

Nitrile hydratases from metagenome libraries

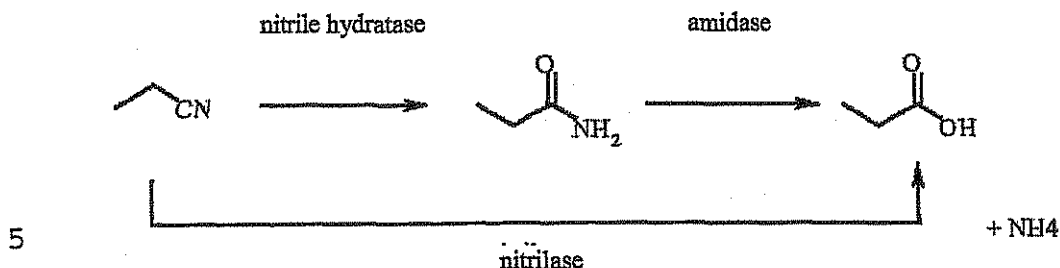
The present invention is directed toward specific degenerate primers. These latter are preferably employed in
5 a process for preparing nitrile hydratases. The present invention therefore also relates to the nitrile hydratases which are prepared by the process which is performed using the primers and to other proteins which are required for the nitrile hydratases to be active. The nucleic acids
10 encoding these protein sequences, and expression systems comprising these nucleic acids, likewise form an additional part of the subject matter of the application. The use of the nitrile hydratases, and of the underlying nucleic acid sequences, constitutes an additional aspect of the present
15 invention.

The amide and carboxylic acid structural classes are becoming ever more important as precursors of fine chemicals. Specific aminoamides and (proteinogenic and
20 nonproteinogenic) amino acids are key intermediates for synthesizing pharmaceutical and agrochemical products as well as in the foodstuffs field. Enantiomerically pure amides and amino acids, in particular, play an ever greater role in the abovementioned areas of application.

25 Aminonitrile precursors, as are required for preparing the abovementioned compound classes, can be readily obtained in racemic form by way of what is termed Strecker synthesis. The nitriles which have been obtained in this way can then
30 be converted into the corresponding amides and carboxylic acids by means of chemical or enzymic hydrolysis.

Three enzymes which are capable of being involved in the enzymic hydrolysis of nitriles are known. Nitrilases
35 convert a nitrile function directly into the acid whereas nitrile hydratases (E.C. 4.2.1.84) in this case form the corresponding amide. This latter can then finally be

converted into the corresponding carboxylic acid by means of an amidase (E.C. 3.5.1.4) (scheme 1).



Scheme 1:

10 Using isolated enzymes or whole-cell catalysts to hydrolyze nitriles to give the corresponding amides and acids helps to save large quantities of salt which would otherwise accrue in the neutralization step following the chemical hydrolysis of nitriles. For this reason, the enzymic hydrolysis of nitriles to give, for example, aminoamides
15 and/or amino acids represents a more sustainable production process.

In their active form, nitrile hydratases consist of 2 nonhomologous, α and β subunits. These latter form
20 heterodimers and tetramers, while even decamers have been demonstrated to be present in the case of *Rhodococcus rhodochromus* J1. While the α and β subunits are of approximately the same size, they do not otherwise share any similarities. Nitrile hydratases are metalloproteins
25 which contain Fe^{3+} or Co^{3+} ions (Bunch A. W. (1998), Nitriles, in: Biotechnology, Volume 8a, Biotransformations I, Chapter 6, Eds.: Rehm HJ, Reed G, Wiley-VCH, p. 277-324; Shearer J, Kung IY, Lovell S, Kaminsky W, Kovacs JA (2001) Why is there an "inert" metal center in the active site of
30 nitrile hydratase? Reactivity and ligand dissociation from a five-coordinate Co(III) nitrile hydratase model. J Am Chem Soc 123: 463-468; Kobayashi M, Shimizu S (2000) Nitrile hydrolases. Current Opinion in Chemical Biology 4: 95-102).

- One of the greatest challenges thus far has been that of heterologously preparing nitrile hydratases in a suitable host, preferably *E. coli*. This Gram-negative bacterium is known to express heterologous proteins at high rates. An additional advantage is the yield of biomass in high cell density fermentations using *E. coli*. In such fermentations, it is possible to achieve productivities of more than 100 g of dry biomass (DBM) in a period of from 24 to 44 hours (Lee SY (1996) High cell-density culture of *Escherichia coli*. TIBTECH 14:98-105; Riesenbergr D, Guthke R (1999) High-cell-density cultivation of microorganisms. Appl Microbiol Biotechnol 51:422-430).
- Most nitrile hydratase α and β subunit sequences are known from the genus *Rhodococcus*. However, it is precisely the nitrile hydratases from this genus which it has thus far only been possible to express in *E. coli* with particular difficulty (Ikehata O, Nishiyama M, Horinouchi S, Beppu T (1989) Primary structure of nitrile hydratase deduced from the nucleotide sequence of a *Rhodococcus* species and its expression in *Escherichia coli*. Eur J Biochem 181: 563-570).
- The literature describes systems for expressing nitrile hydratases whose specific activities are between 4.2 and 12.2 U/mg of total protein in the case of Co-dependent nitrile hydratases from *R. rhodochrous* J1 (Kobayasjhi M, Nishiyama M, Nagasawa T, Horinouchi S, Beppu T, Yamada H (1991) Cloning, nucleotide sequence and expression in *Escherichia coli* of two cobalt-containing nitrole hydratase genes from *Rhodococcus rhodochrous*. Biochim Biophys Acta 1129: 23-33) and 452 U/mg of total protein in the case of an iron-dependent nitrile hydratase from *Rhodococcus spec.* N-771 (Njori M, Yohda M, Odaka M, Matsushita Y, Tsujimura M, Yoshida T, Dohmae N, Takio K Endo I (1999) Functional expression of Nitrile hydratases in *E. coli*: Requirement of

a nitrile hydratase activator and a post-translational modification of a ligand cysteine. J Biochem 125: 696-704), with this corresponding to about approx. 248 U/mg of DBM (dry biomass) (calculation in accordance with Goodsell DS (1991) Inside a cell. TIBS 16: 203-206). Interestingly, it was not possible to reproduce the latter activity using nitrile hydratases from *R. erythropolis*, which is closely related to *Rhodococcus spec. N-711*, and employing similar vector systems and arrangements of the structural genes.

10 There was, therefore, still a need for processes and systems which enable the enzymes in question to be made available in a manner which is adequate for the industrial scale.

15 Methods which have thus far been described for screening for nitrile hydratases have been restricted to isolating microorganisms which exhibit a corresponding enzyme activity. These microorganisms have either been taken from existing strain collections or selectively propagated in

20 what are termed enrichment media (Colquhoun JA, Heald SC, Li L, Tamaoka J, Kato C, Horikoshi K and Bull AT (1998a) Taxonomy and biotransformation activities of some deep-sea actinomycetes. Extremophiles 2: 269-277; Colquhoun JA, Mexson J, Goodfellow M, Ward AC, Horikoshi K and Bull AT (1998b) Novel rhodococci and other mycolate actinomycetes from the deep sea. Antonie van Leeuwenhoek 74: 27-40). A disadvantage of these screening methods is that they have in the main found microorganisms of the genera *Rhodococcus*, *Bacillus* or *Pseudomonas*, or the like, and the diversity of

30 the corresponding nitrile hydratases in regard to activity or substrate specificity is also limited as a consequence (Bunch A. W. (1998), Nitriles, in: Biotechnology, Volume 8a, Biotransformations I, Chapter 6, Eds.: Rehm H.J., Reed G., Wiley-VCH, p. 277-324; Cowan D, Cramp R, Pereira R, Graham D, Almathawa Q (1998) Biochemistry and biotechnology of mesophilic and thermophilic nitrile metabolising enzymes. Extremophiles 2: 207-216; Yamada H, Kobayashi M

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(1996) Nitrile hydratase and its application to industrial production of acrylamide. *Biosci Biotechnol Biochem* 60: 1391-141).

5 Current estimates assume that, as a rule, only 0.01 - 1% of the microorganisms in a habitat can be cultured and thereby made available for screening in accordance with the above-described method (Amann RI, Ludwig W, Schleifer KH (1995) Phylogenetic identification and in situ detection of
10 individual microbial cells without cultivation. *Microbiol. Rev.* 59: 143-169; Pace NR (1997) A molecular view of microbial diversity and the biosphere. *Science* 276: 734-740; Cowan DA (2000) Microbial genomes-the untapped resource. *Trends Biotechnol.* 18: 14-16). "Directly cloning"
15 the genomic DNA of, if at all possible, all the organisms in a soil sample (i.e. the "metagenome"), and making this DNA available, in the form of metagenome gene libraries, for a genetic screening, is therefore increasingly gaining importance for identifying novel industrial enzymes. In
20 this approach, the genetic screening for enzyme-encoding genes can either be effected on the basis of sequence homology, based on conserved sequence motifs, or, when suitable enzyme tests/indicator media are available, on the basis of activity homology (Lorenz P, Köhler B, Wolf M, Eck
25 J, Zinke H (2000) Expression Cloning of Metagenome DNA from Soil. *Biotechnol. 2000, Book of Abstr. Vol 2: 306*).

While nitrile hydratases have already been amplified from metagenomic DNA by means of a PCR-based screening method
30 using degenerate primers, the nature of the sequences is such that they exhibit very high degrees of similarity (90 - 99%) with those for known nitrile hydratases (Precigou S, Goulas P, Duran R, (2001) Rapid and specific identification of nitrile hydratase encoding genes in soil
35 samples by polymerase chain reaction, *FEMS Microbiol. Letters* 204: 155-161). It is not possible to make any proper assessment of the primers which are used in this

reference since the authors do not reveal the sequences of the degenerate primers or their degree of degeneracy. The high degree of similarity of the sequences to those for the *Rhodococcus rhodochrous* J1 nitrile hydratases suggests that
5 the substrate specificity is not markedly different from that of this latter enzyme, either.

There was, therefore, still a need for processes and systems which enable additional enzymes of the type in
10 question to be made available in a manner which is adequate for the industrial scale.

The object of the present invention was therefore to specify an additional process for preparing nitrile
15 hydratases. In particular, the process should be able to identify nitrile hydratases which are present in what are termed nonculturable organisms. An additional object of the present invention was to prepare nitrile hydratases which are superior to those in the prior art.

20 These objects, and additional objects which are not specified in detail but which ensue in an obvious manner from the prior art, are achieved by the specification of specific primer constituents as claimed in claim 1 and
25 their use in a process which has the features of claim 2. Claims 3 to 5 relate to preferred embodiments of the process in question. Claim 6 protects the protein sequences according to the invention while claim 7 is directed toward the nucleic acid sequences which encode them and claim 8
30 relates to the expression systems which are provided with these nucleic acid sequences. Claim 9 deals with novel nitrile hydratases which have been prepared in accordance with the invention. Claims 10 and 11 are directed toward specific uses.

35 A prerequisite for finding novel nitrile hydratases is to specify nucleic acid sequences which are able to serve as

probes for nitrile hydratase genes which are present in metagenome DNA libraries. By means of specifying degenerate primer constituents from the group consisting of

- 5 A-01f : gcsmrsgcstgg (Seq. ID NO. 1)
- B-01f : ggsctscscsc (Seq. ID NO. 2)
- B-01r : ggsagsagscc (Seq. ID NO. 3)
- C-01r : ggnccgcwbsgg (Seq. ID NO. 4)
- A-01f : gcnmrrgcntgg (Seq. ID NO. 5)
- 10 B-01f : ggnytnccncc (Seq. ID NO. 6)
- B-01r : ggnggnarncc (Seq. ID NO. 7)
- C-01r : gwngwrtccca (Seq. ID NO. 8)
- A-01f : gcntggrynga (Seq. ID NO. 9)
- B-01f : ggnytsccncc (Seq. ID NO. 10)
- 15 B-01r : ggnggsarncc (Seq. ID NO. 11)
- C-01r : swns wrtccca (Seq. ID NO. 12)

the skilled person obtains, completely surprisingly but no less advantageously for that, special nucleic acid sequences which help to construct specific probes for screening for nitrile hydratase genes in metagenome DNA libraries. These nucleic acid sequences are degenerate sequences which, on the one hand, are specific enough for finding only nitrile hydratase genes but which, on the other hand, are so nonspecific that, if at all possible, all the nitrile hydratase genes which are present are detected. At the time of the invention, it was not possible to deduce their preparation from the prior art in an obvious manner.

30 The present invention accordingly also related to a process for preparing protein sequences which are required for constructing the activity of a nitrile hydratase, such that

- a) a metagenome DNA library of a habitat is prepared,
- 35 b) this library is contacted with in each case at least one forward(f) primer and one reverse(r) primer exhibiting a

degenerate nucleic acid sequence in accordance with sequences 1 to 12,

c) a PCR is carried out using these primers,

d) the full-length sequences of the nucleic acids encoding

5 protein sequences which are required for constructing the activity of a nitrile hydratase are generated from the part sequences which are obtained, and

e) these full-length sequences are cloned into a host organism and expressed.

10

As already noted at the outset, nitrile hydratases consist of at least two different subunits (α and β subunits).

However, further protein sequences, in addition to these two subunits, may also be required to enable the nitrile

15 hydratases to be active. The presence of particular putative "activators" (e.g. folding proteins, etc.) may

sometimes be required in order to enable the corresponding nitrile hydratases to become active. The nucleic acid

20 sequences which encode these activators are frequently located in the immediate vicinity of the nucleic acid

sequences which encode the corresponding nitrile hydratase subunits. It is consequently possible, by screening for

nucleic acid sequences encoding nitrile hydratases, also at the same time to detect all the protein sequences which are

25 required for the nitrile hydratases to be active. According to the process in accordance with the invention, the first

step is that of preparing a metagenome DNA library of a particular habitat. The skilled person is familiar with the

30 way in which this library is prepared (Knietzsch, AW, Tanja; BS; Henne, ADR (2003) Metagenomes of Complex Microbial

Consortia Derived from Different Soils as Sources for Novel Genes Conferring Formation of Carbonyls from Short-Chain

Polyols on *Escherichia coli*. Journal of Molecular Microbiology and Biotechnology 5(1): 46-56; Rondon, MR;

35 August, PR; Bettermann AD; Brady, SF; Grossman, T H; Liles, MR; Loiacono, KA; Lynch, BA; MacNeil, IA; Minor C; Tiong, CL; Gilman, M; Osburne, MS; Clardy, J; Handelsman, J;

Goodman, RM (2000) Cloning the soil metagenome: a strategy for accessing the genetic and functional diversity of uncultured microorganisms. *Applied and environmental microbiology* 66(6):2541-7). Primers which exhibit the
 5 degenerate nucleic acid sequences according to the invention (Seq. ID Nos. 1-12) are then added to this library. A PC reaction is subsequently carried out, with this reaction giving rise to part sequences of the nucleic acid sequences encoding nitrile hydratase subunits.
 10 Employing methods of the prior art, the skilled person can then use these part sequences to identify the corresponding full-length nucleic acid sequences (Schloss, PD; Handelsman, J(2003) *Biotechnological prospects from metagenomics. Current Opinion in Biotechnology*, 14(3): 303-
 15 310; Rondon, MR; August, PR; Bettermann AD; Brady, SF; Grossman, T H; Liles, MR; Loiacono, KA; Lynch, BA; MacNeil, IA; Minor C; Tiong, CL; Gilman, M; Osburne, MS; Clardy, J; Handelsman, J; Goodman, RM (2000) Cloning the soil metagenome: a strategy for accessing the genetic and
 20 functional diversity of uncultured microorganisms. *Applied and environmental microbiology* 66(6):2541-7). Finally, the nucleic acid sequences which have been found are recombinantly prepared in particular expression systems. The methods for doing this are likewise known to the
 25 skilled person (lit. see above).

In a preferred embodiment, the degenerate nucleic acid sequences according to the invention (Seq. ID Nos. 1-12) are employed in the present process such that in each case
 30 primer pairs composed of primers exhibiting the nucleic acid sequences A-01f (Seq. ID Nos. 1, 5 and 9) and B-01r (Seq. ID Nos. 3, 7 and 11) or C-01r (Seq. ID Nos. 4, 8 and 12) and also B-01f (Seq. ID Nos. 2, 6 and 10) and C-01r (Seq. ID Nos. 4, 8 and 12) are used in the PCR. When these
 35 combinations are employed, the nucleic acid sequences encoding the protein sequences which are required for

nitrile hydratase activity are detected in a preferred and efficient manner.

Preference is furthermore given to locating certain other nucleic acid sequences (e.g. "stabilizing regions") upstream of the above-described degenerate primer constituents (Kwok S, Chang SY, Sninsky JJ, Wang A, 1995, "Design and use of mismatched and degenerate primers" In: "PCR Primer, A laboratory Manual" Dieffenbach CW & Dveksler GS (Editors), Cold Spring Harbor Laboratory Press, pp143-155; Compton T, 1990, "Degenerate Primers for DNA Amplification" In: "PCR Protocols, A Guide to Methods and Applications", Innis MA, Gelfand DH, Sninsky JJ, White TJ (Editors) Academic Press, San Diego, pp39-45). In this case, the primers employed in the PC reaction consist of degenerate nucleic acid sequences of the above-described type (sequences ID Nos. 1 to 12) and the nucleic acid sequences mentioned in the sequences having the ID Nos. 13 to 23. Very particular preference is given, therefore, to a process in which nucleic acid sequences selected from the group consisting of:

	GCCAAGGTCGTC	(Seq. ID NO. 13)
	GGCCGGTCCTG	(Seq. ID NO. 14)
	TCCTTGTACCAGGTC	(Seq. ID NO. 15)
	GCCCCGC	(Seq. ID NO. 16)
25	GGCGCTAATGTTGTT	(Seq. ID NO. 17)
	TGGCCGGTTCTG	(Seq. ID NO. 18)
	CAAATTCTTTATACCAAGTC	(Seq. ID NO. 19)
	CCATATATCGCATTTTCAGCT	(Seq. ID NO. 20)
	GGTCGTGGCCAAG	(Seq. ID NO. 21)
30	GGCCGGTCCTG	(Seq. ID NO. 22)
	TCCTTGTACCAGGTC	(Seq. ID NO. 23)
	GCGCATTTTCGGCG	(Seq. ID NO.

are placed upstream of the degenerate nucleic acid sequences (Seq. ID Nos. 1-12). These upstream sequences are likewise derived from conserved nitrile hydratase regions and adapted to the codon usage of organisms having a different GC content.

A process as described at the outset, in which use is made of primers which are selected from the group consisting of

	<u>GCCAAGGTCGTCgcsmrsgcstgg</u>	(Seq. ID NO. 25)
5	<u>GGCCGGTCCTGggsctscsc</u>	(Seq. ID NO. 26)
	<u>TCCTTGTACCAGGTCggsagsc</u>	(Seq. ID NO. 27)
	<u>GCCCGCCggnccgwbssg</u>	(Seq. ID NO. 28)
	<u>GGCGCTAAAGTTGTTgcnmrrgcntgg</u>	(Seq. ID NO. 29)
	<u>TGGCCGGTTCTGggnytncncc</u>	(Seq. ID NO. 30)
10	<u>CAAATTCCTTTATACCAAGTCggnnggnarncc</u>	(Seq. ID NO. 31)
	<u>CCATATATCGCATTTTCAGCTgwngrwtccca</u>	(Seq. ID NO. 32)
	<u>GGTCGTGGCCAAGgcntggrynga</u>	(Seq. ID NO. 33)
	<u>GGCCGGTCCTGggnytscncc</u>	(Seq. ID NO. 34)
	<u>TCCTTGTACCAGGTCggnnggsarncc</u>	(Seq. ID NO. 35)
15	<u>GCGCATTTTCGGCGswnsrwtccca</u>	(Seq. ID NO. 36)

is therefore very particularly advantageous.

Using these primers, it was possible to detect nucleic acid sequences encoding nitrile hydratases, as well as other genes for putative "activators", in metagenome DNA libraries.

Accordingly, a next part of the subject matter of the present invention is constituted by the protein sequences which are required for constructing the activity of a nitrile hydratase, with these sequences possessing less than 100% homology, preferably less than 97%, more preferably less than 96%, even more preferably less than 95%, yet more preferably less than 90%, very preferably less than 85%, and extremely preferably less than 80%, homology, at the amino acid level, with such known protein sequences, and with the nucleic acid sequences encoding them being generated from part sequences which give a positive hybridization signal, under stringent conditions, with the primers according to the invention exhibiting the nucleic acid sequences having the sequences ID Nos. 1 to 12.

The positive hybridization is a prerequisite for it to be possible to find corresponding nucleic acid sequences using the PC reaction-based screening. Methods with which the skilled person is familiar can then be used to obtain the corresponding recombinant protein sequences from these nucleic acid sequences.

These recombinant techniques are used to obtain organisms which are able to provide the protein sequence in question in a quantity which is adequate for an industrial process.

- 10 The rec-protein sequences according to the invention are prepared using recombinant DNA methods which are known to the skilled person (Sambrook, J.; Fritsch, E. F. and Maniatis, T. (1989), *Molecular cloning: a laboratory manual*, 2nd ed., Cold Spring Harbor Laboratory Press, New York; Balbas, P. and Bolivar, F. (1990), *Design and construction of expression plasmid vectors in E.coli*, *Methods Enzymol.* 185, 14-37; Rodriguez, R.L. and Denhardt, D. T (eds) (1988), *Vectors: a survey of molecular cloning vectors and their uses*, 205-225, Butterworth, Stoneham).
- 15 With regard to the general procedure (PCR, cloning, expression, etc.), the reader may be referred to the following literature and the references cited therein: Universal GenomeWalker™ Kit User Manual, Clontech, 3/2000, and the literature cited therein; Triglia T.; Peterson, M. G. and Kemp, D.J. (1988), A procedure for in vitro amplification of DNA segments that lie outside the boundaries of known sequences, *Nucleic Acids Res.* 16, 8186; Sambrook, J.; Fritsch, E. F. and Maniatis, T. (1989), *Molecular cloning: a laboratory manual*, 2nd ed., Cold Spring Harbor Laboratory Press, New York; Rodriguez, R.L. and Denhardt, D. T (eds) (1988), *Vectors: a survey of molecular cloning vectors and their uses*, Butterworth, Stoneham.

- 20 The corresponding nucleic acid sequences, which encode the protein sequences which have just been described, also constitute a next part of the present invention. These nucleic acid sequences are therefore all those which,
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- 30
- 35

within the bounds of the degeneracy of the genetic code, encode the same protein sequence. This consequently likewise also encompasses those nucleic acid sequences which possess, at the nucleic acid level, a homology with the nucleic acid sequences found in accordance with the invention of at least 70 percent, or corresponding fragments of these nucleic acid sequences, which in turn encode protein sequences which are involved in constructing the activity of a nitrile hydratase. Preference is given to these nucleic acid sequences encoding protein sequences which are improved as compared with the protein sequences found in accordance with the invention.

Examples of nucleic acid sequences according to the invention are those of the odd-numbered Seq. ID Nos. 37 to 85. It is possible, as described above, to use the nucleic acid sequences which have been found to obtain the protein sequences according to the invention in high yields from rapidly growing host organisms, e.g. *E. coli*.

This takes place by incorporating (cloning) the nucleic acid sequences according to the invention into special expression systems which can be used to obtain the corresponding protein sequences recombinantly from preferred host organisms. A next aspect of the present invention is therefore constituted by a (artificially prepared) expression system which exhibits one or more of the nucleic acid sequences according to the invention. In principle, all the systems which are familiar to the skilled person for this purpose are suitable for use as the expression system. These systems are preferably plasmids or vectors and microorganisms.

In principle, all the embodiments which are available to the skilled person for this purpose are suitable for use as plasmids or vectors. These plasmids and vectors can be found, for example, in Studier and coworkers (Studier, W. F.; Rosenberg A. H.; Dunn J. J.; Dubendroff J. W.; (1990),

Use of the T7 RNA polymerase to direct expression of cloned genes, *Methods Enzymol.* 185, 61-89) or in the brochures supplied by the companies Novagen, Promega, New England Biolabs, Clontech or Gibco BRL. Other preferred plasmids and vectors can be found in: Glover, D. M. (1985), DNA cloning: a practical approach, Vol. I-III, IRL Press Ltd., Oxford; Rodriguez, R.L. and Denhardt, D. T (eds) (1988), Vectors: a survey of molecular cloning vectors and their uses, 179-204, Butterworth, Stoneham; Goeddel, D. V. (1990), Systems for heterologous gene expression, *Methods Enzymol.* 185, 3-7; Sambrook, J.; Fritsch, E. F. and Maniatis, T. (1989), *Molecular cloning: a laboratory manual*, 2nd ed., Cold Spring Harbor Laboratory Press, New York.

Plasmids which can very preferably be used to clone the construct containing the nucleic acid sequences according to the invention into the host organism are: pUC18 (Roche Biochemicals), pKK-177-3H (Roche Biochemicals), pBTac2 (Roche Biochemicals), pKK223-3 (Amersham Pharmacia Biotech), pKK-233-3 (Stratagene) or pET (Novagen). Extreme preference is given to plasmids of the pET series.

As has been said, the recombinant microorganism into which the plasmids or vectors which contain the nucleic acid sequences according to the invention are cloned is used for replicating and isolating an adequate quantity of the recombinant protein sequence. The methods for doing this are well known to the skilled person (Sambrook, J.; Fritsch, E. F. and Maniatis, T. (1989), *Molecular cloning: a laboratory manual*, 2nd ed., Cold Spring Harbor Laboratory Press, New York). In principle, the microorganisms which are used can be any organisms which the skilled person takes into consideration for this purpose, for example yeasts, such as *Hansenula polymorpha*, *Pichia* sp., *Saccharomyces cerevisiae*, prokaryotes, such as *E. coli* or *Bacillus subtilis*, or eukaryotes, such as mammalian cells, insect cells or plant cells. Preference is given to using *E. coli* strains for this purpose. The following are very

particularly preferred: *E. coli* XL1 Blue, NM 522, JM101, JM109, JM105, RR1, DH5 α , TOP 10⁻ or HB101, BL21, BL21 (DE3), or BL21 (DE3) codon plus RIL.

5 Furthermore, the nucleic acid sequences which are in accordance with the invention (uneven-numbered Seq. ID Nos. 37 to 85) and which are additionally further improved, and which encode the protein sequences which are required for constructing the activity of a nitrile hydratase, are
10 preferably suitable for preparing what are termed whole-cell catalysts. In principle, whole-cell catalysts of this type are recombinant microorganisms such as those which have just been mentioned. However, in addition to the cloned genes encoding a nitrile hydratase, these whole-cell
15 catalysts also comprise additional enzymes which are required for decomposing nitriles into acids. As explained at the outset, these enzymes are enzymes which exhibit amidase activity.

For this reason, whole-cell catalysts which comprise at
20 least one cloned gene for a protein sequence possessing (D- or L-)amidase activity and cloned genes encoding an active nitrile hydratase are also regarded as being recombinant microorganisms of the abovementioned type. Optionally, the whole-cell catalyst can comprise additional nucleic acid
25 sequences which encode enzymes which are advantageous for decomposing a nitrile function into an acid function. These enzymes are, in particular, enzymes which are selected from the group of protein sequences possessing α -aminonitrile racemase activity, possessing cyanohydrin racemase acti-
30 vity, possessing α -hydroxycarboxylic acid racemase activity or possessing (α - or β -)-amino acid amide racemase activity.

In addition to producing the protein sequences according to the invention, which are required for constructing a
35 nitrile hydratase activity, the whole-cell catalyst according to the invention preferably produces a protein sequence possessing L-amidase activity from rhizobium,

preferably *R. huautlense* DSM 14983 (WO2004/005517) or possessing D-amidase activity, e.g. that from *Variovorax* (EP 1318193).

Corresponding racemases are known, for example, from
5 *Pseudomonas putida* and *Rhodococcus* sp. (Godtfredsen, S. E.; Clausen, K.; Ingvorsen, K.; Hermes, H. F.; Van Balken, J. A.; Meijer, E. M. (1989, EP 0 307 023; WO 8 901 525). Other amino acid amide racemases have been described in *Klebsiella oxytoca* by Hermes and coworkers (Hermes, H. F.
10 M.; Peeters, W. P.; Peters, P. J. (1990), EP 0 383 403), as well as in *Agrobacterium rhizogenes* and *Ochrobacterium anthropi* (Boesten, W. H. J.; Raemakers-Franken, P. C.; Sonke, T.; Euverink, G. J. W.; WO 03106691). The advantage of using corresponding racemases is based on the fact that
15 100% of a racemic nitrile can be converted into the corresponding enantiomerically enriched acid.

An organism which is mentioned in DE10155928 as being a host organism is preferably used as whole-cell catalyst. The advantage of such an organism is that several enzyme
20 systems are expressed simultaneously, which means that it is only necessary to grow one rec-organism for reacting a readily preparable nitrile or cyanohydrin or α -aminonitrile to give the corresponding enantiomerically enriched acid.

In order to adjust the expression of the nucleic acid
25 sequences in question with regard to the turnover rates of the protein sequences (enzymes) which they encode, the corresponding nucleic acid sequences can be installed on different plasmids possessing different copy numbers and/or promoters of differing strength can be used so as to ensure
30 that the nucleic acid sequences are expressed at different strengths. In enzyme systems which have been adjusted in this way, there is, advantageously, no accumulation of an intermediate compound, which might possibly have an inhibitory effect, and the reaction under consideration can
35 take place at an optimal overall rate. However, this is sufficiently well known to the skilled person (Gellissen, G.; Piontek, M.; Dahlems, U.; Jenzelewski, V.; Gavanagan, J.

W.; DiCosimo, R.; Anton, D. L.; Janowicz, Z. A. (1996), Recombinant *Hansenula polymorpha* as a biocatalyst. Co-expression of the spinach glycolate oxidase (GO) and the *S. cerevisiae* catalase T (CTT1) gene, Appl. Microbiol. Biotechnol. 46, 46-54; Farwick, M.; London, M.; Dohmen, J.; Dahlems, U.; Gellissen, G.; Strasser, A. W.; DE19920712).

It is consequently possible to use the present process to prepare the α and β subunits of nitrile hydratases using metagenome DNA libraries as the starting material. Accordingly, another part of the subject matter of the present invention is constituted by the nitrile hydratases which exhibit the protein sequences according to the invention for nitrile hydratase α subunits and β subunits and which can be prepared from the nucleic acid sequences which are made available by this process and which encode the α and β subunits according to the invention. As is demonstrated in the examples, active nitrile hydratases are also formed, in this connection, when any arbitrary α subunits are combined with any arbitrary β subunits. This thereby makes it possible to increase still further the diversity of possible nitrile hydratases.

Other parts of the subject matter of the present invention relate to the use of the nucleic acid sequences which are prepared by the process according to the invention for producing improved protein sequences which are required for constructing the activity of a nitrile hydratase. The way in which the skilled person proceeds when improving protein sequences on the basis of altering nucleic acid sequences is well known. In general, this is effected by using methods of mutagenesis. Any methods which are available to the skilled person for this purpose are suitable for being used as mutagenesis methods. In particular, these methods are saturation mutagenesis, random mutagenesis, in-vitro recombination methods and site-directed mutagenesis (Eigen, M. and Gardiner, W. (1984), Evolutionary molecular

engineering based on RNA replication, Pure Appl. Chem. 56, 967-978; Chen, K. and Arnold, F. (1991), Enzyme engineering for nonaqueous solvents: random mutagenesis to enhance activity of subtilisin E in polar organic media. Bio/Technology 9, 1073-1077; Horwitz, M. and Loeb, L. (1986), Promoters Selected From Random DNA-Sequences, Proc Natl Acad Sci USA 83, 7405-7409; Dube, D. and L. Loeb (1989), Mutants Generated By The Insertion Of Random Oligonucleotides Into The Active-Site Of The Beta-Lactamase Gene, Biochemistry 28, 5703-5707; Stemmer, P.C. (1994), Rapid evolution of a protein *in vitro* by DNA shuffling, Nature 370, 389-391 and Stemmer, P.C. (1994), DNA shuffling by random fragmentation and reassembly: *In vitro* recombination for molecular evolution. Proc Natl Acad Sci USA 91, 10747-10751).

The novel nucleic acid sequences which are obtained in this way are cloned into a host organism, and expressed, using the above-described methods, and the protein sequences which have been prepared in this way are detected by means of suitable screening methods and then isolated. In principle, any detection reactions which are possible for the molecules which are formed are suitable for the detection. Nitrile hydratase activities can be detected in a coupled enzymic test using amidases, with ammonium being formed as a by product. In principle, any possible reactions for detecting ammonia or ammonium ions, such as Nessler reagent (Vogel, A., I., (1989) Vogel's textbook of quantitative chemical analysis, John Wiley & Sons, Inc., 5th ed., 679-698, New York), the indophenol reaction, also termed Berthelot's reaction (Wagner, R., (1969) Neue Aspekte zur Stickstoffanalytik in der Wasserchemie, Vom Wasser, [Novel aspects of nitrogen analysis in water chemistry, about water] VCH-Verlag, vol. 36, 263-318, Weinheim), in particular the enzymic determination using glutamate dehydrogenase (Bergmeyer, H., U., and Beutler, H.-O. (1985) Ammonia, in: Methods of Enzymatic Analysis, VCH-Verlag, 3rd Edition, Vol. 8: 454-461, Weinheim), or else

detection using ammonium-sensitive electrodes, are suitable for detecting it. Furthermore, HPLC methods are used for detecting amino acids, for example a derivative method based on o-phthaldialdehyde and N-isobutyryl-cysteine for
5 enantiomerically separating amino acids (Brückner, H., Wittner R., and Godel H., (1991) Fully automated high-performance liquid chromatographic separation of DL-amino acids derivatized with o-Phthaldialdehyde together with N-isopropyl-cysteine. Application to food samples, Anal.
10 Biochem. 144, 204-206). The amide which is directly formed by the nitrile hydratase reaction can likewise be detected using HPLC methods (e.g. reverse phase).

In a last embodiment of the present invention, the latter
15 relates to the use of the nitrile hydratases according to the invention for preparing organic acid amides and acids, in particular enantiomerically enriched α -hydroxy acids or α -amino acids.

For this application, the nitrile hydratase in question can
20 be used in free form, as a homogeneously purified compound or as a recombinantly prepared enzyme. Furthermore, the enzyme can also be employed as a constituent of an intact host organism or in combination with the disrupted, and, if desired, highly purified, cell mass of the host organism.
25 It is likewise possible to use the enzyme in immobilized form (Sharma B. P.; Bailey L. F. and Messing R. A. (1982), Immobilisierte Biomaterialien - Techniken und Anwendungen [Immobilized biomaterials - techniques and applications], Angew. Chem. 94, 836-852). The immobilization is
30 advantageously effected by means of lyophilization (Paradkar, V. M.; Dordick, J. S. (1994), Aqueous-Like Activity of α -Chymotrypsin Dissolved in Nearly Anhydrous Organic Solvents, J. Am. Chem. Soc. 116, 5009-5010; Mori, T.; Okahata, Y. (1997), A variety of lipi-coated glycoside
35 hydrolases as effective glycosyl transfer catalysts in homogeneous organic solvents, Tetrahedron Lett. 38, 1971-1974; Otamiri, M.; Adlercreutz, P.; Matthiasson, B. (1992),

Complex formation between chymotrypsin and ethyl cellulose as a means to solubilize the enzyme in active form in toluene, *Biocatalysis* 6, 291-305). Very particular preference is given to lyophilization in the presence of
5 surface-active substances such as Aerosol OT or polyvinylpyrrolidone or polyethylene glycol (PEG) or Brij 52 (diethylene glycol monocetyl ether) (Kamiya, N.; Okazaki, S.-Y.; Goto, M. (1997), *Surfactant-horseradish peroxidase complex catalytically active in anhydrous*
10 *benzene, Biotechnol. Tech.* 11, 375-378).

Greatest preference is given to immobilization on Eupergit®, in particular Eupergit C® and Eupergit 250L® (Röhm) (Eupergit.RTM. C, a carrier for immobilization of enzymes of industrial potential. Katchalski-Katzir, E.;
15 Kraemer, D. M. *Journal of Molecular Catalysis B: Enzymatic* (2000), 10(1-3), 157-176).

Preference is also given to immobilization on Ni-NTA in combination with the polypeptide being provided with a His tag (hexahistidine) (Purification of proteins using
20 polyhistidine affinity tags. Bornhorst, Joshua A.; Falke, Joseph J. *Methods in Enzymology* (2000), 326, 245-254).

It is likewise possible to conceive of using the polypeptides as CLECs (St. Clair, N.; Wang, Y.-F.; Margolin, A. L. (2000), *Cofactor-bound cross-linked enzyme*
25 *crystals (CLEC) of alcohol dehydrogenase, Angew. Chem. Int. Ed.* 39, 380-383).

By using these procedures, it is possible to successfully generate polypeptides which are able to function in mixtures of aqueous and organic solvents, or in media which
30 are entirely organic, from polypeptides which are rendered unstable by organic solvents.

The present invention describes the isolation of nitrile hydratases and their corresponding genes by using molecular
35 genetic methods to tap the uncultured microbial diversity of different habitats. Using degenerate primers in connection with the PCR-based screening of metagenome DNA

libraries identifies nitrile hydratase genes and elucidates the part sequences of the PCR products which are obtained in this way. In a subsequent step, the complete DNA sequences of the genes are determined in order, after
 5 cloning and heterologous expression, to provide enzyme samples for activity profiling and investigating applications.

By means of rationally selecting soil samples, which might
 10 possibly contain nitriles, for preparing metagenome libraries (metagenome DNA libraries) and focusing on application-relevant substrates within the context of enrichment cultures, it is possible to enrich nitrile-converting microorganisms. In any case, the genetic
 15 screening in metagenome libraries provides a pool of corresponding nitrile hydratase genes for subsequent expression, which pool can, however, also serve as the basis for enzyme optimization by means of directional evolution.

20 Metagenome libraries which were prepared from four different habitats and sites (grassland, forest, sandy ecosystem and biofilm), and which contained more than 83 000 clones, were screened for identifying novel nitrile
 25 hydratases. The skilled person is familiar with the construction of such metagenome libraries (Schloss, PD; Handelsman, J (2003) Biotechnological prospects from metagenomics. Current Opinion in Biotechnology, 14(3): 303-310; Knietsch, AW, Tanja; BS; Henne, ADR (2003) Metagenomes
 30 of Complex Microbial Consortia Derived from Different Soils as Sources for Novel Genes Conferring Formation of Carbonyls from Short-Chain Polyols on Escherichia coli. Journal of Molecular Microbiology and Biotechnology 5(1): 46-56; Rondon, MR; August, PR; Bettermann AD; Brady, SF;
 35 Grossman, T H; Liles, MR; Loiacono, KA; Lynch, BA; MacNeil, IA; Mino, r C; Tiong, CL; Gilman, M; Osburne, MS; Clardy, J; Handelsman, J; Goodman, RM (2000) Cloning the soil

metagenome: a strategy for accessing the genetic and functional diversity of uncultured microorganisms. Applied and environmental microbiology 666): 2541-7). These clones together contain about 3000 MBp of DNA. The
5 metagenome DNA libraries were screened for novel nitrile hydratases by means of a PCR screening which used degenerate oligonucleotides which were derived from conserved primary structural motifs of known nitrile hydratases. The degenerate primers employed were those
10 depicted in sequence ID Nos. 25 to 36.

In this connection, the primers having the primer constituent A-01f can, in particular, in each case be combined with those having the primer constituents B-01r
15 (size of the PCR product to be expected, approx. 210 bp) or C-01r (size of the PCR product to be expected, approx. 350 bp), while the primers having the primer constituent B-01f can be combined with those having the primer constituent C-01r (size of the PCR product to be expected,
20 approx. 180 bp). In particular, the degenerate "core" part of the primers (underlined in sequence ID Nos. 25 to 36) is important for identifying novel nitrile hydratases while the nondegenerate segment can be varied.

25 The corresponding PC reaction can be carried out using known methods. When special polymerases are used, the PCR must be amended in accordance with the manufacturer's instructions.

30 It was possible to identify clones carrying genes for nitrile hydratases in all 5 of the metagenome libraries. The skilled person is familiar with methods for using the part sequences of the nitrile hydratase genes for identifying the gene-carrying clone and elucidating the
35 full-length sequence (Schloss, PD; Handelsman, J (2003) Biotechnological prospects from metagenomics. Current Opinion in Biotechnology, 14(3): 303-310; Duran, R;

Nishiyama, M; Horinouchi, S; Beppy, T (1993)
Characterization of nitrile hydratase genes cloned by DNA
screening from *Rhodococcus erythropolis*. *Biosci Biotech
Biochem* 57(8): 1323-1328).

5

It was thus possible to determine the full-length sequences
(nucleic acid sequences) of a total of 12 genes encoding α
subunits and a total of 10 genes encoding β subunits
(figure 1). The sequences of three genes which could encode
10 putative nitrile hydratase "activators" were also
determined.

In order to prepare the nitrile hydratases, the genes
encoding the α and β subunits were in each case placed
15 separately under the control of the phage T7 gene 10
promoter. The vectors pET22b and, respectively, pET26b
(Novagen, CN Bioscience, Inc.) were used for this purpose
(figs. 5 and 6). Using a two-vector expression system made
it possible to simply combine the nitrile hydratase
20 subunits from different enzymes. Corresponding constructs
were expressed in *E. coli* strain BL21 (DE3) CodonPlus RIL
(Novagen, CN Bioscience, Inc.). The cells were incubated at
26°C using LB medium and induced with 0.5 mM IPTG on
reaching a cell density of O.D.580 = 1.0. It was possible
25 to deal with the problem, which occasionally arose, of
protein aggregation following overexpression at 37°C by
reducing the temperature down to 26°C. Coexpressing various
chaperones (trigger factor, GroEL/GroES and Dnak/DnaJ/GrpE)
did not, at 26°C, show any additional effect which went
30 beyond that achieved by reducing the temperature.

The yields with regard to activity against benzonitrile
varied considerably (fig. 2) with it not being possible to
demonstrate any clear correlation between the protein
35 quantity which was prepared and the activity which was
produced. Thus, at 30 U/g of DBM, the yield in the case of
the clone containing the nitrile hydratase M49bD9 (Seq. ID

No. 49/71) was relatively high even though it was not possible to detect any proteins for the nitrile hydratase in the SDS-PAGE analysis. This contrasts with the clone M12K24 (Seq. ID No. 39/63), which was determined to have an activity of about 2.5 U/g of DBM even though it was possible to overexpress both subunits relatively strongly and a substantial proportion could be found in the soluble fraction. The magnitude of the activity consequently depends to a high degree on the particular enzyme in question.

In the case of a variety of nitrile hydratases, an additional small open reading frame, which encodes a protein sequence which appears to be involved in activating the nitrile hydratase, was identified in the immediate vicinity of the genes for the subunits of the enzyme. These proteins were designated P12K (Seq. ID No. 81/83) or P14K (Seq. ID No. 85) since their molecular weight was about 12 or 14 kDa. While it was possible to prepare some nitrile hydratases in active form even without these P12K homologs, the presence of these proteins was essential for other nitrile hydratases to be expressed in active form. In order to investigate the influence of P12K homologs, whose genes were also found in three metagenome clones, on the expression of the corresponding nitrile hydratases, and also on that of nitrile hydratases from other clones, two of these genes were cloned into the vector pBBR1MCS5 (Kovach et al., 1995, Four new derivatives of the broad host range cloning vector pBBR1MCS, carrying different antibiotic-resistance cassettes, Gene 166: 175-176) and, in this vector, placed under the control of the lac promoter. The corresponding constructs were designated pBBR5-P12K-M49bD9 and pBBR5-P12K-M3aG10 (figs. 6 and 7).

The nitrile hydratase genes M49bD9 (Seq. ID No. 49/71) and M3aG10 (Seq. ID No. 57/79) were expressed in the presence of the corresponding P12K homologs (Seq. ID No. 83 -

M49bD9; Seq. ID No. 85 - M3aG10) at 26°C in *E. coli* strain BL21 CodonPlus RIL. In both cases, it was possible to markedly overexpress the subunits of the nitrile hydratases.

- 5 The presence of the P12K homolog (Seq. ID No. 83) increased the activity of the clone M49bD9 nitrile hydratases (Seq. ID No. 49/71) by about a factor of 27, up to approx. 830 U/g of DBM (table 1). In the case of the clone M3aG10 enzyme (Seq. ID No. 57/79), it was possible to detect
10 activity (approx. 23 U/g of DBM) for the first time under these conditions. These results verify that the presence of the P12K homologs (Seq. ID No. 85) can be crucial for increasing the activity yield.

- 15 Table 1: Activity of nitrile hydratases following coexpression with P12K homologs (Seq. ID Nos.83 and 85)

Clone	Without P12K	With P12K
M49bD9 (α , β)	30 U/g of DBM	826 U/g of DBM
M3aG10	0 U/g of DBM	23 U/g of DBM

- Combining subunits from different metagenome clones opens up the possibility of generating nitrile hydratases which
20 possess potentially novel substrate specificities. By combining α subunits with different β subunits, it would be possible to generate a great diversity of combinations of novel nitrile hydratases. Such a combination of subunits from unrelated nitrile hydratases has not thus far been
25 disclosed in the literature.

- In order to evaluate this possibility, the α subunits of clones M73dC9 and M15aA6 (Seq. ID Nos. 59 and 45), for which it was not possible to find any β subunits, were
30 expressed with the β subunit of clone M12K24 (Seq. ID No. 63) in *E. coli* strain BL21 codon plus RIL.

Whereas both subunits could be expressed at approximately equal strengths when the a-M73dC9a/b-M12K24 (Seq. ID Nos. 59 and 63) pair was expressed, the α subunit of clone M15aA6 (Seq. ID No. 45) appears to have been expressed more strongly than does the β subunit of clone M12K24 (Seq. ID No. 63).

Combining the α subunit of clone M73dC9 (Seq. ID No. 59) and the β subunit of clone M12K24 (Seq. ID No. 63) surprisingly led to the formation of an active nitrile hydratase having an activity of approx. 0.07 U/g of DBM (fig. 3). This result verifies that it is in principle possible to prepare active enzymes by combining nitrile hydratase subunits from different clones. At the time of the invention, it was not possible to deduce this, as such, from the prior art.

Stringent conditions: in this present document, the expression "under stringent conditions" is understood as described in Sambrook et al. (Sambrook, J.; Fritsch, E. F. and Maniatis, T. (1989), Molecular cloning: a laboratory manual, 2nd ed., Cold Spring Harbor Laboratory Press, New York). A stringent hybridization in accordance with the present invention preferably exists when a positive hybridization signal is still observed after washing for 1 hour with 1 x SSC (150 mM sodium chloride, 15 mM sodium citrate, pH 7.0) and 0.1% SDS (sodium dodecyl sulfate) at 50°C, preferably at 55°C, more preferably at 62°C and most preferably at 68°C, and more preferably for 1 hour with 0.2 x SSC and 0.1% SDS at 50°C, more preferably at 55°C, yet more preferably at 62°C and most preferably at 68°C.

Within the context of the invention, optically enriched (enantiomerically enriched, enantiomer enriched) compounds are understood as meaning the presence of one optical antipode at > 50 mol% in a mixture with the other antipode.

All types of single-stranded DNA, and DNA which is complementary thereto or double-stranded DNA (e.g. genomic DNA or cDNA), as well as RNA (e.g. mRNA), or mixtures thereof, are subsumed under the term nucleic acid sequences.

Improved nucleic acid sequences encode improved protein sequences. Improved protein sequences are those which exhibit an improvement, as compared with the original sequences, in regard to activity and/or selectivity and/or stability. According to the invention, this means that the proteins are more active and/or more selective or less selective, or are more stable under the reaction conditions employed. While the activity and the stability of the proteins should naturally be as high as possible for the industrial application, an improvement with regard to selectivity is said to have taken place when the substrate selectivity decreases but the enantioselectivity of the proteins is increased. This also applies to proteins as constituents of nitrile hydratases insofar as they help to confer the corresponding improved properties on the enzyme.

According to the invention, the claimed protein sequences and nucleic acid sequences also encompass those sequences which exhibit a homology (excluding the natural degeneracy) which is greater than 70% (with regard to the nucleic acid sequence) or 80% (with regard to the protein sequences), preferably greater than 90%, 91%, 92%, 93% or 94%, more preferably greater than 95% or 96% and particularly preferably greater than 97%, 98% or 99%, with one of these sequences, provided the mode of action or purpose of such a sequence is preserved. The expression "homology" (or identity), as used herein, can be defined by the equation $H(\%) = [1 - V/X] \times 100$, in which H denotes homology, X is the total number of nucleobases/amino acids in the comparison sequence and V is the number of different nucleobases/amino acids in the sequence to be considered, based on the

comparison sequence. In any case, the term nucleic acid sequences which encode polypeptides encompasses all the sequences which appear possible in accordance with the degeneracy of the genetic code.

5

Description of the figures:

Figure 1: Homology among the nitrile hydratase α and β subunits which were found by means of a genetic screening.

10

Figure 2: Activity of different metagenome nitrile hydratases toward benzonitrile following expression in *E. coli* BL21 (DE3) codon plus RIL, with $\square$ and $\blacksquare$ without coexpression of the trigger factor (n.d.: not determined).

15

Figure 3: Activity of nitrile hydratases when subunits from different metagenome clones are combined.

Figure 4: The vector map shows the general arrangement of the α subunits in plasmid pET22, taking clone M49bD9 as an example.

20

Figure 5: The vector map shows the general arrangement of the β subunits in plasmid pET26 taking clone M49bD9 as an example.

25

Figure 6: The vector map shows the arrangement of the P12K protein from clone M49bD9 in plasmid pBBR5.

Figure 7: The vector map shows the arrangement of the P12K protein from clone M3aG10 in plasmid pBBR5.

30

Experimental section:

General PCR protocol:

5

Culturing microorganisms

The E. coli cells were cultured and stored as described in
Sambrock et al. (Sambrock, J.; Fritsch, E. F. and Maniatis,
10 T. (1989), Molecular cloning: a laboratory manual, 2nd ed.,
Cold Spring Harbor Laboratory Press, New York).

PCR protocol:

15

A typical PCR protocol is described below, with it being
necessary to adapt the protocol in accordance with the
manufacturer's instructions when using a different
polymerase.

20

25 µl of HotStarTaq mastermix, Qiagen
(2.5 U of polymerase, 200 µM dNTPs, 1 × PCR buffer)

25 pmol of primer 01f

25

25 pmol of primer 01r

2 µl of template DNA (20 - 200 ng)

30 dist. water to 50 µl

Program:

Initial denaturation: 1 x 15 min at 95°C

5

Cycle program:

35 × { 1 min at 95°C
1 min at 55 - 69°C
1 min at 72°C

10

Final elongation: 1 x 7 min at 72°C

Last extension: 7 min at 72°C

15 Digestion with restriction enzymes

The DNA to be cut is provided with 5 U of restriction enzyme and the appurtenant buffer and, unless otherwise
20 required, incubated at 37°C. Chromosomal DNA is digested with 10 U of enzyme. The incubation period is 1.5 - 2.5 hours.

Treatment with alkaline phosphatase

25 In order to prevent vectors which have only been cut with
one restriction endonuclease from religating with
themselves, the phosphate residue protruding at the 5' end
is removed with alkaline phosphatase. Circular DNA can only
30 be formed once again by inserting a DNA fragment.

The vector which has been cut with a restriction endonuclease is incubated at 65°C for 15 minutes in order to stop the restriction endonuclease. The dephosphorylation 35 buffer is then added and the vector is incubated with 1 U of shrimp alkaline phosphatase at 37°C for 10 min. The

enzyme is then separated off from the vector DNA by means of a subsequent gel electrophoresis.

5 Treatment with T4 DNA ligase

Vector and insert are used for the ligation in a ratio of 1:3. The volume is chosen to be as small as possible (7 - 20 μ l). The mixture is incubated overnight at 16°C in ligation buffer and in the presence of 1 U of ligase.

Transformation

15 100 μ l of competent cells are added by pipette to the ligation mixture, the constituents of which are mixed by being repeatedly drawn up into the pipette. After 30 min of incubation on ice, a heat shock step at 42°C is carried out for 45 sec and the mixture is incubated once again on ice for 2 min. 120-900 μ l of SOC medium are added and the mixture is incubated at 37°C for 45 min while being agitated. The mixture is subsequently plated out and incubated overnight at 37°C.

25

Expression of metagenome nitrile hydratases

The constructs containing T7 promoters were expressed in accordance with the following protocol:

30 50 ml of LB_{amp100} medium containing 2 mM Fe citrate and in each case 50 μ g of kanamycin and ampicillin/ml were inoculated in a ratio of 99:1 with an overnight culture. After an OD₆₀₀ of approx. 0.5 had been reached, expression of the nitrile hydratases was induced with 1 mM IPTG (isopropylthiogalactoside). The cells were harvested 35 approx. 24 hours after induction at 26°C.

Detection of activity using benzonitrile as substrate

The biotransformation was carried out on a 10 ml scale using approx. 100 mg of moist biomass ($OD_{600} = 5$) in 5 potassium phosphate buffer (100 mM), pH 7.0. The incubation took place at 30°C and the substrate concentration was approx. 5 mM benzonitrile. Samples were taken every 5 - 10 min over a period of at most 1 hour. The sample volume was 100 μ l and the reaction was stopped by adding 10 μ l of 10 50% phosphoric acid.

The concentrations of benzonitrile and benzamide were then determined by means of HPLC:

- 15 Column: RP18 Phenomenex Hypersil ODS 5 μ column (with precolumn)
Mobile phase: 10 mM K_2HPO_4 , (pH 2.3)
Flow rate: 1 ml/min
Wavelength: 202 nm
- 20 Injection volume: 20 μ l
Duration of HPLC run: 12-15 min

The activity was calculated by calculating a μ mol turnover after one minute, with one μ mol corresponding to one U 25 (unit). Specific activities are given in U per g of DBM or mg of protein.

Claims

1. A protein sequence comprising (a) the sequence set forth in SEQ ID NO: 50; (b) the sequence set forth in SEQ ID NO: 72; and (c) the sequence set forth in SEQ ID NO: 84.

2. A nucleic acid sequence which encodes a protein sequence as claimed in claim 1.

3. The nucleic acid sequence of claim 2, wherein the nucleic acid sequence comprises (a) the sequence set forth in SEQ ID No: 49; (b) the sequence set forth in SEQ ID NO: 71; or (c) the sequence set forth in SEQ ID NO: 83.

4. An expression system which exhibits one or more nucleic acid sequences as claimed in claim 2.

5. A nucleic acid sequence having at least 70% homology with the sequence set forth in SEQ ID NO: 49, 71 or 83, wherein said nucleic acid sequence or fragment thereof encodes a protein sequence which is involved in constructing the activity of a nitrile hydratase.

6. A nitrile hydratase which exhibits protein sequences for α subunits and β subunits as claimed in claim 1(a) and (b), respectively.

7. The use of the nucleic acid sequences as claimed in claim 2 for producing improved protein sequences which are required for constructing the activity of a nitrile hydrates.

8. The use of the nitrile hydratases as claimed in claim 6 for preparing organic acid amides and acids.

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    <223> a or g or c or t/u

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    11

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25   <213> Artificial

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40   <210> 13
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    15

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    7

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45

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    <212> DNA
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    <223> Primer
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    20
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25 <220>
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25   <223> Primer

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30   <223> a or c or g or t/u

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 22

55 <210> 35
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    <223> a or c or g or t/u

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    26

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    24

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    48
    Met Ser Lys His Val His Asp Tyr Tyr Ala Lys Lys Lys His Asp His
    1 5 10 15

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gat cat gac cac gac gtc cac gag gcg atc gag gac cgg gac gag ggt
 96
 5 Asp His Asp His Asp Val His Glu Ala Ile Glu Asp Arg Asp Glu Gly
 20 25 30
 ccg ccg tcg gaa ttc gag atc atg agc cgc gcc atg cag gag ctg ctg
 144
 10 Pro Pro Ser Glu Phe Glu Ile Met Ser Arg Ala Met Gln Glu Leu Leu
 35 40 45
 gaa gag aag ggc gtc gtc acc gcg gag cag gtc cgg cgc agc atg gag
 192
 15 Glu Glu Lys Gly Val Val Thr Ala Glu Gln Val Arg Arg Ser Met Glu
 50 55 60
 aag ttc gag gaa gag ctg ccc tac cgg ggg gcg cgg gtc gtc gcg cac
 240
 20 Lys Phe Glu Glu Glu Leu Pro Tyr Arg Gly Ala Arg Val Val Ala His
 65 70 75 80
 gcc tgg acc gac ccg gaa ttc aag aag cgg ctg ctg gcg gac ggc aag
 288
 25 Ala Trp Thr Asp Pro Glu Phe Lys Lys Arg Leu Leu Ala Asp Gly Lys
 85 90 95
 gcc gcc gtc tcg gag ttc ggc atc gat ttg gaa gcc gag cgg ctg atc
 336
 30 Ala Ala Val Ser Glu Phe Gly Ile Asp Leu Glu Ala Glu Arg Leu Ile
 100 105 110
 gcg gtc gcg aac aca acg gac gtg cac aac gtc atc gtg tgc acg ctg
 384
 35 Ala Val Ala Asn Thr Thr Asp Val His Asn Val Ile Val Cys Thr Leu
 115 120 125
 tgc tcg tgc tac ccg cgc acg ctg ctc ggc atg ccg ccg acc tgg tac
 432
 40 Cys Ser Cys Tyr Pro Arg Thr Leu Leu Gly Met Pro Pro Thr Trp Tyr
 130 135 140
 aag agc gat aac tac cgc tcg cgc gtg gtc tac gaa ccg cgc gcg gtg
 480
 45 Lys Ser Asp Asn Tyr Arg Ser Arg Val Val Tyr Glu Pro Arg Ala Val
 145 150 155 160
 ctg aag gaa ttc ggc acc gtg ctg ccg gag cgc gtc acc gtg cgc gtg
 528
 50 Leu Lys Glu Phe Gly Thr Val Leu Pro Glu Arg Val Thr Val Arg Val
 165 170 175
 cac gac tcc aac gcc gac atg cgc tac gtg gtg atc ccc atg cgc ccg
 576
 55 His Asp Ser Asn Ala Asp Met Arg Tyr Val Val Ile Pro Met Arg Pro
 180 185 190
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 624
 Gln Gly Thr Glu Gly Trp Ser Glu Glu Arg Leu Ala Glu Leu Leu Thr

	195	200	205
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	672		
5	Arg Asp Thr Leu Val Gly Val Thr Val Pro Lys Val Glu Val Gly Ser		
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	705		
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	20	25	30
30	Pro Pro Ser Glu Phe Glu Ile Met Ser Arg Ala Met Gln Glu Leu Leu		
	35	40	45
35	Glu Glu Lys Gly Val Val Thr Ala Glu Gln Val Arg Arg Ser Met Glu		
	50	55	60
40	Lys Phe Glu Glu Glu Leu Pro Tyr Arg Gly Ala Arg Val Val Ala His		
	65	70	75 80
	Ala Trp Thr Asp Pro Glu Phe Lys Lys Arg Leu Leu Ala Asp Gly Lys		
45	85	90	95
	Ala Ala Val Ser Glu Phe Gly Ile Asp Leu Glu Ala Glu Arg Leu Ile		
	100	105	110
50	Ala Val Ala Asn Thr Thr Asp Val His Asn Val Ile Val Cys Thr Leu		
	115	120	125
55	Cys Ser Cys Tyr Pro Arg Thr Leu Leu Gly Met Pro Pro Thr Trp Tyr		
	130	135	140

Lys Ser Asp Asn Tyr Arg Ser Arg Val Val Tyr Glu Pro Arg Ala Val
 145 150 155 160

5 Leu Lys Glu Phe Gly Thr Val Leu Pro Glu Arg Val Thr Val Arg Val
 165 170 175

10 His Asp Ser Asn Ala Asp Met Arg Tyr Val Val Ile Pro Met Arg Pro
 180 185 190

15 Gln Gly Thr Glu Gly Trp Ser Glu Glu Arg Leu Ala Glu Leu Leu Thr
 195 200 205

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 210 215 220

20 Arg Lys Ser Lys Gly Gly Ser Lys Thr Arg
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25 <210> 39
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 1 5 10 15

45 tca gcc atg aac att cca gct cgt gaa ttc gcc ctc cag cgc act gcg
 96
 Ser Ala Met Asn Ile Pro Ala Arg Glu Phe Ala Leu Gln Arg Thr Ala
 20 25 30

50 ccg gtc gag cag cgt gtc gac gcg atc cag gcg gcg ctc gac gaa cgc
 144
 Pro Val Glu Gln Arg Val Asp Ala Ile Gln Ala Ala Leu Asp Glu Arg
 35 40 45

55 ggt ttg aac gcc agt gac gca gtc cag gaa ttg agc cac ctg gcg gag
 192
 Gly Leu Asn Ala Ser Asp Ala Val Gln Glu Leu Ser His Leu Ala Glu
 50 55 60

gag caa tgg att ccg cgc aat ggc gcg cgg gtc gtc gcc aaa gcc tgg
 240
 Glu Gln Trp Ile Pro Arg Asn Gly Ala Arg Val Val Ala Lys Ala Trp
 65 70 75 80
 5 gtc gac ccg gaa ttc cgc gcg cgg ctt ctg gcc gac ggt cgc gcc gcc
 288
 Val Asp Pro Glu Phe Arg Ala Arg Leu Leu Ala Asp Gly Arg Ala Ala
 85 90 95
 10 gtt gcc gaa ctg ggc ctc tcg atg ccg aag cat cac cgg cac ctc gtg
 336
 Val Ala Glu Leu Gly Leu Ser Met Pro Lys His His Arg His Leu Val
 100 105 110
 15 gtg ctg gag aac acg ccg agc gtg cag aac gtc atc tgc tgc acg cag
 384
 Val Leu Glu Asn Thr Pro Ser Val Gln Asn Val Ile Cys Cys Thr Gln
 115 120 125
 20 tgc tcg tgc acg gcg ttc acg atc atc gga ctg ccg ccc gac tgg tac
 432
 Cys Ser Cys Thr Ala Phe Thr Ile Ile Gly Leu Pro Pro Asp Trp Tyr
 130 135 140
 25 aag gac ctg gaa tac cgc gcg cga gtc gtc cgg gag tcg cgc acc gtg
 480
 Lys Asp Leu Glu Tyr Arg Ala Arg Val Val Arg Glu Ser Arg Thr Val
 145 150 155 160
 30 ctg aag gag atg gga ctg gat ctg cct cgg gat gtc gaa att cgc gtc
 528
 Leu Lys Glu Met Gly Leu Asp Leu Pro Arg Asp Val Glu Ile Arg Val
 165 170 175
 35 tgg gat acc act gcc gac acg cgc tac atg gta ttg ccg gta cag ccg
 576
 Trp Asp Thr Thr Ala Asp Thr Arg Tyr Met Val Leu Pro Val Gln Pro
 180 185 190
 40 ccg gaa acc atc ggc tgg ccc gag gag aaa ctg gtc gac atc gtg acg
 624
 Pro Glu Thr Ile Gly Trp Pro Glu Glu Lys Leu Val Asp Ile Val Thr
 195 200 205
 45 cgc gac ggc atg atc ggc gtc gcg cgg gta tag
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 Arg Asp Gly Met Ile Gly Val Ala Arg Val
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 55 <213> Unknown
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 20 25 30
 10 Pro Val Glu Gln Arg Val Asp Ala Ile Gln Ala Ala Leu Asp Glu Arg
 35 40 45
 15 Gly Leu Asn Ala Ser Asp Ala Val Gln Glu Leu Ser His Leu Ala Glu
 50 55 60
 20 Glu Gln Trp Ile Pro Arg Asn Gly Ala Arg Val Val Ala Lys Ala Trp
 65 70 75 80
 Val Asp Pro Glu Phe Arg Ala Arg Leu Leu Ala Asp Gly Arg Ala Ala
 85 90 95
 25 Val Ala Glu Leu Gly Leu Ser Met Pro Lys His His Arg His Leu Val
 100 105 110
 30 Val Leu Glu Asn Thr Pro Ser Val Gln Asn Val Ile Cys Cys Thr Gln
 115 120 125
 35 Cys Ser Cys Thr Ala Phe Thr Ile Ile Gly Leu Pro Pro Asp Trp Tyr
 130 135 140
 40 Lys Asp Leu Glu Tyr Arg Ala Arg Val Val Arg Glu Ser Arg Thr Val
 145 150 155 160
 Leu Lys Glu Met Gly Leu Asp Leu Pro Arg Asp Val Glu Ile Arg Val
 165 170 175
 45 Trp Asp Thr Thr Ala Asp Thr Arg Tyr Met Val Leu Pro Val Gln Pro
 180 185 190
 50 Pro Glu Thr Ile Gly Trp Pro Glu Glu Lys Leu Val Asp Ile Val Thr
 195 200 205
 55 Arg Asp Gly Met Ile Gly Val Ala Arg Val
 210 215

<210> 41

<211> 699
 <212> DNA
 <213> Unknown

5 <220>
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 ggt gcg acg gct gca ccc gtt gga ctc gcc aag atc gct cac tcc cac
 20 96
 Gly Ala Thr Ala Ala Pro Val Gly Leu Ala Lys Ile Ala His Ser His
 20 25 30
 gag cac cag gcc gtt cct tcc gac ctc acg ctc cgg gtc aag tcc ctc
 25 144
 Glu His Gln Ala Val Pro Ser Asp Leu Thr Leu Arg Val Lys Ser Leu
 35 40 45
 gaa tcg ctg ctg gtc gag aag ggt ctc gtg gac cgt gag gcc ctc gac
 30 192
 Glu Ser Leu Leu Val Glu Lys Gly Leu Val Asp Arg Glu Ala Leu Asp
 50 55 60
 gtg ctc gtc gat acc tac gag aac aag atc ggt ccg cga aac ggc gct
 35 240
 Val Leu Val Asp Thr Tyr Glu Asn Lys Ile Gly Pro Arg Asn Gly Ala
 65 70 75 80
 cgc gtc gtc gcg cgg gcg tgg gtc gat ccc gcg tac aaa gag cgc ctg
 40 288
 Arg Val Val Ala Arg Ala Trp Val Asp Pro Ala Tyr Lys Glu Arg Leu
 85 90 95
 ctg aaa gac gcc acc tcg gcg atc gcc gag ctc ggt tac acc gga gcc
 45 336
 Leu Lys Asp Ala Thr Ser Ala Ile Ala Glu Leu Gly Tyr Thr Gly Ala
 100 105 110
 cag ggt gag cac atg gtg gcg ctc gag aat acc ccc gcg gtg cac aac
 50 384
 Gln Gly Glu His Met Val Ala Leu Glu Asn Thr Pro Ala Val His Asn
 115 120 125
 ctc gtc gtt tgc acg ctc tgc tcc tgc tat cca tgg ccg gtg ctc ggt
 55 432
 Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly
 130 135 140

ctg ccc ccg gtc tgg tac aaa tcg gcg ccc tac cga tcg cgc tcg gtc
 480
 Leu Pro Pro Val Trp Tyr Lys Ser Ala Pro Tyr Arg Ser Arg Ser Val
 145 150 155 160
 5
 atc gat ccg cgc gcc gtt ctc gcc gag ttc ggg ctc gag ctg ccg gaa
 528
 Ile Asp Pro Arg Gly Val Leu Gly Glu Phe Gly Leu Glu Leu Pro Glu
 165 170 175
 10
 ggg gtc gag gtg cgc gtc tgg gac tcg acg gcg gag ctc ccg tat ctc
 576
 Gly Val Glu Val Arg Val Trp Asp Ser Thr Ala Glu Leu Arg Tyr Leu
 180 185 190
 15
 gtt ttg ccg gag ccg ccc gaa gcc acg gcg caa ctg agc gaa gaa gcg
 624
 Val Leu Pro Glu Arg Pro Glu Gly Thr Ala Gln Leu Ser Glu Glu Ala
 195 200 205
 20
 ctc gcg gat ctc gtc acc ccg gat gcc atg atc gcc gtc gcg aaa gtc
 672
 Leu Ala Asp Leu Val Thr Arg Asp Ala Met Ile Gly Val Ala Lys Val
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 Ser Leu Pro Ala Gly Gly Ala Glu
 225 230
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 35 <213> Unknown
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 Glu His Gln Ala Val Pro Ser Asp Leu Thr Leu Arg Val Lys Ser Leu
 35 40 45
 55
 Glu Ser Leu Leu Val Glu Lys Gly Leu Val Asp Arg Glu Ala Leu Asp
 50 55 60
 Val Leu Val Asp Thr Tyr Glu Asn Lys Ile Gly Pro Arg Asn Gly Ala
 65 70 75 80

5 Arg Val Val Ala Arg Ala Trp Val Asp Pro Ala Tyr Lys Glu Arg Leu
 85 90 95
 Leu Lys Asp Ala Thr Ser Ala Ile Ala Glu Leu Gly Tyr Thr Gly Ala
 100 105 110
 10 Gln Gly Glu His Met Val Ala Leu Glu Asn Thr Pro Ala Val His Asn
 115 120 125
 15 Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly
 130 135 140
 20 Leu Pro Pro Val Trp Tyr Lys Ser Ala Pro Tyr Arg Ser Arg Ser Val
 145 150 155 160
 Ile Asp Pro Arg Gly Val Leu Gly Glu Phe Gly Leu Glu Leu Pro Glu
 165 170 175
 25 Gly Val Glu Val Arg Val Trp Asp Ser Thr Ala Glu Leu Arg Tyr Leu
 180 185 190
 30 Val Leu Pro Glu Arg Pro Glu Gly Thr Ala Gln Leu Ser Glu Glu Ala
 195 200 205
 35 Leu Ala Asp Leu Val Thr Arg Asp Ala Met Ile Gly Val Ala Lys Val
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 40 Ser Leu Pro Ala Gly Gly Ala Glu
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 45 <210> 43
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	96			
5	Leu Ser Glu Met Asp Ile Arg Val Arg Ala Leu Glu Thr Ile Leu Thr	20	25	30
	gag aag ggc tat gtc gat ccg gcg gcg ctc gac cgg atc gtc gag gcg			
	144			
10	Glu Lys Gly Tyr Val Asp Pro Ala Ala Leu Asp Arg Ile Val Glu Ala	35	40	45
	ttc gag acc agg atc ggc ccg cat atc ggc gcc cgt atc gtg gca cgg			
	192			
15	Phe Glu Thr Arg Ile Gly Pro His Ile Gly Ala Arg Ile Val Ala Arg	50	55	60
	gct tgg gcc gac gcc gaa ttc aag cgg cgg ctg ctc gcc gac gcg acc			
	240			
20	Ala Trp Ala Asp Ala Glu Phe Lys Arg Arg Leu Leu Ala Asp Ala Thr	65	70	75
				80
	gag gcg gcg aat tcg ctg ggt cat gcg agc ccg gtc ggc agc cat ctg			
	288			
25	Glu Ala Ala Asn Ser Leu Gly His Ala Ser Pro Val Gly Ser His Leu	85	90	95
	atc gcg gtc gag aac acg ccg cag acc cac aac ctc gtc gtc tgc act			
	336			
30	Ile Ala Val Glu Asn Thr Pro Gln Thr His Asn Leu Val Val Cys Thr	100	105	110
	ttg tgc tgg tgt tat ccg tgg gag gtg ctg gga ttg ccg ccg gtc tgg			
	384			
35	Leu Cys Ser Cys Tyr Pro Trp Glu Val Leu Gly Leu Pro Pro Val Trp	115	120	125
	tac aaa tcc gct gcc tac cgc tgg cgc gtg gtg atc gac ccc aag ggc			
	432			
40	Tyr Lys Ser Ala Ala Tyr Arg Ser Arg Val Val Ile Asp Pro Lys Gly	130	135	140
	gtc ctc gcc gag ttc ggc ctg acc ctg cca ccg gag acc ggg atc cgc			
	480			
45	Val Leu Ala Glu Phe Gly Leu Thr Leu Pro Pro Glu Thr Gly Ile Arg	145	150	155
				160
	atc tgg gat tgg acc gcc gag acc cgg ttt ctg gtg gtg ccg atg cgg			
	528			
50	Ile Trp Asp Ser Thr Ala Glu Thr Arg Phe Leu Val Val Pro Met Arg	165	170	175
	ccc ccc ggc acc gca gcc tgg agc gag gaa cgg ctc gcc gaa ctc gtc			
	576			
55	Pro Pro Gly Thr Ala Gly Trp Ser Glu Glu Arg Leu Ala Glu Leu Val	180	185	190
	acc cgc gac agc atg atc ggc act ggt ctg gcc ggg gcg ccg cag gag			
	624			

Thr Arg Asp Ser Met Ile Gly Thr Gly Leu Ala Gly Ala Pro Gln Glu
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5 atg gcc tcg gca tga
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 Met Ala Ser Ala
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 20 25 30

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 35 40 45

Phe Glu Thr Arg Ile Gly Pro His Ile Gly Ala Arg Ile Val Ala Arg
 50 55 60

35 Ala Trp Ala Asp Ala Glu Phe Lys Arg Arg Leu Leu Ala Asp Ala Thr
 65 70 75 80

40 Glu Ala Ala Asn Ser Leu Gly His Ala Ser Pro Val Gly Ser His Leu
 85 90 95

45 Ile Ala Val Glu Asn Thr Pro Gln Thr His Asn Leu Val Val Cys Thr
 100 105 110

50 Leu Cys Ser Cys Tyr Pro Trp Glu Val Leu Gly Leu Pro Pro Val Trp
 115 120 125

Tyr Lys Ser Ala Ala Tyr Arg Ser Arg Val Val Ile Asp Pro Lys Gly
 130 135 140

55 Val Leu Ala Glu Phe Gly Leu Thr Leu Pro Pro Glu Thr Gly Ile Arg
 145 150 155 160

Ile Trp Asp Ser Thr Ala Glu Thr Arg Phe Leu Val Val Pro Met Arg
165 170 175

5 Pro Pro Gly Thr Ala Gly Trp Ser Glu Glu Arg Leu Ala Glu Leu Val
180 185 190

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195 200 205

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96
Leu His Asp Arg Ala Gly Gly Val Arg Ser Leu Arg Gly Lys Arg Ser
20 25 30

40 cat cgc gcc gga tcg cat cct cgg ggc gct cgc gca tcc gtc gcc aca
144
His Arg Ala Gly Ser His Pro Arg Gly Ala Arg Ala Ser Val Ala Thr
35 40 45

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192
Gly Trp Phe Val Pro Phe Ser Ala Arg Leu Ala Arg Lys Gly Ile Ala
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240
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65 70 75 80

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288
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85 90 95

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 336
 5 Phe Arg Thr Leu Leu Leu Glu Asp Gly Thr Arg Ala Ala Glu Ser Leu
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 384
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 115 120 125

ccc gag att cat cac ctc gtc gtt tgc acg ctg tgc agt tgt tac ccg
 432
 15 Pro Glu Ile His His Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro
 130 135 140

cgc gcg gtg ctg ggc tat ccg ccg ttc tgg ttc aaa tcc gcc gcc tac
 480
 20 Arg Ala Val Leu Gly Tyr Pro Pro Phe Trp Phe Lys Ser Ala Ala Tyr
 145 150 155 160

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 528
 25 Arg Ala Arg Ala Val Arg Asp Pro Arg Gly Leu Ile Ala Glu Trp Gly
 165 170 175

acc atg ctg ccc gac gat gtc cgc gtg cga gtg gtg gac agt acg gcc
 576
 30 Thr Met Leu Pro Asp Asp Val Arg Val Arg Val Val Asp Ser Thr Ala
 180 185 190

gac tat cgc tgg atg gtt ctg ccg gtg cgg ccg gcc ggc act gcg ggc
 624
 35 Asp Tyr Arg Trp Met Val Leu Pro Val Arg Pro Ala Gly Thr Ala Gly
 195 200 205

tgg gat gag gag cgc ctc gcc gca atc gta cgc gaa ggc gat atg atc
 672
 40 Trp Asp Glu Glu Arg Leu Ala Ala Ile Val Arg Glu Gly Asp Met Ile
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 35 40 45
 10 Gly Trp Phe Val Pro Phe Ser Ala Arg Leu Ala Arg Lys Gly Ile Ala
 50 55 60
 15 Pro Pro Ala Glu Ile Ala Glu Arg Ile Ala Val Thr Asp Arg Ala Ser
 65 70 75 80
 20 Pro Ala Met Gly Ala Arg Met Val Ala Lys Ala Trp Thr Asp Pro Ala
 85 90 95
 Phe Arg Thr Leu Leu Leu Glu Asp Gly Thr Arg Ala Ala Glu Ser Leu
 100 105 110
 25 Gly Ile Met Met Arg Gly Ala Pro Pro Leu Gly Val Leu Glu Asn Thr
 115 120 125
 30 Pro Glu Ile His His Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro
 130 135 140
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 145 150 155 160
 40 Arg Ala Arg Ala Val Arg Asp Pro Arg Gly Leu Ile Ala Glu Trp Gly
 165 170 175
 Thr Met Leu Pro Asp Asp Val Arg Val Arg Val Val Asp Ser Thr Ala
 180 185 190
 45 Asp Tyr Arg Trp Met Val Leu Pro Val Arg Pro Ala Gly Thr Ala Gly
 195 200 205
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tat ctc gat ccc gcc gcg ctt gat gcg atg atc gaa gcc tac gag acg
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 Tyr Leu Asp Pro Ala Ala Leu Asp Ala Met Ile Glu Ala Tyr Glu Thr
 20 25 30

cgg att ggg ccg cat aac ggc gcg cgc gtc gtc gcc aag gcc tgg tcc
 25 144
 Arg Ile Gly Pro His Asn Gly Ala Arg Val Val Ala Lys Ala Trp Ser
 35 40 45

gac gcc gca ttc aag cga gcg ctg gtc gag gat gcg acc aag gcc gtg
 30 192
 Asp Ala Ala Phe Lys Arg Ala Leu Val Glu Asp Ala Thr Lys Ala Val
 50 55 60

cag tcg ttc ggc gtg gtc aat cgc gtc ggc gat cac ctg atc gcg gtc
 35 240
 Gln Ser Phe Gly Val Val Asn Arg Val Gly Asp His Leu Ile Ala Val
 65 70 75 80

gag aac acg ccc acg ctg cac aac atc atc gtg tgc acg ttg tgc tcc
 40 288
 Glu Asn Thr Pro Thr Leu His Asn Ile Ile Val Cys Thr Leu Cys Ser
 85 90 95

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 45 336
 Cys Tyr Pro Trp Glu Val Leu Gly Leu Pro Pro Val Trp Tyr Lys Ser
 100 105 110

gcg ccc tac cgc tcg cgc gcg gtc aac gac ccg cgc ggg gta ctc gcc
 50 384
 Ala Pro Tyr Arg Ser Arg Ala Val Asn Asp Pro Arg Gly Val Leu Ala
 115 120 125

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 55 432
 Asp Phe Gly Leu Lys Leu Ala Pro Asp Met Gln Ile Arg Val Trp Asp
 130 135 140

tcg acc gcc gag acg cgc ttc atc gtg ttg ccg atg cgc ccg gcc gga
 480
 Ser Thr Ala Glu Thr Arg Phe Ile Val Leu Pro Met Arg Pro Ala Gly
 145 150 155 160
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 acc gac gcc tgg agc gaa gaa aag ctc gcc gcg ctg gtg aca cgc gat
 528
 Thr Asp Gly Trp Ser Glu Glu Lys Leu Ala Ala Leu Val Thr Arg Asp
 165 170 175
 10
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 576
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 35 40 45
 40 Asp Ala Ala Phe Lys Arg Ala Leu Val Glu Asp Ala Thr Lys Ala Val
 50 55 60
 Gln Ser Phe Gly Val Val Asn Arg Val Gly Asp His Leu Ile Ala Val
 65 70 75 80
 45
 Glu Asn Thr Pro Thr Leu His Asn Ile Ile Val Cys Thr Leu Cys Ser
 85 90 95
 50
 Cys Tyr Pro Trp Glu Val Leu Gly Leu Pro Pro Val Trp Tyr Lys Ser
 100 105 110
 55 Ala Pro Tyr Arg Ser Arg Ala Val Asn Asp Pro Arg Gly Val Leu Ala
 115 120 125
 Asp Phe Gly Leu Lys Leu Ala Pro Asp Met Gln Ile Arg Val Trp Asp

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40	aag ggg ctg atc gcg acc gac gtg atc gat cgc gtg gta gca acg tac 144 Lys Gly Leu Ile Ala Thr Asp Val Ile Asp Arg Val Val Ala Thr Tyr 35 40 45		
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55	gcg att gcc gag ctg gga ttc ggc ggc ttg cag ggc gaa cac atg atg 288 Ala Ile Ala Glu Leu Gly Phe Gly Gly Leu Gln Gly Glu His Met Met 85 90 95		

gtc gtg gaa aac acg ccg tcc gta cat aac gtg atc tgt tgc acg cta
 336
 Val Val Glu Asn Thr Pro Ser Val His Asn Val Ile Cys Cys Thr Leu
 100 105 110
 5 tgc tca tgc tat ccg tgg ccg gtc ctg gga ctt ccg ccg agc tgg tac
 384
 Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly Leu Pro Pro Ser Trp Tyr
 115 120 125
 10 aag tcg ctg gcg tat cgt tgc cga atc gtg cgc gag ccg cgc gcc gtc
 432
 Lys Ser Leu Ala Tyr Arg Ser Arg Ile Val Arg Glu Pro Arg Ala Val
 130 135 140
 15 ctc ggc gaa ttc ggc ctc gaa ttg ccc gaa acg gtg gaa gtc cgc gta
 480
 Leu Gly Glu Phe Gly Leu Glu Leu Pro Glu Thr Val Glu Val Arg Val
 145 150 155 160
 20 tgg gat agc agt gct gag atg cgc tat ctc gtg ttg ccg gag cgt cca
 528
 Trp Asp Ser Ser Ala Glu Met Arg Tyr Leu Val Leu Pro Glu Arg Pro
 165 170 175
 25 gcg gga acg acg gag ttg agc gaa gcg gaa ttg gct tca ttg atc acg
 576
 Ala Gly Thr Thr Glu Leu Ser Glu Ala Glu Leu Ala Ser Leu Ile Thr
 180 185 190
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 35 40 45
 Glu Lys Glu Val Gly Pro Leu Asn Gly Ala Lys Val Val Ala Arg Ala

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20	Cys	Ser	Cys	Tyr	Pro	Trp	Pro	Val	Leu	Gly	Leu	Pro	Pro	Ser	Trp	Tyr			
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25	Lys	Ser	Leu	Ala	Tyr	Arg	Ser	Arg	Ile	Val	Arg	Glu	Pro	Arg	Ala	Val			
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40	Ala	Gly	Thr	Thr	Glu	Leu	Ser	Glu	Ala	Glu	Leu	Ala	Ser	Leu	Ile	Thr			
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 Ala Lys Ala Leu Ala Glu Ala Leu Ser Lys Gln Gly Leu Val Pro Glu
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 144
 Gly Phe Leu Asp Gln Val Gly Ser His Ala Ala Glu Ala Trp Ser Pro
 35 40 45
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 192
 Arg Asn Gly Ala Arg Val Val Ala Arg Ala Trp Val Asp Pro Glu Tyr
 50 55 60
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 cgg acg cgc ttg ctc gcc gac ggc acc gcc gcg tgc gcc gcg ctc ggc
 240
 Arg Thr Arg Leu Leu Ala Asp Gly Thr Ala Ala Cys Ala Ala Leu Gly
 65 70 75 80
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 288
 Tyr Ala Gly Pro Gln Gly Glu Tyr Ile Val Val Leu Glu Asp Thr Leu
 85 90 95
 25
 gcc gtt cac aac gtg atc gtg tgt acg caa tgc tcg tgt act gcg tgg
 336
 Ala Val His Asn Val Ile Val Cys Thr Gln Cys Ser Cys Thr Ala Trp
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 384
 Pro Val Leu Gly Leu Pro Pro Asp Trp Tyr Lys Ser Pro Glu Tyr Arg
 115 120 125
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 gcc cgc gtc gtg cgg gag ccg cga cgg gtg ctt cgc gaa atg ggc ctc
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 Glu Leu Ser Glu Ser Val Thr Ile Arg Val Trp Asp Thr Thr Ala Glu
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 Thr Arg Phe Leu Val Leu Pro Leu Arg Pro Ala Gly Thr Glu Gly Trp
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 576
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 25 Arg Asn Gly Ala Arg Val Val Ala Arg Ala Trp Val Asp Pro Glu Tyr
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 30 Arg Thr Arg Leu Leu Ala Asp Gly Thr Ala Ala Cys Ala Ala Leu Gly
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 Tyr Ala Gly Pro Gln Gly Glu Tyr Ile Val Val Leu Glu Asp Thr Leu
 85 90 95
 35
 Ala Val His Asn Val Ile Val Cys Thr Gln Cys Ser Cys Thr Ala Trp
 100 105 110
 40
 Pro Val Leu Gly Leu Pro Pro Asp Trp Tyr Lys Ser Pro Glu Tyr Arg
 115 120 125
 45 Ala Arg Val Val Arg Glu Pro Arg Arg Val Leu Arg Glu Met Gly Leu
 130 135 140
 Glu Leu Ser Glu Ser Val Thr Ile Arg Val Trp Asp Thr Thr Ala Glu
 145 150 155 160
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 Thr Arg Phe Leu Val Leu Pro Leu Arg Pro Ala Gly Thr Glu Gly Trp
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 cat ccg ggc gtc gcc aag gac gag aag gtc cac ggc tat tac caa ttg
 96
 His Pro Gly Val Ala Lys Asp Glu Lys Val His Gly Tyr Tyr Gln Leu
 20 25 30

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 ctc ggc ctc gcc atc aaa gag ctg ctg atc gaa aaa ggc gtc atc acc
 144
 Leu Gly Leu Ala Ile Lys Glu Leu Leu Ile Glu Lys Gly Val Ile Thr
 35 40 45

35
 gcc gcc gag gtg cgc caa gcg atc gag gcg cgc gac gcg atc acg ccg
 192
 Ala Ala Glu Val Arg Gln Ala Ile Glu Ala Arg Asp Ala Ile Thr Pro
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 240
 Ser Leu Gly Gly Lys Val Val Ala Arg Ala Trp Thr Asp Pro Ala Tyr
 65 70 75 80

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 Lys Ala Arg Leu Ile Ala Asp Pro Ala Ala Ala Met Met Glu Met Gly
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50
 gtc gat ctc ggc ccc acc gga ctc gcc atc gcc gag aac acg ccg gag
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 Val Asp Leu Gly Pro Thr Gly Leu Ala Ile Ala Glu Asn Thr Pro Glu
 100 105 110

55
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 384
 Ala His Asn Val Ile Val Cys Thr Leu Cys Ser Cys Tyr Pro Arg Ala
 115 120 125

gtg ctc ggc ctg ccg ccc tcc tgg tac aag gac cgc gat tac cgg tcg
 432
 Val Leu Gly Leu Pro Pro Ser Trp Tyr Lys Asp Arg Asp Tyr Arg Ser
 130 135 140
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 480
 Arg Val Val Arg Glu Pro Arg Ala Val Leu Lys Glu Phe Gly Thr Glu
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 528
 Leu Pro Asp Asp Val Asp Val Arg Val His Asp Ser Thr Ala Asp Leu
 165 170 175
 15 cgc tat ctc gtg ctg ccg atg cgc ccg gcc ggc acc gag ggc atg agc
 576
 Arg Tyr Leu Val Leu Pro Met Arg Pro Ala Gly Thr Glu Gly Met Ser
 180 185 190
 20 gag gcg gag ctg gcc gag atc gtg acg cgc gac tgc atg atc ggc gtg
 624
 Glu Ala Glu Leu Ala Glu Ile Val Thr Arg Asp Cys Met Ile Gly Val
 195 200 205
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 35 40 45
 Ala Ala Glu Val Arg Gln Ala Ile Glu Ala Arg Asp Ala Ile Thr Pro
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 Ser Leu Gly Gly Lys Val Val Ala Arg Ala Trp Thr Asp Pro Ala Tyr
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5 Lys Ala Arg Leu Ile Ala Asp Pro Ala Ala Ala Met Met Glu Met Gly
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 10 Ala His Asn Val Ile Val Cys Thr Leu Cys Ser Cys Tyr Pro Arg Ala
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 130 135 140
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 25 Leu Pro Asp Asp Val Asp Val Arg Val His Asp Ser Thr Ala Asp Leu
 165 170 175
 Arg Tyr Leu Val Leu Pro Met Arg Pro Ala Gly Thr Glu Gly Met Ser
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 96

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	Lys	Gly	Tyr	Val	Asp	Pro	Ala	Ala	Leu	Asp	Ala	Ile	Ile	Asp	Thr	Tyr	
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	Glu	Thr	Lys	Ile	Gly	Pro	Arg	Asn	Gly	Ala	Arg	Val	Val	Ala	Lys	Ala	
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	Ala	Val	Ala	Glu	Leu	Gly	Tyr	Gly	Gly	Arg	Gly	Gly	Glu	His	Ile	Val	
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	Cys	Ser	Cys	Tyr	Pro	Trp	Pro	Val	Leu	Gly	Leu	Pro	Pro	Val	Trp	Tyr	
			115				120						125				
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	Lys	Ser	Pro	Pro	Tyr	Arg	Ser	Lys	Ala	Val	Ile	Asp	Pro	Arg	Gly	Val	
		130					135					140					
40	ctg	gcc	gat	ttc	ggc	gtg	acc	ctg	ccg	gag	gga	caa	agg	atc	cgc	gtc	480
	Leu	Ala	Asp	Phe	Gly	Val	Thr	Leu	Pro	Glu	Gly	Gln	Arg	Ile	Arg	Val	
		145				150					155				160		
45	tgg	gat	toc	acc	gcc	gaa	acc	cgc	ttc	att	gtc	atc	ccc	ctg	cgc	ccg	528
	Trp	Asp	Ser	Thr	Ala	Glu	Thr	Arg	Phe	Ile	Val	Ile	Pro	Leu	Arg	Pro	
					165				170						175		
50	gcc	ggg	acg	gaa	ggc	tgg	tcg	gaa	gaa	cag	ctg	gcg	gcg	atc	gtg	acg	576
	Ala	Gly	Thr	Glu	Gly	Trp	Ser	Glu	Glu	Gln	Leu	Ala	Ala	Ile	Val	Thr	
				180					185					190			
55	cgt	gac	agc	atg	atc	ggc	acc	ggc	gtg	gtc	agc	gcg	gag	gct	tcg	cga	624
	Arg	Asp	Ser	Met	Ile	Gly	Thr	Gly	Val	Val	Ser	Ala	Glu	Ala	Ser	Arg	
			195					200					205				

tga
627

5 <210> 56
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 <212> PRT
 <213> Unknown

10 <220>
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 <400> 56

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20 Thr Pro Ile Glu Ala Arg Val Lys Ala Leu Glu Ser Leu Leu Val Ala
 20 25 30

25 Lys Gly Tyr Val Asp Pro Ala Ala Leu Asp Ala Ile Ile Asp Thr Tyr
 35 40 45

30 Glu Thr Lys Ile Gly Pro Arg Asn Gly Ala Arg Val Val Ala Lys Ala
 50 55 60

35 Trp Thr Asp Pro Glu Phe Ala Ala Arg Leu Lys Gln Asp Gly Ser Ala
 65 70 75 80

40 Ala Val Ala Glu Leu Gly Tyr Gly Gly Arg Gly Gly Glu His Ile Val
 85 90 95

45 Ala Cys Phe Asn Thr Pro Glu Glu His Asn Leu Ile Val Cys Thr Leu
 100 105 110

50 Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly Leu Pro Pro Val Trp Tyr
 115 120 125

55 Lys Ser Pro Pro Tyr Arg Ser Lys Ala Val Ile Asp Pro Arg Gly Val
 130 135 140

60 Leu Ala Asp Phe Gly Val Thr Leu Pro Glu Gly Gln Arg Ile Arg Val
 145 150 155 160

65 Trp Asp Ser Thr Ala Glu Thr Arg Phe Ile Val Ile Pro Leu Arg Pro
 165 170 175

Ala Gly Thr Glu Gly Trp Ser Glu Glu Gln Leu Ala Ala Ile Val Thr

	180	185	190
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5 Arg Asp Ser Met Ile Gly Thr Gly Val Val Ser Ala Glu Ala Ser Arg
 195 200 205

10 <210> 57
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 <212> DNA
 <213> Unknown

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 48
 Met Asp Pro Thr Arg Arg Ser Phe Leu Ala Ser Thr Val Ala Leu Thr
 1 5 10 15

30 ggc ggc gca gct atc ccc gat ctg gct cat gcg gca gac cac gac cac
 96
 Gly Gly Ala Ala Ile Pro Asp Leu Ala His Ala Ala Asp His Asp His
 20 25 30

35 cag cat caa gat ttg ccg tcc gat ctg gcg ctg cgg gtg aag tcg ctc
 144
 Gln His Gln Asp Leu Pro Ser Asp Leu Ala Leu Arg Val Lys Ser Leu
 35 40 45

40 gaa tcg ctg ctt gtc gag aag ggg ctg gtg gag cga gca gcg ctc gac
 192
 Glu Ser Leu Leu Val Glu Lys Gly Leu Val Glu Arg Ala Ala Leu Asp
 50 55 60

45 gcg ctg gtg gac acc tac gag cac aaa gtc ggg ccg cga aac gga gcg
 240
 Ala Leu Val Asp Thr Tyr Glu His Lys Val Gly Pro Arg Asn Gly Ala
 65 70 75 80

50 cgc gtt gtc gcg cgg gcc tgg gtt gac ccg gac tac aag caa cgg tta
 288
 Arg Val Val Ala Arg Ala Trp Val Asp Pro Asp Tyr Lys Gln Arg Leu
 85 90 95

55 ttc gcg aac ggt acc gcc gca gtc gcg gag ttc ggc tac tcc ggc tcg
 336
 Phe Ala Asn Gly Thr Ala Ala Val Ala Glu Phe Gly Tyr Ser Gly Ser
 100 105 110

cag ggc gct gac atc cgg gtc gtc gaa aac acg gcc act gta cat aac
 384
 Gln Gly Ala Asp Ile Arg Val Val Glu Asn Thr Ala Thr Val His Asn

	115	120	125
	ctc gtc gtg tgc acg ctg tgc tct tgt tat ccc tgg cgg gtg ctg ggc		
	432		
5	Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly		
	130	135	140
	ttg ccg ccg gtc tgg tac aag tcc gcg ccc tat cgg tct cgc gtg gtg		
	480		
10	Leu Pro Pro Val Trp Tyr Lys Ser Ala Pro Tyr Arg Ser Arg Val Val		
	145	150	155 160
	atc gat ccg cga ggt gtg ctg cgc gag ttc ggc gtg gtg ctg ccg gac		
	528		
15	Ile Asp Pro Arg Gly Val Leu Arg Glu Phe Gly Val Val Leu Pro Asp		
	165	170	175
	cat atc gaa gtg cgt gtc tat gac agc acg gcg gag caa cgc tat cta		
	576		
20	His Ile Glu Val Arg Val Tyr Asp Ser Thr Ala Glu Gln Arg Tyr Leu		
	180	185	190
	gtg ctg ccg gag cgg ccg gcc gga acc gaa aac ctg aca gaa gaa gcg		
	624		
25	Val Leu Pro Glu Arg Pro Ala Gly Thr Glu Asn Leu Thr Glu Glu Ala		
	195	200	205
	ctg gcg ctg ctg gtg acg cgc gac gcg atg att ggc gtg gcc aag gtc		
	672		
30	Leu Ala Leu Leu Val Thr Arg Asp Ala Met Ile Gly Val Ala Lys Val		
	210	215	220
	gcg ccg ccg gga ggc cgc gga tga		
	696		
35	Ala Pro Pro Gly Gly Arg Gly		
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40	<211> 231		
	<212> PRT		
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	<400> 58		
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	Gly Gly Ala Ala Ile Pro Asp Leu Ala His Ala Ala Asp His Asp His		
	20	25	30
55	Gln His Gln Asp Leu Pro Ser Asp Leu Ala Leu Arg Val Lys Ser Leu		
	35	40	45

Glu Ser Leu Leu Val Glu Lys Gly Leu Val Glu Arg Ala Ala Leu Asp
 50 55 60

5

Ala Leu Val Asp Thr Tyr Glu His Lys Val Gly Pro Arg Asn Gly Ala
 65 70 75 80

10

Arg Val Val Ala Arg Ala Trp Val Asp Pro Asp Tyr Lys Gln Arg Leu
 85 90 95

15

Phe Ala Asn Gly Thr Ala Ala Val Ala Glu Phe Gly Tyr Ser Gly Ser
 100 105 110

20

Gln Gly Ala Asp Ile Arg Val Val Glu Asn Thr Ala Thr Val His Asn
 115 120 125

Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly
 130 135 140

25

Leu Pro Pro Val Trp Tyr Lys Ser Ala Pro Tyr Arg Ser Arg Val Val
 145 150 155 160

30

Ile Asp Pro Arg Gly Val Leu Arg Glu Phe Gly Val Val Leu Pro Asp
 165 170 175

35

His Ile Glu Val Arg Val Tyr Asp Ser Thr Ala Glu Gln Arg Tyr Leu
 180 185 190

40

Val Leu Pro Glu Arg Pro Ala Gly Thr Glu Asn Leu Thr Glu Glu Ala
 195 200 205

Leu Ala Leu Leu Val Thr Arg Asp Ala Met Ile Gly Val Ala Lys Val
 210 215 220

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Ala Pro Pro Gly Gly Arg Gly
 225 230

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<210> 59
 <211> 609
 <212> DNA
 <213> Unknown

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<222> (1)..(609)

<223>

<400> 59

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Met Ser Ser Lys Pro Thr Glu Asp Leu Gly Thr Tyr Gln Pro Leu Thr
1 5 10 15

10 tac tac cag atg atg gaa gtg agc ctg cgc gag ctg ctg gtg gag aag
96
Tyr Tyr Gln Met Met Glu Val Ser Leu Arg Glu Leu Leu Val Glu Lys
20 25 30

15 ggc gtg atc acc gaa gcg gaa gtc gcc cgc gcg atg ggc gag atc ggc
144
Gly Val Ile Thr Glu Ala Glu Val Ala Arg Ala Met Gly Glu Ile Gly
35 40 45

20 gcg aga agc ccg gag cgc ggc gcg aag atg gtc gcg cgc gcg tgg gtg
192
Ala Arg Ser Pro Glu Arg Gly Ala Lys Met Val Ala Arg Ala Trp Val
50 55 60

25 gac ccg gcg tac aag gcg cgc atg ctt gcc gac ggc agc aag gcc gcc
240
Asp Pro Ala Tyr Lys Ala Arg Met Leu Ala Asp Gly Ser Lys Ala Ala
65 70 75 80

30 gag gag ctc ggg ttc gag gtg ccg ggc ctc aag ctg atc gtg gtc gag
288
Glu Glu Leu Gly Phe Glu Val Pro Gly Leu Lys Leu Ile Val Val Glu
85 90 95

35 aac acc gcg gac acg cac aac gtg gtc gtg tgc acg ctg tgc tcg tgc
336
Asn Thr Ala Asp Thr His Asn Val Val Val Cys Thr Leu Cys Ser Cys
100 105 110

40 tac ccg cgc atc ctg ctc ggc atc ccg ccc gag tgg tac aag tcg cgc
384
Tyr Pro Arg Ile Leu Leu Gly Ile Pro Pro Glu Trp Tyr Lys Ser Arg
115 120 125

45 agc tac cgc agc cgc aca gtg cgc gag ccg cgc gcg gtg ctc gcc gaa
432
Ser Tyr Arg Ser Arg Thr Val Arg Glu Pro Arg Ala Val Leu Ala Glu
130 135 140

50 ttc ggc acg acc atc ccg gag aac gtc gcg atc cga gtg cac gac agc
480
Phe Gly Thr Thr Ile Pro Glu Asn Val Ala Ile Arg Val His Asp Ser
145 150 155 160

55 act gcg gac atg cgc tac ctc gtg atg ccg atg cgg cct gcg ggc acc
528
Thr Ala Asp Met Arg Tyr Leu Val Met Pro Met Arg Pro Ala Gly Thr
165 170 175

gaa aat ttc acc gaa gag cag ctc gct gca ttg gtg acg cgc gac agc
 576
 Glu Asn Phe Thr Glu Glu Gln Leu Ala Ala Leu Val Thr Arg Asp Ser
 180 185 190
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 ctg atc ggt gtt tcc tta gca acg ctt ccg tag
 609
 Leu Ile Gly Val Ser Leu Ala Thr Leu Pro
 195 200
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 15 <213> Unknown
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 20 25 30
 30 Gly Val Ile Thr Glu Ala Glu Val Ala Arg Ala Met Gly Glu Ile Gly
 35 40 45
 35 Ala Arg Ser Pro Glu Arg Gly Ala Lys Met Val Ala Arg Ala Trp Val
 50 55 60
 Asp Pro Ala Tyr Lys Ala Arg Met Leu Ala Asp Gly Ser Lys Ala Ala
 65 70 75 80
 40 Glu Glu Leu Gly Phe Glu Val Pro Gly Leu Lys Leu Ile Val Val Glu
 85 90 95
 45 Asn Thr Ala Asp Thr His Asn Val Val Val Cys Thr Leu Cys Ser Cys
 100 105 110
 50 Tyr Pro Arg Ile Leu Leu Gly Ile Pro Pro Glu Trp Tyr Lys Ser Arg
 115 120 125
 55 Ser Tyr Arg Ser Arg Thr Val Arg Glu Pro Arg Ala Val Leu Ala Glu
 130 135 140
 Phe Gly Thr Thr Ile Pro Glu Asn Val Ala Ile Arg Val His Asp Ser
 145 150 155 160

Thr Ala Asp Met Arg Tyr Leu Val Met Pro Met Arg Pro Ala Gly Thr
 165 170 175
 5
 Glu Asn Phe Thr Glu Glu Gln Leu Ala Ala Leu Val Thr Arg Asp Ser
 180 185 190
 10
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 195 200
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 <212> DNA
 <213> Unknown
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 25 <222> (1)..(825)
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 30 48
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 1 5 10 15
 gct gcc aac tgg caa ctt atc agt cgc cct tca tgg tca gct tgt aat
 35 96
 Ala Ala Asn Trp Gln Leu Ile Ser Arg Pro Ser Trp Ser Ala Cys Asn
 20 25 30
 ata ttg gtc ctc atg agc gcc acg cac ccc aaa aag cgc gcc gcc gac
 40 144
 Ile Leu Val Leu Met Ser Ala Thr His Pro Lys Lys Arg Ala Ala Asp
 35 40 45
 atc ggc ggc aac aaa gcc ggc gcg gtg gac acc gcg gat cac ggc atg
 45 192
 Ile Gly Gly Asn Lys Ala Gly Ala Val Asp Thr Ala Asp His Gly Met
 50 55 60
 aag ttc tgg gag cgg cag gcc aac gcc ctg cgc acc gcg ctg cgg cgc
 50 240
 Lys Phe Trp Glu Arg Gln Ala Asn Ala Leu Arg Thr Ala Leu Arg Arg
 65 70 75 80
 aat gga ctg atg agc gta gat gag ctg cgc cgc gca gcg gag gac ctg
 55 288
 Asn Gly Leu Met Ser Val Asp Glu Leu Arg Arg Ala Ala Glu Asp Leu
 85 90 95

gga gac cgc tac gcg aag ctt gag tac ttc gag cgc acg acg ttc gcg
 336
 Gly Asp Arg Tyr Ala Lys Leu Glu Tyr Phe Glu Arg Thr Thr Phe Ala
 100 105 110
 5
 ctg cgc acg gtc ctg ctc gaa aag ggc tac ttc acg gag gag tcg ctc
 384
 Leu Arg Thr Val Leu Leu Glu Lys Gly Tyr Phe Thr Glu Glu Ser Leu
 115 120 125
 10
 gcg gcg aag atg gcc gag gtg cgg aag ccg ctt cga tgt gcc gcg caa
 432
 Ala Ala Lys Met Ala Glu Val Arg Lys Pro Leu Arg Cys Ala Ala Gln
 130 135 140
 15
 gaa gga att gcc ggt gaa gaa gaa agt gaa gcg atg aac cca gcg acg
 480
 Glu Gly Ile Ala Gly Glu Glu Glu Ser Glu Ala Met Asn Pro Ala Thr
 145 150 155 160
 20
 ggc aag cag gac ggc caa cgg ctg cca tct acg tat acc gcg gcg ccc
 528
 Gly Lys Gln Asp Gly Gln Arg Leu Pro Ser Thr Tyr Thr Ala Ala Pro
 165 170 175
 25
 ggg cac cga ttc gat gtc ggt gac cgc gtt gtg gtc aag cgc tca aat
 576
 Gly His Arg Phe Asp Val Gly Asp Arg Val Val Val Lys Arg Ser Asn
 180 185 190
 30
 ccg ccc ggc cac cgc cgc acg cct cat tac atc cgc ggc aag acg ggc
 624
 Pro Pro Gly His Arg Arg Thr Pro His Tyr Ile Arg Gly Lys Thr Gly
 195 200 205
 35
 gtg atc gag cgc atc tgc ggc gcc ttc ccc aac ccg gaa gag ctg gca
 672
 Val Ile Glu Arg Ile Cys Gly Ala Phe Pro Asn Pro Glu Glu Leu Ala
 210 215 220
 40
 tac gga ttc gac ggc gaa ccg aag aag gtg ctc tac cgc gtg cga ttc
 720
 Tyr Gly Phe Asp Gly Glu Pro Lys Lys Val Leu Tyr Arg Val Arg Phe
 225 230 235 240
 45
 cgg caa aaa gag gtg tgg ccg gcc tat cgc ggc ccg gcg cac gac gtg
 768
 Arg Gln Lys Glu Val Trp Pro Ala Tyr Arg Gly Pro Ala His Asp Val
 245 250 255
 50
 atc gag atg gag att ttc gag cat tgg ctc gag ccg gcg cag agc cag
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 Ile Glu Met Glu Ile Phe Glu His Trp Leu Glu Pro Ala Gln Ser Gln
 260 265 270
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 aaa acc tga
 825
 Lys Thr

5 <210> 62
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 <400> 62

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 1 5 10 15

Ala Ala Asn Trp Gln Leu Ile Ser Arg Pro Ser Trp Ser Ala Cys Asn
 20 25 30

20 Ile Leu Val Leu Met Ser Ala Thr His Pro Lys Lys Arg Ala Ala Asp
 35 40 45

25 Ile Gly Gly Asn Lys Ala Gly Ala Val Asp Thr Ala Asp His Gly Met
 50 55 60

30 Lys Phe Trp Glu Arg Gln Ala Asn Ala Leu Arg Thr Ala Leu Arg Arg
 65 70 75 80

Asn Gly Leu Met Ser Val Asp Glu Leu Arg Arg Ala Ala Glu Asp Leu
 85 90 95

35 Gly Asp Arg Tyr Ala Lys Leu Glu Tyr Phe Glu Arg Thr Thr Phe Ala
 100 105 110

40 Leu Arg Thr Val Leu Leu Glu Lys Gly Tyr Phe Thr Glu Glu Ser Leu
 115 120 125

45 Ala Ala Lys Met Ala Glu Val Arg Lys Pro Leu Arg Cys Ala Ala Gln
 130 135 140

50 Glu Gly Ile Ala Gly Glu Glu Glu Ser Glu Ala Met Asn Pro Ala Thr
 145 150 155 160

Gly Lys Gln Asp Gly Gln Arg Leu Pro Ser Thr Tyr Thr Ala Ala Pro
 165 170 175

55 Gly His Arg Phe Asp Val Gly Asp Arg Val Val Val Lys Arg Ser Asn
 180 185 190

Pro Pro Gly His Arg Arg Thr Pro His Tyr Ile Arg Gly Lys Thr Gly
 195 200 205

5

Val Ile Glu Arg Ile Cys Gly Ala Phe Pro Asn Pro Glu Glu Leu Ala
 210 215 220

10

Tyr Gly Phe Asp Gly Glu Pro Lys Lys Val Leu Tyr Arg Val Arg Phe
 225 230 235 240

15

Arg Gln Lys Glu Val Trp Pro Ala Tyr Arg Gly Pro Ala His Asp Val
 245 250 255

20

Ile Glu Met Glu Ile Phe Glu His Trp Leu Glu Pro Ala Gln Ser Gln
 260 265 270

Lys Thr

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<210> 63
 <211> 627
 <212> DNA
 <213> Unknown

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 <222> (1)..(627)
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 Met Asp Gly Met His Asp Leu Gly Gly Arg Gln Gly Phe Gly Pro Val
 1 5 10 15

45

cgc tac acg atc gac gcg ccc gca ttc cat tcg ccg tgg gaa gtg cgc
 96
 Arg Tyr Thr Ile Asp Ala Pro Ala Phe His Ser Pro Trp Glu Val Arg
 20 25 30

50

gcg aat tcg ctc tat gcg ttc gcg gtg cgc ctc ggc atc ttc aac atg
 144
 Ala Asn Ser Leu Tyr Ala Phe Ala Val Arg Leu Gly Ile Phe Asn Met
 35 40 45

55

gac gaa tac cgc cat gcg atc gag cgg atg gag ccg cgc cat tac ctc
 192
 Asp Glu Tyr Arg His Ala Ile Glu Arg Met Glu Pro Arg His Tyr Leu
 50 55 60

ggc gcc ggc tat tac gaa cgc tcg ttg acc ggc ctc gcg acc ttg ctg
 240
 Gly Ala Gly Tyr Tyr Glu Arg Ser Leu Thr Gly Leu Ala Thr Leu Leu
 65 70 75 80
 5
 gtc gag aag ggc gtc gtg acg cgc gag gaa ctc gag acc cgg gcg cag
 288
 Val Glu Lys Gly Val Val Thr Arg Glu Glu Leu Glu Thr Arg Ala Gln
 85 90 95
 10
 ggc cgc tac ccg ctg gcg atg ccc agc gcg cct ggc cgc acc aat gcg
 336
 Gly Arg Tyr Pro Leu Ala Met Pro Ser Ala Pro Gly Arg Thr Asn Ala
 100 105 110
 15
 cag gca cgc gag cgt ttc cag ccg ggc gac cgg gtt cgc gtc aag gcg
 384
 Gln Ala Arg Glu Arg Phe Gln Pro Gly Asp Arg Val Arg Val Lys Ala
 115 120 125
 20
 gat ttc gtg tcg ggg cac gtg cgg atg ccg gcg tac atc cgc ggc aag
 432
 Asp Phe Val Ser Gly His Val Arg Met Pro Ala Tyr Ile Arg Gly Lys
 130 135 140
 25
 acc ggc gtg gtc gtc agc gag tcc ccg gac tat ccg ttt ccc gat gcg
 480
 Thr Gly Val Val Val Ser Glu Ser Pro Asp Tyr Pro Phe Pro Asp Ala
 145 150 155 160
 30
 cat gcg cac tcg gtc gat gcc cag gac gag cca acc tac gac gtg cgc
 528
 His Ala His Ser Val Asp Ala Gln Asp Glu Pro Thr Tyr Asp Val Arg
 165 170 175
 35
 ttc cgc agc gag gat cta tgg ccg gat tcc gcc gat tcc gca ctc gtt
 576
 Phe Arg Ser Glu Asp Leu Trp Pro Asp Ser Ala Asp Ser Ala Leu Val
 180 185 190
 40
 cac gtc ggc gta ttc cag agc tac ctc gag cgg gag tcg acg cca gga
 624
 His Val Gly Val Phe Gln Ser Tyr Leu Glu Arg Glu Ser Thr Pro Gly
 195 200 205
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 627
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 5 Arg Tyr Thr Ile Asp Ala Pro Ala Phe His Ser Pro Trp Glu Val Arg
 20 25 30
 10 Ala Asn Ser Leu Tyr Ala Phe Ala Val Arg Leu Gly Ile Phe Asn Met
 35 40 45
 15 Asp Glu Tyr Arg His Ala Ile Glu Arg Met Glu Pro Arg His Tyr Leu
 50 55 60
 Gly Ala Gly Tyr Tyr Glu Arg Ser Leu Thr Gly Leu Ala Thr Leu Leu
 65 70 75 80
 20 Val Glu Lys Gly Val Val Thr Arg Glu Glu Leu Glu Thr Arg Ala Gln
 85 90 95
 25 Gly Arg Tyr Pro Leu Ala Met Pro Ser Ala Pro Gly Arg Thr Asn Ala
 100 105 110
 30 Gln Ala Arg Glu Arg Phe Gln Pro Gly Asp Arg Val Arg Val Lys Ala
 115 120 125
 35 Asp Phe Val Ser Gly His Val Arg Met Pro Ala Tyr Ile Arg Gly Lys
 130 135 140
 Thr Gly Val Val Val Ser Glu Ser Pro Asp Tyr Pro Phe Pro Asp Ala
 145 150 155 160
 40 His Ala His Ser Val Asp Ala Gln Asp Glu Pro Thr Tyr Asp Val Arg
 165 170 175
 45 Phe Arg Ser Glu Asp Leu Trp Pro Asp Ser Ala Asp Ser Ala Leu Val
 180 185 190
 50 His Val Gly Val Phe Gln Ser Tyr Leu Glu Arg Glu Ser Thr Pro Gly
 195 200 205
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10 Met Asn Gly Val His Asp Met Gly Gly Met His Gly Met Gly Ala Ile
1 5 10 15

cgc cgc gag gag aac gag ccc gct ttc cac gag ccc tgg gag ggg cgg
96
15 Arg Arg Glu Glu Asn Glu Pro Ala Phe His Glu Pro Trp Glu Gly Arg
20 25 30

gtt ttc gct ctg acc acg gcg gtc gag gcc tgg ggt cgg tgg acc ctg
144
20 Val Phe Ala Leu Thr Thr Ala Val Glu Ala Trp Gly Arg Trp Thr Leu
35 40 45

gat gct tcc cga cac cgg atc gag cgg atg aat gcg gcg gac tat ctg
192
25 Asp Ala Ser Arg His Arg Ile Glu Arg Met Asn Ala Ala Asp Tyr Leu
50 55 60

cgg gtg agc tat tac gag aag tgg ctc gag tcg ctt ctc gct ctc ctg
240
30 Arg Val Ser Tyr Tyr Glu Lys Trp Leu Glu Ser Leu Leu Ala Leu Leu
65 70 75 80

tcc gag acc gga atg gcg agt ccg gcg gag ata cgg agt ggg gag cgt
288
35 Ser Glu Thr Gly Met Ala Ser Pro Ala Glu Ile Arg Ser Gly Glu Arg
85 90 95

gcc gac ggc aca ccg aaa gcg acc ccg ccg ctc ccg gcc gac cac gtg
336
40 Ala Asp Gly Thr Pro Lys Ala Thr Pro Pro Leu Pro Ala Asp His Val
100 105 110

acg gcg att ctc gcc agc ggg ttt ccc gcg agc cgg gag gcg gga gct
384
45 Thr Ala Ile Leu Ala Ser Gly Phe Pro Ala Ser Arg Glu Ala Gly Ala
115 120 125

gcg cct cgc ttc cga gtg agc gag cgg gtg cgc acc cgg aac atc aac
432
50 Ala Pro Arg Phe Arg Val Ser Glu Arg Val Arg Thr Arg Asn Ile Asn
130 135 140

ccg acg act cac acg gcg ctt ccg cga tac gcc cgg cgg aag ctc ggg
480
55 Pro Thr Thr His Thr Arg Leu Pro Arg Tyr Ala Arg Arg Lys Leu Gly
145 150 155 160

acg atc gag cgc gac cac gga gtg ttc gtc ttc ccg gat acg aac gcg
528

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Thr Ile Glu Arg Asp His Gly Val Phe Val Phe Pro Asp Thr Asn Ala
 165 170 175
 5 cac gct ctc ggg gag aaa ccg cag cac gtc tat tcg gtt cgt ttc gag
 576
 His Ala Leu Gly Glu Lys Pro Gln His Val Tyr Ser Val Arg Phe Glu
 180 185 190
 10 gcg cgt gag ctc tgg ggc gag act gcc agg cca gag gat tcc gtc tac
 624
 Ala Arg Glu Leu Trp Gly Glu Thr Ala Arg Pro Glu Asp Ser Val Tyr
 195 200 205
 15 atc gat ctt tgg gac gag tac ctt gaa ccc gtg tag
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 Ile Asp Leu Trp Asp Glu Tyr Leu Glu Pro Val
 210 215
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 20 25 30
 40 Val Phe Ala Leu Thr Thr Ala Val Glu Ala Trp Gly Arg Trp Thr Leu
 35 40 45
 Asp Ala Ser Arg His Arg Ile Glu Arg Met Asn Ala Ala Asp Tyr Leu
 50 55 60
 45 Arg Val Ser Tyr Tyr Glu Lys Trp Leu Glu Ser Leu Leu Ala Leu Leu
 65 70 75 80
 50 Ser Glu Thr Gly Met Ala Ser Pro Ala Glu Ile Arg Ser Gly Glu Arg
 85 90 95
 55 Ala Asp Gly Thr Pro Lys Ala Thr Pro Pro Leu Pro Ala Asp His Val
 100 105 110
 Thr Ala Ile Leu Ala Ser Gly Phe Pro Ala Ser Arg Glu Ala Gly Ala
 115 120 125

	Ala	Pro	Arg	Phe	Arg	Val	Ser	Glu	Arg	Val	Arg	Thr	Arg	Asn	Ile	Asn	
	130						135							140			
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	Pro	Thr	Thr	His	Thr	Arg	Leu	Pro	Arg	Tyr	Ala	Arg	Arg	Lys	Leu	Gly	
	145					150					155					160	
10																	
	Thr	Ile	Glu	Arg	Asp	His	Gly	Val	Phe	Val	Phe	Pro	Asp	Thr	Asn	Ala	
					165					170					175		
15																	
	His	Ala	Leu	Gly	Glu	Lys	Pro	Gln	His	Val	Tyr	Ser	Val	Arg	Phe	Glu	
				180					185					190			
20																	
	Ala	Arg	Glu	Leu	Trp	Gly	Glu	Thr	Ala	Arg	Pro	Glu	Asp	Ser	Val	Tyr	
			195					200					205				
25																	
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	Met	Thr	Asn	Ser	Leu	His	Asp	Met	Gly	Gly	Met	His	Gly	Phe	Gly	Arg	
	1				5					10					15		
50																	
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	96																
	Val	Glu	Pro	Glu	Pro	Asn	Glu	Pro	Pro	Phe	His	Gln	Arg	Trp	Glu	Gly	
				20					25					30			
55																	
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ctg ggt tgg tcc tac tat cgg cgc tgg ttt ctt ggc ctg gag agc cgg
 240
 5 Leu Gly Ser Ser Tyr Tyr Arg Arg Trp Phe Leu Gly Leu Glu Ser Arg
 65 70 75 80

 ctg ctg ctg cgc ggc ctc gtt ggc gag gac gag atc gcg gca ggc cgt
 288
 10 Leu Leu Leu Arg Gly Leu Val Gly Glu Asp Glu Ile Ala Ala Gly Arg
 85 90 95

 tgg atg cgc gcc ggc gcc atg ttg ccg cgc acc ctg acc cag gcc gat
 336
 15 Ser Met Arg Ala Gly Ala Met Leu Pro Arg Thr Leu Thr Gln Ala Asp
 100 105 110

 gtg gag aaa acc ctg acc cgc ggc gac ttc gcc cgc ccg acc aac acc
 384
 20 Val Glu Lys Thr Leu Thr Arg Gly Asp Phe Ala Arg Pro Thr Asn Thr
 115 120 125

 ccg gcg cgt ttc cag ccg ggc gac ccg gtg caa acg aag aac atc aac
 432
 25 Pro Ala Arg Phe Gln Pro Gly Asp Arg Val Gln Thr Lys Asn Ile Asn
 130 135 140

 ccg gcg acc cac acc cgc ctg ccg cgc tat gcc cgc ggc aag act ggc
 480
 30 Pro Ala Thr His Thr Arg Leu Pro Arg Tyr Ala Arg Gly Lys Thr Gly
 145 150 155 160

 acg gtc gag gcg gtc cgc ggc gtt cac gtc ttt ccc gac acc gcc gcg
 528
 35 Thr Val Glu Ala Val Arg Gly Val His Val Phe Pro Asp Thr Ala Ala
 165 170 175

 ctc ggc gcc ggc gac gac ccg caa tgg ctc tac gcc gtg gtc ttc ccg
 576
 40 Leu Gly Ala Gly Asp Asp Pro Gln Trp Leu Tyr Ala Val Val Phe Pro
 180 185 190

 gcg cgc gag ttg tgg gga gag gcg gcc gat ccc gcg atc aaa atc tgg
 624
 45 Ala Arg Glu Leu Trp Gly Glu Ala Ala Asp Pro Ala Ile Lys Ile Ser
 195 200 205

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 660
 50 Ile Glu Ala Phe Glu Pro Tyr Ile Asp Pro Ala
 210 215

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 55 <212> PRT
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15	Arg	Val	Leu	Gly	Met	Gln	Arg	Ala	Met	Gly	Phe	Thr	Gly	Leu	Trp	Thr	35	40	45	
20	Ile	Asp	Ala	Gly	Arg	Ala	Ser	Leu	Glu	Ala	Leu	Pro	Pro	Leu	Ala	Tyr	50	55	60	
25	Leu	Gly	Ser	Ser	Tyr	Tyr	Arg	Arg	Trp	Phe	Leu	Gly	Leu	Glu	Ser	Arg	65	70	75	
30	Leu	Leu	Leu	Arg	Gly	Leu	Val	Gly	Glu	Asp	Glu	Ile	Ala	Ala	Gly	Arg	85	90	95	
35	Ser	Met	Arg	Ala	Gly	Ala	Met	Leu	Pro	Arg	Thr	Leu	Thr	Gln	Ala	Asp	100	105	110	
40	Val	Glu	Lys	Thr	Leu	Thr	Arg	Gly	Asp	Phe	Ala	Arg	Pro	Thr	Asn	Thr	115	120	125	
45	Pro	Ala	Arg	Phe	Gln	Pro	Gly	Asp	Arg	Val	Gln	Thr	Lys	Asn	Ile	Asn	130	135	140	
50	Pro	Ala	Thr	His	Thr	Arg	Leu	Pro	Arg	Tyr	Ala	Arg	Gly	Lys	Thr	Gly	145	150	155	
55	Thr	Val	Glu	Ala	Val	Arg	Gly	Val	His	Val	Phe	Pro	Asp	Thr	Ala	Ala	165	170	175	
	Leu	Gly	Ala	Gly	Asp	Asp	Pro	Gln	Trp	Leu	Tyr	Ala	Val	Val	Phe	Pro	180	185	190	
	Ala	Arg	Glu	Leu	Trp	Gly	Glu	Ala	Ala	Asp	Pro	Ala	Ile	Lys	Ile	Ser	195	200	205	
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20

gag ccg gaa gcg aac gag ccc gcc ttc cat gcg gaa tgg gaa ggc cgc
 96
 Glu Pro Glu Ala Asn Glu Pro Ala Phe His Ala Glu Trp Glu Gly Arg
 20 25 30

25

tgc ctc gcg ctc aac cgc gcc atg ggt gcg atc ggc gcc tgg acc atc
 144
 Cys Leu Ala Leu Asn Arg Ala Met Gly Ala Ile Gly Ala Trp Thr Ile
 35 40 45

30

gat gaa ggc cgt gcc ggc atc gag atc ctg ccg ccg gag att tat ctt
 192
 Asp Glu Gly Arg Ala Gly Ile Glu Ile Leu Pro Pro Glu Ile Tyr Leu
 50 55 60

35

ggc agt tcg tac tat gga aaa tgg gcg cgg cgg ctg gag aat atg gtg
 240
 Gly Ser Ser Tyr Tyr Gly Lys Trp Ala Arg Arg Leu Glu Asn Met Val
 65 70 75 80

40

gtc gca cgc ggg ttc gcg ggc gcc gat gaa ctc gcc gcg ggt cgc gca
 288
 Val Ala Arg Gly Phe Ala Gly Ala Asp Glu Leu Ala Ala Gly Arg Ala
 85 90 95

45

gcg cgt ccc ggc aga tcg gtg aaa cga aag ctt acg gtc gcc gaa gtg
 336
 Ala Arg Pro Gly Arg Ser Val Lys Arg Lys Leu Thr Val Ala Glu Val
 100 105 110

50

ccg cgc acg ctg acg cgc ggt tca ttt ttc cgc gag gca aca aag ccg
 384
 Pro Arg Thr Leu Thr Arg Gly Ser Phe Phe Arg Glu Ala Thr Lys Pro
 115 120 125

55

gca cga ttt gcg gtc ggc gaa cgc gtg cgc acc agg aac att cat ccg
 432
 Ala Arg Phe Ala Val Gly Glu Arg Val Arg Thr Arg Asn Ile His Pro
 130 135 140

gcg acg cac act cgg ttg ccg cga tat gcg cgc ggc cat gtc ggc gtg
 480
 Ala Thr His Thr Arg Leu Pro Arg Tyr Ala Arg Gly His Val Gly Val
 145 150 155 160
 5
 atc gag gcg atc cgc ggt tgc cac gta ttt ccc gac tcg gtt gcg atc
 528
 Ile Glu Ala Ile Arg Gly Cys His Val Phe Pro Asp Ser Val Ala Ile
 165 170 175
 10
 ggc gcc ggc gag aac ccg caa tgg ctt tat acg gtg gtg ttc gaa ggc
 576
 Gly Ala Gly Glu Asn Pro Gln Trp Leu Tyr Thr Val Val Phe Glu Gly
 180 185 190
 15
 cgc acg ctg tgg ggc gat agc gcc gat ccg acg ctt aag gtc tcg atc
 624
 Arg Thr Leu Trp Gly Asp Ser Ala Asp Pro Thr Leu Lys Val Ser Ile
 195 200 205
 20
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 663
 Glu Ala Phe Glu Pro Tyr Leu Glu Pro Ala Gln Pro
 210 215 220
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 30 <213> Unknown
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 20 25 30
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 Cys Leu Ala Leu Asn Arg Ala Met Gly Ala Ile Gly Ala Trp Thr Ile
 35 40 45
 50
 Asp Glu Gly Arg Ala Gly Ile Glu Ile Leu Pro Pro Glu Ile Tyr Leu
 50 55 60
 55
 Gly Ser Ser Tyr Tyr Gly Lys Trp Ala Arg Arg Leu Glu Asn Met Val
 65 70 75 80
 Val Ala Arg Gly Phe Ala Gly Ala Asp Glu Leu Ala Ala Gly Arg Ala
 85 90 95

Ala Arg Pro Gly Arg Ser Val Lys Arg Lys Leu Thr Val Ala Glu Val
100 105 110

5 Pro Arg Thr Leu Thr Arg Gly Ser Phe Phe Arg Glu Ala Thr Lys Pro
115 120 125

10 Ala Arg Phe Ala Val Gly Glu Arg Val Arg Thr Arg Asn Ile His Pro
130 135 140

15 Ala Thr His Thr Arg Leu Pro Arg Tyr Ala Arg Gly His Val Gly Val
145 150 155 160

20 Ile Glu Ala Ile Arg Gly Cys His Val Phe Pro Asp Ser Val Ala Ile
165 170 175

Gly Ala Gly Glu Asn Pro Gln Trp Leu Tyr Thr Val Val Phe Glu Gly
180 185 190

25 Arg Thr Leu Trp Gly Asp Ser Ala Asp Pro Thr Leu Lys Val Ser Ile
195 200 205

30 Glu Ala Phe Glu Pro Tyr Leu Glu Pro Ala Gln Pro
210 215 220

35 <210> 71
<211> 888
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40 <220>
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45 <223>

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1 5 10 15

atg gcg gag gcg gac gag ccg gtc ttt cat gag ccc tgg gaa ggt cgc
96
55 Met Ala Glu Ala Asp Glu Pro Val Phe His Glu Pro Trp Glu Gly Arg
20 25 30

gtg ttt gcg ctc aac atg ctc ggc atc ggg cgc gag ccc att ccg gtg
144

Val Phe Ala Leu Asn Met Leu Gly Ile Gly Arg Glu Pro Ile Pro Val
 35 40 45
 5 gac gcg ctg cgc cat cgc att gag cgg atc gag ccg tgg cgc tat ctg
 192
 Asp Ala Leu Arg His Arg Ile Glu Arg Ile Glu Pro Trp Arg Tyr Leu
 50 55 60
 10 acg tcg agc tat tac gaa cga tgg ctg gcc gaa atg gag cag gcc atc
 240
 Thr Ser Ser Tyr Tyr Glu Arg Trp Leu Ala Glu Met Glu Gln Ala Ile
 65 70 75 80
 15 atc gat gcg ggc acg ctg act cct ggt gaa atc gat gcg cga atg ggc
 288
 Ile Asp Ala Gly Thr Leu Thr Pro Gly Glu Ile Asp Ala Arg Met Gly
 85 90 95
 20 gag ctc gaa acg gat cct gat cgc cca ctg cca agg act gat aac cct
 336
 Glu Leu Glu Thr Asp Pro Asp Arg Pro Leu Pro Arg Thr Asp Asn Pro
 100 105 110
 25 gag cat gcc gat ggg gtg gcg gcg gcg ttg cgc gcc ggc agt ccc gta
 384
 Glu His Ala Asp Gly Val Ala Ala Ala Leu Arg Ala Gly Ser Pro Val
 115 120 125
 30 acg cgc aag att cgc aag cag ccg cgc ttc aca atc ggc gat cgg gtc
 432
 Thr Arg Lys Ile Arg Lys Gln Pro Arg Phe Thr Ile Gly Asp Arg Val
 130 135 140
 35 gta acg cgc aat ctt aat ccg cac ggc cat acg cgg ctg ccg cgc tat
 480
 Val Thr Arg Asn Leu Asn Pro His Gly His Thr Arg Leu Pro Arg Tyr
 145 150 155 160
 40 gcg cgc ggc aag cgc ggc gtc gta acg ctg cac cat ggc gca cat gtc
 528
 Ala Arg Gly Lys Arg Gly Val Val Thr Leu His His Gly Ala His Val
 165 170 175
 45 ttt ccg gat acg aac gcg cac ggg ctg ggc gag cat ccg cag cat ctc
 576
 Phe Pro Asp Thr Asn Ala His Gly Leu Gly Glu His Pro Gln His Leu
 180 185 190
 50 tat acg gtg cga ttt cct gcg cgc gag tta tgg agc gac gcg gcc gag
 624
 Tyr Thr Val Arg Phe Pro Ala Arg Glu Leu Trp Ser Asp Ala Ala Glu
 195 200 205
 55 ccg aaa gaa tcg ata atg atc gat ttg tgg gag agc tat ctt caa ccc
 672
 Pro Lys Glu Ser Ile Met Ile Asp Leu Trp Glu Ser Tyr Leu Gln Pro
 210 215 220

gat atc ggc agc aaa gcg tcg tcg tcc gcg aaa ggc aaa gcg acg ccg
 720
 Asp Ile Gly Ser Lys Ala Ser Ser Ser Ala Lys Gly Lys Ala Thr Pro
 225 230 235 240
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 768
 Lys Val Lys Pro Ala Met Ala Lys Ala Thr Ala Lys Val Ser Val Ser
 245 250 255
 10
 gcc aag gcc aaa act cgg gga aag gcg gcg ccg aag gag cgt cca aaa
 816
 Ala Lys Ala Lys Thr Arg Gly Lys Ala Ala Pro Lys Glu Arg Pro Lys
 260 265 270
 15
 ctg aaa cct gcg cga gcg gcg acc tca gca gca tcc ggc ggc gaa aaa
 864
 Leu Lys Pro Ala Arg Ala Ala Thr Ser Ala Ala Ser Gly Gly Glu Lys
 275 280 285
 20
 gct aag cga aag gcc aaa cga tga
 888
 Ala Lys Arg Lys Ala Lys Arg
 290 295
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 30 <213> Unknown
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 20 25 30
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 35 40 45
 50
 Asp Ala Leu Arg His Arg Ile Glu Arg Ile Glu Pro Trp Arg Tyr Leu
 50 55 60
 55
 Thr Ser Ser Tyr Tyr Glu Arg Trp Leu Ala Glu Met Glu Gln Ala Ile
 65 70 75 80
 Ile Asp Ala Gly Thr Leu Thr Pro Gly Glu Ile Asp Ala Arg Met Gly
 85 90 95

Glu Leu Glu Thr Asp Pro Asp Arg Pro Leu Pro Arg Thr Asp Asn Pro
 100 105 110

5

Glu His Ala Asp Gly Val Ala Ala Ala Leu Arg Ala Gly Ser Pro Val
 115 120 125

10

Thr Arg Lys Ile Arg Lys Gln Pro Arg Phe Thr Ile Gly Asp Arg Val
 130 135 140

15

Val Thr Arg Asn Leu Asn Pro His Gly His Thr Arg Leu Pro Arg Tyr
 145 150 155 160

20

Ala Arg Gly Lys Arg Gly Val Val Thr Leu His His Gly Ala His Val
 165 170 175

Phe Pro Asp Thr Asn Ala His Gly Leu Gly Glu His Pro Gln His Leu
 180 185 190

25

Tyr Thr Val Arg Phe Pro Ala Arg Glu Leu Trp Ser Asp Ala Ala Glu
 195 200 205

30

Pro Lys Glu Ser Ile Met Ile Asp Leu Trp Glu Ser Tyr Leu Gln Pro
 210 215 220

35

Asp Ile Gly Ser Lys Ala Ser Ser Ser Ala Lys Gly Lys Ala Thr Pro
 225 230 235 240

40

Lys Val Lys Pro Ala Met Ala Lys Ala Thr Ala Lys Val Ser Val Ser
 245 250 255

Ala Lys Ala Lys Thr Arg Gly Lys Ala Ala Pro Lys Glu Arg Pro Lys
 260 265 270

45

Leu Lys Pro Ala Arg Ala Ala Thr Ser Ala Ala Ser Gly Gly Glu Lys
 275 280 285

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Ala Lys Arg Lys Ala Lys Arg
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 <212> DNA
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<223> Metagenome - beta unit nitrile hydratase - M6dE2

<220>

<221> CDS

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<223>

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   1           5           10           15

15 gag atc gag ccc gat gag ccg gtg ttc cac gag ccc tgg gag gcg ctg
   96
   Glu Ile Glu Pro Asp Glu Pro Val Phe His Glu Pro Trp Glu Ala Leu
           20           25           30

20 gtt ttc gct ctg atg act ctc ggt atc ggg aag ctc ggc gcg tac aac
   144
   Val Phe Ala Leu Met Thr Leu Gly Ile Gly Lys Leu Gly Ala Tyr Asn
           35           40           45

25 gcc gat gag tac cgc cac tcg atc gag cgg atg gat ccg gcc cac tac
   192
   Ala Asp Glu Tyr Arg His Ser Ile Glu Arg Met Asp Pro Ala His Tyr
           50           55           60

30 ctt gcg gcg acg tac tac gag cgc atg ctc acc ggc gtc gca acg ctc
   240
   Leu Ala Ala Thr Tyr Tyr Glu Arg Met Leu Thr Gly Val Ala Thr Leu
   65           70           75           80

35 ctc gtc gag aag aac gtc gtt gcc cgc gac gag ctc gaa gcg cgc gcc
   288
   Leu Val Glu Lys Asn Val Val Ala Arg Asp Glu Leu Glu Ala Arg Ala
           85           90           95

40 ggc ggg ccc ttc ccg ctg tca cgg ccg gtg gcc gag cgg ccg acg gcg
   336
   Gly Gly Pro Phe Pro Leu Ser Arg Pro Val Ala Glu Arg Pro Thr Ala
           100           105           110

45 gac ctt cgg ccc cag cca caa cca cgc ttc gcg gtc ggg gat cgg gtc
   384
   Asp Leu Arg Pro Gln Pro Gln Pro Arg Phe Ala Val Gly Asp Arg Val
           115           120           125

50 gtc gtg cgc gac atc cac ccg gcc ggg cat act cgt gtg ccg cgc tac
   432
   Val Val Arg Asp Ile His Pro Ala Gly His Thr Arg Val Pro Arg Tyr
           130           135           140

55 gtg cgg ggc aag cgc ggg acc gtc gtg cac gtc gcg ccg aaa ttc tcg
   480
   Val Arg Gly Lys Arg Gly Thr Val Val His Val Ala Pro Lys Phe Ser
   145           150           155           160

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ttc ccc gac acg gcc gcg cac ggg ctg acc cat cgg agc gag cac acg
 528
 Phe Pro Asp Thr Ala Ala His Gly Leu Thr His Arg Ser Glu His Thr
 165 170 175
 5
 tat cac gtg gaa ttc gtc gcg agt gac ctt tgg gcc gac gtg gcc ggg
 576
 Tyr His Val Glu Phe Val Ala Ser Asp Leu Trp Ala Asp Val Ala Gly
 180 185 190
 10
 agc aat gag agc gta ctc gtg gac ctg tgg gac ggc tat ctg gag ggc
 624
 Ser Asn Glu Ser Val Leu Val Asp Leu Trp Asp Gly Tyr Leu Glu Gly
 195 200 205
 15
 gca tga
 630
 Ala
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 Glu Ile Glu Pro Asp Glu Pro Val Phe His Glu Pro Trp Glu Ala Leu
 20 25 30
 40 Val Phe Ala Leu Met Thr Leu Gly Ile Gly Lys Leu Gly Ala Tyr Asn
 35 40 45
 45 Ala Asp Glu Tyr Arg His Ser Ile Glu Arg Met Asp Pro Ala His Tyr
 50 55 60
 Leu Ala Ala Thr Tyr Tyr Glu Arg Met Leu Thr Gly Val Ala Thr Leu
 65 70 75 80
 50
 Leu Val Glu Lys Asn Val Val Ala Arg Asp Glu Leu Glu Ala Arg Ala
 85 90 95
 55
 Gly Gly Pro Phe Pro Leu Ser Arg Pro Val Ala Glu Arg Pro Thr Ala
 100 105 110

Asp Leu Arg Pro Gln Pro Gln Pro Arg Phe Ala Val Gly Asp Arg Val
 115 120 125

5 Val Val Arg Asp Ile His Pro Ala Gly His Thr Arg Val Pro Arg Tyr
 130 135 140

10 Val Arg Gly Lys Arg Gly Thr Val Val His Val Ala Pro Lys Phe Ser
 145 150 155 160

Phe Pro Asp Thr Ala Ala His Gly Leu Thr His Arg Ser Glu His Thr
 165 170 175

15 Tyr His Val Glu Phe Val Ala Ser Asp Leu Trp Ala Asp Val Ala Gly
 180 185 190

20 Ser Asn Glu Ser Val Leu Val Asp Leu Trp Asp Gly Tyr Leu Glu Gly
 195 200 205

25 Ala

30 <210> 75
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 <212> DNA
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 1 5 10 15

acc gct ccc cac gaa ccg acc ttc tgg gaa aag cag gtg gac gcg atc
 96

50 Thr Ala Pro His Glu Pro Thr Phe Trp Glu Lys Gln Val Asp Ala Ile
 20 25 30

cac ggc ctg ctc ggc gat tcc aag cgc cgc atc acg ctg cgc gac gag
 144

55 His Gly Leu Leu Gly Asp Ser Lys Arg Arg Ile Thr Leu Arg Asp Glu
 35 40 45

aac cgc ctc tat atc gaa tcg ctc ggc gac gac gtc tac aac acg ctc
 192

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Asn Arg Leu Tyr Ile Glu Ser Leu Gly Asp Asp Val Tyr Asn Thr Leu
 50          55          60

5  ggc tat tac gag cgc tgg acc gcc gcc atg tgc cgc cag ctc atc gac
   240
   Gly Tyr Tyr Glu Arg Trp Thr Ala Ala Met Cys Arg Gln Leu Ile Asp
   65          70          75          80

10 aag ggc gtg ctg acg cag gac gag atc gac gcc aag atc gcc gag ctg
   288
   Lys Gly Val Leu Thr Gln Asp Glu Ile Asp Ala Lys Ile Ala Glu Leu
           85          90          95

15 cgc gcc cgc ggc gtc ggc gcg gga cga cga cga aac ggc ctg caa acc
   336
   Arg Ala Arg Gly Val Gly Ala Gly Arg Arg Arg Asn Gly Leu Gln Thr
           100          105          110

20 gtg agc gcc gat ctg gcc gcc gat ctg gcc atc gcg ccg cgc ttc gcc
   384
   Val Ser Ala Asp Leu Ala Ala Asp Leu Ala Ile Ala Pro Arg Phe Ala
           115          120          125

25 gcc ggc gac cgc gtg cgg gtg cgc gac gat tat ccg ccc ggg cac atc
   432
   Ala Gly Asp Arg Val Arg Val Arg Asp Asp Tyr Pro Pro Gly His Ile
   130          135          140

30 cgc acg ccg gtc tat gtg cgc ggc aag acg ggc gtg gtg acg cgc tgc
   480
   Arg Thr Pro Val Tyr Val Arg Gly Lys Thr Gly Val Val Thr Arg Cys
   145          150          155          160

35 ttc ggc gcg ttc aag aac ccg gaa ttg ctc gcc atc ggc aag gac ggc
   528
   Phe Gly Ala Phe Lys Asn Pro Glu Leu Leu Ala Ile Gly Lys Asp Gly
           165          170          175

40 ctg ccc aag aaa att ctc tac gaa gtg cgc ttc aag cag acc gat ctc
   576
   Leu Pro Lys Lys Ile Leu Tyr Glu Val Arg Phe Lys Gln Thr Asp Leu
           180          185          190

45 tgg ccc gac tat gcc ggg ccg gcg acc gat acg ctg ctg atc gac atc
   624
   Trp Pro Asp Tyr Ala Gly Pro Ala Thr Asp Thr Leu Leu Ile Asp Ile
           195          200          205

50 tac gaa cat tgg ctg agc gac gcg tga
   651
   Tyr Glu His Trp Leu Ser Asp Ala
   210          215

55 <210> 76
   <211> 216
   <212> PRT
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15 His Gly Leu Leu Gly Asp Ser Lys Arg Arg Ile Thr Leu Arg Asp Glu
35 40 45

20 Asn Arg Leu Tyr Ile Glu Ser Leu Gly Asp Asp Val Tyr Asn Thr Leu
50 55 60

Gly Tyr Tyr Glu Arg Trp Thr Ala Ala Met Cys Arg Gln Leu Ile Asp
65 70 75 80

25 Lys Gly Val Leu Thr Gln Asp Glu Ile Asp Ala Lys Ile Ala Glu Leu
85 90 95

30 Arg Ala Arg Gly Val Gly Ala Gly Arg Arg Arg Asn Gly Leu Gln Thr
100 105 110

35 Val Ser Ala Asp Leu Ala Ala Asp Leu Ala Ile Ala Pro Arg Phe Ala
115 120 125

40 Ala Gly Asp Arg Val Arg Val Arg Asp Asp Tyr Pro Pro Gly His Ile
130 135 140

Arg Thr Pro Val Tyr Val Arg Gly Lys Thr Gly Val Val Thr Arg Cys
145 150 155 160

45 Phe Gly Ala Phe Lys Asn Pro Glu Leu Leu Ala Ile Gly Lys Asp Gly
165 170 175

50 Leu Pro Lys Lys Ile Leu Tyr Glu Val Arg Phe Lys Gln Thr Asp Leu
180 185 190

55 Trp Pro Asp Tyr Ala Gly Pro Ala Thr Asp Thr Leu Leu Ile Asp Ile
195 200 205

Tyr Glu His Trp Leu Ser Asp Ala
210 215

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   cag atc gag aag gac gag tcg ccc ttc cat gcg cgc tgg gaa ggc cgg
   96
   Gln Ile Glu Lys Asp Glu Ser Pro Phe His Ala Arg Trp Glu Gly Arg
25      20      25      30

   gcg caa gcg atg tac aac gcc att gcg gcc acg ggc aga ctg gtg ctt
   144
   Ala Gln Ala Met Tyr Asn Ala Ile Ala Ala Thr Gly Arg Leu Val Leu
30      35      40      45

   ggc ggt aga ccc aca cgg gaa ggg ttc ccg ccg gcc gaa tat ctc cgc
   192
   Gly Gly Arg Pro Thr Arg Glu Gly Phe Pro Pro Ala Glu Tyr Leu Arg
35      50      55      60

   atg agc tac tat gaa ttg ggt ttc agg gtg ctg gtc gag gac ttg gtc
   240
   Met Ser Tyr Tyr Glu Leu Gly Phe Arg Val Leu Val Glu Asp Leu Val
40  65      70      75      80

   ctg aac ggt ttg gtg acg cgc gcg gaa atc acg agc ggc cgt ccg gca
   288
   Leu Asn Gly Leu Val Thr Arg Ala Glu Ile Thr Ser Gly Arg Pro Ala
45      85      90      95

   aag ggg gct gca aag tcg acg ccc gca atc acc gcc gcc acc gcg cag
   336
   Lys Gly Ala Ala Lys Ser Thr Pro Ala Ile Thr Ala Ala Thr Ala Gln
50      100      105      110

   gca tat atg ttc gcg ctc aaa tcg acc cgg cga gac gta ccg gtc acg
   384
   Ala Tyr Met Phe Ala Leu Lys Ser Thr Arg Arg Asp Val Pro Val Thr
55      115      120      125

   gcg cgt ttc caa gtc ggt cag cgt gtg cgc gcg cgc aac atc aat ccg
   432
   Ala Arg Phe Gln Val Gly Gln Arg Val Arg Ala Arg Asn Ile Asn Pro

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	130	135	140
	gtc acc cat acg cgc ctg ccc cgt tac gcg cgc ggc aaa ttc ggc gtt 480		
5	Val Thr His Thr Arg Leu Pro Arg Tyr Ala Arg Gly Lys Phe Gly Val 145	150	155 160
	atc gaa cgt gac cac ggt gtt tac agg ttc gac gat tcc ttt gcc acg 528		
10	Ile Glu Arg Asp His Gly Val Tyr Arg Phe Asp Asp Ser Phe Ala Thr 165	170	175
	tcc ggc gac gag aag ccc cag cac gtt tat tct gtg cgc ttc gcg gcg 576		
15	Ser Gly Asp Glu Lys Pro Gln His Val Tyr Ser Val Arg Phe Ala Ala 180	185	190
	cgc gaa cta tgg ggc gaa gcc gcg ccg ccg cga gat gct gtc tat atc 624		
20	Arg Glu Leu Trp Gly Glu Ala Ala Pro Pro Arg Asp Ala Val Tyr Ile 195	200	205
	gaa atc tgg gat gac aac ctt gag cca gcg tga 657		
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45	Gln Ile Glu Lys Asp Glu Ser Pro Phe His Ala Arg Trp Glu Gly Arg 20	25	30
	Ala Gln Ala Met Tyr Asn Ala Ile Ala Ala Thr Gly Arg Leu Val Leu 35	40	45
50	Gly Gly Arg Pro Thr Arg Glu Gly Phe Pro Pro Ala Glu Tyr Leu Arg 50	55	60
55	Met Ser Tyr Tyr Glu Leu Gly Phe Arg Val Leu Val Glu Asp Leu Val 65	70	75 80
	Leu Asn Gly Leu Val Thr Arg Ala Glu Ile Thr Ser Gly Arg Pro Ala		

	85	90	95
5	Lys Gly Ala Ala Lys Ser Thr Pro Ala Ile Thr Ala Ala Thr Ala Gln 100 105 110		
10	Ala Tyr Met Phe Ala Leu Lys Ser Thr Arg Arg Asp Val Pro Val Thr 115 120 125		
15	Ala Arg Phe Gln Val Gly Gln Arg Val Arg Ala Arg Asn Ile Asn Pro 130 135 140		
	Val Thr His Thr Arg Leu Pro Arg Tyr Ala Arg Gly Lys Phe Gly Val 145 150 155 160		
20	Ile Glu Arg Asp His Gly Val Tyr Arg Phe Asp Asp Ser Phe Ala Thr 165 170 175		
25	Ser Gly Asp Glu Lys Pro Gln His Val Tyr Ser Val Arg Phe Ala Ala 180 185 190		
30	Arg Glu Leu Trp Gly Glu Ala Ala Pro Pro Arg Asp Ala Val Tyr Ile 195 200 205		
	Glu Ile Trp Asp Asp Asn Leu Glu Pro Ala 210 215		
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45	<220> <221> CDS <222> (1)..(696) <223>		
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55	ggc ggc gca gct atc ccc gat ctg gct cat gcg gca gac cac gac cac 96 Gly Gly Ala Ala Ile Pro Asp Leu Ala His Ala Ala Asp His Asp His 20 25 30		

5 cag cat caa gat ttg ccg tcc gat ctg gcg ctg cgg gtg aag tcg ctc
 144
 Gln His Gln Asp Leu Pro Ser Asp Leu Ala Leu Arg Val Lys Ser Leu
 35 40 45
 10 gaa tcg ctg ctt gtc gag aag ggg ctg gtg gag cga gca gcg ctc gac
 192
 Glu Ser Leu Leu Val Glu Lys Gly Leu Val Glu Arg Ala Ala Leu Asp
 50 55 60
 15 gcg ctg gtg gac acc tac gag cac aaa gtc ggg ccg cga aac gga gcg
 240
 Ala Leu Val Asp Thr Tyr Glu His Lys Val Gly Pro Arg Asn Gly Ala
 65 70 75 80
 20 cgc gtt gtc gcg cgg gcc tgg gtt gac ccg gac tac aag caa cgg tta
 288
 Arg Val Val Ala Arg Ala Trp Val Asp Pro Asp Tyr Lys Gln Arg Leu
 85 90 95
 25 ttc gcg aac ggt acc gcc gca gtc gcg gag ttc ggc tac tcc ggc tcg
 336
 Phe Ala Asn Gly Thr Ala Ala Val Ala Glu Phe Gly Tyr Ser Gly Ser
 100 105 110
 30 cag ggc gct gac atc cgg gtc gtc gaa aac acg gcc act gta cat aac
 384
 Gln Gly Ala Asp Ile Arg Val Val Glu Asn Thr Ala Thr Val His Asn
 115 120 125
 35 ctc gtc gtg tgc acg ctg tgc tct tgt tat ccc tgg ccg gtg ctg ggc
 432
 Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly
 130 135 140
 40 ttg ccg ccg gtc tgg tac aag tcc gcg ccc tat cgg tct cgc gtg gtg
 480
 Leu Pro Pro Val Trp Tyr Lys Ser Ala Pro Tyr Arg Ser Arg Val Val
 145 150 155 160
 45 atc gat ccg cga ggt gtg ctg cgc gag ttc ggc gtg gtg ctg ccg gac
 528
 Ile Asp Pro Arg Gly Val Leu Arg Glu Phe Gly Val Val Leu Pro Asp
 165 170 175
 50 cat atc gaa gtg cgt gtc tat gac agc acg gcg gag caa cgc tat cta
 576
 His Ile Glu Val Arg Val Tyr Asp Ser Thr Ala Glu Gln Arg Tyr Leu
 180 185 190
 55 gtg ctg ccg gag cgg ccg gcc gga acc gaa aac ctg aca gaa gaa gcg
 624
 Val Leu Pro Glu Arg Pro Ala Gly Thr Glu Asn Leu Thr Glu Glu Ala
 195 200 205
 60 ctg gcg ctg ctg gtg acg cgc gac gcg atg att ggc gtg gcc aag gtc
 672
 Leu Ala Leu Leu Val Thr Arg Asp Ala Met Ile Gly Val Ala Lys Val
 210 215 220

gcg ccg ccg gga ggc cgc gga tga
 696
 Ala Pro Pro Gly Gly Arg Gly
 5 225 230

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 10 <212> PRT
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Gly Gly Ala Ala Ile Pro Asp Leu Ala His Ala Ala Asp His Asp His
 20 25 30

25 Gln His Gln Asp Leu Pro Ser Asp Leu Ala Leu Arg Val Lys Ser Leu
 35 40 45

30 Glu Ser Leu Leu Val Glu Lys Gly Leu Val Glu Arg Ala Ala Leu Asp
 50 55 60

35 Ala Leu Val Asp Thr Tyr Glu His Lys Val Gly Pro Arg Asn Gly Ala
 65 70 75 80

Arg Val Val Ala Arg Ala Trp Val Asp Pro Asp Tyr Lys Gln Arg Leu
 85 90 95
 40

Phe Ala Asn Gly Thr Ala Ala Val Ala Glu Phe Gly Tyr Ser Gly Ser
 100 105 110

45 Gln Gly Ala Asp Ile Arg Val Val Glu Asn Thr Ala Thr Val His Asn
 115 120 125

50 Leu Val Val Cys Thr Leu Cys Ser Cys Tyr Pro Trp Pro Val Leu Gly
 130 135 140

55 Leu Pro Pro Val Trp Tyr Lys Ser Ala Pro Tyr Arg Ser Arg Val Val
 145 150 155 160

Ile Asp Pro Arg Gly Val Leu Arg Glu Phe Gly Val Val Leu Pro Asp
 165 170 175

5

His Ile Glu Val Arg Val Tyr Asp Ser Thr Ala Glu Gln Arg Tyr Leu
180 185 190

Val Leu Pro Glu Arg Pro Ala Gly Thr Glu Asn Leu Thr Glu Glu Ala
195 200 205

10

Leu Ala Leu Leu Val Thr Arg Asp Ala Met Ile Gly Val Ala Lys Val
210 215 220

15

Ala Pro Pro Gly Gly Arg Gly
225 230

20

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<212> DNA
<213> Unknown

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<220>
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<220>
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30 <223>

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48

35 Met Lys Asp Ser Pro Val Phe Arg Glu Pro Trp Glu Ala Gln Ala Phe
1 5 10 15

gcg ttg gcg atc tcg ttg caa gac cgt ggc gtg ttc acg cga gac gaa
96

40 Ala Leu Ala Ile Ser Leu Gln Asp Arg Gly Val Phe Thr Arg Asp Glu
20 25 30

tgg gcg gcg gca ctc ggc gat gaa atc aag aag gcg caa gct gcc ggc
144

45 Trp Ala Ala Ala Leu Gly Asp Glu Ile Lys Lys Ala Gln Ala Ala Gly
35 40 45

gat ccc gat acg ggc gag act tat tac cat cat tgg atg gca gcg ctc
192

50 Asp Pro Asp Thr Gly Glu Thr Tyr Tyr His His Trp Met Ala Ala Leu
50 55 60

gaa cgg ctg att gca gcc aag ggt gtt gcc gat acg cag acg ctc gcg
240

55 Glu Arg Leu Ile Ala Ala Lys Gly Val Ala Asp Thr Gln Thr Leu Ala
65 70 75 80

cgc aca cgc gac gcc tgg cag cac gcc tgt gcg cga acg ccg cat gcc
288

Arg Thr Arg Asp Ala Trp Gln His Ala Cys Ala Arg Thr Pro His Gly
 85 90 95

5 gcg cca atc gag cta aga ccg gac gac ttc agg aat tga
 327
 Ala Pro Ile Glu Leu Arg Pro Asp Asp Phe Arg Asn
 100 105

10 <210> 82
 <211> 108
 <212> PRT
 <213> Unknown

15 <220>
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 1 5 10 15

25 Ala Leu Ala Ile Ser Leu Gln Asp Arg Gly Val Phe Thr Arg Asp Glu
 20 25 30

30 Trp Ala Ala Ala Leu Gly Asp Glu Ile Lys Lys Ala Gln Ala Ala Gly
 35 40 45

35 Asp Pro Asp Thr Gly Glu Thr Tyr Tyr His His Trp Met Ala Ala Leu
 50 55 60

40 Glu Arg Leu Ile Ala Ala Lys Gly Val Ala Asp Thr Gln Thr Leu Ala
 65 70 75 80

45 Arg Thr Arg Asp Ala Trp Gln His Ala Cys Ala Arg Thr Pro His Gly
 85 90 95

50 Ala Pro Ile Glu Leu Arg Pro Asp Asp Phe Arg Asn
 100 105

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 48
 5 Met Arg Thr Val Ala Glu Gln Ile Ala Ala Asp Leu Ala Ser Pro Ala
 1 5 10 15

 gcg att ccg cgc cgc aac ggc gag ccg gtc ttc gac gag cct tgg gaa
 96
 10 Ala Ile Pro Arg Arg Asn Gly Glu Pro Val Phe Asp Glu Pro Trp Glu
 20 25 30

 agt cgt gcg ttt ggg ata gcg gtc gcc ctt tcc gag ggt ggc ctc tat
 144
 15 Ser Arg Ala Phe Gly Ile Ala Val Ala Leu Ser Glu Gly Gly Leu Tyr
 35 40 45

 tca tgg gat gaa ttt cgc gat tgc ctg att gct gaa atc aca gcg gcg
 192
 20 Ser Trp Asp Glu Phe Arg Asp Cys Leu Ile Ala Glu Ile Thr Ala Ala
 50 55 60

 gat gcg cgc ggc gag cat acg agc tat tac gaa cgg ttt ctc gcc gcc
 240
 25 Asp Ala Arg Gly Glu His Thr Ser Tyr Tyr Glu Arg Phe Leu Ala Ala
 65 70 75 80

 ctg cag cat ctg ctc gcg gcc aaa cgc ctc tgc act ccc gat gaa gtc
 288
 30 Leu Gln His Leu Leu Ala Ala Lys Arg Leu Cys Thr Pro Asp Glu Val
 85 90 95

 gag cgg cgg atg aac act agc gca ggc acc tga
 321
 35 Glu Arg Arg Met Asn Thr Ser Ala Gly Thr
 100 105

 <210> 84
 40 <211> 106
 <212> PRT
 <213> Unknown

 <220>
 45 <223> Metagenome - p12K unit

 <400> 84

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 Ala Ile Pro Arg Arg Asn Gly Glu Pro Val Phe Asp Glu Pro Trp Glu
 20 25 30
 55

 Ser Arg Ala Phe Gly Ile Ala Val Ala Leu Ser Glu Gly Gly Leu Tyr
 35 40 45

Ser Trp Asp Glu Phe Arg Asp Cys Leu Ile Ala Glu Ile Thr Ala Ala
 50 55 60

5
 Asp Ala Arg Gly Glu His Thr Ser Tyr Tyr Glu Arg Phe Leu Ala Ala
 65 70 75 80

10 Leu Gln His Leu Leu Ala Ala Lys Arg Leu Cys Thr Pro Asp Glu Val
 85 90 95

15 Glu Arg Arg Met Asn Thr Ser Ala Gly Thr
 100 105

20 <210> 85
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 <213> Unknown

25 <220>
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30 <220>
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35 <400> 85
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 48
 Met Thr Thr Leu Ser Gln Arg Glu Ala Ala Pro Ser Ala Glu Leu Leu
 1 5 10 15

40 gac cta ccg caa ctt cca agc gac acc gac ggc ccc gtc ttc gcg gaa
 96
 Asp Leu Pro Gln Leu Pro Ser Asp Thr Asp Gly Pro Val Phe Ala Glu
 20 25 30

45 cct tgg gaa gcg gaa gcg ttt gcg ctt gcc gta agt ctt tca gag caa
 144
 Pro Trp Glu Ala Glu Ala Phe Ala Leu Ala Val Ser Leu Ser Glu Gln
 35 40 45

50 gga cat ttc acg tgg aag gaa tgg gca gca acg ctc gcc gat gaa ctg
 192
 Gly His Phe Thr Trp Lys Glu Trp Ala Ala Thr Leu Ala Asp Glu Leu
 50 55 60

55 gag ggc gcc gcc aat cgc ggc gag ccg gat gac ggt acg cat tat tat
 240
 Glu Gly Ala Ala Asn Arg Gly Glu Pro Asp Asp Gly Thr His Tyr Tyr
 65 70 75 80

gag tac tgg ctg acg gcc ctg gaa agg ctg gtt acg atc aag ggc ctg
 288
 Glu Tyr Trp Leu Thr Ala Leu Glu Arg Leu Val Thr Ile Lys Gly Leu

	85	90	95
	aca gat cag caa gcg atg cgc gag cgc aaa gag gcg tgg gaa gaa gcc		
	336		
5	Thr Asp Gln Gln Ala Met Arg Glu Arg Lys Glu Ala Trp Glu Glu Ala		
	100	105	110
	tat cgc cat acc ccg cat ggc gcg cca gtt gaa ctt atg tct ccg ctc		
	384		
10	Tyr Arg His Thr Pro His Gly Ala Pro Val Glu Leu Met Ser Pro Leu		
	115	120	125
	gat caa agc cgg ata gcc gaa gag gac agc gaa tcc tca tag		
	426		
15	Asp Gln Ser Arg Ile Ala Glu Glu Asp Ser Glu Ser Ser		
	130	135	140
	<210> 86		
20	<211> 141		
	<212> PRT		
	<213> Unknown		
	<220>		
25	<223> Metagenome - p12K unit		
	<400> 86		
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	Asp Leu Pro Gln Leu Pro Ser Asp Thr Asp Gly Pro Val Phe Ala Glu		
	20	25	30
35	Pro Trp Glu Ala Glu Ala Phe Ala Leu Ala Val Ser Leu Ser Glu Gln		
	35	40	45
40	Gly His Phe Thr Trp Lys Glu Trp Ala Ala Thr Leu Ala Asp Glu Leu		
	50	55	60
	Glu Gly Ala Ala Asn Arg Gly Glu Pro Asp Asp Gly Thr His Tyr Tyr		
45	65	70	75 80
	Glu Tyr Trp Leu Thr Ala Leu Glu Arg Leu Val Thr Ile Lys Gly Leu		
50	85	90	95
	Thr Asp Gln Gln Ala Met Arg Glu Arg Lys Glu Ala Trp Glu Glu Ala		
	100	105	110
55	Tyr Arg His Thr Pro His Gly Ala Pro Val Glu Leu Met Ser Pro Leu		
	115	120	125

Asp	Gln	Ser	Arg	Ile	Ala	Glu	Glu	Asp	Ser	Glu	Ser	Ser
130						135					140	