RNA-GUIDED GENE DRIVES

RELATED APPLICATION DATA

This application claims priority to U.S. Provisional Application No. 61/924,735, filed on January 8, 2014 and U.S. Provisional Application No. 62/024,642, filed on July 15, 2014 each of which is hereby incorporated herein by reference in their entirety for all purposes.

BACKGROUND

Gene drives are generally known as genetic elements that skew the natural odds in their favor of being inherited and passed on by progeny. Examples include homing endonuclease genes that copy themselves into chromosomes lacking them, segregation distorters that destroy competing chromosomes during meiosis, transposons that insert copies of themselves elsewhere in the genome, Medea elements that eliminate competing siblings who do not inherit them, and maternally heritable microorganisms such as Wolbachia that induce cytoplasmic incompatibility to favor the spread of infected individuals. Because they circumvent the normal rules of natural selection, all of these elements have been considered as potential "gene drive" systems capable of spreading engineered modifications through insect vector populations to block the spread of disease. Homing endonuclease based gene drives have been proposed as a means of genetically controlling malaria mosquito populations. See Windbichler et al., Nature, doi:10.1038/nature09937 (2011). Site-specific selfish genes have been proposed as tools for the control and genetic engineering of natural populations. See Burt, Proc. R. Soc. Lond. B (2003) 270, 921-928 (2003). However, such proposed gene drives are limited in their site specificity or difficult to express in various organisms. A need therefore exists to develop gene drives which can target any desired gene and can be utilized across a broad spectrum of organisms.

SUMMARY

Aspects of the present disclosure are directed to RNA guided gene drives, and in particular, a foreign nucleic acid sequence which is stably introduced into a germline cell of a desired organism. The term foreign nucleic acid sequence as described herein and the term RNA guided gene drive may be used interchangeable herein. A resulting transgenic organism may be developed from the germline cell. The resulting transgenic organism can then be introduced into a wild type population, and through mating by the transgenic organism with a wild type organism, the foreign nucleic acid sequence is transferred to resulting offspring or progeny. As a result, methods of the present disclosure are directed to the production of a population of transgenic

organisms having desired traits from an initial transgenic organism and a wild type organism. When the transgenic organism is introduced into a wild type population and mates with a wild type organism, the resulting progeny may be referred to as an altered wild type population. As a result of the foreign nucleic acid sequence being stably entered into the genome of the transgenic offspring or transgenic progeny, transgenic offspring or transgenic progeny may have one or more desired traits resulting from expression of the foreign nucleic acid. According to certain aspects, methods described herein may be used to create an altered wild type population of transgenic organisms where the transgenic organisms exhibit one or more desired traits resulting from the expression of the foreign nucleic acid.

According to certain aspects, the foreign nucleic acid sequence encodes at least an RNA guided DNA binding protein, such as one or more of an RNA guided DNA binding protein nuclease, an RNA guided DNA binding protein nickase or a nuclease null RNA guided DNA binding protein, and one or more or a plurality of guide RNAs (ribonucleic acids). A guide RNA is complementary to DNA (deoxyribonucleic acid), such as a target DNA in the genome of a germline cell. The foreign nucleic acid sequence also encodes at least one or more promoters such that the germline cell may express the RNA guided DNA binding protein and the guide RNAs or any other nucleic acid sequence or gene which may be in the foreign nucleic acid sequence. One of skill will readily be able to identify suitable promoters based on the present disclosure and the particular germline cell. The foreign nucleic acid sequence may also include any other nucleic acid sequence or sequences known to those of skill in the art to be required for expression of the foreign nucleic acid sequence by a germline cell. The foreign nucleic acid sequence may also include any other gene sequence or gene sequences desired to be expressed by the germline cell. Such a gene sequence or such gene sequences may be referred to as "cargo DNA." It is to be understood that one of skill will readily be able to identify one or more gene sequences depending upon the desired trait one of skill wishes to be exhibited by the germline cell or the organism developed from the germline cell when the cell expresses the foreign nucleic acid sequence. The foreign nucleic acid sequence also encodes at least two flanking sequences which flank at least the RNA guided DNA binding protein nuclease and the one or more guide RNAs. As known to those of skill in the art, flanking sequences are placed at opposite ends of a particular nucleic acid sequence such that the particular nucleic acid sequence is between the flanking sequences. According to one aspect, the flanking sequences include at least a sequence which is identical to a corresponding sequence on a selected chromosome. According to one aspect, such flanking sequences allow a cell to insert the foreign nucleic acid sequence into its genomic DNA at a cut site using well-understood mechanisms such as homologous recombination or nonhomologous end joining.

According to certain aspects, when the foreign nucleic acid sequence is expressed by the germline cell, one or more of an RNA guided DNA binding protein and one or more or a plurality

of guide RNAs are produced. The RNA guided DNA binding protein and a guide RNA produces a complex of the RNA guided DNA binding protein, the guide RNA and a double stranded DNA target sequence. In this aspect, the RNA is said to guide the DNA binding protein to the double stranded DNA target sequence for binding thereto. This aspect of the present disclosure may be referred to as co-localization of the RNA and DNA binding protein to or with the double stranded DNA.

DNA binding proteins within the scope of the present disclosure may include those which create a double stranded break (which may be referred to as a DNA binding protein nuclease), those which create a single stranded break (referred to as a DNA binding protein nickase) or those which have no nuclease activity (referred to as a nuclease null DNA binding protein) but otherwise bind to target DNA. In this manner, a DNA binding protein-guide RNA complex may be used to create a double stranded break at a target DNA site, to create a single stranded break at a target DNA site or to localize a transcriptional regulator protein or domain, which may be expressed by the cell, at a target DNA site so as to regulate expression of target DNA. According to certain aspects, the foreign nucleic acid sequence may encode one or more of a DNA binding protein nuclease, a DNA binding protein nickase or a nuclease null DNA binding protein. The foreign nucleic acid sequence may also encode one or more transcriptional regulator proteins or domains or one or more donor nucleic acid sequences that are intended to be inserted into the genomic DNA. According to one aspect, the foreign nucleic acid sequence encoding an RNA guided nuclease-null DNA binding protein further encodes the transcriptional regulator protein or domain fused to the RNA guided nuclease-null DNA binding protein. According to one aspect, the foreign nucleic acid sequence encoding one or more RNAs further encodes a target of an RNA-binding domain and the foreign nucleic acid encoding the transcriptional regulator protein or domain further encodes an RNA-binding domain fused to the transcriptional regulator protein or domain.

Accordingly, expression of a foreign nucleic acid sequence by a germline cell may result in a double stranded break, a single stranded break and/or transcriptional activation or repression of the genomic DNA. Donor DNA may be inserted at the break site by cell mechanisms such as homologous recombination or nonhomologous end joining. It is to be understood that expression of a foreign nucleic acid sequence as described herein may result in a plurality of double stranded breaks or single stranded breaks at various locations along target genomic DNA, including one or more or a plurality of gene sequences, as desired.

Aspects of the present disclosure are directed to using the foreign nucleic acid sequence as a gene drive. The concept of a gene drive is known to those of skill in the art and refers to a foreign nucleic acid sequence which when expressed is capable of inserting itself into the genome of the cell into which it has been introduced. The concept of a gene drive is provided in Windbichler et al., *Nature,* doi:10.1038/nature09937 (2011) and Burt, *Proc. R. Soc. Lond.* B (2003)

270, 921-928 (2003) each of which is hereby incorporated by reference in their entireties.

According to one aspect of the present disclosure, the foreign nucleic acid sequences described herein act as gene drives when introduced into a germline cell. In one aspect, the foreign nucleic acid sequence is expressed by the germline cell to produce an RNA guided DNA binding protein and a guide RNA. The guide RNA is complementary to a target DNA sequence on a chromosome. The RNA guided DNA binding protein and the guide RNA co-localize to the target DNA, and the target DNA is cleaved in a site specific manner. The target DNA may be a target DNA site on one or both chromosomes of a chromosome pair. The foreign nucleic acid sequence is then inserted into the genomic DNA at the target DNA cut site, for example, by homologous recombination. The foreign nucleic acid sequence may be inserted into the genomic DNA at one or both chromosomes of a chromosome pair if each chromosome has been cleaved in a site specific manner by the RNA guided DNA binding protein. If inserted into both chromosomes of a chromosome pair, then the germline cell is homozygous for the foreign nucleic acid sequence. In an alternate embodiment, the foreign nucleic acid sequence is inserted into a first chromosome of a chromosome pair. The inserted foreign nucleic acid sequence is then expressed by the cell and the RNA guided DNA binding protein and the guide RNA co-localize at or to a second chromosome of a chromosome pair which is then cleaved in a site specific manner, just as was the first chromosome. The cleaved target DNA in the second chromosome is then repaired, for example by homologous recombination, using the first chromosome as a template. In this manner, the second chromosome is repaired to include the foreign nucleic acid sequence resulting in a germline cell that is homozygous for the foreign nucleic sequence, i.e., the foreign nucleic acid sequence is present in both the first and second chromosome of the chromosome pair. The mechanisms by which cells repair damaged, cleaved or cut genomic DNA are well known. Aspects of the present disclosure take advantage of these cell mechanisms in combination with DNA binding protein nucleases or nickases to create a gene drive with desired foreign genetic material that inserts into the genomic DNA of cells wherein the cell becomes homozygous for the foreign genetic material. The foreign genetic material is then passed on to progeny to create a population of transgenic organisms having one or more desired traits.

Using the concept of a foreign nucleic acid sequence as described herein as a gene drive, methods are provided for incorporating foreign genetic material into a wild population of organisms. Methods are provided for making a cell homozygous for foreign genetic material. Methods are provided for spreading a genetic modification through a wild population of organisms. Methods are provided for spreading a human designed genetic modification through a wild population of organisms. Methods are provided for pan genome engineering by spreading a human designed genetic modification through a wild population of organisms. Methods are provided for editing the genome of a wild species. Methods are provided for editing multiple loci of genomic

DNA of an organism. Methods are provided for the multiplexed editing of genomic loci. Methods are provided for reversibly editing a locus or multiple loci of genomic DNA of an organism.

Based upon the desired function of the gene drive described herein, methods are provided for controlling gene flow through a wild population of organisms. Methods are provided for suppressing expansion of a target population of an organism. Methods are provided for decreasing or eliminating a target population of an organism. Methods are providing for increasing a target population of an organism. Methods are provided for reducing or eliminating vector born diseases, such as malaria. Methods are provided for decreasing the spread of disease by a target organism, such as an insect vector population. Methods are provided for disrupting a gene responsible for disease transmission by a target organism. Methods are provided for disrupting a Y chromosome in a germline cell. Methods are provided for disrupting an X chromosome in a germline cell. Methods are provided for controlling invasive pests. Methods are provided for preserving species threatened by ecological change.

According to certain aspects, a method of blocking gene flow from an engineered organism to the wild-type population is provided which includes 1) the insertion of a recoded copy of a gene whose phenotype when disrupted is dominant negative lethal into a distal region of the same chromosome in which the wild-type copy is encoded and 2) the insertion of a selfish genetic element, i.e. RNA guided gene drive as described herein and as the term gene drive is understood by those of skill in the art, that copies itself in place of the wild-type version of the same gene, such that any offspring from an engineered organism and an engineered organism contain two functional copies of the recoded gene, while the offspring of an engineered organism and a wild-type organism will possess only a single copy of the recoded gene after the selfish genetic element replaces the wild-type copy and cannot copy the recoded gene onto the wild-type chromosome.

According to certain aspects, a method of blocking gene flow between a subpopulation bearing a unique sequence and the remaining population is provided which includes 1) releasing a first selfish genetic element that spreads exclusively using the unique sequence and inserts part of a gene whose phenotype when disrupted is dominant negative elsewhere in the genome, 2) releasing a second selfish genetic element that exclusively spreads using the partial gene sequences and inserts a recoded version of the gene and also disrupts the wild-type copy of the gene such that 1) all offspring resulting from matings between an organism bearing the first selfish genetic element and an organism bearing the second selfish genetic element contain the second selfish genetic element in which both wild-type copies of the gene are disrupted but are replaced by recoded copies elsewhere in the genome, and 2) any cross between an organism containing the second selfish genetic element and a wild-type organism generates no progeny because the wild-type copy of the gene is lost and not replaced by a recoded copy.

According to certain aspects, a method of biasing the sex ratio of offspring is provided

including using one or more chromosomes that together 1) encode an RNA-guided nuclease that is expressed exclusively during pre-meiosis and 2) express guide RNAs that target the nuclease to cleave sequences uniquely found on one of the sex chromosomes such that viable gametes contain fewer instances of the targeted chromosome than is typical of an unmodified organism.

According to certain aspects, a method of biasing the sex ratio of a population is provided including using a sex chromosome that encodes an RNA-guided nuclease that is expressed exclusively during pre-meiosis and also expresses guide RNAs that target the nuclease to cleave sequences uniquely found on the opposite sex chromosome such that viable gametes predominantly contain the sex chromosome encoding the RNA-guided nuclease.

According to certain aspects, a method of biasing the sex ratio of offspring towards the heterogametic sex (for example XY) is provided including using a chromosome engineered with a copy of an essential gene normally present on the X chromosome and a selfish genetic element that copies itself in place of the wild-type essential gene on the X chromosome such that female progeny are not developmentally viable due to loss of the essential gene while male progeny survive due to the copy on the engineered chromosome.

According to certain aspects, a method of biasing the sex ratio of a population towards the heterogametic sex (for example XY) is provided including using a Y chromosome engineered with a copy of an essential gene normally present on the X chromosome and a selfish genetic element that copies itself in place of the wild-type essential gene on the X chromosome such that female progeny are not developmentally viable due to loss of the essential gene while male progeny survive due to the copy on the engineered Y chromosome.

According to certain aspects, a method of biasing the sex ratio of offspring towards the homogametic sex (for example XX in mammals) including using one or more chromosomes that together 1) encode an RNA-guided nuclease and 2) express guide RNAs that target the nuclease to cleave sequences uniquely found on the heterogametic sex chromosome (for example XY in mammals) such that the heterogametic sex chromosome of any offspring that would normally develop as the heterogametic sex (e.g. males in mammals) is destroyed.

According to certain aspects, a method of biasing the sex ratio of a population towards the homogametic sex (for example XX in mammals) including using a homogametic sex chromosome that 1) encodes an RNA-guided nuclease and 2) expresses guide RNAs that target the nuclease to cleave sequences uniquely found on the heterogametic sex chromosome (for example XY in mammals) such that the heterogametic sex chromosome of any offspring that would normally develop as the heterogametic sex (e.g. males in mammals) is destroyed.

According to certain aspects, a method of biasing the sex ratio of a population towards males includes the release of engineered males that encode a selfish genetic element that copies itself in place of a gene required for female fertility. According to one aspect, the selfish genetic

element is encoded on the Y chromosome. According to one aspect, the Y chromosome also encodes a selfish genetic element targeting an essential gene present on the X chromosome.

According to certain aspects, a method of population control is provided including the release into a population to be removed of engineered male organisms containing a Y chromosome that encodes 1) a selfish genetic element that copies itself in place of a gene required for female fertility and 2) a second selfish genetic element that copies itself in place of an essential gene on the X chromosome such that female offspring are nonviable, and the release into a population to be protected of organisms encoding a third selfish genetic element that changes the sequence of the essential gene on the X chromosome such that the offspring of matings between males containing the first and second selfish genetic elements and females containing the third selfish genetic element give rise to fertile male but sterile female offspring.

According to certain aspects, a method is provided of altering a eukaryotic germline cell of an organism including introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell, wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence, wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA, wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA, wherein at least one copy of the sequence located between the first flanking sequence and the second flanking sequence that is cut and replaced by the foreign nucleic acid sequence is required for the organism to survive or to produce viable offspring, expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner, inserting the first foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the first foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell

homozygous for the foreign nucleic acid sequence, and performing the above expression and insertion steps at developmental stage at which the sequence located between the first flanking sequence and the second flanking sequence that is cut and replaced by the foreign nucleic acid sequence to produce a genetic load is no longer required for that organism to survive or produce fertile offspring.

According to certain aspects, a method for targeted population suppression or extinction according to such methods described herein includes releasing an RNA-guided genetic load drive into the targeted population.

A genetic load as that term is used herein may refer to the difference between the fitness of the theoretically optimal genotype in a population and the fitness of the observed average genotype in a population. The term genetic load may also refer to a reduction in the mean fitness for a population compared to the maximum fitness.

Further features and advantages of certain embodiments of the present invention will become more fully apparent in the following description of embodiments and drawings thereof, and from the claims.

BRIEF DESCRIPTION OF THE DRAWINGS

The patent or application file contains at least one drawing executed in color. Copies of this patent or patent application publication with color drawing(s) will be provided by the Office upon request and payment of the necessary fee. The foregoing and other features and advantages of the present embodiments will be more fully understood from the following detailed description of illustrative embodiments taken in conjunction with the accompanying drawings in which:

Fig. 1A is a schematic of a standard RNA-guided Cas9 gene drive showing expression of Cas9 and guide RNA, co-localization, cutting of a target gene to create a break and insertion of the RNA guided Cas9 gene drive into the target gene at the cut site by homologous recombination. Fig. 1B depicts a chromosomal view of gene drive inheritance. Fig. 1C depicts a mutation preventing the co-localization of Cas9 and a guide RNA so as to prevent cutting of the target DNA and insertion of the gene drive. Fig. 1D depicts the use of a plurality of guide RNAs and cut sites to avoid a mutation from preventing insertion of the gene drive. Fig. 1E depicts NHEJ deleting all recognition sites to create a drive resistant allele. Fig. 1F depicts NHEJ creating a drive resistant allele that deleteriously disrupts an essential gene.

Fig. 2A depicts the use of a single drive to edit multiple distal sites by including appropriate guide RNAs. Fig. 2B depicts the use of subsequent drives to update changes made by a first drive. Fig. 2C depicts the use of a restorative drive to all changes back to the wild-type, leaving only the Cas9 gene and guide RNAs.

Fig. 3 depicts the use of two gene drives that target the same essential gene using different

guide RNAs that become incompatible when crossed due to the loss of both copies. Releasing this type of drive into a population will artificially split it into as many species as there are drives.

Fig. 4A depicts the use of meiotic gene drives to eliminate the competing sex chromosome during meiosis, producing a biased pool of gametes. Drive expression is limited to pre-meiosis to prevent the drive from being lethal. Fig. 4B depicts that meiotic sex chromosome silencing can be circumvented using two orthogonal Cas9 nucleases. Cas9 A causes the cassette encoding Cas9 B to be copied from the sex chromosome to an autosome early in development, where it is free to eliminate the competing sex chromosome during meiosis. Fig. 4C depicts that zygotic X-drives can target the Y chromosome for destruction in the fertilized egg, yielding exclusively female progeny. Fig. 4D depicts that zygotic Y-drives can encode an essential gene normally found on the X chromosome and utilize a gene drive cassette to eliminate that gene from the X. The drive copies itself from the paternal to the maternal X, leaving female zygotes without any copies of the essential gene.

Fig. 5A depicts population suppression using dual-function Y-drives where the A Y-drive that also contains a sterile-daughter capability (Y-Drive-SD) acts as a zygotic drive in the wild-type population, which can lead to extinction. Fig. 5B depicts that upon encountering a standard drive that recodes the essential gene, the Y-Drive-SD will no longer drive. The sterile-daughter effect will prevent mutual drive invasion until the invasive population collapses.

Fig. 6A depicts a method for controlling low- and high-mobility invasive species where Y-drive male Asian carp released throughout invaded habitats such as the Mississippi can be used to eradicate Asian carp populations. A protective recoding drive can be available to protect Asian populations in the event of deliberate human transport of Y-drive males. Fig. 6B depicts releasing Y-Drive-SD male rats into invasive rat populations which will initiate local eradication while a standard drive protectively recodes the native rat populations of Eurasia. The Y-Drive-SD construct will eradicate invasive rats from most islands and many continental areas. Stowaway-mediated gene flow will be limited by the sterile-daughter effect of Y-Drive-SD in recoded populations. The process can be repeated with new drives once recoded stowaways successfully invade rat-free habitats.

Fig. 7 depicts drive-mediated regulating of endogenous genes where RNA-guided transcriptional regulators can modulate the activity of distant genes without editing their sequence. Regulation will be evolutionarily stable during the lifetime of the drive if the drive nuclease requires the regulator for proper expression.

Fig. 8 depicts inducing speciation from an unmodified wild-type population. Two successive gene drives targeting a subpopulation bearing a unique allele (depicted as a unique gene) can induce genetic incompatibility with the remaining wild-type population. The first drive spreads using the unique allele and recodes a gene that displays a dominant lethal phenotype when

disrupted. The second drive spreads using the recoded gene and eliminates the first drive. Any mating with a wild-type organism causes the second drive to disrupt the wild-type copy, causing lethality in all progeny.

Fig. 9 depicts a method described herein wherein Drive 1 recodes a male essential gene on the Y chromosome (if present) and a recessive essential gene on the X chromosome while inserting itself (i.e., being inserted by a cell mechanism such as homologous recombination after expression of Cas9 and guide RNA and localization of the Cas9 and guide RNA to a target nucleic acid sequence) adjacent to the unique gene. Drive 2 spreads through the modified X chromosome, eliminates Drive 1 from the autosome, and also jumps into the modified Y chromosome if present. Upon mating with a wild-type organism, Drive 2 eliminates the essential genes from the X or Y chromosomes. The only viable progeny are hybrid females that retain a single recoded copy of the recessive essential gene on the X chromosome. These can mate freely with males that have been modified with Drive 2, permitting wild-type genes to spread into the engineered population. Mating with a wild-type male yields only more hybrid females, preventing gene flow back into the wild-type population.

Fig. 10 depicts a method for making a zygotic Y-drive as described herein.

Fig. 11A depicts one method of making a sterile-daughter Y chromosome that eliminates a gene required for female fertility. Y chromosome (Y-SD) is not a true drive, but incorporates a limited gene drive mechanism that replaces a gene essential for female fertility with the drive. Fig. 11B depicts that in the daughters of Y-SD males, the drive similar replaces the maternal copy of the female fertility gene, causing sterility. If no gene specific for female fertility is known, a fertility gene required for both sexes can be target and complemented with an extra copy elsewhere on the Y chromosome.

Fig. 12A depicts a method of making a Y-drive that includes sterile-daughter capability (Y-drive-SD) (Y-drive + sterile-daughter Y chromosomes. Upon mating with a wild-type female, Y-drive-SD replaces or disrupts both an essential gene and a female fertility gene. Males survive due to complementation from the extra copy on Y-drive-SD. As depicted in Fig. 12B, in female progeny, the drives similarly replace the maternal copies, leaving daughters embryonically nonviable. Y-drive-SD consequently exhibits drive. As depicted in Fig. 12C, females modified with a protective drive have the sites flanking the essential gene recoded. Y-Drive-SD can only replace the fertility gene. As depicted in Fig. 12D, female progeny are still viable, but lack any copies of the fertility gene and are consequently infertile. Because males do not gain any fitness benefit from the absence of sisters, Y-drive-SD will not drive in a protected population. However, its only fitness cost is that incurred by the drive itself, so it will be removed from the population only slowly. The loss of female reproductive output will result in potent population suppression. Because of this, protected individuals will find it difficult to invade populations where Y-drive-SD

is common. Similarly, Y-Drive-SD will not be able to invade protected populations.

Fig. 13A is a schematic showing that biased inheritance of *ADE2* is readily visible in *S. cerevisiae*. Mutations in *ADE2* generate a red phenotype on adenine-limiting media due to the buildup of red pigments. Mating a red mutant haploid to a wild-type haploid produce cream-colored diploids, which yield 50% red and 50% cream-colored progeny upon sporulation. Biased inheritance of *ADE2* is readily visible in *S. cerevisiae*. Fig. 13B is a schematic showing that when haploids with a gene drive targeting *ADE2* mate with wild-type haploids in the presence of Cas9, cutting and subsequent replacement or disruption of *ADE2* will lead to red diploids that yield exclusively red progeny. Fig. 13C is an image of *S. cerevisiae* colonies. Diploids produced by mating wild-type and *ADE2*::sgRNA gene drive haploids yield cream-colored colonies in the absence of Cas9 or when the target site is removed by recoding but uniformly red colonies when both are present, demonstrating Cas9-dependent disruption of the wild-type *ADE2* copy. Fig. 13D are images of spores from 15 dissected tetrads that produce uniformly red colonies on adenine-limited plates, confirming disruption of the *ADE2* gene inherited from the wild-type parent. In the absence of the target site or Cas9, normal 2:2 segregation is observed.

Fig. 14A is a schematic showing that the *ADE2*-targeting gene drive was modified to carry *URA3* as a cargo gene. Fig. 14 B is an image of diploids produced by mating wild-type *URA3*$^-$ haploid yeast with haploids encoding the gene drive carrying *URA3* which were sporulated and tetrads dissected to isolate colonies arising from individual spores. All of these grew when replica-plated onto plates lacking uracil, demonstrating that the drive successfully copied *URA3* in all diploids. Fig. 14C is a schematic showing that the *ABD1*-targeting gene drive cuts and recodes the tail end of the essential *ABD1* gene. Gene drives and cargo genes remain intact upon copying and can spread by targeting both non-essential and essential genes.

Fig. 15A is a schematic of a phylogenomic tree indicating the relationships between wild-type strains selected for gene drive testing. Adapted with permission from Macmillan Publishers Ltd: *Nature* 458:337-341, copyright 2009. Fig. 15 B depicts quantitative PCR results of the extent of inheritance-biasing in diverse yeast strains depicting the relative abundance of wild-type and drive-containing alleles in diploids arising from matings between SK1 haploids bearing gene drives and diverse wild-type haploid strains. "No Cas9" and "No Target" refer to the haploid cells containing the *ADE2* drive element mated to wild-type haploids in the absence of Cas9 or to a wild-type strain with a mutation in the targeted sequence that prevents cutting. "2nd gen" refers to the haploid progeny of an earlier mating.

DETAILED DESCRIPTION

Embodiments of the present disclosure are based on the use of RNA guided DNA binding proteins to co-localize with guide RNA at a target DNA site and act as gene drives. Such DNA

binding proteins are readily known to those of skill in the art to bind to DNA for various purposes. Such DNA binding proteins may be naturally occurring. DNA binding proteins included within the scope of the present disclosure include those which may be guided by RNA, referred to herein as guide RNA. According to one aspect, the guide RNA is between about 10 to about 500 nucleotides. According to one aspect, the RNA is between about 20 to about 100 nucleotides. According to this aspect, the guide RNA and the RNA guided DNA binding protein form a co-localization complex at the DNA.

DNA binding proteins having nuclease activity are known to those of skill in the art, and include naturally occurring DNA binding proteins having nuclease activity, such as Cas9 proteins present, for example, in Type II CRISPR systems. Such Cas9 proteins and Type II CRISPR systems are well documented in the art. See Makarova et al., *Nature Reviews, Microbiology*, Vol. 9, June 2011, pp. 467-477 including all supplementary information hereby incorporated by reference in its entirety.

Bacterial and archaeal CRISPR-Cas systems rely on short guide RNAs in complex with Cas proteins to direct degradation of complementary sequences present within invading foreign nucleic acid. See Deltcheva, E. et al. CRISPR RNA maturation by trans-encoded small RNA and host factor RNase III. *Nature* 471, 602-607 (2011); Gasiunas, G., Barrangou, R., Horvath, P. & Siksnys, V. Cas9-crRNA ribonucleoprotein complex mediates specific DNA cleavage for adaptive immunity in bacteria. *Proceedings of the National Academy of Sciences of the United States of America* 109, E2579-2586 (2012); Jinek, M. et al. A programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity. *Science* 337, 816-821 (2012); Sapranauskas, R. et al. The Streptococcus thermophilus CRISPR/Cas system provides immunity in Escherichia coli. *Nucleic acids research* 39, 9275-9282 (2011); and Bhaya, D., Davison, M. & Barrangou, R. CRISPR-Cas systems in bacteria and archaea: versatile small RNAs for adaptive defense and regulation. *Annual review of genetics* 45, 273-297 (2011). A recent *in vitro* reconstitution of the *S. pyogenes* type II CRISPR system demonstrated that crRNA ("CRISPR RNA") fused to a normally trans-encoded tracrRNA ("trans-activating CRISPR RNA") is sufficient to direct Cas9 protein to sequence-specifically cleave target DNA sequences matching the crRNA. Expressing a gRNA homologous to a target site results in Cas9 recruitment and degradation of the target DNA. See H. Deveau *et al.*, Phage response to CRISPR-encoded resistance in Streptococcus thermophilus. *Journal of Bacteriology* 190, 1390 (Feb, 2008).

Three classes of CRISPR systems are generally known and are referred to as Type I, Type II or Type III). According to one aspect, a particular useful enzyme according to the present disclosure to cleave dsDNA is the single effector enzyme, Cas9, common to Type II. See K. S. Makarova *et al.*, Evolution and classification of the CRISPR-Cas systems. *Nature reviews. Microbiology* 9, 467 (Jun, 2011) hereby incorporated by refernece in its entirety. Within bacteria,

the Type II effector system consists of a long pre-crRNA transcribed from the spacer-containing CRISPR locus, the multifunctional Cas9 protein, and a tracrRNA important for gRNA processing. The tracrRNAs hybridize to the repeat regions separating the spacers of the pre-crRNA, initiating dsRNA cleavage by endogenous RNase III, which is followed by a second cleavage event within each spacer by Cas9, producing mature crRNAs that remain associated with the tracrRNA and Cas9.

According to one aspect, the enzyme of the present disclosure, such as Cas9 unwinds the DNA duplex and searches for sequences matching the crRNA to cleave. Target recognition occurs upon detection of complementarity between a "protospacer" sequence in the target DNA and the remaining spacer sequence in the crRNA. Importantly, Cas9 cuts the DNA only if a correct protospacer-adjacent motif (PAM) is also present at the 3' end. According to certain aspects, different protospacer-adjacent motif can be utilized. For example, the *S. pyogenes* system requires an NGG sequence, where N can be any nucleotide. *S. thermophilus* Type II systems require NGGNG (see P. Horvath, R. Barrangou, CRISPR/Cas, the immune system of bacteria and archaea. *Science* 327, 167 (Jan 8, 2010) hereby incorporated by reference in its entirety and NNAGAAW (see H. Deveau *et al.*, Phage response to CRISPR-encoded resistance in Streptococcus thermophilus. *Journal of bacteriology* 190, 1390 (Feb, 2008) hereby incorporatd by reference in its entirety), respectively, while different *S. mutans* systems tolerate NGG or NAAR (see J. R. van der Ploeg, Analysis of CRISPR in Streptococcus mutans suggests frequent occurrence of acquired immunity against infection by M102-like bacteriophages. *Microbiology* 155, 1966 (Jun, 2009) hereby incorporated by refernece in its entirety. Bioinformatic analyses have generated extensive databases of CRISPR loci in a variety of bacteria that may serve to identify additional useful PAMs and expand the set of CRISPR-targetable sequences (see M. Rho, Y. W. Wu, H. Tang, T. G. Doak, Y. Ye, Diverse CRISPRs evolving in human microbiomes. *PLoS genetics* 8, e1002441 (2012) and D. T. Pride *et al.*, Analysis of streptococcal CRISPRs from human saliva reveals substantial sequence diversity within and between subjects over time. *Genome research* 21, 126 (Jan, 2011) each of which are hereby incorporated by reference in their entireties.

Exemplary DNA binding proteins having nuclease activity function to nick or cut double stranded DNA. Such nuclease activity may result from the DNA binding protein having one or more polypeptide sequences exhibiting nuclease activity. Such exemplary DNA binding proteins may have two separate nuclease domains with each domain responsible for cutting or nicking a particular strand of the double stranded DNA. Exemplary polypeptide sequences having nuclease activity known to those of skill in the art include the McrA-HNH nuclease related domain and the RuvC-like nuclease domain. Accordingly, exemplary DNA binding proteins are those that in nature contain one or more of the McrA-HNH nuclease related domain and the RuvC-like nuclease domain.

In *S. pyogenes*, Cas9 generates a blunt-ended double-stranded break 3bp upstream of the protospacer-adjacent motif (PAM) via a process mediated by two catalytic domains in the protein: an HNH domain that cleaves the complementary strand of the DNA and a RuvC-like domain that cleaves the non-complementary strand. See Jinke et al., *Science* 337, 816-821 (2012) hereby incorporated by reference in its entirety. Cas9 proteins are known to exist in many Type II CRISPR systems including the following as identified in the supplementary information to Makarova et al., Nature Reviews, Microbiology, Vol. 9, June 2011, pp. 467-477: Methanococcus maripaludis C7; Corynebacterium diphtheriae; Corynebacterium efficiens YS-314; Corynebacterium glutamicum ATCC 13032 Kitasato; Corynebacterium glutamicum ATCC 13032 Bielefeld; Corynebacterium glutamicum R; Corynebacterium kroppenstedtii DSM 44385; Mycobacterium abscessus ATCC 19977; Nocardia farcinica IFM10152; Rhodococcus erythropolis PR4; Rhodococcus jostii RHA1; Rhodococcus opacus B4 uid36573; Acidothermus cellulolyticus 11B; Arthrobacter chlorophenolicus A6; Kribbella flavida DSM 17836 uid43465; Thermomonospora curvata DSM 43183; Bifidobacterium dentium Bd1; Bifidobacterium longum DJO10A; Slackia heliotrinireducens DSM 20476; Persephonella marina EX H1; Bacteroides fragilis NCTC 9434; Capnocytophaga ochracea DSM 7271; Flavobacterium psychrophilum JIP02 86; Akkermansia muciniphila ATCC BAA 835; Roseiflexus castenholzii DSM 13941; Roseiflexus RS1; Synechocystis PCC6803; Elusimicrobium minutum Pei191; uncultured Termite group 1 bacterium phylotype Rs D17; Fibrobacter succinogenes S85; Bacillus cereus ATCC 10987; Listeria innocua;Lactobacillus casei; Lactobacillus rhamnosus GG; Lactobacillus salivarius UCC118; Streptococcus agalactiae A909; Streptococcus agalactiae NEM316; Streptococcus agalactiae 2603; Streptococcus dysgalactiae equisimilis GGS 124; Streptococcus equi zooepidemicus MGCS10565; Streptococcus gallolyticus UCN34 uid46061; Streptococcus gordonii Challis subst CH1; Streptococcus mutans NN2025 uid46353; Streptococcus mutans; Streptococcus pyogenes M1 GAS; Streptococcus pyogenes MGAS5005; Streptococcus pyogenes MGAS2096; Streptococcus pyogenes MGAS9429; Streptococcus pyogenes MGAS10270; Streptococcus pyogenes MGAS6180; Streptococcus pyogenes MGAS315; Streptococcus pyogenes SSI-1; Streptococcus pyogenes MGAS10750; Streptococcus pyogenes NZ131; Streptococcus thermophiles CNRZ1066; Streptococcus thermophiles LMD-9; Streptococcus thermophiles LMG 18311; Clostridium botulinum A3 Loch Maree; Clostridium botulinum B Eklund 17B; Clostridium botulinum Ba4 657; Clostridium botulinum F Langeland; Clostridium cellulolyticum H10; Finegoldia magna ATCC 29328; Eubacterium rectale ATCC 33656; Mycoplasma gallisepticum; Mycoplasma mobile 163K; Mycoplasma penetrans; Mycoplasma synoviae 53; Streptobacillus moniliformis DSM 12112; Bradyrhizobium BTAi1; Nitrobacter hamburgensis X14; Rhodopseudomonas palustris BisB18; Rhodopseudomonas palustris BisB5; Parvibaculum lavamentivorans DS-1; Dinoroseobacter shibae DFL 12; Gluconacetobacter diazotrophicus Pal 5

FAPERJ; Gluconacetobacter diazotrophicus Pal 5 JGI; Azospirillum B510 uid46085; Rhodospirillum rubrum ATCC 11170; Diaphorobacter TPSY uid29975; Verminephrobacter eiseniae EF01-2; Neisseria meningitides 053442; Neisseria meningitides alpha14; Neisseria meningitides Z2491; Desulfovibrio salexigens DSM 2638; Campylobacter jejuni doylei 269 97; Campylobacter jejuni 81116; Campylobacter jejuni; Campylobacter lari RM2100; Helicobacter hepaticus; Wolinella succinogenes; Tolumonas auensis DSM 9187; Pseudoalteromonas atlantica T6c; Shewanella pealeana ATCC 700345; Legionella pneumophila Paris; Actinobacillus succinogenes 130Z; Pasteurella multocida; Francisella tularensis novicida U112; Francisella tularensis holarctica; Francisella tularensis FSC 198; Francisella tularensis tularensis; Francisella tularensis WY96-3418; and Treponema denticola ATCC 35405. The Cas9 protein may be referred by one of skill in the art in the literature as Csn1. An exemplary *S. pyogenes* Cas9 protein sequence is shown below. See Deltcheva et al., *Nature* 471, 602-607 (2011) hereby incorporated by reference in its entirety.

MDKKYSIGLDIGTNSVGWAVITDEYKVPSKKFKVLGNTDRHSIKKNLIGALLFDSGETAE
ATRLKRTARRRYTRRKNRICYLQEIFSNEMAKVDDSFFHRLEESFLVEEDKKHERHPIFG
NIVDEVAYHEKYPTIYHLRKKLVDSTDKADLRLIYLALAHMIKFRGHFLIEGDLNPDNSD
VDKLFIQLVQTYNQLFEENPINASGVDAKAILSARLSKSRRLENLIAQLPGEKKNGLFGN
LIALSLGLTPNFKSNFDLAEDAKLQLSKDTYDDDLDNLLAQIGDQYADLFLAAKNLSDAI
LLSDILRVNTEITKAPLSASMIKRYDEHHQDLTLLKALVRQQLPEKYKEIFFDQSKNGYA
GYIDGGASQEEFYKFIKPILEKMDGTEELLVKLNREDLLRKQRTFDNGSIPHQIHLGELH
AILRRQEDFYPFLKDNREKIEKILTFRIPYYVGPLARGNSRFAWMTRKSEETITPWNFEE
VVDKGASAQSFIERMTNFDKNLPNEKVLPKHSLLYEYFTVYNELTKVKYVTEGMRKPAFL
SGEQKKAIVDLLFKTNRKVTVKQLKEDYFKKIECFDSVEISGVEDRFNASLGTYHDLLKI
IKDKDFLDNEENEDILEDIVLTLTLFEDREMIEERLKTYAHLFDDKVMKQLKRRRYTGWG
RLSRKLINGIRDKQSGKTILDFLKSDGFANRNFMQLIHDDSLTFKEDIQKAQVSGQGDSL
HEHIANLAGSPAIKKGILQTVKVVDELVKVMGRHKPENIVIEMARENQTTQKGQKNSRER
MKRIEEGIKELGSQILKEHPVENTQLQNEKLYLYYLQNGRDMYVDQELDINRLSDYDVDH
IVPQSFLKDDSIDNKVLTRSDKNRGKSDNVPSEEVVKKMKNYWRQLLNAKLITQRKFDNL
TKAERGGLSELDKAGFIKRQLVETRQITKHVAQILDSRMNTKYDENDKLIREVKVITLKS
KLVSDFRKDFQFYKVREINNYHHAHDAYLNAVVGTALIKKYPKLESEFVYGDYKVYDVR
K
MIAKSEQEIGKATAKYFFYSNIMNFFKTEITLANGEIRKRPLIETNGETGEIVWDKGRDF
ATVRKVLSMPQVNIVKKTEVQTGGFSKESILPKRNSDKLIARKKDWDPKKYGGFDSPTVA
YSVLVVAKVEKGKSKKLKSVKELLGITIMERSSFEKNPIDFLEAKGYKEVKKDLIIKLPK
YSLFELENGRKRMLASAGELQKGNELALPSKYVNFLYLASHYEKLKGSPEDNEQKQLFVE
QHKHYLDEIIEQISEFSKRVILADANLDKVLSAYNKHRDKPIREQAENIIHLFTLTNLGA
PAAFKYFDTTIDRKRYTSTKEVLDATLIHQSITGLYETRIDLSQLGGD

According to one aspect, the specificity of gRNA-directed Cas9 cleavage is used as a mechanism for genome engineering and as a drive gene. According to one aspect, hybridization of the gRNA need not be 100 percent in order for the enzyme to recognize the gRNA/DNA hybrid and affect cleavage. Some off-target activity could occur. For example, the *S. pyogenes* system tolerates mismatches in the first 6 bases out of the 20bp mature spacer sequence *in vitro*. According to one aspect, greater stringency may be beneficial *in vivo* when potential off-target

sites matching (last 14 bp) NGG exist within the human reference genome for the gRNAs.

According to certain aspects, specificity may be improved. When interference is sensitive to the melting temperature of the gRNA-DNA hybrid, AT-rich target sequences may have fewer off-target sites. Carefully choosing target sites to avoid pseudo-sites with at least 14bp matching sequences elsewhere in the genome may improve specificity. The use of a Cas9 variant requiring a longer PAM sequence may reduce the frequency of off-target sites. Directed evolution may improve Cas9 specificity to a level sufficient to completely preclude off-target activity, ideally requiring a perfect 20bp gRNA match with a minimal PAM. Accordingly, modification to the Cas9 protein is a representative embodiment of the present disclosure. CRISPR systems useful in the present disclosure are described in R. Barrangou, P. Horvath, CRISPR: new horizons in phage resistance and strain identification. *Annual review of food science and technology* 3, 143 (2012) and B. Wiedenheft, S. H. Sternberg, J. A. Doudna, RNA-guided genetic silencing systems in bacteria and archaea. *Nature* 482, 331 (Feb 16, 2012) each of which are hereby incorporated by reference in their entireties.

According to certain aspects, the DNA binding protein is altered or otherwise modified to inactivate the nuclease activity. Such alteration or modification includes altering one or more amino acids to inactivate the nuclease activity or the nuclease domain. Such modification includes removing the polypeptide sequence or polypeptide sequences exhibiting nuclease activity, i.e. the nuclease domain, such that the polypeptide sequence or polypeptide sequences exhibiting nuclease activity, i.e. nuclease domain, are absent from the DNA binding protein. Other modifications to inactivate nuclease activity will be readily apparent to one of skill in the art based on the present disclosure. Accordingly, a nuclease-null DNA binding protein includes polypeptide sequences modified to inactivate nuclease activity or removal of a polypeptide sequence or sequences to inactivate nuclease activity. The nuclease-null DNA binding protein retains the ability to bind to DNA even though the nuclease activity has been inactivated. Accordingly, the DNA binding protein includes the polypeptide sequence or sequences required for DNA binding but may lack the one or more or all of the nuclease sequences exhibiting nuclease activity. Accordingly, the DNA binding protein includes the polypeptide sequence or sequences required for DNA binding but may have one or more or all of the nuclease sequences exhibiting nuclease activity inactivated.

According to one aspect, a DNA binding protein having two or more nuclease domains may be modified or altered to inactivate all but one of the nuclease domains. Such a modified or altered DNA binding protein is referred to as a DNA binding protein nickase, to the extent that the DNA binding protein cuts or nicks only one strand of double stranded DNA. When guided by RNA to DNA, the DNA binding protein nickase is referred to as an RNA guided DNA binding protein nickase.

An exemplary DNA binding protein is an RNA guided DNA binding protein nuclease of a

Type II CRISPR System, such as a Cas9 protein or modified Cas9 or homolog of Cs9. An exemplary DNA binding protein is a Cas9 protein nickase. An exemplary DNA binding protein is an RNA guided DNA binding protein of a Type II CRISPR System which lacks nuclease activity. An exemplary DNA binding protein is a nuclease-null Cas9 protein.

According to certain aspects of methods of RNA-guided genome regulation described herein, Cas9 is altered to reduce, substantially reduce or eliminate nuclease activity. According to one aspect, Cas9 nuclease activity is reduced, substantially reduced or eliminated by altering the RuvC nuclease domain or the HNH nuclease domain. According to one aspect, the RuvC nuclease domain is inactivated. According to one aspect, the HNH nuclease domain is inactivated. According to one aspect, the RuvC nuclease domain and the HNH nuclease domain are inactivated. According to an additional aspect, Cas9 proteins are provided where the RuvC nuclease domain and the HNH nuclease domain are inactivated. According to an additional aspect, nuclease-null Cas9 proteins are provided insofar as the RuvC nuclease domain and the HNH nuclease domain are inactivated. According to an additional aspect, a Cas9 nickase is provided where either the RuvC nuclease domain or the HNH nuclease domain is inactivated, thereby leaving the remaining nuclease domain active for nuclease activity. In this manner, only one strand of the double stranded DNA is cut or nicked.

According to an additional aspect, nuclease-null Cas9 proteins are provided where one or more amino acids in Cas9 are altered or otherwise removed to provide nuclease-null Cas9 proteins. According to one aspect, the amino acids include D10 and H840. See Jinke et al., *Science* 337, 816-821 (2012). According to an additional aspect, the amino acids include D839 and N863. According to one aspect, one or more or all of D10, H840, D839 and H863 are substituted with an amino acid which reduces, substantially eliminates or eliminates nuclease activity. According to one aspect, one or more or all of D10, H840, D839 and H863 are substituted with alanine. According to one aspect, a Cas9 protein having one or more or all of D10, H840, D839 and H863 substituted with an amino acid which reduces, substantially eliminates or eliminates nuclease activity, such as alanine, is referred to as a nuclease-null Cas9 or Cas9N and exhibits reduced or eliminated nuclease activity, or nuclease activity is absent or substantially absent within levels of detection. According to this aspect, nuclease activity for a Cas9N may be undetectable using known assays, i.e. below the level of detection of known assays.

According to one aspect, the Cas9 protein, Cas9 protein nickase or nuclease null Cas9 includes homologs and orthologs thereof which retain the ability of the protein to bind to the DNA and be guided by the RNA. According to one aspect, the Cas9 protein includes the sequence as set forth for naturally occurring Cas9 from *S. pyogenes* and protein sequences having at least 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, 98% or 99% homology thereto and being a DNA binding protein, such as an RNA guided DNA binding protein.

According to one aspect, an engineered Cas9-gRNA system is provided which enables RNA-guided genome regulation in cells by tethering transcriptional activation domains to either a nuclease-null Cas9 or to guide RNAs. According to one aspect of the present disclosure, one or more transcriptional regulatory proteins or domains (such terms are used interchangeably) are joined or otherwise connected to a nuclease-deficient Cas9 or one or more guide RNA (gRNA). The transcriptional regulatory domains correspond to targeted loci. Accordingly, aspects of the present disclosure include methods and materials for localizing transcriptional regulatory domains to targeted loci by fusing, connecting or joining such domains to either Cas9N or to the gRNA. According to certain aspects, methods are provided for regulating endogenous genes using Cas9N, one or more gRNAs and a transcriptional regulatory protein or domain. According to one aspect, an endogenous gene can be any desired gene, referred to herein as a target gene.

According to one aspect, a Cas9N-fusion protein capable of transcriptional activation is provided. According to one aspect, a VP64 activation domain (see Zhang et al., *Nature Biotechnology* 29, 149-153 (2011) hereby incorporated by reference in its entirety) is joined, fused, connected or otherwise tethered to the C terminus of Cas9N. According to one method, the transcriptional regulatory domain is provided to the site of target genomic DNA by the Cas9N protein. According to one method, a Cas9N fused to a transcriptional regulatory domain is provided within a cell along with one or more guide RNAs. The Cas9N with the transcriptional regulatory domain fused thereto bind at or near target genomic DNA. The one or more guide RNAs bind at or near target genomic DNA. The transcriptional regulatory domain regulates expression of the target gene. According to a specific aspect, a Cas9N-VP64 fusion activated transcription of reporter constructs when combined with gRNAs targeting sequences near the promoter, thereby displaying RNA-guided transcriptional activation.

According to one aspect, a gRNA-fusion protein capable of transcriptional activation is provided. According to one aspect, a VP64 activation domain is joined, fused, connected or otherwise tethered to the gRNA. According to one method, the transcriptional regulatory domain is provided to the site of target genomic DNA by the gRNA. According to one method, a gRNA fused to a transcriptional regulatory domain is provided within a cell along with a Cas9N protein. The Cas9N binds at or near target genomic DNA. The one or more guide RNAs with the transcriptional regulatory protein or domain fused thereto bind at or near target genomic DNA. The transcriptional regulatory domain regulates expression of the target gene. According to a specific aspect, a Cas9N protein and a gRNA fused with a transcriptional regulatory domain activated transcription of reporter constructs, thereby displaying RNA-guided transcriptional activation.

According to one aspect, the transcriptional regulator protein or domain is a transcriptional activator. According to one aspect, the transcriptional regulator protein or domain upregulates

expression of the target nucleic acid. According to one aspect, the transcriptional regulator protein or domain is a transcriptional repressor. According to one aspect, the transcriptional regulator protein or domain downregulates expression of the target nucleic acid. Transcriptional activators and transcriptional repressors can be readily identified by one of skill in the art based on the present disclosure.

According to one aspect, the foreign nucleic acid sequence encodes two or more guide RNAs with each RNA being complementary to an adjacent site in the DNA target nucleic acid and also encodes at least one RNA guided DNA binding protein nickase and being guided by the two or more RNAs, wherein the two or more RNAs and the at least one RNA guided DNA binding protein nickase are expressed and wherein the at least one RNA guided DNA binding protein nickase co-localizes with the two or more RNAs to the DNA target nucleic acid and nicks the DNA target nucleic acid resulting in two or more adjacent nicks. According to certain aspects, the two or more adjacent nicks are on the same strand of the double stranded DNA. According to one aspect, the two or more adjacent nicks are on the same strand of the double stranded DNA and result in homologous recombination. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA and create double stranded breaks. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA and create double stranded breaks resulting in nonhomologous end joining. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA and are offset with respect to one another. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA and are offset with respect to one another and create double stranded breaks. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA and are offset with respect to one another and create double stranded breaks resulting in nonhomologous end joining. According to one aspect, the two or more adjacent nicks are on different strands of the double stranded DNA and create double stranded breaks resulting in fragmentation of the target nucleic acid thereby preventing expression of the target nucleic acid.

According to certain aspects, binding specificity of the RNA guided DNA binding protein may be increased according to methods described herein. According to one aspect, off-set nicks are used in methods of genome-editing. A large majority of nicks seldom result in NHEJ events, (see Certo et al., *Nature Methods* 8, 671-676 (2011) hereby incorporated by reference in its entirety) thus minimizing the effects of off-target nicking. In contrast, inducing off-set nicks to generate double stranded breaks (DSBs) is highly effective at inducing gene disruption. According to certain aspects, 5' overhangs generate more significant NHEJ events as opposed to 3' overhangs. Similarly, 3' overhangs favor HR over NHEJ events, although the total number of HR

events is significantly lower than when a 5' overhang is generated. Accordingly, methods are provided for using nicks for homologous recombination and off-set nicks for generating double stranded breaks to minimize the effects of off-target Cas9-gRNA activity.

Germline cells according to the present disclosure include any germline cell into which foreign nucleic acids can be introduced and expressed as described herein. It is to be understood that the basic concepts of the present disclosure described herein are not limited by cell type. Germline cells according to the present disclosure include eukaryotic germline cells, prokaryotic germline cells, animal germline cells, mammalian germline cells, plant germline cells, insect germline cells, fungal germline cells, archael germline cells, eubacterial germline cells and the like. Further, germline cells include any in which it would be beneficial or desirable to introduce a foreign nucleic acid sequence described herein.

Target nucleic acids include any nucleic acid sequence to which a co-localization complex as described herein can be useful to either cut, nick or regulate. Target nucleic acids include genes. For purposes of the present disclosure, DNA, such as double stranded DNA, can include the target nucleic acid and a co-localization complex can bind to or otherwise co-localize with the DNA at or adjacent or near the target nucleic acid and in a manner in which the co-localization complex may have a desired effect on the target nucleic acid. Such target nucleic acids can include endogenous (or naturally occurring) nucleic acids and exogenous (or foreign) nucleic acids. One of skill based on the present disclosure will readily be able to identify or design guide RNAs and Cas9 proteins which co-localize to a DNA including a target nucleic acid. One of skill will further be able to identify transcriptional regulator proteins or domains which likewise co-localize to a DNA including a target nucleic acid. DNA includes genomic DNA, mitochondrial DNA, viral DNA or exogenous DNA.

Foreign nucleic acids (i.e. those which are not part of a cell's natural nucleic acid composition) may be introduced into a cell using any method known to those skilled in the art for such introduction. Such methods include transfection, transduction, viral transduction, microinjection, lipofection, nucleofection, nanoparticle bombardment, transformation, conjugation and the like. One of skill in the art will readily understand and adapt such methods using readily identifiable literature sources.

Transcriptional regulator proteins or domains which are transcriptional activators or transcriptional repressors may be readily identifiable by those skilled in the art based on the present disclosure and the particular germline cell.

The following examples are set forth as being representative of the present disclosure. These examples are not to be construed as limiting the scope of the present disclosure as these and other equivalent embodiments will be apparent in view of the present disclosure, figures and accompanying claims.

EXAMPLE I

Cas9 Gene Drives

Fig. 1A is a schematic showing the colocalization of a guide RNA and a Cas9 nuclease to a target DNA. The target DNA sites include a "protospacer" sequence that matches the "spacer" of the guide RNA and a short protospacer-adjacent motif (PAM) required for Cas9 binding. An exemplary Cas9 gene drive shown in Fig. 1A is a foreign nucleic acid sequence as described herein (which may be referred to as a "cassette" as that term is understood by those of skill in the art) that encodes at least an RNA guided Cas9 nuclease, one or more guide RNAs that are complementary to one or more target DNA sites, such as within an adjacent essential gene, and flanking sequences. The cassette may also include any cargo genetic material, such as one or more genes to be expressed by the germline cell into which the cargo genetic material is to be inserted or to be expressed by the descendants of that cell. The Cas9 gene drive is expressed to produce Cas9 and the one or more guide RNAs which then co-localize at one or more target DNA sites on a first chromosome of a chromosome pair and where the Cas9 then cleaves the target DNA on the first chromosome in a site specific manner producing a double stranded break. Homology-directed repair of Cas9-induced double-strand breaks then inserts the Cas9 gene drive into the first chromosome at the cut site. According to certain aspects, the inserted Cas9 gene drive is then expressed to produce Cas9 and the one or more guide RNAs which then co-localize at the one or more target DNA sites on the second chromosome of the chromosome pair and then cleaves the target DNA in a site specific manner producing a double stranded break. Homology directed repair of the Cas9-induced double stranded breaks uses the intact Cas9 drive-containing chromosome as a template thereby inserting the Cas9 drive into the second chromosome of the chromosome pair to create a chromosome pair that is homozygous for the Cas9 gene drive. This is depicted in Fig. 1B.

Fig. 1C depicts that a mutation at the target DNA site may prevent cutting by a single Cas9 protein. Accordingly, aspects of the present disclosure depicted in Fig. 1D provide multiple guide RNAs for multiple target DNA sites. In this manner, RNA-guided gene drives can overcome resistant alleles by targeting multiple adjacent sequences such that no single mutation can confer resistance. The chances that a target site will not be cleaved due to a mutation are greatly reduced as the target DNA site will likely be cleaved in multiple locations thereby providing a cut site into which the RNA guided gene drive can be inserted. As shown in Fig. 1E, repair using the alternative non-homologous end-joining (NHEJ) pathway can delete all recognition sites, creating a drive-resistant allele that outcompetes the drive. Fig. 1F depicts that NHEJ events that disrupt an essential gene are highly deleterious and will be selected against in favor of the drive. Choosing target sites within an essential gene ensures that incorrect repair events cannot create resistant alleles by deleting all of the targeted sequences. Non-homologous end joining can create small

insertions or deletions at the junction of the repair which can prevent binding of a guide RNA and Cas9 to the repair site. As a result, the RNA guided gene drive may not be able to be inserted into this site on the chromosome if there is only a single guide RNA for this site encoded in the RNA guided gene drive. However, having the RNA guided gene drive encode for multiple guide RNAs targeting multiple sites and therefore making multiple cut sites will increase the odds that the RNA guided gene drive will be inserted into the chromosome. Further, aspects of the present disclosure are directed to targeting multiple sites for Cas9 within an essential gene as a method for improving efficiency of inserting the RNA guided gene drive into a chromosome.

EXAMPLE II

Editing and Regulating Multiple Endogenous Loci

According to certain aspects of the present disclosure, an RNA-guided gene drive can edit multiple endogenous alleles distal to the drive itself by incorporating guide RNAs that target the respective wild-type alleles. As shown in Fig. 2A, multiple guide RNAs are provided for a plurality of gene sequences, i.e., Gene A WT, Gene B WT and Essential Gene WT. Expression of the multiple guide RNAs and the Cas9 protein results in each of the genes being modified by Cas9. According to one aspect, the Cas9 drive may also include one or more orthogonal nuclease null Cas9 proteins and associated guide RNAs with transcriptional regulators which can colocalize to a target DNA site so that the transcriptional regulator can regulate the gene. See Fig. 7. As shown in Fig. 2B, additional subsequent Cas9 drives can be used to update the changes that have been previously made by a prior Cas9 drive. For example, if a particular drive causes unexpected side effects, a second drive released as an update can overwrite one or all modifications encoded by the first drive. According to this aspect, the second drive can be designed to remove the existing inserted drive and then itself be inserted into the chromosome. In addition, one or more genes that have been modified by a prior RNA guided gene drive may be removed and replaced with a desired gene. Similarly, as depicted in Fig. 2C, a third restorative drive can restore the wild type sequence in all genomic locations except for the presence of the Cas9 gene and guide RNAs required to spread the restorative drive. In this manner, a population can be modified to revert back to its wild type. Accordingly, aspects of the present disclosure are directed to the modification or replacing of a previously inserted RNA guided Cas9 gene drive by introducing a second RNA guided Cas9 gene drive into a previously modified cell using the methods described herein. The second gene drive is designed to make further modifications beyond those made by the first gene drive or to eliminate the modifications made by the first gene drive.

EXAMPLE III

Controlling Spread

According to certain aspects, the spread of RNA-guided gene drives can be limited to a single target species or even a subpopulation by targeting a unique gene or sequence polymorphism associated with the single target species or subpopulation. Because the drive can only cut the unique sequence, it does not spread through non-target populations. Accordingly, aspects of the present disclosure are directed to methods of designing and using RNA guided gene drives that are specific for a unique gene sequence or sequence polymorphism. In this manner one or more guide RNA are designed to be complementary to the unique gene sequence or sequence polymorphism. In this manner, the DNA binding protein is restricted to localizing at the unique gene sequence or sequence polymorphism.

EXAMPLE IV

Inducing Artificial Speciation

According to certain aspects, methods are provided of using RNA-guided gene drives to block gene flow by causing genetic incompatibility between two populations. According to this aspect and as shown in Fig. 3, two gene drives that target different sequences within the same essential gene using different guide RNAs are released into two populations. The two populations become incompatible because of the loss of each copy of the essential gene within each population. Releasing this type of drive into a population will artificially split it into as many species as there are drives. As shown in exemplary Fig. 3, drive A cuts sequences 1, 2, 3, and 4 within essential gene X. As drive A spreads, it replaces gene X with a version that has sequences 1, 2, 3, and 4 recoded so they can not be cut. Drive B cuts sequences 5, 6, 7, and 8 within gene X, and its version of gene X has these sequences recoded so they can not be cut. Crossing a male with drive A and a female with drive B results in progeny that inherit one chromosome with drive A and gene X with 1/2/3/4 recoded, and one chromosome with drive B and gene X with 5/6/7/8 recoded. Drive A cuts the second copy of gene X at sites 1/2/3/4, because the second copy doesn't have those sites recoded - it only has 5/6/7/8 recoded. Similarly, drive B cuts the first copy of gene X at sites 5/6/7/8 because the first copy doesn't have those sites recoded, it only has 1/2/3/4 recoded. The organism ends up with both copies of gene X cut. Since gene X is essential and the organism lacks an intact copy of gene X to fix them with, the organism dies. Drive A and drive B are accordingly incompatible.

According to this aspect, each drive encodes a different version of the essential gene - a first foreign nucleic acid sequence and a second foreign nucleic acid sequence - that each preserve the amino acid sequence of the essential protein encoded by that gene, but which have different

sets of target locations recoded - the first drive has sites 1/2/3/4 recoded to be noncomplementary to guides 1/2/3/4, whereas the second drive has sites 5/6/7/8 recoded to be noncomplementary to guides 5/6/7/8. When the drives are crossed, the first drive cleaves the second drive's copy of the essential gene at 1/2/3/4 because it doesn't have those sites recoded, while the second drive cleaves the first drive's copy of the essential gene at 5/6/7/8 because it doesn't have those sites recoded.

More complex gene drives can block gene flow between an engineered subpopulation and the wild-type population without modifying the latter (see Fig. 8). This barrier can even be one-way, allowing the engineered population to receive but not donate genetic material to the wild-type population (see Fig. 9). These or related designs might be used to prevent other gene drives or genetic modifications from spreading to unmodified members of the same or related species. According to this aspect, only one of the two populations are modified and two separate RNA guided gene drives are used. The first drive creates an insertion site for the second drive. The second drive 1) includes a copy of an essential gene, such that it copies that gene as it spreads into the insertion site created by the first drive, and 2) kills the wild-type copy of that essential gene. When an organism with the second drive mates with an organism with the first drive, the second drive is copied into the insertion site created by the first drive, adding a copy of the essential gene and removing the wild-type copy. Net change in essential genes is 0, so the cell is viable. When an organism with the second drive is mated with a wild-type organism, the drive and its essential gene is unable to be inserted into the wild-type chromosome because it lacks an insertion site. However, the second drive still kills the wild-type copy. Net change in essential genes is -1, so the cell dies. Accordingly, any matings between organisms containing the second drive and wild-type organisms produce no progeny. This aspect of the disclosure provides a method of making transgenic organisms incompatible with the wild-type population to prevent the spread of transgenes.

EXAMPLE V

Population Suppression and Extinction

According to certain aspects, RNA-guided nucleases can create several different forms of sex-biasing gene drives that may be useful for population control. As shown in Fig. 4A, expressing Cas9 and guide RNAs targeting the X chromosome exclusively during male meiosis will result in a classical "meiotic" Y-drive analogous to those found in nature. The meiotic gene drive eliminates the competing sex chromosome during meiosis, producing a biased pool of gametes. Drive expression must be strictly limited to pre-meiosis or the drive will be lethal. This type of drive can bias the gametes of organisms employing XY, ZW, and to a lesser extent XO sex-determination systems in favor of either sex. Further with respect to Fig. 4A, an RNA guided gene drive is introduced into the Y chromosome of a spermatocyte. The RNA guided gene drive includes a

plurality of guide RNA complementary to target sequences on the X chromosome. When the RNA guided gene drive is expressed, the guide RNAs and the Cas9 nuclease colocalize at the X chromosome where the Cas9 nuclease creates double stranded breaks rendering the X chromosome inoperable. After meiosis, only sperm having the Y chromosome will be viable.

Importantly, orthogonal Cas9 nucleases should be capable of circumventing meiotic chromosome silencing. Aspects of the present disclosure are directed to the use of a first RNA guided gene drive as described herein on a Y chromosome to express Cas9 with suitable guide RNA prior to generation of sperm where the expressed Cas9 and the suitable guide RNA cuts an autosome and a second RNA guided gene drive encoding a second Cas9 and suitable guide RNA is inserted into the autosome. During spermatogenesis, the second RNA guided gene drive in the autosome is expressed and the expressed second Cas9 and suitable guide RNA cuts the X chromosome thereby rendering it inoperable. This method allows the cutting of the X chromosome even in species in which the X and Y chromosomes are not expressed during spermatogenesis. As shown in Fig. 4B, Cas9 causes the cassette encoding Cas9B to be copied from the sex chromosome to an autosome early in development, where it is free to eliminate the competing sex chromosome during meiosis. According to one aspect, RNA guided gene drives that bias for a particular sex are provided on the sex chromosome which they bias the population towards. For example, males have Y chromosomes in mammals, so a drive that biases populations towards males has to be on the Y chromosome, as it then assists the Y chromosome make more copies of itself. As mammalian sex chromosomes are not expressed during meiosis (sperm and egg formation), two copies of Cas9 can be used to facilitate proper insertion of the drives into a chromosome which when expressed will produce a Cas9 that will cut the sex chromosome. The first Cas9 acts early in development, well before sperm formation, and it cuts an autosome (non sex chromosomes that are not silenced during meiosis) thereby allowing insertion of an RNA guided gene drive encoding a second Cas9 into the autosome. This second Cas9 is expressed from the autosome during sperm formation and cuts the X chromosome, thereby ensuring that most viable sperm contain the Y chromosome. Thy above described method therefore uses two RNA guided gene drives where one is inserted into an expressable autosome to bias populations towards males in the situation where sex chromosomes can't express genes during meiosis.

As shown in Fig. 4C, "zygotic" sex-biasing drives eliminate opposite-sex siblings during the zygotic or post-zygotic stage (see Rice and Friberg 2008 hereby incorporated by reference in its entirety). Importantly, they do not require highly specific expression patterns in order to function. X-drives eliminate male zygotes by simply targeting sites on the Y chromosome (Fig. 4C). In Y-drives, a gene drive cassette copies itself in place of an essential gene on the X chromosome, the loss of which is complemented in males by a copy inserted elsewhere on the driving Y chromosome (Fig. 4D). It may be necessary to adjust expression of the transplanted essential gene

to account for dosage compensation in some species.

With respect to Fig. 4C, a wild type spermatocyte creates a wild type sperm with an X chromosome and a wild type sperm with a Y chromosome. An oocyte is provided with an RNA guided gene drive with one or more guide RNAs complementary to the Y chromosome, which is referred to as an "X drive oocyte". When the wild type sperm with the Y chromosome is combined with the oocyte, the RNA guided gene drive is expressed, the guide RNA and the Cas9 nuclease colocalize to the Y chromosome where the Cas9 nuclease creates double stranded breaks in the Y chromosome rendering the Y chromosome inoperable and thereby producing an inoperable zygote. In contrast, when the wild type sperm with the X chromosome is combined with the oocyte, the RNA guided gene drive is expressed, guide RNA complementary to the X chromosome and the Cas9 nuclease colocalize to the X chromosome where the Cas9 nuclease creates double stranded breaks in the X chromosome which is repaired by homologous recombination thereby inserting the RNA guided gene drive into the X chromosome rendering the zygote homozygous for the X chromosome. This copy of the X chromosome remains viable because the double strand breaks are repaired when the RNA guided gene drive is copied. The resulting progeny are all female, because the males are not viable due to loss of the Y chromosome. These females have an advantage over other females that do not encode an RNA guided gene drive because they do not have to compete with male siblings for resources. Consequently, the X chromosome with an RNA guided gene drive has a fitness advantage. With respect to Fig. 4D, zygotic Y-drives encode an essential gene normally found on the X chromosome and utilize a gene drive cassette to eliminate that essential gene from the X chromosome. The Y drive is inserted from the paternal to the maternal X chromosome, leaving female zygotes without any copies of the essential gene. In particular, a spermatocyte is provided with a Y chromosome having an essential gene from the X chromosome and an RNA guided gene drive encoding one or more guide RNAs complementary to the essential gene on the X chromosome. The X chromosome of the spermatocyte also includes the same RNA guided gene drive. The spermatocyte produces a sperm with the Y chromosome and a sperm with the X chromosome. The sperm with the Y chromosome is combined with a wild type oocyte having an X chromosome with the essential gene. When the RNA guided gene drive is expressed, the one or more guide RNAs colocalize with the Cas9 nuclease to the X chromosome where the Cas9 nuclease cuts out the essential gene from the X chromosome. However, since the essential gene from the X chromosome is present on the Y chromosome, the zygote will be viable. In contrast, when the sperm with the X chromosome including the RNA guided gene drive combines with the wild type oocyte with the X chromosome, the expressed one or more guide RNAs colocalize with the Cas9 nuclease to the X chromosome where the Cas9 nuclease cuts out the essential gene from the X chromosome. Since the essential gene no longer exists in the oocyte, the oocyte is not viable.

The fitness benefit of zygotic drives will vary depending on the degree of sibling competition, the extent of parental investment in offspring, and the mating dynamics in adults. All zygotic Y-drives should produce at least as many if not more sons than competing wild-type Y chromosomes, and none of those sons will have to compete with sisters. Unlike other drive types, the host range of zygotic Y-drives can be restricted by ensuring that matings with specific at-risk related species or non-targeted subpopulations are sterile. This can be accomplished by incorporating guide RNAs that cleave unique sequences on the X-chromosomes to eliminate hybrid males.

Releasing a Y-drive will render the local population extinct if mutations capable of blocking the drive do not quickly arise on the X chromosome or one of the autosomes. The drive can be halted and eventually eliminated by releasing organisms bearing sex chromosomes that are immune to cutting. Close monitoring and control can prevent total extinction.

RNA-guided nucleases can also create less aggressive "sterile-daughter" Y chromosomes that can suppress populations but do not spread like a drive (Fig. 10A). A copy of Cas9 on the Y chromosome that targets an X-linked gene essential for female fertility rather than viability will still eliminate the reproductive potential of female progeny, but will provide no advantage to sons. Due to the minor cost of Cas9 activity, a sterile-daughter Y will eventually be outcompeted by the wild-type Y chromosome, but not before causing severe population suppression if introduced in sufficiently large numbers. This approach is closely related to the female-specific lethality and sterility methods being developed for the control of insect and fish populations, but is likely to be both more straightforward to construct and more evolutionarily stable due to its location on the Y.

Hybrid approaches to population control might combine the advantages of the Y-drive and sterile-daughter methods (Fig. 5A-B). For example, a Y chromosome that targets both an essential gene and a female fertility gene will quickly drive itself through the population because it is a Y-drive. However, it would lose its fitness advantage upon encountering a subsequently released standard gene drive that recodes the essential gene, but not the female fertility gene, on the X chromosome. Hybrid females will grow up to be infertile instead of dying embryonically, causing continued suppression without extinction.

A related approach to population control utilizes RNA-guided gene drives to disrupt one or more genes that are 1) required for fertility or viability, but 2) one intact copy largely suffices for function. Such a drive would cut and replace the target gene in the germline cells of organisms that inherit one copy of the drive and one copy of the wild-type gene at any time *after* the gene's function is required. For example, a gene required for gonad development and subsequent fertility might be cut and replaced by an RNA-guided gene drive just prior to meiosis. Since an organism that inherited only one copy would already have undergone correct gonad development because the

wild-type copy suffices for this purpose, it would have normal fertility. However, most or all offspring would inherit the drive due to drive copying just prior to sperm and/or egg production. This design allows the drive to spread through the population when rare because most mating events will produce more individuals that inherit one copy of the drive and one copy of the wild-type. Later on, matings between two individuals that carry the drive will produce sterile offspring, leading to a population crash. This scenario has been modeled extensively as described in: Burt, A. Site-specific selfish genes as tools for the control and genetic engineering of natural populations. Proc. Biol. Sci. 270, 921–928 (2003). Deredec, A., Burt, A. & Godfray, H. C. J. The population genetics of using homing endonuclease genes in vector and pest management. Genetics 179, 2013–2026 (2008). North, A., Burt, A. & Godfray, H. C. J. Modelling the spatial spread of a homing endonuclease gene in a mosquito population. J. Appl. Ecol. 50, 1216–1225 (2013), each of which is hereby incorporated by reference in their entireties. Methods of restricting gene expression to germline cells and/or the time period prior to meiosis are known to those in the art.

EXAMPLE VI

Applications

The gene drives described herein have particular practical utility in the eradication of infectious diseases and the control of invasive species. Such RNA guided Cas9 gene drives may be used to quickly spread protective alleles through threatened or soon-to-be-threatened species such as amphibians. Such RNA guided Cas9 gene drives may also be used to immunize wild populations that commonly serve as reservoirs for human viruses by targeting dsDNA viruses with Cas9 and RNA viruses with RNAi machinery from a foreign species carried by the drive. Disease vectors can be engineered to be unable to acquire the pathogen or can be eliminated using a Y drive described herein, i.e. an RNA guided Cas9 gene drive which inhibits propogation of an X chromosome. Similarly, invasive and ecologically destructive pests could be locally controlled or eradicated using Y-drives. Domesticated animals may be prevented from contributing genes to related wild species and feral populations controlled to minimize ecological damage and reduce the need for shelters and rescue agencies. Similarly, transgenic crops and animals could be genetically separated from their unmodified cousins, as could threatened species endangered by genetic dilution from more abundant or invasive relatives. Finally, gene drives could be used to directly test hypotheses concerning the evolutionary and ecological importance of genes, sex ratios, and speciation in natural environments.

The gene drives described herein have particular practical utility with vector born diseases. The human toll of vector-borne infectious disease is staggering. Malaria alone kills over 650,000 people each year, most of them children, while afflicting 200 million more with debilitating fevers that economically devastate their societies. Dengue, yellow fever, trypanosomiasis, leishmaniasis,

Chagas disease, and Lyme disease are caused by others pathogens that spread using vectors. All of these can potentially be reduced or even eliminated by driving changes in the vector that prevent transmission. Scientists have identified several candidate gene disruptions or inserted genes that interfere with the transmission of malaria (Ito et al Nature 2002, PMID:12024215: Dong et al PloS Pathogens 2012, PMID: 22216006; Isaacs et al., PNAS 2012, PMID: 22689959) and other well-studied diseases (Franz et al PNAS 2006, PMID: 16537508), but not for many other pathogens. Hence, aspects of the present disclosure are directed to directly eliminating the vector species with a Y-drive. In the case of malaria, this strategy is particularly promising against emerging mosquito vectors that prefer to bite and rest outdoors, as these behaviors are highly resistant to current control strategies focused around indoor insecticide spraying and bednets. Although all vector species must be targeted in a given area in order to stop transmission, the disease will be permanently eradicated if the newly vacated ecological niches are filled by competing non-vector species. Significantly, this strategy requires little or no understanding of the vector's molecular biology, but unavoidably entails the local or possibly global extinction of the vector species.

The gene drives described herein have particular practical utility in controlling invasive species. One of the most environmentally damaging consequences of global economic activity is the transport of invasive species, which often causes ecological disruption and the extinction of native species. Isolated ecosystems such as those on small islands are especially vulnerable. Cas9 Y-drives have tremendous potential to promote biodiversity by controlling or even eradicating these species from individual islands or possibly entire continents. Designing drive sequences unique to the invasive species, incorporating guide RNAs that will degrade the X chromosomes of at-risk relatives, and using speciation drives to render the target species genetically incompatible with its relatives target the drive to the selected species and reduce the risk of cross-species transfer without directly modifying all related species. Related species can be protected by releasing species-specific standard gene drives that recode the sites required for Y-drive function. The risk that the Y-drive might spread from the invasive population back into the native habitat is negligible for species that only invade through intentional human action, such as freshwater fish or cane toads, but Y-drive spread is all but certain for rats and other invasive stowaways. Native populations can always be protected from extinction by releasing an X-recoding drive or even a resistant X chromosome. As an example, invasive populations of low-mobility species such as Asian carp are excellent candidates for straightforward elimination by Y-drive. (Fig. 6A) Because gene flow would require deliberate human introduction, local eradication is likely to be permanent and achievable without being forced to recode the native population. In contrast, rats and other highly mobile invasive species cannot be permanently eliminated barring total extinction of the species. However, they might be controlled by periodically releasing a standard recoding drive in native populations and a Y-drive with sterile-daughter effect (Y-Drive-SD) throughout invasive

populations (Fig. 6B).

EXAMPLE VII

Agricultural Safety and Sustainability

Aspects of the present disclosure are directed to the use of the Cas9 methods described herein to control, reduce, or eliminate weeds and pests associated with agriculture. A pest generally refers to a plant or animal detrimental to humans or human concerns such as with agriculture or livestock production. The term pest may include any living organism which is invasive or prolific, detrimental, troublesome, noxious, destructive, a nuisance to either plants or animals, human or human concerns, livestock, human structures, wild ecosystems etc. The term pest may also include vermin, weed, plant and animal parasites and pathogens. A weed according to the present disclosure may refer to a plant considered undesirable in a particular situation. In a broad sense, a weed may be considered a pest that is a plant. Examples commonly are plants unwanted in human-controlled settings, such as farm fields, gardens, lawns, and parks. The term "weed" may also include any plant that grows or reproduces aggressively, or is invasive outside its native habitat.

According to one embodiment of the present disclosure, the Cas9 methods are used to insert into the genome of a germline cell of a weed or pest one or more RNA-guided "sensitizing drives." A sensitizing drive is a gene drive as described herein and may also be referred to as a "sensitizing gene drive." Accordingly, a "sensitizing drive" is a gene drive that is inserted into genomic DNA and which is transferred to progeny and renders the progeny sensitive to an external stimulus. According to one aspect, the RNA-guided "sensitizing drive" confers to the weed or pest as a result of being incorporated into genomic DNA a harmful sensitivity to a compound or chemical, such as toxicity. In this manner, the growth and proliferation of the weed or pest may be controlled by contacting the weed or pest with a compound or chemical or condition which ordinarily would not be toxic to the weed or pest. In this manner, the phenotype of the weed or pest has been altered by the sensitizing gene drive as a result of insertion of the gene drive into the germline cell and transfer to progeny. In this manner, the reference to a weed or pest may refer to the population that results from initial insertion of the sensitizing gene drive into the germline cell and mating with a wild type population. The resulting population may be referred to as an altered population or sensitized population or genetically altered population.

According to the present disclosure, the term "sensitivity" as it relates to a weed or pest means a harmful reaction, such as toxicity, to a compound or chemical or condition to which the weed or pest is exposed. The sensitizing drive is a gene drive as described herein that alters genomic DNA to result in a harmful reaction, such as toxicity, in response to a compound, chemical or condition to which the weed or pest is exposed. "Sensitizing drives" may also be

referred to herein as "toxicity drives" or "toxicity gene drive" to the extent that a chemical, compound or condition is toxic to the weed or pest as a result of having the sensitizing drive or toxicity drive present in its genome. According to one aspect, the sensitizing drive or toxicity drive is exogenously added to the germline organism. In this manner, the sensitizing gene drive or toxicity gene drive is a foreign nucleic acid, even though, according to certain application, it may include a sequence native to the weed or pest species, but not present in the germline cell into which it is to be introduced. According to one aspect, the sensitizing drive or toxicity drive is a foreign nucleic acid that is exogenously added to the germline organism using an RNA guided DNA binding protein nuclease and associated guide RNA as described herein.

According to one aspect, the weed or pest may be resistant to a particular herbicide or pesticide or the weed or pest may have developed over time a resistance to the particular herbicide or pesticide. The evolution of weeds or pests to herbicides or pesticides is a major problem for agriculture. Methods described herein use RNA-guided "sensitizing drives" to confer sensitivity of the organism to a chemical, compound or condition that was previously nontoxic or of low toxicity to the organism, so that progeny of the organism will be sensitive to the chemical, compound or condition and the organism will die as a result of toxicity or proliferation will be lowered, or the organism will be rendered sterile so that it cannot reproduce. In this manner, a nucleic acid sequence within the sensitizing gene drive is expressed by the organism, such as a weed or pest, and the expression of the nucleic acid alters the phenotype of the organism rendering it vulnerable to the chemical, compound or condition.

According to an additional aspect, a sensitizing drive or drives are used to replace resistant alleles (such as those with resistance mutations) with their ancestral (or non-mutated) equivalents to restore sensitivity. Accordingly, an organism that has developed resistance to an herbicide or pesticide through mutation of an allele may be rendered sensitive to the herbicide or pesticide by removing the mutant form of the allele having resistance and replacing it with the non-mutant form having sensitivity. In this manner, the non-mutant form of the allele is foreign to the organism or population of organisms as it includes the mutant form of the allele. Accordingly, a sensitizing gene drive might reverse known mutations that confer resistance to existing pesticides or herbicides. RNA-guided gene drives are used to reverse genome alterations that have spread through populations of an organism. According to this aspect, releasing a second drive could overwrite one or all changes caused by a first drive, as described herein. In this manner, the population of organisms is characterized by the presence of a first gene drive and first nucleic acid sequence imparting a phenotype such as toxicity that was first introduced into a germline cell to produce that phenotype in progeny and which was then transferred to progeny. However, through mutations or otherwise, the phenotype is lost. In this context, a second gene drive with a second nucleic acid sequence imparting the same phenotype is introduced into a germline cell with or

without the first nucleic acid being removed and the second nucleic acid sequence is carried through into progeny. In this manner, the first gene drive nucleic acid is overwritten by the second gene drive nucleic acid, for example, to result in toxicity, by reversing the mutation creating resistance.

According to an additional aspect, a sensitizing drive may carry a prodrug-converting enzyme that would render a prodrug molecule toxic to organisms that express the enzyme. In this manner, the enzyme is produced and should the organism be contacted with the prodrug, the enzyme will convert the prodrug into a compound or chemical that is toxic to the organism, such as a weed or pest. Various prodrug/enzyme combinations will be apparent to one of skill in the art based on the present disclosure.

According to a still additional aspect, a sensitizing drive may replace an essential gene for a version that is strongly inhibited by a particular small molecule. Accordingly, either expression of the gene would be inhibited or the expression product of the gene would be inhibited. Such inhibition may result in death of the organism or a lower proliferation. Because, in some embodiments, sensitizing drives would have no effect in the absence of their associated molecule - and in some cases vice versa - they could grant very fine control over the geography and extent of population suppression with minimal ecological risk.

According to one exemplary aspect, a sensitizing drive is used to reverse mutations allowing the western corn rootworm to resist Bt toxins (see Gassman et al PNAS, vol. 111 no. 14, pages 5141–5146, doi: 10.1073/pnas.1317179111 (2014) hereby incorporated by reference in its entirety) or horseweed and pigweed to resist the herbicide glyphosate (See Gaines, T. A. et al. Proc. Natl. Acad. Sci. 107, 1029–1034 (2010) and Ge, X., d' Avignon, D. A., Ackerman, J. J. & Sammons, R. D. Pest Manag. Sci. 66, 345–348 (2010) each of which are hereby incorporated by reference in their entireties), an herbicide currently essential to environmentally sustainable no-till agriculture. According to this aspect, organisms having the sensitizing drive are released into the wild population, individual organisms of which may or may not have resistance, and the sensitizing drive is carried through into progeny, the result of which is to decrease resistance of the progeny population (where resistance is present in the wild population) and increase sensitivity of the progeny population to a particular herbicide or pesticide (where resistance is present in the wild population.) It is to be understood that only some members of the wild type population may have resistance but that the sensitizing drive will spread through both sensitive and resistant members of the wild type population resulting in a progeny population having the sensitizing drive. According to one aspect, organisms including the sensitizing drive or drives would be released into areas not treated with herbicide or pesticide thereby creating reservoirs of sensitizing drives that could spread into adjacent areas which are treated with an herbicide or pesticide.

According to one aspect, methods are provided where a sensitizing drive or drives are used

to counter mutations which lead to resistance. According to this aspect, introducing such sensitizing drives into the genome of a wild type population, initially and over subsequent time periods, that has developed resistance to a pesticide or herbicide as a result of a mutation or mutations reverses the effect of the mutation or mutations to provide resistance and renders the pesticide or herbicide toxic to the population. Such a method allows the use of any given herbicide or pesticide indefinitely because mutations conferring resistance are countered or replaced thereby "rolling back" whatever resistance may occur.

According to an additional aspect, a plant or animal may be rendered vulnerable to a chemical or compound or condition by the inclusion into its genome (and therefore its progeny) of a sensitizing drive or drives which renders the chemical, compound or condition toxic to the plant or animal, such as a weed or pest. Accordingly, compounds which may be considered safe to humans may be toxic to plants or animals as a result of the inclusion of a sensitizing drive or drives into the genome of the plant or animal thereby rendering the chemical or compound toxic to the plant or animal. According to this aspect, existing compounds considered safe and/or effective might be applied against organisms that are not presently vulnerable to them if a sensitizing drive were to deliver a sensitive gene from an affected species or laboratory isolate or replace a gene important for fitness with a sensitive gene.

According to a further aspect, methods are provided where sensitizing drives are used to render pest populations vulnerable to molecules, compounds or chemicals that are harmless to other forms of life. Whereas current pesticides and herbicides - even those designated as "organic" - are chosen for their toxicity towards insect pests and weeds, they often harm non-pest species or even humans because the affected pathways are conserved across species. According to the present disclosure, methods are provided where a gene or genes are delivered that confer sensitivity to an organism to a normally harmless molecule. Introduction of the sensitivity gene drive effectively converts that molecule into a pesticide or herbicide highly specific for the particular pest or weed species modified by the drive. The combination of the drive and the molecule is lethal to the organism. One exemplary embodiment is the use of an enzyme/prodrug combination where the enzyme is introduced in the genome of the organism and is expressed. When the organism is exposed to the prodrug, the enzyme converts the prodrug into an active herbicide or pesticide. Analogous candidates demonstrating the principle for this aspect of the present disclosure are antiviral or anticancer therapies in which a locally produced viral or tumor-specific enzyme activates a prodrug. Examples include the cytosine deaminase/5-fluorocytosine pairing and the nitroreductase/CB1954 pairing. In the case of sensitizing drives, the enzyme would be delivered to the target species by the gene drive, causing the prodrug to become a specific pesticide. Another exemplary pairing includes primary metabolic enzymes altered to become strongly inhibited by particular chemicals. For example, an engineered invertase may become

nonfunctional in the presence of a mostly biologically inert xenobiotic chemical such as sucralose or related halogenated polysaccharides. A sensitizing drive would replace the organism's natural invertase gene with the engineered version, rendering it sensitive to the otherwise mostly inert compound. One of skill would readily be able to identify useful enzyme/chemical pairings suitable for this purpose based on the sensitizing drive methods described herein.

One of skill will readily be able to identify weeds within the scope of the present disclosure as including those weed plants harmful to agricultural crops. Such weeds may or may not be designated as "noxious weed" under federal or state law. For example, horseweed and pigweed are considered weeds that are harmful to agricultural crops but may not be designed as noxious weeds. Exemplary weeds designated as noxious weeds by the USDA include the following.

Aquatic

Latin Name	Common Name(s)
Azolla pinnata	Mosquito fern, water velvet
Caulerpa taxifolia (Mediterranean strain)	Killer algae
Eichhornia azurea	Anchored waterhyacinth, rooted waterhyacinth
Hydrilla verticillata	Hydrilla
Hygrophila polysperma	Miramar weed
Ipomoea aquatica	Water-spinach, swamp morning glory
Lagarosiphon major	African elodea
Limnophila sessiliflora	Ambulia
Melaleuca quinquenervia	Broadleaf paper bark tree
Monochoria hastata	Arrowleaf false pickerelweed
Monochoria vaginalis	Heartshape false pickerelweed
Ottelia alismoides	Duck lettuce
Sagittaria sagittifolia	Arrowhead
Salvinia auriculata	Giant salvinia
Salvinia biloba	Giant salvinia
Salvinia herzogii	Giant salvinia
Salvinia molesta	Giant salvinia
Solanum tampicense	Wetland nightshade
Sparganium erectum	Exotic bur-reed

Parasitic

Latin Name	Common Name(s)
Aeginetia spp.	Varies by species
Alectra spp.	Varies by species

Cuscuta spp.(except for natives)	Dodders
Orobanche spp. (except for natives)	Broomrapes
Striga spp.	Witchweeds

Terrestrial

Latin Name	Common Name(s)
Acacia nilotica	Prickly acacia
Ageratina adenophora	Crofton weed
Ageratina riparia	Mistflower, spreading snakeroot
Alternanthera sessilis	Sessile joyweed
Arctotheca calendula	Capeweed
Asphodelus fistulosis	Onionweed
Avena sterilis	Animated oat, wild oat
Carthamus oxyacantha	Wild safflower
Chrysopogon aciculatus	Pilipiliula
Commelina benghalensis	Benghal dayflower
Crupina vulgaris	Common crupina
Digitaria scalarum	African couchgrass, fingergrass
Digitaria velutina	Velvet fingergrass, annual couchgrass
Drymaria arenariodes	Lightning weed
Emex australis	Three-corned jack
Emex spinosa	Devil's thorn
Euphorbia terracina	False caper, Geraldton carnation weed
Galega officinalis	Goatsrue
Heracleum mantegazzianum	Giant hogweed
Imperata brasiliensis	Brazilian satintail
Imperata cylindrica	Cogongrass
Inula britannica	British yellowhead
Ischaemum rugosum	Murainograss
Leptochloa chinensis	Asian sprangletop
Lycium ferocissimum	African boxthorn
Lygodium flexuosum	Maidenhair creeper
Lygodium microphyllum	Old world climbing fern
Melastoma malabathricum	Malabar melastome
Mikania cordata	Mile-a-minute
Mikania micrantha	Bittervine
Mimosa invisa	Giant sensitive plant
Mimosa pigra	Catclaw mimosa
Moraea collina	Cape tulip
Moraea flaccida	One leaf cape tulip
Moraea miniata	Two leaf cape tulip
Moraea ochroleuca	Apricot tulip
Moraea pallida	Yellow tulip

Nassella trichotoma	Serrated tussock
Onopordum acaulon	Stemless thistle
Onopordum illyricum	Illyricum thistle
Opuntia aurantiaca	Jointed prickly pear
Oryza longistaminata	Red rice
Oryza punctata	Red rice

Oryza rufipogon	Red rice
Paspalum scrobiculatum	Kodo-millet
Pennisetum clandestinum	Kikuyugrass
Pennisetum macrourum	African feathergrass
Pennisetum pedicellatum	Kyasumagrass
Pennisetum polystachion	Missiongrass, thin napiergrass
Prosopis alpataco	Mesquite
Prosopis argentina	Mesquite
Prosopis articulata	Velvet mesquite
Prosopis burkartii	Mesquite
Prosopis caldenia	Calden
Prosopis calingastana	Cusqui
Prosopis campestris	Mesquite
Prosopis castellanosii	Mesquite
Prosopis denudans	Mesquite
Prosopis elata	Mesquite
Prosopis farcta	Syrian mesquite
Prosopis ferox	Mesquite
Prosopis fiebrigii	Mesquite
Prosopis hassleri	Mesquite
Prosopis humilis	Algaroba
Prosopis kuntzei	Mesquite
Prosopis pallida	Kiawe, algarroba
Prosopis palmeri	Mesquite
Prosopis reptans	Tornillo
Prosopis rojasiana	Mesquite
Prosopis ruizlealii	Mesquite
Prosopis ruscifolia	Mesquite
Prosopis sericantha	Mesquite
Prosopis strombulifera	Argentine screwbean
Prosopis torquata	Mesquite
Rottboellia cochinchinensis	Itchgrass
Rubus fruticosis	Wild blackberry
Rubus moluccanus	Wild raspberry
Saccharum spontaneum	Wild sugarcane
Sagittaria sagittifolia	Arrowhead
Salsola vermiculata	Wormleaf salsola

Senecio inaequidens	South African ragwort
Senecio madagascariensis	Fireweed
Setaria pumila	Cattail grass
Solanum torvum	Turkeyberry
Solanum viarum	Tropical soda apple
Spermacoce alata	Winged false buttonweed

Tridax procumbens	Coat buttons
Urochloa panicoides	Liverseed grass

Additional weeds within the scope of the present disclosure include Atriplex, Spreading; Beggarsticks, Nodding; Brome, Downy; Carrot, wild; Chamomile, scentless; Chickweed, common; Cucumber, bur; Dandelion; Fleabane, Canada; Flixweed; Grass, Stink; Grass, Tufted love; Groundcherry, Smooth; Hedge-nettle, Marsh; Horse-nettle; Horsetail, Field; Lettuce, Prickly; Mercury, Three-seeded; Muhly, Wire-stemmed; Nipplewort; Redtop; Sandbur, Long-spined; Smartweed, Swamp; Sow-thistle, Annual; Sow-thistle, Perennial; Speedwell, Corn; Vetch, Tufted; Violet, Field; Waterhemp, Common; Wood-sorrel species; Bermuda grass; Bindweed; Broadleaf plantain; Burdock; Common lambsquarters; Creeping Charlie; Dandelion; Goldenrod; Japanese Knotweed; Kudzu; Leafy spurge; Milk thistle; Poison ivy; Ragweed; Sorrel; Striga; St John's wort; Sumac; Tree of heaven; Wild carrot; Wood sorrel and Yellow nutsedge.

Additional weeds identified by scientific name include Acalypha rhomboidea Raf.; Agrostis gigantea Roth; Amaranthus rudis L.; Atriplex patula L.; Bidens cernua L.; Bromus tectorum L.; Cenchrus longispinus Hack.; Conyza Canadensis; Daucus carota L.; Descurainia sophia L.; Equisetium arvense L.; Eragrostis spp.; Lactuca scariola L.; Lapsana communis L.; Matricaria perforata Merat.; Muhlenbergia frondosa Poir.; Oxallis dillenii Jacq; Physalis virginiana Mill.; Polygonum coccineum Muhl.; Sicyos angulatus L.; Solanum carolinense L; Sonchus arvensis L.; Sonchus oleraceus L.; Stachys palustris L.; Stellaria media; Taraxacum officinale Weber.; Veronica avensis L.; Vicia cracca L.; and Viola avensis L.

It is to be understood that additional weeds within the scope of the present disclosure may be identified by sources readily available to those of skill in the art.

Common herbicides to which a weed may be resistant or develop resistance include the following. One of skill in the art will readily be able to identify herbicides toxic to particular weed species based on the present disclosure.

Common or Code Name*	Trade Name	Manufacturer
Acetochlor	Harness	Monsanto
Acifluorfen	Blazer	BASF
Aclonifen	Challenge	Bayer CropScience
Acrolein	Magnacide	Baker
Alachlor	Lasso	Monsanto
Alloxydim	Fervin	Bayer CropScience
Ametryn	Evik	Syngenta
Amicarbazone		Bayer CropScience
Amidosulfuron*	Adret, Gratil	Bayer CropScience
Aminocyclopyrachlor	MAT28/KJM44	DuPont
Aminopyralid	Milestone	Dow AgroSciences
Amitrole	Amitrol T	Bayer CropScience
Anilofos	Aniloguard	Ghardat
Asulam	Asulox	Bayer CropScience
Atrazine	Aatrex	Syngenta
	Atrazine	Terra
Azimsulfuron	Gulliver	DuPont
Beflubutamid	Herbaflex	Ube Industries
Benazolin	Asset	Bayer CropScience
Benefin	Balan	Dow AgroSciences
Bensulfuron	Londax	DuPont
Bensulide	Prefar	Gowan
Bentazon	Basagran	BASF
Benzofenap	Taipan	Bayer CropScience
Bifenox	Modown	Bayer CropScience
Bispyribac	Regiment, Velocity	Valent
Bromacil	Hyvar	DuPont
Bromoxynil	Buctril	Bayer CropScience
Butachlor	Machete	Monsanto
Butafenacil	Rebin, Inspire	Syngenta
Butroxydim	Falcon	Syngenta
Butylate	Sutan	Cedar
Cacodylic acid	Cotton-Aide HC	Monterey
Carbetamide	Carbetamex	Feinchemie
Carfentrazone	Aim, Shark, QuickSilver	FMC
Chlorsulfuron	Glean, Telar	DuPont
Chlortoluron	Dicuran	Syngenta
Cinmethylin	Argold, Cinch	BASF
Clethodim	Select, Prism	Valent
Clodinfop	Topik	Syngenta
Clodinafop-propargy*	Discover	Syngenta
Clomazone	Command	FMC
Clopyralid	Stinger, Lontrel	Dow AgroSciences
Cloransulam-methyl	Firstrate	Dow AgroSciences
Cyanazine	Bladex	DuPont
Cycloate	RoNeet	Cedar
Cyclosulfamuron	Invest	BASF
Cycloxydim	Focus	BASF
Cyhalofop	Clincher	Dow AgroSciences
DCPA	Dacthal	Amvac
2,4-D	Several	Several

Common or Code Name*	Trade Name	Manufacturer
2,4-DB	Several	Several
Dazomet	Basamid	BASF
Desmedipham	Betamix	Bayer CropScience
Desmetryn	Semeron	Syngenta
Dicamba	Banvel, Clarity	BASF
Dichlobenil	Casoron	Uniroyal
Dichlorprop	Several	Bayer CropScience
Diclofop	Hoelon	Bayer CropScience
Diclosulam	Strongarm	Dow AgroSciences
Difenzoquat	Avenge	BASF
Diflufenican	Javelin	Bayer CropScience
Diflufenzopyr	Distinct	BASF
Dimethenamid	Frontier	BASF
Diquat	Reward, Reglone	Syngenta
Dithiopyr	Dimension	Monsanto
Diuron	Karmex	DuPont
DSMA	Several	several
Endothall	Several	Atochem
EPTC	Eptam	Syngenta
Ethalfluralin	Sonalan	Dow AgroSciences
	Curbit	UAP
Ethametsulfuron	Muster	DuPont
Ethofumesate	Nortron, Prograss	Bayer Crop Science
Ethoxysulfuron	Sunrice	Bayer Crop Science
Fenoxaprop	Acclaim, Whip	Bayer CropScience
Fentrazamide	Lecs	Bayer CropScience
Flazasulfuron	Katana	Ishihara Sangyo
Florasulam	Frontline, Boxer	Dow AgroSciences
Fluazifop-P	Fusilade DX, Fusilade II	Syngenta
Flucarbazone-sodium	Everest	Arvesta
Flufenacet	Axiom (flufenacet + metribuzin)	Bayer CropScience
	Epic (flufenacet + isoxaflutole)	
Flufenpyr-ethyl*	S-3153	Valent
Flumetsulam	Python, Broadstrike	Dow AgroSciences
Flumiclorac	Resource	Valent
Flumioxazin	BroadStar, Valor, Chateau	Valent
Fluometuron	Cotoran	Syngenta
	Others	others
Flupyrsulfuron	Lexus	DuPont
Flurchloridone	Racer	Syngenta
Fluridone	Sonar	Dow AgroSciences
Fluroxypyr	Starane, Vista, Spotlight	Dow AgroSciences
Fluthiacet	Action	Syngenta
Fomesafen	Reflex, Flexstar	Syngenta
Foramsulfuron	Equip, Revolver	Bayer CropScience
Fosamine	Krenite	Dupont
Glufosinate	Liberty, Rely	Bayer CropScience
Glyphosate	Roundup, Ultra, Rodeo, Touchdown Pro	Monsanto, Syngenta
Halosulfuron	Permit, Battalion	Monsanto
	SedgeHammer	Gowan
Haloxyfop	Verdict	Dow AgroSciences
Hexazinone	Velpar	DuPont
Imazamethabenz	Assert	BASF
Imazamox	Raptor, Clearcast	BASF

Common or Code Name*	Trade Name	Manufacturer
Imazapic	Cadre, Plateau	BASF
Imazapyr	Arsenal, Habitat	BASF
Imazaquin	Image, Scepter	BASF
Imazethapyr	Pursuit	BASF
Iodosulfuron	Husar	Bayer CropScience
Isoxaben	Gallery	Dow AgroSciences
Isoxaflutole	Balance	Bayer CropScience
Lactofen	Cobra	Valent
Linuron	Lorox	DuPont
	Others	others
MCPA	Several	several
MCPB	Several	several
Mecoprop	MCPP, Mecopex	several
Mesosulfuron	Osprey	Bayer CropScience
Mefluidide	Enable	3M Company
Mesotrione	Callisto, Tenacity	Syngenta
Metham	Metham Sodium	Amvac
Metolachlor	Dual	Syngenta
	Dual Magnum, Pennant Magnum	
Metribuzin	Sencor, Lexone	Bayer CropScience
Metsulfuron	Ally, Escort	DuPont
MSMA	Several	several
Napropamide	Devrinol	United Phosphorus
Naptalam	Alanap	Uniroyal
Nicosulfuron	Accent	DuPont
Norflurazon	Zorial, Solicam	Syngenta
Oryzalin	Surflan	Dow AgroSciences
Oxadiazon	Ronstar	Lesco
Oxasulfuron*	Expert	Syngenta
Oxyfluorfen	Goal	Dow AgroSciences
Paraquat	Gramoxone Inteon, Gramoxone Max	Syngenta
Pebulate	Tillam	Monterey
Pelargonic acid	Scythe	Dow AgroSciences
Pendimethalin	Prowl	BASF
Penoxsulam	Grasp, Granite	Dow AgroSciences
Phenmedipham	Spin-Aid	Bayer CropScience
Picloram	Tordon	Dow AgroSciences
Picolinafen*	Pico	BASF
Pinoxaden	Axial	Syngenta
Primisulfuron	Beacon	Syngenta
Prodiamine	Barricade	Syngenta
Prometon	Pramitol	Agriliance
Prometryn	Caparol	Syngenta
Pronamide	Kerb	Dow AgroSciences
Propanil	Stam	Dow AgroSciences
Propaquizafop	Shogun	Syngenta
Propoxycarbazone	Olympus	Bayer CropScience
Propazine	MiloPro	Griffin
Prosulfocarb	Boxer (in France)	Syngenta
Pyrasulfotole	Huskie	Bayer CropScience
Pyrazon	Pyramin	BASF
Pyrazosulfuron-ethyl*	Sirius	Bayer CropScience
Pyridate	Tough	Syngenta
Pyrithiobac	Staple	DuPont
Pyroxsulam	Powerflex	Dow AgroSciences
Quinclorac	Drive, Facet, Paramount	BASF
Quinmerac*	Fiesta	BASF
Quizalofop	Assure	DuPont

Common or Code Name*	Trade Name	Manufacturer
Rimsulfuron	Matrix, Basis, TranXit	DuPont
Saflufenacil	Kixor	BASF
Sethoxydim	Poast Poast Plus	BASF
Siduron	Tupersan	DuPont
Simazine	Princep, others	Syngenta, others
Sulcotrione	Mikado, Galleon	Bayer CropScience
Sulfentrazone	Dismiss, Spartan, Authority	FMC
Sulfometuron	Oust	DuPont
Sulfosulfuron	Certainty, Monitor, Outrider, Maverick	Monsanto
Tebuthiuron	Spike	Dow AgroSciences
Tembotrione	Laudis	Bayer CropScience
Tepaloxydim	Equinox	BASF
Terbacil	Sinbar	DuPont
Terbutryn	Igran	Syngenta
Thiazopyr	Visor	Dow AgroSciences
Thifensulfuron	Pinnacle, Harmony	DuPont
Thiencarbazone		Bayer CropScience
Thiobencarb	Bolero	Valent
Topramezone	Impact	Amvac
Tralkoxydim	Achieve	Syngenta
Triallate	FarGo	Monsanto
Triasulfuron	Amber, Fuego	Syngenta
Tribenuron	Express	DuPont
Triclopyr	Renovate Garlon,	SeaPro Dow AgroSciences
Trifloxysulfuron	Envoke, Monument	Syngenta
Triflusulfuron	Upbeet	DuPont
Vernolate	Vernam	Drexel

It is to be understood that additional herbicides within the scope of the present disclosure may be identified by sources readily available to those of skill in the art.

Pests within the scope of the present disclosure associated with corn include the following.

Scientific Name	Common Name
Autographa gamma	Silver Y moth
Chilo suppressalis	Asiatic rice borer
Diabrotica speciosa	Cucurbit beetle
Harpophora maydis	Late wilt of corn
Helicoverpa armigera	Old world bollworm
Heteronychus arator	Black maize beetle
Peronosclerospora maydis	Java downy mildew

Peronosclerospora philippinensis	Philippine downy mildew
Punctodera chalcoensis	Mexican corn cyst nematode
Sclerophthora rayssiae var. *zeae*	Brown stripe downy mildew
Spodoptera littoralis	Egyptian cottonworm
Spodoptera litura	Cotton cutworm
Thaumatotibia leucotreta	False codling moth

Pests within the scope of the present disclosure associated with cotton include the following.

Scientific Name	Common Name
Anthonomus grandis	Boll weevil
Autographa gamma	Silver Y moth
Eutetranychus orientalis	Citrus brown mite
Helicoverpa armigera	Old World bollworm
Oxycarenus hyalinipennis	Cotton seed bug
Pectinophora gossypiella	Pink bollworm
Spodoptera littoralis	Egyptian cottonworm
Spodoptera litura	Cotton cutworm
Thaumatotibia leucotreta	False codling moth

Pests within the scope of the present disclosure associated with oak include the following.

Scientific Name	Common Name
Adoxophyes orana	Summer fruit tortrix moth
Aeolesthes sarta	City longhorned beetle
Agrilus biguttatus	Oak splendour beetle
Archips xylosteanus	Variegated golden tortrix
Epiphyas postvittana	Light brown apple moth
Lymantria dispar asiatica	Asian gypsy moth
Lymantria mathura	Rosy moth
Massicus raddei	Mountain oak longhorned beetle
Phytophthora quercina	Oak decline
Platypus quercivorus	Oak ambrosia beetle

Raffaelea quercivora	Japanese oak wilt
Scolytus intricatus	European oak bark beetle
Spodoptera littoralis	Egyptian cottonworm
Thaumatotibia leucotreta	False codling moth
Thaumetopoea processionea	Oak processionary moth
Tortrix viridana	Green oak tortrix
Tremex fuscicornis	Tremex woodwasp

Pests within the scope of the present disclosure associated with pine include the following.

Scientific Name	Common Name
Candidatus Phytoplasma pini 16SrXXI-A	Pine witches' broom
Cronartium flaccidum	Scots pine blister rust
Dendroctonus micans	Great spruce bark beetle
Dendrolimus pini	Pine-tree lappet
Dendrolimus punctatus	Masson pine moth
Dendrolimus sibiricus	Siberian silk moth
Diprion pini	Pine sawfly
Hylobius abietis	Large pine weevil
Lymantria mathura	Rosy moth
Monochamus saltuarius	Japanese pine sawyer
Monochamus sutor	Small white-marmorated longhorned beetle
Mycosphaerella gibsonii	Needle blight of pine
Panolis flammea	Pine beauty moth
Tomicus destruens	No common name, a pine shoot beetle

Pests within the scope of the present disclosure associated with small grains include the following.

Scientific Name	Common Name
Autographa gamma	Silver Y moth
Cernuella virgata	Maritime garden snail
Cochlicella spp.	Exotic species
Diabrotica speciosa	Cucurbit beetle

Helicoverpa armigera	Old world bollworm
Heterodera filipjevi	Cereal cyst nematode
Heterodera latipons	Mediterranean cereal cyst nematode
Heteronychus arator	Black maize beetle
Lobesia botrana	European grape vine moth
Meloidogyne artiellia	British root-knot nematode
Nysius huttoni	Wheat bug
Peronosclerospora philippinensis	Philippine downy mildew
Spodoptera littoralis	Egyptian cottonworm
Spodoptera litura	Cotton cutworm

Pests within the scope of the present disclosure associated with soybean include the following.

Scientific Name	Common Name
Adoxophyes orana	Summer fruit tortrix moth
Alectra vogelii	Yellow witchweed
Autographa gamma	Silver Y moth
Cernuella virgata	Maritime garden snail
Chrysodeixis chalcites	Golden twin spot moth
Crocidosema aporema	Bud borer
Diabrotica speciosa	Cucurbit beetle
Eutetranychus orientalis	Citrus brown mite
Helicoverpa armigera	Old world bollworm
Spodoptera littoralis	Egyptian cottonworm

Pests within the scope of the present disclosure associated with grape include the following.

Scientific Name	Common Name
Adoxophyes orana	Summer fruit tortrix moth
Candidatus Phytoplasma austra	licnse

Cryptoblabes gnidiella		Honeydew moth
Epiphyas postvittana		
Eupoecilia ambiguella		Light brown apple moth
Candidatus Phytoplasma vitis 1		
Heteronychus arator Lobesia		European grape berry moth
botrana Pseudopezicula		
tracheiphila Spodoptera	6SrV-C	Flavescence dorée Black
littoralis Spodoptera litura		maize beetle European grape
Thaumatotibia leucotreta		vine moth Rotbrenner
		Egyptian cottonworm

Pests within the scope of the present disclosure associated with palm include the following.

Scientific Name	Common Name
Bursaphelenchus cocophilus	Red ring nematode
Candidatus Phytoplasma palmae	
16SrIV	Palm lethal yellowing
Cocadviroid Coconut cadang cadang	
Darna pallivitta	Nettle caterpillar
Haplaxius crudus	American palm cixiid
Metamasius hemipterus	West Indian cane weevil
Oryctes rhinoceros	Coconut rhinoceros beetle
Paysandisia archon	No common name, a palm borer
Raoiella indica	Red palm mite
Rhabdoscelus obscurus	New Guinea sugarcane weevil
Rhynchophorus ferrugineus	Red palm weevil
Rhynchophorus palmarum	South American palm weevil

Pests within the scope of the present disclosure associated with solanaceous plants include the following.

Scientific Name	Common Name
Autographa gamma	Silver-Y moth
Candidatus Phytoplasma australiense	
16SrXII-B	Australian grapevine yellows

Chrysodeixis chalcites	Golden twin spot moth
Globodera pallida	Pale cyst nematode
Globodera rostochiensis	Golden nematode
Helicoverpa armigera	Old world bollworm
Meloidogyne fallax	False Columbia root-knot nematode
Meloidogyne minor	Root-knot nematode
Neoleucinodes elegantalis	Tomato fruit borer
Ralstonia solanacearum race 3 biovar 2	Bacterial wilt/Southern bacterial Wilt
Spodoptera littoralis	Egyptian cottonworm
Spodoptera litura	Cotton cutworm
Synchytrium endobioticum	Potato wart
Tecia solanivora	Guatemalan potato tuber moth
Thaumatotibia leucotreta	False codling moth
Tuta absoluta	Tomato leaf miner

Pests within the scope of the present disclosure associated with stone fruit include the following.

Scientific Name	Common Name
Adoxophyes orana	Summer fruit tortrix
Argyresthia pruniella	Cherry blossom moth
Bactrocera zonata	Peach fruit fly
Candidatus Phytoplasma prunorum 16SrX-F	European stone fruit yellows
Enarmonia formosana	Cherry bark tortrix
Epiphyas postvittana	Light brown apple moth
Grapholita funebrana (Syn.: *Cydia funebrana*)	Plum fruit moth
Leucoptera malifoliella	Pear leaf blister moth
Lobesia botrana	European grape vine moth
Monilia polystroma	Asiatic brown rot
Monilinia fructigena	Brown rot, Apple brown rot
Potyvirus Plum Pox Virus	Plum pox
Rhagoletis cerasi	European cherry fruit fly
Thaumatotibia leucotreta	False codling moth

Additional agricultural pests include the following cyst nematodes.

Scientific Name	Common Name
Globodera pallida	Pale cyst nematode
Globodera rostochiensis	Golden nematode
Heterodera cajani	Pigeonpea cyst nematode
Heterodera ciceri	Chickpea cyst nematode
Heterodera filipjevi	Cereal cyst nematode
Heterodera latipons	Mediterranean cereal cyst nematode
Heterodera sacchari	Sugarcane cyst nematode
Punctodera chalcoensis	Mexican corn cyst nematode

Additional agricultural pests include the following exotic wood borer or bark beetles.

Scientific Name	Common Name
Agrilus auroguttatus	Goldspotted oak borer
Agrilus biguttatus	Oak splendour beetle
Agrilus planipennis	Emerald ash borer
Anoplophora chinensis	Citrus longhorned beetle
Anoplophora glabripennis	Asian longhorned beetle
Chlorophorus annularis	Bamboo borer
Chlorophorus strobilicola	Slender-banded pine cone longhorn beetle
Dendroctonus micans	Great spruce bark beetle
Ips sexdentatus	Six-toothed bark beetle
Ips typographus	European spruce bark beetle
Megaplatypus mutatus	No common name, an ambrosia beetle
Monochamus alternatus	Japanese pine sawyer
Monochamus saltuarius	Japanese pine sawyer
Monochamus sutor	Small white-marmorated longhorned beetle
Orthotomicus erosus	Mediterraneran pine engraver
Pityogenes chalcographus	Sixtoothed spruce bark beetle
Platypus quercivorus	Oak ambrosia beetle
Scolytus intricatus	European oak bark beetle

Tetropium castaneum	Black spruce beetle
Tetropium fuscum	Brown spruce longhorned beetle
Tomicus destruens	No common name, a pine shoot beetle
Tomicus minor	Lesser pine shoot beetle

Scientific Name	Common Name
Tomicus piniperda	Pine shoot beetle
Trichoferus campestris	
Trypodendron domesticum	Velvet longhorned beetle
Xyleborus glabratus	

Additional agricultural pests include the following mollusks.

Scientific Name	Common Name
Belocaulus spp.	No common name, leatherleaf slugs
Cernuella spp.	No common name, hygromiid snails
Cochlicella spp.	No common name, cochlicellid snails
Colosius spp.	No common name, leatherleaf slugs
Laevicaulis spp.	No common name, leatherleaf slugs
Lissachatina fulica	Giant African snail
Meghimatium pictum	Chinese slug
Monacha spp.	No common name, hygromiid snails
Sarasinula spp.	No common name, leatherleaf slugs
Semperula spp.	No common name, leatherleaf slugs
Veronicella spp.	No common name, leatherleaf slugs

Additional agricultural pests include the following moths.

Scientific Name	Common Name
Dendrolimus pini	Pine-tree lappet
Dendrolimus punctatus	Masson pine moth
Dendrolimus sibiricus	Siberian silk moth
Lymantria albescens	Okinawa gypsy moth
Lymantria dispar asiatica	Asian gypsy moth

Lymantria dispar japonica	Japanese gypsy moth
Lymantria mathura	Rosy moth
Lymantria monacha	Nun moth
Lymantria postalba	White-winged gypsy moth
Lymantria umbrosa	Hokkaido gypsy moth
Lymantria xylina	Casuarina tussock moth

It is to be understood that additional pests within the scope of the present disclosure may be identified by sources readily available to those of skill in the art.

Common pesticides in general within the scope of the present disclosure to which a pest may be resistant or develop resistance include the following: algicides, antifouling agents, antimicrobials, attractants, biopesticides, biocides, disinfectants, fungicides, fumigants, insecticides, miticides, microbial pesticides, molluscicides, nematicides, pheromones, repellants and rodenticides.

The following pesticide species are useful within the scope of the present disclosure: Glyphosate, Atrazine, Metam Sodium, Metolachlor-S, Acetochlor, Dichloropropene, 2,4-D, Methyl Bromide, Chloropicrin, Pendimenthalin, Ethephon, Chlorothalonil, Metam Potassium, Chlorpyrifos, Copper Hydroxide, Copper Sulfate, Simazine, Trifluralin, Propanil, Mancozeb, Aldicarb, Acephate, Diuron, MCPA, Paraquat, Dimethenamid, Carbaryl, MCPP, MSMA, Pyrethroids, Malathion, Dicamba, Pelarganoc Acid, Sulfuryl fluoride, Triclopyr, Paradiclorobenzene, Naphthalene, Chlorpyrifos, Naled, Dicrotophos, Phosmet, Phorate, Diazinon, Dimethoate, Azinphos-Methyl, and N,N-diethyl-meta-toluamide (insect repellant). One of skill in the art will readily be able to identify additional pesticide species using publicly available information of databases, for example the EPA list of registered pesticides available at world wide website iaspub.epa.gov/apex/pesticides/f?p=chemicalsearch:1 and with reference to the most commonly used conventional pesticide active ingredients in the U.S. Agricultural, Home and Garden, Industry, Commercial, and Government Market Sectors as compiled by the EPA and publicly available at world wide website www.epa.gov/opp00001/pestsales/.

One of skill in the art will readily be able to identify pesticides toxic to particular pest species based on the present disclosure.

Accordingly, aspect of the present disclosure are directed to a method of altering a eukaryotic germline cell of an organism including introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, and including a sensitizing nucleic acid the expression of which is

harmful to the organism when the organism is exposed to a chemical, compound or condition, wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell, wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence, wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA, wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA, expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner, inserting the first foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the first foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the foreign nucleic acid sequence, and expressing the sensitizing nucleic acid rendering the resulting organism sensitive to the chemical, compound or condition such that the resulting organism perishes or is rendered sterile when exposed to the chemical, compound or condition.

According to one aspect, expression of the sensitizing nucleic acid increases toxicity of the chemical, compound or condition to the organism. According to one aspect, the germline cell is grown into an organism and the sensitizing nucleic acid is transferred to progeny to create a population of organisms including the sensitizing nucleic acid and where the sensitizing nucleic acid increases toxicity of the chemical, compound or condition to the organism. According to one aspect, the organism is a weed or pest. According to one aspect, the sensitizing nucleic acid is a sensitizing gene which replaces an existing gene. According to one aspect, the sensitizing gene is the exact or codon-altered ancestral version of an existing mutant gene in wild populations, such that the current mutated version is replaced with the ancestral form. According to one aspect, the existing gene has acquired a mutation contributing to resistance to a pesticide, herbicide, or fungicide. According to one aspect, the chemical or compound is a pesticide, herbicide or fungicide. According to one aspect, the pesticide, herbicide, or fungicide is one of: a Bt toxin produced by Cry1A.105, Cry1Ab, Cry1F, Cry2Ab, Cry3Bb1, Cry34Ab1, Cry35Ab1, mCry3A, or

VIP, 2,4-D or glyphosate. According to one aspect, the sensitizing gene replaces an existing gene whose function is required for the organism to survive or reproduce. According to one aspect, the chemical is a prodrug and the sensitizing gene encodes a corresponding prodrug-converting enzyme. According to one aspect, the enzyme/chemical pairing is cytosine deaminase/5-fluorocytosine, or nitroreductase/CB1954.

According to certain aspect of the present disclosure, a method of controlling a weed or pest population including a sensitivity gene drive in the genome of the weed or pest population wherein the sensitivity gene drive renders the weed or pest population vulnerable to toxicity when in the presence of a chemical, compound or condition is provided including contacting the weed or pest population with the chemical, compound or condition in an effective amount to kill the weed or pest, lower proliferation of the weed or pest or render the weed or pest sterile to inhibit proliferation. According to one aspect, the chemical or compound is an herbicide, or pesticide or fungicide.

According to certain aspect of the present disclosure, a method of altering a eukaryotic germline cell of an organism including a first sensitizing gene drive is provided including introducing into the germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, and including a second sensitizing nucleic acid sequence the expression of which is harmful to the organism when the organism is exposed to a chemical, compound or condition, wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome including the first sensitizing gene drive and a second chromosome of a chromosome pair of the germline cell including the first sensitizing gene drive, wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs of the second foreign nucleic acid sequence are between the first flanking sequence and the second flanking sequence, wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA, wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA, expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner,

wherein the first foreign nucleic acid sequence is removed, inserting the second foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the second foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the second foreign nucleic acid sequence, and expressing the second sensitizing nucleic acid sequence rendering the resulting organism sensitive to the chemical, compound or condition such that the resulting organism perishes or is rendered sterile when exposed to the chemical, compound or condition. According to one aspect, the resulting organism is introduced into a wild type population such as that progeny of the resulting organism and a wild type organism includes the second sensitizing nucleic acid sequence. According to one aspect, the organism is a weed or pest.

According to certain aspect of the present disclosure, a method of altering a eukaryotic germline cell of an organism including a first sensitizing gene drive is provided including introducing into the germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, and including a second sensitizing nucleic acid sequence the expression of which is harmful to the organism when the organism is exposed to a chemical, compound or condition, wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome including the first sensitizing gene drive and a second chromosome of a chromosome pair of the germline cell including the first sensitizing gene drive, wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs of the second foreign nucleic acid sequence are between the first flanking sequence and the second flanking sequence, wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA, wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA, expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner, inserting the second foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the second foreign nucleic acid

sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the second foreign nucleic acid sequence, and expressing the second sensitizing nucleic acid sequence rendering the resulting organism sensitive to the chemical, compound or condition such that the resulting organism perishes or is rendered sterile when exposed to the chemical, compound or condition. According to certain aspects, the resulting organism is introduced into a wild type population such as that progeny of the resulting organism and a wild type organism includes the second sensitizing nucleic acid sequence. According to certain aspects, the organism is a weed or pest.

EXAMPLE I

Plasmids and Genomic Cassettes

Gene drive cassettes were synthesized from gBlocks (Integrated DNA Technologies, Coralville, IA) and inserted into SK1 cells via Cas9-mediated genome modification as follows. Guide RNAs for each drive were cloned into p416-Cas9 containing plasmids with expression driven by the SNR52 promoter. See DiCarlo, J. E. *et al.* Genome engineering in Saccharomyces cerevisiae using CRISPR-Cas systems, *Nucleic Acids Res.* 41, 4336–4343 (2013). 60 base pair homology arms to the target locus were added on both ends of the gene drive cassette via PCR and 5 ug of PCR product was co-transformed with the p416-Cas9-gRNA plasmids. Correctly integrated gene drives were verified by sequencing and p416-Cas9-gRNA plasmids were cured using 5-Fluoroorotic Acid (FOA) selection.

To create the *URA3*-containing *ADE2* gene drive, the *ADE2* gene drive was cloned next to the *Candida albicans URA3* gene in the pAG60 plasmid. The entire URA3 cassette and gene drive were PCR amplified and inserted using Cas9-mediated genome modification into the *ADE2* locus of haploid SK1 cells.

The recoded C-terminus of the *ABD1* gene and corresponding gene drive were synthesized as a gBlock to remove homology and generate mutations in the seed sequence via synonymous changes. The TEF1 terminator was inserted at the 3'end of the recoded *ABD1* gene between the gene and the gRNA as *ABD1* shares a terminator with the *VHC1* gene. The entire cassette was integrated into the haploid SK1 genome using Cas9-mediated genome modification.

The p416-Cas9-gRNA plasmid (conferring uracil prototrophy) is a variant of the previously described p414-Cas9-gRNA plasmid (conferring tryptophan prototrophy) (see DiCarlo, J. E. *et al.* Genome engineering in Saccharomyces cerevisiae using CRISPR-Cas systems, *Nucleic Acids Res.* 41, 4336–4343 (2013)) (Addgene #43802). One or the other was used in each mating experiment. The pRS413 vector was transformed into select cell types to confer histidine prototrophy as a marker to select for diploid cells. Strain genotypes are provided in Table 1 below.

Table 1

Strain	Genotype
SK1 A	MATa ho::LYS2 lys2 ura3 leu2::hisG his3::hisG trp1::hisG
SK1 α	MATα ho::LYS2 lys2 ura3 leu2::hisG his3::hisG trp1::hisG
Y12A	MATa ho::HygMX ura3::KanMX
YPS128	MATa ho::HygMX ura3::KanMX
YJM981	MATa ho::HygMX ura3::KanMX
Y55	MATa ho::HygMX ura3::KanMX
UWOPS05-217.3	MATa ho::HygMX ura3::KanMX
DBVPG 6044	MATa ho::HygMX ura3::KanMX
273614N	MATa ho::HygMX ura3::KanMX

EXAMPLE II

Yeast Mating Experiments

Haploid drive-containing SK1 yeast and haploid wild-type strains of the opposite mating type were mixed in equal amounts in YPAD liquid media and incubated overnight. The resulting diploids were washed in sterile water and plated on selective media for both parental genotypes. Table 2 below details the specific crosses.

Table 2

MATa Genotype	MATα Genotype	Selection
SK1 pRS414 – Cas9	SK1 ade2::gRNA (gene drive), pRS413	SC-histidine - tryptophan
SK1 ade2::gRNA + URA3 (gene drive), p414-Cas9	SK1 pRS413	SC-histidine - tryptophan
SK1 p414-Cas9	SK1 abd1::ABD1 recoded +gRNA (gene drive), pRS413	SC-histidine - tryptophan
Y12A Hygromycin B resistance (HygR)	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B
YPS128 Hygromycin B resistance (HygR)	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B
YJM981 Hygromycin B resistance (HygR)	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B
Y55 Hygromycin B resistance (HygR)	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B
UWOPS05-217.3 Hygromycin B	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B

resistance (HygR)		
DBVPG 6044 Hygromycin B resistance (HygR)	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B
273614N Hygromycin B resistance (HygR)	SK1 ade2::gRNA (gene drive), p416-Cas9	SC-uracil+300 ug/mL Hygromycin B
SK1 ADE2:: ADE2 silently recoded genomic target seed sequence, p414-Cas9	SK1 ade2::gRNA (gene drive), pRS413	SC-histidine - tryptophan
SK1, p414-empty	SK1 ade2::gRNA (gene drive), pRS413	SC-histidine - tryptophan

EXAMPLE III

Sporulation and Tetrad Dissection

After mating in liquid YPAD and selection for diploids on selection plates, the selection plates were scraped into 10 mL selective media and grown overnight at 30°C. A fresh 5 mL YPAD culture was then inoculated to and OD=0.1 and grown 4-5 hours at 30°C. The entire culture was then washed twice in 10 mL water, inoculated into 2 mL of sporulation media(1% potassium acetate), and incubated at room-temperature for 3 days or until spores were visible. Sporulated cells were suspended in 50 µL of a stock solution of zymolyase (50 µg/mL in 1M sorbitol) and incubated at 30C for 5 minutes, transferred to ice, diluted with 150 µL cold H_2O, microdissected using a Zeiss tetrad dissection microscope, and isolated spores grown on YPAD plates.

EXAMPLE IV

Selection for URA3 Function

Dissected spores were grown in synthetic complete (SC) media and then spotted onto SC medium as well as SC medium without uracil. To enhance red color, all SC solid media used for plate images contained 0.5 X adenine hemisulfate (final concentration of 0.08 mM).

EXAMPLE V

Quantitative PCR

Candidate primer pairs were designed to amplify short regions specific to each drive or the wild-type sequence replaced by the drive, as well as the ACT1 gene as a control. All sequences are

included in the supplementary information. Genomic DNA was extracted using Method A as described in Looke et al.31

KAPA SYBR FAST qPCR Master Mix (2X) was used to perform the qPCR reaction along with 25 ng of genomic DNA. The amplification efficiency and relative specificity of each primer pair were measured by amplifying dilutions of genomic DNA from wild-type and drive haploids, respectively, and the best-performing and well-matched pairs selected for use (see below for all primers used). Quantitative PCR reactions were performed on genomic DNA isolated from each parental haploid as well as from diploids arising from three independent mating events. Three reactions (technical replicates) were performed per sample on a LightCycler 96 machine by Roche.

EXAMPLE VI

Calculations

Results from three technical replicates were averaged for calculations. In order to directly calculate the ratio of alleles before PCR amplification, the efficiencies of the different primer pairs were first determined. Efficiencies were calculated from qPCR runs of serial dilutions (6 orders of magnitude) as:

$\text{Efficiency}=10^{-1/\text{slope}}$

R^2 values were higher than 0.99 in all cases except for one pair (adc2::URA3+sgRNA).

The allelic ratios were calculated as:

$x_a \cdot E_a^{Ct,a} = x_b \cdot E_b^{Ct,b}$

$x_a / x_b = E_b^{Ct,b} / E_a^{Ct,a}$

with x_a and x_b being the initial concentration of drive and wt DNA,

E_a and E_b the efficiency of the respective primer pairs and

Ct,a and Ct,b the Ct values for each sample.

EXAMPLE VII

Efficiency of CRISPR/Cas9 Gene Drives in Yeast

To directly measure the efficiency of CRISPR/Cas9 gene drives in yeast, a system was developed using the red color that builds up in yeast lacking functional copies of the ADE2 gene. See Chamberlain, N., Cutts, N. S. & Rainbow, C. The formation of pigment and arylamine by yeasts. *J. Gen. Microbiol.* 7, 54–60 (1952). As depicted in Fig. 13A, if red $ADE2^-$ haploids are mated with cream-colored wild-type haploids, the resulting heterozygous diploids inherit one functional copy and are therefore cream-colored. When these diploids undergo meiosis and reproduce via sporulation, half of the resulting haploids inherit the broken copy and are consequently red; the other half inherit the intact copy and are cream-colored (Fig. 13A).

As depicted in Fig. 13B, if the red haploids encode a functional gene drive that cuts and

replaces the intact ADE2 locus inherited from the wild-type parent, their diploid progeny will be red. Because these red diploids will have two broken copies of ADE2, all of their sporulated haploid offspring will inherit a broken copy and consequently will also be red. Thus, the cutting efficiency of a gene drive that targets and replaces ADE2 can be assessed by simply plating diploids and counting the fraction that are red.

A gene drive construct targeting ADE2 was made. To prevent accidental escape of the gene drive into the wild, Cas9 and guide RNAs were separated to avoid creating a self-sufficient inheritance-biasing cassette. Consequently, constructs encoded a guide RNA targeting ADE2, while Cas9 was provided from an episomal plasmid. Red haploids were mated to wild-type yeast of the opposite mating type in the presence or absence of the plasmid and plated on media that selects for diploids. As shown in Fig. 13C, nearly all colonies were red in the presence of the plasmid, indicating highly efficient cutting of the ADE2 copy inherited from the wild-type parent. In the absence of Cas9, no red diploid colonies were observed, demonstrating that the drive can only spread in laboratory yeast populations that provide it.

To verify that the ADE2 alleles from wild-type parents were lost, the mated diploids were sporulated and their resultant haploid progeny were examined. As shown in Fig, 13D, upon dissecting 18 cas9+ diploids, a perfect 4:0 ratio of red:cream haploids was observed, confirming that all copies of the ADE2 locus were disrupted. In contrast, 18 cream-colored cas9— diploids yielded a 2:2 red:cream ratio, indicating normal inheritance of the inactivated drive and the wild-type alleles.

To determine whether the ADE2 disruptions in red diploids were the result of successful copying of the drive element by homologous recombination, 72 haploids derived from dissected cas9+ diploids were sequenced. All sequenced colonies contained intact drives without additional mutations, indicating that drive mobilization was efficient and occurred at high fidelity.

An ADE2 gene drive was modified as shown in schematic in Fig. 14A to contain an *in cis* URA3 allele, which allows laboratory modified yeast strains to grow in the absence of uracil supplementation. This drive element was tested for the ability of gene drives to "carry" an associated cargo element when copied into a target locus. URA3-containing drive haploids were mated with wild-type haploids in the presence of an episomal Cas9 plasmid, diploids were selected (all of which were red) and sporulated, and 18 tetrads were dissected. As was the case for the original ADE2 gene drive, all of the sporulated haploid cells formed red colonies. As shown in Fig. 14B, all grew normally when replica plated on uracil deficient media, indicating that the URA3 cargo element was efficiently copied with the drive.

Gene drives which target and recode an essential gene could avoid drive resistance in even large populations since error prone repair events that modify the target site will cause lethality. Non-essential but nonetheless important genes could similarly be edited because mutants created

by NHEJ events would still be less fit than the drive itself. To test essential gene recoding during drive insertion, a third gene drive targeting ABD1 shown in schematic in Fig. 14C was constructed. See Mao, X., Schwer, B. & Shuman, S. Mutational analysis of the Saccharomyces cerevisiae ABD1 gene: cap methyltransferase activity is essential for cell growth, *Mol. Cell. Biol.* 16, 475–480 (1996).

A haploid strain containing a recoded ABD1 allele upstream of a guide RNA targeting the natural ABD1 coding sequence was mated to wild-type cells in the presence of Cas9. Diploid cells were selected of which 18 were sporulated 18 of them. 72 segregants were sequenced. All contained the recoded ABD1 locus and the guide RNAs, thereby demonstrating gene drives based on essential gene recoding.

Gene drives were copied from a laboratory strain into a diverse group of native *S. cerevisiae* strains. *ADE2* drive-containing haploids were mated with 6 phylogenetically and phenotypically diverse wild-type strains of haploid *S. cerevisiae*. See Liti, G. *et al.* Population genomics of domestic and wild yeasts, *Nature* 458, 337–341 (2009). See also Fig. 15A. To quantitatively measure the efficiency of gene drive copying for each cross, quantitative PCR was performed on populations of all diploids using one set of primers specific to the drive and another set designed to amplify either wild-type or NHEJ-disrupted alleles. As shown In Fig. 15B which was generated using BoxPlot (see Spitzer, M., Wildenhain, J., Rappsilber, J. & Tyers, M. BoxPlotR: a web tool for generation of box plots. *Nat. Methods* 11, 121–122 (2014)), the fraction of diploid chromosomes containing the *ADE2* gene drive was over 99% regardless of wild-type parent, demonstrating the use of the drive in diverse backgrounds. Addition of the *URA3* cargo gene did not appreciably change this efficiency. The *ABD1* drive was copied at an equivalent rate.

The stability of the drive over successive copying events was investigated. As shown in Fig. 15B, several haploid offspring of the first-round *ADE2* gene drive diploids were mated to wild-type haploids containing the Cas9-expressing plasmid. All of the second-generation gene drive constructs biased inheritance at the same efficiency, demonstrating a continued ability to spread through sexually reproducing populations over generations.

The following genome modification primer and gBlock sequences are provided.

Genome modification primers and gBlocks	Sequence
ADE2.sgRNA.ade2.1 insert.F	TACGAACCGGGTAATACTAAGTGATTGACTCTTGCTGACCTTTTAT TAAGAACTAAATGGtctttgaaaagataatgtatgattatgctttc
ADE2.sgRNA.ade2.1 insert.R	TAATAAGTGATCTTATGTATGAAATTCTTAAAAAAGGACACCTGTA AGCGTTGATTTCTAagacataaaaaacaaaaaaagcaccac

gRNA+CaURA3.ade2.F	TACGAACCGGGTAATACTAAGTGATTGACTCTTGCTGACCTTTTAT TAAGAACTAAATGGagacataaaaaacaaaaaaagcaccaccg
gRNA+CaURA3.ade2.R	TAATAAGTGATCTTATGTATGAAATTCTTAAAAAAGGACACCTGTA AGCGTTGATTTCTAtcgacactggatggcggcgttagtatc
ABD1.recode+gRNA	AGCCAGATGCCATTCAACAAGTTCTTCGTGCAGGAGATACCAAA GTGGATAGAACGTTTCAGCCCAAAGATGCGTGAGGGGCTTCAGC GTAGCGACGGGCGTTACGGGGTGGAGGGTGACGAGAAAGAGGC TGCTAGCTACTTTTACACGATGTTCGCTTTTAGAAAAGTTAAGCA ATACATAGAGCCTGAGTCAGTTAAACCAAATTGAACGGCTCCTCG CTGCAGACCTGCGAGCAGGGAAACGCTCCCCTCACAGACGCGTT GAATTGTCCCCACGCCGCGCCCCTGTAGAGAAATATAAAAGGTTA GGATTTGCCACTGAGGTTCTTCTTTCATATACTTCCTTTTAAAATC TTGCTAGGATACAGTTCTCACATCACATCCGAACATAAACAACCA TGGGTATGACCGACCAAGCGACGCCCAACCTGCCATCACGAGAT TTCGATCCCACCGCCGCCTTCTATGAAAGGtctttgaaaagataatgtatgattat gctttcactcatatttatacagaaacttgatgttttctttcgagtatatacaaggtgattacatgtacgtttgaagta caactctagattttgtagtgccctcttgggctagcggtaaaggtgcgcatttttttcacaccctacaatgttctgtt caaaagattttggtcaaacgctgtagaagtgaaagttggtgcgcatgtttcggcgttcgaaacttctccgcag tgaaagataaatgatcTGAAGGGGATGAAAAGGAAGGTTTTAGAGCTAGA AATAGCAAGTTAAAATAAGGCTAGTCCGTTATCAACTTGAAAAAG TGGCACCGAGTCGGTGGTGCTTTTTTTGTTTTTTATGTCT
ABD1.recode+gRNA.int.F	tatgttgtgccattcgaaaccttaagaagtttggctgatgaatacggtttggaactagtaAGCCAGATG CCATTCAACAAGTTC
ABD1.recode+gRNA.int.R	gtaatacggccgaaatacagatgctttatagtagggttattgtttctattcatttttattAGACATAAAA AACAAAAAAAGCACCACC
ADE2.gRNA	tctttgaaaagataatgtatgattatgctttcactcatatttatacagaaacttgatgttttctttcgagtatatacaa ggtgattacatgtacgtttgaagtacaactctagattttgtagtgccctcttgggctagcggtaaaggtgcgca ttttttcacaccctacaatgttctgttcaaaagattttggtcaaacgctgtagaagtgaaagttggtgcgcatgtt tcggcgttcgaaacttctccgcagtgaaagataaatgatcACTTGAAGATTCTTTAGTGT GTTTTAGAGCTAGAAATAGCAAGTTAAAATAAGGCTAGTCCGTTA TCAACTTGAAAAAGTGGCACCGAGTCGGTGGTGCTTTTTTTGTTT TTTATGTCT
ADE2.1.silent.seed.90mer.F	TGATGTGCTAACGATTGAGATTGAGCATGTTGATGTTCCTACCCTG AAAAACCTGCAAGTAAAACATCCCAAATTAAAAATTTACCCTTC
ADE2.1.silent.seed.90mer.	GAAGGGTAAATTTTTAATTTGGGATGTTTTACTTGCAGGTTTTTCA

F	GGGTAGGAACATCAACATGCTCAATCTCAATCGTTAGCACATCA
ABD1.ver.F	ATAGATAATGTTCCTGAATATGTTGTGCCA
ABD1.ver.R	TTACTACATATAGAAGTCTTGTAATACGGCCG
ADE2.ver.F	GCTACGAACCGGGTAATACTAAGTGATTG
ADE2.ver.R	CAGGTAATTATTCCTTGCTTCTTGTTACTGG
qPCR Primers	Sequence
ade2.WT.qPCR.F	TACGAACCGGGTAATACTAAGTGATTGACTC
ade2.gRNA.qPCR.R	CGCTAGCCCAAGAGGGCACTACA
ade2.WT.qPCR.R	TACCAACTGTTCTAGAATCCATACTTGATTGTTT
URA3.genedrive.ade2.WT. qPCR.F	TACGAACCGGGTAATACTAAGTGATTGACTC
URA3.genedrive.ade2.WT. qPCR.R	CCTCCTAATATACCAACTGTTCTAGAATCCAT
URA3.genedrive. ade2.gRNA.qPCR.R	AAACTTCTCCGCAGTGAAAGATAAATGATC
ABD1_rec_qPCR.R	CGAGGAGCCGTTCAATTTGGTTTAACTGAC
ABD1_rec_qPCR.F	AGATGCGTGAGGGGCTTCAGC
ABD1_WT_qPCR (JDwt1.4).F	GAAGGGGATGAAAAGGAAGC
ABD1_WT_qPCR (JDwt1.3).R	CGCTTTCCGGTTCGATATAC
ACT1.qPCR.F	CGAAAGATTCAGAGCCCCAGAAGCT
ACT1.qPCR.R	CGGTGATTTCCTTTTGCATTCTTTCG

Claims:

1. A method of altering a eukaryotic germline cell of an organism comprising

introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner,

inserting the first foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the first foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the foreign nucleic acid sequence.

2. The method of claim 1 wherein the RNA guided DNA binding protein nuclease creates a double stranded break.

3. The method of claim 1 wherein the RNA guided DNA binding protein nuclease is Cas9.

4. The method of claim 1 wherein the RNA guided DNA binding protein nuclease creates a single stranded break.

5. The method of claim 1 wherein the RNA guided DNA binding protein nuclease is Cas9 nickase.

6. The method of claim 1 wherein at least one target location is within an essential gene and the foreign nucleic acid sequence is inserted adjacent to and replacing part of the essential gene.

7. The method of claim 1 wherein the first foreign nucleic acid is inserted by homologous recombination.

8. The method of claim 1 wherein the germline cell is a fungal cell, a plant cell, an insect cell or a mammalian cell.

9. The method of claim 1 wherein the RNA includes between about 10 to about 250 nucleotides.

10. The method of claim 1 wherein the RNA includes between about 20 to about 100 nucleotides.

11. The method of claim 1 wherein the RNA guided DNA binding protein nuclease co-localizes with an associated guide RNA at a plurality of associated target locations to cleave either the first chromosome or the second chromosome at a plurality of cleavage sites.

12. The method of claim 11 wherein cleaving at the plurality of cleavage sites removes a DNA sequence which is replaced with the first foreign nucleic acid sequence by homologous recombination.

13. The method of claim 1 further including the step of growing the germline cell into the organism.

14. The method of claim 13 further including the step of mating the organism with a wild-type organism of the same species to produce an offspring germline cell including a first chromosome of a chromosome pair with the first foreign nucleic acid sequence.

15. The method of claim 14 wherein the first foreign nucleic acid sequence is expressed by the offspring germline cell to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on a second chromosome of the chromosome pair and the RNA guided DNA binding protein nuclease cleaves the second chromosome at the target location in a cleavage site specific manner and,

inserting the first foreign nucleic acid sequence into the second chromosome of the chromosome pair at the cleavage site to render the offspring germline cell homozygous for the foreign nucleic acid sequence.

16. The method of claim 15 wherein the steps of growing the germline cell into the organism and mating the organism with a wild type organism of the same species is repeated to produce a population of organisms of the same species that are homozygous for the first foreign nucleic acid sequence.

17. The method of claim 1 wherein the first foreign nucleic acid sequence includes one or more selected gene sequences.

18. The method of claim 16, wherein the first foreign nucleic acid sequence includes one or more selected gene sequences and the one or more selected gene sequences is within the genome of

the organisms within the population of organisms.

19. The method of claim 16 wherein an altered organism is introduced into the population of organisms and that includes within its genome a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to one or more target locations on the first foreign nucleic acid sequence and wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first foreign nucleic acid sequence and wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first foreign nucleic acid sequence,

wherein the altered organism mates with a member of the population of organisms to produce an offspring germline cell including the second foreign nucleic acid sequence and the first foreign nucleic acid sequence,

wherein the offspring germline cell expresses the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first foreign nucleic acid sequence, the RNA guided DNA binding protein nuclease cleaves the first foreign nucleic acid sequence at the target location in a cleavage site specific manner and,

wherein the second foreign nucleic acid sequence is inserted into the genome of the germline cell at the cleavage site to render the offspring germline cell homozygous for the second foreign nucleic acid sequence.

20. The method of claim 16 wherein altered wild type organisms are introduced into the population that have the one or more target locations on genomic DNA altered such that the one or more guide RNAs are noncomplementary to the one or more target locations and are prevented from binding to the one or more target locations such that the first foreign nucleic acid sequence does not enter the genome of the altered wild type organisms.

21. The method of claim 20 wherein the altered wild type organisms are altered by expression of a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs which co-localize to the first foreign nucleic acid sequence to cleave the first foreign nucleic acid sequence in a cleavage site specific manner resulting in a recoded allele.

22. The method of claim 1 further comprising introducing into the germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to the one or more target locations that were altered by the first foreign nucleic acid sequence,

expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the chromosome of the genomic DNA at the target location in a cleavage site specific manner to remove the first foreign nucleic acid sequence, and

inserting the second foreign nucleic acid sequence into the chromosome of the genomic DNA at the cleavage site.

23. The method of claim 22 wherein the second foreign nucleic acid sequence includes genetic modifications different from the first foreign nucleic acid sequence.

24. The method of claim 23 wherein the second foreign nucleic acid sequence includes the wild type sequence of the chromosome that was modified by the first foreign nucleic acid sequence.

25. The method of claim 1 wherein at least one target location is within an essential protein-coding gene and the foreign nucleic acid sequence is inserted into the essential gene such that the amino acid sequence of the essential protein is preserved but the nucleotide sequence of the gene is altered such that the one or more guide RNAs are noncomplementary to the one or more target locations

26. A method of altering a eukaryotic germline cell comprising

introducing into the germline cell a foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome of the genomic DNA,

expressing the foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein and an

associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner,

inserting the foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site,

expressing the inserted foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the genomic DNA at the target location in a cleavage site specific manner,

inserting the foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the foreign nucleic acid sequence.

27. A method of rendering two populations of a species unable to produce viable progeny comprising

altering a first population of the species to include a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more first guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, wherein the one or more first guide RNAs are complementary to one or more target locations on an essential gene, and the nucleotide sequence of the essential gene has been altered such that the one or more guide RNAs are noncomplementary to the one or more target locations but the amino acid sequence of the encoded essential protein is preserved,

altering a second population of the species to include a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more second guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, wherein the one or more second guide RNAs are complementary to one or more target locations on the essential gene that are different from the target locations of the one or more first guide RNAs, and the nucleotide sequence of the essential gene has been altered such that the one or more second guide RNAs are noncomplementary to the one or more target locations, but the amino acid sequence of the encoded essential protein is preserved,

mating the first population with the second population to produce germline cells where the first foreign nucleic acid sequence is expressed to cleave the essential gene contributed by second parent and the second foreign nucleic acid sequence are expressed to cleave the essential gene contributed by the first parent.

28. A method of replacing an allele in a eukaryotic germline cell having a mutation in a target nucleic acid sequence comprising

introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and a plurality of guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the plurality of guide RNAs are complementary to a plurality of target locations on genomic DNA of a chromosome that are adjacent to a mutation on an allele wherein the mutation prevents binding of a corresponding guide RNA to the genomic DNA,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the plurality of guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the chromosome of the genomic DNA,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the plurality of RNAs wherein the RNA guided DNA binding protein nuclease and associated guide RNAs co-localize to a plurality of associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the plurality of target locations in a cleavage site specific manner so as to remove a portion of the genomic DNA including the mutation, and

inserting the first foreign nucleic acid sequence into the chromosome of the genomic DNA at the cleavage site.

29. The method of claim 28 wherein at least one of the target locations are within a gene important to the fitness of the organism such as an essential gene, such that a deletion event that removes all target locations is highly deleterious and will be removed from the population by natural selection.

30. A method of altering a eukaryotic germline cell of an organism at a plurality of genes comprising

introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and a plurality of guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, along with one or more additional foreign nucleic acid sequences corresponding to the plurality of genes,

wherein the plurality of guide RNAs include one or more gene specific guide RNAs that are complementary to one or more target locations on a plurality of genes in the genomic DNA of a chromosome of the germline cell,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the plurality of guide RNAs are between the first

flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of one of the target locations on the chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the same target location on the chromosome of the genomic DNA,

wherein each of the one or more nucleic acid sequences contain a first flanking sequence and a second flanking sequence corresponding to a first portion and a second portion of the remaining target locations,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the plurality of RNAs wherein the RNA guided DNA binding protein nuclease and associated guide RNAs co-localize to a plurality of associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at a plurality of genes in a cleavage site specific manner, and

inserting the first foreign nucleic acid sequence into the chromosome of the genomic DNA at the cleavage site.

31. The method of claim 30 further comprising introducing into the germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and a plurality of guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the plurality of guide RNAs are complementary to the plurality of target locations that were altered by the first foreign nucleic acid sequence,

expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the plurality of guide RNAs wherein the RNA guided DNA binding protein nuclease and associated guide RNAs co-localize to associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the chromosome of the genomic DNA at the associated target locations in a cleavage site specific manner to remove the first foreign nucleic acid sequence, and

inserting the second foreign nucleic acid sequence into the chromosome of the genomic DNA at the cleavage site.

32. The method of claim 31 wherein the second foreign nucleic acid sequence includes genetic modifications different from the first foreign nucleic acid sequence.

33. The method of claim 32 wherein the second foreign nucleic acid sequence includes the wild type sequence of the chromosome that was modified by the first foreign nucleic acid sequence.

34. A method of rendering two populations unable to produce offspring with one another

comprising

introducing into a first germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and a first set of guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the first set of guide RNAs are complementary to a first set of target locations on genomic DNA of a chromosome,

wherein the first set of target locations are within an essential protein-coding gene and the first foreign nucleic acid sequence is inserted into the essential gene such that the amino acid sequence of the essential protein is preserved but the nucleotide sequence of the gene is altered such that the first set of guide RNAs are noncomplementary to the one or more target locations

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the first set of guide RNAs and the first foreign nucleic acid sequence preserving the amino acid sequence of the essential protein are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the chromosome of the genomic DNA,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the first set of guide RNAs wherein the RNA guided DNA binding protein nuclease and associated guide RNAs co-localize to a first set of associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the first set of target locations in a cleavage site specific manner so as to remove a portion of the genomic DNA within the essential gene, and

inserting the first foreign nucleic acid sequence into the chromosome of the genomic DNA at the cleavage site, and

introducing into a second germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and a second set of guide RNAs and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the second set of guide RNAs are complementary to a second set of target locations on genomic DNA of a chromosome,

wherein the second set of target locations are within the same essential protein-coding gene as the first set of target locations but are not identical to any of the first set of target locations and the second foreign nucleic acid sequence is inserted into the essential gene such that the amino acid sequence of the essential protein is preserved but the nucleotide sequence of the gene is altered such that the second set of guide RNAs are noncomplementary to the second set of target locations,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the second set of guide RNAs and the second foreign nucleic acid sequence preserving the amino acid sequence of the essential protein are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the chromosome of the genomic DNA,

expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the second set of guide RNAs wherein the RNA guided DNA binding protein nuclease and associated second set of guide RNAs co-localize to a second set of associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the second set of target locations in a cleavage site specific manner so as to remove a portion of the genomic DNA within the essential, and

inserting the second foreign nucleic acid sequence into the chromosome of the genomic DNA at the cleavage site,

wherein any mutual progeny of organisms arising from the first germline cell and the second germline cell are not viable where the first set of guide RNAs causes the RNA guided DNA binding protein nuclease to cleave the first set of target locations within the second foreign nucleic acid sequence and the second set of guide RNAs causes the RNA guided DNA binding protein nuclease to cleave the second set of target locations within the first foreign nucleic acid sequence.

35. A method of producing only Y chromosome sperm from a spermatocyte comprising

introducing into the spermatocyte a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to one or more target locations on an X chromosome of the spermatocyte and one or more target locations on a Y chromosome of the spermatocyte,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the Y chromosome,

wherein the second flanking sequence includes a second sequence identical to a second

portion of the target location on the Y chromosome,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs complementary to one or more target locations on the X chromosome of the spermatocyte and one or more target locations on the Y chromosome of the spermatocyte wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a target locations on the Y chromosome co-localize to an associated target location on the Y chromosome and the RNA guided DNA binding protein nuclease cleaves the Y chromosome at the target location in a cleavage site specific manner,

inserting the first foreign nucleic acid sequence into the Y chromosome at the cleavage site, and

wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a target location on the X chromosome co-localize to an associated target location on the X chromosome and the RNA guided DNA binding protein nuclease cleaves the X chromosome at the target location in a cleavage site specific manner so as to render the X chromosome inoperable.

36. A method of producing only Y chromosome sperm from a spermatocyte comprising

expressing an RNA guided DNA binding protein nuclease and one or more guide RNAs, from sequences located on the Y chromosome using corresponding promoter sequences,

wherein the one or more guide RNAs are complementary to one or more target locations on an X chromosome of the spermatocyte,

wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to one or more target locations on the X chromosome co-localize to the one or more associated target locations on the X chromosome and the RNA guided DNA binding protein nuclease cleaves the X chromosome at the one or more target locations in a cleavage site specific manner so as to render the X chromosome inoperable.

37. A method of producing only X chromosome sperm from a spermatocyte comprising

expressing an RNA guided DNA binding protein nuclease and one or more guide RNAs, from sequences located on the X chromosome using corresponding promoter sequences,

wherein the one or more guide RNAs are complementary to one or more target locations on an Y chromosome of the spermatocyte,

wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to one or more target locations on the Y chromosome co-localize to the one or more associated target locations on the Y chromosome and the RNA guided DNA binding protein nuclease cleaves the Y chromosome at the one or more target locations in a cleavage site specific manner so as to render the Y chromosome inoperable.

38. A method of producing only Y chromosome-containing sperm from a spermatocyte

comprising

modifying a Y chromosome to encode a first foreign nucleic acid sequence and a second foreign nucleic acid sequence,

wherein the first foreign nucleic acid sequence encodes a first RNA guided DNA binding protein nuclease and a first set of guide RNAs and corresponding promoter sequences,

wherein the second foreign nucleic acid sequence encodes an RNA guided DNA binding protein nuclease and a second set of guide RNAs and corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the second nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the second set of guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first RNA guided DNA binding protein nuclease cannot recognize the second set of guide RNAs and the second RNA guided DNA binding protein nuclease cannot recognize the first set of guide RNAs,

wherein the first set of guide RNAs are complementary to a plurality of target locations on genomic DNA of a chromosome that is not a sex chromosome,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the chromosome of the genomic DNA,

wherein the first foreign nucleic acid sequence is expressed in the germline before meiosis to produce the first RNA guided DNA binding protein nuclease and the first set of RNAs wherein the RNA guided DNA binding protein nuclease and associated guide RNAs co-localize to a first set of associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the chromosome of the genomic DNA at the first set of target locations in a cleavage site specific manner such that cellular repair mechanisms copy the second foreign nucleic acid sequence from the Y chromosome onto the chromosome of the genomic DNA,

wherein the copy of the second foreign nucleic acid sequence located on the chromosome of the genomic DNA is expressed prior to or during meiosis to produce the second RNA-guided DNA binding protein and the second set of guide RNAs,

wherein the second RNA guided DNA binding protein nuclease and the second set of guide RNAs complementary to one or more target locations on the X chromosome co-localize to the one or more associated target locations on the X chromosome and the RNA guided DNA binding protein nuclease cleaves the X chromosome at the one or more target locations in a cleavage site specific manner so as to render the X chromosome inoperable.

39. A method of producing only X chromosome-containing sperm from a spermatocyte comprising

modifying an X chromosome to encode a first foreign nucleic acid sequence and a second foreign nucleic acid sequence,

wherein the first foreign nucleic acid sequence encodes a first RNA guided DNA binding protein nuclease and a first set of guide RNAs and corresponding promoter sequences,

wherein the second foreign nucleic acid sequence encodes an RNA guided DNA binding protein nuclease and a second set of guide RNAs and corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the second nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the second set of guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first RNA guided DNA binding protein nuclease cannot recognize the second set of guide RNAs and the second RNA guided DNA binding protein nuclease cannot recognize the first set of guide RNAs,

wherein the first set of guide RNAs are complementary to a plurality of target locations on genomic DNA of a chromosome that is not a sex chromosome,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the chromosome of the genomic DNA,

wherein the first foreign nucleic acid sequence is expressed in the germline before meiosis to produce the first RNA guided DNA binding protein nuclease and the first set of RNAs wherein the RNA guided DNA binding protein nuclease and associated guide RNAs co-localize to a first set of associated target locations on the chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the chromosome of the genomic DNA at the first set of target locations in a cleavage site specific manner such that cellular repair mechanisms copy the second foreign nucleic acid sequence from the X chromosome onto the chromosome of the genomic DNA,

wherein the copy of the second foreign nucleic acid sequence located on the chromosome of the genomic DNA is expressed prior to or during meiosis to produce the second RNA-guided DNA binding protein and the second set of guide RNAs,

wherein the second RNA guided DNA binding protein nuclease and the second set of guide RNAs complementary to one or more target locations on the Y chromosome co-localize to the one or more associated target locations on the Y chromosome and the RNA guided DNA binding protein nuclease cleaves the Y chromosome at the one or more target locations in a

cleavage site specific manner so as to render the Y chromosome inoperable.

40. A method of producing female progeny to the exclusion of male progeny with a modified X chromosome having a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs complementary to one or more target nucleic acid sequences on the Y chromosome of a sperm and one or more guide RNAs complementary to one or more target nucleic acid sequences on the X chromosome of a sperm and a first flanking sequence and a second flanking sequence, expressing the first foreign nucleic acid sequence in the oocyte to produce an RNA guided DNA binding protein nuclease and one or more guide RNAs complementary to one or more target nucleic acid sequences on the Y chromosome and one or more target nucleic acid sequences on the X chromosome, wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence, wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the X chromosome of the genomic DNA, wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the X chromosome of the genomic DNA, expressing the first foreign nucleic acid sequence to produce an RNA guided DNA binding protein nuclease and one or more guide RNAs complementary to the one or more target nucleic acid sequences on the Y chromosome and the one or more target nucleic acid sequences on the X chromosome, wherein combining a wild type sperm having a Y chromosome with an oocyte containing the modified X chromosome results in colocalizing the RNA guided DNA binding protein nuclease and the one or more guide RNAs targeting the Y chromosome to the one or more target nucleic acid sequences on the Y chromosome and the RNA guided DNA binding protein nuclease cuts the Y chromosome rendering subsequent development of the oocyte into a male organism inoperable, and

combining a wild type sperm having a X chromosome with an oocyte containing the modified X chromosome results in colocalizing the RNA guided DNA binding protein nuclease and the one or more guide RNAs to the one or more target nucleic acid sequences on the X chromosome where the RNA guided DNA binding protein nuclease cuts the X chromosome and the first foreign nucleic acid sequence is inserted into the X chromosome.

41. A method of producing male progeny to the exclusion of female progeny with a modified Y chromosome having a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs complementary to one or more target nucleic acid sequences on the X chromosome located within a gene that is essential for development, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, wherein the first flanking sequence includes a first sequence identical to a first portion of the essential gene on the X chromosome, wherein the second flanking sequence

includes a second sequence identical to a second portion of the essential gene on the X chromosome, and the modified Y chromosome further includes a second nucleic acid sequence encoding the essential gene on the X chromosome that is altered such that the one or more target locations recognized by the one or more guide RNAs are noncomplementary to the sequences recognized by one or more the guide RNAs, such that when an oocyte having a wild-type X chromosome is combined with a sperm having the modified Y chromosome results in expressing the first foreign nucleic acid sequence produce an RNA guided DNA binding protein nuclease and one or more guide RNAs complementary to one or more target nucleic acid sequences on the X chromosome, colocalizing the RNA guided DNA binding protein nuclease and the one or more guide RNAs to the one or more target nucleic acid sequences on the X chromosome where the RNA guided DNA binding protein nuclease cuts the X chromosome to remove the essential gene and to insert the first nucleic acid sequence therein, creating a modified X chromosome that remains viable due to the additional copy of the essential gene on the Y chromosome, and

combining an oocyte having a wild-type X chromosome with a sperm having the modified X chromosome results in expressing the first foreign nucleic acid sequence to produce an RNA guided DNA binding protein nuclease and one or more guide RNAs complementary to one or more target nucleic acid sequences on the X chromosome, colocalizing the RNA guided DNA binding protein nuclease and the one or more guide RNAs to the one or more target nucleic acid sequences on the X chromosome where the RNA guided DNA binding protein nuclease cuts the X chromosome to remove the essential gene and to insert the first foreign nucleic acid sequence into the X chromosome, wherein the zygote consequently lacks the essential X chromosome gene.

42. A method for generating fertile males and infertile females, wherein the method of claim 41 is modified in that the essential gene on the X chromosome is only essential for fertility, such that organisms lacking that gene are infertile.

43. The method of claim 41, wherein the first nucleic acid sequence additionally contains a third flanking sequence and a fourth flanking sequence, wherein the third flanking sequence includes a third sequence identical to a first portion of a gene essential for fertility, wherein the fourth flanking sequence includes a fourth sequence identical to a second portion of a gene essential for fertility, wherein the RNA-guided DNA-binding protein nuclease is located between the third flanking sequence and the fourth flanking sequence, and the first nucleic acid sequence additionally contains a second set of guide RNAs complementary to one or more target sequences located within the gene essential for fertility, expressing the RNA-guided DNA binding protein nuclease and second set of guide RNAs, colocalizing the RNA guided DNA binding protein nuclease and the one or more guide RNAs to the one or more target nucleic acid sequences within the gene essential for fertility, wherein the RNA guided DNA binding protein nuclease cuts the gene essential for fertility and inserts the first nucleic acid sequence therein.

44. A method for controlled population suppression wherein the Y chromosome of claim 43 is released into the population to be suppressed and the RNA guided gene drive of claim 30 that alters the one or more target sequences located within the gene essential for fertility that is targeted by the Y chromosome of claim 43 such that they are noncomplementary to the guide RNAs employed by the Y chromosome of claim 43 but the protein encoded by the gene retains the same amino acid sequence is released in a population to be protected such that the Y chromosome produces only sons in the population to be suppressed and produces fertile sons and sterile daughters upon mating with females that have been modified by the RNA guided gene drive released in the population to be protected.

45. A method for targeted population suppression or extinction comprising releasing the Y chromosome of claim 44 into a population to be suppressed or eradicated.

46. A method for targeted population suppression or extinction comprising releasing a sex-biasing chromosome into the targeted population.

47. A method of blocking gene flow from an engineered organism to a wild type organism of the same species comprising first modifying the genome of the wild-type organism such that sequences corresponding to those from the middle of a gene that must be present as two functional copies for the organism to be viable that is hereafter termed the essential gene are inserted into another chromosomal region distal to the wild-type copy of the gene, and subsequently

introducing into the germline of the engineered organism a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence and a third flanking sequence and a fourth flanking sequence and a sequence encoding the same protein as the essential gene but does not utilize the same nucleic acid sequence and is hereafter termed the recoded essential gene,

wherein the one or more guide RNAs are complementary to the one or more target locations inserted into the distal region of the chromosome and also to the sequences within the essential gene,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs and the third flanking sequence and the fourth flanking sequence and the recoded essential gene are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location in the distal region of the chromosome,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location in the distal region of the chromosome,

wherein the third flanking sequence includes a third sequence identical to a third portion of

the essential gene,

wherein the fourth flanking sequence includes a fourth sequence identical to a fourth portion of the essential gene,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs complementary to one or more target locations in the distal region of the chromosome and to the one or more target sequences within the essential gene, wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a target location co-localize to an associated target location in the distal region of the chromosome, and the RNA guided DNA binding protein nuclease cleaves the target sequence in the distal region of the chromosome in a cleavage site specific manner,

inserting the first foreign nucleic acid sequence into the distal region of the chromosome,

wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a target location co-localize to an associated target location in the wild-type essential gene, and the RNA guided DNA binding protein nuclease cleaves the wild-type essential gene in a cleavage site specific manner,

inserting the sequence between the third flanking region and the fourth flanking region into the wild-type copy of the essential gene thereby disrupting its function, such that

the engineered organism has two recoded copies of the essential gene in a distal region of the chromosome and no functional copies of the wild-type essential gene and crossing an engineered organism with a wild-type organism causes the disruption of the wild-type copy of the essential gene without inserting an additional copy of the recoded essential gene into the distal region of the chromosome.

48. The method of claim 47, where the essential gene is located on the X chromosome and does require two copies for viability, and the distal region is also on the X chromosome, and the first modification of the organism also modifies the Y chromosome such that a gene required for male fertility or viability hereafter termed the male essential gene is recoded in its nucleic acid sequence in a manner that preserves the amino acid sequence of the encoded male essential protein, and an additional copy of the male essential gene that is nonfunctional is inserted next to the recoded copy of the male essential gene, and the first nucleic acid sequence also includes a fifth flanking sequence that includes a fifth sequence identical to a fifth portion of the male essential gene, and the first nucleic acid sequence also includes a sixth flanking sequence that includes a sixth sequence identical to a sixth portion of the male essential gene, and the first nucleic acid sequence also includes a second set of guide RNAs corresponding to sequences within the wild-type male essential gene that were recoded to render the gene nonfunctional, and the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the guide RNAs and the third flanking sequence and the fourth flanking sequence and the recoded

essential gene are between the fifth flanking sequence and the sixth flanking sequence which are between the first flanking sequence and the second flanking sequence, a female with two modified X chromosomes mated with a male with a modified Y chromosome yields viable male offspring because the RNA-guided DNA-binding protein nuclease does not cut the recoded or the nonfunctional male essential gene and the modified Y chromosome does not cut the essential gene on the modified X chromosome, and an oocyte from a female with two modified X chromosomes that is fertilized by a sperm with a wild-type Y chromosome results in the RNA-guided DNA-binding protein nuclease cutting the wild-type male essential gene and copying the sequence between the fifth flanking sequence and the sixth flanking sequence in its place and resulting in a sterile or nonviable male, and an oocyte from a wild-type female that is fertilized by a sperm with a modified Y chromosome results in the RNA-guided DNA-binding protein nuclease cutting the essential gene and resulting in nonviable male progeny, and the combination of a sperm and an egg that together include a modified and a wild-type X chromosome result in the RNA-guided DNA-binding protein nuclease cutting the wild-type essential gene without copying the recoded essential gene, resulting in hybrid female progeny with one modified X chromosome and one disrupted X chromosome, and these hybrid females mated with wild-type males produce nonviable or sterile males or hybrid females, and half the male progeny of hybrid females and males bearing a modified Y chromosome are viable.

49. A method of blocking gene flow from a subpopulation bearing a unique genetic polymorphism to the remainder of the wild-type population comprising releasing the RNA guided gene drive described in claim 30 that can spread only through populations bearing a unique polymorphism in the target location between the first flanking sequence and the second flanking sequence which additionally inserts the target sites into a distal region of the chromosome wherein the RNA guided gene drive encodes for the first and second flanking sequences for insertion of the RNA guided gene drive into the unique polymorphism region and which encodes third and fourth flanking sequences for inserting the target sites into the distal region, and subsequently

releasing an organism that has been secondarily modified by introducing into the germline of an organism so first modified a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and a second set of guide RNAs and a third set of guide RNAs, and including corresponding promoter sequences and a fifth flanking sequence and a sixth flanking sequence and a seventh flanking sequence and an eighth flanking sequence and a ninth flanking sequence and a tenth flanking sequence and a sequence encoding the same protein as the essential gene but that does not utilize the same nucleic acid sequence and is hereafter termed the recoded essential gene,

wherein the second set of guide RNAs are complementary to the first set of target locations inserted into the distal region of the chromosome and also to the sequences within the essential

gene,

wherein the third set of guide RNAs are complementary to a set of target sequences that are uniquely found within the first nucleic acid sequence,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs and the seventh flanking sequence and the eighth flanking sequence and the ninth flanking sequence and the tenth flanking sequence and the recoded essential gene are between the fifth flanking sequence and the sixth flanking sequence,

wherein the fifth flanking sequence includes a fifth sequence identical to a fifth portion of the target location in the distal region of the chromosome,

wherein the sixth flanking sequence includes a sixth sequence identical to a sixth portion of the target location in the distal region of the chromosome,

wherein the seventh flanking sequence includes a seventh sequence identical to a seventh portion of the essential gene,

wherein the eighth flanking sequence includes a eighth sequence identical to a eighth portion of the essential gene,

wherein the ninth flanking sequence includes a ninth sequence identical to an ninth portion of the first flanking sequence,

wherein the tenth flanking sequence includes a tenth sequence identical to a tenth portion of the second flanking sequence,

expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the second set RNAs complementary to the second set of target locations in the distal region of the chromosome and to the second set target sequences within the essential gene, wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a target location co-localize to an associated target location in the distal region of the chromosome, and the RNA guided DNA binding protein nuclease cleaves the target sequence in the distal region of the chromosome in a cleavage site specific manner,

inserting the first foreign nucleic acid sequence into the distal region of the chromosome,

wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a target location co-localize to an associated target location in the wild-type essential gene, and the RNA guided DNA binding protein nuclease cleaves the wild-type essential gene in a cleavage site specific manner,

inserting the sequence between the seventh flanking region and the eighth flanking region into the wild-type copy of the essential gene thereby disrupting its function,

expressing the third set of guide RNAs complementary to sequences uniquely located within the first nucleic acid sequence,

wherein the RNA guided DNA binding protein nuclease and an associated guide RNA complementary to a sequence within the first nucleic acid sequence co-localize to a target sequence within the first nucleic acid sequence, and the RNA guided DNA binding protein nuclease cleaves the target sequence within the first nucleic acid sequence in a cleavage site specific manner,

replacing the first nucleic acid sequence with the sequence between the ninth flanking region and the tenth flanking region, such that

the engineered organism has no copies of the first nucleic acid sequence and two recoded copies of the essential gene in a distal region of the chromosome and no functional copies of the wild-type essential gene and crossing an engineered organism with a wild-type organism causes the disruption of the wild-type copy of the essential gene without inserting an additional copy of the recoded essential gene into the distal region of the chromosome.

51. A germline cell comprising

a foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA.

52. A method of altering a eukaryotic germline cell of an organism comprising

introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, and including a sensitizing nucleic acid the expression of which is harmful to the organism when the organism is exposed to a chemical, compound or condition,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein

nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner,

inserting the first foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the first foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the foreign nucleic acid sequence, and

expressing the sensitizing nucleic acid rendering the resulting organism sensitive to the chemical, compound or condition such that the resulting organism perishes or is rendered sterile when exposed to the chemical, compound or condition.

53. The method of claim 52 wherein expression of the sensitizing nucleic acid increases toxicity of the chemical, compound or condition to the organism.

54. The method of claim 52 wherein the germline cell is grown into an organism and the sensitizing nucleic acid is transferred to progeny to create a population of organisms including the sensitizing nucleic acid and where the sensitizing nucleic acid increases toxicity of the chemical, compound or condition to the organism.

55. The method of claim 52 wherein the organism is a weed or pest.

56. The method of claim 52, where the sensitizing nucleic acid is a sensitizing gene which replaces an existing gene.

57. The method of claim 56, where the sensitizing gene is the exact or codon-altered ancestral version of an existing mutant gene in wild populations, such that the current mutated version is replaced with the ancestral form.

58. The method of claim 56, where the existing gene has acquired a mutation contributing to resistance to a pesticide, herbicide, or fungicide.

59. The method of claim 52 wherein the chemical or compound is a pesticide, herbicide or

fungicide.

60. The method of claim 59, where the pesticide, herbicide, or fungicide is one of: a Bt toxin produced by Cry1A.105, CryIAb, CryIF, Cry2Ab, Cry3Bb1, Cry34Ab1, Cry35Ab1, mCry3A, or VIP, 2,4-D or glyphosate.

61. The method of claim 56, where the sensitizing gene replaces an existing gene whose function is required for the organism to survive or reproduce.

62. The method of claim 52, where the chemical is a prodrug and the sensitizing gene encodes a corresponding prodrug-converting enzyme.

63. The method of claim 62, where the enzyme/chemical pairing is cytosine deaminase/5-fluorocytosine, or nitroreductase/CB1954.

64. A method of controlling a weed or pest population including a sensitivity gene drive in the genome of the weed or pest population wherein the sensitivity gene drive renders the weed or pest population vulnerable to toxicity when in the presence of a chemical, compound or condition comprising

contacting the weed or pest population with the chemical, compound or condition in an effective amount to kill the weed or pest, lower proliferation of the weed or pest or render the weed or pest sterile to inhibit proliferation.

65. The method of claim 64 wherein the chemical or compound is an herbicide, or pesticide or fungicide.

66. A method of altering a eukaryotic germline cell of an organism including a first sensitizing gene drive comprising

introducing into the germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, and including a second sensitizing nucleic acid sequence the expression of which is harmful to the organism when the organism is exposed to a chemical, compound or condition,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome including the first sensitizing gene drive and a second chromosome of a chromosome pair of the germline cell including the first sensitizing gene drive,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs of the second foreign nucleic acid sequence are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second

portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner, wherein the first foreign nucleic acid sequence is removed,

inserting the second foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the second foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the second foreign nucleic acid sequence, and

expressing the second sensitizing nucleic acid sequence rendering the resulting organism sensitive to the chemical, compound or condition such that the resulting organism perishes or is rendered sterile when exposed to the chemical, compound or condition.

67. The method of claim 66 wherein the resulting organism is introduced into a wild type population such as that progeny of the resulting organism and a wild type organism includes the second sensitizing nucleic acid sequence.

68. The method of claim 66 wherein the organism is a weed or pest.

69. A method of altering a eukaryotic germline cell of an organism including a first sensitizing gene drive comprising

introducing into the germline cell a second foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence, and including a second sensitizing nucleic acid sequence the expression of which is harmful to the organism when the organism is exposed to a chemical, compound or condition,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome including the first sensitizing gene drive and a second chromosome of a chromosome pair of the germline cell including the first sensitizing gene drive,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs of the second foreign nucleic acid sequence are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

expressing the second foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner,

inserting the second foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the second foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the second foreign nucleic acid sequence, and

expressing the second sensitizing nucleic acid sequence rendering the resulting organism sensitive to the chemical, compound or condition such that the resulting organism perishes or is rendered sterile when exposed to the chemical, compound or condition.

70. The method of claim 69 wherein the resulting organism is introduced into a wild type population such as that progeny of the resulting organism and a wild type organism includes the second sensitizing nucleic acid sequence.

71. The method of claim 69 wherein the organism is a weed or pest.

72. The method of altering a eukaryotic germline cell of an organism comprising

introducing into the germline cell a first foreign nucleic acid sequence encoding an RNA guided DNA binding protein nuclease and one or more guide RNAs, and including corresponding promoter sequences and a first flanking sequence and a second flanking sequence,

wherein the one or more guide RNAs are complementary to one or more target locations on genomic DNA of a first chromosome and a second chromosome of a chromosome pair of the germline cell,

wherein the nucleic acid sequence encoding the RNA guided DNA binding protein nuclease and the nucleic acid sequence encoding the one or more guide RNAs are between the first flanking sequence and the second flanking sequence,

wherein the first flanking sequence includes a first sequence identical to a first portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein the second flanking sequence includes a second sequence identical to a second portion of the target location on the first chromosome or the second chromosome of the genomic DNA,

wherein at least one copy of the sequence located between the first flanking sequence and the second flanking sequence that is cut and replaced by the foreign nucleic acid sequence is required for the organism to survive or to produce viable offspring,

expressing the first foreign nucleic acid sequence to produce the RNA guided DNA binding protein nuclease and the one or more RNAs wherein the RNA guided DNA binding protein nuclease and an associated guide RNA co-localize to an associated target location on the first chromosome of the genomic DNA and the second chromosome of the genomic DNA and the RNA guided DNA binding protein nuclease cleaves the first chromosome of the genomic DNA at the target location in a cleavage site specific manner and cleaves the second chromosome of the genomic DNA at the target location in a cleavage site specific manner,

inserting the first foreign nucleic acid sequence into the first chromosome of the chromosome pair of the genomic DNA at the cleavage site, and inserting the first foreign nucleic acid sequence into the second chromosome of the chromosome pair of the genomic DNA at the cleavage site to render the germline cell homozygous for the foreign nucleic acid sequence, and

performing the above expression and insertion steps at developmental stage at which the sequence located between the first flanking sequence and the second flanking sequence that is cut and replaced by the foreign nucleic acid sequence to produce a genetic load is no longer required for that organism to survive or produce fertile offspring.

73. A method for targeted population suppression or extinction according to any previous claim comprising releasing an RNA-guided genetic load drive into the targeted population.