(OTC):c.793T>C (p.Trp265Arg)	72558445	OTC	[]	[]	['not provided']
NM_000493.3(COL10A1):c.1841T>C (p.Leu614Pro)	111033545	-	[]	[]	['Metaphyseal chondrodysplasia, Schmid type']
NM_000493.3(COL10A1):c.1771T>C (p.Cys591Arg)	111033546	-	[]	[]	['Metaphyseal chondrodysplasia, Schmid type']
NM_000493.3(COL10A1):c.2011T>C (p.Ser671Pro)	111033552	-	[]	[]	['Metaphyseal chondrodysplasia, Schmid type']
NM_004614.4(TK2):c.173A>G (p.Asn58Ser)	138439950	TK2	[]	[]	['Mitochondrial DNA depletion syndrome 2']
NM_004614.4(TK2):c.644T>C (p.Leu215Pro)	281865497	TK2	[]	[]	['Mitochondrial DNA depletion syndrome 2']
NM_004614.4(TK2):c.156+2T>C	281865499	TK2	[]	[]	['Mitochondrial DNA depletion syndrome 2']
NM_153026.2(PRICKLE1):c.1414T>C (p.Tyr472His)	281865564	PRICKLE1	[]	[]	['Progressive myoclonus epilepsy with ataxia']
NM_017882.2(CLN6):c.767A>G (p.Asp256Gly)	143781303	CLN6	[]	[]	['not provided']
NM_130838.1(UBE3A):c.389T>C (p.Ile130Thr)	111033597	UBE3A	[]	[]	['Angelman syndrome']
NM_001198799.2(ASCC1):c.953A>G (p.Asn318Ser)	146370051	ASCC1	[]	[]	[]
NM_000174.4(GP9):c.70T>C (p.Cys24Arg)	28933378	GP9	['CCCAYGTACCTGCCGCGCCCTGG']	['CCCAYGTACCTGCCGCGCCCTGG']	['Bernard Soulier syndrome', 'Bernard-Soulier syndrome type C']
NM_001173464.1(KIF21A):c.3029T>C (p.Ile1010Thr)	121912587	KIF21A	[]	[]	['Fibrosis of extraocular muscles, congenital, 1']
NM_001302946.1(TRNT1):c.668T>C (p.Ile223Thr)	370011798	TRNT1	[]	['GCAAYTGCAGAAAATGCAAAAGG']	['Sideroblastic anemia with B-cell immunodeficiency, periodic fevers, and developmental delay']
NM_000371.3(TTR):c.250T>C (p.Phe84Leu)	121918091	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis']
m.5814T>C	200077222	MT-TC	[]	[]	['Juvenile myopathy,

					encephalopathy, lactic acidosis AND stroke']
NM_000277.1(PAH):c.293T>C (p.Leu98Ser)	62517167	PAH	[]	['AAGATCTYGAGGCATGACATTGG']	['Mild non-PKU hyperphenylalanemia', 'not provided']
NM_017882.2(CLN6):c.486+2T>C	796052355	CLN6	[]	[]	['not provided']
NM_001813.2(CENPE):c.4063A>G (p.Lys1355Glu)	141488085	CENPE	[]	[]	['Primary autosomal recessive microcephaly 13']
NM_001918.3(DBT):c.1150G>A (p.Gly384Ser)	12021720	DBT	[]	['GACYCACAGAGCCCAATTTCTGG']	['Intermediate maple syrup urine disease type 2']
NM_000495.4(COL4A5):c.4699T>C (p.Cys1567Arg)	104886288	COL4A5	['AGTAYGTGAAGCTCCAGCTGTGG']	['AGTAYGTGAAGCTCCAGCTGTGG']	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.4756T>C (p.Cys1586Arg)	104886289	COL4A5	[]	['TCCCCATYGTCCTCAGGGATGGG']	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.5032T>C (p.Cys1678Arg)	104886310	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000155.3(GALT):c.482T>C (p.Leu161Pro)	111033700	GALT	['AGCYGGGTGCCCAGTACCCTTGG']	['AGCYGGGTGCCCAGTACCCTTGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_001199252.2(SGOL1):c.67A>G (p.Lys23Glu)	199815268	-	[]	[]	['Chronic atrial and intestinal dysrhythmia']
NC_012920.1:m.5559A>G	370471013	MT-TW	[]	['CAACYTACTGAGGGCTTTGAAGG']	['Leigh disease']
NM_000487.5(ARSA):c.410T>C (p.Leu137Pro)	121434215	ARSA	[]	['GCCTTCCYGCCCCCCCATCAGGG']	['Metachromatic leukodystrophy, adult type']
NM_000051.3(ATM):c.7967T>C (p.Leu2656Pro)	121434218	-	[]	[]	[]
NM_000096.3(CP):c.650T>C (p.Phe217Ser)	386134125	CP	[]	[]	['Deficiency of ferroxidase']
NM_000096.3(CP):c.1123T>C (p.Tyr375His)	386134128	CP	[]	['ACACTACYACATTGCCGCTGAGG']	['Deficiency of ferroxidase']
NM_000268.3(NF2):c.185T>C (p.Phe62Ser)	121434261	NF2	[]	[]	['Neurofibromatosis, type 2']
NM_000495.4(COL4A5):c.4803+121T>C	104886423	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_024529.4(CDC73):c.191T>C (p.Leu64Pro)	121434264	CDC73	[]	[]	['Hyperparathyroidism 1']
NM_001061.4(TBXAS1):c.1463T>C (p.Leu488Pro)	199422114	TBXAS1	[]	[]	[]
NM_001127328.2(ACADM):c.1136T>C (p.Ile379Thr)	121434275	ACADM	[]	['GTGCAGAYACTTGGAGGCAATGG']	['Medium-chain acyl-coenzyme A dehydrogenase deficiency']

NM_001127328.2(ACADM):c.742T>C (p.Cys248Arg)	121434276	ACADM	[]	['CAGCGAYGTTCAGATACTAGAGG']	['Medium-chain acyl-coenzyme A dehydrogenase deficiency']
NM_000016.5(ACADM):c.199T>C (p.Tyr67His)	121434280	ACADM	[]	[]	['Medium-chain acyl-coenzyme A dehydrogenase deficiency', 'not provided']
NM_002225.3(IVD):c.134T>C (p.Leu45Pro)	121434284	IVD	[]	['ATGGGCYAAGCGAGGAGCAGAGG']	['ISOVALERIC ACIDEMIA, TYPE I']
NM_005957.4(MTHFR):c.968T>C (p.Leu323Pro)	121434297	MTHFR	[]	[]	['Homocystinuria due to MTHFR deficiency']
NM_000136.2(FANCC):c.1661T>C (p.Leu554Pro)	104886458	-	[]	[]	['Fanconi anemia, complementation group C', 'not provided']
NM_005908.3(MANBA):c.1513T>C (p.Ser505Pro)	121434334	MANBA	[]	['ATTACGYCCAGTCCTACAAATGG', 'TTACGYCCAGTCCTACAAATGGG', 'TACGYCCAGTCCTACAAATGGGG']	['Beta-D-mannosidosis']
NM_000244.3(MEN1):c.518T>C (p.Leu173Pro)	386134256	MEN1	[]	[]	['Multiple endocrine neoplasia, type 1']
NM_199242.2(UNC13D):c.1208T>C (p.Leu403Pro)	121434353	UNC13D	[]	[]	['Hemophagocytic lymphohistiocytosis, familial, 3']
NM_152783.4(D2HGDH):c.1331T>C (p.Val444Ala)	121434360	D2HGDH	[]	[]	['D-2-hydroxyglutaric aciduria 1']
NM_207118.2(GTF2H5):c.62T>C (p.Leu21Pro)	121434365	GTF2H5	[]	[]	['Photosensitive trichothiodystrophy']
NM_000159.3(GCDH):c.883T>C (p.Tyr295His)	121434366	GCDH	[]	['CGCCCGGYACGGCATCGCGTGGG', 'GCCCGGYACGGCATCGCGTGGGG']	['Glutaric aciduria, type 1']
NM_018668.4(VPS33B):c.89T>C (p.Leu30Pro)	121434385	VPS33B	[]	[]	['Arthrogryposis renal dysfunction cholestasis syndrome']
NM_000424.3(KRT5):c.541T>C (p.Ser181Pro)	60715293	KRT5	[]	['GTTTGCCYCCTTCATCGACAAGG']	['Epidermolysis bullosa herpetiformis, Dowling-Meara', 'not provided']
NM_001003722.1(GLE1):c.2051T>C (p.Ile684Thr)	121434409	GLE1	[]	['AAGGACAYTCCTGTCCCCAAGGG']	['Lethal arthrogryposis with anterior horn cell disease']
NM_003659.3(AGPS):c.1406T>C (p.Leu469Pro)	121434413	AGPS	[]	[]	['Rhizomelic chondrodysplasia punctata type 3']
NM_004550.4(NDUFS2):c.1237T>C	121434429	NDUFS2	[]	[]	['Mitochondrial complex I

(p.Ser413Pro)					deficiency', 'not provided']
NM_001287.5(CLCN7):c.2297T>C (p.Leu766Pro)	121434434	CLCN7	[]	['GGGCCYGCGGCACCTGGTGGTGG']	['Osteopetrosis autosomal recessive 4']
m.14709T>C	121434453	MT-TE	[]	[]	['Diabetes-deafness syndrome maternally transmitted']
NM_000466.2(PEX1):c.1991T>C (p.Leu664Pro)	121434455	PEX1	[]	['GATGACCYTGACCTCATTGCTGG']	['Zellweger syndrome']
NM_198253.2(TERT):c.3043T>C (p.Cys1015Arg)	199422307	TERT	[]	[]	['Aplastic anemia']
m.4290T>C	121434469	MT-TI	['ACTYTGATAGAGTAAATAATAGG']	['ACTYTGATAGAGTAAATAATAGG']	[]
m.4291T>C	121434471	MT-TI	['ACTTYGATAGAGTAAATAATAGG']	['ACTTYGATAGAGTAAATAATAGG']	['Hypertension, hypercholesterolemia, and hypomagnesemia, mitochondrial']
m.9997T>C	121434475	MT-TG	[]	[]	['Primary familial hypertrophic cardiomyopathy']
NM_001099274.1(TINF2):c.860T>C (p.Leu287Pro)	199422316	TINF2	[]	[]	['Dyskeratosis congenita autosomal dominant']
NM_001099274.1(TINF2):c.862T>C (p.Phe288Leu)	199422317	TINF2	[]	['CTGYTTCCCTTTAGGAATCTCGG']	['Aplastic anemia']
NM_000430.3(PAFAH1B1):c.505T>C (p.Ser169Pro)	121434484	PAFAH1B1	[]	[]	['Subcortical band heterotopia']
NM_000430.3(PAFAH1B1):c.92T>C (p.Phe31Ser)	121434486	PAFAH1B1	[]	[]	['Lissencephaly 1']
NM_005535.2(IL12RB1):c.592T>C (p.Cys198Arg)	121434495	IL12RB1	[]	[]	['Immunodeficiency 30']
NM_030662.3(MAP2K2):c.400T>C (p.Tyr134His)	121434499	MAP2K2	[]	[]	['Cardiofaciocutaneous syndrome 4', 'Rasopathy', 'Noonan syndrome and Noonan-related syndrome']
NM_001065.3(TNFRSF1A):c.349T>C (p.Cys117Arg)	104895221	TNFRSF1A	[]	['CTCTTCTYGCACAGTGGACCGGG']	['TNF receptor-associated periodic fever syndrome (TRAPS)']
NM_000123.3(ERCC5):c.2573T>C (p.Leu858Pro)	121434575	-	[]	[]	['Xeroderma pigmentosum, group G']
NM_001493.2(GDI1):c.275T>C (p.Leu92Pro)	121434607	GDI1	[]	[]	['X-Linked Mental Retardation 41']

NM_020061.5(OPN1LW):c.607T>C (p.Cys203Arg)	121434621	OPN1LW	[]	[]	['Cone monochromatism']
NM_024420.2(PLA2G4A):c.331T>C (p.Ser111Pro)	121434634	PLA2G4A	[]	[]	[]
NM_005557.3(KRT16):c.395T>C (p.Leu132Pro)	60944949	KRT16	[]	[]	['Pachyonychia congenita, type 1', 'not provided']
NM_000485.2(APRT):c.407T>C (p.Met136Thr)	28999113	APRT	[]	[]	['Adenine phosphoribosyltransferase deficiency', 'APRT deficiency, Japanese type']
NM_005270.4(GLI2):c.4663T>C (p.Ser1555Pro)	144372453	GLI2	[]	[]	['Holoprosencephaly 9', 'not specified']
NM_024753.4(TTC21B):c.2384T>C (p.Leu795Pro)	387907060	TTC21B	[]	[]	['Asphyxiating thoracic dystrophy 4']
NM_000155.3(GALT):c.680T>C (p.Leu227Pro)	111033846	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000138.4(FBN1):c.4987T>C (p.Cys1663Arg)	137854459	FBN1	[]	['GGGACAYGTTACAACACCGTTGG']	['Marfan syndrome']
NM_032446.2(MEGF10):c.976T>C (p.Cys326Arg)	387907073	MEGF10	[]	[]	['Myopathy, areflexia, respiratory distress, and dysphagia, early-onset, mild variant']
NM_024027.4(COLEC11):c.505T>C (p.Ser169Pro)	387907075	COLEC11	[]	['CAGCTGYCCTGCCAGGGCCGCGG', 'AGCTGYCCTGCCAGGGCCGCGGG', 'GCTGYCCTGCCAGGGCCGCGGGG', 'CTGYCCTGCCAGGGCCGCGGGGG']	['Carnevale syndrome']
NM_001946.3(DUSP6):c.566A>G (p.Asn189Ser)	143946794	DUSP6	['CACTAYTGGGGTCTCGGTCAAGG']	['CACTAYTGGGGTCTCGGTCAAGG']	['Hypogonadotropic hypogonadism 19 with or without anosmia']
NM_000138.4(FBN1):c.3793T>C (p.Cys1265Arg)	137854474	FBN1	['CTTGYGTTATGATGGATTCATGG']	['CTTGYGTTATGATGGATTCATGG']	['Marfan syndrome']
NM_022068.3(PIEZO2):c.2134A>G (p.Met712Val)	587777453	PIEZO2	[]	[]	['Oculomelic amyoplasia']
NM_000570.4(FCGR3B):c.316A= (p.Ile106=)	2290834	FCGR3B	[]	[]	[]
NM_000352.4(ABCC8):c.674T>C (p.Leu225Pro)	1048095	ABCC8	[]	['TGCYGTCCAAAGGCACCTACTGG']	['Permanent neonatal diabetes mellitus']

NM_153490.2(KRT13):c.332T>C (p.Leu111Pro)	59897026	KRT13	[]	[]	['White sponge nevus 2', 'not provided']
NM_000132.3(F8):c.1174T>C (p.Ser392Pro)	28933669	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000492.3(CFTR):c.1021T>C (p.Ser341Pro)	397508144	CFTR	[]	[]	['Cystic fibrosis']
NM_000133.3(F9):c.1058T>C (p.Val353Ala)	137852255	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_001202.3(BMP4):c.362A>G (p.His121Arg)	376960358	BMP4	['TTCGTGGYGGAAGCTCCTCACGG']	['TTCGTGGYGGAAGCTCCTCACGG']	['Microphthalmia syndromic 6']
NM_000133.3(F9):c.1328T>C (p.Ile443Thr)	137852268	F9	['GAAYATATACCAAGGTATCCCGG']	['GAAYATATACCAAGGTATCCCGG']	['Hereditary factor IX deficiency disease']
NM_000133.3(F9):c.1357T>C (p.Trp453Arg)	137852269	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_000435.2(NOTCH3):c.1363T>C (p.Cys455Arg)	28933698	NOTCH3	['ACCYGTATCTGTATGGCAGGTGG']	['TTCACCYGTATCTGTATGGCAGG', 'ACCYGTATCTGTATGGCAGGTGG']	['Cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy']
NM_019074.3(DLL4):c.1168T>C (p.Cys390Arg)	796065347	DLL4	[]	['GAAYGTCCCCCCAACTTCACCGG']	['Adams-Oliver syndrome', 'ADAMS-OLIVER SYNDROME 6']
NM_000032.4(ALAS2):c.595T>C (p.Tyr199His)	137852310	ALAS2	[]	[]	['Hereditary sideroblastic anemia']
NM_019074.3(DLL4):c.583T>C (p.Phe195Leu)	796065351	DLL4	[]	[]	['Adams-Oliver syndrome']
NM_000402.4(G6PD):c.1054T>C (p.Tyr352His)	137852347	G6PD	[]	['AGGGYACCTGGACGACCCCACGG']	['Anemia, nonspherocytic hemolytic, due to G6PD deficiency']
NM_007325.4(GRIA3):c.2117T>C (p.Met706Thr)	137852352	GRIA3	[]	[]	['Mental retardation, X-linked, syndromic, wu type']
NM_000132.3(F8):c.6554T>C (p.Leu2185Ser)	137852365	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.5372T>C (p.Met1791Thr)	137852375	F8	['TCAYGGTGAGTTAAGGACAGTGG']	['TCAYGGTGAGTTAAGGACAGTGG']	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.1754T>C (p.Ile585Thr)	137852376	F8	['AACAGAYAA	['AACAGAYAATGTCAGACAAGAGG'	['Hereditary factor VIII deficiency

			TGTCA GACAA GAGG']	]	disease']
NM_001127695.1(CTSA):c.707T>C (p.Leu236Pro)	137854546	CTSA	[]	[]	['Galactosialidosis, early infantile']
NM_000132.3(F8):c.935T>C (p.Phe312Ser)	137852405	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.980T>C (p.Leu327Pro)	137852407	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000308.2(CTSA):c.1271T>C (p.Met424Thr)	137854548	CTSA	[]	[]	['Galactosialidosis, late infantile']
NM_000132.3(F8):c.1481T>C (p.Ile494Thr)	137852413	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_001250.5(CD40):c.247T>C (p.Cys83Arg)	28931586	CD40	[]	[]	['Immunodeficiency with hyper IgM type 3']
NM_000165.4(GJA1):c.32T>C (p.Leu11Pro)	121912969	GJA1	[]	[]	['Oculodentodigital dysplasia']
NM_000132.3(F8):c.1958T>C (p.Val653Ala)	137852430	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_001972.2(ELANE):c.211T>C (p.Cys71Arg)	28931611	ELANE	[]	[]	['Severe congenital neutropenia autosomal dominant']
NM_000098.2(CPT2):c.1342T>C (p.Phe448Leu)	74315297	CPT2	[]	[]	['CARNITINE PALMITOYLTRANSFERASE II DEFICIENCY, LATE-ONSET', 'not provided']
NM_213653.3(HFE2):c.842T>C (p.Ile281Thr)	74315326	HFE2	[]	[]	['Hemochromatosis type 2A']
NM_213653.3(HFE2):c.302T>C (p.Leu101Pro)	74315327	HFE2	[]	['GGACCYCGCCTTCCATTCGGCGG']	['Hemochromatosis type 2A']
NM_000194.2(HPRT1):c.122T>C (p.Leu41Pro)	137852480	HPRT1	[]	[]	['Lesch-Nyhan syndrome']
NM_000261.1(MYOC):c.1297T>C (p.Cys433Arg)	74315338	MYOC	[]	[]	['Primary open angle glaucoma juvenile onset 1']
NM_000194.2(HPRT1):c.170T>C (p.Met57Thr)	137852495	HPRT1	[]	[]	['Lesch-nyhan syndrome, neurologic variant']
NM_000267.3(NF1):c.3728T>C (p.Leu1243Pro)	137854564	NF1	[]	[]	['Neurofibromatosis, type 1']
NM_000291.3(PGK1):c.263T>C (p.Leu88Pro)	137852531	PGK1	[]	[]	['Phosphoglycerate kinase 1 deficiency']
NM_000291.3(PGK1):c.946T>C	137852533	PGK1	[]	[]	['Phosphoglycerate kinase 1 deficiency']

(p.Cys316Arg)					
NM_000291.3(PGK1):c.758T>C (p.Ile253Thr)	137852534	PGK1	[]	[]	['Phosphoglycerate kinase 1 deficiency']
NM_170784.2(MKKS):c.830T>C (p.Leu277Pro)	74315398	MKKS	[]	[]	['Bardet-Biedl syndrome 6', 'not provided']
NM_000451.3(SHOX):c.877T>C (p.Ter293Arg)	137852559	SHOX	[]	[]	['Leri Weill dyschondrosteosis']
NM_001029871.3(RSPO4):c.319T>C (p.Cys107Arg)	74315421	RSPO4	[]	[]	['Anonychia']
NM_000044.3(AR):c.2033T>C (p.Leu678Pro)	137852579	AR	[]	['GTCCYGGAAGCCATTGAGCCAGG']	[]
NM_000044.3(AR):c.2423T>C (p.Met808Thr)	137852592	AR	[]	[]	['Reifenstein syndrome']
NM_000044.3(AR):c.2596T>C (p.Ser866Pro)	137852597	AR	[]	[]	['Androgen resistance syndrome']
NM_172201.1(KCNE2):c.161T>C (p.Met54Thr)	74315447	KCNE2	[]	[]	['Long QT syndrome 6', 'Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_172201.1(KCNE2):c.170T>C (p.Ile57Thr)	74315448	KCNE2	[]	[]	['Long QT syndrome 6', 'Cardiac arrhythmia', 'not provided']
NM_000211.4(ITGB2):c.446T>C (p.Leu149Pro)	137852611	ITGB2	['AGCYAGGTGGCGACCTGCTCCGG']	['AGCYAGGTGGCGACCTGCTCCGG']	['Leukocyte adhesion deficiency']
NM_000211.4(ITGB2):c.412T>C (p.Ser138Pro)	137852617	ITGB2	[]	[]	['Leukocyte adhesion deficiency']
NM_000023.2(SGCA):c.524T>C (p.Val175Ala)	137852622	SGCA	[]	[]	['Limb-girdle muscular dystrophy, type 2D']
NM_001166107.1(HMGCS2):c.520T>C (p.Phe174Leu)	137852636	HMGCS2	[]	['CCCTCYTCAATGCTGCCAACTGG']	['mitochondrial 3-hydroxy-3-methylglutaryl-CoA synthase deficiency']
NM_001886.2(CRYBA4):c.281T>C (p.Phe94Ser)	74315486	CRYBA4	[]	[]	['Cataract 23']
NM_001886.2(CRYBA4):c.206T>C (p.Leu69Pro)	74315487	CRYBA4	[]	[]	['Cataract 23']
NM_002047.2(GARS):c.548T>C (p.Leu183Pro)	137852644	GARS	[]	[]	['Distal hereditary motor neuronopathy type 5']
NM_000268.3(NF2):c.1604T>C (p.Leu535Pro)	74315493	NF2	[]	[]	['Neurofibromatosis, type 2']
NM_000095.2(COMP):c.982T>C	137852653	COMP	[]	[]	['Pseudoachondroplastic

(p.Cys328Arg)					spondyloepiphyseal dysplasia syndrome']
NM_000095.2(COMP):c.1042T>C (p.Cys348Arg)	137852656	COMP	[]	[]	['Pseudoachondroplastic spondyloepiphyseal dysplasia syndrome']
NM_017929.5(PEX26):c.2T>C (p.Met1Thr)	74315506	PEX26	[]	[]	['Peroxisome biogenesis disorder 7B']
NM_033163.3(FGF8):c.118T>C (p.Phe40Leu)	137852661	FGF8	[]	['TTCCCTGYTCCGGGCTGGCCGGG']	['Kallmann syndrome 6']
NM_007315.3(STAT1):c.2117T>C (p.Leu706Ser)	137852677	STAT1	[]	[]	['Immunodeficiency 31a']
NM_007315.3(STAT1):c.1799T>C (p.Leu600Pro)	137852678	STAT1	[]	[]	['Mycobacterial and viral infections, susceptibility to, autosomal recessive']
NM_000336.2(SCNN1B):c.1858T>C (p.Tyr620His)	137852707	SCNN1B	[]	[]	['Pseudoprimary hyperaldosteronism']
NM_005215.3(DCC):c.503T>C (p.Met168Thr)	121912967	DCC	[]	['AGCCCAYGCCAACAATCCACTGG']	[]
NM_001034850.2(FAM134B):c.873+2T>C	137852738	FAM134B	[]	[]	['Hereditary sensory and autonomic neuropathy type IIA']
NM_000182.4(HADHA):c.1025T>C (p.Leu342Pro)	137852772	HADHA	[]	[]	['Mitochondrial trifunctional protein deficiency']
NM_001165974.1(UROC1):c.209T>C (p.Leu70Pro)	137852796	UROC1	[]	[]	['Urocanate hydratase deficiency']
NM_000405.4(GM2A):c.412T>C (p.Cys138Arg)	137852797	GM2A	[]	[]	['Tay-Sachs disease, variant AB']
NM_001039523.2(CHRNA1):c.901T>C (p.Phe301Leu)	137852806	CHRNA1	[]	['TGTGYTCCTTCTGGTCATCGTGG']	['Myasthenic syndrome, congenital, fast-channel']
NM_003688.3(CASK):c.802T>C (p.Tyr268His)	137852817	CASK	[]	[]	['FG syndrome 4']
NM_003688.3(CASK):c.2740T>C (p.Trp914Arg)	137852819	CASK	['CACAGYGGGTCCCTGTCTCCTGG', 'ACAGYGGGTCCCCTGTCTCCTGGG']	['CACAGYGGGTCCCTGTCTCCTGG', 'ACAGYGGGTCCCTGTCTCCTGGG']	['FG syndrome 4']
NM_182760.3(SUMF1):c.1006T>C (p.Cys336Arg)	137852848	SUMF1	[]	[]	['Multiple sulfatase deficiency']

NM_182760.3(SUMF1):c.463T>C (p.Ser155Pro)	137852850	SUMF1	[]	['GGCGACYCCTTTGTCTTTGAAGG']	['Multiple sulfatase deficiency', 'not provided']
NM_000158.3(GBE1):c.671T>C (p.Leu224Pro)	137852886	GBE1	[]	['AATGTACYACCAAGAATCAAAGG']	['Glycogen storage disease, type IV', 'GLYCOGEN STORAGE DISEASE IV, NONPROGRESSIVE HEPATIC']
m.8356T>C	118192099	MT-TK	[]	[]	['Myoclonus with epilepsy with ragged red fibers', 'MERRF/MELAS overlap syndrome']
NM_024312.4(GNPTAB):c.1120T>C (p.Phe374Leu)	137852900	GNPTAB	[]	[]	['Pseudo-Hurler polydystrophy', 'I cell disease']
NM_031924.4(RSPH3):c.631-2A>G	142800871	RSPH3	[]	[]	['Kartagener syndrome']
NM_058172.5(ANTXR2):c.566T>C (p.Ile189Thr)	137852905	ANTXR2	[]	[]	['Hyaline fibromatosis syndrome']
NM_000419.3(ITGA2B):c.641T>C (p.Leu214Pro)	137852911	ITGA2B	[]	['CTGGTGCYTGGGGCTCCTGGCGG']	['Glanzmann thrombasthenia']
NM_001171507.2(MCFD2):c.407T>C (p.Ile136Thr)	137852914	MCFD2	[]	[]	['Factor v and factor viii, combined deficiency of, 2']
NM_000540.2(RYR1):c.1205T>C (p.Met402Thr)	118192117	RYR1	['CGCAYGATCCACAGCACCAATGG']	['CGCAYGATCCACAGCACCAATGG']	['Congenital myopathy with fiber type disproportion', 'Central core disease', 'not provided']
NM_000540.2(RYR1):c.13703T>C (p.Leu4568Pro)	118192131	RYR1	[]	[]	['Central core disease', 'not provided']
NM_000540.2(RYR1):c.13949T>C (p.Leu4650Pro)	118192138	RYR1	[]	[]	['Central core disease', 'not provided']
NM_138694.3(PKHD1):c.10658T>C (p.Ile3553Thr)	137852948	PKHD1	[]	['GAGCCCAYTGAAATACGCTCAGG']	['Polycystic kidney disease, infantile type']
NM_012464.4(TLL1):c.713T>C (p.Val238Ala)	137852952	TLL1	['GGGATTGYTGTTCATGAATTGGG']	['GGGATTGYTGTTCATGAATTGGG']	['Atrial septal defect 6']
NM_000540.2(RYR1):c.14762T>C (p.Phe4921Ser)	118192156	RYR1	[]	[]	['Central core disease', 'not provided']
NM_024960.4(PANK2):c.178T>C (p.Ser60Pro)	137852964	PANK2	[]	['ATTGACYCAGTCGGATTCAATGG']	[]
NM_001001486.1(ATP2C1):c.1751T>C (p.Leu584Pro)	137853015	ATP2C1	[]	[]	['Familial benign pemphigus']
NM_014363.5(SACS):	137853017	SACS	[]	[]	['Spastic ataxia

c.5836T>C (p.Trp1946Arg)					Charlevoix-Saguenay type']
NM_014363.5(SACS): c.9742T>C (p.Trp3248Arg)	137853018	SACS	[]	[]	['Spastic ataxia Charlevoix-Saguenay type']
NM_014363.5(SACS): c.3161T>C (p.Phe1054Ser)	137853019	SACS	[]	[]	['Spastic ataxia Charlevoix-Saguenay type']
NM_006899.3(IDH3B):c.395T>C (p.Leu132Pro)	137853020	IDH3B	[]	['TGCGGCYGAGG TAGGTGGTCTGG', 'GCGGCYGAGGTA GGTGGTCTGGG']	['Retinitis pigmentosa 46']
NM_001139.2(ALOX 12B):c.1277T>C (p.Leu426Pro)	137853023	ALOX12B	[]	[]	['Autosomal recessive congenital ichthyosis 2']
NM_001080463.1(DY NC2H1):c.3719T>C (p.Ile1240Thr)	137853028	DYNC2H1	[]	[]	['Short-rib thoracic dysplasia 3 with or without polydactyly']
NM_006009.3(TUBA1 A):c.1190T>C (p.Leu397Pro)	137853048	TUBA1A	[]	[]	['Lissencephaly 3']
NM_004519.3(KCNQ 3):c.925T>C (p.Trp309Arg)	118192249	KCNQ3	[]	[]	['Benign familial neonatal seizures 2']
NM_000531.5(OTC):c .332T>C (p.Leu111Pro)	1800324	OTC	[]	[]	['Ornithine carbamoyltransferas e deficiency']
NM_001172567.1(MY D88):c.317T>C (p.Leu106Pro)	137853065	MYD88	[]	[]	['Myd88 deficiency']
NM_002241.4(KCNJ1 0):c.418T>C (p.Cys140Arg)	137853068	KCNJ10	[]	[]	['SeSAME syndrome']
NM_000455.4(STK11) :c.200T>C (p.Leu67Pro)	137853077	STK11	[]	[]	['Peutz-Jeghers syndrome']
NM_000518.4(HBB):c .332T>C (p.Leu111Pro)	35256489	HBB	[]	[]	['Beta thalassemia major']
NM_005094.3(SLC27 A4):c.739T>C (p.Ser247Pro)	137853133	SLC27A4	[]	[]	['Ichthyosis prematurity syndrome']
NM_001145308.4(LR TOMT):c.313T>C (p.Trp105Arg)	137853186	LRTOMT	[]	[]	['Deafness, autosomal recessive 63']
NM_178012.4(TUBB2 B):c.514T>C (p.Ser172Pro)	137853194	TUBB2B	[]	[]	['Polymicrogyria, asymmetric']
NM_178012.4(TUBB2 B):c.683T>C (p.Leu228Pro)	137853195	TUBB2B	[]	[]	['Polymicrogyria, asymmetric']
NM_178012.4(TUBB2 B):c.793T>C (p.Phe265Leu)	137853196	TUBB2B	[]	[]	['Polymicrogyria, asymmetric']
NM_001082971.1(DD C):c.925T>C (p.Phe309Leu)	137853209	DDC	[]	[]	['Deficiency of aromatic-L-amino-acid decarboxylase']
NM_006121.3(KRT1):	137853225	KRT1	[]	[]	['Bullous

c.1424T>C (p.Leu475Pro)					ichthyosiform erythroderma']
NM_033500.2(HK1):c.1550T>C (p.Leu517Ser)	137853249	HK1	[]	['GACTTCTYGGCCCTGGATCTTGG', 'TTCTYGGCCCTGGATCTTGGAGG']	['Hemolytic anemia due to hexokinase deficiency']
NM_000249.3(MLH1):c.229T>C (p.Cys77Arg)	63749859	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_000444.5(PHEX):c.1664T>C (p.Leu555Pro)	137853270	PHEX	[]	['AGCYCCAGAAGCCTTTCTTTTGG']	['Familial X-linked hypophosphatemic vitamin D refractory rickets']
NM_000321.2(RB1):c.2134T>C (p.Cys712Arg)	137853296	RB1	[]	[]	['Retinoblastoma']
NM_007313.2(ABL1):c.988T>C (p.Phe330Leu)	137853304	ABL1	[]	[]	[]
NM_003639.4(IKBKG):c.1249T>C (p.Cys417Arg)	137853325	IKBKG	[]	['TGGAGYGCATTGAGTAGGGCCGG']	['Hypohidrotic ectodermal dysplasia with immune deficiency', 'Hyper-IgM immunodeficiency, X-linked, with hypohidrotic ectodermal dysplasia']
NM_017534.5(MYH2):c.5609T>C (p.Leu1870Pro)	786201023	-	[]	[]	['Inclusion body myopathy 3']
NM_000314.6(PTEN):c.406T>C (p.Cys136Arg)	786201044	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_001257988.1(TYMP):c.665A>G (p.Lys222Arg)	149977726	TYMP	['CACGAGTYTCTTACTGAGAATGG']	['CACGAGTYTCTTACTGAGAATGG', 'GAGTYTCTTACTGAGAATGGAGG']	[]
NM_016725.2(FOLR1):c.493+2T>C	144637717	FOLR1	['AGGYGAGGGCTGGGGTGGGCAGG']	['CTTCAGGYGAGGGCTGGGGTGGG', 'AGGYGAGGGCTGGGGTGGGCAGG']	['not provided']
NM_002225.3(IVD):c.295+2T>C	748026507	IVD	[]	[]	['not provided']
NM_000175.3(GPI):c.1016T>C (p.Leu339Pro)	137853587	GPI	[]	[]	[]
NM_002055.4(GFAP):c.1055T>C (p.Leu352Pro)	28932769	GFAP	[]	['GGACCYGCTCAATGTCAAGCTGG']	['Alexander disease', 'not provided']
NM_020921.3(NIN):c.3665A>G (p.Gln1222Arg)	187464517	NIN	[]	[]	['Seckel syndrome 7']
NM_005603.4(ATP8B1):c.2097+2T>C	387906381	ATP8B1	[]	[]	['Progressive intrahepatic

					cholestasis']
NM_005144.4(HR):c.-320T>C	387906382	HR	[]	[]	['Marie Unna hereditary hypotrichosis 1']
NM_001303.3(COX10):c.2T>C (p.Met1Thr)	387906383	COX10	[]	[]	['Congenital myasthenic syndrome, acetazolamide-responsive']
NM_004415.2(DSP):c.4961T>C (p.Leu1654Pro)	749730642	DSP	[]	[]	['not provided']
NM_000392.4(ABCC2):c.1967+2T>C	387906396	ABCC2	[]	[]	['Dubin-Johnson syndrome']
NM_002769.4(PRSS1):c.116T>C (p.Val39Ala)	397507439	-	[]	['TACCAGGYGTCCCTGAATTCTGG']	['Hereditary pancreatitis']
NM_130439.3(MXI1):c.552+2T>C	387906417	MXI1	[]	[]	['Malignant tumor of prostate']
m.8528T>C	387906422	-	[]	[]	['Cardiomyopathy, infantile hypertrophic']
NM_000132.3(F8):c.985T>C (p.Cys329Arg)	387906430	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.1417T>C (p.Tyr473His)	387906443	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.1729T>C (p.Ser577Pro)	387906446	F8	[]	['AAAGAAYCTGTAGATCAAAGAGG']	['Hereditary factor VIII deficiency disease']
NM_004333.4(BRAF):c.1403T>C (p.Phe468Ser)	397507473	BRAF	['ATCATYTGGAACAGTCTACAAGG', 'TCATYTGGAACAGTCTACAAGGG']	['ATCATYTGGAACAGTCTACAAGG', 'TCATYTGGAACAGTCTACAAGGG']	['Cardiofaciocutaneous syndrome', 'Rasopathy']
NM_004333.4(BRAF):c.1454T>C (p.Leu485Ser)	397507475	BRAF	[]	[]	['Cardiofaciocutaneous syndrome', 'Rasopathy']
NM_005188.3(CBL):c.1201T>C (p.Cys401Arg)	397507492	CBL	[]	[]	['Rasopathy']
NM_000133.3(F9):c.52T>C (p.Cys18Arg)	387906474	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_005249.4(FOXG1):c.700T>C (p.Ser234Pro)	786205008	FOXG1	[]	[]	['Rett syndrome, congenital variant', 'not provided']
NM_000133.3(F9):c.82T>C (p.Cys28Arg)	387906481	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_000133.3(F9):c.1031T>C (p.Ile344Thr)	387906482	F9	[]	['ACGAACAYCTTCCTCAAATTTGG']	['Hereditary factor IX deficiency disease']

m.11253T>C	200145866	MT-ND4	[]	[]	['Leber optic atrophy']
NM_000131.4(F7):c.38T>C (p.Leu13Pro)	387906507	F7	[]	[]	['Factor VII deficiency']
NM_000131.4(F7):c.983T>C (p.Phe328Ser)	387906508	F7	[]	['GACGTYCTCTGAGAGGACGCTGG']	['Factor VII deficiency']
NM_000422.2(KRT17):c.296T>C (p.Leu99Pro)	28933089	KRT17	[]	[]	['Pachyonychia congenita type 2', 'not provided']
NM_001040113.1(MYH11):c.3791T>C (p.Leu1264Pro)	387906532	MYH11	[]	['GAAGCYGGAGGCGCAGGTGCAGG']	['Aortic aneurysm, familial thoracic 4']
NM_013246.2(CLCF1):c.46T>C (p.Cys16Arg)	137853934	-	[]	[]	['Cold-induced sweating syndrome 2']
NM_013246.2(CLCF1):c.676T>C (p.Ter226Arg)	137853935	-	[]	[]	['Cold-induced sweating syndrome 2']
NM_001077620.2(PRCD):c.2T>C (p.Met1Thr)	527236092	-	[]	[]	['Retinitis pigmentosa']
NM_000488.3(SERPINC1):c.68T>C (p.Leu23Pro)	387906575	SERPINC1	[]	[]	['Antithrombin III deficiency']
NM_206933.2(USH2A):c.9751T>C (p.Cys3251Arg)	527236118	USH2A	[]	[]	['Retinitis pigmentosa']
NM_001458.4(FLNC):c.752T>C (p.Met251Thr)	387906586	FLNC	[]	[]	['Myopathy, distal, 4']
NM_000781.2(CYP11A1):c.665T>C (p.Leu222Pro)	387906601	CYP11A1	[]	[]	['Adrenal insufficiency, congenital, with 46,XY sex reversal, partial or complete']
NM_000548.3(TSC2):c.2410T>C (p.Cys804Arg)	137853995	TSC2	[]	[]	['Tuberous sclerosis syndrome', 'Tuberous sclerosis 2', 'not provided']
NM_001145661.1(GATA2):c.1117T>C (p.Cys373Arg)	387906633	GATA2	[]	[]	['Lymphedema, primary, with myelodysplasia']
NM_002693.2(POLG):c.3470A>G (p.Asn1157Ser)	548076633	POLG	['CAAGAGGYTGGTGATCTGCAAGG']	['CAAGAGGYTGGTGATCTGCAAGG']	['not provided']
NM_002465.3(MYBPC1):c.706T>C (p.Trp236Arg)	387906657	MYBPC1	[]	[]	['Distal arthrogryposis type 1B']
NM_002465.3(MYBPC1):c.2566T>C (p.Tyr856His)	387906658	MYBPC1	[]	['CAAACCYATATCCGCAGAGTTGG']	['Distal arthrogryposis type 1B']
NM_003392.4(WNT5A):c.544T>C (p.Cys182Arg)	387906663	WNT5A	[]	[]	['Robinow syndrome']
NM_005188.3(CBL):c.1186T>C (p.Cys396Arg)	387906665	CBL	[]	[]	['Rasopathy']

NM_006902.4(PRRX1):c.338T>C (p.Phe113Ser)	387906667	PRRX1	[]	[]	['Dysgnathia complex']
NM_001111035.1(ACP5):c.602T>C (p.Leu201Pro)	387906672	-	[]	[]	['Spondyloenchondrodysplasia with immune dysregulation']
NM_002734.4(PRKAR1A):c.1117T>C (p.Tyr373His)	387906693	PRKAR1A	[]	[]	['Acrodysostosis 1 with or without hormone resistance']
NM_002734.4(PRKAR1A):c.980T>C (p.Ile327Thr)	387906695	PRKAR1A	[]	[]	['Acrodysostosis 1 with or without hormone resistance']
NM_003491.3(NAA10):c.109T>C (p.Ser37Pro)	387906701	NAA10	[]	['TGGCCTTYCCTGGCCCCAGGTGG', 'GGCCTTYCCTGGCCCCAGGTGGG']	['N-terminal acetyltransferase deficiency']
NM_006306.3(SMC1A):c.2351T>C (p.Ile784Thr)	387906702	SMC1A	['AGAYTGGTGTGCGCAACATCCGG']	['AGAYTGGTGTGCGCAACATCCGG']	['Congenital muscular hypertrophy-cerebral syndrome']
NM_000377.2(WAS):c.814T>C (p.Ser272Pro)	387906716	WAS	[]	[]	['Severe congenital neutropenia X-linked']
NM_000377.2(WAS):c.881T>C (p.Ile294Thr)	387906717	WAS	[]	['GACTTCAYTGAGGACCAGGGTGG', 'ACTTCAYTGAGGACCAGGGTGGG']	['Severe congenital neutropenia X-linked']
m.12201T>C	387906733	MT-TH	[]	[]	['Deafness, nonsyndromic sensorineural, mitochondrial']
NM_139125.3(MASP1):c.1888T>C (p.Cys630Arg)	387906753	MASP1	[]	[]	['Michels syndrome']
NM_007315.3(STAT1):c.520T>C (p.Cys174Arg)	387906763	STAT1	[]	[]	['Immunodeficiency 31C']
NM_053025.3(MYLK):c.5275T>C (p.Ser1759Pro)	387906781	-	[]	[]	['Aortic aneurysm, familial thoracic 7']
NM_000287.3(PEX6):c.1601T>C (p.Leu534Pro)	387906809	PEX6	[]	['CTTCYGGGCCGGGACCGTGATGG', 'TTCYGGGCCGGGACCGTGATGGG']	['Peroxisome biogenesis disorder 4B']
NM_000174.4(GP9):c.167T>C (p.Leu56Pro)	28933377	GP9	[]	[]	['Bernard-Soulier syndrome type C']
NM_004153.3(ORC1):c.266T>C (p.Phe89Ser)	387906827	ORC1	[]	[]	['Meier-Gorlin syndrome 1']
NM_021252.4(RAB18):c.619T>C (p.Ter207Gln)	387906833	RAB18	[]	[]	['Warburg micro syndrome 3']
NM_000702.3(ATP1A2):c.2291T>C (p.Leu764Pro)	28933398	ATP1A2	[]	[]	['Familial hemiplegic migraine type 2']
NM_000702.3(ATP1A2):c.2659T>C	28933399	ATP1A2	[]	[]	['Familial hemiplegic migraine

(p.Trp887Arg)					type 2']
NM_000702.3(ATP1A2):c.2192T>C (p.Met731Thr)	28933400	ATP1A2	[]	[]	['Familial hemiplegic migraine type 2']
NM_020433.4(JPH2):c.421T>C (p.Tyr141His)	387906897	JPH2	[]	[]	['Familial hypertrophic cardiomyopathy 17']
NM_173170.1(IL36RN):c.80T>C (p.Leu27Pro)	387906914	IL36RN	[]	[]	['Pustular psoriasis, generalized']
NM_004990.3(MARS):c.1568T>C (p.Ile523Thr)	201555303	MARS	[]	[]	['Interstitial lung and liver disease']
NM_020191.2(MRPS22):c.644T>C (p.Leu215Pro)	387906924	MRPS22	['ATCYTAGGGTAAGGTGACTTAGG']	['ATCYTAGGGTAAGGTGACTTAGG']	['Combined oxidative phosphorylation deficiency 5']
NM_022445.3(TPK1):c.119T>C (p.Leu40Pro)	387906936	TPK1	[]	[]	['THIAMINE METABOLISM DYSFUNCTION SYNDROME 5 (EPISODIC ENCEPHALOPATHY TYPE)']
NM_020634.1(GDF3):c.914T>C (p.Leu305Pro)	387906945	GDF3	[]	[]	['Congenital ocular coloboma', 'Microphthalmia, isolated 7']
NM_024513.3(FYCO1):c.4127T>C (p.Leu1376Pro)	387906965	FYCO1	[]	['CAGCCYGATCCCCATCACTGTGG']	['Cataract, autosomal recessive congenital 2']
NM_006147.3(IRF6):c.65T>C (p.Leu22Pro)	387906967	IRF6	[]	['GCCYCTACCCTGGGCTCATCTGG']	['Van der Woude syndrome', 'Popliteal pterygium syndrome']
NM_025132.3(WDR19):c.2129T>C (p.Leu710Ser)	387906980	WDR19	[]	[]	['Cranioectodermal dysplasia 4', 'SENIOR-LOKEN SYNDROME 8']
NM_025132.3(WDR19):c.20T>C (p.Leu7Pro)	387906982	WDR19	[]	['TCTCACYGCTAGAAAAGACTTGG']	['Asphyxiating thoracic dystrophy 5']
NM_014874.3(MFN2):c.647T>C (p.Phe216Ser)	387906990	MFN2	[]	[]	[]
NM_016097.4(IER3IP1):c.233T>C (p.Leu78Pro)	387907012	IER3IP1	[]	[]	['Microcephaly, epilepsy, and diabetes syndrome']
NM_022489.3(INF2):c.310T>C (p.Cys104Arg)	387907034	INF2	[]	[]	['Charcot-Marie-Tooth disease, dominant intermediate E']
NM_022489.3(INF2):c.383T>C (p.Leu128Pro)	387907037	INF2	[]	[]	['Charcot-Marie-Tooth disease, dominant intermediate E']
NM_003235.4(TG):c.3229T>C	137854433	TG	[]	[]	['Iodotyrosyl coupling defect']

(p.Cys1077Arg)					
NM_058246.3(DNAJB6):c.277T>C (p.Phe93Leu)	387907046	DNAJB6	[]	[]	['Limb-girdle muscular dystrophy, type 1E']
NM_032446.2(MEGF10):c.2320T>C (p.Cys774Arg)	387907072	MEGF10	[]	['GGGCAGYGTACTTGCCGCACTGG']	['Myopathy, areflexia, respiratory distress, and dysphagia, early-onset', 'Myopathy, areflexia, respiratory distress, and dysphagia, early-onset, mild variant']
NM_005502.3(ABCA1):c.4429T>C (p.Cys1477Arg)	137854494	ABCA1	['CCTGTGYGTCCCCAGGGGCAGG', 'CTGTGYGTCCCCAGGGGCAGGG']	['CCTGTGYGTCCCCAGGGGCAGG', 'CTGTGYGTCCCCCAGGGGCAGGG', 'TGTGYGTCCCCCAGGGGCAGGGG', 'GTGYGTCCCCCAGGGGCAGGGGG']	['Tangier disease']
NM_005502.3(ABCA1):c.6026T>C (p.Phe2009Ser)	137854499	ABCA1	[]	['GAGTYCTTTGCCCTTTTGAGAGG']	['Familial hypoalphalipoproteinemia']
NM_015175.2(NBEAL2):c.1163T>C (p.Leu388Pro)	387907113	NBEAL2	[]	[]	['Gray platelet syndrome']
NM_000196.3(HSD11B2):c.1012T>C (p.Tyr338His)	387907117	HSD11B2	[]	['CCGCCGCYATTACCCCGGCCAGG', 'CGCCGCYATTACCCCGGCCAGGG']	['Apparent mineralocorticoid excess']
NM_024599.5(RHBDF2):c.557T>C (p.Ile186Thr)	387907129	RHBDF2	['AGAYTGTGGATCCGCTGGCCCGG']	['AGAYTGTGGATCCGCTGGCCCGG']	['Howel-Evans syndrome']
NM_001077488.3(GNAS):c.299T>C (p.Leu100Pro)	137854531	GNAS	[]	[]	['Pseudohypoparathyroidism type 1A']
NM_001256714.1(DNAAF3):c.386T>C (p.Leu129Pro)	387907151	DNAAF3	[]	[]	['Kartagener syndrome', 'Ciliary dyskinesia, primary, 2']
NM_020894.2(UVSSA):c.94T>C (p.Cys32Arg)	387907164	UVSSA	['AAAATTYGCAAGTATGTCTTAGG']	['AAAATTYGCAAGTATGTCTTAGG', 'AAATTYGCAAGTATGTCTTAGGG']	['UV-sensitive syndrome 3']
NM_000267.3(NF1):c.1523T>C (p.Leu508Pro)	137854558	NF1	[]	[]	['Neurofibromatosis, type 1']
NM_000267.3(NF1):c.6200T>C (p.Leu2067Pro)	137854561	NF1	[]	[]	['Neurofibromatosis, familial spinal']
NM_004453.3(ETFDH):c.1130T>C (p.Leu377Pro)	387907170	ETFDH	[]	['CCAAAACYCACCTTTCCTGGTGG']	[]

NM_000267.3(NF1):c.1070T>C (p.Leu357Pro)	137854563	NF1	[]	[]	['Neurofibromatosis, type 1', 'Neurofibromatosis, familial spinal']
NM_024306.4(FA2H):c.707T>C (p.Phe236Ser)	387907172	FA2H	[]	[]	['Spastic paraplegia 35']
NM_001004127.2(ALG11):c.1142T>C (p.Leu381Ser)	387907182	-	[]	[]	['Congenital disorder of glycosylation type 1P']
NM_021167.4(GATAD1):c.304T>C (p.Ser102Pro)	387907188	GATAD1	[]	[]	['Cardiomyopathy, dilated, 2b']
NM_033360.3(KRAS):c.211T>C (p.Tyr71His)	387907205	KRAS	[]	['GGACCAGYACATGAGGACTGGGG', 'CCAGYACATGAGGACTGGGGAGG', 'CAGYACATGAGGACTGGGGAGGG']	['Cardiofaciocutaneous syndrome 2']
NM_006265.2(RAD21):c.1753T>C (p.Cys585Arg)	387907213	-	[]	[]	['Cornelia de Lange syndrome 4']
NM_000222.2(KIT):c.1859T>C (p.Val620Ala)	387907217	KIT	[]	[]	[]
NM_000335.4(SCN5A):c.5380T>C (p.Tyr1794His)	137854615	SCN5A	[]	[]	['Brugada syndrome', 'Brugada syndrome 1']
NM_000076.2(CDKN1C):c.827T>C (p.Phe276Ser)	387907224	CDKN1C	[]	[]	['Intrauterine growth retardation, metaphyseal dysplasia, adrenal hypoplasia congenita, and genital anomalies']
NM_005691.3(ABCC9):c.3058T>C (p.Ser1020Pro)	387907229	ABCC9	[]	[]	['Hypertrichotic osteochondrodysplasia']
NM_012343.3(NNT):c.2930T>C (p.Leu977Pro)	387907233	NNT	[]	[]	['Glucocorticoid deficiency 4']
NM_024110.4(CARD14):c.467T>C (p.Leu156Pro)	387907240	CARD14	[]	['CAGCAGCYGCAGGAGCACCTGGG']	['Pityriasis rubra pilaris']
NM_002501.3(NFIX):c.179T>C (p.Leu60Pro)	387907254	NFIX	[]	[]	['Sotos syndrome 2']
NM_001165963.1(SCN1A):c.121A>T (p.Lys41Ter)	764444350	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_005022.3(PFN1):c.341T>C (p.Met114Thr)	387907265	PFN1	[]	[]	['Amyotrophic lateral sclerosis 18']
NM_001172567.1(MYD88):c.818T>C (p.Leu273Pro)	387907272	MYD88	[]	[]	['Macroglobulinemia, waldenstrom, somatic']
NM_152296.4(ATP1A3):c.2431T>C (p.Ser811Pro)	387907282	ATP1A3	[]	['TGCCATCYCACTGGCGTACGAGG']	['Alternating hemiplegia of childhood 2']

NM_022787.3(NMNAT1):c.838T>C (p.Ter280Gln)	387907290	NMNAT1	[]	[]	['Leber congenital amaurosis 9']
NM_005120.2(MED12):c.3493T>C (p.Ser1165Pro)	387907361	MED12	[]	['AGGACYCTGAGCCAGGGGCCCGG']	['Ohdo syndrome, X-linked']
NM_006194.3(PAX9):c.62T>C (p.Leu21Pro)	28933970	PAX9	[]	['GGCCGCYGCCCAACGCCATCCGG']	['Tooth agenesis, selective, 3']
NM_000492.3(CFTR):c.2780T>C (p.Leu927Pro)	397508435	CFTR	[]	[]	['Cystic fibrosis']
NM_177976.2(ARL6):c.272T>C (p.Ile91Thr)	137854907	ARL6	[]	[]	['Bardet-Biedl syndrome']

Table 8. Human gene mutations that may be corrected by changing a guanine (G) to an adenine (A). The gene name, gene symbol,and dbSNP database reference number (RS#) are indicated. Also indicated are exemplary protospacers with their PAM sequences (gRNAs and gRNAall) and the base to be edited *e.g.*, a C, indicated by a "Y". The "gRNAs" sequences, from top to bottom, correspond to SEQ ID NOs: 3434-3601. The "gRNA all" sequences, from top to bottom, correspond to SEQ ID NOs: 3602-4266.

Name	**RS# (dbSNP)**	**GeneSymbol**	**gRNAs**	**gRNAall**	**Phenotypes**
NM_000138.4(FBN1):c.3128A>G (p.Lys1043Arg)	137854472	FBN1	[]	['TGCACYTGCCGTGGGTGCAGAGG']	[]
NM_000237.2(LPL):c.953A>G (p.Asn318Ser)	268	LPL	[]	[]	['Hyperlipidemia, familial combined']
NM_000257.3(MYH7):c.2708A>G (p.Glu903Gly)	727504261	MYH7	[]	['AGCGCYCCTCAGCATCTGCCAGG']	['Cardiomyopathy', 'not specified']
NM_000059.3(BRCA2):c.476-2A>G	81002853	BRCA2	[]	['ACCACYGGGGGTAAAAAAAGGGG', 'TACCACYGGGGGTAAAAAAAGGG', 'ATACCACYGGGGGTAAAAAAAGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2', 'Hereditary cancer-predisposing syndrome']
NM_000059.3(BRCA2):c.9118-2A>G	81002862	BRCA2	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2']
NM_000059.3(BRCA2):c.9649-2A>G	81002895	BRCA2	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2']
NM_000387.5(SLC25A20):c.713A>G (p.Gln238Arg)	28934589	SLC25A20	[]	[]	['Carnitine acylcarnitine translocase deficiency', 'not provided']
NM_000060.3(BTD):c.755A>G (p.Asp252Gly)	28934601	BTD	[]	[]	['Biotinidase deficiency']
NM_172107.2(KCNQ2):c.851A>G	28939683	KCNQ2	[]	[]	['Benign familial neonatal seizures 1']

(p.Tyr284Cys)					
NM_006158.4(NEFL):c.293A>G (p.Asn98Ser)	58982919	NEFL	[]	[]	['Charcot-Marie-Tooth disease, type IF', 'not provided']
NM_000019.3(ACAT1):c.473A>G (p.Asn158Ser)	199524907	ACAT1	[]	[]	['Deficiency of acetyl-CoA acetyltransferase']
NM_007294.3(BRCA1):c.4987-2A>G	397509212	BRCA1	[]	[]	['Familial cancer of breast']
NM_007294.3(BRCA1):c.213-12A>G	80358163	BRCA1	[]	[]	['Familial cancer of breast', 'Hereditary breast and ovarian cancer syndrome', 'Breast-ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
NM_001382.3(DPAGT1):c.509A>G (p.Tyr170Cys)	28934876	DPAGT1	['ACAYAGTACAGGATTCCTGCGGG', 'GACAYAGTACAGGATTCCTGCGG']	['ACAYAGTACAGGATTCCTGCGGG', 'GACAYAGTACAGGATTCCTGCGG']	['Congenital disorder of glycosylation type 1J']
NM_032237.4(POMK):c.773A>G (p.Gln258Arg)	397509386	POMK	[]	[]	['Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies, type A12']
NM_201647.2(STAMBP):c.125A>G (p.Glu42Gly)	397509387	STAMBP	[]	[]	['Microcephaly-capillary malformation syndrome']
NM_006876.2(B4GAT1):c.1168A>G (p.Asn390Asp)	397509397	B4GAT1	['CTGATYTTCAGCCTCCTTTTGGG', 'GCTGATYTTCAGCCTCCTTTTGG']	['TGATYTTCAGCCTCCTTTTGGGG', 'CTGATYTTCAGCCTCCTTTTGGG', 'GCTGATYTTCAGCCTCCTTTTGG']	['Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies, type A13']
NM_004315.4(ASAH1):c.965+4A>G	397509415	ASAH1	[]	[]	['Farber lipogranulomatosis']
NM_000257.3(MYH7):c.1477A>G (p.Met493Val)	730880875	MYH7	[]	[]	['Cardiomyopathy']
NM_021020.3(LZTS1):c.355A>G (p.Lys119Glu)	119473032	LZTS1	[]	['CCCTYCTCGGAGCCCTGTAGAGG']	[]
NM_022455.4(NSD1):c.5893-2A>G	587784163	NSD1	[]	[]	['Sotos syndrome 1']
NM_006894.5(FMO3):c.182A>G (p.Asn61Ser)	72549322	FMO3	[]	[]	['Trimethylaminuria']
NM_000403.3(GALE):c.101A>G (p.Asn34Ser)	121908046	GALE	['TGGAAGYTATCGATGACCACAGG	['TGGAAGYTATCGATGACCACAGG']	['UDPglucose-4-epimerase deficiency']

			']		
NM_000314.6(PTEN):c.527A>G (p.Tyr176Cys)	757498880	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000053.3(ATP7B):c.2305A>G (p.Met769Val)	193922103	ATP7B	[]	[]	['Wilson disease', 'not specified', 'not provided']
NM_000173.6(GP1BA):c.763A>G (p.Met255Val)	121908064	GP1BA	[]	[]	['Pseudo von Willebrand disease']
NM_000422.2(KRT17):c.275A>G (p.Asn92Ser)	59151893	KRT17	['GTCAYTGAGGTTCTGCATGGTGG']	['GTCAYTGAGGTTCTGCATGGTGG', 'GCGGTCAYTGAGGTTCTGCATGG']	['Pachyonychia congenita type 2', 'not provided']
NM_000288.3(PEX7):c.854A>G (p.His285Arg)	62653611	PEX7	[]	[]	['Rhizomelic chondrodysplasia punctata type 1']
NM_003742.2(ABCB11):c.890A>G (p.Glu297Gly)	11568372	ABCB11	[]	[]	['Progressive familial intrahepatic cholestasis 2', 'Benign recurrent intrahepatic cholestasis 2']
NM_012243.2(SLC35A3):c.886A>G (p.Ser296Gly)	141952252	SLC35A3	[]	[]	['Arthrogryposis, mental retardation, and seizures']
NM_000061.2(BTK):c.1082A>G (p.Tyr361Cys)	28935478	BTK	['GATGGYAGTTAATGAGCTCAGGG', 'TGATGGYAGTTAATGAGCTCAGG']	['GATGGYAGTTAATGAGCTCAGGG', 'TGATGGYAGTTAATGAGCTCAGG']	[]
NM_000169.2(GLA):c.815A>G (p.Asn272Ser)	28935495	-	[]	[]	['Fabry disease']
NM_016069.9(PAM16):c.226A>G (p.Asn76Asp)	786203989	-	['TCATAGTYCTGCAGAGGAGAGGG']	['CATAGTYCTGCAGAGGAGAGGGG', 'TCATAGTYCTGCAGAGGAGAGGG']	['Chondrodysplasia, megarbane-dagher-melki type']
NM_058163.1(TSR2):c.191A>G (p.Glu64Gly)	786203996	TSR2	[]	[]	['Diamond-Blackfan anemia with microtia and cleft palate', 'DIAMOND-BLACKFAN ANEMIA 14 WITH MANDIBULOFACIAL DYSOSTOSIS']
NM_001098398.1(COPA):c.728A>G (p.Asp243Gly)	794727994	-	[]	[]	[]
NM_005957.4(MTHFR):c.1114A>G (p.Lys372Glu)	786204024	MTHFR	[]	[]	['Homocysteinemia due to MTHFR deficiency']
NM_012193.3(FZD4):c.1024A>G (p.Met342Val)	80358293	-	[]	[]	['Exudative vitreoretinopathy 1']
NM_000186.3(CFH):c	460184	CFH	['CAGYTGA	['CAGYTGAATTTG	['Atypical

.3590T>C (p.Val1197Ala)			ATTTGTGTGTAAACGG']	TGTGTAAACGG']	hemolytic-uremic syndrome 1']
NM_014846.3(KIAA0196):c.1411A>G (p.Asn471Asp)	80338865	KIAA0196	[]	[]	['Spastic paraplegia 8']
NM_014946.3(SPAST):c.1165A>G (p.Thr389Ala)	786204132	SPAST	['AGCATTGYCTTCCCATTCCCAGG']	['ATTGYCTTCCCATTCCCAGGTGG', 'AGCATTGYCTTCCCATTCCCAGG']	['Spastic paraplegia 4, autosomal dominant']
NM_000925.3(PDHB):c.395A>G (p.Tyr132Cys)	28935769	PDHB	[]	[]	['Pyruvate dehydrogenase E1-beta deficiency']
NM_000090.3(COL3A1):c.2338-2A>G	794728050	COL3A1	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_000540.2(RYR1):c.97A>G (p.Lys33Glu)	193922746	RYR1	[]	[]	['King Denborough syndrome', 'not provided']
NM_000540.2(RYR1):c.7043A>G (p.Glu2348Gly)	193922801	RYR1	[]	['TTCYCCTCCACGCTCTCGCCTGG']	['not provided']
NM_000218.2(KCNQ1):c.652A>G (p.Lys218Glu)	36210419	KCNQ1	[]	['GCCCCTYGGAGCCCACGCAGAGG']	['Torsades de pointes', 'Cardiac arrhythmia']
NM_000540.2(RYR1):c.14647-1449A>G	193922886	RYR1	[]	[]	['Minicore myopathy with external ophthalmoplegia', 'not provided']
NM_004429.4(EFNB1):c.472A>G (p.Met158Val)	28936071	EFNB1	[]	[]	['Craniofrontonasal dysplasia']
NM_007254.3(PNKP):c.1029+2T>C	199919568	PNKP	['CCGGYGAGGCCCTGGGGCGGGGG', 'TCCGGYGAGGCCCTGGGGCGGGG']	['CCGGYGAGGCCCTGGGGCGGGGG', 'TCCGGYGAGGCCCTGGGGCGGGG', 'ATCCGGYGAGGCCCTGGGGCGGG', 'GATCCGGYGAGGCCCTGGGGCGG']	['not provided']
NM_005448.2(BMP15):c.704A>G (p.Tyr235Cys)	104894765	BMP15	['ATTGAAAYAGAGTAACAAGAAGG']	['ATTGAAAYAGAGTAACAAGAAGG']	['Ovarian dysgenesis 2']
NM_004415.2(DSP):c.1141-2A>G	794728111	DSP	[]	[]	['not provided']
NM_016035.4(COQ4):c.155T>C (p.Leu52Ser)	786204770	COQ4	['GCTGTYGGCCGCCGGCTCCGCGG']	['GCTGTYGGCCGCCGGCTCCGCGG']	['COENZYME Q10 DEFICIENCY, PRIMARY, 7']
NM_015717.4(CD207):c.790T>C (p.Trp264Arg)	200837270	CD207	[]	[]	['Birbeck granule deficiency']
NM_004771.3(MMP20):c.611A>G (p.His204Arg)	786204826	MMP20	['CGAAAYGTGTATCTCCTCCCAGG']	['CGAAAYGTGTATCTCCTCCCAGG']	['Amelogenesis imperfecta, hypomaturation type, IIA2']
NM_003392.4(WNT5	786204836	WNT5A	[]	[]	['Robinow

A):c.257A>G (p.Tyr86Cys)					syndrome']
NM_000918.3(P4HB): c.1178A>G (p.Tyr393Cys)	786204843	P4HB	[]	[]	['Cole Carpenter syndrome']
NM_000314.6(PTEN): c.139A>G (p.Arg47Gly)	786204855	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000314.6(PTEN): c.320A>G (p.Asp107Gly)	786204858	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000447.2(PSEN2):c.715A>G (p.Met239Val)	28936379	PSEN2	[]	[]	['Alzheimer disease, type 4', 'not provided']
NM_005263.3(GFI1):c .1145A>G (p.Asn382Ser)	28936381	GFI1	[]	[]	['Severe congenital neutropenia 2, autosomal dominant']
NM_005263.3(GFI1):c .1208A>G (p.Lys403Arg)	28936382	GFI1	[]	[]	['Neutropenia, nonimmune chronic idiopathic, of adults']
NM_000314.6(PTEN): c.512A>G (p.Gln171Arg)	786204865	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000314.6(PTEN): c.254-2A>G	786204926	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_001103.3(ACTN2):c.1883A>G (p.Glu628Gly)	786204951	ACTN2	[]	[]	['Familial hypertrophic cardiomyopathy 23']
NM_003159.2(CDKL5):c.-162-2A>G	786204973	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2']
NM_001159287.1(TPI 1):c.622A>G (p.Ile208Val)	121964849	TPI1	[]	[]	['Triosephosphate isomerase deficiency']
NM_003159.2(CDKL5):c.2277-2A>G	786204979	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2']
NM_003159.2(CDKL5):c.458A>G (p.Asp153Gly)	786204985	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2']
NM_003159.2(CDKL5):c.91A>G (p.Arg31Gly)	786204991	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2']
NM_001007792.1(NT RK1):c.986A>G (p.Tyr329Cys)	121964869	NTRK1	[]	[]	['Hereditary insensitivity to pain with anhidrosis']
NM_001007792.1(NT RK1):c.1651A>G (p.Met551Val)	121964870	NTRK1	[]	[]	['Hereditary insensitivity to pain with anhidrosis']
NM_004360.3(CDH1): c.2512A>G (p.Ser838Gly)	121964872	CDH1	[]	[]	['Hereditary diffuse gastric cancer', 'Hereditary cancer-predisposing syndrome', 'Neoplasm of ovary']
NM_003122.4(SPINK	17107315	SPINK1	[]	[]	['Hereditary

1):c.101A>G (p.Asn34Ser)					pancreatitis', 'Tropical calcific pancreatitis', 'Pancreatitis, chronic, susceptibility to']
NM_021098.2(CACNA1H):c.4645A>G (p.Met1549Val)	786205050	CACNA1H	[]	[]	['Primary hyperaldosteronism']
NM_001385.2(DPYS):c.1001A>G (p.Gln334Arg)	121964923	DPYS	[]	[]	['Dihydropyrimidinase deficiency']
NM_003494.3(DYSF):c.3443-33A>G	786205083	DYSF	['GCCAGAGYGAGTGGCTGGAGTGG']	['GCCAGAGYGAGTGGCTGGAGTGG']	['Limb-girdle muscular dystrophy, type 2B']
NM_003816.2(ADAM9):c.411-8A>G	786205086	ADAM9	[]	[]	[]
NM_000532.4(PCCB):c.1304A>G (p.Tyr435Cys)	121964961	PCCB	[]	[]	['Propionic acidemia']
NM_000071.2(CBS):c.1150A>G (p.Lys384Glu)	121964967	CBS	['AACTYGGTCCTGCGGGATGGGGG']	['AACTYGGTCCTGCGGGATGGGGG', 'GAACTYGGTCCTGCGGGATGGGG', 'GGAACTYGGTCCTGCGGGATGGG', 'AGGAACTYGGTCCTGCGGGATGG']	['Homocystinuria, pyridoxine-responsive']
NM_000093.4(COL5A1):c.655-2A>G	786205101	COL5A1	[]	[]	['Ehlers-Danlos syndrome, classic type']
NM_000481.3(AMT):c.125A>G (p.His42Arg)	121964983	AMT	['GCCAGGYGGAAGTCATAGAGCGG']	['GCCAGGYGGAAGTCATAGAGCGG']	['Non-ketotic hyperglycinemia']
NM_000108.4(DLD):c.214A>G (p.Lys72Glu)	121964987	DLD	[]	[]	['Maple syrup urine disease, type 3']
NM_000108.4(DLD):c.1483A>G (p.Arg495Gly)	121964989	DLD	[]	['TTCTCYAAAAGCTTCTGATAAGG']	['Maple syrup urine disease, type 3']
NM_000108.4(DLD):c.1081A>G (p.Met361Val)	121964993	DLD	[]	[]	['Maple syrup urine disease, type 3']
NM_001918.3(DBT):c.1355A>G (p.His452Arg)	121965002	DBT	[]	[]	[]
NM_030813.5(CLPB):c.1850A>G (p.Tyr617Cys)	786205138	CLPB	[]	[]	['3-METHYLGLUTACONIC ACIDURIA WITH CATARACTS, NEUROLOGIC INVOLVEMENT, AND NEUTROPENIA']
NM_006610.3(MASP2):c.359A>G (p.Asp120Gly)	72550870	MASP2	[]	[]	['MASP2 deficiency']

NM_000398.6(CYB5R3):c.719A>G (p.Asp240Gly)	121965018	CYB5R3	[]	[]	['METHEMOGLOBINEMIA, TYPE I']
NM_000095.2(COMP):c.1358A>G (p.Asn453Ser)	28936668	COMP	[]	[]	[]
NM_000095.2(COMP):c.1418A>G (p.Asp473Gly)	28936669	COMP	[]	['ATTGYCGTCGTCGTCGTCGCAGG']	[]
NM_003816.2(ADAM9):c.1396-2A>G	786205151	ADAM9	[]	[]	[]
NM_001204830.1(LIPT1):c.535A>G (p.Thr179Ala)	786205156	-	[]	[]	['LIPOYLTRANSFERASE 1 DEFICIENCY']
NM_000274.3(OAT):c.734A>G (p.Tyr245Cys)	121965046	OAT	[]	[]	['Ornithine aminotransferase deficiency']
NM_018488.2(TBX4):c.1592A>G (p.Gln531Arg)	28936696	TBX4	[]	['GTACYGTAAGGAAGATTCTCGGG', 'GGTACYGTAAGGAAGATTCTCGG']	['Ischiopatellar dysplasia']
NM_001110556.1(FLNA):c.1829-2A>G	786205183	FLNA	[]	[]	['X-linked periventricular heterotopia']
NM_003865.2(HESX1):c.541A>G (p.Thr181Ala)	28936704	HESX1	[]	[]	['Growth hormone deficiency with pituitary anomalies']
NM_000137.2(FAH):c.1141A>G (p.Arg381Gly)	121965077	FAH	[]	['TCCYGGTCTGACCATTCCCCAGG']	['Tyrosinemia type I']
NM_000137.2(FAH):c.836A>G (p.Gln279Arg)	121965078	FAH	[]	[]	['Tyrosinemia type I']
NM_006129.4(BMP1):c.808A>G (p.Met270Val)	786205219	BMP1	[]	[]	['Osteogenesis imperfecta type 13']
NM_001987.4(ETV6):c.1252A>G (p.Arg418Gly)	786205226	ETV6	[]	[]	['Thrombocytopenia 5']
NM_014423.3(AFF4):c.760A>G (p.Thr254Ala)	786205233	AFF4	[]	[]	['CHOPS SYNDROME']
NM_000140.3(FECH):c.68_194del	786205247	FECH	[]	[]	['Erythropoietic protoporphyria']
NM_000138.4(FBN1):c.3344A>G (p.Asp1115Gly)	794728203	FBN1	[]	['ACTCAYCAATATCTGCAAAATGG']	['Thoracic aortic aneurysms and aortic dissections']
NM_198270.3(NHS):c.853-2A>G	786205257	NHS	[]	[]	['Nance-Horan syndrome']
NM_005502.3(ABCA1):c.1790A>G (p.Gln597Arg)	2853578	ABCA1	[]	[]	['Tangier disease']
NM_003002.3(SDHD):c.275A>G (p.Asp92Gly)	786205436	SDHD	[]	['GAATAGYCCATCGCAGAGCAAGG']	['Fatal infantile mitochondrial cardiomyopathy']
NM_005259.2(MSTN):c.458A>G (p.Lys153Arg)	1805086	-	[]	[]	[]
NM_198056.2(SCN5A	1805124	SCN5A	[]	[]	['Progressive

):c.1673A>G (p.His558Arg)					familial heart block type 1A', 'not specified', 'not provided']
NM_000784.3(CYP27A1):c.776A>G (p.Lys259Arg)	72551317	CYP27A1	[]	['AGTCCACYTGGGGAGGAAGGTGG']	['Cholestanol storage disease']
NM_000784.3(CYP27A1):c.1061A>G (p.Asp354Gly)	72551320	CYP27A1	[]	[]	['Cholestanol storage disease']
NM_000463.2(UGT1A1):c.992A>G (p.Gln331Arg)	72551348	-	[]	[]	['Crigler-Najjar syndrome, type II']
NM_000463.2(UGT1A1):c.1070A>G (p.Gln357Arg)	72551351	-	[]	[]	['Crigler Najjar syndrome, type 1']
NM_016218.2(POLK):c.1385A>G (p.Asn462Ser)	786205687	POLK	[]	['ATTCACAYTCTTCAACTTAATGG']	['Malignant tumor of prostate']
NM_016218.2(POLK):c.181A>G (p.Asn61Asp)	786205689	POLK	[]	[]	['Malignant tumor of prostate']
NM_016218.2(POLK):c.1477A>G (p.Lys493Glu)	786205692	POLK	[]	[]	['Malignant tumor of prostate']
NM_000138.4(FBN1):c.7916A>G (p.Tyr2639Cys)	794728280	FBN1	[]	['TGTTCAYACTGGAAGCCGGCGGG', 'CTGTTCAYACTGGAAGCCGGCGG']	['Thoracic aortic aneurysms and aortic dissections']
NM_000132.3(F8):c.1331A>G (p.Lys444Arg)	28937272	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000531.5(OTC):c.268A>G (p.Ser90Gly)	72554340	OTC	[]	[]	['not provided']
NM_005502.3(ABCA1):c.2804A>G (p.Asn935Ser)	28937313	ABCA1	['TCCAYTGTGGCCCAGGAAGGAGG']	['TCCAYTGTGGCCCAGGAAGGAGG', 'CGCTCCAYTGTGGCCCAGGAAGG']	['Tangier disease']
NM_004380.2(CREBBP):c.3524A>G (p.Tyr1175Cys)	28937315	CREBBP	[]	[]	['Rubinstein-Taybi syndrome']
NM_000335.4(SCN5A):c.3971A>G (p.Asn1324Ser)	28937317	SCN5A	[]	['GCAYTGACCACCACCTCAAGTGG']	['Long QT syndrome 3', 'Congenital long QT syndrome']
NM_000891.2(KCNJ2):c.901A>G (p.Met301Val)	786205818	KCNJ2	[]	[]	['Cardiac arrhythmia']
NM_144499.2(GNAT1):c.386A>G (p.Asp129Gly)	786205854	GNAT1	[]	['CGGAGYCCTTCCACAGCCGCTGG']	['NIGHT BLINDNESS, CONGENITAL STATIONARY, TYPE 1G']
NM_000523.3(HOXD13):c.974A>G (p.Gln325Arg)	104893635	HOXD13	[]	[]	['Syndactyly type 5']
NM_013953.3(PAX8):c.160A>G (p.Ser54Gly)	104893660	-	[]	[]	['Thyroid agenesis']
NM_014585.5(SLC40	104893663	SLC40A1	[]	[]	['Hemochromatosis

A1):c.470A>G (p.Asp157Gly)					type 4']
NM_003124.4(SPR):c.448A>G (p.Arg150Gly)	104893665	SPR	[]	[]	['Sepiapterin reductase deficiency']
NM_000258.2(MYL3):c.445A>G (p.Met149Val)	104893748	MYL3	[]	[]	['Familial hypertrophic cardiomyopathy 8', 'Cardiomyopathy', 'Hypertrophic cardiomyopathy']
NM_000539.3(RHO):c.533A>G (p.Tyr178Cys)	104893776	RHO	[]	['GGATGYACCTGAGGACAGGCAGG']	['Retinitis pigmentosa 4']
NM_000539.3(RHO):c.569A>G (p.Asp190Gly)	104893777	RHO	[]	[]	['Retinitis pigmentosa 4']
NM_000539.3(RHO):c.44A>G (p.Asn15Ser)	104893786	RHO	[]	[]	['Retinitis pigmentosa', 'Retinitis pigmentosa 4']
NM_001814.4(CTSC):c.1235A>G (p.Tyr412Cys)	28937571	CTSC	[]	[]	['Periodontitis, aggressive, 1']
NM_001701.3(BAAT):c.226A>G (p.Met76Val)	28937579	BAAT	[]	[]	['Hypercholanemia, familial']
NM_001257342.1(BCS1L):c.232A>G (p.Ser78Gly)	28937590	BCS1L	[]	['GACACYGAGGTGCTGAGTACGGG', 'CGACACYGAGGTGCTGAGTACGG']	['GRACILE syndrome']
NM_004407.3(DMP1):c.1A>G (p.Met1Val)	104893834	DMP1	[]	[]	['Autosomal recessive hypophosphatemic vitamin D refractory rickets']
NM_000406.2(GNRHR):c.317A>G (p.Gln106Arg)	104893836	GNRHR	[]	[]	['Hypogonadotropic hypogonadism']
NM_172250.2(MMAA):c.620A>G (p.Tyr207Cys)	104893849	MMAA	[]	[]	['Methylmalonic aciduria cblA type']
NM_000320.2(QDPR):c.449A>G (p.Tyr150Cys)	104893866	QDPR	[]	['TGCCGYACCCGATCATACCTGGG', 'ATGCCGYACCCGATCATACCTGG']	['Dihydropteridine reductase deficiency']
NM_015629.3(PRPF31):c.527+3A>G	587776590	PRPF31	[]	['GACAYACCCCTGGGTGGTGGAGG', 'GCGGACAYACCCCTGGGTGGTGG']	['Retinitis pigmentosa 11']
NM_000112.3(SLC26A2):c.1273A>G (p.Asn425Asp)	104893920	SLC26A2	[]	[]	['Diastrophic dysplasia', 'Achondrogenesis, type IB']
NM_000344.3(SMN1):c.815A>G (p.Tyr272Cys)	104893922	SMN1	[]	[]	['Werdnig-Hoffmann disease']
NM_000344.3(SMN1):c.784A>G	104893932	SMN1	[]	[]	['Kugelberg-Welander disease']

(p.Ser262Gly)					
NM_000409.3(GUCA1A):c.296A>G (p.Tyr99Cys)	104893967	GUCA1A	[]	[]	['Cone dystrophy 3']
NM_182548.3(LHFPL5):c.380A>G (p.Tyr127Cys)	104893975	LHFPL5	[]	[]	['Deafness, autosomal recessive 67']
NM_001024630.3(RUNX2):c.598A>G (p.Thr200Ala)	104893993	RUNX2	[]	[]	['Cleidocranial dysostosis', 'Cleidocranial dysplasia, forme fruste, dental anomalies only']
NM_000162.3(GCK):c.641A>G (p.Tyr214Cys)	104894015	GCK	[]	['GTAGYAGCAGGAGATCATCGTGG']	['Hyperinsulinemic hypoglycemia familial 3']
NM_001002010.2(NT5C3A):c.686A>G (p.Asn229Ser)	104894028	NT5C3A	[]	[]	['Uridine 5-prime monophosphate hydrolase deficiency, hemolytic anemia due to']
NM_203288.1(RP9):c.509A>G (p.Asp170Gly)	104894039	RP9	[]	[]	['Retinitis pigmentosa 9']
NM_000474.3(TWIST1):c.466A>G (p.Ile156Val)	104894059	TWIST1	[]	[]	['Saethre-Chotzen syndrome']
NM_000532.4(PCCB):c.1606A>G (p.Asn536Asp)	202247823	PCCB	[]	['ATATYTGCATGTTTTCTCCAAGG']	['Propionic acidemia', 'not provided']
NM_021615.4(CHST6):c.521A>G (p.Lys174Arg)	28937877	CHST6	[]	[]	['Macular corneal dystrophy Type I']
NM_178138.4(LHX3):c.332A>G (p.Tyr111Cys)	104894117	LHX3	['CAGGTGGYACACGAAGTCCTGGG']	['CAGGTGGYACACGAAGTCCTGGG']	['Pituitary hormone deficiency, combined 3']
NM_004897.4(MINPP1):c.809A>G (p.Gln270Arg)	104894171	MINPP1	[]	[]	['Thyroid cancer, follicular']
NM_000073.2(CD3G):c.1A>G (p.Met1Val)	104894199	CD3G	[]	['CCAYGTCAGTCTCTGTCCTCCGG']	['Immunodeficiency 17']
NM_001885.2(CRYAB):c.358A>G (p.Arg120Gly)	104894201	CRYAB	[]	[]	['Alpha-B crystallinopathy']
NM_001814.4(CTSC):c.857A>G (p.Gln286Arg)	104894208	CTSC	[]	['CTCCYGAGGGCTTAGGATTGGGG', 'CCTCCYGAGGGCTTAGGATTGGG', 'ACCTCCYGAGGGCTTAGGATTGG']	['Papillon-Lef\xc3\xa8vre syndrome', 'Haim-Munk syndrome']
NM_001814.4(CTSC):c.1040A>G (p.Tyr347Cys)	104894211	CTSC	[]	['TCCTACAYAGTGGTACTCAGAGG']	['Papillon-Lef\xc3\xa8vre syndrome', 'Periodontitis, aggressive, 1']
NM_012193.3(FZD4):c.766A>G	104894223	-	['GAAATAYGATGGGGC	['GAAATAYGATGGGGCGCTCAGGG',	['Retinopathy of prematurity']

(p.Ile256Val)			GCTCAGGG', 'AGAAATAYGATGGGGCGCTCAGG']	'AGAAATAYGATGGGGCGCTCAGG']	
NM_005343.2(HRAS):c.350A>G (p.Lys117Arg)	104894227	-	[]	[]	['Costello syndrome']
NM_145014.2(HYLS1):c.632A>G (p.Asp211Gly)	104894232	-	[]	[]	['Hydrolethalus syndrome']
NM_000525.3(KCNJ11):c.776A>G (p.His259Arg)	104894248	KCNJ11	['GACAYGGTAGATGATCAGCGGGG']	['GACAYGGTAGATGATCAGCGGGG', 'TGACAYGGTAGATGATCAGCGGG', 'ATGACAYGGTAGATGATCAGCGG']	['Islet cell hyperplasia']
NM_000317.2(PTS):c.155A>G (p.Asn52Ser)	104894275	PTS	['TAAYTGTGCCCATGGCCATTTGG']	['TAAYTGTGCCCATGGCCATTTGG']	['6-pyruvoyl-tetrahydropterin synthase deficiency']
NM_000317.2(PTS):c.139A>G (p.Asn47Asp)	104894278	PTS	[]	[]	['Hyperphenylalaninemia, bh4-deficient, a, due to partial pts deficiency']
NM_000317.2(PTS):c.347A>G (p.Asp116Gly)	104894279	PTS	[]	[]	['Hyperphenylalaninemia, bh4-deficient, a, due to partial pts deficiency']
NM_022051.2(EGLN1):c.1121A>G (p.His374Arg)	119476045	EGLN1	[]	[]	['Erythrocytosis, familial, 3']
NM_015915.4(ATL1):c.773A>G (p.His258Arg)	119476048	ATL1	[]	[]	['Spastic paraplegia 3']
NM_000448.2(RAG1):c.2735A>G (p.Tyr912Cys)	104894290	RAG1	[]	['CTGYACTGGCAGAGGGATTCTGG']	['Histiocytic medullary reticulosis']
NM_000448.2(RAG1):c.1286A>G (p.Asp429Gly)	104894292	RAG1	[]	[]	['Histiocytic medullary reticulosis']
NM_003002.3(SDHD):c.341A>G (p.Tyr114Cys)	104894304	SDHD	[]	[]	['Hereditary Paraganglioma-Pheochromocytoma Syndromes', 'Paragangliomas 1']
NM_015141.3(GPD1L):c.370A>G (p.Ile124Val)	72552293	GPD1L	[]	[]	['Brugada syndrome 2', 'Primary familial hypertrophic cardiomyopathy', 'Long QT syndrome', 'Sudden infant death syndrome', 'Cardiomyopathy']
NM_020661.2(AICDA):c.415A>G (p.Met139Val)	104894322	AICDA	[]	[]	['Immunodeficiency with hyper IgM type 2']

NM_014365.2(HSPB8):c.421A>G (p.Lys141Glu)	104894351	HSPB8	[]	[]	['Charcot-Marie-Tooth disease', 'Distal hereditary motor neuronopathy type 2A']
NM_000217.2(KCNA1):c.676A>G (p.Thr226Ala)	104894354	KCNA1	[]	['GCGYTTCCACGATGAAGAAGGGG', 'AGCGYTTCCACGATGAAGAAGGG', 'CAGCGYTTCCACGATGAAGAAGG']	['Episodic ataxia type 1']
NM_014239.3(EIF2B2):c.638A>G (p.Glu213Gly)	104894425	EIF2B2	[]	['AGTTGTCYCAATACCTGCTTTGG']	['Leukoencephalopathy with vanishing white matter', 'Ovarioleukodystrophy']
NM_002408.3(MGAT2):c.785A>G (p.His262Arg)	104894447	MGAT2	[]	[]	['Carbohydrate-deficient glycoprotein syndrome type II']
NM_002408.3(MGAT2):c.952A>G (p.Asn318Asp)	104894448	MGAT2	[]	[]	['Carbohydrate-deficient glycoprotein syndrome type II']
NM_000270.3(PNP):c.383A>G (p.Asp128Gly)	104894450	PNP	[]	['ATAYCTCCAACCTCAAACTTGGG', 'GATAYCTCCAACCTCAAACTTGG']	['Purine-nucleoside phosphorylase deficiency']
NM_000270.3(PNP):c.575A>G (p.Tyr192Cys)	104894452	PNP	[]	[]	['Purine-nucleoside phosphorylase deficiency']
NM_005982.3(SIX1):c.386A>G (p.Tyr129Cys)	104894478	SIX1	[]	[]	['Melnick-Fraser syndrome', 'Branchiootic syndrome 3']
NM_000101.3(CYBA):c.281A>G (p.His94Arg)	104894510	CYBA	[]	[]	['Granulomatous disease, chronic, autosomal recessive, cytochrome b-negative']
NM_024887.3(DHDDS):c.124A>G (p.Lys42Glu)	147394623	DHDDS	[]	['GGCACTYCTTGGCATAGCGACGG']	['Retinitis pigmentosa 59']
NM_024006.5(VKORC1):c.172A>G (p.Arg58Gly)	104894541	VKORC1	[]	[]	['Warfarin response']
NM_001128085.1(ASPA):c.692A>G (p.Tyr231Cys)	104894550	-	[]	[]	['Spongy degeneration of central nervous system']
NM_001128085.1(ASPA):c.71A>G (p.Glu24Gly)	104894551	-	[]	[]	['Spongy degeneration of central nervous system']
NM_000019.3(ACAT1):c.472A>G (p.Asn158Asp)	148639841	ACAT1	[]	[]	['Deficiency of acetyl-CoA acetyltransferase', 'not provided']
NM_014254.2(TMEM5):c.1016A>G	150736997	TMEM5	[]	[]	['Congenital muscular dystrophy-

(p.Tyr339Cys)					dystroglycanopathy with brain and eye anomalies, type A10']
NM_005557.3(KRT16):c.374A>G (p.Asn125Ser)	60723330	KRT16	[]	['GCGGTCAYTGAGGTTCTGCATGG']	['Pachyonychia congenita, type 1', 'Palmoplantar keratoderma, nonepidermolytic, focal', 'not provided']
NM_005450.4(NOG):c.665A>G (p.Tyr222Cys)	104894602	NOG	[]	[]	['Tarsal carpal coalition syndrome', 'Cushing symphalangism']
NM_030665.3(RAI1):c.4685A>G (p.Gln1562Arg)	104894634	RAI1	[]	['CTGCTGCYGTCGTCGTCGCTTGG']	['Smith-Magenis syndrome']
NM_000346.3(SOX9):c.517A>G (p.Lys173Glu)	104894647	SOX9	[]	[]	['Acampomelic campomelic dysplasia']
NM_024301.4(FKRP):c.926A>G (p.Tyr309Cys)	104894679	FKRP	[]	[]	['Congenital muscular dystrophy-dystroglycanopathy without mental retardation, type B5']
NM_000495.4(COL4A5):c.2394A>G (p.Lys798=)	281874691	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_001604.5(PAX6):c.1075-2A>G	794726661	PAX6	[]	[]	['Congenital aniridia']
NM_000363.4(TNNI3):c.569A>G (p.Asp190Gly)	104894728	TNNI3	[]	[]	['Familial restrictive cardiomyopathy 1', 'Familial hypertrophic cardiomyopathy 7']
NM_000363.4(TNNI3):c.532A>G (p.Lys178Glu)	104894730	TNNI3	[]	['CCTYCTTCACCTGCTTGAGGTGG', 'CCTCCTYCTTCACCTGCTTGAGG']	['Familial restrictive cardiomyopathy 1']
NM_000054.4(AVPR2):c.839A>G (p.Tyr280Cys)	104894752	AVPR2	[]	[]	['Nephrogenic diabetes insipidus, X-linked']
NM_000074.2(CD40LG):c.386A>G (p.Glu129Gly)	104894772	CD40LG	[]	[]	['Immunodeficiency with hyper IgM type 1']
NM_000495.4(COL4A5):c.4977-2A>G	281874752	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_001165963.1(SCN1A):c.5264A>G (p.Asp1755Gly)	794726722	-	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000495.4(COL4A5):c.547-2A>G	281874756	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.610-2A>G	281874758	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_032520.4(GNPTG):c.610-2A>G	193302855	GNPTG	['CCCYGAAGGTGGAGGATGCAGGG']	['CCCYGAAGGTGGAGGATGCAGGG', 'GCCCYGAAGGTG	['Mucolipidosis III Gamma']

				GAGGATGCAGG']	
NM_002049.3(GATA1):c.653A>G (p.Asp218Gly)	104894816	GATA1	[]	['GTCCTGYCCCTCCGCCACAGTGG']	['GATA-1-related thrombocytopenia with dyserythropoiesis']
NM_000157.3(GBA):c.680A>G (p.Asn227Ser)	364897	GBA	['CCAYTGGTCTTGAGCCAAGTGGG', 'TCCAYTGGTCTTGAGCCAAGTGG']	['CCAYTGGTCTTGAGCCAAGTGGG', 'TCCAYTGGTCTTGAGCCAAGTGG']	['Gaucher disease', 'Subacute neuronopathic Gaucher disease', 'Gaucher disease, type 1']
NM_001097642.2(GJB1):c.194A>G (p.Tyr65Cys)	104894819	GJB1	[]	[]	['X-linked hereditary motor and sensory neuropathy']
NM_000166.5(GJB1):c.614A>G (p.Asn205Ser)	104894822	GJB1	[]	[]	['X-linked hereditary motor and sensory neuropathy']
NM_001165963.1(SCN1A):c.747T>G (p.Asp249Glu)	773407463	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000169.2(GLA):c.886A>G (p.Met296Val)	104894830	-	[]	[]	['Fabry disease', 'Fabry disease, cardiac variant']
NM_000169.2(GLA):c.101A>G (p.Asn34Ser)	104894835	-	[]	[]	['Fabry disease']
NM_001165963.1(SCN1A):c.1662+3A>G	794726773	SCN1A	[]	['GTGCCAYACCTGGTGTGGGGAGG']	['Severe myoclonic epilepsy in infancy']
NM_000169.2(GLA):c.1228A>G (p.Thr410Ala)	104894852	-	[]	[]	['Fabry disease']
NM_000202.6(IDS):c.404A>G (p.Lys135Arg)	104894861	IDS	[]	['AAAGACTYTTCCCACCGACATGG']	['Mucopolysaccharidosis, MPS-II']
NM_001165963.1(SCN1A):c.383+1A>G	794726803	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000266.3(NDP):c.131A>G (p.Tyr44Cys)	104894870	NDP	[]	[]	['Atrophia bulborum hereditaria']
NM_000266.3(NDP):c.125A>G (p.His42Arg)	104894874	NDP	[]	['TGGYGCCTCATGCAGCGTCGAGG']	[]
NM_001128227.2(GNE):c.604A>G (p.Met202Val)	121908634	GNE	[]	[]	['Inclusion body myopathy 2']
NM_001165963.1(SCN1A):c.3880-2A>G	794726816	-	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_001165963.1(SCN1A):c.1046A>G (p.Tyr349Cys)	794726844	SCN1A	['ACATAYATCCCTCTGGACATTGG']	['ACATAYATCCCTCTGGACATTGG']	['Severe myoclonic epilepsy in infancy']
NM_001015877.1(PHF6):c.700A>G (p.Lys234Glu)	104894917	PHF6	[]	[]	['Borjeson-Forssman-Lehmann syndrome']
NM_001015877.1(PHF6):c.686A>G (p.His229Arg)	104894918	PHF6	[]	[]	['Borjeson-Forssman-Lehmann syndrome']
NM_001015877.1(PHF6):c.769A>G (p.Arg257Gly)	104894919	PHF6	['CTCYTGATGTTGTTGTGAGCTGG']	['CTCYTGATGTTGTTGTGAGCTGG']	['Borjeson-Forssman-Lehmann syndrome']

NM_000307.4(POU3F4):c.1000A>G (p.Lys334Glu)	104894922	POU3F4	[]	[]	['Deafness, X-linked 2']
NM_000330.3(RS1):c.667T>C (p.Cys223Arg)	104894929	-	[]	[]	['Juvenile retinoschisis']
NM_003413.3(ZIC3):c.1213A>G (p.Lys405Glu)	104894962	ZIC3	['TGTGTTYGCGCAGGGAGCTCGGG', 'ATGTGTTYGCGCAGGGAGCTCGG']	['TGTGTTYGCGCAGGGAGCTCGGG', 'ATGTGTTYGCGCAGGGAGCTCGG']	['Heterotaxy, visceral, X-linked']
NM_002420.5(TRPM1):c.296T>C (p.Leu99Pro)	191205969	TRPM1	[]	['AAGCYCTTAATATCTGTGCATGG']	['Congenital stationary night blindness, type 1C']
NM_004006.2(DMD):c.1150-2A>G	794727030	DMD	[]	[]	['Duchenne muscular dystrophy', 'Becker muscular dystrophy']
NM_203290.2(POLR1C):c.221A>G (p.Asn74Ser)	371802902	POLR1C	[]	[]	['LEUKODYSTROPHY, HYPOMYELINATING, 11']
NM_019109.4(ALG1):c.1188-2A>G	794727073	ALG1	[]	['TAAACYGCAGAGAGAACCAAGGG', 'GTAAACYGCAGAGAGAACCAAGG']	['Congenital disorder of glycosylation type 1K']
NM_004463.2(FGD1):c.2016-2A>G	794727099	FGD1	[]	[]	['Aarskog syndrome']
NM_024110.4(CARD14):c.425A>G (p.Glu142Gly)	281875213	CARD14	[]	[]	['Psoriasis susceptibility 2', 'not provided']
NM_001004334.3(GPR179):c.659A>G (p.Tyr220Cys)	281875236	GPR179	[]	['CCCACAYATCCATCTGCCTGCGG']	['Congenital stationary night blindness, type 1E', 'not provided']
NM_018965.3(TREM2):c.401A>G (p.Asp134Gly)	28939079	TREM2	['CCGGTGAYCCAGGGGGTCTATGG']	['TGAYCCAGGGGGTCTATGGGAGG', 'CGGTGAYCCAGGGGGTCTATGGG', 'CCGGTGAYCCAGGGGGTCTATGG']	['Polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy']
NM_015915.4(ATL1):c.1222A>G (p.Met408Val)	28939094	ATL1	[]	['CACCCAYCTTCTTCACCCCTCGG']	['Spastic paraplegia 3']
NM_002437.4(MPV17):c.186+2T>C	147952488	MPV17	['TGGYAAGTTCTCCCCTCAACAGG']	['TGGYAAGTTCTCCCCTCAACAGG']	['Navajo neurohepatopathy', 'not provided']
NM_005359.5(SMAD4):c.1500A>G (p.Ile500Met)	281875320	SMAD4	['TGAGYATGCATAAGCGACGAAGG']	['TGAGYATGCATAAGCGACGAAGG']	['Myhre syndrome', 'not provided']
NM_005359.5(SMAD4):c.1498A>G (p.Ile500Val)	281875322	SMAD4	['TGAGTAYGCATAAGCGACGAAGG']	['TGAGTAYGCATAAGCGACGAAGG']	['Hereditary cancer-predisposing syndrome', 'Myhre syndrome', 'not provided']

NM_005359.5(SMAD4):c.989A>G (p.Glu330Gly)	281875324	SMAD4	[]	['ATCCATTYCAAAGTAAGCAATGG']	['Juvenile polyposis syndrome', 'Hereditary cancer-predisposing syndrome', 'not provided']
NM_000342.3(SLC4A1):c.166A>G (p.Lys56Glu)	5036	SLC4A1	[]	[]	[]
NM_000518.4(HBB):c.*113A>G	33985472	HBB	[]	[]	[]
NM_001127255.1(NLRP7):c.2738A>G (p.Asn913Ser)	104895503	-	['TCTGGYTGATACTCAAGTCCAGG']	['TCTGGYTGATACTCAAGTCCAGG']	['Hydatidiform mole']
NM_000037.3(ANK1):c.-108T>C	77173848	ANK1	[]	['GGGCCYGGCCCGCACGTCACAGG']	['Spherocytosis, type 1, autosomal recessive']
NM_201631.3(TGM5):c.763T>C (p.Trp255Arg)	115677373	TGM5	['TGCGGAGYGGACGGGCAGCGTGG']	['TGCGGAGYGGACGGGCAGCGTGG']	['Peeling skin syndrome, acral type']
NM_020435.3(GJC2):c.-167A>G	587776888	GJC2	[]	[]	['Leukodystrophy, hypomyelinating, 2']
NM_130466.3(UBE3B):c.1A>G (p.Met1Val)	672601304	UBE3B	[]	[]	['Kaufman oculocerebrofacial syndrome']
NM_022124.5(CDH23):c.146-2A>G	794727649	-	[]	[]	['Usher syndrome, type 1D']
NM_014191.3(SCN8A):c.667A>G (p.Arg223Gly)	672601319	SCN8A	[]	[]	['Early infantile epileptic encephalopathy 13']
NM_001164405.1(BHLHA9):c.211A>G (p.Asn71Asp)	672601337	BHLHA9	[]	[]	['Syndactyly type 9']
NM_021830.4(C10orf2):c.1754A>G (p.Asn585Ser)	672601360	C10orf2	[]	[]	['Perrault syndrome 5']
NM_002887.3(RARS):c.5A>G (p.Asp2Gly)	672601372	RARS	[]	[]	['Leukodystrophy, hypomyelinating, 9']
NM_002887.3(RARS):c.1A>G (p.Met1Val)	672601375	RARS	[]	[]	['Leukodystrophy, hypomyelinating, 9']
NM_001943.3(DSG2):c.1880-2A>G	397514038	DSG2	[]	[]	['Arrhythmogenic right ventricular cardiomyopathy, type 10', 'Cardiomyopathy']
NM_024422.4(DSC2):c.631-2A>G	397514042	DSC2	[]	[]	['Arrhythmogenic right ventricular cardiomyopathy, type 11', 'Cardiomyopathy']
NM_001159772.1(CANT1):c.671T>C (p.Leu224Pro)	150181226	CANT1	[]	['CGTCYGTACGTGGGCGGCCTGGG', 'GCGTCYGTACGTGGGCGGCCTGG']	['Desbuquois syndrome']
NM_031418.2(ANO3):c.2053A>G (p.Ser685Gly)	587776923	ANO3	[]	[]	['Dystonia 24']

NM_014191.3(SCN8A):c.5302A>G (p.Asn1768Asp)	202151337	SCN8A	[]	[]	['Early infantile epileptic encephalopathy 13']
NM_000367.3(TPMT):c.719A>G (p.Tyr240Cys)	1142345	TPMT	[]	[]	['Thiopurine methyltransferase deficiency']
NM_003907.2(EIF2B5):c.271A>G (p.Thr91Ala)	28939717	EIF2B5	['AAATGYTTCCTGTACACCTGTGG']	['AAATGYTTCCTGTACACCTGTGG']	['Leukoencephalopathy with vanishing white matter']
NM_004006.2(DMD):c.10554-2A>G	794727890	DMD	[]	[]	['Duchenne muscular dystrophy', 'Becker muscular dystrophy', 'Dilated cardiomyopathy 3B']
NM_000084.4(CLCN5):c.815A>G (p.Tyr272Cys)	273585644	CLCN5	[]	[]	['Dent disease 1']
NM_000084.4(CLCN5):c.1637A>G (p.Lys546Arg)	273585649	CLCN5	[]	[]	['Dent disease 1']
NM_000041.3(APOE):c.237-2A>G	397514253	APOE	[]	['CGCCCYGCGGCCGAGAGGGCGGG', 'GCGCCCYGCGGCCGAGAGGGCGG']	['Familial type 3 hyperlipoproteinemia']
NM_000155.3(GALT):c.940A>G (p.Asn314Asp)	2070074	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase', 'not provided']
NM_005045.3(RELN):c.2288A>G (p.Asp763Gly)	794727998	RELN	[]	[]	['EPILEPSY, FAMILIAL TEMPORAL LOBE, 7']
NM_001914.3(CYB5A):c.130-2A>G	794728010	CYB5A	[]	[]	['Methemoglobinemia type 4']
NM_001613.2(ACTA2):c.1A>G (p.Met1Val)	794728019	ACTA2	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_000109.3(DMD):c.1700T>C (p.Leu567Pro)	370644567	DMD	[]	[]	['Becker muscular dystrophy', 'Exertional myalgia, muscle stiffness and myoglobinuria', 'not provided']
NM_000060.3(BTD):c.194A>G (p.His65Arg)	397514341	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c.278A>G (p.Tyr93Cys)	397514348	BTD	[]	['GTTCAYAGATGTCAAGGTTCTGG']	['Biotinidase deficiency']
NM_000060.3(BTD):c.356A>G (p.Asn119Ser)	397514353	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c.364A>G (p.Arg122Gly)	397514354	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c.515A>G (p.Asn172Ser)	397514366	BTD	[]	[]	['Biotinidase deficiency']
NM_000060.3(BTD):c	397514370	BTD	[]	[]	['Biotinidase

.583A>G (p.Asn195Asp)					deficiency']
NM_000060.3(BTD):c.584A>G (p.Asn195Ser)	397514371	BTD	[]	[]	['Biotinidase deficiency', 'not provided']
NM_000060.3(BTD):c.641A>G (p.Asn214Ser)	397514377	BTD	['AGAGGYTGTGTTTACGGTAGCGG']	['AGAGGYTGTGTTTACGGTAGCGG']	['Biotinidase deficiency']
NM_000061.2(BTK):c.1288A>G (p.Lys430Glu)	128620184	BTK	['TCTYGATGGCCACGTCGTACTGG']	['TCTYGATGGCCACGTCGTACTGG']	['X-linked agammaglobulinemia']
NM_001002294.2(FMO3):c.923A>G (p.Glu308Gly)	2266780	FMO3	[]	[]	['Trimethylaminuria']
m.15579A>G	207460002	MT-CYB	[]	[]	['Multisystem disorder']
NM_000060.3(BTD):c.1313A>G (p.Tyr438Cys)	397514415	BTD	[]	['GGCAYACAGCTCTTTGGATAAGG']	['Biotinidase deficiency']
NM_000060.3(BTD):c.1619A>G (p.Tyr540Cys)	397514431	BTD	[]	[]	['Biotinidase deficiency']
NM_000023.2(SGCA):c.410A>G (p.Glu137Gly)	397514451	SGCA	[]	[]	['Limb-girdle muscular dystrophy, type 2D']
NM_004813.2(PEX16):c.992A>G (p.Tyr331Cys)	397514472	PEX16	['AAGYAGATTTTCTGCCAGGTGGG', 'GAAGYAGATTTTCTGCCAGGTGG', 'GTAGAAGYAGATTTTCTGCCAGG']	['AAGYAGATTTTCTGCCAGGTGGG', 'GAAGYAGATTTTCTGCCAGGTGG', 'GTAGAAGYAGATTTTCTGCCAGG']	['Peroxisome biogenesis disorder 8B']
NM_000933.3(PLCB4):c.1868A>G (p.Tyr623Cys)	397514480	PLCB4	[]	[]	['Auriculocondylar syndrome 1', 'Auriculocondylar syndrome 2']
NM_005340.6(HINT1):c.152A>G (p.His51Arg)	397514491	HINT1	['AAAAYGTGTTGGTGCTTGAGGGG', 'GAAAAYGTGTTGGTGCTTGAGGG', 'AGAAAAYGTGTTGGTGCTTGAGG']	['AAAAYGTGTTGGTGCTTGAGGGG', 'GAAAAYGTGTTGGTGCTTGAGGG', 'AGAAAAYGTGTTGGTGCTTGAGG']	['Gamstorp-Wohlfart syndrome']
NM_007171.3(POMT1):c.430A>G (p.Asn144Asp)	397514501	POMT1	[]	['GAGCATYCTCTGTTTCAAAGAGG']	['Limb-girdle muscular dystrophy-dystroglycanopathy, type C1']

NM_003863.3(DPM2):c.68A>G (p.Tyr23Cys)	397514503	DPM2	['TGTAGYAGGTGAAGATGATCAGG']	['TGTAGYAGGTGAAGATGATCAGG']	['Congenital disorder of glycosylation type 1u']
NM_174917.4(ACSF3):c.1A>G (p.Met1Val)	370382601	ACSF3	[]	['GGCAGCAYTGCACTGACAGGCGG']	['not provided']
NM_183075.2(CYP2U1):c.1139A>G (p.Glu380Gly)	397514514	CYP2U1	[]	[]	['Spastic paraplegia 56, autosomal recessive']
NM_000344.3(SMN1):c.389A>G (p.Tyr130Cys)	397514517	SMN1	[]	[]	['Kugelberg-Welander disease']
NM_000138.4(FBN1):c.4337-2A>G	794728216	FBN1	[]	[]	['Thoracic aortic aneurysms and aortic dissections']
NM_012082.3(ZFPM2):c.2209A>G (p.Lys737Glu)	397514521	-	[]	[]	['Double outlet right ventricle']
NM_001168272.1(ITPR1):c.1759A>G (p.Asn587Asp)	397514536	ITPR1	[]	[]	['Spinocerebellar ataxia 29']
NM_178012.4(TUBB2B):c.767A>G (p.Asn256Ser)	397514568	TUBB2B	[]	[]	['Polymicrogyria, asymmetric']
NM_000531.5(OTC):c.155A>G (p.Glu52Gly)	72554317	OTC	[]	[]	['not provided']
NM_001866.2(COX7B):c.41-2A>G	397514584	COX7B	[]	[]	['Aplasia cutis congenita, reticulolinear, with microcephaly, facial dysmorphism, and other congenital anomalies']
NM_001099922.2(ALG13):c.339A>G (p.Ala113=)	397514587	ALG13	[]	[]	['Congenital disorder of glycosylation type 1s']
NM_000531.5(OTC):c.238A>G (p.Lys80Glu)	72554332	OTC	[]	['AAGGACTYCCCTTGCAATAAAGG']	['Ornithine carbamoyltransferase deficiency', 'not provided']
NM_001083614.1(EARS2):c.502A>G (p.Arg168Gly)	397514591	EARS2	[]	[]	['Combined oxidative phosphorylation deficiency 12']
NM_001083614.1(EARS2):c.193A>G (p.Lys65Glu)	397514595	EARS2	[]	[]	['Combined oxidative phosphorylation deficiency 12']
NM_198578.3(LRRK2):c.3364A>G (p.Ile1122Val)	34805604	LRRK2	[]	[]	['Parkinson disease 8, autosomal dominant']
NM_033109.4(PNPT1):c.1160A>G (p.Gln387Arg)	397514598	PNPT1	[]	[]	['Combined oxidative phosphorylation deficiency 13']
NM_033109.4(PNPT1):c.1424A>G (p.Glu475Gly)	397514599	PNPT1	[]	['GACTYCAGATGTAACTCTTATGG']	['Deafness, autosomal recessive 70']

NM_000531.5(OTC):c.277A>G (p.Thr93Ala)	72554344	OTC	[]	[]	['not provided']
NM_000390.2(CHM):c.1520A>G (p.His507Arg)	397514603	CHM	[]	[]	['Choroideremia']
NM_181690.2(AKT3):c.686A>G (p.Asn229Ser)	397514605	AKT3	[]	[]	['Megalencephaly-polymicrogyria-polydactyly-hydrocephalus syndrome 2']
NM_006567.3(FARS2):c.431A>G (p.Tyr144Cys)	397514610	FARS2	[]	[]	['Mitochondrial encephalomyopathy', 'Combined oxidative phosphorylation deficiency 14', 'Global developmental delay']
NM_000531.5(OTC):c.377A>G (p.Asp126Gly)	72554358	OTC	[]	[]	['not provided']
NM_005609.2(PYGM):c.152A>G (p.Asp51Gly)	397514631	PYGM	[]	[]	['Glycogen storage disease, type V']
NM_000130.4(F5):c.1601G>A (p.Arg534Gln)	6025	F5	[]	[]	['Recurrent abortion', 'Thrombophilia due to factor V Leiden']
NM_000108.4(DLD):c.1444A>G (p.Arg482Gly)	397514650	DLD	[]	['GACTCYAGCTATATCTTCACAGG']	['Maple syrup urine disease, type 3']
NM_138554.4(TLR4):c.896A>G (p.Asp299Gly)	4986790	TLR4	[]	[]	['Endotoxin hyporesponsiveness']
NM_003156.3(STIM1):c.251A>G (p.Asp84Gly)	397514675	STIM1	[]	['TTCCACAYCCACATCACCATTGG']	['Myopathy with tubular aggregates']
NM_003156.3(STIM1):c.326A>G (p.His109Arg)	397514677	STIM1	[]	[]	['Myopathy with tubular aggregates']
NM_000238.3(KCNH2):c.1900A>G (p.Thr634Ala)	794728377	KCNH2	[]	[]	['Cardiac arrhythmia']
NM_000238.3(KCNH2):c.1913A>G (p.Lys638Arg)	794728378	KCNH2	[]	['ATCYTCTCTGAGTTGGTGTTGGG', 'GATCYTCTCTGAGTTGGTGTTGG']	['Cardiac arrhythmia']
NM_001457.3(FLNB):c.604A>G (p.Met202Val)	121908895	FLNB	[]	[]	['Atelosteogenesis type 1']
NM_001893.4(CSNK1D):c.137A>G (p.His46Arg)	397514693	CSNK1D	[]	[]	['Advanced sleep phase syndrome, familial, 2']
NM_002163.2(IRF8):c.322A>G (p.Lys108Glu)	397514710	IRF8	[]	[]	['Monocyte and dendritic cell deficiency, autosomal recessive']

NM_002163.2(IRF8):c.238A>G (p.Thr80Ala)	397514711	IRF8	[]	['AACCTCGYCTTCCAAGTGGCTGG']	['Autosomal dominant CD11C+/CD1C+ dendritic cell deficiency']
NM_001127217.2(SMAD9):c.127A>G (p.Lys43Glu)	397514715	SMAD9	[]	[]	['Primary pulmonary hypertension 2']
NM_001035.2(RYR2):c.12290A>G (p.Asn4097Ser)	794728784	RYR2	[]	[]	['not provided']
NM_000388.3(CASR):c.85A>G (p.Lys29Glu)	397514729	CASR	[]	['CCCCCTYCTTTTGGGCTCGCTGG']	['Hypocalcemia, autosomal dominant 1, with bartter syndrome']
NM_003793.3(CTSF):c.962A>G (p.Gln321Arg)	397514731	CTSF	[]	[]	['Ceroid lipofuscinosis, neuronal, 13']
NM_173076.2(ABCA12):c.4139A>G (p.Asn1380Ser)	28940269	ABCA12	[]	[]	['Autosomal recessive congenital ichthyosis 4A']
NM_017890.4(VPS13B):c.8978A>G (p.Asn2993Ser)	28940272	VPS13B	['TCAYTGATAAGCAGGGCCCAGGG', 'TTCAYTGATAAGCAGGGCCCAGG']	['TCAYTGATAAGCAGGGCCCAGGG', 'TTCAYTGATAAGCAGGGCCCAGG']	['Cohen syndrome', 'not specified']
NM_022114.3(PRDM16):c.2447A>G (p.Asn816Ser)	397514743	PRDM16	[]	['GCCGCCGYTTTGGCTGGCACGGG']	['Left ventricular noncompaction 8']
NM_005689.2(ABCB6):c.508A>G (p.Ser170Gly)	397514757	ABCB6	[]	['TGGGCYGTTCCAAGACACCAGGG', 'GTGGGCYGTTCCAAGACACCAGG']	['Dyschromatosis universalis hereditaria 3']
NM_015335.4(MED13L):c.752A>G (p.Glu251Gly)	28940309	MED13L	[]	[]	['Transposition of great arteries']
NM_152443.2(RDH12):c.677A>G (p.Tyr226Cys)	28940313	RDH12	[]	['CACTGCGYAGGTGGTGACCCCGG']	['Leber congenital amaurosis 13']
NM_000517.4(HBA2):c.96-2A>G	41457746	HBA2	[]	[]	[]
NM_000218.2(KCNQ1):c.1787A>G (p.Glu596Gly)	794728538	KCNQ1	[]	['GTCTYCTACTCGGTTCAGGCGGG', 'TGTCTYCTACTCGGTTCAGGCGG']	['Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.605A>G (p.Asp202Gly)	794728569	KCNQ1	[]	['AGGYCTGTGGAGTGCAGGAGAGG']	['Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.1515-2A>G	794728573	KCNQ1	[]	['GCCYGCAGTGGAGAGAGGAGAGG']	['Cardiac arrhythmia']
NM_000498.3(CYP11B2):c.1492A>G (p.Thr498Ala)	72554626	-	[]	[]	['Corticosterone methyloxidase type 2 deficiency']
NM_000169.2(GLA):c.644A>G (p.Asn215Ser)	28935197	-	[]	[]	['Fabry disease', 'not provided']

NM_000218.2(KCNQ1):c.1085A>G (p.Lys362Arg)	12720458	KCNQ1	[]	[]	['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_001035.2(RYR2):c.12533A>G (p.Asn4178Ser)	794728787	RYR2	[]	[]	['not provided']
NM_003494.3(DYSF):c.3349-2A>G	370874727	DYSF	[]	['CCGCCCYGGAGACACGAAGCTGG']	['Limb-girdle muscular dystrophy, type 2B']
NM_001035.2(RYR2):c.568A>G (p.Arg190Gly)	794728814	RYR2	[]	[]	['not provided']
NM_198056.2(SCN5A):c.2788-2A>G	794728859	SCN5A	[]	['ACCYGTCGAGATAATGGGTCAGG']	['not provided']
NM_198056.2(SCN5A):c.4453A>G (p.Ile1485Val)	794728886	SCN5A	[]	[]	['not provided']
NM_198056.2(SCN5A):c.4462A>G (p.Thr1488Ala)	794728887	SCN5A	[]	['CCTCTGYCATGAAGATGTCCTGG']	['not provided']
NM_001927.3(DES):c.1324A>G (p.Thr442Ala)	794728995	DES	[]	[]	['not provided']
NM_001613.2(ACTA2):c.145A>G (p.Met49Val)	397515325	ACTA2	[]	[]	['Aortic aneurysm, familial thoracic 6']
NM_000782.4(CYP24A1):c.1226T>C (p.Leu409Ser)	6068812	CYP24A1	[]	[]	['Idiopathic hypercalcemia of infancy']
NM_000372.4(TYR):c.125A>G (p.Asp42Gly)	28940878	TYR	[]	['CTCCTGYCCCCGCTCCACGGTGG']	['Tyrosinase-negative oculocutaneous albinism', 'not provided']
NM_000372.4(TYR):c.1A>G (p.Met1Val)	28940881	TYR	[]	[]	['Tyrosinase-negative oculocutaneous albinism', 'Oculocutaneous albinism type 1B', 'not provided']
NM_000403.3(GALE):c.770A>G (p.Lys257Arg)	28940884	GALE	[]	[]	['UDPglucose-4-epimerase deficiency']
NM_000529.2(MC2R):c.761A>G (p.Tyr254Cys)	28940892	MC2R	['ACATGYAGCAGGCGCAGTAGGGG', 'GACATGYAGCAGGCGCAGTAGGG']	['ACATGYAGCAGGCGCAGTAGGGG', 'GACATGYAGCAGGCGCAGTAGGG', 'AGACATGYAGCAGGCGCAGTAGG']	['ACTH resistance']
NM_000061.2(BTK):c.919A>G (p.Arg307Gly)	128621195	BTK	[]	[]	['X-linked agammaglobulinemia']
NM_000061.2(BTK):c.1766A>G	128621206	BTK	[]	[]	['X-linked agammaglobulinemi

(p.Glu589Gly)					a']
NM_018486.2(HDAC8):c.539A>G (p.His180Arg)	397515416	HDAC8	[]	[]	['Cornelia de Lange syndrome 5']
NM_018486.2(HDAC8):c.1001A>G (p.His334Arg)	397515418	HDAC8	['CTCAYGATCTGGGATCTCAGAGG']	['CTCAYGATCTGGGATCTCAGAGG']	['Cornelia de Lange syndrome 5']
NM_172107.2(KCNQ2):c.1636A>G (p.Met546Val)	397515420	KCNQ2	[]	['GCAYGACACTGCAGGGGGGTGGG', 'CGCAYGACACTGCAGGGGGGTGG', 'AACCGCAYGACACTGCAGGGGGG']	['Early infantile epileptic encephalopathy 7']
NM_001410.2(MEGF8):c.7099A>G (p.Ser2367Gly)	397515428	MEGF8	[]	['GACYCCCGTGAAATGATTCCCGG']	['Carpenter syndrome 2']
NM_004247.3(EFTUD2):c.623A>G (p.His208Arg)	397515431	EFTUD2	[]	[]	['Growth and mental retardation, mandibulofacial dysostosis, microcephaly, and cleft palate']
NM_004572.3(PKP2):c.1171-2A>G	794729133	PKP2	[]	[]	['not provided']
NM_018972.2(GDAP1):c.368A>G (p.His123Arg)	397515442	GDAP1	[]	[]	['Charcot-Marie-Tooth disease type 2K']
NM_014795.3(ZEB2):c.3134A>G (p.His1045Arg)	397515449	ZEB2	[]	[]	['Mowat-Wilson syndrome']
NM_002336.2(LRP6):c.1298A>G (p.Asn433Ser)	397515473	LRP6	[]	[]	['Coronary artery disease, autosomal dominant 2']
NM_001015879.1(AURKC):c.379-2A>G	397515484	AURKC	[]	[]	['Infertility associated with multi-tailed spermatozoa and excessive DNA']
NM_000495.4(COL4A5):c.3107-4A>G	397515497	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_201631.3(TGM5):c.122T>C (p.Leu41Pro)	143601447	TGM5	[]	['TCAACCYCACCCTGTACTTCAGG']	['Peeling skin syndrome, acral type']
NM_013254.3(TBK1):c.1201A>G (p.Lys401Glu)	756751089	TBK1	[]	[]	['FRONTOTEMPORAL DEMENTIA AND/OR AMYOTROPHIC LATERAL SCLEROSIS 4']
NM_000207.2(INS):c.*59A>G	397515519	-	[]	['GGGCYTTATTCCATCTCTCTCGG']	['Permanent neonatal diabetes mellitus']
NM_000370.3(TTPA):c.191A>G (p.Asp64Gly)	397515523	TTPA	[]	['CAGGYCCAGATCGAAATCCCGGG', 'CCAGGYCCAGATCGAAATCCCGG']	['Ataxia with vitamin E deficiency']
NM_001006657.1(WDR35):c.2912A>G	397515535	WDR35	[]	[]	['Cranioectodermal dysplasia 2']

(p.Tyr971Cys)					
NM_000424.3(KRT5):c.1424A>G (p.Glu475Gly)	61348633	KRT5	[]	[]	['Epidermolysis bullosa herpetiformis, Dowling-Meara', 'not provided']
NM_004595.4(SMS):c.443A>G (p.Gln148Arg)	397515551	SMS	[]	[]	['Snyder Robinson syndrome']
NM_001256850.1(TTN):c.45629-2A>G	794729266	-	[]	[]	['not provided']
NM_000404.2(GLB1):c.947A>G (p.Tyr316Cys)	72555361	GLB1	[]	[]	['Infantile GM1 gangliosidosis']
NM_000404.2(GLB1):c.1498A>G (p.Thr500Ala)	72555368	GLB1	[]	[]	['Mucopolysaccharidosis, MPS-IV-B']
NM_000404.2(GLB1):c.1772A>G (p.Tyr591Cys)	72555371	GLB1	[]	[]	['GM1-GANGLIOSIDOSIS, TYPE I, WITH CARDIAC INVOLVEMENT']
NM_000487.5(ARSA):c.1055A>G (p.Asn352Ser)	2071421	ARSA	[]	[]	['Metachromatic leukodystrophy', 'not provided']
NM_001037811.2(HSD17B10):c.713A>G (p.Asn238Ser)	122461163	HSD17B10	[]	[]	['2-methyl-3-hydroxybutyric aciduria']
NM_000138.4(FBN1):c.1148-2A>G	397515756	FBN1	[]	[]	['Marfan syndrome']
NM_001875.4(CPS1):c.1010A>G (p.His337Arg)	28940283	CPS1	[]	[]	['Congenital hyperammonemia, type I']
NM_000169.2(GLA):c.1153A>G (p.Thr385Ala)	397515869	-	['AGCTGTGYGATGAAGCAGGCAGG']	['AGCTGTGYGATGAAGCAGGCAGG']	['not specified', 'not provided']
NM_000256.3(MYBPC3):c.1224-2A>G	397515891	MYBPC3	[]	['TACTTGCYGTAGAACAGAAGGGG']	['Familial hypertrophic cardiomyopathy 4', 'Cardiomyopathy']
NM_000048.3(ASL):c.857A>G (p.Gln286Arg)	28941472	ASL	[]	[]	['Argininosuccinate lyase deficiency', 'not provided']
NM_000256.3(MYBPC3):c.1928-2A>G	397515937	MYBPC3	[]	[]	['Primary familial hypertrophic cardiomyopathy', 'Familial hypertrophic cardiomyopathy 4', 'Cardiomyopathy']
NM_002693.2(POLG):c.1283T>C (p.Leu428Pro)	774610098	POLG	[]	[]	['not provided']
NM_004628.4(XPC):c.413-24A>G	794729657	XPC	[]	[]	['Xeroderma pigmentosum, group C']
NM_030973.3(MED25):c.116A>G	794729668	MED25	[]	[]	['BASEL-VANAGAITE-

(p.Tyr39Cys)					SMIRIN-YOSEF SYNDROME']
NM_001955.4(EDN1): c.271A>G (p.Lys91Glu)	587777231	EDN1	[]	[]	['Auriculocondylar syndrome 3']
NM_003002.3(SDHD) :c.149A>G (p.His50Arg)	11214077	SDHD	[]	[]	['Pheochromocytoma', 'Merkel cell carcinoma', 'Hereditary cancer-predisposing syndrome', 'Carcinoid tumor of intestine', 'Cowden syndrome 3', 'not specified', 'not provided']
NM_001003722.1(GLE1):c.433-10A>G	386833693	GLE1	[]	[]	['Lethal arthrogryposis with anterior horn cell disease']
NM_000371.3(TTR):c.185A>G (p.Glu62Gly)	11541796	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis']
NM_000256.3(MYBPC3):c.927-2A>G	397516082	MYBPC3	[]	['GTCCCYGTGTCCCGCAGTCTAGG']	['Familial hypertrophic cardiomyopathy 4', 'Cardiomyopathy']
NM_001148.4(ANK2) :c.4373A>G (p.Glu1458Gly)	72544141	ANK2	[]	[]	['Long QT syndrome', 'Congenital long QT syndrome', 'Long QT syndrome 4', 'Cardiac arrhythmia, ankyrin B-related', 'Cardiac arrhythmia']
NM_000257.3(MYH7) :c.2206A>G (p.Ile736Val)	397516138	MYH7	[]	['TATCAAYGAACTGTCCCTCAGGG', 'CTATCAAYGAACTGTCCCTCAGG']	['Familial hypertrophic cardiomyopathy 1', 'Cardiomyopathy', 'not specified']
NM_000356.3(TCOF1):c.149A>G (p.Tyr50Cys)	28941769	TCOF1	['GTGTGTAYAGATGTCCAGAAGGG']	['GTGTGTAYAGATGTCCAGAAGGG']	['Treacher collins syndrome 1']
NM_002150.2(HPD):c.97G>A (p.Ala33Thr)	1154510	HPD	[]	['ATGACGYGGCCTGAATCACAGGG', 'AATGACGYGGCCTGAATCACAGG']	['4-Alpha-hydroxyphenylpyruvate hydroxylase deficiency']
NM_000375.2(UROS) :c.184A>G (p.Thr62Ala)	28941775	UROS	[]	[]	['Congenital erythropoietic porphyria']
NM_001008216.1(GALE):c.308A>G (p.Asp103Gly)	28940883	GALE	[]	[]	['UDPglucose-4-epimerase deficiency']
NM_000260.3(MYO7A):c.6439-2A>G	397516330	MYO7A	[]	['ATATCCYGGGGGAGCAGAAAGGG', 'GATATCCYGGGGGAGCAGAAAGG']	['Usher syndrome, type 1']

NM_033071.3(SYNE1):c.15705-12A>G	606231134	SYNE1	[]	[]	['Spinocerebellar ataxia, autosomal recessive 8']
NM_000187.3(HGD):c.1112A>G (p.His371Arg)	120074172	HGD	[]	[]	['Alkaptonuria']
NM_000053.3(ATP7B):c.3443T>C (p.Ile1148Thr)	60431989	ATP7B	['TGCTGAYTGGAAACCGTGAGTGG']	['TGCTGAYTGGAAACCGTGAGTGG']	['Wilson disease']
NM_000441.1(SLC26A4):c.-3-2A>G	397516411	-	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_003041.3(SLC5A2):c.1961A>G (p.Asn654Ser)	61742739	-	[]	[]	['Familial renal glucosuria']
NM_000551.3(VHL):c.467A>G (p.Tyr156Cys)	397516441	VHL	[]	[]	['Von Hippel-Lindau syndrome']
NM_000531.5(OTC):c.481A>G (p.Asn161Asp)	72556270	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.482A>G (p.Asn161Ser)	72556271	OTC	[]	['CAGCCCAYTGATAATTGGGATGG']	['not provided']
NM_000531.5(OTC):c.541-2A>G	72556289	OTC	['TCCYAAAAGGCACGGGATGAAGG']	['TCCYAAAAGGCACGGGATGAAGG']	['not provided']
NM_000531.5(OTC):c.542A>G (p.Glu181Gly)	72556290	OTC	['ATAGTGTYCCTAAAAGGCACGGG']	['ATAGTGTYCCTAAAAGGCACGGG']	['not provided']
NM_000527.4(LDLR):c.2483A>G (p.Tyr828Cys)	28942085	LDLR	[]	[]	['Familial hypercholesterolemia', 'not provided']
NM_000271.4(NPC1):c.3467A>G (p.Asn1156Ser)	28942105	NPC1	[]	[]	['Niemann-Pick disease type C1']
NM_000271.4(NPC1):c.3263A>G (p.Tyr1088Cys)	28942106	NPC1	[]	[]	['NIEMANN-PICK DISEASE, TYPE C1, JUVENILE FORM']
NM_020247.4(ADCK3):c.1541A>G (p.Tyr514Cys)	119468008	ADCK3	[]	[]	['Coenzyme Q10 deficiency, primary, 4']
NM_001063.3(TF):c.956A>G (p.His319Arg)	41295774	TF	[]	[]	[]
NM_172201.1(KCNE2):c.281A>G (p.Glu94Gly)	74424227	KCNE2	[]	[]	['Congenital long QT syndrome']
NM_002294.2(LAMP2):c.65-2A>G	397516743	LAMP2	[]	[]	['Danon disease']
NM_002880.3(RAF1):c.524A>G (p.His175Arg)	397516822	RAF1	[]	[]	['Noonan syndrome 5']
NM_033360.3(KRAS):c.13A>G (p.Lys5Glu)	193929331	KRAS	[]	[]	['Noonan syndrome 3', 'Rasopathy']
NM_000525.3(KCNJ1	193929337	KCNJ11	[]	[]	['Permanent neonatal

1):c.155A>G (p.Gln52Arg)					diabetes mellitus']
NM_000525.3(KCNJ11):c.544A>G (p.Ile182Val)	193929348	KCNJ11	['AGAYGAGGGTCTCAGCCCTGCGG']	['AGAYGAGGGTCTCAGCCCTGCGG']	['Permanent neonatal diabetes mellitus']
NM_000525.3(KCNJ11):c.989A>G (p.Tyr330Cys)	193929356	KCNJ11	[]	[]	['Permanent neonatal diabetes mellitus', 'Neonatal insulin-dependent diabetes mellitus']
NM_001288953.1(TTC7A):c.1715A>G (p.Lys572Arg)	139010200	TTC7A	[]	[]	['Multiple gastrointestinal atresias']
NM_024809.4(TCTN2):c.1506-2A>G	374349989	TCTN2	[]	[]	['Meckel syndrome type 8']
NM_012275.2(IL36RN):c.104A>G (p.Lys35Arg)	187015338	IL36RN	[]	[]	['Pustular psoriasis, generalized']
NM_178517.3(PIGW):c.499A>G (p.Met167Val)	200024253	PIGW	[]	[]	['Hyperphosphatasia with mental retardation syndrome 5']
NM_015662.2(IFT172):c.5179T>C (p.Cys1727Arg)	149614625	-	[]	[]	['Short-rib thoracic dysplasia 10 with or without polydactyly']
NM_000226.3(KRT9):c.469A>G (p.Met157Val)	58597584	KRT9	[]	[]	['Epidermolytic palmoplantar keratoderma', 'not provided']
NM_023073.3(C5orf42):c.3290-2A>G	606231260	C5orf42	[]	['ATCYATCAAATACAAAAATTTGG']	['Orofaciodigital syndrome 6']
NM_005633.3(SOS1):c.508A>G (p.Lys170Glu)	397517172	SOS1	[]	[]	['Noonan syndrome 4', 'Rasopathy', 'not provided']
NM_006306.3(SMC1A):c.3254A>G (p.Tyr1085Cys)	587784418	SMC1A	['CTTAYAGATCTCATCAATGTTGG']	['CTTAYAGATCTCATCAATGTTGG']	['Congenital muscular hypertrophy-cerebral syndrome']
NM_006218.2(PIK3CA):c.1637A>G (p.Gln546Arg)	397517201	PIK3CA	[]	[]	['Neoplasm of ovary']
NM_006218.2(PIK3CA):c.3073A>G (p.Thr1025Ala)	397517202	PIK3CA	[]	[]	['Non-small cell lung cancer']
NM_002354.2(EPCAM):c.492-2A>G	606231281	EPCAM	[]	[]	['Diarrhea 5, with tufting enteropathy, congenital']
NM_033056.3(PCDH15):c.1998-2A>G	397517452	PCDH15	[]	[]	['Usher syndrome, type 1F']
NM_000301.3(PLG):c.112A>G (p.Lys38Glu)	73015965	PLG	[]	[]	['Plasminogen deficiency, type I']
NM_000155.3(GALT):c.424A>G (p.Met142Val)	111033692	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000096.3(CP):c.2953A>G	386134132	CP	[]	[]	['Deficiency of ferroxidase']

(p.Met985Val)					
NM_000197.1(HSD17B3):c.703A>G (p.Met235Val)	119481074	HSD17B3	[]	[]	['Testosterone 17-beta-dehydrogenase deficiency']
NM_000197.1(HSD17B3):c.389A>G (p.Asn130Ser)	119481079	HSD17B3	[]	[]	['Testosterone 17-beta-dehydrogenase deficiency']
NM_015474.3(SAMHD1):c.1106T>C (p.Leu369Ser)	515726139	SAMHD1	[]	[]	['Aicardi Goutieres syndrome 5']
NM_004817.3(TJP2):c.1992-2A>G	587777521	TJP2	[]	['CAGCTCYGAGAAGAAACCACGGG', 'TCAGCTCYGAGAAGAAACCACGG']	['Progressive familial intrahepatic cholestasis 4']
NM_000257.3(MYH7):c.617A>G (p.Lys206Arg)	730880846	MYH7	[]	['CTTCYTGCTGCGGTCCCCAATGG']	['Cardiomyopathy']
NM_020919.3(ALS2):c.2980-2A>G	386134184	ALS2	[]	[]	['Juvenile primary lateral sclerosis']
m.10044A>G	41362547	MT-TG	[]	[]	['Sudden death']
NM_002977.3(SCN9A):c.406A>G (p.Ile136Val)	80356468	SCN9A	[]	[]	['Primary erythromelalgia']
NM_001128425.1(MUTYH):c.536A>G (p.Tyr179Cys)	34612342	MUTYH	[]	[]	['MYH-associated polyposis', 'Hereditary cancer-predisposing syndrome', 'Endometrial carcinoma', 'Carcinoma of colon', 'not specified', 'not provided']
NM_206933.2(USH2A):c.12067-2A>G	397517978	USH2A	[]	['TTCCCYGTAAGAAAATTAACAGG']	['Usher syndrome, type 2A', 'Retinitis pigmentosa 39']
NM_000216.2(ANOS1):c.1A>G (p.Met1Val)	606231409	ANOS1	[]	['GCACCAYGGCTGCGGGTCGAGGG', 'GGCACCAYGGCTGCGGGTCGAGG']	['Kallmann syndrome 1']
NM_206933.2(USH2A):c.1841-2A>G	397518003	USH2A	[]	[]	['Usher syndrome, type 2A']
NM_000368.4(TSC1):c.1760A>G (p.Lys587Arg)	118203576	TSC1	[]	[]	['Tuberous sclerosis syndrome', 'Tuberous sclerosis 1', 'Hereditary cancer-predisposing syndrome', 'not specified']
NM_021830.4(C10orf2):c.1523A>G (p.Tyr508Cys)	80356540	C10orf2	[]	[]	['Mitochondrial DNA depletion syndrome 7 (hepatocerebral type)', 'not provided']
NM_003334.3(UBA1):	80356546	UBA1	[]	['TGGCYTGTCACC	['Arthrogryposis

c.1639A>G (p.Ser547Gly)				CGGATATGTGG']	multiplex congenita, distal, X-linked']
NM_206933.2(USH2A):c.8559-2A>G	397518039	USH2A	['ATCYAAAGCAAAAGACAAGCAGG']	['ATCYAAAGCAAAAGACAAGCAGG']	['Retinitis pigmentosa', 'Usher syndrome, type 2A']
NM_000038.5(APC):c.1744-2A>G	587783035	APC	['TCCYAGTAAGAAACAGAATATGG']	['TCCYAGTAAGAAACAGAATATGG']	['Familial adenomatous polyposis 1']
NM_194248.2(OTOF):c.766-2A>G	80356584	OTOF	[]	['GACCYGCAGGCAGGAGAAGGGGG', 'TGACCYGCAGGCAGGAGAAGGGG', 'CTGACCYGCAGGCAGGAGAAGGG', 'GCTGACCYGCAGGCAGGAGAAGG']	['Deafness, autosomal recessive 9']
NM_000525.3(KCNJ11):c.509A>G (p.Lys170Arg)	80356621	KCNJ11	[]	[]	['Permanent neonatal diabetes mellitus']
NM_000352.4(ABCC8):c.215A>G (p.Asn72Ser)	80356634	ABCC8	[]	[]	['Permanent neonatal diabetes mellitus']
NM_000352.4(ABCC8):c.4270A>G (p.Ile1424Val)	80356653	ABCC8	[]	[]	['Permanent neonatal diabetes mellitus']
NM_000207.2(INS):c.323A>G (p.Tyr108Cys)	80356672	-	[]	[]	['Permanent neonatal diabetes mellitus']
NM_000083.2(CLCN1):c.382A>G (p.Met128Val)	80356699	CLCN1	[]	[]	['Myotonia congenita', 'Congenital myotonia, autosomal dominant form']
NM_001008211.1(OPTN):c.1433A>G (p.Glu478Gly)	267606929	OPTN	[]	[]	['Amyotrophic lateral sclerosis type 12']
NM_007375.3(TARDBP):c.506A>G (p.Asp169Gly)	80356717	TARDBP	[]	[]	['Amyotrophic lateral sclerosis type 10']
NM_007375.3(TARDBP):c.1009A>G (p.Met337Val)	80356730	TARDBP	[]	[]	['Amyotrophic lateral sclerosis type 10']
NM_007375.3(TARDBP):c.1028A>G (p.Gln343Arg)	80356731	TARDBP	[]	[]	['Amyotrophic lateral sclerosis type 10']
NM_001701.3(BAAT):c.967A>G (p.Ile323Val)	80356747	BAAT	['CAAYGAAGAGGAATTGCCCCTGG']	['CAAYGAAGAGGAATTGCCCCTGG']	['Atypical hemolytic-uremic syndrome 1']
NM_012463.3(ATP6V0A2):c.732-2A>G	80356753	ATP6V0A2	[]	[]	['Cutis laxa with osteodystrophy']
NM_001876.3(CPT1A):c.1361A>G (p.Asp454Gly)	80356778	CPT1A	[]	[]	['Carnitine palmitoyltransferase I deficiency']
NM_001876.3(CPT1A):c.1079A>G	80356787	CPT1A	[]	[]	['Carnitine palmitoyltransferase

(p.Glu360Gly)					I deficiency']
NM_001876.3(CPT1A):c.1493A>G (p.Tyr498Cys)	80356791	CPT1A	[]	[]	['Carnitine palmitoyltransferase I deficiency']
NM_003159.2(CDKL5):c.211A>G (p.Asn71Asp)	587783072	CDKL5	[]	[]	['Atypical Rett syndrome', 'not provided']
NM_000257.3(MYH7):c.1615A>G (p.Met539Val)	730880930	MYH7	[]	['GGAACAYGCACTCCTCTTCCAGG']	['Cardiomyopathy']
m.5843A>G	118203894	MT-TY	[]	[]	[]
NM_000130.4(F5):c.1000A>G (p.Arg334Gly)	118203905	F5	[]	[]	[]
NM_000130.4(F5):c.5189A>G (p.Tyr1730Cys)	118203907	F5	['GTAGYAGGCCCAAGCCCGACAGG']	['GTAGYAGGCCCAAGCCCGACAGG']	['Factor V deficiency']
NM_000052.6(ATP7A):c.3911A>G (p.Asn1304Ser)	151340632	ATP7A	[]	[]	['Menkes kinky-hair syndrome', 'Cutis laxa, X-linked']
NM_007294.3(BRCA1):c.5053A>G (p.Thr1685Ala)	80356890	BRCA1	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
NM_000046.3(ARSB):c.629A>G (p.Tyr210Cys)	118203943	ARSB	[]	[]	['Mucopolysaccharidosis type VI', 'not provided']
NM_013319.2(UBIAD1):c.305A>G (p.Asn102Ser)	118203945	UBIAD1	['GTAAGTGYTGACCAAATTACCGG']	['GTAAGTGYTGACCAAATTACCGG']	['Schnyder crystalline corneal dystrophy']
NM_013319.2(UBIAD1):c.355A>G (p.Arg119Gly)	118203947	UBIAD1	[]	['TCCYGTCATCACTCTTTTTGTGG']	['Schnyder crystalline corneal dystrophy']
NM_013319.2(UBIAD1):c.695A>G (p.Asn232Ser)	118203949	UBIAD1	['GGTGTTGYTGGAATGGAGAATGG']	['GGTGTTGYTGGAATGGAGAATGG']	['Schnyder crystalline corneal dystrophy']
NM_013319.2(UBIAD1):c.335A>G (p.Asp112Gly)	118203950	UBIAD1	[]	[]	['Schnyder crystalline corneal dystrophy']
NM_024334.2(TMEM43):c.271A>G (p.Ile91Val)	144811578	TMEM43	[]	[]	['Emery-Dreifuss muscular dystrophy 7, autosomal dominant', 'not provided']
NM_012073.4(CCT5):c.440A>G (p.His147Arg)	118203986	CCT5	[]	[]	['Neuropathy, hereditary sensory, with spastic paraplegia, autosomal recessive']
NM_000033.3(ABCD1):c.443A>G (p.Asn148Ser)	128624216	ABCD1	['CACTGYTGACGAAGGTAGCAGG	['CACTGYTGACGAAGGTAGCAGGG', 'GCACTGYTGACG	['Adrenoleukodystrophy']

			G']	AAGGTAGCAGG']	
NM_000146.3(FTL):c.1A>G (p.Met1Val)	139732572	FTL	['CTCAYGGTTGGTTGGCAAGAAGG']	['CTCAYGGTTGGTTGGCAAGAAGG']	['L-ferritin deficiency']
NM_000785.3(CYP27B1):c.566A>G (p.Glu189Gly)	118204012	CYP27B1	[]	[]	['Vitamin D-dependent rickets, type 1']
NM_000252.2(MTM1):c.1261-10A>G	397518445	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_139281.2(WDR36):c.1064A>G (p.Asn355Ser)	118204022	WDR36	[]	[]	['Glaucoma 1, open angle, G']
NM_000392.4(ABCC2):c.4145A>G (p.Gln1382Arg)	72558202	ABCC2	[]	[]	['Dubin-Johnson syndrome']
NM_000165.4(GJA1):c.617A>G (p.Lys206Arg)	397518464	GJA1	[]	[]	['Oculodentodigital dysplasia']
NM_000833.4(GRIN2A):c.1123-2A>G	397518469	GRIN2A	[]	[]	['Focal epilepsy with speech disorder with or without mental retardation']
NM_015702.2(MMADHC):c.746A>G (p.Tyr249Cys)	118204046	MMADHC	[]	[]	['Homocystinuria, cblD type, variant 1']
NM_000526.4(KRT14):c.368A>G (p.Asn123Ser)	60171927	KRT14	[]	['GCGGTCAYTGAGGTTCTGCATGG']	['Epidermolysis bullosa herpetiformis, Dowling-Meara', 'not provided']
NM_000237.2(LPL):c.548A>G (p.Asp183Gly)	118204064	LPL	['AGCTGGAYCGAGGCCTTAAAAGG']	['GCTGGAYCGAGGCCTTAAAAGGG', 'AGCTGGAYCGAGGCCTTAAAAGG']	['Hyperlipoproteinemia, type I']
NM_016247.3(IMPG2):c.370T>C (p.Phe124Leu)	201893545	IMPG2	['ACTYTTTGGGATCGACTTCCTGG']	['ACTYTTTGGGATCGACTTCCTGG']	['Macular dystrophy, vitelliform, 5']
NM_004035.6(ACOX1):c.832A>G (p.Met278Val)	118204090	ACOX1	[]	[]	['Pseudoneonatal adrenoleukodystrophy']
NM_004035.6(ACOX1):c.926A>G (p.Gln309Arg)	118204092	ACOX1	[]	[]	['Pseudoneonatal adrenoleukodystrophy']
NM_000190.3(HMBS):c.1A>G (p.Met1Val)	118204118	HMBS	[]	[]	['Porphyria, acute intermittent, nonerythroid variant']
NM_001363.4(DKC1):c.941A>G (p.Lys314Arg)	199422248	DKC1	[]	['AATCYTGGCCCCATAGCAGATGG']	['Dyskeratosis congenita X-linked']
NM_000078.2(CETP):c.1376A>G (p.Asp459Gly)	2303790	CETP	[]	[]	['Hyperalphalipoproteinemia']
NM_000531.5(OTC):c.595A>G (p.Asn199Asp)	72558405	OTC	[]	[]	['not provided']

NM_000531.5(OTC):c.596A>G (p.Asn199Ser)	72558406	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.613A>G (p.Met205Val)	72558411	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.717+3A>G	72558432	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.718-2A>G	72558433	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.788A>G (p.Asp263Gly)	72558443	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.790A>G (p.Thr264Ala)	72558444	OTC	[]	[]	['not provided']
NM_000531.5(OTC):c.929A>G (p.Glu310Gly)	72558467	OTC	[]	['TCCACTYCTTCTGGCTTTCTGGG', 'ATCCACTYCTTCTGGCTTTCTGG']	['not provided']
NM_000531.5(OTC):c.988A>G (p.Arg330Gly)	72558478	OTC	[]	['ACTTTCYGTTTTCTGCCTCTGGG', 'CACTTTCYGTTTTCTGCCTCTGG']	['not provided']
NM_000488.3(SERPINC1):c.655A>G (p.Asn219Asp)	121909571	SERPINC1	[]	[]	['Antithrombin III deficiency']
NM_007294.3(BRCA1):c.122A>G (p.His41Arg)	80357276	BRCA1	['AAATATGYGGTCACACTTTGTGG']	['AAATATGYGGTCACACTTTGTGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1']
NM_007294.3(BRCA1):c.1A>G (p.Met1Val)	80357287	BRCA1	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
NM_198056.2(SCN5A):c.1134T>A (p.Tyr378Ter)	373172185	SCN5A	[]	[]	['not provided']
m.7526A>G	121434454	MT-TD	[]	[]	[]
NM_007294.3(BRCA1):c.211A>G (p.Arg71Gly)	80357382	BRCA1	[]	[]	['Familial cancer of breast', 'Hereditary breast and ovarian cancer syndrome', 'Breast-ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
NM_000512.4(GALNS):c.1460A>G (p.Asn487Ser)	118204440	GALNS	['ACGYTGAGCTGGGGCTGCGCGGG', 'CACGYTGAGCTGGGGCTGCGCGG']	['ACGYTGAGCTGGGGCTGCGCGGG', 'CACGYTGAGCTGGGGCTGCGCGG']	['Mucopolysaccharidosis, MPS-IV-A']

NM_000505.3(F12):c.158A>G (p.Tyr53Cys)	118204455	F12	[]	['GGTGGYACTGGAAGGGGAAGTGG']	[]
NM_007294.3(BRCA1):c.5453A>G (p.Asp1818Gly)	80357477	BRCA1	[]	['TTGYCCTCTGTCCAGGCATCTGG']	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1']
NM_032492.3(JAGN1):c.485A>G (p.Gln162Arg)	587777730	JAGN1	[]	[]	['Severe congenital neutropenia', 'Severe congenital neutropenia 6, autosomal recessive']
NM_000257.3(MYH7):c.2087A>G (p.Asn696Ser)	730880732	MYH7	[]	[]	['Cardiomyopathy']
NM_000430.3(PAFAH1B1):c.446A>G (p.His149Arg)	121434482	PAFAH1B1	[]	[]	['Lissencephaly 1']
NM_000363.4(TNNI3):c.547A>G (p.Lys183Glu)	730881077	TNNI3	[]	[]	['Cardiomyopathy']
NM_018105.2(THAP1):c.266A>G (p.Lys89Arg)	267607111	THAP1	[]	[]	['Dystonia 6, torsion']
NM_016599.4(MYOZ2):c.738A>G (p.Ile246Met)	140126678	MYOZ2	[]	[]	['Familial hypertrophic cardiomyopathy 16', 'not specified', 'not provided']
NM_000161.2(GCH1):c.671A>G (p.Lys224Arg)	41298442	GCH1	[]	[]	['Dystonia 5, Dopa-responsive type', 'Dystonia, dopa-responsive, with or without hyperphenylalaninemia, autosomal recessive']
NM_017415.2(KLHL3):c.926A>G (p.Gln309Arg)	199469627	KLHL3			['Pseudohypoaldosteronism, type 2']
NM_017415.2(KLHL3):c.1670A>G (p.Tyr557Cys)	199469645	KLHL3			['Pseudohypoaldosteronism, type 2', 'Pseudohypoaldosteronism type 2D']
NM_003590.4(CUL3):c.1207-26A>G	199469650	CUL3			['Pseudohypoaldosteronism, type 2']
NM_003590.4(CUL3):c.1238A>G (p.Asp413Gly)	199469656	CUL3			['Pseudohypoaldosteronism, type 2', 'Pseudohypoaldosteronism type 2E']
NM_003590.4(CUL3):c.1376A>G (p.Lys459Arg)	199469658	CUL3			['Pseudohypoaldosteronism, type 2']
NM_003590.4(CUL3):c.1377+3A>G	199469661	CUL3			['Pseudohypoaldosteronism, type 2']
NM_007294.3(BRCA1):c.4096+3A>G	80358015	BRCA1	[]	[]	['Hereditary breast and ovarian cancer syndrome', 'Breast-'

					ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
NM_007294.3(BRCA1):c.135-2A>G	80358065	BRCA1	[]	[]	['Breast-ovarian cancer, familial 1']
NM_024426.4(WT1):c.1391A>G (p.Asp464Gly)	121907902	WT1	[]	[]	['Drash syndrome']
NM_007294.3(BRCA1):c.212+3A>G	80358083	BRCA1	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 1']
NM_024426.4(WT1):c.1021A>G (p.Ser341Gly)	121907908	WT1	[]	['CGCYCTCGTACCCTGTGCTGTGG']	['Mesothelioma']
NM_007294.3(BRCA1):c.4676-2A>G	80358096	BRCA1	[]	[]	['Breast-ovarian cancer, familial 1', 'Hereditary cancer-predisposing syndrome']
NM_000280.4(PAX6):c.1171A>G (p.Thr391Ala)	121907926	PAX6	[]	['GTGGYGCCCGAGGTGCCCATTGG']	['Optic nerve aplasia, bilateral']
NM_000520.4(HEXA):c.611A>G (p.His204Arg)	121907976	HEXA	[]	[]	['Tay-Sachs disease']
NM_000159.3(GCDH):c.1213A>G (p.Met405Val)	141437721	GCDH	[]	[]	['Glutaric aciduria, type 1']
NM_024740.2(ALG9):c.860A>G (p.Tyr287Cys)	121908023	ALG9	[]	['TTAYACAAAACAATGTTGAGTGG']	['Congenital disorder of glycosylation type 1L']
NM_003051.3(SLC16A1):c.610A>G (p.Lys204Glu)	80358222	SLC16A1	[]	[]	['Erythrocyte lactate transporter defect']
NM_000229.1(LCAT):c.463A>G (p.Asn155Asp)	121908057	LCAT	[]	[]	['Fish-eye disease']
NM_000639.2(FASLG):c.466A>G (p.Arg156Gly)	80358238	FASLG	[]	[]	['Autoimmune lymphoproliferative syndrome']
NM_001369.2(DNAH5):c.1121T>C (p.Ile374Thr)	147499872	DNAH5	[]	[]	['Ciliary dyskinesia, primary, 3']
NM_138691.2(TMC1):c.1960A>G (p.Met654Val)	121908074	TMC1	[]	[]	['Deafness, autosomal recessive 7']
NM_024301.4(FKRP):c.1387A>G (p.Asn463Asp)	121908110	FKRP	[]	[]	['Congenital muscular dystrophy-dystroglycanopathy (with or without mental retardation) type B5', 'Limb-girdle muscular dystrophy-dystroglycanopathy, type C5', 'Muscular

					dystrophy', 'Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies type A5', 'Congenital muscular dystrophy-dystroglycanopathy without mental retardation, type B5', 'not provided']
NM_175073.2(APTX):c.602A>G (p.His201Arg)	121908133	APTX	['GCCAAYGGTAACGGGCCTTTGGG']	['GCCAAYGGTAACGGGCCTTTGGG', 'AGCCAAYGGTAACGGGCCTTTGG']	['Adult onset ataxia with oculomotor apraxia']
NM_001243133.1(NLRP3):c.1880A>G (p.Glu627Gly)	121908148	NLRP3	[]	['ACAATYCCAGCTGGCTGGGCTGG']	['Familial cold urticaria']
NM_006492.2(ALX3):c.608A>G (p.Asn203Ser)	121908166	ALX3	[]	['CGGYTCTGGAACCAGACCTGGGG', 'GCGGYTCTGGAACCAGACCTGGG', 'TGCGGYTCTGGAACCAGACCTGG']	['Frontonasal dysplasia 1']
NM_020451.2(SEPN1):c.1A>G (p.Met1Val)	121908184	SEPN1	[]	['CCCAYGGCTGCGGCTGGCGGCGG', 'CGGCCCAYGGCTGCGGCTGGCGG']	['Eichsfeld type congenital muscular dystrophy']
NM_001127221.1(CACNA1A):c.4151A>G (p.Tyr1384Cys)	121908219	CACNA1A	[]	[]	['Familial hemiplegic migraine type 1']
NM_005249.4(FOXG1):c.686T>A (p.Ile229Asn)	199502880	FOXG1	[]	[]	['not provided']
NM_130468.3(CHST14):c.878A>G (p.Tyr293Cys)	121908258	CHST14	[]	['AAGTCAYAGTGCACGGCACAAGG']	['Ehlers-Danlos syndrome, musculocontractural type']
NM_013391.3(DMGDH):c.326A>G (p.His109Arg)	121908331	DMGDH	[]	[]	['Dimethylglycine dehydrogenase deficiency']
NM_015166.3(MLC1):c.422A>G (p.Asn141Ser)	121908344	MLC1	[]	[]	['Megalencephalic leukoencephalopathy with subcortical cysts 1']
NM_000441.1(SLC26A4):c.1105A>G (p.Lys369Glu)	121908361	SLC26A4	[]	[]	['Enlarged vestibular aqueduct syndrome']
NM_000441.1(SLC26A4):c.2168A>G (p.His723Arg)	121908362	SLC26A4	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_015560.2(OPA1):c.1745A>G (p.Tyr582Cys)	121908376	OPA1	[]	[]	['Optic Atrophy Type 1']
NM_001128425.1(MUTYH):c.1241A>G (p.Gln414Arg)	121908383	MUTYH	[]	['AAGCYGCTCTGAGGGCTCCCAGG']	['Neoplasm of stomach']
NM_015247.2(CYLD):c.2240A>G	121908389	CYLD	[]	[]	['Familial multiple trichoepitheliomata',

(p.Glu747Gly)					'Spiegler-Brooke syndrome']
NM_021102.3(SPINT2):c.488A>G (p.Tyr163Cys)	121908403	SPINT2	['TCCAYAGATGAAGTTATTGCAGG']	['TCCAYAGATGAAGTTATTGCAGG']	['Diarrhea 3, secretory sodium, congenital, syndromic']
NM_004924.4(ACTN4):c.763A>G (p.Lys255Glu)	121908415	ACTN4	[]	[]	['Focal segmental glomerulosclerosis 1']
NM_004795.3(KL):c.578A>G (p.His193Arg)	121908423	KL	['CAGYGGTACAGGGTGACCACGGG', 'CCAGYGGTACAGGGTGACCACGG']	['CAGYGGTACAGGGTGACCACGGG', 'CCAGYGGTACAGGGTGACCACGG']	[]
NM_005682.6(ADGRG1):c.263A>G (p.Tyr88Cys)	121908466	ADGRG1	['TGGYAGAGGCCCCTGGGGTCAGG']	['TGGYAGAGGCCCCTGGGGTCAGG']	['Polymicrogyria, bilateral frontoparietal']
NM_139025.4(ADAMTS13):c.1582A>G (p.Arg528Gly)	121908473	ADAMTS13	[]	[]	['Upshaw-Schulman syndrome']
NM_014270.4(SLC7A9):c.695A>G (p.Tyr232Cys)	121908487	SLC7A9	[]	[]	['Cystinuria']
NM_004211.3(SLC6A5):c.1472A>G (p.Tyr491Cys)	121908494	SLC6A5	[]	[]	['Hyperekplexia 3']
NM_004211.3(SLC6A5):c.1526A>G (p.Asn509Ser)	121908497	SLC6A5	[]	[]	['Hyperekplexia 3']
NM_182643.2(DLC1):c.2875A>G (p.Thr959Ala)	121908500	DLC1	[]	[]	['Carcinoma of colon']
NM_014946.3(SPAST):c.1322A>G (p.Asp441Gly)	121908512	SPAST	[]	[]	['Spastic paraplegia 4, autosomal dominant']
NM_014946.3(SPAST):c.1157A>G (p.Asn386Ser)	121908514	SPAST	[]	[]	['Spastic paraplegia 4, autosomal dominant']
NM_000026.2(ADSL):c.736A>G (p.Lys246Glu)	119450944	ADSL	[]	[]	['Adenylosuccinate lyase deficiency']
NM_000334.4(SCN4A):c.3478A>G (p.Ile1160Val)	121908549	SCN4A	['TGAYGGAGGGGATGGCGCCTAGG']	['TGAYGGAGGGGATGGCGCCTAGG']	[]
NM_000334.4(SCN4A):c.421A>G (p.Ile141Val)	121908561	SCN4A	[]	[]	['Paramyotonia congenita of von Eulenburg']
NM_004328.4(BCS1L):c.148A>G (p.Thr50Ala)	121908580	BCS1L	[]	['GTGYGATCATGTAATGGCGCCGG']	['Mitochondrial complex III deficiency']
NM_152384.2(BBS5):c.547A>G (p.Thr183Ala)	121908582	BBS5	[]	[]	['Bardet-Biedl syndrome 5']
NM_016417.2(GLRX5	121908584	GLRX5	[]	['CCTGACCYTGTC	['Anemia,

):c.294A>G (p.Gln98=)				GGAGCTCCGGG']	sideroblastic, pyridoxine-refractory, autosomal recessive']
NM_006206.4(PDGFRA):c.1664A>G (p.Tyr555Cys)	121908589	PDGFRA	[]	[]	[]
NM_002755.3(MAP2K1):c.389A>G (p.Tyr130Cys)	121908595	MAP2K1	['CCAYAGAAGCCCACGATGTACGG']	['CCAYAGAAGCCCACGATGTACGG']	['Cardiofaciocutaneous syndrome 3', 'Rasopathy']
NM_012082.3(ZFPM2):c.89A>G (p.Glu30Gly)	121908601	ZFPM2	[]	[]	['Double outlet right ventricle', 'Tetralogy of Fallot', 'Diaphragmatic hernia 3']
NM_012082.3(ZFPM2):c.2527A>G (p.Thr843Ala)	121908604	-	[]	[]	['Diaphragmatic hernia 3']
NM_022817.2(PER2):c.1984A>G (p.Ser662Gly)	121908635	PER2	[]	['GCCACACYCTCTGCCTTGCCCGG']	['Advanced sleep phase syndrome, familial']
NM_030761.4(WNT4):c.647A>G (p.Glu216Gly)	121908650	WNT4	[]	[]	['Mullerian aplasia and hyperandrogenism']
NM_003839.3(TNFRSF11A):c.508A>G (p.Arg170Gly)	121908655	TNFRSF11A	[]	['GGGTCYGCATTTGTCCGTGGAGG']	['Osteopetrosis autosomal recessive 7']
NM_000539.3(RHO):c.886A>G (p.Lys296Glu)	29001653	RHO	[]	['CGCTCTYGGCAAAGAACGCTGGG', 'GCGCTCTYGGCAAAGAACGCTGG']	['Retinitis pigmentosa 4']
NM_004006.2(DMD):c.2317A>G (p.Lys773Glu)	128626244	DMD	[]	[]	['Duchenne muscular dystrophy']
NM_003722.4(TP63):c.697A>G (p.Lys233Glu)	121908838	TP63	['AGCTTYTTTGTAGACAGGCATGG']	['AGCTTYTTTGTAGACAGGCATGG']	['Split-hand/foot malformation 4']
NM_003722.4(TP63):c.1052A>G (p.Asp351Gly)	121908844	TP63	[]	[]	['Ectrodactyly, ectodermal dysplasia, and cleft lip/palate syndrome 3']
NM_003722.4(TP63):c.1054A>G (p.Arg352Gly)	121908847	TP63	[]	[]	['ADULT syndrome', 'Orofacial cleft 8']
NM_000369.2(TSHR):c.1856A>G (p.Asp619Gly)	121908859	TSHR	[]	[]	['Thyroid adenoma, hyperfunctioning']
NM_000369.2(TSHR):c.548A>G (p.Lys183Arg)	121908879	TSHR	[]	[]	['Hyperthyroidism, familial gestational']
NM_003060.3(SLC22A5):c.632A>G (p.Tyr211Cys)	121908888	SLC22A5	[]	[]	['Renal carnitine transport defect', 'not provided']
NM_006502.2(POLH):c.1603A>G	56307355	POLH	[]	['AGACTTTYCTGCTTAAAGAAGGG']	['Xeroderma pigmentosum,

(p.Lys535Glu)					variant type']
NM_002977.3(SCN9A):c.1964A>G (p.Lys655Arg)	121908919	-	[]	['CCTTTTCYTGTGTATTTGATTGG']	['Generalized epilepsy with febrile seizures plus, type 7', 'not specified']
NM_002977.3(SCN9A):c.184A>G (p.Ile62Val)	121908920	SCN9A	[]	[]	['Febrile seizures, familial, 3b']
NM_006892.3(DNMT3B):c.2450A>G (p.Asp817Gly)	121908939	DNMT3B	[]	['GACACGYCTGTGTAGTGCACAGG']	['Centromeric instability of chromosomes 1,9 and 16 and immunodeficiency']
NM_001130978.1(DYSF):c.3892A>G (p.Ile1298Val)	121908954	DYSF	[]	[]	['Miyoshi muscular dystrophy 1', 'Limb-girdle muscular dystrophy, type 2B', 'not specified']
NM_001130978.1(DYSF):c.5264A>G (p.Glu1755Gly)	121908961	DYSF	[]	[]	['Limb-girdle muscular dystrophy, type 2B']
NM_016203.3(PRKAG2):c.1148A>G (p.His383Arg)	121908988	PRKAG2	[]	[]	['Familial hypertrophic cardiomyopathy 6']
NM_000492.3(CFTR):c.2738A>G (p.Tyr913Cys)	121909008	CFTR	['CACATAAYACGAACTGGTGCTGG']	['CACATAAYACGAACTGGTGCTGG']	['Cystic fibrosis']
NM_000492.3(CFTR):c.326A>G (p.Tyr109Cys)	121909031	CFTR	[]	[]	['Cystic fibrosis']
NM_000492.3(CFTR):c.650A>G (p.Glu217Gly)	121909046	CFTR	[]	[]	['Cystic fibrosis']
NM_001040667.2(HSF4):c.256A>G (p.Ile86Val)	121909050	HSF4	[]	[]	['Cataract, zonular']
NM_005025.4(SERPINI1):c.1013A>G (p.His338Arg)	121909052	SERPINI1	[]	[]	['Familial encephalopathy with neuroserpin inclusion bodies']
NM_005422.2(TECTA):c.5609A>G (p.Tyr1870Cys)	121909058	TECTA	[]	[]	['Deafness, autosomal dominant 12']
NM_170695.3(TGIF1):c.838A>G (p.Thr280Ala)	121909068	TGIF1	[]	[]	['Holoprosencephaly 4']
NM_001005360.2(DNM2):c.1684A>G (p.Lys562Glu)	121909088	DNM2	[]	['ACTYCTTCTCTTTCTCCTGAGGG', 'TACTYCTTCTCTTTCTCCTGAGG']	['Charcot-Marie-Tooth disease, dominant intermediate b, with neutropenia']
NM_000483.4(APOC2):c.1A>G (p.Met1Val)	120074112	-	[]	['GCCCAYAGTGTCCAGAGACCTGG']	['Apolipoprotein C2 deficiency']
NM_000543.4(SMPD1):c.1154A>G (p.Asn385Ser)	120074123	SMPD1	[]	[]	['Niemann-Pick disease, type B']
NM_000019.3(ACAT1):c.278A>G	120074145	ACAT1	[]	[]	['Deficiency of acetyl-CoA

(p.Asn93Ser)					acetyltransferase']
NM_138477.2(CDAN1):c.1796A>G (p.Asn599Ser)	120074166	CDAN1	[]	[]	['Congenital dyserythropoietic anemia, type I']
NM_000187.3(HGD):c.1102A>G (p.Met368Val)	120074173	HGD	[]	[]	['Alkaptonuria']
NM_001089.2(ABCA3):c.1702A>G (p.Asn568Asp)	121909184	ABCA3	['ACCGTYGTGGCCCAGCAGGACGG']	['ACCGTYGTGGCCCAGCAGGACGG']	['Surfactant metabolism dysfunction, pulmonary, 3']
NM_000503.5(EYA1):c.1639A>G (p.Arg547Gly)	121909197	EYA1	[]	[]	[]
NM_000218.2(KCNQ1):c.418A>G (p.Ser140Gly)	120074192	KCNQ1	['CGCYGAAGATGAGGCAGACCAGG']	['CGCYGAAGATGAGGCAGACCAGG']	['Atrial fibrillation, familial, 3', 'Atrial fibrillation']
NM_000314.6(PTEN):c.368A>G (p.His123Arg)	121909222	PTEN	[]	[]	['Cowden syndrome 1']
NM_000314.6(PTEN):c.278A>G (p.His93Arg)	121909238	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome', 'Macrocephaly/autism syndrome']
NM_000314.6(PTEN):c.755A>G (p.Asp252Gly)	121909239	PTEN	[]	['ATAYCACCACACACAGGTAACGG']	['Macrocephaly/autism syndrome']
NM_198217.2(ING1):c.515A>G (p.Asn172Ser)	121909251	ING1	[]	['TGGYTGCACAGACAGTACGTGGG', 'CTGGYTGCACAGACAGTACGTGG']	['Squamous cell carcinoma of the head and neck']
NM_012338.3(TSPAN12):c.734T>C (p.Leu245Pro)	200519776	TSPAN12	[]	[]	['Exudative vitreoretinopathy 5']
NM_001001557.2(GDF6):c.1271A>G (p.Lys424Arg)	121909353	GDF6	[]	[]	['Klippel-Feil syndrome 1, autosomal dominant']
NM_000163.4(GHR):c.594A>G (p.Glu198=)	121909360	GHR	[]	[]	['Laron-type isolated somatotropin defect']
NM_000256.3(MYBPC3):c.175A>G (p.Thr59Ala)	121909375	MYBPC3	[]	[]	['Familial hypertrophic cardiomyopathy 4']
NM_001174089.1(SLC4A11):c.2518A>G (p.Met840Val)	121909396	SLC4A11	[]	['GATCAYCTTCATGTAGGGCAGGG', 'AGATCAYCTTCATGTAGGGCAGG']	['Corneal dystrophy and perceptive deafness']
NM_001100.3(ACTA1):c.350A>G (p.Asn117Ser)	121909520	ACTA1	['GCGGYTGGCCTTGGGATTGAGGG', 'CGCGGYTGGCCTTGGGATTGAGG']	['CGGYTGGCCTTGGGATTGAGGGG', 'GCGGYTGGCCTTGGGATTGAGGG', 'CGCGGYTGGCCTTGGGATTGAGG']	['Nemaline myopathy 3']
NM_000034.3(ALDO	121909533	ALDOA	[]	['CCAYCCAACCCT	['HNSHA due to

A):c.386A>G (p.Asp129Gly)				AAGAGAAGAGG']	aldolase A deficiency']
NM_000495.4(COL4A5):c.3455-9A>G	104886388	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_001145.4(ANG):c.208A>G (p.Ile70Val)	121909541	-	[]	[]	['Amyotrophic lateral sclerosis type 9']
NM_000051.3(ATM):c.3118A>G (p.Met1040Val)	3092857	ATM	[]	[]	['Hereditary cancer-predisposing syndrome', 'not specified']
NM_023110.2(FGFR1):c.1121A>G (p.Tyr374Cys)	121909631	FGFR1	[]	[]	['Osteoglophonic dysplasia']
NM_182925.4(FLT4):c.3104A>G (p.His1035Arg)	121909653	FLT4	['CTGYGGATGCACTGGGGTGCGGG', 'TCTGYGGATGCACTGGGGTGCGG']	['CTGYGGATGCACTGGGGTGCGGG', 'TCTGYGGATGCACTGGGGTGCGG']	[]
NM_000145.3(FSHR):c.1345A>G (p.Thr449Ala)	121909663	FSHR	[]	[]	['Ovarian hyperstimulation syndrome']
NM_001017420.2(ESCO2):c.1132-7A>G	80359862	ESCO2	[]	[]	['Roberts-SC phocomelia syndrome']
NM_001017420.2(ESCO2):c.1674-2A>G	80359869	ESCO2	[]	[]	['Roberts-SC phocomelia syndrome']
NM_024577.3(SH3TC2):c.505T>C (p.Tyr169His)	80359890	SH3TC2	[]	[]	['Charcot-Marie-Tooth disease, type 4C', 'Charcot-Marie-Tooth disease, type IV', 'Mononeuropathy of the median nerve, mild']
NM_032119.3(ADGRV1):c.18131A>G (p.Tyr6044Cys)	121909763	ADGRV1	[]	[]	['Usher syndrome, type 2C']
NM_001360.2(DHCR7):c.839A>G (p.Tyr280Cys)	121909766	DHCR7	[]	[]	['Smith-Lemli-Opitz syndrome']
NM_000517.4(HBA2):c.1A>G (p.Met1Val)	121909803	HBA2	[]	[]	['Hemoglobin H disease, nondeletional']
NM_004006.2(DMD):c.835A>G (p.Thr279Ala)	128627255	DMD	[]	['TGACCGYGATCTGCAGAGAAGGG', 'CTGACCGYGATCTGCAGAGAAGG']	['Dilated cardiomyopathy 3B']
NM_015896.3(ZMYND10):c.797T>C (p.Leu266Pro)	200913791	ZMYND10	[]	[]	['Kartagener syndrome', 'Ciliary dyskinesia, primary, 22']
NM_001085.4(SERPINA3):c.1240A>G (p.Met414Val)	116929575	SERPINA3	[]	['GCTCAYGAAGAAGATGTTCTGGG', 'TGCTCAYGAAGA	[]

				AGATGTTCTGG']	
NM_058216.2(RAD51C):c.1027-2A>G	587780835	RAD51C	[]	[]	['Fanconi anemia, complementation group O']
NM_006231.3(POLE):c.4444+3A>G	398122515	POLE	[]	[]	['Facial dysmorphism, immunodeficiency, livedo, and short stature']
NM_002769.4(PRSS1):c.161A>G (p.Asn54Ser)	144422014	-	[]	[]	['Hereditary pancreatitis']
NM_001204316.1(PRLR):c.635A>G (p.His212Arg)	398122522	PRLR	[]	[]	['Hyperprolactinemia']
NM_004992.3(MECP2):c.410A>G (p.Glu137Gly)	61748392	MECP2	[]	['CAACYCCACTTTAGAGCGAAAGG']	['Mental retardation, X-linked, syndromic 13']
NM_020366.3(RPGRIP1):c.3749-2A>G	376517859	RPGRIP1	[]	[]	['Cone-rod dystrophy 13']
NM_000552.3(VWF):c.2384A>G (p.Tyr795Cys)	61748478	VWF	['GTCAYAGTTCTGGCACGTTTTGG']	['GTCAYAGTTCTGGCACGTTTTGG']	['von Willebrand disease type 2N', 'not provided']
NM_001040613.2(TMEM70):c.*7-2A>G	183973249	TMEM70	[]	[]	['Nuclearly-encoded mitochondrial complex V (ATP synthase) deficiency 2']
NM_001005741.2(GBA):c.1049A>G (p.His350Arg)	78198234	GBA	[]	[]	['Gaucher disease, perinatal lethal']
NM_000218.2(KCNQ1):c.332A>G (p.Tyr111Cys)	199472678	KCNQ1	[]	[]	['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.344A>G (p.Glu115Gly)	199472679	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ1):c.440A>G (p.Gln147Arg)	199472689	KCNQ1			['Atrial fibrillation']
NM_000218.2(KCNQ1):c.592A>G (p.Ile198Val)	199472700	KCNQ1	[]	[]	['Congenital long QT syndrome']
NM_001943.3(DSG2):c.880A>G (p.Lys294Glu)	752432726	DSG2	[]	[]	['Cardiomyopathy']
NM_000218.2(KCNQ1):c.820A>G (p.Ile274Val)	199472728	KCNQ1	[]	[]	['Sudden infant death syndrome', 'Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000218.2(KCNQ1):c.842A>G (p.Tyr281Cys)	199472732	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ	199472750	KCNQ1			['Congenital long

1):c.950A>G (p.Asp317Gly)					QT syndrome']
NM_000218.2(KCNQ 1):c.964A>G (p.Thr322Ala)	199472754	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ 1):c.1138A>G (p.Arg380Gly)	199472770	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.1193A>G (p.Lys398Arg)	199472777	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000218.2(KCNQ 1):c.1640A>G (p.Gln547Arg)	199472798	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.1669A>G (p.Lys557Glu)	199472801	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.1705A>G (p.Lys569Glu)	199472808	KCNQ1			['Congenital long QT syndrome']
NM_001005741.2(GB A):c.667T>C (p.Trp223Arg)	61748906	GBA	[]	['CCCACTYGGCTC AAGACCAATGG']	['Gaucher disease, type 1', 'not provided']
NM_000218.2(KCNQ 1):c.1756A>G (p.Asn586Asp)	199472812	KCNQ1			['Congenital long QT syndrome', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ 1):c.1793A>G (p.Lys598Arg)	199472817	KCNQ1			['Sudden infant death syndrome']
NM_000238.3(KCNH 2):c.82A>G (p.Lys28Glu)	199472829	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.128A>G (p.Tyr43Cys)	199472836	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH 2):c.301A>G (p.Lys101Glu)	199472856	KCNH2	[]	[]	['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH 2):c.652A>G (p.Met218Val)	199472869	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1424A>G (p.Tyr475Cys)	199472907	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1777A>G (p.Ile593Val)	199472930	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1783A>G (p.Lys595Glu)	199472932	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1790A>G (p.Tyr597Cys)	199472934	KCNH2			['Long QT syndrome', 'Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1826A>G	199472940	KCNH2			['Congenital long QT syndrome']

(p.Asp609Gly)					
NM_000238.3(KCNH2):c.1847A>G (p.Tyr616Cys)	199472946	KCNH2			['Long QT syndrome', 'Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.1885A>G (p.Asn629Asp)	199472956	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.1897A>G (p.Asn633Asp)	199472960	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1903A>G (p.Asn635Asp)	199472963	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.1910A>G (p.Glu637Gly)	199472967	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.2510A>G (p.Asp837Gly)	199473004	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.2591A>G (p.Asp864Gly)	199473008	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH2):c.3118A>G (p.Ser1040Gly)	199473024	KCNH2	[]	['CTGCYCTCCACGTCGCCCCGGGG', 'CCTGCYCTCCACGTCGCCCCGGG', 'GCCTGCYCTCCACGTCGCCCCGG']	['Sudden infant death syndrome']
NM_000238.3(KCNH2):c.3233A>G (p.Tyr1078Cys)	199473029	KCNH2			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.343A>G (p.Ser115Gly)	199473057	SCN5A			['Congenital long QT syndrome']
m.827A>G	28358569	MT-RNR1	[]	[]	['Aminoglycoside-induced deafness', 'Deafness, nonsyndromic sensorineural, mitochondrial']
NM_000335.4(SCN5A):c.688A>G (p.Ile230Val)	199473074	SCN5A	['ATAYAGTTTTCAGGGCCCGGAGG', 'CTGATAYAGTTTTCAGGGCCCGG']	['ATAYAGTTTTCAGGGCCCGGAGG', 'CTGATAYAGTTTTCAGGGCCCGG']	['Brugada syndrome']
NM_000335.4(SCN5A):c.715A>G (p.Ile239Val)	199473075	SCN5A			['Congenital long QT syndrome']
NM_000252.2(MTM1):c.575A>G (p.Tyr192Cys)	587783838	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_004572.3(PKP2):c.275T>A (p.Leu92Ter)	763639737	PKP2	[]	[]	['not provided']

NM_000335.4(SCN5A):c.1502A>G (p.Asp501Gly)	199473117	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.2249A>G (p.Gln750Arg)	199473152	SCN5A			['Congenital long QT syndrome']
NM_198056.2(SCN5A):c.2527A>G (p.Thr843Ala)	199473165	SCN5A			['Congenital long QT syndrome', 'not provided']
NM_001165963.1(SCN1A):c.1277A>G (p.Tyr426Cys)	796052973	SCN1A	[]	[]	['not provided']
NM_000335.4(SCN5A):c.3755A>G (p.Glu1252Gly)	199473214	SCN5A			['Brugada syndrome']
NM_198056.2(SCN5A):c.4000A>G (p.Ile1334Val)	199473226	SCN5A	[]	[]	['Congenital long QT syndrome', 'not provided']
NM_000335.4(SCN5A):c.4252A>G (p.Lys1418Glu)	199473242	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4291A>G (p.Arg1431Gly)	199473245	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4412A>G (p.Asn1471Ser)	199473255	SCN5A			['Congenital long QT syndrome']
NM_198056.2(SCN5A):c.4478A>G (p.Lys1493Arg)	199473260	SCN5A	[]	[]	['Atrial fibrillation', 'Congenital long QT syndrome', 'not provided']
NM_000335.4(SCN5A):c.4489A>G (p.Met1497Val)	199473264	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.4577A>G (p.Lys1526Arg)	199473270	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.5161A>G (p.Asn1721Asp)	199473299	SCN5A			['Brugada syndrome']
NM_198056.2(SCN5A):c.5302A>G (p.Ile1768Val)	199473311	SCN5A			['Congenital long QT syndrome', 'not provided']
NM_000335.4(SCN5A):c.5318A>G (p.Asn1773Ser)	199473313	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.5366A>G (p.Asp1789Gly)	199473317	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.5402A>G (p.Asp1801Gly)	199473318	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.5513A>G (p.Asp1838Gly)	199473321	SCN5A			['Congenital long QT syndrome']
NM_198056.2(SCN5A):c.5726A>G (p.Gln1909Arg)	199473326	SCN5A			['Congenital long QT syndrome', 'not provided']
NM_172201.1(KCNE2	199473366	KCNE2			['Atrial fibrillation']

):c.269A>G (p.Glu90Gly)					
NM_000891.2(KCNJ2):c.223A>G (p.Thr75Ala)	199473370	KCNJ2			['Congenital long QT syndrome']
NM_000891.2(KCNJ2):c.233A>G (p.Asp78Gly)	199473371	KCNJ2			['Andersen Tawil syndrome', 'Congenital long QT syndrome']
NM_000891.2(KCNJ2):c.574A>G (p.Thr192Ala)	199473382	KCNJ2			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.548A>G (p.Lys183Arg)	199473396	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.1061A>G (p.Lys354Arg)	199473404	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.1070A>G (p.Gln357Arg)	199473405	KCNQ1			['Congenital long QT syndrome', 'Cardiac arrhythmia', 'Long QT syndrome, LQT1 subtype']
NM_000218.2(KCNQ 1):c.1078A>G (p.Arg360Gly)	199473406	KCNQ1	[]	[]	['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.209A>G (p.His70Arg)	199473419	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH 2):c.1724A>G (p.Glu575Gly)	199473424	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1747A>G (p.Ile583Val)	199473427	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1762A>G (p.Asn588Asp)	199473431	KCNH2			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.430A>G (p.Thr144Ala)	199473451	KCNQ1			['Congenital long QT syndrome']
NM_000218.2(KCNQ 1):c.931A>G (p.Thr311Ala)	199473469	KCNQ1			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.286A>G (p.Ile96Val)	199473496	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1205A>G (p.His402Arg)	199473506	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1259A>G (p.Tyr420Cys)	199473507	KCNH2			['Congenital long QT syndrome']
NM_000238.3(KCNH 2):c.1502A>G (p.Asp501Gly)	199473513	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH 2):c.1912A>G (p.Lys638Glu)	199473528	KCNH2			['Congenital long QT syndrome']

NM_000238.3(KCNH2):c.2131A>G (p.Ile711Val)	199473532	KCNH2	[]	[]	['Long QT syndrome', 'Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000238.3(KCNH2):c.2266A>G (p.Met756Val)	199473534	KCNH2			['Acquired long QT syndrome']
NM_000238.3(KCNH2):c.3343A>G (p.Met1115Val)	199473546	KCNH2			['Congenital long QT syndrome', 'Cardiac arrhythmia']
NM_000335.4(SCN5A):c.89A>G (p.Glu30Gly)	199473551	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.1217A>G (p.Asn406Ser)	199473568	SCN5A			['Brugada syndrome']
NM_000249.3(MLH1):c.791-2A>G	267607794	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms', 'Hereditary cancer-predisposing syndrome']
NM_000335.4(SCN5A):c.2780A>G (p.Asn927Ser)	199473589	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.3164A>G (p.Asp1055Gly)	199473593	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.4223A>G (p.Tyr1408Cys)	199473610	SCN5A			['Brugada syndrome']
NM_198056.2(SCN5A):c.4346A>G (p.Tyr1449Cys)	199473613	SCN5A			['Brugada syndrome', 'not provided']
NM_198056.2(SCN5A):c.4978A>G (p.Ile1660Val)	199473625	SCN5A	['CGAYGTTGAAGAGGGCAGGCAGG']	['CGAYGTTGAAGAGGGCAGGCAGG', 'AGCCCGAYGTTGAAGAGGGCAGG']	['Brugada syndrome', 'not provided']
NM_000335.4(SCN5A):c.5138A>G (p.Asp1713Gly)	199473628	SCN5A			['Brugada syndrome']
NM_000335.4(SCN5A):c.5297A>G (p.Tyr1766Cys)	199473632	SCN5A			['Congenital long QT syndrome']
NM_000335.4(SCN5A):c.5317A>G (p.Asn1773Asp)	199473633	SCN5A			['Congenital long QT syndrome']
NM_000531.5(OTC):c.527A>G (p.Tyr176Cys)	72556283	OTC	['TGAGGYAATCAGCCAGGATCTGG']	['TGAGGYAATCAGCCAGGATCTGG']	['not provided']
NM_001130823.1(DNMT1):c.1532A>G (p.Tyr511Cys)	199473690	DNMT1			['Hereditary sensory neuropathy type IE']
NM_000303.2(PMM2):c.563A>G	80338704	PMM2	[]	[]	['Carbohydrate-deficient

(p.Asp188Gly)					glycoprotein syndrome type I', 'not provided']
NM_000051.3(ATM): c.8030A>G (p.Tyr2677Cys)	28942103	-	[]	[]	['Ataxia-telangiectasia variant']
NM_175053.3(KRT74):c.821T>C (p.Phe274Ser)	147962513	KRT74	[]	[]	["Ectodermal dysplasia, 'pure' hair-nail type", 'Ectodermal dysplasia 7, hair/nail type']
NM_000059.3(BRCA2):c.426-2A>G	398122779	BRCA2	[]	[]	['Familial cancer of breast', 'Breast-ovarian cancer, familial 2']
NM_133433.3(NIPBL) :c.5428-2A>G	587783974	NIPBL	[]	[]	['Cornelia de Lange syndrome 1']
NM_000238.3(KCNH 2):c.1129-2A>G	794728365	KCNH2	[]	['GGACCYGCACC CGGGGAAGGCGG']	['Cardiac arrhythmia']
NM_000260.3(MYO7 A):c.6029A>G (p.Asp2010Gly)	111033175	MYO7A	[]	[]	['Usher syndrome, type 1']
NM_000531.5(OTC):c .548A>G (p.Tyr183Cys)	72556293	OTC	[]	['AGAGCTAYAGT GTTCCTAAAAGG']	['not provided']
NM_000441.1(SLC26 A4):c.1151A>G (p.Glu384Gly)	111033244	SLC26A4	[]	['TGAATYCCTAAG GAAGAGACTGG']	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_004004.5(GJB2): c.617A>G (p.Asn206Ser)	111033294	GJB2	[]	[]	['Deafness, autosomal recessive 1A', 'Hearing impairment']
NM_000441.1(SLC26 A4):c.919-2A>G	111033313	SLC26A4	[]	[]	['Pendred syndrome', 'Enlarged vestibular aqueduct syndrome']
NM_001363.4(DKC1): c.115A>G (p.Lys39Glu)	121912296	DKC1	[]	[]	['Dyskeratosis congenita X-linked']
NM_001363.4(DKC1): c.196A>G (p.Thr66Ala)	121912297	DKC1	[]	[]	['Dyskeratosis congenita X-linked']
NM_001363.4(DKC1): c.361A>G (p.Ser121Gly)	121912305	DKC1	[]	[]	['Dyskeratosis congenita X-linked', 'Hoyeraal Hreidarsson syndrome']
NM_178151.2(DCX):c .413A>G (p.Tyr138Cys)	587783552	DCX	[]	[]	['Heterotopia']
NM_024675.3(PALB2):c.212-2A>G	730881879	PALB2	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000260.3(MYO7 A):c.1344-2A>G	111033415	MYO7A	[]	['AGCYGCAGGGG CACAGGGATGGG' , 'AAGCYGCAGGGG CACAGGGATGG']	['Usher syndrome, type 1']

NM_000454.4(SOD1):c.131A>G (p.His44Arg)	121912435	SOD1	[]	[]	['Amyotrophic lateral sclerosis type 1']
NM_000454.4(SOD1):c.302A>G (p.Glu101Gly)	121912439	SOD1	[]	['AGAATCTYCAATAGACACATCGG']	['Amyotrophic lateral sclerosis type 1']
NM_000454.4(SOD1):c.140A>G (p.His47Arg)	121912443	SOD1	[]	[]	['Amyotrophic lateral sclerosis type 1']
NM_133433.3(NIPBL):c.737A>G (p.Asp246Gly)	587784042	NIPBL	[]	[]	['Cornelia de Lange syndrome 1']
NM_000454.4(SOD1):c.242A>G (p.His81Arg)	121912458	SOD1	[]	[]	['Amyotrophic lateral sclerosis type 1']
NM_001754.4(RUNX1):c.328A>G (p.Lys110Glu)	121912498	RUNX1	[]	[]	['Familial platelet disorder with associated myeloid malignancy']
NM_000238.3(KCNH2):c.1408A>G (p.Asn470Asp)	121912505	KCNH2	[]	[]	['Long QT syndrome 2', 'Congenital long QT syndrome']
NM_000233.3(LHCGR):c.1733A>G (p.Asp578Gly)	121912518	-	[]	[]	['Gonadotropin-independent familial sexual precocity']
NM_000493.3(COL10A1):c.1790A>G (p.Tyr597Cys)	111033554	-	[]	[]	['Metaphyseal chondrodysplasia, Schmid type']
NM_000233.3(LHCGR):c.1691A>G (p.Asp564Gly)	121912540	-	[]	[]	['Gonadotropin-independent familial sexual precocity']
NM_002769.4(PRSS1):c.68A>G (p.Lys23Arg)	111033567	-	[]	['ATCYTGTCATCATCATCAAAGGG', 'GATCYTGTCATCATCATCAAAGG']	['Hereditary pancreatitis']
NM_004999.3(MYO6):c.737A>G (p.His246Arg)	121912560	MYO6	[]	[]	['Sensorineural deafness with hypertrophic cardiomyopathy']
NM_000901.4(NR3C2):c.2327A>G (p.Gln776Arg)	121912565	NR3C2	[]	['TCATCYGTTTGCCTGCTAAGCGG']	['Pseudohypoaldosteronism type 1 autosomal dominant']
NM_000901.4(NR3C2):c.2915A>G (p.Glu972Gly)	121912574	NR3C2	[]	['CCGACYCCACCTTGGGCAGCTGG']	['Pseudohypoaldosteronism type 1 autosomal dominant']
NM_001173464.1(KIF21A):c.2839A>G (p.Met947Val)	121912589	KIF21A	[]	['ATTCAYATCTGCCTCCATGTTGG']	['Fibrosis of extraocular muscles, congenital, 1']
NM_000155.3(GALT):c.67A>G (p.Thr23Ala)	111033635	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_001041.3(SI):c.350A>G (p.Gln117Arg)	121912612	SI	[]	[]	['Sucrase-isomaltase deficiency']
NM_000155.3(GALT):c.1A>G (p.Met1Val)	111033639	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate

					uridylyltransferase']
NM_021625.4(TRPV4):c.998A>G (p.Asp333Gly)	121912634	TRPV4	[]	[]	['Spondylometaphyseal dysplasia, Kozlowski type']
NM_000155.3(GALT):c.253-2A>G	111033661	GALT	[]	['ATTCACCYACCGACAAGGATAGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase', 'not provided']
NM_000155.3(GALT):c.290A>G (p.Asn97Ser)	111033669	GALT	[]	['GAAGTCGYTGTCAAACAGGAAGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.379A>G (p.Lys127Glu)	111033682	GALT	[]	['TGACCTYACTGGGTGGTGACGGG', 'ATGACCTYACTGGGTGGTGACGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000343.3(SLC5A1):c.83A>G (p.Asp28Gly)	121912669	SLC5A1	[]	[]	['Congenital glucose-galactose malabsorption']
NM_005159.4(ACTC1):c.1088A>G (p.Glu363Gly)	121912674	-	[]	[]	['Dilated cardiomyopathy 1R']
NM_005159.4(ACTC1):c.373A>G (p.Met125Val)	121912677	-	[]	[]	['Atrial septal defect 5']
NM_000155.3(GALT):c.565-2A>G	111033731	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_001681.3(ATP2A2):c.2300A>G (p.Asn767Ser)	121912732	ATP2A2	[]	[]	['Darier disease, acral hemorrhagic type']
NM_000155.3(GALT):c.821-2A>G	111033767	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000342.3(SLC4A1):c.2509A>G (p.Thr837Ala)	121912750	SLC4A1	[]	[]	['Spherocytosis type 4']
NM_000155.3(GALT):c.950A>G (p.Gln317Arg)	111033786	GALT	[]	['CAGCYGCCAATGGTTCCAGTTGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_001202.3(BMP4):c.278A>G (p.Glu93Gly)	121912765	BMP4	[]	['CCTCCYCCCCAGACTGAAGCCGG']	['Microphthalmia syndromic 6']
NM_000155.3(GALT):c.1001A>G (p.Lys334Arg)	111033809	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.1048A>G (p.Thr350Ala)	111033817	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.1132A>G (p.Ile378Val)	111033819	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']

m.3243A>G	199474657	MT-TL1	[]	[]	['Leigh disease', 'Cyclical vomiting syndrome', 'Juvenile myopathy, encephalopathy, lactic acidosis AND stroke', 'Myoclonus with epilepsy with ragged red fibers', 'Cytochrome-c oxidase deficiency', 'Diabetes-deafness syndrome maternally transmitted', '3-Methylglutaconic aciduria', 'Age-related macular degeneration 2', 'MERRF/MELAS overlap syndrome']
m.3252A>G	199474661	MT-TL1	[]	[]	['Mitochondrial encephalomyopathy']
m.3251A>G	199474662	MT-TL1	[]	[]	[]
NM_000258.2(MYL3):c.517A>G (p.Met173Val)	199474708	MYL3	[]	[]	['Cardiomyopathy', 'not specified', 'not provided']
NM_000094.3(COL7A1):c.425A>G (p.Lys142Arg)	121912856	COL7A1	[]	['CACCYTGGGGACACCAGGTCGGG', 'TCACCYTGGGGACACCAGGTCGG']	['Epidermolysis bullosa dystrophica inversa, autosomal recessive']
NM_152263.3(TPM3):c.505A>G (p.Lys169Glu)	199474715	TPM3	[]	['CCAACTYACGAGCCACCTACAGG']	['Congenital myopathy with fiber type disproportion', 'not provided']
NM_152263.3(TPM3):c.733A>G (p.Arg245Gly)	199474718	TPM3	[]	['ATCYCTCAGCAAACTCAGCACGG']	['Congenital myopathy with fiber type disproportion', 'not provided']
NM_001844.4(COL2A1):c.4172A>G (p.Tyr1391Cys)	121912889	COL2A1	['GCAGTGGYAGGTGATGTTCTGGG']	['GCAGTGGYAGGTGATGTTCTGGG']	['Spondyloperipheral dysplasia', 'Platyspondylic lethal skeletal dysplasia Torrance type']
NM_001844.4(COL2A1):c.2974A>G (p.Arg992Gly)	121912895	COL2A1	[]	['CCTCYCTCACCACGTTGCCCAGG']	['Spondyloepimetaphyseal dysplasia Strudwick type']
NM_001848.2(COL6A1):c.362A>G (p.Lys121Arg)	121912936	COL6A1	[]	[]	['Ullrich congenital muscular dystrophy', 'Bethlem myopathy', 'not provided']
NM_004004.5(GJB2):c.218A>G (p.His73Arg)	121912968	GJB2	[]	[]	['Keratoderma palmoplantar deafness']
NM_000941.2(POR):c.1733A>G (p.Tyr578Cys)	121912975	POR	[]	[]	['Antley-Bixler syndrome with genital anomalies

					and disordered steroidogenesis']
NM_001943.3(DSG2):c.797A>G (p.Asn266Ser)	121913011	DSG2	[]	[]	['Arrhythmogenic right ventricular cardiomyopathy, type 10']
NM_000129.3(F13A1):c.851A>G (p.Tyr284Cys)	121913074	F13A1	[]	['ATAGGCAYAGATATTGTCCCAGG']	['Factor xiii, a subunit, deficiency of']
NM_000043.4(FAS):c.695A>G (p.Tyr232Cys)	121913079	FAS	[]	[]	['Autoimmune lymphoproliferative syndrome, type 1a']
NM_000043.4(FAS):c.763A>G (p.Asn255Asp)	121913082	FAS	[]	[]	[]
NM_000043.4(FAS):c.353A>G (p.Asn118Ser)	121913083	FAS	[]	[]	[]
NM_206933.2(USH2A):c.14020A>G (p.Arg4674Gly)	80338904	USH2A	[]	[]	['Retinitis pigmentosa', 'Retinitis pigmentosa 39']
NM_000142.4(FGFR3):c.833A>G (p.Tyr278Cys)	121913115	FGFR3	[]	[]	['Hypochondroplasia']
NM_000183.2(HADHB):c.788A>G (p.Asp263Gly)	121913131	HADHB	[]	[]	['Mitochondrial trifunctional protein deficiency']
NM_001079817.1(INSR):c.1459A>G (p.Lys487Glu)	121913136	INSR	[]	[]	['Leprechaunism syndrome']
NM_000208.2(INSR):c.707A>G (p.His236Arg)	121913145	INSR	[]	['GCTGYGGCAACAGAGGCCTTCGG']	['Leprechaunism syndrome']
NM_000208.2(INSR):c.1466A>G (p.Asn489Ser)	121913147	INSR	[]	[]	['Insulin-resistant diabetes mellitus AND acanthosis nigricans']
NM_000208.2(INSR):c.1372A>G (p.Asn458Asp)	121913160	INSR	[]	[]	['Leprechaunism syndrome']
NM_000016.5(ACADM):c.797A>G (p.Asp266Gly)	201375579	ACADM	[]	[]	['not provided']
NM_024577.3(SH3TC2):c.530-2A>G	80338920	SH3TC2	[]	[]	['Charcot-Marie-Tooth disease, type 4C']
NM_001127500.1(MET):c.3743A>G (p.Tyr1248Cys)	121913246	MET	[]	[]	['Renal cell carcinoma, papillary, 1']
NM_000517.4(HBA2):c.*92A>G	63750067	HBA2	['ACTTYATTCAAAGACCAGGAAGG']	['CTTYATTCAAAGACCAGGAAGGG', 'ACTTYATTCAAAGACCAGGAAGG']	['Hemoglobin H disease, nondeletional']
NM_199440.1(HSPD1):c.86A>G (p.Asp29Gly)	72466451	HSPD1	[]	[]	['Leukodystrophy, hypomyelinating, 4']
NM_000249.3(MLH1):c.544A>G	63750211	MLH1	[]	[]	['Hereditary Nonpolyposis

(p.Arg182Gly)					Colorectal Neoplasms']
NM_025137.3(SPG11):c.1457-2A>G	312262726	SPG11	[]	[]	['Spastic paraplegia 11, autosomal recessive']
NM_025137.3(SPG11):c.2608A>G (p.Ile870Val)	312262745	SPG11	[]	['ACTTAYCCTGGGGAGAAGGATGG']	['Spastic paraplegia 11, autosomal recessive']
NM_025137.3(SPG11):c.2833A>G (p.Arg945Gly)	312262748	SPG11	[]	[]	['Spastic paraplegia 11, autosomal recessive']
NM_003867.3(FGF17):c.560A>G (p.Asn187Ser)	398123026	FGF17	['CGTGGYTGGGGAAGGGCAGCTGG']	['CGTGGYTGGGGAAGGGCAGCTGG']	['Hypogonadotropic hypogonadism 20 with or without anosmia']
NM_025137.3(SPG11):c.6477+4A>G	312262780	SPG11	[]	[]	['Spastic paraplegia 11, autosomal recessive']
NM_000141.4(FGFR2):c.1124A>G (p.Tyr375Cys)	121913478	FGFR2	[]	[]	['Cutis Gyrata syndrome of Beare and Stevenson', 'Endometrial carcinoma']
NM_000142.4(FGFR3):c.1118A>G (p.Tyr373Cys)	121913485	FGFR3	[]	[]	['Thanatophoric dysplasia type 1']
NM_003611.2(OFD1):c.290A>G (p.Glu97Gly)	312262820	OFD1	[]	[]	['Oral-facial-digital syndrome']
NM_000222.2(KIT):c.1924A>G (p.Lys642Glu)	121913512	KIT	['GACTTYGAGTTCAGACATGAGGG']	['GACTTYGAGTTCAGACATGAGGG', 'GGACTTYGAGTTCAGACATGAGG']	[]
NM_003611.2(OFD1):c.382-2A>G	312262829	OFD1	[]	[]	['Oral-facial-digital syndrome']
NM_000391.3(TPP1):c.857A>G (p.Asn286Ser)	119455958	TPP1	[]	[]	['Ceroid lipofuscinosis, neuronal, 2']
NM_005912.2(MC4R):c.508A>G (p.Ile170Val)	121913560	MC4R	[]	[]	['Obesity']
NM_005912.2(MC4R):c.821A>G (p.Asn274Ser)	121913561	MC4R	[]	[]	['Obesity']
NM_005912.2(MC4R):c.289A>G (p.Asn97Asp)	121913565	MC4R	[]	[]	['Obesity']
NM_005912.2(MC4R):c.185A>G (p.Asn62Ser)	121913566	MC4R	[]	[]	['Obesity']
NM_000530.6(MPZ):c.286A>G (p.Lys96Glu)	121913583	MPZ	[]	[]	['Charcot-Marie-Tooth disease type 1B']
NM_000095.2(COMP):c.1760A>G (p.His587Arg)	312262901	COMP	[]	[]	['Pseudoachondroplastic spondyloepiphyseal dysplasia syndrome']
NM_000530.6(MPZ):c.242A>G (p.His81Arg)	121913594	MPZ	['GGCATAGYGGAAGAT	['GGCATAGYGGAAGATCTATGAGG']	['Charcot-Marie-Tooth disease type

			CTATGAGG']		1B']
NM_000484.3(APP):c.2146A>G (p.Ile716Val)	63750399	APP	[]	[]	['Alzheimer disease, type 1', 'not provided']
NM_000329.2(RPE65):c.1292A>G (p.Tyr431Cys)	62636300	RPE65	[]	[]	['Leber congenital amaurosis 2', 'not provided']
NM_002470.3(MYH3):c.1385A>G (p.Asp462Gly)	121913622	MYH3	[]	[]	['Distal arthrogryposis type 2B']
NM_000257.3(MYH7):c.2333A>G (p.Asp778Gly)	121913634	MYH7	[]	[]	['Familial hypertrophic cardiomyopathy 1', 'not specified']
NM_001127500.1(MET):c.3785A>G (p.Lys1262Arg)	121913677	MET	[]	[]	['Childhood hepatocellular carcinoma']
NM_000222.2(KIT):c.2459A>G (p.Asp820Gly)	121913682	KIT	[]	['AGAAYCATTCTTGATGTCTCTGG']	['Mast cell disease, systemic']
NM_000222.2(KIT):c.2386A>G (p.Arg796Gly)	121913684	KIT	[]	[]	[]
NM_006005.3(WFS1):c.1385A>G (p.Glu462Gly)	398123066	WFS1	[]	[]	['Cataract, nuclear total']
NM_000495.4(COL4A5):c.3925-2A>G	587776400	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NC_012920.1:m.13514A>G	587776440	MT-ND5	[]	[]	['Leigh disease']
NM_000021.3(PSEN1):c.488A>G (p.His163Arg)	63750590	PSEN1	[]	[]	['Alzheimer disease, type 3', 'not provided']
NM_000484.3(APP):c.2140A>G (p.Thr714Ala)	63750643	APP	[]	[]	['Alzheimer disease, type 1', 'not provided']
NM_173560.3(RFX6):c.224-12A>G	587776515	RFX6	[]	[]	['Mitchell-Riley syndrome']
NM_014043.3(CHMP2B):c.85A>G (p.Ile29Val)	63750818	CHMP2B	[]	[]	['Frontotemporal Dementia, Chromosome 3-Linked', 'Amyotrophic lateral sclerosis 17', 'not provided']
NM_000057.3(BLM):c.1088-2A>G	367543015	BLM	[]	[]	['Bloom syndrome']
NM_001011658.3(TRAPPC2):c.238+4T>C	587776753	-	[]	[]	['Spondyloepiphyseal dysplasia tarda']
NM_000151.3(G6PC):c.230+4A>G	587776757	G6PC	[]	['GTTCYTACCACTTAAAGACGAGG']	['Glycogen storage disease type 1A']
NM_000463.2(UGT1A1):c.1085-2A>G	587776766	-	['ACCYGAGATGCAAAATAGGGAGG']	['ACCYGAGATGCAAAATAGGGAGG', 'GTGACCYGAGATGCAAAATAGGG', 'GGTGACCYGAGATGCAAAATAGG']	['Crigler Najjar syndrome, type 1']

NM_000330.3(RS1):c.286T>C (p.Trp96Arg)	61752063	-	[]	['TTCTTCGYGGACTGCAAACAAGG']	['Juvenile retinoschisis', 'not provided']
NM_001024847.2(TGFBR2):c.1472-2A>G	587776770	TGFBR2	[]	[]	['Loeys-Dietz syndrome 2']
NM_000257.3(MYH7):c.5807A>G (p.Ter1936Trp)	367543053	MYH7	[]	[]	['Congenital myopathy with fiber type disproportion']
NM_000321.2(RB1):c.2490-1398A>G	587776791	RB1	[]	[]	['Retinoblastoma']
NM_024549.5(TCTN1):c.221-2A>G	367543065	TCTN1	[]	['AGCAACYGCAGAAAAAAGAGGGG', 'CAGCAACYGCAGAAAAAAGAGGG']	['Joubert syndrome 13']
NM_000228.2(LAMB3):c.565-3T>C	587776813	LAMB3	[]	[]	['Adult junctional epidermolysis bullosa']
NM_015884.3(MBTPS2):c.1523A>G (p.Asn508Ser)	587776867	MBTPS2	[]	[]	['Keratosis pilaris decalvans']
NM_000174.4(GP9):c.182A>G (p.Asn61Ser)	5030764	GP9	['GGCTGYTGTTGGCCAGCAGAAGG']	['GGCTGYTGTTGGCCAGCAGAAGG']	['Bernard-Soulier syndrome type C']
NM_000894.2(LHB):c.221A>G (p.Gln74Arg)	5030773	LHB	[]	['CCACCYGAGGCAGGGGCGGCAGG']	['Isolated lutropin deficiency']
NM_000264.3(PTCH1):c.2479A>G (p.Ser827Gly)	199476092	-	[]	['CGTTACYGAAACTCCTGTGTAGG']	['Gorlin syndrome', 'Holoprosencephaly 7', 'not specified', 'not provided']
NM_000021.3(PSEN1):c.415A>G (p.Met139Val)	63751037	PSEN1	[]	[]	['Alzheimer disease, type 3', 'not provided']
NM_000484.3(APP):c.2078A>G (p.Glu693Gly)	63751039	APP	[]	[]	['Alzheimer disease', 'Alzheimer disease, type 1', 'Cerebral amyloid angiopathy, APP-related', 'not provided']
NM_000117.2(EMD):c.450-2A>G	398123158	EMD	[]	['CGTTCCCYGAGGCAAAAGAGGGG']	['not provided']
RMRP:n.71A>G	199476103	RMRP	[]	['ACTTYCCCCTAGGCGGAAAGGGG', 'GACTTYCCCCTAGGCGGAAAGGG', 'GGACTTYCCCCTAGGCGGAAAGG']	['Metaphyseal chondrodysplasia, McKusick type', 'Metaphyseal dysplasia without hypotrichosis']
m.14495A>G	199476106	MT-ND6	[]	[]	['Leber optic atrophy']
m.11084A>G	199476113	MT-ND4	[]	[]	['Juvenile myopathy, encephalopathy, lactic acidosis AND stroke']
NM_000551.3(VHL):c.233A>G (p.Asn78Ser)	5030804	VHL	['TGCGAYTGCAGAAGATGACCTGG']	['GCGAYTGCAGAAGATGACCTGGG', 'TGCGAYTGCAGAAGATGACCTGG']	['Von Hippel-Lindau syndrome']

m.3397A>G	199476120	MT-ND1	[]	[]	['Alzheimer disease', 'Parkinson disease, late-onset']
m.4136A>G	199476121	MT-ND1	[]	[]	['Leber optic atrophy']
NM_003094.3(SNRPE):c.1A>G (p.Met1Val)	587776924	SNRPE	[]	[]	['Hypotrichosis 11']
NM_001310338.1(MGME1):c.743A>G (p.Tyr248Cys)	587776944	MGME1	[]	[]	['Mitochondrial DNA depletion syndrome 11']
NM_000249.3(MLH1):c.122A>G (p.Asp41Gly)	63751094	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_138425.3(C12orf57):c.1A>G (p.Met1Val)	587776954	C12orf57	[]	[]	['Temtamy syndrome', 'Seizures', 'Corpus callosum abnormalities', 'Colobomatous microphthalmia', 'Global developmental delay']
NM_000277.1(PAH):c.1169A>G (p.Glu390Gly)	5030856	PAH	[]	['CTCYCTGCCACGTAATACAGGGG', 'ACTCYCTGCCACGTAATACAGGG', 'AACTCYCTGCCACGTAATACAGG']	['Phenylketonuria', 'Hyperphenylalaninemia, non-pku', 'not provided']
NM_000277.1(PAH):c.1241A>G (p.Tyr414Cys)	5030860	PAH	[]	['GGGTCGYAGCGAACTGAGAAGGG', 'TGGGTCGYAGCGAACTGAGAAGG']	['Phenylketonuria', 'Hyperphenylalaninemia, non-pku', 'not provided']
NM_000155.3(GALT):c.308A>G (p.Gln103Arg)	367543252	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_207352.3(CYP4V2):c.1091-2A>G	199476183	CYP4V2	[]	[]	['Bietti crystalline corneoretinal dystrophy']
NM_000518.4(HBB):c.*111A>G	63751128	HBB	[]	[]	[]
NM_000155.3(GALT):c.857A>G (p.Tyr286Cys)	367543262	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_000155.3(GALT):c.854A>G (p.Lys285Arg)	367543263	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_207352.3(CYP4V2):c.761A>G (p.His254Arg)	199476193	CYP4V2	[]	[]	['Bietti crystalline corneoretinal dystrophy']
NM_000155.3(GALT):c.968A>G (p.Tyr323Cys)	367543267	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']

NM_001142519.1(FAM111A):c.1012A>G (p.Thr338Ala)	587777014	FAM111A	[]	[]	['Gracile bone dysplasia']
NM_000132.3(F8):c.1660A>G (p.Ser554Gly)	137852419	F8	['AACYAGAGTAATAGCGGGTCAGG']	['AACYAGAGTAATAGCGGGTCAGG']	['Hereditary factor VIII deficiency disease']
NM_020988.2(GNAO1):c.521A>G (p.Asp174Gly)	587777055	GNAO1	[]	['GGATGYCCTGCTCGGTGGGCTGG']	['Early infantile epileptic encephalopathy 17']
NM_000155.3(GALT):c.905-2A>G	398123187	GALT	[]	[]	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_015662.2(IFT172):c.4607T>C (p.Leu1536Pro)	587777080	IFT172	[]	[]	[]
NM_014754.2(PTDSS1):c.1058A>G (p.Gln353Arg)	587777088	PTDSS1	[]	[]	['Lenz-Majewski hyperostosis syndrome']
NM_003859.1(DPM1):c.742T>C (p.Ser248Pro)	587777114	-	[]	[]	['Congenital disorder of glycosylation type 1E']
NM_001018005.1(TPM1):c.742A>G (p.Lys248Glu)	199476319	TPM1	[]	[]	['Left ventricular noncompaction 9', 'not provided']
NM_004826.3(ECEL1):c.2278T>C (p.Cys760Arg)	587777129	ECEL1	[]	[]	['Arthrogryposis, distal, type 5d']
NM_014908.3(DOLK):c.2T>C (p.Met1Thr)	587777137	DOLK	[]	[]	['Congenital disorder of glycosylation type 1M']
NM_000350.2(ABCA4):c.4540-2A>G	61752435	ABCA4	[]	[]	['Stargardt disease 1', 'not provided']
NM_001128085.1(ASPA):c.433-2A>G	63751297	-	[]	[]	['Spongy degeneration of central nervous system']
NM_176787.4(PIGN):c.808T>C (p.Ser270Pro)	587777186	PIGN	[]	[]	['Multiple congenital anomalies-hypotonia-seizures syndrome 1']
NM_001165899.1(PDE4D):c.1850T>C (p.Ile617Thr)	587777188	PDE4D	['CTATAYTGTTCATCCCCTCTGGG', 'ACTATAYTGTTCATCCCCTCTGG']	['CTATAYTGTTCATCCCCTCTGGG', 'ACTATAYTGTTCATCCCCTCTGG']	['Acrodysostosis 2, with or without hormone resistance']
NM_005017.3(PCYT1A):c.571T>C (p.Phe191Leu)	587777195	PCYT1A	['GCATGYTTGCTCCAACACAGAGG']	['GCATGYTTGCTCCAACACAGAGG']	['Spondylometaphyseal dysplasia with cone-rod dystrophy']
NM_024301.4(FKRP):c.1A>G (p.Met1Val)	587777223	FKRP	[]	['CCGCAYGGGGCCGAAGTCTGGGG', 'GCCGCAYGGGGCCGAAGTCTGGG', 'AGCCGCAYGGGGCCGAAGTCTGG']	['Congenital muscular dystrophy-dystroglycanopathy with brain and eye anomalies type A5']

NM_198947.3(FAM111B):c.1879A>G (p.Arg627Gly)	587777237	FAM111B	[]	[]	['Poikiloderma, hereditary fibrosing, with tendon contractures, myopathy, and pulmonary fibrosis']
NM_003638.2(ITGA8):c.2982+2T>C	587777279	ITGA8	[]	[]	['Renal adysplasia']
NM_199189.2(MATR3):c.1864A>G (p.Thr622Ala)	587777301	MATR3	['CGGYTGAACTCTCAGTCTTCTGG']	['CGGYTGAACTCTCAGTCTTCTGG']	['Myopathy, distal, 2']
NM_001739.1(CA5A):c.697T>C (p.Ser233Pro)	587777316	CA5A	[]	[]	['Carbonic anhydrase VA deficiency, hyperammonemia due to']
NM_005051.2(QARS):c.169T>C (p.Tyr57His)	587777333	QARS	[]	[]	['Microcephaly, progressive, with seizures and cerebral and cerebellar atrophy']
NM_002234.3(KCNA5):c.143A>G (p.Glu48Gly)	587777336	KCNA5	[]	[]	['Atrial fibrillation, familial, 7']
NM_021803.3(IL21):c.146T>C (p.Leu49Pro)	587777338	IL21	[]	[]	['Common variable immunodeficiency 11']
NM_000132.3(F8):c.6794A>G (p.Gln2265Arg)	137852470	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_178014.3(TUBB):c.895A>G (p.Met299Val)	587777355	TUBB	[]	[]	['Cortical dysplasia, complex, with other brain malformations 6']
NM_005957.4(MTHFR):c.1969T>C (p.Ter657Arg)	768434408	MTHFR	[]	[]	['Homocysteinemia due to MTHFR deficiency']
NM_005359.5(SMAD4):c.425-6A>G	377767327	SMAD4	[]	[]	['Juvenile polyposis syndrome']
NM_022068.3(PIEZO2):c.8215T>C (p.Ser2739Pro)	587777454	PIEZO2	[]	[]	['Oculomelic amyoplasia']
NM_003108.3(SOX11):c.347A>G (p.Tyr116Cys)	587777479	SOX11	[]	['GTACTTGYAGTCGGGGTAGTCGG']	['Mental retardation, autosomal dominant 27']
NM_021072.3(HCN1):c.814T>C (p.Ser272Pro)	587777493	HCN1	[]	[]	['Epileptic encephalopathy, early infantile, 24']
NM_020435.3(GJC2):c.-170A>G	587777496	GJC2	[]	['TTGYTCCCCCCTCGGCCTCAGGG', 'ATTGYTCCCCCCTCGGCCTCAGG']	['Leukodystrophy, hypomyelinating, 2']
NM_022552.4(DNMT3A):c.1943T>C (p.Leu648Pro)	587777507	DNMT3A	[]	['CTCCYGGTGCTGAAGGACTTGGG', 'GCTCCYGGTGCTGAAGGACTTGG']	['Tatton-Brown-rahman syndrome']
NM_022552.4(DNMT3A):c.2705T>C	587777510	DNMT3A	[]	[]	['Tatton-Brown-rahman syndrome']

(p.Phe902Ser)					
NM_000223.3(KRT12):c.403A>G (p.Arg135Gly)	58410481	KRT12	[]	[]	['Meesman corneal dystrophy', 'not provided']
NM_000232.4(SGCB):c.1A>G (p.Met1Val)	398123262	SGCB	[]	[]	['Limb-girdle muscular dystrophy, type 2E', 'not provided']
NM_020630.4(RET):c.2342A>G (p.Gln781Arg)	377767416	RET	[]	[]	['MEN2 phenotype: Unclassified']
NM_018400.3(SCN3B):c.482T>C (p.Met161Thr)	587777557	SCN3B	[]	['AATCAYGATGTACATCCTTCTGG']	['Atrial fibrillation, familial, 16']
NM_001030001.2(RPS29):c.149T>C (p.Ile50Thr)	587777569	RPS29	[]	['GATAYCGGTTTCATTAAGGTAGG']	['Diamond-Blackfan anemia 13']
NM_177550.4(SLC13A5):c.1463T>C (p.Leu488Pro)	587777578	SLC13A5	[]	[]	['Epileptic encephalopathy, early infantile, 25']
NM_002880.3(RAF1):c.1808T>C (p.Leu603Pro)	587777586	RAF1	[]	[]	['Cardiomyopathy, dilated, 1NN']
NM_025150.4(TARS2):c.695+3A>G	587777594	TARS2	[]	[]	['Combined oxidative phosphorylation deficiency 21']
NM_001759.3(CCND2):c.838A>G (p.Thr280Ala)	587777618	CCND2	[]	[]	['Megalencephaly-polymicrogyria-polydactyly-hydrocephalus syndrome 3']
NM_153334.6(SCARF2):c.190T>C (p.Cys64Arg)	587777657	SCARF2	[]	['CCACGYGCTGCGCTGGCTGGAGG']	['Marden Walker like syndrome']
NM_005726.5(TSFM):c.57+4A>G	587777689	TSFM	[]	['ACTTCYCACCGGGTAGCTCCCGG']	['Combined oxidative phosphorylation deficiency 3']
NM_000255.3(MUT):c.329A>G (p.Tyr110Cys)	796052005	MUT	[]	['GCAYACTGGCGGATGGTCCAGGG', 'AGCAYACTGGCGGATGGTCCAGG']	['not provided']
NM_021870.2(FGG):c.1210T>C (p.Ser404Pro)	587777720	FGG	[]	[]	['Hypodysfibrinogenemia']
NM_017617.3(NOTCH1):c.1285T>C (p.Cys429Arg)	587777736	NOTCH1	['GGCAAGYGCATCAACACGCTGGG']	['GGCAAGYGCATCAACACGCTGGG', 'GGGCAAGYGCATCAACACGCTGG']	['Adams-Oliver syndrome 1', 'Adams-Oliver syndrome 5']
NM_014946.3(SPAST):c.1688-2A>G	587777752	SPAST	['TTCYGTAAAACATAAAAGTCAGG']	['TTCYGTAAAACATAAAAGTCAGG']	['Spastic paraplegia 4, autosomal dominant']
NM_014946.3(SPAST):c.1245+4A>G	587777755	SPAST	[]	[]	['Spastic paraplegia 4, autosomal dominant']
NM_014946.3(SPAST):c.1216A>G	587777757	SPAST	[]	[]	['Spastic paraplegia 4, autosomal

(p.Ile406Val)					dominant']
NM_144596.3(TTC8): c.115-2A>G	587777809	TTC8	[]	['GTTCCYGGAAA GCATTAAGAAGG']	['Retinitis pigmentosa 51']
NM_170784.2(MKKS) :c.110A>G (p.Tyr37Cys)	74315396	MKKS	[]	[]	['Bardet-Biedl syndrome 6', 'McKusick Kaufman syndrome']
NM_000252.2(MTM1) :c.566A>G (p.Asn189Ser)	132630302	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1) :c.1190A>G (p.Tyr397Cys)	132630303	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_152384.2(BBS5): c.522+3A>G	587777828	BBS5	[]	[]	['Bardet-Biedl syndrome 5']
NM_001205019.1(GK):c.880A>G (p.Asn294Asp)	132630331	GK	[]	[]	['Deficiency of glycerol kinase']
NM_000166.5(GJB1): c.580A>G (p.Met194Val)	587777878	GJB1	[]	['TAGCAYGAAGA CGGTGAAGACGG']	['X-linked hereditary motor and sensory neuropathy']
NM_000311.3(PRNP): c.547A>G (p.Thr183Ala)	74315411	PRNP	[]	[]	['Genetic prion diseases', 'Spongiform encephalopathy with neuropsychiatric features']
NM_144773.2(PROK R2):c.629A>G (p.Gln210Arg)	74315417	PROKR2	[]	[]	['Kallmann syndrome 3']
NM_000531.5(OTC):c .919A>G (p.Lys307Glu)	796052013	OTC	[]	[]	['not provided']
NM_001029871.3(RS PO4):c.194A>G (p.Gln65Arg)	74315420	RSPO4	[]	['CGTACYGGCGG ATGCCTTCCCGG']	['Anonychia']
NM_004333.4(BRAF): c.770A>G (p.Gln257Arg)	180177035	BRAF	[]	[]	['Noonan syndrome 7', 'Cardiofaciocutaneo us syndrome', 'Rasopathy', 'not provided']
NM_004333.4(BRAF): c.1495A>G (p.Lys499Glu)	180177037	BRAF	[]	[]	['Cardiofaciocutaneo us syndrome', 'Rasopathy']
NM_198056.2(SCN5A):c.5297T>A (p.Met1766Lys)	752476527	SCN5A	[]	[]	['not provided']
NM_000030.2(AGXT) :c.248A>G (p.His83Arg)	180177186	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_000030.2(AGXT) :c.424-2A>G (p.Gly_142Gln145del)	180177219	AGXT	[]	['AGGCCCYGAGG AAGCAGGGACGG']	['Primary hyperoxaluria, type I']
NM_198578.3(LRRK2):c.5096A>G (p.Tyr1699Cys)	35801418	LRRK2	[]	[]	['Parkinson disease 8, autosomal dominant']
NM_002693.2(POLG):	367610201	POLG	[]	['CTCAYGGCACTT	['not provided']

c.1808T>C (p.Met603Thr)				ACCTGGGATGG']	
NM_000030.2(AGXT):c.596-2A>G	180177245	AGXT	[]	[]	['Primary hyperoxaluria, type I']
NM_020223.3(FAM20C):c.1364-2A>G	796051853	FAM20C	[]	[]	['Raine syndrome']
NM_012203.1(GRHPR):c.84-2A>G	180177319	GRHPR	[]	['TCACAGCYGCGGGGAAAGGGAGG']	['Primary hyperoxaluria, type II']
NM_006017.2(PROM1):c.2077-521A>G	796051882	PROM1	[]	[]	['Cone-rod dystrophy 2']
NM_012203.1(GRHPR):c.934A>G (p.Asn312Asp)	180177324	GRHPR	['CAAGTYGTTAGCTGCCAACAAGG']	['CAAGTYGTTAGCTGCCAACAAGG']	['Primary hyperoxaluria, type II']
NM_000016.5(ACADM):c.329A>G (p.Glu110Gly)	796051900	ACADM	[]	[]	['not provided']
NM_004453.3(ETFDH):c.929A>G (p.Tyr310Cys)	796051958	ETFDH	[]	[]	['not provided']
NM_000255.3(MUT):c.1885A>G (p.Arg629Gly)	796052004	MUT	[]	[]	['not provided']
NM_012434.4(SLC17A5):c.548A>G (p.His183Arg)	119491109	SLC17A5	[]	[]	['Sialic acid storage disease, severe infantile type']
NM_000328.2(RPGR):c.155-2A>G	62638632	RPGR	[]	[]	['Retinitis pigmentosa 15', 'not provided']
NM_005557.3(KRT16):c.373A>G (p.Asn125Asp)	58608173	KRT16	[]	[]	['Pachyonychia congenita, type 1', 'not provided']
NM_000532.4(PCCB):c.655-2A>G	796052020	PCCB	[]	[]	['not provided']
NM_000030.2(AGXT):c.777-2A>G	796052068	AGXT	[]	['GGTACCYGGAAGACACGAGGGGG', 'TGGTACCYGGAAGACACGAGGGG']	['Primary hyperoxaluria, type I']
NM_000121.3(EPOR):c.1460A>G (p.Asn487Ser)	62638745	EPOR	['AGGGYTGGAGTAGGGGCCATCGG']	['AGGGYTGGAGTAGGGGCCATCGG']	['Acute myeloid leukemia, M6 type', 'Familial erythrocytosis, 1']
NM_000552.3(VWF):c.1583A>G (p.Asn528Ser)	61754010	VWF	[]	['TGCCAYTGTAATTCCCACACAGG']	['von Willebrand disease, type 2a', 'not provided']
NM_001918.3(DBT):c.1017_1018insNC_000001.11:g.100207187_100207312	796052135	DBT	[]	[]	['Intermediate maple syrup urine disease type 2']
NM_001243473.1(B9D1):c.400+2T>C	143149764	B9D1	[]	[]	['Meckel syndrome, type 9', 'not provided']
NM_001165963.1(SCN1A):c.4766T>G (p.Val1589Gly)	764037830	-	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_000321.2(RB1):c.	587778866	RB1	[]	['ATTYCAATGGCT	['Retinoblastoma']

1927A>G (p.Lys643Glu)				TCTGGGTCTGG']	
NM_006331.7(EMG1):c.257A>G (p.Asp86Gly)	74435397	EMG1	[]	['ATAYCTGGCCGCGCTTCCCCAGG']	['Bowen-Conradi syndrome']
NM_000249.3(MLH1):c.113A>G (p.Asn38Ser)	587778888	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_017777.3(MKS1):c.1382A>G (p.Tyr461Cys)	730882120	MKS1	[]	[]	['Bardet-Biedl syndrome 13']
NM_000261.1(MYOC):c.1010A>G (p.Gln337Arg)	74315335	MYOC	[]	[]	['Primary open angle glaucoma juvenile onset 1']
NM_152515.4(CKAP2L):c.2T>C (p.Met1Thr)	548949031	CKAP2L	[]	[]	['Filippi syndrome']
NM_000156.5(GAMT):c.1A>G (p.Met1Val)	796052527	GAMT	[]	['CGCTCAYGCTGCAGGCTGGACGG']	['not provided']
NM_000833.4(GRIN2A):c.1930A>G (p.Ser644Gly)	796052544	GRIN2A	[]	[]	['not provided']
NM_000144.4(FXN):c.385-2A>G	140987490	FXN	[]	[]	['Friedreich ataxia']
NM_172107.2(KCNQ2):c.297-2A>G	796052615	KCNQ2	[]	[]	['not provided']
NM_172107.2(KCNQ2):c.611A>G (p.Gln204Arg)	796052624	KCNQ2	[]	[]	['not provided']
NM_172107.2(KCNQ2):c.848A>G (p.Lys283Arg)	796052637	KCNQ2	[]	['GTACYTGTCCCCGTAGCCAATGG']	['not provided']
NM_052859.3(RFT1):c.887T>A (p.Ile296Lys)	772820136	RFT1	[]	[]	['Congenital disorder of glycosylation type 1N']
NM_000553.4(WRN):c.561A>G (p.Lys187=)	775802030	WRN	[]	[]	['Werner syndrome']
NM_194277.2(FRMD7):c.556A>G (p.Met186Val)	786205896	FRMD7	[]	[]	['Infantile nystagmus, X-linked']
NM_000535.5(PMS2):c.989-2A>G	587779347	PMS2	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms', 'Hereditary cancer-predisposing syndrome']
NM_000203.4(IDUA):c.1874A>G (p.Tyr625Cys)	587779401	IDUA	[]	[]	['Hurler syndrome']
NM_001105243.1(PCDH19):c.1019A>G (p.Asn340Ser)	796052839	PCDH19	[]	[]	['not provided']
NM_002693.2(POLG):c.2840A>G (p.Lys947Arg)	796052891	POLG	[]	[]	['not provided']
NM_032228.5(FAR1):c.1094A>G	724159963	FAR1	[]	['GATAYCATACAGGAATGCTGGGG',	['Peroxisomal fatty acyl-coa reductase 1

(p.Asp365Gly)				'AGATAYCATACAGGAATGCTGGG', 'TAGATAYCATACAGGAATGCTGG']	disorder']
NM_000090.3(COL3A1):c.2284-2A>G (p.Gly762_Lys779del)	587779558	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_014305.3(TGDS):c.269A>G (p.Glu90Gly)	724160004	TGDS	[]	[]	['Catel Manzke syndrome']
NM_014305.3(TGDS):c.892A>G (p.Asn298Asp)	724160005	TGDS	[]	[]	['Catel Manzke syndrome']
NM_000090.3(COL3A1):c.997-2A>G (p.Gly333_Lys350del+)	587779602	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_002185.3(IL7R):c.197T>C (p.Ile66Thr)	1494558	IL7R	[]	[]	['Severe combined immunodeficiency, autosomal recessive, T cell-negative, B cell-positive, NK cell-positive', 'not specified']
NM_000277.1(PAH):c.974A>G (p.Tyr325Cys)	62508578	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_000090.3(COL3A1):c.997-10A>G (p.Pro332_Gly333insFFQ)	587779670	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_000090.3(COL3A1):c.3202-2A>G (p.Gly1068_Pro1085del)	587779682	COL3A1	[]	[]	['Ehlers-Danlos syndrome, type 4']
NM_000090.3(COL3A1):c.1762-2A>G (p.Gly588_Gln605del)	587779722	COL3A1	[]	['CACCCYAAAGAAGAAGTGGTCGG']	['Ehlers-Danlos syndrome, type 4']
NM_021007.2(SCN2A):c.4036A>G (p.Ile1346Val)	796053135	SCN2A	[]	[]	['not provided']
m.13637A>G	200855215	MT-ND5	[]	[]	['Leber optic atrophy']
NM_021007.2(SCN2A):c.387-2A>G	796053169	SCN2A	['AATAAAGYAGAATATCGTCAAGG']	['AATAAAGYAGAATATCGTCAAGG']	['not provided']
NM_021007.2(SCN2A):c.851A>G (p.Asp284Gly)	796053173	SCN2A	[]	[]	['not provided']
NM_006516.2(SLC2A1):c.848A>G (p.Gln283Arg)	796053251	SLC2A1	[]	[]	['not provided']
NM_006516.2(SLC2A1):c.19-2A>G	796053272	SLC2A1	[]	[]	['not provided']
NM_000136.2(FANCC):c.-78-2A>G	587779898	FANCC	[]	[]	['Hereditary cancer-predisposing syndrome']
m.8296A>G	118192102	MT-TK	[]	['TTTACAGYGGGC	['Diabetes-deafness

				TCTAGAGGGGG']	syndrome maternally transmitted']
NM_005360.4(MAF):c.172A>G (p.Thr58Ala)	727502767	MAF	[]	[]	['Cataracts, congenital, with sensorineural deafness, down syndrome-like facial appearance, short stature, and mental retardation']
NM_001145901.1(SARS2):c.1175A>G (p.Asp392Gly)	727502784	SARS2	[]	[]	['Hyperuricemia, pulmonary hypertension, renal failure, and alkalosis']
NM_001077494.3(NFKB2):c.2594A>G (p.Asp865Gly)	727502787	NFKB2	[]	['CTGYCTTCCTTCACCTCTGCTGG']	['Common variable immunodeficiency 10']
NM_002238.3(KCNH1):c.1399A>G (p.Ile467Val)	727502819	KCNH1	[]	[]	['Zimmermann-Laband syndrome', 'Temple-Baraitser syndrome']
NM_172362.2(KCNH1):c.1508A>G (p.Gln503Arg)	727502821	KCNH1	[]	[]	['Temple-Baraitser syndrome']
NM_000546.5(TP53):c.701A>G (p.Tyr234Cys)	587780073	TP53	[]	[]	['Li-Fraumeni syndrome', 'Hereditary cancer-predisposing syndrome']
NM_003060.3(SLC22A5):c.694A>C (p.Thr232Pro)	188698686	SLC22A5	[]	[]	['not provided']
NM_000540.2(RYR1):c.14591A>G (p.Tyr4864Cys)	118192146	RYR1	[]	[]	['Central core disease', 'not provided']
NM_058216.2(RAD51C):c.706-2A>G	587780259	RAD51C	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000501.3(ELN):c.800-2A>G	727503027	ELN	[]	[]	['Supravalvar aortic stenosis', 'not provided']
NM_000117.2(EMD):c.266-2A>G	727503036	EMD	[]	['AGCCYTGGGAAGGGGGGCAGCGG']	['Emery-Dreifuss muscular dystrophy 1, X-linked']
NM_003242.5(TGFBR2):c.1273A>G (p.Met425Val)	104893817	TGFBR2	[]	[]	['Loeys-Dietz syndrome 2']
NM_153638.2(PANK2):c.700A>G (p.Thr234Ala)	137852965	PANK2	[]	[]	[]
NM_005861.3(STUB1):c.194A>G (p.Asn65Ser)	690016544	STUB1	[]	['GGCCCGGYTGGTGTAATACACGG']	['Spinocerebellar ataxia, autosomal recessive 16']
NM_005360.4(MAF):c.890A>G (p.Lys297Arg)	121917736	MAF	[]	[]	['Cataract, pulverulent, juvenile-onset']
NM_005211.3(CSF1R	690016554	CSF1R	[]	['GTATCYGGGAG	['Hereditary diffuse

):c.2655-2A>G				ATAGGACAGAGG']	leukoencephalopathy with spheroids']
NM_003361.3(UMOD):c.383A>G (p.Asn128Ser)	121917770	UMOD	['CACAYTGACACATGTGGCCAGGG']	['CACAYTGACACATGTGGCCAGGG', 'CCACAYTGACACATGTGGCCAGG']	['Familial juvenile gout']
NM_000256.3(MYBPC3):c.2234A>G (p.Asp745Gly)	727503190	MYBPC3	[]	[]	['Familial hypertrophic cardiomyopathy 4', 'Familial hypertrophic cardiomyopathy 1', 'not specified']
NM_172107.2(KCNQ2):c.1A>G (p.Met1Val)	118192185	KCNQ2	[]	['GCACCAYGGTGCCTGGCGGGAGG']	['Benign familial neonatal seizures 1']
NM_000256.3(MYBPC3):c.1213A>G (p.Met405Val)	727503207	MYBPC3	[]	[]	['Cardiomyopathy', 'not specified']
NM_000021.3(PSEN1):c.998A>G (p.Asp333Gly)	121917809	PSEN1	[]	[]	['Primary dilated cardiomyopathy', 'Cardiomyopathy, dilated, 1u', 'Heart failure']
NM_021954.3(GJA3):c.188A>G (p.Asn63Ser)	121917823	GJA3	[]	[]	['Zonular pulverulent cataract 3']
NM_000322.4(PRPH2):c.422A>G (p.Tyr141Cys)	61755781	PRPH2	[]	[]	['Macular dystrophy, vitelliform, adult-onset', 'Patterned dystrophy of retinal pigment epithelium', 'not provided']
NM_007035.3(KERA):c.740A>G (p.Asn247Ser)	121917858	KERA	[]	[]	['Cornea plana 2']
NM_002181.3(IHH):c.284A>G (p.Glu95Gly)	121917859	IHH	[]	[]	['Brachydactyly type A1']
NM_000257.3(MYH7):c.1157A>G (p.Tyr386Cys)	727503269	MYH7	[]	[]	['Primary familial hypertrophic cardiomyopathy']
NM_000097.5(CPOX):c.1210A>G (p.Lys404Glu)	121917868	CPOX	[]	[]	['Harderoporphyria']
NM_012064.3(MIP):c.401A>G (p.Glu134Gly)	121917869	MIP	[]	['AGATCYCCACTGTGGTTGCCTGG']	['Cataract 15, multiple types']
NM_025243.3(SLC19A3):c.1264A>G (p.Thr422Ala)	121917884	SLC19A3	[]	[]	['Basal ganglia disease, biotin-responsive']
NM_000373.3(UMPS):c.286A>G (p.Arg96Gly)	121917890	UMPS	[]	[]	['Orotic aciduria']
NM_000536.3(RAG2):c.115A>G (p.Arg39Gly)	121917897	RAG2	[]	[]	['Histiocytic medullary reticulosis']
NM_130838.1(UBE3A):c.1694-2A>G	587780579	UBE3A	[]	[]	['Angelman syndrome']
NM_016335.4(PROD	2904551	PRODH	[]	[]	['Proline

H):c.1322T>C (p.Leu441Pro)					dehydrogenase deficiency', 'Schizophrenia 4']
NM_006920.4(SCN1A):c.4352A>G (p.Tyr1451Cys)	121917962	-	[]	[]	['Severe myoclonic epilepsy in infancy', 'not provided']
NM_006920.4(SCN1A):c.1876A>G (p.Ser626Gly)	121917990	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy', 'Generalized epilepsy']
NM_000478.4(ALPL):c.1250A>G (p.Asn417Ser)	121918014	ALPL	[]	['AGGCCCAYTGCCATACAGGATGG']	['Infantile hypophosphatasia']
NM_000174.4(GP9):c.110A>G (p.Asp37Gly)	121918036	GP9	[]	['GCAGYCCACCCACAGCCCCATGG']	['Bernard-Soulier syndrome type C']
NM_002693.2(POLG):c.2591A>G (p.Asn864Ser)	121918050	POLG	[]	[]	['Mitochondrial DNA depletion syndrome 4B, MNGIE type']
NM_000374.4(UROD):c.932A>G (p.Tyr311Cys)	121918061	UROD	[]	[]	['Hepatoerythropoietic porphyria']
NM_000217.2(KCNA1):c.763A>G (p.Asn255Asp)	121918067	KCNA1	[]	[]	[]
NM_000371.3(TTR):c.238A>G (p.Thr80Ala)	121918070	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis', 'Cardiomyopathy']
NM_000371.3(TTR):c.401A>G (p.Tyr134Cys)	121918075	TTR	['GGAGYAGGGGCTCAGCAGGGCGG', 'ATAGGAGYAGGGGCTCAGCAGGG']	['GGAGYAGGGGCTCAGCAGGGCGG', 'ATAGGAGYAGGGGCTCAGCAGGG']	['Amyloidogenic transthyretin amyloidosis']
NM_000371.3(TTR):c.205A>G (p.Thr69Ala)	121918081	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis']
NM_000371.3(TTR):c.379A>G (p.Ile127Val)	121918089	TTR	[]	['CGGCAAYGGTGTAGCGGCGGGGG', 'GCGGCAAYGGTGTAGCGGCGGGG']	['Amyloidogenic transthyretin amyloidosis']
NM_000371.3(TTR):c.113A>G (p.Asp38Gly)	121918098	TTR	[]	[]	['Amyloidogenic transthyretin amyloidosis', 'AMYLOIDOSIS, LEPTOMENINGEAL, TRANSTHYRETIN-RELATED']
NM_000823.3(GHRHR):c.985A>G (p.Lys329Glu)	121918121	GHRHR	[]	['CGACTYGGAGAGACGCCTGCAGG']	['Isolated growth hormone deficiency type 1B']
NM_000275.2(OCA2):c.1465A>G (p.Asn489Asp)	121918170	OCA2	['GACATYTGGAGGGTCCCCGATGG']	['GACATYTGGAGGGTCCCCGATGG']	['Tyrosinase-positive oculocutaneous albinism']

NM_000181.3(GUSB):c.1484A>G (p.Tyr495Cys)	121918178	GUSB	[]	[]	['Mucopolysaccharidosis type VII']
NM_018122.4(DARS2):c.133A>G (p.Ser45Gly)	121918209	DARS2	[]	[]	['Leukoencephalopathy with Brainstem and Spinal Cord Involvement and Lactate Elevation']
NM_015697.7(COQ2):c.890A>G (p.Tyr297Cys)	121918230	COQ2	[]	[]	['Coenzyme Q10 deficiency, primary 1']
NM_015697.7(COQ2):c.683A>G (p.Asn228Ser)	121918232	COQ2	[]	[]	['Coenzyme Q10 deficiency, primary 1']
NM_015384.4(NIPBL):c.7289A>G (p.Tyr2430Cys)	121918265	NIPBL	[]	[]	['Cornelia de Lange syndrome 1']
NM_004183.3(BEST1):c.707A>G (p.Tyr236Cys)	121918291	BEST1	[]	[]	['Vitreoretinochoroidopathy dominant']
NM_014362.3(HIBCH):c.365A>G (p.Tyr122Cys)	121918329	HIBCH	[]	[]	['Beta-hydroxyisobutyryl-CoA deacylase deficiency']
NM_015335.4(MED13L):c.6068A>G (p.Asp2023Gly)	121918333	MED13L	[]	['ATATCAYCTAGAGGGAAGGGGGG', 'CATATCAYCTAGAGGGAAGGGGG']	['Transposition of great arteries']
NM_015040.3(PIKFYVE):c.3308A>G (p.Lys1103Arg)	121918336	PIKFYVE	[]	[]	['Fleck corneal dystrophy']
NM_006306.3(SMC1A):c.2974-2A>G	727503774	SMC1A	[]	[]	['Congenital muscular hypertrophy-cerebral syndrome']
NM_002633.2(PGM1):c.343A>G (p.Thr115Ala)	121918371	PGM1	[]	[]	['Congenital disorder of glycosylation type 1t']
NM_000040.1(APOC3):c.280A>G (p.Thr94Ala)	121918381	APOC3	['CTGAAGYTGGTCTGACCTCAGGG', 'GCTGAAGYTGGTCTGACCTCAGG']	['CTGAAGYTGGTCTGACCTCAGGG', 'GCTGAAGYTGGTCTGACCTCAGG']	[]
NM_000040.1(APOC3):c.232A>G (p.Lys78Glu)	121918382	APOC3	[]	[]	['Hyperalphalipoproteinemia 2']
NM_001146040.1(GLRA1):c.910A>G (p.Lys304Glu)	121918412	GLRA1	[]	[]	['Hyperekplexia hereditary']
NM_000171.3(GLRA1):c.523A>G (p.Met175Val)	121918414	GLRA1	[]	[]	['Hyperekplexia hereditary']
NM_021957.3(GYS2):c.116A>G (p.Asn39Ser)	121918423	GYS2	[]	[]	['Hypoglycemia with deficiency of glycogen synthetase in the liver']

NM_002834.3(PTPN11):c.188A>G (p.Tyr63Cys)	121918459	PTPN11	[]	[]	['Noonan syndrome', 'Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_013382.5(POMT2):c.1726-2A>G	727503873	POMT2	[]	[]	['not provided']
NM_002834.3(PTPN11):c.236A>G (p.Gln79Arg)	121918466	PTPN11	[]	[]	['Noonan syndrome', 'Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_000313.3(PROS1):c.773A>G (p.Asn258Ser)	121918473	PROS1	[]	[]	['Protein S deficiency']
NM_000313.3(PROS1):c.586A>G (p.Lys196Glu)	121918474	PROS1	[]	[]	['Protein S deficiency']
NM_000141.4(FGFR2):c.983A>G (p.Tyr328Cys)	121918493	FGFR2	[]	[]	['Crouzon syndrome']
NM_000141.4(FGFR2):c.874A>G (p.Lys292Glu)	121918500	FGFR2	['TGCTYGATCCACTGGATGTGGGG']	['TGCTYGATCCACTGGATGTGGGG', 'GTGCTYGATCCACTGGATGTGGG', 'CGTGCTYGATCCACTGGATGTGG']	['Crouzon syndrome']
NM_000141.4(FGFR2):c.1576A>G (p.Lys526Glu)	121918507	FGFR2	[]	[]	['Crouzon syndrome', 'Scaphocephaly, maxillary retrusion, and mental retardation']
NM_002739.3(PRKCG):c.380A>G (p.Gln127Arg)	121918515	PRKCG	[]	[]	['Spinocerebellar ataxia 14']
NM_002739.3(PRKCG):c.1081A>G (p.Ser361Gly)	121918517	PRKCG	[]	[]	['Spinocerebellar ataxia 14']
NM_000098.2(CPT2):c.359A>G (p.Tyr120Cys)	121918528	CPT2	['GATAGGYACATATCAAACCAGGG', 'AGATAGGYACATATCAAACCAGG']	['GATAGGYACATATCAAACCAGGG', 'AGATAGGYACATATCAAACCAGG']	['Carnitine palmitoyltransferase II deficiency, infantile']
NM_005587.2(MEF2A):c.788A>G (p.Asn263Ser)	121918530	MEF2A	['CCAAGAYTACCACCACCTGGTGG']	['AGAYTACCACCACCTGGTGGAGG', 'CCAAGAYTACCACCACCTGGTGG']	[]
NM_006204.3(PDE6C):c.1363A>G (p.Met455Val)	121918539	PDE6C	[]	[]	['Achromatopsia 5']
NM_017654.3(SAMD9):c.4483A>G (p.Lys1495Glu)	121918554	SAMD9	[]	[]	['Tumoral calcinosis, familial, normophosphatemic']
NM_000191.2(HMGCL):c.698A>G (p.His233Arg)	727503963	HMGCL	[]	[]	['not provided']

NM_020166.4(MCCC1):c.137-2A>G	727504006	MCCC1	[]	[]	['3 Methylcrotonyl-CoA carboxylase 1 deficiency', 'not provided']
NM_001035.2(RYR2):c.12602A>G (p.Gln4201Arg)	121918605	RYR2	[]	['CGCCAGCYGCATTTCAAAGATGG']	['Catecholaminergic polymorphic ventricular tachycardia']
NM_002764.3(PRPS1):c.343A>G (p.Met115Val)	587781262	PRPS1	[]	['TAGCAYATTTGCAACAAGCTTGG']	['Charcot-Marie-Tooth disease, X-linked recessive, type 5', 'Deafness, high-frequency sensorineural, X-linked']
NM_001161766.1(AHCY):c.344A>G (p.Tyr115Cys)	121918608	AHCY	[]	['GCGGGYACTTGGTGTGGATGAGG']	['Hypermethioninemia with s-adenosylhomocysteine hydrolase deficiency']
NM_000702.3(ATP1A2):c.1033A>G (p.Thr345Ala)	121918613	ATP1A2	[]	['CTGYCAGGGTCAGGCACACCTGG']	['Familial hemiplegic migraine type 2']
NM_003126.2(SPTA1):c.143A>G (p.Lys48Arg)	121918644	SPTA1	[]	[]	['Hereditary pyropoikilocytosis']
NM_001024858.2(SPTB):c.1A>G (p.Met1Val)	121918651	SPTB	[]	[]	['Elliptocytosis 3']
NM_000899.4(KITLG):c.107A>G (p.Asn36Ser)	121918653	KITLG	[]	[]	['Familial progressive hyperpigmentation with or without hypopigmentation']
NM_198253.2(TERT):c.2315A>G (p.Tyr772Cys)	121918663	TERT	[]	[]	['Aplastic anemia', 'PULMONARY FIBROSIS AND/OR BONE MARROW FAILURE, TELOMERE-RELATED, 1']
NM_001063.3(TF):c.1936A>G (p.Lys646Glu)	121918678	TF	[]	[]	[]
NM_000535.5(PMS2):c.904-2A>G	587781339	PMS2	[]	['GCAGACCYGCACAAAATACAAGG']	['Hereditary cancer-predisposing syndrome']
NM_001128177.1(THRB):c.1324A>G (p.Met442Val)	121918691	THRB	[]	['CTTCAYGTGCAGGAAGCGGCTGG']	['Thyroid hormone resistance, generalized, autosomal dominant']
NM_001128177.1(THRB):c.1327A>G (p.Lys443Glu)	121918692	THRB	[]	['CCACCTYCATGTGCAGGAAGCGG']	['Thyroid hormone resistance, generalized, autosomal dominant']
NM_001128177.1(THRB):c.1009A>G	121918709	THRB	[]	[]	['Thyroid hormone resistance, selective

(p.Thr337Ala)					pituitary']
NM_004612.3(TGFBR1):c.1199A>G (p.Asp400Gly)	121918711	TGFBR1	['ATAGATGYCAGCACGTTTGAAGG']	['ATAGATGYCAGCACGTTTGAAGG']	['Loeys-Dietz syndrome 1']
NM_000359.2(TGM1):c.1469A>G (p.Asp490Gly)	121918724	TGM1	[]	[]	['Autosomal recessive congenital ichthyosis 1']
NM_000257.3(MYH7):c.1727A>G (p.His576Arg)	727504238	MYH7	[]	[]	['Familial hypertrophic cardiomyopathy 1', 'Cardiomyopathy', 'not specified']
NM_000257.3(MYH7):c.1954A>G (p.Arg652Gly)	727504239	MYH7	[]	[]	['Primary familial hypertrophic cardiomyopathy', 'Familial hypertrophic cardiomyopathy 1']
NM_000257.3(MYH7):c.1496A>G (p.Glu499Gly)	727504270	MYH7	[]	[]	['Cardiomyopathy', 'not specified']
NM_000257.3(MYH7):c.2539A>G (p.Lys847Glu)	727504310	MYH7	[]	[]	['Familial hypertrophic cardiomyopathy 1', 'Cardiomyopathy', 'not specified']
NM_000256.3(MYBPC3):c.2906-2A>G	727504333	MYBPC3	[]	['CCGTTCYGTGGGTATAGAGTGGG', 'GCCGTTCYGTGGGTATAGAGTGG']	['Familial hypertrophic cardiomyopathy 4']
NM_001128425.1(MUTYH):c.1187-2A>G	587781628	MUTYH	['ACCYGAGAGGGAGGGCAGCCAGG']	['ACCYGAGAGGGAGGGCAGCCAGG']	['Hereditary cancer-predisposing syndrome', 'Carcinoma of colon']
NM_005188.3(CBL):c.1228-2A>G	727504426	CBL	[]	[]	['Juvenile myelomonocytic leukemia', 'Noonan syndrome-like disorder with or without juvenile myelomonocytic leukemia', 'Rasopathy']
NM_000501.3(ELN):c.890-2A>G	727504434	ELN	['GCCYGAAAACACAGCCACAGAGG']	['GCCYGAAAACACAGCCACAGAGG']	['Supravalvar aortic stenosis']
NM_001165963.1(SCN1A):c.2877T>A (p.Cys959Ter)	775214722	SCN1A	[]	[]	['not provided']
NM_000833.4(GRIN2A):c.2449A>G (p.Met817Val)	796052551	GRIN2A	['CCAYGTTGTCAATGTCCAGCTGG']	['CCAYGTTGTCAATGTCCAGCTGG']	['not provided']
NM_000314.6(PTEN):c.493-2A>G	587781784	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']

NM_000498.3(CYP11B2):c.1157T>C (p.Val386Ala)	61757294	-	[]	[]	['Corticosterone methyloxidase type 2 deficiency', 'Corticosterone methyloxidase type 1 deficiency']
NM_006204.3(PDE6C):c.1483-2A>G	786200910	PDE6C	[]	['CTTTCYGTTGAAATAAGGATGGG', 'TCTTTCYGTTGAAATAAGGATGG']	['Achromatopsia 5']
NM_003588.3(CUL4B):c.901-2A>G	786200913	CUL4B	[]	[]	['Syndromic X-linked mental retardation, Cabezas type']
NM_000397.3(CYBB):c.302A>G (p.His101Arg)	137854591	CYBB	[]	[]	['Granulomatous disease, chronic, X-linked, variant', 'not provided']
NM_000311.3(PRNP):c.385A>G (p.Met129Val)	1799990	PRNP	[]	[]	['Jakob-Creutzfeldt disease', 'Genetic prion diseases', 'Fatal familial insomnia', 'not specified']
NM_000051.3(ATM):c.3994-2A>G	587782276	ATM	[]	[]	['Ataxia-telangiectasia syndrome', 'Hereditary cancer-predisposing syndrome']
NM_005211.3(CSF1R):c.1754-2A>G	281860267	CSF1R	[]	[]	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_004646.3(NPHS1):c.1756A>G (p.Arg586Gly)	730880174	NPHS1	[]	[]	['Finnish congenital nephrotic syndrome']
NM_005211.3(CSF1R):c.2320-2A>G	281860272	CSF1R	['CACYGAGGGAAAGCACTGCAGGG']	['CACYGAGGGAAAGCACTGCAGGG', 'GCACYGAGGGAAAGCACTGCAGG']	['Hereditary diffuse leukoencephalopathy with spheroids']
NM_000256.3(MYBPC3):c.3392T>C (p.Ile1131Thr)	370890951	MYBPC3	[]	[]	['Cardiomyopathy', 'Cardiac arrest', 'not specified']
NM_000551.3(VHL):c.586A>T (p.Lys196Ter)	281860296	VHL	[]	['GGTCTTYCTGCACATTTGGGTGG']	['Von Hippel-Lindau syndrome']
NM_005247.2(FGF3):c.146A>G (p.Tyr49Cys)	281860300	FGF3	['CAGYAGAGCTTGCGGCGCCGGGG', 'GCAGYAGAGCTTGCGGCGCCGGG', 'CGCAGYAGAGCTTGCGGCGCCGG']	['CAGYAGAGCTTGCGGCGCCGGGG', 'GCAGYAGAGCTTGCGGCGCCGGG', 'CGCAGYAGAGCTTGCGGCGCCGG']	['Deafness with labyrinthine aplasia microtia and microdontia (LAMM)']
NM_005247.2(FGF3):c.317A>G	281860306	FGF3	[]	[]	['Deafness with labyrinthine aplasia

(p.Tyr106Cys)					microtia and microdontia (LAMM)']
NM_000314.6(PTEN): c.403A>G (p.Ile135Val)	587782360	PTEN	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_004990.3(MARS) :c.1031A>G (p.Tyr344Cys)	766466297	MARS	[]	[]	['Pulmonary alveolar proteinosis', 'Interstitial lung and liver disease']
NM_006343.2(MERT K):c.1605-2A>G	730880273	MERTK	[]	[]	['Retinitis pigmentosa 38']
NM_003611.2(OFD1): c.935+706A>G	730880283	OFD1	[]	[]	['Retinitis Pigmentosa 23']
NM_004793.3(LONP1):c.2353A>G (p.Arg785Gly)	730880293	LONP1	[]	[]	['CODAS syndrome']
NM_001698.2(AUH):c .263-2A>G	730880311	AUH	[]	[]	['3-Methylglutaconic aciduria']
NM_001698.2(AUH):c .943-2A>G	730880312	AUH	[]	[]	['3-Methylglutaconic aciduria']
NM_000169.2(GLA):c .370-2A>G	730880444	-	[]	['GTGAACCYGAA ATGAGAGGGAGG']	['not provided']
NM_001110792.1(ME CP2):c.520A>G (p.Arg174Gly)	727505391	MECP2	[]	[]	['Rett disorder']
NM_030662.3(MAP2 K2):c.181A>G (p.Lys61Glu)	730880517	MAP2K2	[]	[]	['Cardiofaciocutaneo us syndrome', 'Rasopathy']
NM_000256.3(MYBP C3):c.1227-2A>G	730880531	MYBPC3	[]	['GTACCYGGGTG GGGGCCGCAGGG' , 'TGTACCYGGGTG GGGGCCGCAGG']	['Familial hypertrophic cardiomyopathy 4', 'Cardiomyopathy']
NM_000642.2(AGL):c .4260-12A>G	369973784	AGL	[]	[]	['Glycogen storage disease type III', 'Glycogen storage disease IIIa', 'Glycogen storage disease IIIb']
NM_000267.3(NF1):c. 1642-8A>G	267606602	NF1	[]	[]	['Neurofibromatosis, type 1', 'Juvenile myelomonocytic leukemia']
NM_000267.3(NF1):c. 5944-5A>G	267606604	NF1	[]	[]	['Neurofibromatosis, type 1', 'Neurofibromatosis, familial spinal']
m.1555A>G	267606617	MT-RNR1	[]	[]	['Aminoglycoside-induced deafness', 'Cardiomyopathy, restrictive', 'Deafness, nonsyndromic sensorineural, mitochondrial']
NM_022458.3(LMBR 1):c.423+5252A>G	606231150	LMBR1	[]	[]	['Triphalangeal thumb', 'Preaxial

					polydactyly 2']
NM_000642.2(AGL):c.3439A>G (p.Arg1147Gly)	267606639	AGL	[]	[]	['Glycogen storage disease IIIc']
NM_013411.4(AK2):c.494A>G (p.Asp165Gly)	267606643	AK2	[]	['TCAYCTTTCATGGGCTCTTTTGG']	['Reticular dysgenesis']
NM_001142800.1(EYS):c.9209T>C (p.Ile3070Thr)	183589498	EYS	[]	[]	['Retinitis pigmentosa']
NM_004183.3(BEST1):c.680A>G (p.Tyr227Cys)	267606677	BEST1	[]	[]	['Vitelliform dystrophy', 'Retinitis pigmentosa, concentric', 'not provided']
NM_005188.3(CBL):c.1144A>G (p.Lys382Glu)	267606705	CBL	[]	['TATTTYACATAGTTGGAATGTGG']	['Noonan syndrome-like disorder with or without juvenile myelomonocytic leukemia']
NM_001017361.2(KHDC3L):c.1A>G (p.Met1Val)	606231235	KHDC3L	[]	[]	['Hydatidiform mole, recurrent, 2']
NM_144577.3(CCDC114):c.487-2A>G	606231239	CCDC114	[]	[]	['Ciliary dyskinesia, primary, 20']
NM_000277.1(PAH):c.916A>G (p.Ile306Val)	62642934	PAH	[]	['GGCCAAYTTCCTGTAATTGGGGG', 'AGGCCAAYTTCCTGTAATTGGGG']	['Phenylketonuria', 'Hyperphenylalaninemia, non-pku', 'not provided']
NM_000257.3(MYH7):c.2792A>G (p.Glu931Gly)	730880760	MYH7	[]	[]	['Cardiomyopathy']
NM_207034.2(EDN3):c.335A>G (p.His112Arg)	267606778	EDN3	[]	[]	['Waardenburg syndrome type 4B']
NM_000117.2(EMD):c.1A>G (p.Met1Val)	267606782	EMD	[]	['TCCAYGGCGGGTGCGGGCTCAGG']	['Emery-Dreifuss muscular dystrophy, X-linked']
NM_003937.2(KYNU):c.592A>G (p.Thr198Ala)	606231307	KYNU	[]	[]	['Hydroxykynureninuria']
NM_004387.3(NKX2-5):c.461A>G (p.Glu154Gly)	587782928	NKX2-5	[]	[]	['Atrial septal defect 7 with or without atrioventricular conduction defects']
NM_000142.4(FGFR3):c.1454A>G (p.Gln485Arg)	267606808	FGFR3	[]	[]	['Thanatophoric dysplasia type 1']
NM_014053.3(FLVCR1):c.361A>G (p.Asn121Asp)	267606820	FLVCR1	[]	['AGGCGTYGACCAGCGAGTACAGG']	['Posterior column ataxia with retinitis pigmentosa']
NM_000257.3(MYH7):c.4664A>G (p.Glu1555Gly)	730880805	-	[]	['GCCCYCCTCGTGCTCCAGGGAGG', 'CTTGCCCYCCTCGTGCTCCAGGG']	['Cardiomyopathy']
NM_138387.3(G6PC3):c.346A>G (p.Met116Val)	267606834	G6PC3	[]	['TGATCAYGCAGTGTCCAGAAGGG', 'GTGATCAYGCAGTGTCCAGAAGG']	['Dursun syndrome']

NM_020347.3(LZTFL1):c.260T>C (p.Leu87Pro)	515726135	LZTFL1	[]	[]	['Bardet-Biedl syndrome', 'Bardet-Biedl syndrome 17']
NM_000175.3(GPI):c.1028A>G (p.Gln343Arg)	267606851	GPI	[]	['GTACYGGTCATAGGGCAGCATGG']	['Hemolytic anemia, nonspherocytic, due to glucose phosphate isomerase deficiency']
NM_005859.4(PURA):c.289A>G (p.Lys97Glu)	587782994	PURA	[]	[]	['Neonatal hypotonia', 'Intellectual disability', 'Seizures', 'Delayed speech and language development', 'Global developmental delay', 'Mental retardation, autosomal dominant 31']
NM_005144.4(HR):c.-218A>G	267606869	HR	['CTCYAGGGCCGCAGGTTGGAGGG']	['CTCYAGGGCCGCAGGTTGGAGGG', 'GCTCYAGGGCCGCAGGTTGGAGG', 'GGCGCTCYAGGGCCGCAGGTTGG']	['Marie Unna hereditary hypotrichosis 1']
NM_000257.3(MYH7):c.789A>G (p.Ile263Met)	730880855	MYH7	[]	[]	['Cardiomyopathy']
NM_000060.3(BTD):c.683A>G (p.Asp228Gly)	587783004	BTD	[]	[]	['Biotinidase deficiency']
NM_000257.3(MYH7):c.1051A>G (p.Lys351Glu)	730880864	MYH7	[]	[]	['Cardiomyopathy']
NM_015713.4(RRM2B):c.190T>C (p.Trp64Arg)	515726182	RRM2B	[]	['TTCCTTCYGGACAGCAGAAGAGG']	['RRM2B-related mitochondrial disease']
NM_005957.4(MTHFR):c.971A>G (p.Asn324Ser)	267606887	MTHFR	['CGCGGYTGAGGGTGTAGAAGTGG']	['CGCGGYTGAGGGTGTAGAAGTGG']	['Homocystinuria due to MTHFR deficiency']
NM_015713.4(RRM2B):c.368T>C (p.Phe123Ser)	515726187	RRM2B	[]	[]	['RRM2B-related mitochondrial disease']
m.12770A>G	267606894	MT-ND5	[]	[]	['Juvenile myopathy, encephalopathy, lactic acidosis AND stroke']
NM_000257.3(MYH7):c.1805A>G (p.Asn602Ser)	730880880	MYH7	[]	[]	['Cardiomyopathy']
NM_018109.3(MTPAP):c.1432A>G (p.Asn478Asp)	267606900	MTPAP	['AATGGATYCTGAATGTACAGAGG']	['AATGGATYCTGAATGTACAGAGG']	['Ataxia, spastic, 4, autosomal recessive']
NM_000257.3(MYH7):c.2717A>G (p.Asp906Gly)	267606908	MYH7	[]	[]	['Primary familial hypertrophic cardiomyopathy',

					'Familial hypertrophic cardiomyopathy 1', 'Cardiomyopathy']
NM_003122.4(SPINK1):c.160T>C (p.Tyr54His)	515726207	SPINK1	[]	[]	['Hereditary pancreatitis']
NM_003159.2(CDKL5):c.404-2A>G	587783080	CDKL5	[]	[]	['not provided']
NM_003159.2(CDKL5):c.449A>G (p.Lys150Arg)	587783083	CDKL5	['ACAGTYTTAGGACATCATTGTGG']	['ACAGTYTTAGGACATCATTGTGG']	['not provided']
NM_016203.3(PRKAG2):c.1589A>G (p.His530Arg)	267606977	PRKAG2	[]	[]	['Familial hypertrophic cardiomyopathy 6', 'not provided']
NM_198965.1(PTHLH):c.534A>G (p.Ter178Trp)	267606987	PTHLH	[]	[]	['Brachydactyly type E2']
NM_000531.5(OTC):c.122A>G (p.Asp41Gly)	74518351	OTC	[]	[]	['not provided']
NM_001134363.2(RBM20):c.1909A>G (p.Ser637Gly)	267607005	RBM20	[]	[]	['Dilated cardiomyopathy 1DD']
NM_000553.4(WRN):c.403A>G (p.Lys135Glu)	267607008	WRN	[]	[]	['Werner syndrome']
NM_002880.3(RAF1):c.1279A>G (p.Ser427Gly)	730881002	RAF1	[]	['GCTGCYGCCCTCGCACCACTGGG', 'GGCTGCYGCCCTCGCACCACTGG']	['Rasopathy']
NM_002977.3(SCN9A):c.29A>G (p.Gln10Arg)	267607030	SCN9A	[]	['AAGCTCYGAGGTCCTGGGGGAGG']	['Primary erythromelalgia']
NM_016955.3(SEPSECS):c.1001A>G (p.Tyr334Cys)	267607036	SEPSECS	[]	[]	['Pontocerebellar hypoplasia type 2D']
NM_007373.3(SHOC2):c.4A>G (p.Ser2Gly)	267607048	SHOC2	[]	['TACYCATGGTGACTCAAGCCTGG']	['Noonan-like syndrome with loose anagen hair', 'Rasopathy']
NM_005633.3(SOS1):c.1430A>G (p.Gln477Arg)	730881044	SOS1	[]	[]	['Rasopathy']
NM_007375.3(TARDBP):c.787A>G (p.Lys263Glu)	267607102	TARDBP	[]	[]	['FRONTOTEMPORAL DEMENTIA WITH TDP43 INCLUSIONS, TARDBP-RELATED']
NM_003286.2(TOP1):c.1598A>G (p.Asp533Gly)	267607131	-	[]	[]	[]
NM_021625.4(TRPV4):c.1805A>G (p.Tyr602Cys)	267607150	TRPV4	[]	[]	['Spondyloepiphyseal dysplasia Maroteaux type']
NM_000551.3(VHL):c	267607170	VHL	[]	[]	['Von Hippel-Lindau

.491A>G (p.Gln164Arg)					syndrome']
NM_001006657.1(WDR35):c.1877A>G (p.Glu626Gly)	267607174	WDR35	[]	[]	['Cranioectodermal dysplasia 2']
NM_024884.2(L2HGDH):c.293A>G (p.His98Arg)	267607206	L2HGDH	[]	[]	['L-2-hydroxyglutaric aciduria']
NM_002437.4(MPV17):c.262A>G (p.Lys88Glu)	267607256	MPV17	[]	[]	['Navajo neurohepatopathy']
NM_006888.4(CALM1):c.293A>G (p.Asn98Ser)	267607277	CALM1	[]	[]	['Catecholaminergic polymorphic ventricular tachycardia', 'Ventricular tachycardia, catecholaminergic polymorphic, 4']
NM_000487.5(ARSA):c.*96A>G	6151429	ARSA	[]	[]	['Metachromatic leukodystrophy', 'Arylsulfatase A pseudodeficiency', 'not provided']
NM_003122.4(SPINK1):c.194+2T>C	148954387	SPINK1	[]	[]	['Hereditary pancreatitis']
NM_000552.3(VWF):c.3437A>G (p.Tyr1146Cys)	267607326	VWF	[]	[]	['von Willebrand disease type 2', 'not provided']
NM_000489.4(ATRX):c.4826A>G (p.His1609Arg)	122445093	ATRX	[]	[]	['ATR-X syndrome']
NM_000489.4(ATRX):c.6488A>G (p.Tyr2163Cys)	122445098	ATRX	[]	[]	['ATR-X syndrome']
NM_000489.4(ATRX):c.6811A>G (p.Arg2271Gly)	122445112	ATRX	[]	[]	[]
NM_004380.2(CREBBP):c.3983-2A>G	587783486	CREBBP	[]	['GCAGCCCYAGGAAGTCCAGAAGG']	['Rubinstein-Taybi syndrome']
NM_004380.2(CREBBP):c.4508A>G (p.Tyr1503Cys)	587783497	CREBBP	[]	[]	['Rubinstein-Taybi syndrome']
NM_000051.3(ATM):c.3154-2A>G	730881357	ATM	[]	['AGCCYACGGGAAAAGAACTGTGG']	['Hereditary cancer-predisposing syndrome']
NM_178151.2(DCX):c.1027-2A>G	587783518	DCX	[]	[]	['Heterotopia']
NM_178151.2(DCX):c.520A>G (p.Lys174Glu)	587783557	DCX	[]	[]	['Heterotopia']
NM_178151.2(DCX):c.538A>G (p.Lys180Glu)	587783560	DCX	[]	[]	['Heterotopia']
NM_178151.2(DCX):c.607A>G (p.Thr203Ala)	587783570	DCX	[]	[]	['Heterotopia']
NM_001257235.1(AL	398122394	ALG13	[]	[]	['Congenital disorder

G13):c.8A>G (p.Asn3Ser)					of glycosylation type 1s']
NM_001256864.1(DNAJC6):c.801-2A>G	398122404	DNAJC6	[]	['AGGTATCYGAAACAGAAGGTTGG']	['Parkinson disease 19, juvenile-onset']
NM_001927.3(DES):c.1024A>G (p.Asn342Asp)	267607482	DES	[]	['GAATCGTYCTGCAGGAGAGGGGG']	['Myofibrillar myopathy 1', 'not provided']
NM_001927.3(DES):c.735+3A>G	267607483	DES	[]	[]	['Myofibrillar myopathy 1', 'not provided']
NM_001927.3(DES):c.1333A>G (p.Thr445Ala)	267607498	DES	[]	[]	['not provided']
NM_005249.4(FOXG1):c.757A>G (p.Asn253Asp)	587783641	FOXG1	[]	[]	['Rett syndrome, congenital variant']
NM_000252.2(MTM1):c.1406A>G (p.His469Arg)	587783789	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000391.3(TPP1):c.833A>G (p.Gln278Arg)	796053439	TPP1	[]	['CAGGTACYGCACATCTAGACTGG']	['not provided']
NM_000252.2(MTM1):c.301A>G (p.Ser101Gly)	587783818	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.343-2A>G	587783821	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.529-2A>G	587783831	MTM1	[]	[]	['Severe X-linked myotubular myopathy']
NM_000252.2(MTM1):c.550A>G (p.Arg184Gly)	587783835	MTM1	[]	['GTTATTCYCCAATGGTGATTGGG']	['Severe X-linked myotubular myopathy']
NM_000158.3(GBE1):c.691+2T>C	192044702	GBE1	[]	[]	['Glycogen storage disease, type IV']
NM_000252.2(MTM1):c.629A>G (p.Asp210Gly)	587783842	MTM1	[]	['TCATCAYCTGAGGCACGATACGG']	['Severe X-linked myotubular myopathy']
NM_000249.3(MLH1):c.545+3A>G	267607760	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_000249.3(MLH1):c.589-2A>G	267607767	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms', 'Hereditary cancer-predisposing syndrome']
NM_000249.3(MLH1):c.884+4A>G	267607777	MLH1	[]	['TGCTACAYTACCTGAGGTACAGG']	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_000081.3(LYST):c.10127A>G (p.Asn3376Ser)	80338669	LYST	[]	[]	['Ch\xc3\xa9diak-Higashi syndrome']
NM_000528.3(MAN2	80338678	MAN2B1	[]	[]	['Deficiency of

B1):c.1831-2A>G					alpha-mannosidase']
NM_022132.4(MCCC2):c.569A>G (p.His190Arg)	119103225	MCCC2	[]	[]	['3-methylcrotonyl CoA carboxylase 2 deficiency']
m.3260A>G	199474663	MT-TL1	['TTAAGTTYTATGCGATTACCGGG']	['TTAAGTTYTATGCGATTACCGGG']	['Cardiomyopathy with or without skeletal myopathy']
NM_014874.3(MFN2):c.827A>G (p.Gln276Arg)	119103264	MFN2	[]	[]	['Hereditary motor and sensory neuropathy with optic atrophy']
NM_004525.2(LRP2):c.770-2A>G	80338743	LRP2	[]	[]	['Donnai Barrow syndrome']
NM_005120.2(MED12):c.3020A>G (p.Asn1007Ser)	80338759	MED12	[]	[]	['X-linked mental retardation with marfanoid habitus syndrome']
NM_000834.3(GRIN2B):c.2172-2A>G	398122824	GRIN2B	[]	[]	['Mental retardation, autosomal dominant 6']
NM_000249.3(MLH1):c.1990-2A>G	267607883	MLH1	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms', 'not provided']
NM_000518.4(HBB):c.59A>G (p.Asn20Ser)	33972047	HBB	[]	['CACGYTCACCTTGCCCCACAGGG', 'CCACGYTCACCTTGCCCCACAGG']	['alpha Thalassemia']
NM_003688.3(CASK):c.2168A>G (p.Tyr723Cys)	398122844	CASK	[]	[]	['FG syndrome 4', 'Mental retardation and microcephaly with pontine and cerebellar hypoplasia']
NM_024675.3(PALB2):c.109-2A>G	730881897	PALB2	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000251.2(MSH2):c.1511-2A>G	267607962	MSH2	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms']
NM_003124.4(SPR):c.596-2A>G	398122922	SPR	[]	[]	['Sepiapterin reductase deficiency']
NM_022455.4(NSD1):c.4498-3A>G	587784120	NSD1	[]	[]	['Sotos syndrome 1']
NM_000455.4(STK11):c.889A>G (p.Arg297Gly)	730881978	STK11	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_001927.3(DES):c.1289-2A>G	398122940	DES	[]	[]	['Muscular dystrophy, limb-girdle, type 2r']
NM_000546.5(TP53):c.709A>G (p.Met237Val)	730882004	TP53	[]	['ACACAYGTAGTTGTAGTGGATGG']	['Li-Fraumeni syndrome', 'Hereditary cancer-predisposing syndrome']

NM_024876.3(ADCK4):c.857A>G (p.Asp286Gly)	398122979	ADCK4	[]	[]	['Nephrotic syndrome, type 9']
NM_022455.4(NSD1): c.6059A>G (p.Asn2020Ser)	587784178	NSD1	[]	[]	['Sotos syndrome 1']
NM_022455.4(NSD1): c.6356A>G (p.Asp2119Gly)	587784191	NSD1	[]	[]	['Sotos syndrome 1']
NM_007332.2(TRPA1):c.2564A>G (p.Asn855Ser)	398123010	-	[]	[]	['Familial episodic pain syndrome 1']
NM_001231.4(CASQ1):c.731A>G (p.Asp244Gly)	730882052	CASQ1	[]	['GGCTTGYCTGGGATGGTCACAGG']	['Myopathy, vacuolar, with casq1 aggregates']
NM_004004.5(GJB2): c.487A>G (p.Met163Val)	80338949	GJB2	[]	[]	['Deafness, autosomal recessive 1A', 'not specified']
NM_130466.3(UBE3B):c.545-2A>G	398123022	UBE3B	[]	[]	['Kaufman oculocerebrofacial syndrome']
NM_000334.4(SCN4A):c.4078A>G (p.Met1360Val)	80338959	SCN4A	[]	['GATCAYGATGGTGATGTCGAAGG']	['Hyperkalemic Periodic Paralysis Type 1']
NM_000334.4(SCN4A):c.4108A>G (p.Met1370Val)	80338960	SCN4A	[]	['CCATCAYGGTGACCATGTTGAGG']	['Hyperkalemic Periodic Paralysis Type 1']
NM_000334.4(SCN4A):c.4774A>G (p.Met1592Val)	80338962	SCN4A	[]	['TGTACAYGTTGACCACGATGAGG']	['Hyperkalemic Periodic Paralysis Type 1', 'Familial hyperkalemic periodic paralysis']
NM_015250.3(BICD2):c.2321A>G (p.Glu774Gly)	398123030	BICD2	[]	[]	['Spinal muscular atrophy, lower extremity predominant 2, autosomal dominant']
NM_006012.2(CLPP): c.270+4A>G	398123035	CLPP	[]	[]	['Autosomal recessive hearing impairment with normal menstrual cycles']
NM_000179.2(MSH6):c.3439-2A>G	267608098	MSH6	[]	[]	['Hereditary Nonpolyposis Colorectal Neoplasms', 'Hereditary cancer-predisposing syndrome', 'not provided']
NM_002246.2(KCNK3):c.575A>G (p.Tyr192Cys)	398123043	KCNK3	[]	[]	['Primary pulmonary hypertension 4']
NM_001070.4(TUBG1):c.275A>G (p.Tyr92Cys)	398123046	TUBG1	[]	[]	['Cortical dysplasia, complex, with other brain malformations 4']
NM_000383.3(AIRE):	179363882	AIRE	[]	[]	['Polyglandular

c.254A>G (p.Tyr85Cys)					autoimmune syndrome, type 1', 'not provided']
NM_001651.3(AQP5): c.367A>G (p.Asn123Asp)	398123057	AQP5	[]	[]	['Diffuse palmoplantar keratoderma, Bothnian type']
NM_012160.4(FBXL4):c.1694A>G (p.Asp565Gly)	398123062	FBXL4	[]	['TATGYCCAGCTG CTGTAACCTGG']	['Mitochondrial DNA depletion syndrome 13 (encephalomyopathi c type)']
NM_024531.4(SLC52 A2):c.914A>G (p.Tyr305Cys)	398123068	SLC52A2	[]	[]	['Brown-Vialetto-Van Laere syndrome 2']
NM_001039550.1(DN AJB2):c.14A>G (p.Tyr5Cys)	730882140	DNAJB2	[]	['GATCTCGYAGTA GGATGCCATGG']	['Charcot-Marie-Tooth disease', 'Charcot-Marie-Tooth disease, axonal, type 2T']
NM_052859.3(RFT1): c.1222A>G (p.Met408Val)	796053522	RFT1	[]	['GCAYCACAAAA TTGTACCTGGGG', 'AGCAYCACAAAA TTGTACCTGGG', 'CAGCAYCACAAA ATTGTACCTGG']	['Congenital disorder of glycosylation type 1N']
NM_007294.3(BRCA1):c.5057A>G (p.His1686Arg)	730882166	BRCA1	[]	[]	['Breast-ovarian cancer, familial 1']
NM_000050.4(ASS1): c.496-2A>G	398123130	ASS1	[]	[]	['Citrullinemia type I']
NM_012233.2(RAB3 GAP1):c.649-2A>G	730882183	RAB3GA P1	[]	[]	['Warburg micro syndrome 1']
NM_000159.3(GCDH) :c.542A>G (p.Glu181Gly)	398123194	GCDH	[]	[]	['Glutaric aciduria, type 1', 'not provided']
NM_000288.3(PEX7): c.340-10A>G	267608255	PEX7	[]	[]	['Phytanic acid storage disease', 'Peroxisome biogenesis disorder 9B']
NM_000038.5(APC):c .221-2A>G	786201291	APC	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_000169.2(GLA):c .509A>G (p.Asp170Gly)	398123210	-	[]	[]	['Fabry disease']
NM_000169.2(GLA):c .548-2A>G	398123211	-	[]	['AACCYGTATGA GAAAACAATGGG' , 'TAACCYGTATGA GAAAACAATGG']	['Fabry disease']
NM_006888.4(CALM 1):c.389A>G (p.Asp130Gly)	730882252	CALM1	[]	[]	['Long QT syndrome 14']
NM_000169.2(GLA):c .647A>G (p.Tyr216Cys)	398123217	-	[]	[]	['Fabry disease']
NM_006306.3(SMC1	587784423	SMC1A	[]	['AGCCYGTGCAA	['Congenital

A):c.616-2A>G				ACAGGGGAATGG']	muscular hypertrophy-cerebral syndrome']
NM_000255.3(MUT): c.1445-2A>G	398123276	MUT	[]	[]	['Methylmalonic aciduria due to methylmalonyl-CoA mutase deficiency', 'not provided']
NM_001083962.1(TCF4):c.991-2A>G	587784470	TCF4	[]	[]	['Pitt-Hopkins syndrome']
NM_000060.3(BTD):c.1205A>G (p.Asn402Ser)	201023772	BTD	[]	[]	['Biotinidase deficiency']
NC_000007.14:g.62535490A>G	483352872	-	[]	[]	['Isolated growth hormone deficiency type 1B']
NM_000271.4(NPC1): c.1832A>G (p.Asp611Gly)	483352887	NPC1	[]	[]	['Niemann-Pick disease type C1']
NM_152419.2(HGSNAT):c.372-2A>G	483352896	HGSNAT	[]	[]	['Mucopolysaccharidosis, MPS-III-C']
NM_001199397.1(NEK1):c.869-2A>G	483352906	NEK1	[]	[]	['Short rib-polydactyly syndrome, Majewski type']
NM_000350.2(ABCA4):c.67-2A>G	398123339	ABCA4	[]	[]	['Stargardt disease 1']
NM_004992.3(MECP2):c.27-2A>G	267608412	MECP2	[]	[]	['Rett disorder', 'not provided']
NM_003159.2(CDKL5):c.100-2A>G	267608423	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2', 'not provided']
NM_003159.2(CDKL5):c.125A>G (p.Lys42Arg)	267608429	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2', 'not provided']
NM_000487.5(ARSA):c.1108-2A>G	398123411	ARSA	[]	['GGCTCYGGGGGCAGAGTCAGGGG', 'GGGCTCYGGGGGCAGAGTCAGGG', 'AGGGCTCYGGGGGCAGAGTCAGG']	['Metachromatic leukodystrophy']
NM_003159.2(CDKL5):c.380A>G (p.His127Arg)	267608468	CDKL5	[]	[]	['Atypical Rett syndrome', 'not provided']
NM_000489.4(ATRX):c.134-2A>G	398123420	ATRX	[]	[]	['not provided']
NM_003159.2(CDKL5):c.464-2A>G	267608480	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2', 'not provided']
NM_000489.4(ATRX):c.536A>G (p.Asn179Ser)	398123425	ATRX	[]	[]	['not provided']
NM_000512.4(GALNS):c.1171A>G (p.Met391Val)	398123429	GALNS	[]	['CCGCCAYCAGCGTGTCGCCACGG']	['Mucopolysaccharidosis, MPS-IV-A', 'not provided']

NM_003159.2(CDKL5):c.578A>G (p.Asp193Gly)	267608500	CDKL5	[]	['ATGYCCACGGACTTTCCATAGGG', 'CATGYCCACGGACTTTCCATAGG']	['Early infantile epileptic encephalopathy 2']
NM_000521.3(HEXB):c.1243-2A>G	398123446	HEXB	[]	[]	['Sandhoff disease', 'not provided']
NM_003159.2(CDKL5):c.978-2A>G	267608553	CDKL5	[]	[]	['Early infantile epileptic encephalopathy 2', 'not provided']
NM_001164342.2(ZBTB20):c.1787A>G (p.His596Arg)	483353066	ZBTB20	[]	[]	['Primrose syndrome']
NM_000402.4(G6PD):c.188T>C (p.Ile63Thr)	398123552	-	[]	['ACACACAYATTCATCATCATGGG']	['Anemia, nonspherocytic hemolytic, due to G6PD deficiency', 'not provided']
NM_001893.4(CSNK1D):c.130A>G (p.Thr44Ala)	104894561	CSNK1D	[]	[]	['Advanced sleep phase syndrome, familial, 2']
NM_000155.3(GALT):c.563A>G (p.Gln188Arg)	75391579	GALT	[]	['TTACCYGGCAGTGGGGGTGGGGG', 'CTTACCYGGCAGTGGGGGTGGGG', 'CCTTACCYGGCAGTGGGGGTGGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase', 'not provided']
NM_007055.3(POLR3A):c.2554A>G (p.Met852Val)	267608671	POLR3A	[]	[]	['Hypomyelinating leukodystrophy 7']
NM_001848.2(COL6A1):c.805-2A>G	398123639	COL6A1	[]	['TTCTCCCYGGAACACAAAACAGG']	['Ullrich congenital muscular dystrophy', 'Bethlem myopathy', 'not provided']
NM_001918.3(DBT):c.773-2A>G	398123674	DBT	[]	[]	['Maple syrup urine disease', 'not provided']
NM_001999.3(FBN2):c.3740T>C (p.Met1247Thr)	149054177	FBN2	['GAATGTAYGATAATGAACGGAGG']	['GAATGTAYGATAATGAACGGAGG']	['not specified', 'Macular degeneration, early-onset']
NM_003482.3(KMT2D):c.5645-2A>G	398123750	KMT2D	[]	['GCAGTTCYGTGGGGGAATGAAGG']	['Kabuki make-up syndrome', 'not provided']
NM_003494.3(DYSF):c.1398-2A>G	398123769	DYSF	[]	[]	['Limb-girdle muscular dystrophy, type 2B', 'not provided']
NM_015560.2(OPA1):c.1146A>G (p.Ile382Met)	143319805	OPA1	[]	[]	['Dominant hereditary optic atrophy', 'Optic Atrophy Type 1', 'not specified', 'not provided']
NM_203447.3(DOCK8):c.1418A>G (p.Lys473Arg)	112321280	DOCK8	[]	[]	['Hyperimmunoglobulin E recurrent infection syndrome, autosomal recessive']

NM_001145.4(ANG):c.121A>G (p.Lys41Glu)	121909537	-	['TGGTTYGGCATCATAGTGCTGGG', 'GTGGTTYGGCATCATAGTGCTGG']	['TGGTTYGGCATCATAGTGCTGGG', 'GTGGTTYGGCATCATAGTGCTGG']	['Amyotrophic lateral sclerosis type 9']
NM_004006.2(DMD):c.3432+3A>G	398123938	DMD	[]	[]	['Dilated cardiomyopathy 3B']
NM_006514.3(SCN10A):c.1661T>C (p.Leu554Pro)	138404783	SCN10A	[]	[]	['Episodic pain syndrome, familial, 2']
NM_004006.2(DMD):c.6763-2A>G	398124033	DMD	[]	[]	['Dilated cardiomyopathy 3B']
NM_001079802.1(FKTN):c.1112A>G (p.Tyr371Cys)	119464998	FKTN	[]	[]	[]
NM_004006.2(DMD):c.9224+61934A>G	398124084	DMD	[]	[]	['Dilated cardiomyopathy 3B']
NM_004006.2(DMD):c.9225-647A>G	398124091	DMD	[]	[]	['Duchenne muscular dystrophy', 'Becker muscular dystrophy', 'Dilated cardiomyopathy 3B']
NM_004006.2(DMD):c.9650-2A>G	398124100	DMD	[]	[]	['Duchenne muscular dystrophy', 'Becker muscular dystrophy', 'Dilated cardiomyopathy 3B']
NM_198578.3(LRRK2):c.3342A>G (p.Leu1114=)	35808389	LRRK2	[]	[]	['Parkinson disease 8, autosomal dominant']
NM_031229.2(RBCK1):c.1160A>G (p.Asn387Ser)	566912235	RBCK1	[]	[]	['Polyglucosan body myopathy 1 with or without immunodeficiency']
NM_000178.2(GSS):c.656A>G (p.Asp219Gly)	28938472	GSS	[]	[]	['Glutathione synthetase deficiency of erythrocytes, hemolytic anemia due to']
NM_025132.3(WDR19):c.407-2A>G	374400438	WDR19	[]	[]	['SENIOR-LOKEN SYNDROME 8']
NM_144997.5(FLCN):c.1433-2A>G	398124528	FLCN	[]	['CCCACYGGGGAGAAGGGCAGGGG', 'GCCCACYGGGGAGAAGGGCAGGG', 'GGCCCACYGGGGAGAAGGGCAGG']	['Hereditary cancer-predisposing syndrome', 'not provided']
NM_144997.5(FLCN):c.250-2A>G	398124533	FLCN	[]	[]	['not provided']
NM_000146.3(FTL):c.-160A>G	398124633	FTL	[]	[]	['Hyperferritinemia cataract syndrome']
NM_003184.3(TAF2):c.1945T>C (p.Trp649Arg)	398124645	TAF2	[]	[]	['Mental retardation, autosomal recessive 40']
NM_013281.3(FLRT3	398124654	-	[]	[]	['Hypogonadotropic

):c.1016A>G (p.Lys339Arg)					hypogonadism 21 with or without anosmia']
NM_002834.3(PTPN11):c.767A>G (p.Gln256Arg)	397507523	PTPN11	[]	[]	['Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_000054.4(AVPR2):c.614A>G (p.Tyr205Cys)	104894749	AVPR2	['ACAYAGGTGCGACGGCCCCAGGG', 'GACAYAGGTGCGACGGCCCCAGG']	['ACAYAGGTGCGACGGCCCCAGGG', 'GACAYAGGTGCGACGGCCCCAGG']	['Nephrogenic diabetes insipidus', 'Nephrogenic diabetes insipidus, X-linked']
NM_031157.2(HNRNPA1):c.956A>G (p.Asn319Ser)	397518454	HNRNPA1	[]	[]	['Amyotrophic lateral sclerosis 20']
NM_000277.1(PAH):c.662A>G (p.Glu221Gly)	62514934	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_000219.5(KCNE1):c.176T>C (p.Leu59Pro)	141813529	KCNE1	[]	[]	['Congenital long QT syndrome']
NM_001165963.1(SCN1A):c.1076A>G (p.Asn359Ser)	794726713	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_013339.3(ALG6):c.391T>C (p.Tyr131His)	35383149	ALG6	[]	[]	['Congenital disorder of glycosylation type 1C', 'not specified']
NM_153767.3(KCNJ1):c.1013T>C (p.Met338Thr)	59172778	KCNJ1	[]	[]	['Bartter syndrome antenatal type 2']
NM_176824.2(BBS7):c.968A>G (p.His323Arg)	119466001	BBS7	[]	[]	['Bardet-Biedl syndrome 7']
NM_000199.3(SGSH):c.892T>C (p.Ser298Pro)	138504221	SGSH	[]	[]	['Mucopolysaccharidosis, MPS-III-A', 'not provided']
NM_000891.2(KCNJ2):c.220A>G (p.Thr74Ala)	199473652	KCNJ2			['Congenital long QT syndrome']
NM_001165963.1(SCN1A):c.1048A>G (p.Met350Val)	794726768	SCN1A	['ACAYATATCCCTCTGGACATTGG']	['ACAYATATCCCTCTGGACATTGG']	['Severe myoclonic epilepsy in infancy']
NM_002863.4(PYGL):c.1016A>G (p.Asn339Ser)	113993976	PYGL	[]	[]	['Glycogen storage disease, type VI']
NM_001165963.1(SCN1A):c.2537A>G (p.Glu846Gly)	794726794	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_002693.2(POLG):c.2864A>G (p.Tyr955Cys)	113994099	POLG	[]	[]	['Autosomal dominant progressive external ophthalmoplegia with mitochondrial DNA deletions 1']
NM_000920.3(PC):c.1705A>G	113994144	PC	[]	[]	['Pyruvate carboxylase

(p.Thr569Ala)					deficiency']
NM_025265.3(TSEN2):c.926A>G (p.Tyr309Cys)	113994149	TSEN2	[]	['CAGAGCAYAGACCAAGAAAAAGG']	['Pontocerebellar hypoplasia type 2B']
NM_001039958.1(MESP2):c.271A>G (p.Lys91Glu)	113994156	MESP2	[]	[]	['Spondylocostal dysostosis 2']
NM_024649.4(BBS1):c.1340-2A>G	113994180	-	[]	[]	['Bardet-Biedl syndrome']
NM_033028.4(BBS4):c.157-2A>G	113994192	BBS4	[]	[]	['Bardet-Biedl syndrome', 'Bardet-Biedl syndrome 4']
NM_212472.2(PRKAR1A):c.1A>G (p.Met1Val)	281864779	PRKAR1A	[]	[]	['Carney complex, type 1']
NM_212472.2(PRKAR1A):c.178-2A>G	281864796	PRKAR1A	[]	[]	['Carney complex, type 1']
NM_212472.2(PRKAR1A):c.891+3A>G	281864799	PRKAR1A	[]	[]	['Carney complex, type 1']
NM_001165963.1(SCN1A):c.433A>G (p.Met145Val)	794726849	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy']
NM_005710.2(PQBP1):c.194A>G (p.Tyr65Cys)	121917899	PQBP1	[]	[]	['Renpenning syndrome 1']
NM_000921.4(PDE3A):c.1333A>G (p.Thr445Ala)	794726865	PDE3A	['CGAGGYGGTGGTGGTCCAAGTGG']	['CGAGGYGGTGGTGGTCCAAGTGG']	['Brachydactyly with hypertension']
NM_024312.4(GNPTAB):c.1285-2A>G	281864974	GNPTAB	[]	[]	['Pseudo-Hurler polydystrophy']
NM_024312.4(GNPTAB):c.2783A>G (p.Lys928Arg)	281865003	GNPTAB	[]	[]	['I cell disease']
NM_024312.4(GNPTAB):c.2867A>G (p.His956Arg)	281865005	GNPTAB	[]	[]	['Pseudo-Hurler polydystrophy']
NM_024312.4(GNPTAB):c.3053A>G (p.Asp1018Gly)	281865007	GNPTAB	[]	[]	['I cell disease']
NM_024312.4(GNPTAB):c.3458A>G (p.Asn1153Ser)	281865019	GNPTAB	[]	[]	['Pseudo-Hurler polydystrophy']
NM_024312.4(GNPTAB):c.118-2A>G	281865023	GNPTAB	[]	[]	['I cell disease']
NM_198578.3(LRRK2):c.5605A>G (p.Met1869Val)	281865052	LRRK2	[]	['TCAACAYAATATTTCTAGGCAGG']	['Parkinson disease 8, autosomal dominant']
NM_139241.3(FGD4):c.1762-2A>G	281865065	FGD4	[]	[]	['Charcot-Marie-Tooth disease, type 4H']
NM_006121.3(KRT1):c.1445A>G (p.Tyr482Cys)	58420087	KRT1	[]	[]	['Bullous ichthyosiform erythroderma', 'not provided']
NM_000195.4(HPS1):c.716T>C (p.Leu239Pro)	281865080	HPS1	[]	[]	['Hermansky-Pudlak syndrome 1']
NM_000195.4(HPS1):	281865090	HPS1	[]	[]	['Hermansky-Pudlak

c.2003T>C (p.Leu668Pro)					syndrome 1']
NM_022081.5(HPS4): c.461A>G (p.His154Arg)	281865098	HPS4	[]	[]	['Hermansky-Pudlak syndrome 4']
NM_000277.1(PAH):c .1157A>G (p.Tyr386Cys)	62516141	PAH	[]	[]	['Phenylketonuria', 'not provided']
NM_025114.3(CEP29 0):c.2991+1655A>G	281865192	CEP290	['GAGATAY TCACAATT ACAACTGG ']	['GATAYTCACAAT TACAACTGGGG', 'AGATAYTCACAA TTACAACTGGG', 'GAGATAYTCACA ATTACAACTGG']	['Leber congenital amaurosis 10', 'not provided']
NM_018319.3(TDP1): c.1478A>G (p.His493Arg)	119467003	TDP1	[]	[]	['Spinocerebellar ataxia autosomal recessive with axonal neuropathy']
NM_000051.3(ATM): c.5762_5763insNG_00 9830.1:g.91138_91274	774925473	ATM	[]	[]	['Ataxia-telangiectasia variant']
NM_004614.4(TK2):c. 562A>G (p.Thr188Ala)	281865495	TK2	[]	['AAGYCTCAGGA TTGGTCCGAAGG']	['Mitochondrial DNA depletion syndrome 2']
NM_003494.3(DYSF): c.3041A>G (p.Tyr1014Cys)	756328339	DYSF	[]	['CTAYACTCCCAG CCTGGGGGAGG', 'ATGCTAYACTCC CAGCCTGGGGG', 'GATGCTAYACTC CCAGCCTGGGG']	['Limb-girdle muscular dystrophy, type 2B']
NM_000531.5(OTC):c .1034A>G (p.Tyr345Cys)	72558492	OTC	['AGGTGAG YAATCTGT CAGCAGGG ']	['AGGTGAGYAAT CTGTCAGCAGGG']	['not provided']
NM_000518.4(HBB):c .199A>G (p.Lys67Glu)	34165323	HBB	[]	[]	['Hemoglobinopathy']
NM_153427.2(PITX2) :c.262A>G (p.Lys88Glu)	387906810	PITX2	[]	['TCTYGAACCAAA CCTGGGGGCGG', 'GATTCTYGAACC AAACCTGGGGG', 'CGATTCTYGAAC CAAACCTGGGG']	['Axenfeld-Rieger syndrome type 1']
NM_030964.3(SPRY4):c.530A>G (p.Lys177Arg)	78310959	SPRY4	[]	['AGTGCYTGTCCA GCTCGGGTGGG', 'AAGTGCYTGTCC AGCTCGGGTGG']	['Hypogonadotropic hypogonadism 17 with or without anosmia']
NM_002834.3(PTPN1 1):c.922A>G (p.Asn308Asp)	28933386	PTPN11	[]	[]	['Noonan syndrome', 'Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_000518.4(HBB):c .-50-29A>G	34598529	HBB	[]	[]	['alpha Thalassemia', 'Beta thalassemia intermedia']
NM_207352.3(CYP4V 2):c.1393A>G (p.Arg465Gly)	144109267	CYP4V2	[]	['TTCCYGGGGCCA GCAGAGAAGGG', 'GTTCCYGGGGCC AGCAGAGAAGG']	['Bietti crystalline corneoretinal dystrophy']
NM_001360.2(DHCR 7):c.1A>G	104886033	DHCR7	[]	[]	['Smith-Lemli-Opitz syndrome']

(p.Met1Val)					
NM_000495.4(COL4A5):c.1A>G (p.Met1Val)	104886050	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.2692A>G (p.Met898Val)	104886192	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.2746A>G (p.Ser916Gly)	104886193	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_004572.3(PKP2): c.2062T>C (p.Ser688Pro)	144601090	PKP2	[]	[]	['Arrhythmogenic right ventricular cardiomyopathy', 'not specified', 'not provided']
NM_000495.4(COL4A5):c.4790A>G (p.Tyr1597Cys)	104886298	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.1340-2A>G	104886319	COL4A5	[]	['CACCYGAGTAAGATAAAGAAAGG']	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.2042-18A>G	104886341	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.2147-2A>G	104886344	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.2510-33A>G	104886358	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000495.4(COL4A5):c.3107-2A>G	104886379	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000051.3(ATM): c.7268A>G (p.Glu2423Gly)	121434221	ATM	[]	[]	['Mantle cell lymphoma']
NM_000495.4(COL4A5):c.3605-2A>G	104886385	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_000096.3(CP):c.1209-2A>G	386134137	CP	[]	[]	['Deficiency of ferroxidase']
NM_000348.3(SRD5A2):c.692A>G (p.His231Arg)	121434251	SRD5A2	[]	[]	['3-Oxo-5 alpha-steroid delta 4-dehydrogenase deficiency']
NM_002739.3(PRKCG):c.76A>G (p.Arg26Gly)	386134157	PRKCG	[]	[]	['Spinocerebellar ataxia 14']
NM_000383.3(AIRE): c.247A>G (p.Lys83Glu)	121434255	AIRE	[]	[]	['AUTOIMMUNE POLYENDOCRINOPATHY SYNDROME, TYPE I']
NM_000495.4(COL4A5):c.466-2A>G	104886416	COL4A5	[]	['ACCCYAAAAGAAGCCATCAATGG']	['Alport syndrome, X-linked recessive']
NM_001127328.2(ACADM):c.589A>G (p.Thr197Ala)	121434279	ACADM	[]	[]	['Medium-chain acyl-coenzyme A dehydrogenase deficiency']
NM_000495.4(COL4A5):c.892-2A>G	104886453	COL4A5	[]	[]	['Alport syndrome, X-linked recessive']
NM_020533.2(MCOLN1):c.406-2A>G	104886461	MCOLN1	['TACYGTGGGCAGAGAAGGGGA	['TACYGTGGGCAGAGAAGGGGAGG',	['Ganglioside sialidase deficiency', 'not provided']

			GG']	'AGGTACYGTGGG CAGAGAAGGGG', 'CAGGTACYGTGG GCAGAGAAGGG']	
NM_018136.4(ASPM) :c.2761-25A>G	199422149	ASPM	[]	[]	['Primary autosomal recessive microcephaly 5']
NM_017780.3(CHD7): c.3082A>G (p.Ile1028Val)	121434338	CHD7	[]	[]	['CHARGE association', 'not provided']
NM_017780.3(CHD7): c.164A>G (p.His55Arg)	121434345	CHD7	[]	[]	['Kallmann syndrome 5']
NM_152783.4(D2HG DH):c.1315A>G (p.Asn439Asp)	121434362	D2HGDH	['GCAGGTY ACCATCTC CTGGAGGG ', 'TGCAGGT YACCATCT CCTGGAGG ']	['GCAGGTYACCAT CTCCTGGAGGG', 'TGCAGGTYACCA TCTCCTGGAGG']	['D-2-hydroxyglutaric aciduria 1']
NM_005006.6(NDUF S1):c.755A>G (p.Asp252Gly)	199422224	NDUFS1	[]	[]	['Mitochondrial complex I deficiency', 'not provided']
NM_002894.2(RBBP8):c.1009A>G (p.Lys337Glu)	121434388	RBBP8	[]	[]	['Carcinoma of pancreas']
NM_004621.5(TRPC6):c.428A>G (p.Asn143Ser)	121434391	TRPC6	[]	[]	['Focal segmental glomerulosclerosis 2']
NM_003705.4(SLC25 A12):c.1769A>G (p.Gln590Arg)	121434396	SLC25A1 2	[]	[]	['Hypomyelination, global cerebral']
NM_001363.4(DKC1): c.127A>G (p.Lys43Glu)	199422243	DKC1	[]	[]	['Dyskeratosis congenita X-linked']
NM_001084.4(PLOD3):c.668A>G (p.Asn223Ser)	121434414	PLOD3	[]	[]	['Bone fragility with contractures, arterial rupture, and deafness']
NM_006702.4(PNPLA 6):c.3034A>G (p.Met1012Val)	121434415	PNPLA6	[]	[]	['Spastic paraplegia 39']
NR_001566.1(TERC): n.48A>G	199422262	TERC	[]	[]	['Dyskeratosis congenita autosomal dominant']
NM_004984.2(KIF5A) :c.767A>G (p.Asn256Ser)	121434441	KIF5A	[]	[]	['Spastic paraplegia 10']
NM_004984.2(KIF5A) :c.827A>G (p.Tyr276Cys)	121434443	KIF5A	[]	['GAACAYAGCTTT TCTGGGGGAGG']	['Spastic paraplegia 10']
m.10438A>G	121434456	MT-TR	[]	[]	['Mitochondrial encephalomyopathy']
NM_198253.2(TERT): c.2537A>G (p.Tyr846Cys)	199422302	TERT	[]	[]	['Aplastic anemia']

m.12320A>G	121434463	MT-TL2	['TGGAGTYGCACCAAAATTTTTGG']	['GAGTYGCACCAAAATTTTTGGGG', 'GGAGTYGCACCAAAATTTTTGGG', 'TGGAGTYGCACCAAAATTTTTGG']	['Mitochondrial myopathy']
m.4317A>G	121434465	MT-TI	[]	[]	[]
m.4269A>G	121434466	MT-TI	['ACAYATTTCTTAGGTTTGAGGGG', 'GACAYATTTCTTAGGTTTGAGGG']	['ACAYATTTCTTAGGTTTGAGGGG', 'GACAYATTTCTTAGGTTTGAGGG', 'AGACAYATTTCTTAGGTTTGAGG']	[]
m.4295A>G	121434467	MT-TI	[]	[]	['Primary familial hypertrophic cardiomyopathy', 'Deafness, nonsyndromic sensorineural, mitochondrial']
m.4300A>G	121434470	MT-TI	[]	[]	['Primary familial hypertrophic cardiomyopathy']
NM_001099274.1(TINF2):c.850A>G (p.Thr284Ala)	199422314	TINF2	[]	['TGACTGYGGGGCGCTCCTTATGG']	['Dyskeratosis congenita autosomal dominant']
NM_004044.6(ATIC):c.1277A>G (p.Lys426Arg)	121434478	ATIC	[]	['AGTGTACYTGACAGCAATGGTGG']	['AICAR transformylase/IMP cyclohydrolase deficiency']
NM_001099274.1(TINF2):c.871A>G (p.Arg291Gly)	199422319	TINF2	[]	[]	['Dyskeratosis congenita autosomal dominant']
NM_015474.3(SAMHD1):c.760A>G (p.Met254Val)	121434521	SAMHD1	[]	[]	['Aicardi Goutieres syndrome 5']
NM_001103.3(ACTN2):c.26A>G (p.Gln9Arg)	121434525	ACTN2	[]	[]	['Dilated cardiomyopathy 1AA', 'Cardiomyopathy', 'Dilated cardiomyopathy', 'not specified']
NM_000155.3(GALT):c.812A>G (p.Glu271Gly)	111033765	GALT	[]	['CGCYCAGCAGGGGTCAGCTCAGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase', 'not provided']
NM_000316.2(PTH1R):c.668A>G (p.His223Arg)	121434597	PTH1R	[]	[]	['Metaphyseal chondrodysplasia, Jansen type']
NM_006006.4(ZBTB16):c.1849A>G (p.Met617Val)	121434606	ZBTB16	[]	['GATCAYGGCCGAGTAGTCCCGGG', 'TGATCAYGGCCGAGTAGTCCCGG']	['Skeletal defects, genital hypoplasia, and mental retardation']
NM_014305.3(TGDS):c.700T>C	544436734	TGDS	[]	[]	['Catel Manzke syndrome']

(p.Tyr234His)					
NM_002835.3(PTPN12):c.182A>G (p.Lys61Arg)	121434623	PTPN12	[]	[]	['Carcinoma of colon']
NM_000035.3(ALDOB):c.1027T>C (p.Tyr343His)	369586696	ALDOB	[]	[]	['Hereditary fructosuria']
NM_006180.4(NTRK2):c.2165A>G (p.Tyr722Cys)	121434633	NTRK2	[]	[]	['Obesity, hyperphagia, and developmental delay']
NM_000107.2(DDB2):c.730A>G (p.Lys244Glu)	121434639	DDB2	[]	[]	['Xeroderma pigmentosum, group E']
NM_000017.3(ACADS):c.1108A>G (p.Met370Val)	566325901	ACADS	[]	['AGCCCAYGCCGCCCAGGATCTGG']	['not provided']
NM_001151.3(SLC25A4):c.311A>G (p.Asp104Gly)	28999114	SLC25A4	[]	[]	['Autosomal dominant progressive external ophthalmoplegia with mitochondrial DNA deletions 2']
NM_012079.5(DGAT1):c.751+2T>C	148665132	DGAT1	[]	['ACCGCGGYGAGGACCTCTGTGGG']	['Diarrhea 7']
NM_002036.3(ACKR1):c.-67T>C	2814778	ACKR1	[]	[]	['White blood cell count quantitative trait locus 1']
NM_000492.3(CFTR):c.1666A>G (p.Ile556Val)	75789129	CFTR	[]	[]	['Cystic fibrosis', 'not specified']
NM_000155.3(GALT):c.574A>G (p.Ser192Gly)	111033830	GALT	[]	['TGCYGGCCCATACCTGTCAAGGG', 'CTGCYGGCCCATACCTGTCAAGG']	['Deficiency of UDPglucose-hexose-1-phosphate uridylyltransferase']
NM_174916.2(UBR1):c.407A>G (p.His136Arg)	119477054	UBR1	[]	[]	['Johanson-Blizzard syndrome']
m.3274A>G	199474666	MT-TL1	[]	[]	[]
NM_000060.3(BTD):c.128A>G (p.His43Arg)	146011150	BTD	[]	[]	['Biotinidase deficiency']
NM_172107.2(KCNQ2):c.635A>G (p.Asp212Gly)	118192202	KCNQ2	[]	[]	['Benign familial neonatal seizures 1']
NM_006493.2(CLN5):c.1121A>G (p.Tyr374Cys)	148862100	CLN5	[]	[]	['Ceroid lipofuscinosis neuronal 5']
NM_000060.3(BTD):c.880A>G (p.Ile294Val)	35976361	BTD	[]	[]	['Biotinidase deficiency']
NM_000132.3(F8):c.1226A>G (p.Glu409Gly)	28933671	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.5600A>G (p.His1867Arg)	28933679	F8	[]	['GAGYGCACATCTTTTTCCTAGGG', 'TGAGYGCACATCTTTTTCCTAGG']	['Hereditary factor VIII deficiency disease']
NM_000266.3(NDP):c.1A>G (p.Met1Val)	28933685	NDP	[]	[]	['Atrophia bulborum hereditaria']

NM_000133.3(F9):c.278A>G (p.Asp93Gly)	137852230	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_000133.3(F9):c.329A>G (p.Asp110Gly)	137852234	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_000133.3(F9):c.917A>G (p.Asn306Ser)	137852251	F9	[]	['GCTGCAYTGTAGTTGTGGTGAGG']	['Hereditary factor IX deficiency disease']
NM_000133.3(F9):c.1180A>G (p.Met394Val)	137852262	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_000133.3(F9):c.1231A>G (p.Ser411Gly)	137852277	F9	[]	[]	['Hereditary factor IX deficiency disease']
NM_000292.2(PHKA2):c.896A>G (p.Asp299Gly)	137852289	PHKA2	[]	[]	['Glycogen storage disease type IXa1']
NM_000292.2(PHKA2):c.565A>G (p.Lys189Glu)	137852295	PHKA2	[]	[]	['Glycogen storage disease IXa2']
NM_000032.4(ALAS2):c.1702A>G (p.Ser568Gly)	137852306	ALAS2	[]	[]	['Hereditary sideroblastic anemia']
NM_001287223.1(SCN11A):c.3473T>C (p.Leu1158Pro)	141686175	SCN11A	[]	['CGTGCGCYGTCCCAGTTTGAAGG']	['Episodic pain syndrome, familial, 3']
NM_000402.4(G6PD):c.583A>G (p.Asn195Asp)	137852331	G6PD	[]	['ATGCGGTYCCAGCCTCTGCTGGG']	['Favism, susceptibility to', 'Anemia, nonspherocytic hemolytic, due to G6PD deficiency']
NM_000132.3(F8):c.872A>G (p.Glu291Gly)	137852359	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_018718.2(CEP41):c.107T>C (p.Met36Thr)	368178632	CEP41	[]	[]	['Joubert syndrome 9/15, digenic']
NM_000132.3(F8):c.5183A>G (p.Tyr1728Cys)	137852362	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.5821A>G (p.Asn1941Asp)	137852369	F8	[]	['TAGCCATYGATTGCTGGAGAAGG']	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.328A>G (p.Met110Val)	137852385	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.398A>G (p.Tyr133Cys)	137852389	F8	[]	['TCAYATTCAGCTCCTATAGCAGG']	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.404A>G (p.Asp135Gly)	137852390	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.940A>G (p.Thr314Ala)	137852406	F8	[]	['TGAGCAGYAAGGAAAGTTATTGG']	['Hereditary factor VIII deficiency disease']
NM_000041.3(APOE):c.178A>G	28931576	APOE	[]	['ACAGTGYCTGCACCCAGCGCAGG']	[]

(p.Thr60Ala)					
NM_000132.3(F8):c.1682A>G (p.Asp561Gly)	137852420	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.1892A>G (p.Asn631Ser)	137852429	F8	['ATGYTGGAGGCTTGGAACTCTGG']	['ATGYTGGAGGCTTGGAACTCTGG']	['Hereditary factor VIII deficiency disease']
NM_012082.3(ZFPM2):c.1969A>G (p.Ser657Gly)	28374544	-	[]	[]	['Tetralogy of Fallot']
NM_000098.2(CPT2):c.638A>G (p.Asp213Gly)	74315300	CPT2	[]	[]	['CARNITINE PALMITOYLTRANSFERASE II DEFICIENCY, LATE-ONSET']
NM_000396.3(CTSK):c.990A>G (p.Ter330Trp)	74315301	CTSK	[]	['GAGYCACATCTTGGGGAAGCTGG']	['Pyknodysostosis']
NM_000132.3(F8):c.6113A>G (p.Asn2038Ser)	137852454	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000132.3(F8):c.6278A>G (p.Asp2093Gly)	137852457	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_024009.2(GJB3):c.421A>G (p.Ile141Val)	74315320	GJB3	['CAAYGATGAGCTTGAAGATGAGG']	['CAAYGATGAGCTTGAAGATGAGG']	['Deafness, autosomal recessive']
NM_000132.3(F8):c.104A>G (p.Tyr35Cys)	137852476	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_000194.2(HPRT1):c.602A>G (p.Asp201Gly)	137852479	HPRT1	[]	[]	['Partial hypoxanthine-guanine phosphoribosyltransferase deficiency']
NM_000261.1(MYOC):c.1267A>G (p.Lys423Glu)	74315336	MYOC	[]	[]	['Primary open angle glaucoma juvenile onset 1']
NM_014625.3(NPHS2):c.479A>G (p.Asp160Gly)	74315346	NPHS2	[]	[]	['Nephrotic syndrome, idiopathic, steroid-resistant']
NM_000194.2(HPRT1):c.155A>G (p.Asp52Gly)	137852502	HPRT1	[]	[]	['Partial hypoxanthine-guanine phosphoribosyltransferase deficiency']
NM_002764.3(PRPS1):c.341A>G (p.Asn114Ser)	137852540	PRPS1	[]	['TAGCATAYTTGCAACAAGCTTGG']	['Phosphoribosylpyrophosphate synthetase superactivity']
NM_000055.2(BCHE):c.293A>G (p.Asp98Gly)	1799807	BCHE	[]	[]	['Postanesthetic apnea']
NM_170784.2(MKKS):c.169A>G (p.Thr57Ala)	74315399	MKKS	[]	[]	['Bardet-Biedl syndrome 6']

NM_000311.3(PRNP):c.650A>G (p.Gln217Arg)	74315406	PRNP	[]	[]	['Gerstmann-Straussler-Scheinker syndrome', 'Genetic prion diseases']
NM_000311.3(PRNP):c.560A>G (p.His187Arg)	74315413	PRNP	[]	[]	['Gerstmann-Straussler-Scheinker syndrome', 'Genetic prion diseases', 'Spongiform encephalopathy with neuropsychiatric features']
NM_000044.3(AR):c.2291A>G (p.Tyr764Cys)	137852567	AR	[]	[]	['Reifenstein syndrome']
NM_000044.3(AR):c.2362A>G (p.Met788Val)	137852570	AR	[]	[]	[]
NM_000044.3(AR):c.2632A>G (p.Thr878Ala)	137852578	AR	[]	[]	['Malignant tumor of prostate']
NM_020436.3(SALL4):c.2663A>G (p.His888Arg)	74315429	SALL4	[]	[]	['Duane-radial ray syndrome']
NM_000044.3(AR):c.2708A>G (p.Gln903Arg)	137852582	AR	[]	[]	['Malignant tumor of prostate']
NM_000211.4(ITGB2):c.1052A>G (p.Asn351Ser)	137852613	ITGB2	[]	[]	['Leukocyte adhesion deficiency']
NM_000215.3(JAK3):c.299A>G (p.Tyr100Cys)	137852624	JAK3	[]	['AATCCTGYACAGCAGGACTTGGG']	['Severe combined immunodeficiency, autosomal recessive, T cell-negative, B cell-positive, NK cell-negative']
NM_001166107.1(HMGCS2):c.500A>G (p.Tyr167Cys)	137852640	HMGCS2	[]	['ACCACCGYAGCAGGCATTGGTGG']	['mitochondrial 3-hydroxy-3-methylglutaryl-CoA synthase deficiency']
NM_002047.2(GARS):c.374A>G (p.Glu125Gly)	137852645	GARS	[]	[]	['Charcot-Marie-Tooth disease type 2D', 'Distal hereditary motor neuronopathy type 5']
NM_033163.3(FGF8):c.298A>G (p.Lys100Glu)	137852662	FGF8	[]	[]	['Kallmann syndrome 6']
NM_002180.2(IGHMBP2):c.638A>G (p.His213Arg)	137852666	IGHMBP2	[]	[]	['Werdnig-Hoffmann disease']
NM_004387.3(NKX2-5):c.896A>G (p.Asp299Gly)	137852683	NKX2-5	[]	[]	['Atrial septal defect 7 with or without atrioventricular conduction defects']
NM_004387.3(NKX2-5):c.547A>G (p.Lys183Glu)	137852686	NKX2-5	[]	[]	[]

NM_000310.3(PPT1):c.236A>G (p.Asp79Gly)	137852697	PPT1	[]	[]	['Ceroid lipofuscinosis neuronal 1']
NM_000336.2(SCNN1B):c.863A>G (p.Asn288Ser)	137852712	SCNN1B	[]	[]	['Bronchiectasis']
NM_000579.3(CCR5):c.-301+246A>G	1799987	-	[]	[]	['Human immunodeficiency virus type 1, susceptibility to']
NM_001204.6(BMPR2):c.1454A>G (p.Asp485Gly)	137852745	BMPR2	[]	[]	['Primary pulmonary hypertension']
NM_005591.3(MRE11A):c.350A>G (p.Asn117Ser)	137852760	MRE11A	[]	[]	['Hereditary cancer-predisposing syndrome', 'Ataxia-telangiectasia-like disorder']
NM_003476.4(CSRP3):c.206A>G (p.Lys69Arg)	137852764	CSRP3	[]	[]	['Dilated cardiomyopathy 1M', 'Cardiomyopathy', 'Familial hypertrophic cardiomyopathy 12']
NM_000519.3(HBD):c.-81A>G	35518301	HBD	[]	[]	[]
NM_005633.3(SOS1):c.1654A>G (p.Arg552Gly)	137852814	SOS1	[]	['GCATCCYTTCCAGTGTACTCCGG']	['Noonan syndrome', 'Noonan syndrome 4', 'Rasopathy', 'not provided']
NM_003688.3(CASK):c.2129A>G (p.Asp710Gly)	137852818	CASK	[]	[]	['FG syndrome 4']
NM_031443.3(CCM2):c.1A>G (p.Met1Val)	137852842	CCM2	[]	[]	['Cerebral cavernous malformations 2']
NM_182760.3(SUMF1):c.1A>G (p.Met1Val)	137852855	SUMF1	[]	[]	['Multiple sulfatase deficiency']
NM_001171993.1(HPD):c.362A>G (p.Tyr121Cys)	137852865	HPD	[]	['CCTCAYATCCAGGCAAGAATTGG']	['4-Hydroxyphenylpyruvate dioxygenase deficiency']
NM_024996.5(GFM1):c.521A>G (p.Asn174Ser)	119470018	GFM1	['TTGYTAATAAAAGTTAGAAACGG']	['TTGYTAATAAAAGTTAGAAACGG']	['Combined oxidative phosphorylation deficiency 1']
NM_000158.3(GBE1):c.1634A>G (p.His545Arg)	137852889	GBE1	[]	[]	['Glycogen storage disease, type IV', 'GLYCOGEN STORAGE DISEASE IV, FATAL PERINATAL NEUROMUSCULAR']
NM_000158.3(GBE1):c.1883A>G (p.His628Arg)	137852891	GBE1	[]	[]	['Glycogen storage disease, type IV', 'GLYCOGEN STORAGE DISEASE IV,

					CHILDHOOD NEUROMUSCULAR']
m.8344A>G	118192098	MT-TK	[]	[]	['Parkinson disease, mitochondrial', 'Leigh disease', 'Myoclonus with epilepsy with ragged red fibers']
NM_000540.2(RYR1):c.10100A>G (p.Lys3367Arg)	118192126	RYR1	[]	[]	['Central core disease', 'not provided']
NM_000540.2(RYR1):c.14572A>G (p.Asn4858Asp)	118192144	RYR1	[]	[]	['Central core disease', 'not provided']
NM_012464.4(TLL1):c.1885A>G (p.Ile629Val)	137852953	TLL1	['GGTTAYGGTGCCGTTAAGTTTGG']	['GGTTAYGGTGCCGTTAAGTTTGG']	['Atrial septal defect 6']
NM_025243.3(SLC19A3):c.130A>G (p.Lys44Glu)	137852957	SLC19A3	[]	[]	['Basal ganglia disease, biotin-responsive']
NM_138691.2(TMC1):c.1763+3A>G	370898981	TMC1	[]	['TGGCCYACCAGATCATGCCTTGG']	['Deafness, autosomal recessive 7']
NM_000540.2(RYR1):c.13909A>G (p.Thr4637Ala)	118192166	RYR1	[]	[]	['Central core disease', 'not provided']
NM_000540.2(RYR1):c.14387A>G (p.Tyr4796Cys)	118192167	RYR1	[]	['CCATAYACCAGCCCAGGTACAGG']	['Malignant hyperthermia susceptibility type 1', 'Central core disease', 'not provided']
NM_032667.6(BSCL2):c.263A>G (p.Asn88Ser)	137852972	-	[]	['CGAGACAYTGGCAACAGGGAAGG']	['Distal hereditary motor neuronopathy type 5', 'Silver spastic paraplegia syndrome', 'Charcot-Marie-Tooth disease, type 2']
NM_014795.3(ZEB2):c.3356A>G (p.Gln1119Arg)	137852983	ZEB2	[]	[]	['Mowat-Wilson syndrome']
NM_000540.2(RYR1):c.14740A>G (p.Arg4914Gly)	118192184	RYR1	[]	[]	['Central core disease', 'not provided']
NM_172107.2(KCNQ2):c.356A>G (p.Glu119Gly)	118192193	KCNQ2	[]	['CTTCYCATACTCCTTGATGGTGG', 'GCTCTTCYCATACTCCTTGATGG']	['Benign familial neonatal seizures 1']
NM_172107.2(KCNQ2):c.622A>G (p.Met208Val)	118192201	KCNQ2	[]	['GGATCAYCCGCAGAATCTGCAGG']	['Benign familial neonatal seizures 1']
NM_172107.2(KCNQ2):c.773A>G (p.Asn258Ser)	118192207	KCNQ2	[]	[]	['Benign familial neonatal seizures 1']
NM_004006.2(DMD):c.8734A>G	1800278	DMD	[]	[]	['Duchenne muscular dystrophy',

(p.Asn2912Asp)					'not specified']
NM_004006.2(DMD):c.8762A>G (p.His2921Arg)	1800279	DMD	[]	[]	['Becker muscular dystrophy', 'not specified']
NM_001080463.1(DYNC2H1):c.11284A>G (p.Met3762Val)	137853026	DYNC2H1	[]	[]	['Short-rib thoracic dysplasia 3 with or without polydactyly']
NM_001080463.1(DYNC2H1):c.9044A>G (p.Asp3015Gly)	137853027	DYNC2H1	[]	['ATAYCTCTAATTACATCAGGTGG', 'AGAATAYCTCTAATTACATCAGG']	['Short-rib thoracic dysplasia 3 with or without polydactyly']
NM_001080463.1(DYNC2H1):c.4610A>G (p.Gln1537Arg)	137853033	DYNC2H1	['ACCYGTGAAGGGAACAGAGATGG']	['ACCYGTGAAGGGAACAGAGATGG']	['Short-rib thoracic dysplasia 3 with or without polydactyly']
NM_001080463.1(DYNC2H1):c.5959A>G (p.Thr1987Ala)	137853035	DYNC2H1	[]	[]	['Short-rib thoracic dysplasia 3 with or without polydactyly']
NM_001430.4(EPAS1):c.1603A>G (p.Met535Val)	137853037	EPAS1	[]	[]	['Erythrocytosis, familial, 4']
NM_172107.2(KCNQ2):c.1764-2A>G	118192238	KCNQ2	[]	[]	['Benign familial neonatal seizures 1']
NM_004519.3(KCNQ3):c.914A>G (p.Asp305Gly)	118192248	KCNQ3	[]	[]	['Benign familial neonatal seizures 2']
NM_004519.3(KCNQ3):c.1403A>G (p.Asn468Ser)	118192252	KCNQ3	['TCTTTAYTGTTTAAGCCAACAGG']	['TCTTTAYTGTTTAAGCCAACAGG']	['Benign familial neonatal seizures 2', 'not specified', 'not provided']
NM_004519.3(KCNQ3):c.2462A>G (p.Asn821Ser)	118192254	KCNQ3	[]	[]	['Benign familial neonatal seizures 2', 'not specified', 'not provided']
NM_138701.3(MPLKIP):c.430A>G (p.Met144Val)	137853117	MPLKIP	[]	[]	['Trichothiodystrophy, nonphotosensitive 1']
NM_005094.3(SLC27A4):c.899A>G (p.Gln300Arg)	137853134	SLC27A4	[]	[]	['Ichthyosis prematurity syndrome']
NM_194456.1(KRIT1):c.410A>G (p.Asp137Gly)	137853139	KRIT1	[]	[]	['Cerebral cavernous malformations 1']
NM_000351.4(STS):c.1331A>G (p.His444Arg)	137853169	STS	[]	[]	['X-linked ichthyosis with steryl-sulfatase deficiency']
NM_152416.3(NDUFAF6):c.296A>G (p.Gln99Arg)	137853184	NDUFAF6	[]	[]	['Leigh syndrome due to mitochondrial complex I deficiency']
NM_144573.3(NEXN):c.1955A>G (p.Tyr652Cys)	137853197	NEXN	[]	['ATAYACTCTCCTCCATCTTCTGG']	['Dilated cardiomyopathy 1CC', 'Cardiomyopathy', 'not specified']
NM_000476.2(AK1):c.491A>G	137853203	AK1	[]	['TTCTCAYAGAAGGCGATGACGGG',	['Adenylate kinase deficiency,

(p.Tyr164Cys)				'TTTCTCAYAGAAGGCGATGACGG']	hemolytic anemia due to']
NM_013411.4(AK2):c.1A>G (p.Met1Val)	137853206	AK2	[]	[]	['Reticular dysgenesis']
NM_000308.2(CTSA):c.746+3A>G	786200859	CTSA	[]	['TCCCAYACCTGTTCCCCAGAAGG']	['Galactosialidosis, adult']
NM_002890.2(RASA1):c.1198A>G (p.Lys400Glu)	137853215	RASA1	[]	[]	[]
NM_002890.2(RASA1):c.1201A>G (p.Ile401Val)	137853216	RASA1	[]	[]	[]
NM_000515.4(GH1):c.413A>G (p.Asp138Gly)	137853221	GH1	[]	[]	['Kowarski syndrome']
NM_017636.3(TRPM4):c.2741A>G (p.Lys914Arg)	172151858	TRPM4	[]	[]	['Progressive familial heart block type 1B']
NM_000545.6(HNF1A):c.365A>G (p.Tyr122Cys)	137853237	HNF1A	[]	[]	['Maturity-onset diabetes of the young, type 3']
NM_003494.3(DYSF):c.1285-2A>G	786200897	DYSF	[]	['CAGCYAGAAGACACAGGGAGGGG', 'ACAGCYAGAAGACACAGGGAGGG', 'CACAGCYAGAAGACACAGGGAGG']	['Limb-girdle muscular dystrophy, type 2B']
NM_005055.4(RAPSN):c.-210A>G	786200905	RAPSN	[]	[]	['MYASTHENIC SYNDROME, CONGENITAL, 11, ASSOCIATED WITH ACETYLCHOLINE RECEPTOR DEFICIENCY']
NM_000276.3(OCRL):c.1436A>G (p.Tyr479Cys)	137853262	OCRL	[]	[]	['Dent disease 2']
NM_004463.2(FGD1):c.1396A>G (p.Met466Val)	137853267	FGD1	[]	[]	['Aarskog syndrome']
NM_153252.4(BRWD3):c.4786A>G (p.Lys1596Glu)	137853272	BRWD3	[]	[]	['Mental retardation, X-linked 93']
NM_206933.2(USH2A):c.7595-2144A>G	786200928	USH2A	[]	['CTCTTAYCTTGGGAAAGGAGAGG']	['Usher syndrome, type 2A']
NM_000362.4(TIMP3):c.572A>G (p.Tyr191Cys)	137853299	-	['TGCAGYAGCCGCCCTTCTGCCGG']	['TGCAGYAGCCGCCCTTCTGCCGG']	['Sorsby fundus dystrophy']
NM_006785.3(MALT1):c.1019-2A>G	786200953	MALT1	['CGCYTTGAAAAAAAAAGAAAGGG']	['CGCYTTGAAAAAAAAGAAAGGG', 'TCGCYTTGAAAAAAAAGAAAGG']	['Combined immunodeficiency']
NM_003639.4(IKBKG):c.1259A>G (p.Ter420Trp)	137853321	IKBKG	[]	[]	['Incontinentia pigmenti syndrome', 'Ectodermal

					dysplasia, anhidrotic, with immunodeficiency, osteopetrosis, and lymphedema']
NM_003639.4(IKBKG):c.1219A>G (p.Met407Val)	137853322	IKBKG	[]	['CCAYATCAGGGGCCTGATACTGG']	['Incontinentia pigmenti syndrome']
NM_001014797.2(KCNMA1):c.1301A>G (p.Asp434Gly)	137853333	KCNMA1	[]	[]	['Generalized epilepsy and paroxysmal dyskinesia']
NM_016218.2(POLK):c.1679A>T (p.Glu560Val)	757103131	POLK	[]	[]	['Malignant tumor of prostate']
NM_003000.2(SDHB):c.541-2A>G	786201161	SDHB	[]	[]	['Hereditary cancer-predisposing syndrome']
NM_032383.4(HPS3):c.2482-2A>G	397507168	-	[]	[]	['Hermansky-Pudlak syndrome 3']
NM_000060.3(BTD):c.968A>G (p.His323Arg)	397507176	BTD	[]	[]	['Biotinidase deficiency', 'not provided']
NM_004315.4(ASAH1):c.155A>G (p.Tyr52Cys)	137853595	ASAH1	[]	[]	['Farber lipogranulomatosis']
NM_004315.4(ASAH1):c.1006A>G (p.Asn336Asp)	137853596	ASAH1	[]	[]	['Farber lipogranulomatosis']
NM_000059.3(BRCA2):c.1799A>G (p.Tyr600Cys)	397507276	BRCA2	[]	[]	['Breast-ovarian cancer, familial 2']
NM_022912.2(REEP1):c.183-2A>G	387906264	REEP1	[]	[]	['Spastic paraplegia 31, autosomal dominant']
NM_000022.2(ADA):c.219-2A>G	387906267	ADA	[]	['CCCCYGGGAAGGGAAGAAAGGGG', 'GCCCCYGGGAAGGGAAGAAAGGG', 'AGCCCCYGGGAAGGGAAGAAAGG']	['Severe combined immunodeficiency due to ADA deficiency']
NM_018249.5(CDK5RAP2):c.4005-15A>G	387906274	CDK5RAP2	[]	[]	['Primary autosomal recessive microcephaly 3']
NM_032119.3(ADGRV1):c.14973-2A>G	371981035	ADGRV1	[]	[]	['Usher syndrome, type 2C']
NM_020366.3(RPGRIP1):c.3341A>G (p.Asp1114Gly)	17103671	RPGRIP1	[]	[]	['Leber congenital amaurosis 6', 'not specified']
NM_000492.3(CFTR):c.3717+4A>G	387906362	CFTR	[]	['TCAAATCYCACCCTCTGGCCAGG']	['Cystic fibrosis']
NM_000492.3(CFTR):c.273+4A>G	387906374	CFTR	[]	[]	['Cystic fibrosis']
NM_002769.4(PRSS1):c.65A>G (p.Asp22Gly)	397507442	-	[]	['CTTGYCATCATCATCAAAGGGGG', 'TCTTGYCATCATCATCAAAGGGG', 'ATCTTGYCATCA	['Hereditary pancreatitis']

				TCATCAAAGGG', 'GATCTTGYCATC ATCATCAAAGG']	
NM_004006.2(DMD): c.4675-2A>G	794727575	DMD	[]	[]	['Duchenne muscular dystrophy', 'Becker muscular dystrophy']
NM_000132.3(F8):c.1 418A>G (p.Tyr473Cys)	387906444	F8	[]	[]	['Hereditary factor VIII deficiency disease']
NM_004333.4(BRAF): c.2126A>G (p.Gln709Arg)	397507486	BRAF	[]	[]	['Rasopathy']
NM_002834.3(PTPN1 1):c.124A>G (p.Thr42Ala)	397507501	PTPN11	[]	[]	['Noonan syndrome', 'Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_000834.3(GRIN2 B):c.1238A>G (p.Glu413Gly)	527236034	GRIN2B	[]	[]	['Mental retardation, autosomal dominant 6']
NM_004830.3(MED23):c.3638A>G (p.His1213Arg)	527236035	MED23	[]	[]	['Mental retardation, autosomal recessive 18']
NM_002834.3(PTPN1 1):c.844A>G (p.Ile282Val)	397507529	PTPN11	[]	[]	['Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_000406.2(GNRH R):c.851A>G (p.Tyr284Cys)	28933074	GNRHR	[]	[]	[]
NM_002834.3(PTPN1 1):c.1510A>G (p.Met504Val)	397507547	PTPN11	[]	[]	['Noonan syndrome', 'Noonan syndrome 1', 'Rasopathy', 'not provided']
NM_004629.1(FANC G):c.925-2A>G	397507561	FANCG	[]	[]	['Fanconi anemia, complementation group G']
NM_000492.3(CFTR): c.3140-26A>G	76151804	CFTR	[]	[]	['Cystic fibrosis']
NM_024598.3(USB1): c.502A>G (p.Arg168Gly)	137853971	USB1	[]	['CCACCYGGTTTT CTCTTGATTGG']	['Poikiloderma with neutropenia']
NM_000067.2(CA2):c. 754A>G (p.Asn252Asp)	2228063	CA2	[]	['TGTYCTTCAGTG GCTGAGCTGGG', 'CTGTYCTTCAGT GGCTGAGCTGG']	[]
NM_000138.4(FBN1): c.5096A>G (p.Tyr1699Cys)	387906622	FBN1	[]	[]	['Geleophysic dysplasia 2']
NM_001194958.2(KC NJ18):c.1097A>G (p.Lys366Arg)	527236159	KCNJ18	[]	[]	['Thyrotoxic periodic paralysis', 'Thyrotoxic periodic paralysis 2']
NM_000138.4(FBN1): c.5087A>G (p.Tyr1696Cys)	387906625	FBN1	[]	[]	['Geleophysic dysplasia 2']
NM_000138.4(FBN1): c.5099A>G (p.Tyr1700Cys)	387906626	FBN1	[]	[]	[]
NM_001244710.1(GF	387906638	GFPT1	[]	[]	['Congenital

PT1):c.43A>G (p.Thr15Ala)					myasthenic syndrome with tubular aggregates 1']
NM_002292.3(LAMB2):c.440A>G (p.His147Arg)	387906644	LAMB2	[]	[]	['Nephrotic syndrome, type 5, with or without ocular abnormalities']
NM_005188.3(CBL):c.1112A>G (p.Tyr371Cys)	387906666	CBL	[]	[]	[]
NM_000313.3(PROS1):c.701A>G (p.Tyr234Cys)	387906675	PROS1	['GATTAYATCTGTAGCCTTCGGGG', 'AGATTAYATCTGTAGCCTTCGGG', 'GAGATTAYATCTGTAGCCTTCGG']	['GATTAYATCTGTAGCCTTCGGGG', 'AGATTAYATCTGTAGCCTTCGGG', 'GAGATTAYATCTGTAGCCTTCGG']	['Thrombophilia due to protein S deficiency, autosomal recessive']
NM_032018.6(SPRTN):c.350A>G (p.Tyr117Cys)	527236213	SPRTN	[]	[]	['Ruijs-aalfs syndrome']
NM_022464.4(SIL1):c.645+2T>C	548535414	SIL1	[]	[]	['Marinesco-Sj\xc3\xb6gren syndrome']
NM_001040142.1(SCN2A):c.4419A>G (p.Ile1473Met)	387906685	SCN2A	[]	[]	['Early infantile epileptic encephalopathy 11']
NM_001040142.1(SCN2A):c.754A>G (p.Met252Val)	387906687	SCN2A	[]	[]	['Benign familial neonatal-infantile seizures']
m.10450A>G	387906731	MT-TR	[]	[]	['Mitochondrial encephalomyopathy']
m.5816A>G	387906732	MT-TC	[]	[]	[]
m.608A>G	387906735	MT-TF	['TTCAGYGTATTGCTTTGAGGAGG']	['TTCAGYGTATTGCTTTGAGGAGG']	[]
NM_001376.4(DYNC1H1):c.917A>G (p.His306Arg)	387906738	DYNC1H1	[]	[]	['Charcot-Marie-Tooth disease, axonal, type 2O', 'Charcot-Marie-Tooth disease', 'Spinal muscular atrophy, lower extremity predominant 1, autosomal dominant']
NM_001376.4(DYNC1H1):c.2011A>G (p.Lys671Glu)	387906742	DYNC1H1	[]	[]	['Spinal muscular atrophy, lower extremity predominant 1,

					autosomal dominant']
NM_001376.4(DYNC1H1):c.2909A>G (p.Tyr970Cys)	387906743	DYNC1H1	[]	['ATTCAAGYAGATTACCTGATTGG']	['Spinal muscular atrophy, lower extremity predominant 1, autosomal dominant']
NM_001354.5(AKR1C2):c.235A>G (p.Ile79Val)	387906750	AKR1C2	[]	[]	['46,XY sex reversal 8']
NM_007315.3(STAT1):c.604A>G (p.Met202Val)	387906762	STAT1	[]	[]	['Immunodeficiency 31C']
NM_007315.3(STAT1):c.494A>G (p.Asp165Gly)	387906764	STAT1	[]	[]	['Immunodeficiency 31C']
NM_007315.3(STAT1):c.862A>G (p.Thr288Ala)	387906765	STAT1	[]	[]	['Immunodeficiency 31C']
NM_002052.4(GATA4):c.928A>G (p.Met310Val)	387906772	GATA4	[]	['TCCGCAYTGCAAGAGGCCTGGGG', 'TTCCGCAYTGCAAGAGGCCTGGG']	['Atrial septal defect 2']
NM_021615.4(CHST6):c.329A>G (p.Tyr110Cys)	72547544	CHST6	[]	[]	['Macular corneal dystrophy Type I']
NM_000209.3(PDX1):c.533A>G (p.Glu178Gly)	387906777	PDX1	[]	[]	['Pancreatic agenesis, congenital']
NM_000890.3(KCNJ5):c.472A>G (p.Thr158Ala)	387906778	KCNJ5	[]	[]	['Andersen Tawil syndrome', 'Familial hyperaldosteronism type 3']
NM_001184.3(ATR):c.6431A>G (p.Gln2144Arg)	387906797	ATR	[]	[]	['Cutaneous telangiectasia and cancer syndrome, familial']
NM_000382.2(ALDH3A2):c.1157A>G (p.Asn386Ser)	72547575	ALDH3A2	[]	[]	['Sj\xc3\xb6gren-Larsson syndrome']
NM_001005862.2(ERBB2):c.2480A>G (p.Asn827Ser)	28933370	ERBB2	[]	[]	['Neoplasm of ovary']
NM_006194.3(PAX9):c.271A>G (p.Lys91Glu)	28933373	PAX9	[]	[]	['Tooth agenesis, selective, 3']
NM_001083116.1(PRF1):c.755A>G (p.Asn252Ser)	28933375	PRF1	[]	[]	['Hemophagocytic lymphohistiocytosis, familial, 2', 'Malignant lymphoma, non-Hodgkin']
NM_005257.5(GATA6):c.1354A>G (p.Thr452Ala)	387906817	GATA6	[]	[]	['Pancreatic agenesis and congenital heart disease']
NM_000414.3(HSD17B4):c.650A>G (p.Tyr217Cys)	387906825	HSD17B4	[]	['TGCCACAYACTCTGGCTTCAGGG']	['Gonadal dysgenesis with auditory

					dysfunction, autosomal recessive inheritance']
NM_004153.3(ORC1): c.380A>G (p.Glu127Gly)	387906826	ORC1	[]	[]	['Meier-Gorlin syndrome 1']
NM_002552.4(ORC4): c.521A>G (p.Tyr174Cys)	387906847	ORC4	[]	[]	['Meier-Gorlin syndrome 2']
NM_004544.3(NDUFA10):c.1A>G (p.Met1Val)	387906872	NDUFA10	[]	[]	['Leigh syndrome due to mitochondrial complex I deficiency']
NM_004544.3(NDUFA10):c.425A>G (p.Gln142Arg)	387906873	NDUFA10	[]	[]	['Leigh syndrome due to mitochondrial complex I deficiency']
NM_006796.2(AFG3L2):c.1847A>G (p.Tyr616Cys)	387906889	AFG3L2	['GTAYAGAGGTATTGTTCTTTTGG']	['GTAYAGAGGTATTGTTCTTTTGG']	['Spastic ataxia 5, autosomal recessive']
NM_006587.3(CORIN):c.949A>G (p.Lys317Glu)	387906894	CORIN	[]	[]	['Preeclampsia/eclampsia 5']
NM_006587.3(CORIN):c.1414A>G (p.Ser472Gly)	387906895	CORIN	[]	['GGATAACYTGTACTGTTGTAGGG']	['Preeclampsia/eclampsia 5']
NM_015560.2(OPA1): c.1294A>G (p.Ile432Val)	387906899	OPA1	[]	[]	['Optic Atrophy Type 1']
NM_021625.4(TRPV4):c.590A>G (p.Lys197Arg)	387906903	TRPV4	[]	[]	['Metatrophic dysplasia']
NM_021625.4(TRPV4):c.826A>G (p.Lys276Glu)	387906907	TRPV4	[]	[]	['Metatrophic dysplasia']
NM_024022.2(TMPRSS3):c.308A>G (p.Asp103Gly)	387906915	TMPRSS3	[]	[]	['Deafness, autosomal recessive 8']
NM_019109.4(ALG1): c.1129A>G (p.Met377Val)	387906925	ALG1	[]	[]	['Congenital disorder of glycosylation type 1K']
NM_006886.3(ATP5E):c.35A>G (p.Tyr12Cys)	387906929	-	[]	[]	['Nuclearly-encoded mitochondrial complex V (ATP synthase) deficiency 3']
NM_032578.3(MYPN):c.59A>G (p.Tyr20Cys)	140148105	MYPN	[]	[]	['Primary dilated cardiomyopathy', 'Cardiomyopathy', 'Dilated cardiomyopathy 1KK', 'Familial hypertrophic cardiomyopathy 22', 'not provided']
NM_016013.3(NDUFAF1):c.758A>G (p.Lys253Arg)	387906957	NDUFAF1	[]	['ACCYTGACCTCCTGCCAGTAGGG', 'TACCYTGACCTC	['Mitochondrial complex I deficiency']

				CTGCCAGTAGG']	
NM_032580.3(HES7): c.172A>G (p.Ile58Val)	387906979	HES7	[]	[]	['Spondylocostal dysostosis 5']
NM_024700.3(SNIP1) :c.1097A>G (p.Glu366Gly)	387906986	SNIP1	[]	[]	['Psychomotor retardation, epilepsy, and craniofacial dysmorphism']
NM_016952.4(CDON) :c.2368A>G (p.Thr790Ala)	387906997	CDON	[]	[]	['Holoprosencephaly 11']
NM_031427.3(DNAL 1):c.449A>G (p.Asn150Ser)	387907021	DNAL1	['AGGGAYT GCCTACAA ACACCAGG ']	['AGGGAYTGCCT ACAAACACCAGG']	['Kartagener syndrome', 'Ciliary dyskinesia, primary, 16']
NM_020320.3(RARS2):c.1024A>G (p.Met342Val)	387907048	RARS2	[]	[]	['Pontocerebellar hypoplasia type 6']
NM_000138.4(FBN1): c.6431A>G (p.Asn2144Ser)	137854461	FBN1	[]	[]	['Marfan syndrome']
NM_000138.4(FBN1): c.2261A>G (p.Tyr754Cys)	137854479	FBN1	[]	[]	['Marfan syndrome']
NM_212482.1(FN1):c. 2918A>G (p.Tyr973Cys)	137854488	FN1	['GAAGTAA YAGGTGAC CCCAGGGG ']	['GAAGTAAYAGG TGACCCCAGGGG']	['Glomerulopathy with fibronectin deposits 2']
NM_198994.2(TGM6) :c.980A>G (p.Asp327Gly)	387907098	TGM6	[]	[]	['Spinocerebellar ataxia 35']
NM_001363.4(DKC1): c.1069A>G (p.Thr357Ala)	137854492	DKC1	['GCAGGYA GAGATGAC CGCTGTGG']	['GCAGGYAGAGA TGACCGCTGTGG']	['Dyskeratosis congenita X-linked']
NM_052873.2(IFT43): c.1A>G (p.Met1Val)	387907107	IFT43	[]	[]	['Cranioectodermal dysplasia 3']
NM_201269.2(ZNF64 4):c.2014A>G (p.Ser672Gly)	387907109	ZNF644	[]	[]	['Myopia 21, autosomal dominant']
NM_018699.3(PRDM 5):c.320A>G (p.Tyr107Cys)	387907111	PRDM5	[]	[]	['Brittle cornea syndrome 2']
NM_000132.3(F8):c.5 822A>G (p.Asn1941Ser)	28933682	F8	[]	['TAGCCAYTGATT GCTGGAGAAGG']	['Hereditary factor VIII deficiency disease']
NM_016464.4(TMEM 138):c.287A>G (p.His96Arg)	387907132	TMEM13 8	['GACAYGA AGGGAGAT GCTGAGGG ']	['GACAYGAAGGG AGATGCTGAGGG', 'AGACAYGAAGGG AGATGCTGAGG']	['Joubert syndrome 16']
NM_016464.4(TMEM 138):c.389A>G (p.Tyr130Cys)	387907135	TMEM13 8	[]	['CAGYACAACAC TGCTGCTGTGGG', 'GCAGYACAACAC TGCTGCTGTGG']	['Joubert syndrome 16']
NM_177965.3(C8orf3 7):c.545A>G (p.Gln182Arg)	387907137	C8orf37	[]	[]	['Retinitis pigmentosa 64']
NM_001077488.3(GN AS):c.1A>G (p.Met1Val)	137854530	GNAS	[]	['GCCCAYGGCGG CGGCGGCGGCGG']	['Pseudohypoparathyroidism type 1A']

NM_006920.4(SCN1A):c.2557-2A>G	727504140	SCN1A	[]	[]	['Severe myoclonic epilepsy in infancy', 'Generalized epilepsy with febrile seizures plus, type 2']
NM_000308.2(CTSA):c.200A>G (p.Gln67Arg)	137854541	CTSA	[]	[]	['Combined deficiency of sialidase AND beta galactosidase']
NM_000308.2(CTSA):c.1238A>G (p.Tyr413Cys)	137854543	CTSA	[]	[]	['Combined deficiency of sialidase AND beta galactosidase']
NM_000308.2(CTSA):c.1411A>G (p.Lys471Glu)	137854549	CTSA	[]	[]	['Galactosialidosis, late infantile']
NM_000267.3(NF1):c.4267A>G (p.Lys1423Glu)	137854550	NF1	[]	[]	['Neurofibromatosis, type 1']
NM_000267.3(NF1):c.1466A>G (p.Tyr489Cys)	137854557	NF1	['ACTTAYAGCTTCTTGTCTCCAGG']	['ACTTAYAGCTTCTTGTCTCCAGG']	['Neurofibromatosis, type 1']
NM_018105.2(THAP1):c.70A>G (p.Lys24Glu)	387907176	THAP1	[]	['CCTCACTYGTGGAAAGAAACGGG']	['Dystonia 6, torsion']
NM_000492.3(CFTR):c.1393-2A>G	397508201	CFTR	[]	[]	['Cystic fibrosis']
NM_001172646.1(PLCB4):c.986A>G (p.Asn329Ser)	387907179	PLCB4	[]	[]	['Auriculocondylar syndrome 2']
NM_005850.4(SF3B4):c.1A>G (p.Met1Val)	387907185	SF3B4	[]	[]	['Nager syndrome']
NM_014714.3(IFT140):c.932A>G (p.Tyr311Cys)	387907193	IFT140	[]	[]	['Renal dysplasia, retinal pigmentary dystrophy, cerebellar ataxia and skeletal dysplasia']
NM_005006.6(NDUFS1):c.1783A>G (p.Thr595Ala)	387907199	NDUFS1	[]	[]	['Mitochondrial complex I deficiency']
NM_000397.3(CYBB):c.1499A>G (p.Asp500Gly)	137854593	CYBB	[]	['TCACAYCTTTCTCCTCATCATGG']	['Chronic granulomatous disease, X-linked', 'not provided']
NM_033360.3(KRAS):c.439A>G (p.Lys147Glu)	387907206	KRAS	[]	[]	['Cardiofaciocutaneous syndrome 2']
NM_000335.4(SCN5A):c.5381A>G (p.Tyr1794Cys)	137854614	SCN5A	[]	[]	['Long QT syndrome 3', 'Congenital long QT syndrome']
NM_000076.2(CDKN1C):c.832A>G (p.Lys278Glu)	387907226	CDKN1C	[]	['CGCTYGGCGAAGAAATCTGCGGG', 'GCGCTYGGCGAAGAAATCTGCGG']	['Intrauterine growth retardation, metaphyseal dysplasia, adrenal hypoplasia congenita, and genital anomalies']

NM_022912.2(REEP1):c.304-2A>G	387907242	REEP1	[]	['TCCYGTCAAAGGAAAAACAGAGG']	['Distal hereditary motor neuronopathy type 5B']
NM_198253.2(TERT):c.2705A>G (p.Lys902Arg)	387907250	TERT	[]	[]	['PULMONARY FIBROSIS AND/OR BONE MARROW FAILURE, TELOMERE-RELATED, 1']
NM_005349.3(RBPJ):c.188A>G (p.Glu63Gly)	387907270	RBPJ	[]	[]	['Adams-Oliver syndrome 3']
NM_005349.3(RBPJ):c.505A>G (p.Lys169Glu)	387907271	RBPJ	[]	[]	['Adams-Oliver syndrome 3']
NM_022787.3(NMNAT1):c.817A>G (p.Asn273Asp)	387907291	NMNAT1	[]	['TGTYTCTCTGCAAAGGGGCCAGG']	['Leber congenital amaurosis 9']
NM_000492.3(CFTR):c.1A>G (p.Met1Val)	397508328	CFTR	['TGCAYGGTCTCTCGGGCGCTGGG']	['GCAYGGTCTCTCGGGCGCTGGGG', 'TGCAYGGTCTCTCGGGCGCTGGG', 'CTGCAYGGTCTCTCGGGCGCTGG']	['Cystic fibrosis']
NM_020921.3(NIN):c.5126A>G (p.Asn1709Ser)	387907308	NIN	[]	[]	['Seckel syndrome 7']
NM_021629.3(GNB4):c.265A>G (p.Lys89Glu)	387907341	GNB4	[]	[]	['Charcot-Marie-Tooth disease, dominant intermediate F']
NM_000355.3(TCN2):c.580+624A>T	372866837	TCN2	[]	[]	[]
NM_032415.5(CARD11):c.401A>G (p.Glu134Gly)	387907351	CARD11	[]	[]	['B-CELL EXPANSION WITH NFKB AND T-CELL ANERGY']
NM_005430.3(WNT1):c.624+4A>G	387907354	WNT1	[]	[]	['Osteogenesis imperfecta type 15']
NM_207352.3(CYP4V2):c.367A>G (p.Met123Val)	149684063	CYP4V2	[]	[]	['Bietti crystalline corneoretinal dystrophy', 'not provided']
NM_031885.3(BBS2):c.472-2A>G	137854887	BBS2	[]	[]	['Bardet-Biedl syndrome 2']
NM_015268.3(DNAJC13):c.2564A>G (p.Asn855Ser)	387907571	DNAJC13	[]	[]	['Parkinson disease, late-onset', 'Essential tremor', 'PARKINSON DISEASE 21']
NM_001287.5(CLCN7):c.296A>G (p.Tyr99Cys)	387907576	CLCN7	[]	['TGTCAYAGTCCAAGCTCTGCAGG']	['Osteopetrosis autosomal dominant type 2', 'Osteopetrosis autosomal recessive 4']

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EQUIVALENTS AND SCOPE

[00416] Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents of the embodiments described herein. The scope of the present disclosure is not intended to be limited to the above description, but rather is as set forth in the appended claims.

[00417] Articles such as "a," "an," and "the" may mean one or more than one unless indicated to the contrary or otherwise evident from the context. Claims or descriptions that include "or" between two or more members of a group are considered satisfied if one, more than one, or all of the group members are present, unless indicated to the contrary or otherwise evident from the context. The disclosure of a group that includes "or" between two or more group members provides embodiments in which exactly one member of the group is present, embodiments in which more than one members of the group are present, and embodiments in which all of the group members are present. For purposes of brevity those embodiments have not been individually spelled out herein, but it will be understood that each of these embodiments is provided herein and may be specifically claimed or disclaimed.

[00418] It is to be understood that the invention encompasses all variations, combinations, and permutations in which one or more limitation, element, clause, or descriptive term, from one or more of the claims or from one or more relevant portion of the description, is introduced into another claim. For example, a claim that is dependent on another claim can be modified to include one or more of the limitations found in any other claim that is

dependent on the same base claim. Furthermore, where the claims recite a composition, it is to be understood that methods of making or using the composition according to any of the methods of making or using disclosed herein or according to methods known in the art, if any, are included, unless otherwise indicated or unless it would be evident to one of ordinary skill in the art that a contradiction or inconsistency would arise.

[00419] Where elements are presented as lists, *e.g.*, in Markush group format, it is to be understood that every possible subgroup of the elements is also disclosed, and that any element or subgroup of elements can be removed from the group. It is also noted that the term "comprising" is intended to be open and permits the inclusion of additional elements or steps. It should be understood that, in general, where an embodiment, product, or method is referred to as comprising particular elements, features, or steps, embodiments, products, or methods that consist, or consist essentially of, such elements, features, or steps, are provided as well. For purposes of brevity those embodiments have not been individually spelled out herein, but it will be understood that each of these embodiments is provided herein and may be specifically claimed or disclaimed.

[00420] Where ranges are given, endpoints are included. Furthermore, it is to be understood that unless otherwise indicated or otherwise evident from the context and/or the understanding of one of ordinary skill in the art, values that are expressed as ranges can assume any specific value within the stated ranges in some embodiments, to the tenth of the unit of the lower limit of the range, unless the context clearly dictates otherwise. For purposes of brevity, the values in each range have not been individually spelled out herein, but it will be understood that each of these values is provided herein and may be specifically claimed or disclaimed. It is also to be understood that unless otherwise indicated or otherwise evident from the context and/or the understanding of one of ordinary skill in the art, values expressed as ranges can assume any subrange within the given range, wherein the endpoints of the subrange are expressed to the same degree of accuracy as the tenth of the unit of the lower limit of the range.

[00421] In addition, it is to be understood that any particular embodiment of the present invention may be explicitly excluded from any one or more of the claims. Where ranges are given, any value within the range may explicitly be excluded from any one or more of the claims. Any embodiment, element, feature, application, or aspect of the compositions and/or methods of the invention, can be excluded from any one or more claims. For purposes of brevity, all of the embodiments in which one or more elements, features, purposes, or aspects is excluded are not set forth explicitly herein.

What is claimed is:

1. A fusion protein comprising:

(i) a Cas9 domain, wherein the Cas9 domain when in conjunction with a bound guide RNA (gRNA) specifically binds to a target nucleic acid sequence;

(ii) a cytidine deaminase domain, wherein the cytidine deaminase domain deaminates a cytosine base in a single-stranded portion of the target nucleic acid sequence when in conjunction with the Cas9 domain and the gRNA; and

(iii) an uracil glycosylase inhibitor (UGI) domain, wherein the UGI domain inhibits a uracil-DNA glycosylase.

2. The fusion protein of claim 1, wherein the Cas9 domain comprises the amino acid sequence provided in SEQ ID NO: 674.

3. The fusion protein of claim 1, wherein the Cas9 domain is a Cas9 nickase (nCas9) domain that cuts a nucleotide target strand of a nucleotide duplex, wherein the nucleotide target strand is the strand that binds to the gRNA.

4. The fusion protein of claim 1, wherein the Cas9 domain is a Cas9 nickase (nCas9) domain that comprises a D10A mutation in the amino acid sequence provided in SEQ ID NO: 10, or a corresponding mutation in any of the amino acid sequences provided in SEQ ID NOs: 11-260.

5. The fusion protein of claim 1, wherein the Cas9 domain is a Cas9 nickase (nCas9) domain that comprises one or more of N497A, R661A, Q695A, or Q926A mutations in the amino acid sequence provided in SEQ ID NO: 10, or one or more corresponding mutations in any of the amino acid sequences provided in SEQ ID NOs: 11-260.

6. The fusion protein of claim 1, wherein the cytidine deaminase domain is a deaminase from an apolipoprotein B mRNA-editing complex (APOBEC) family deaminase.

7. The fusion protein of claim 6, wherein the APOBEC family deaminase is selected from the group consisting of APOBEC1 deaminase, APOBEC2 deaminase, APOBEC3A deaminase, APOBEC3B deaminase, APOBEC3C deaminase, APOBEC3D deaminase,

APOBEC3F deaminase, APOBEC3G deaminase, and APOBEC3H deaminase.

8. The fusion protein of claim 1, wherein the cytidine deaminase domain comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 266-284, 607-610, 5724-5736, and 5738-5741.

9. The fusion protein of claim 1, wherein the cytidine deaminase domain is a rat APOBEC1 (rAPOBEC1) deaminase comprising one or more mutations selected from the group consisting of W90Y, R126E, and R132E in SEQ ID NO: 284, or one or more corresponding mutations in another APOBEC deaminase.

10. The fusion protein of claim 1, wherein the cytidine deaminase domain is a human APOBEC1 (hAPOBEC1) deaminase comprising one or more mutations selected from the group consisting of W90Y, Q126E, and R132E in SEQ ID NO: 5724, or one or more corresponding mutations in another APOBEC deaminase.

11. The fusion protein of claim 1, wherein the cytidine deaminase domain is a human APOBEC3G (hAPOBEC3G) deaminase comprising one or more mutations selected from the group consisting of W285Y, R320E, and R326E in SEQ ID NO: 275, or one or more corresponding mutations in another APOBEC deaminase.

12. The fusion protein of claim 1, wherein the cytidine deaminase domain is an activation-induced deaminase (AID).

13. The fusion protein of claim 1, wherein the cytidine deaminase domain is a cytidine deaminase 1 from *Petromyzon marinus* (pmCDA1).

14. The fusion protein of claim 1, wherein the UGI domain comprises the amino acid sequence as set forth in SEQ ID NO: 600.

15. The fusion protein of claim 1, wherein the fusion protein comprises the structure: NH_2-[cytidine deaminase domain]-[Cas9 domain]-[UGI domain]-COOH, and wherein each instance of "-" comprises an optional linker.

16. The fusion protein of claim 1, wherein the cytidine deaminase domain of (ii) and the Cas9 domain of (i) are linked via a linker comprising the amino acid sequence $(GGGS)_n$(SEQ ID NO: 265)$_n$, $(GGGGS)_n$(SEQ ID NO: 5)$_n$, $(G)_n$, $(EAAAK)_n$(SEQ ID NO: 6)$_n$, $(GGS)_n$, $(SGGS)_n$(SEQ ID NO: 4288)$_n$, SGSETPGTSESATPES (SEQ ID NO: 7)$_n$, or $(XP)_n$motif, or a combination thereof, wherein n is independently an integer between 1 and 30, inclusive, and wherein X is any amino acid.

17. The fusion protein of claim 1, wherein the cytidine deaminase domain of (ii) and the Cas9 domain of (i) are linked via a linker comprising the amino acid sequence: SGSETPGTSESATPES (SEQ ID NO: 7).

18. The fusion protein of claim 1 further comprising a nuclear localization sequence (NLS).

19. The fusion protein of claim 18, wherein the NLS comprises the amino acid sequence PKKKRKV (SEQ ID NO: 741) or MDSLLMNRRKFLYQFKNVRWAKGRRETYLC (SEQ ID NO: 742).

20. The fusion protein of claim 18, wherein the fusion protein comprises the structure: NH_2-[cytidine deaminase domain]-[Cas9 domain]-[UGI domain]-[NLS]-COOH, and wherein each instance of "-" comprises an optional linker.

21. The fusion protein of claim 18, wherein the UGI domain and the NLS are linked via a linker comprising the amino acid sequence: SGGS (SEQ ID NO: 4288), or wherein the Cas9 domain and the UGI domain are linked via a linker comprising the amino acid sequence: SGGS (SEQ ID NO: 4288).

22. The fusion protein of claim 1, wherein the fusion protein comprises the amino acid sequence set forth in SEQ ID NO: 594.

23. A complex comprising the fusion protein of claim 1 and the guide RNA bound to the Cas9 domain of the fusion protein.