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(54) Title: NON-VIRAL DNA VECTORS AND USES THEREOF FOR EXPRESSING FACTOR IX THERAPEUTICS

(57) Abstract: The application describes ceDNA vectors having linear and continuous structure for delivery and expression of a transgene. ceDNA vectors comprise an expression cassette flanked by two ITR sequences, where the expression cassette encodes a transgene encoding FIX protein. Some ceDNA vectors further comprise cis-regulatory elements, including regulatory switches. Further provided herein are methods and cell lines for reliable gene expression of FIX protein *in vitro*, *ex vivo* and *in vivo* using the ceDNA vectors. Provided herein are method and compositions comprising ceDNA vectors useful for the expression of FIX protein in a cell, tissue or subject, and methods of treatment of diseases with said ceDNA vectors expressing FIX protein. Such FIX protein can be expressed for treating disease, e.g., hemophilia B.



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NON-VIRAL DNA VECTORS AND USES THEREOF FOR EXPRESSING FACTOR IX THERAPEUTICS

RELATED APPLICATIONS

[0001] The present application claims the benefit of priority to U.S. Provisional Application No. 62/993,857, filed on March 24, 2020, the entire contents of which is incorporated by reference in its entirety herein.

SEQUENCE LISTING

[0002] The instant application contains a Sequence Listing which has been submitted electronically in ASCII format and is hereby incorporated by reference in its entirety. Said ASCII copy, created on March 22, 2021, is named 131698-06520_SL.txt and is 394,694 bytes in size.

TECHNICAL FIELD

[0003] The present disclosure relates to the field of gene therapy, including non-viral vectors for expressing a transgene or isolated polynucleotides in a subject or cell. The disclosure also relates to nucleic acid constructs, promoters, vectors, and host cells including the polynucleotides as well as methods of delivering exogenous DNA sequences to a target cell, tissue, organ or organism. For example, the present disclosure provides methods for using non-viral ceDNA vectors to express FIX, from a cell, *e.g.*, expressing the FIX therapeutic protein for the treatment of a subject with a hemophilia B. The methods and compositions can be applied *e.g.*, for the purpose of treating disease by expressing the FIX protein in a cell or tissue of a subject in need thereof.

BACKGROUND

[0004] Gene therapy aims to improve clinical outcomes for patients suffering from either genetic mutations or acquired diseases caused by an aberration in the gene expression profile. Gene therapy includes the treatment or prevention of medical conditions resulting from defective genes or abnormal gene regulation or expression, *e.g.* underexpression or overexpression, that can result in a disorder, disease, malignancy, etc. For example, a disease or disorder caused by a defective gene might be treated, prevented or ameliorated by delivery of a corrective genetic material to a patient, or might be treated, prevented or ameliorated by altering or silencing a defective gene, *e.g.*, with a corrective genetic material to a patient resulting in the therapeutic expression of the genetic material within the patient.

[0005] The basis of gene therapy is to supply a transcription cassette with an active gene product (sometimes referred to as a transgene), *e.g.*, that can result in a positive gain-of-function effect, a negative loss-of-function effect, or another outcome. Such outcomes can be attributed to expression of an activating antibody or fusion protein or an inhibitory (neutralizing) antibody or fusion protein. Gene

therapy can also be used to treat a disease or malignancy caused by other factors. Human monogenic disorders, disorders caused by variations in a single gene, can be treated by the delivery and expression of a normal gene to the target cells. Delivery and expression of a corrective gene in the patient's target cells can be carried out via numerous methods, including the use of engineered viruses and viral gene delivery vectors. Among the many virus-derived vectors available (*e.g.*, recombinant retrovirus, recombinant lentivirus, recombinant adenovirus, and the like), recombinant adeno-associated virus (rAAV) is gaining popularity as a versatile vector in gene therapy.

[0006] Adeno-associated viruses (AAVs) belong to the *Parvoviridae* family and more specifically constitute the *Dependoparvovirus* genus. Vectors derived from AAV (*i.e.*, rAAV or AAV vectors) are attractive for delivering genetic material because (i) they are able to infect (transduce) a wide variety of non-dividing and dividing cell types including myocytes and neurons; (ii) they are devoid of the virus structural genes, thereby diminishing the host cell responses to virus infection, *e.g.*, interferon-mediated responses; (iii) wild-type viruses are considered non-pathogenic in humans; (iv) in contrast to wild type AAV, which are capable of integrating into the host cell genome, replication-deficient AAV vectors lack the replication (*rep*) gene and generally persist as episomes, thus limiting the risk of insertional mutagenesis or genotoxicity; and (v) in comparison to other vector systems, AAV vectors are generally considered to be relatively poor immunogens and therefore do not trigger a significant immune response (see ii), thus gaining persistence of the vector DNA and potentially, long-term expression of the therapeutic transgenes.

[0007] However, there are several major deficiencies in using AAV particles as a gene delivery vector. One major drawback associated with rAAV is its limited viral packaging capacity of about 4.5 kb of heterologous DNA (Dong et al., 1996; Athanasopoulos et al., 2004; Lai et al., 2010), and as a result, use of AAV vectors has been limited to less than 150,000 Da protein coding capacity. The second drawback is that as a result of the prevalence of wild-type AAV infection in the population, candidates for rAAV gene therapy have to be screened for the presence of neutralizing antibodies that eliminate the vector from the patient. A third drawback is related to the capsid immunogenicity that prevents re-administration to patients that were not excluded from an initial treatment. The immune system in the patient can respond to the vector which effectively acts as a “booster” shot to stimulate the immune system generating high titer anti-AAV antibodies that preclude future treatments. Some recent reports indicate concerns with immunogenicity in high dose situations. Another notable drawback is that the onset of AAV-mediated gene expression is relatively slow, given that single-stranded AAV DNA must be converted to double-stranded DNA prior to heterologous gene expression.

[0008] Additionally, conventional AAV virions with capsids are produced by introducing a plasmid or plasmids containing the AAV genome, *rep* genes, and *cap* genes (Grimm et al., 1998). However, such encapsidated AAV virus vectors were found to inefficiently transduce certain cell and tissue types and the capsids also induce an immune response.

[0009] Accordingly, use of adeno-associated virus (AAV) vectors for gene therapy is limited due to the single administration to patients (owing to the patient immune response), the limited range of transgene genetic material suitable for delivery in AAV vectors due to minimal viral packaging capacity (about 4.5kb), and slow AAV-mediated gene expression.

[0010] There is large unmet need for disease-modifying therapies in hemophilia B. Current therapies are burdensome and require frequent intravenous (IV) administrations. First, these Factor IX injectables do not provide continuous delivery of factors, with trough levels allowing bleeding episodes. Second, there are no approved gene therapies for hemophilia B, and AAV based therapies cannot be used by 25% to 40% of patients due to pre-existing antibodies. AAV can only be administered once, and the resulting Factor IX levels might not be high enough to be efficacious, or may be supranormal, dose levels cannot be titrated. Third, some hemophilia B patients cannot utilize these therapies because of the development of neutralizing antibodies to these exogenous, artificial clotting factors.

[0011] Accordingly, there is need in the field for a technology that permits expression of a therapeutic FIX protein in a cell, tissue or subject for the treatment of hemophilia B.

BRIEF DESCRIPTION

[0012] The technology described herein relates to methods and compositions for treatment of Hemophilia B by expression of Factor IX (FIX) protein from a capsid-free (*e.g.*, non-viral) DNA vector with covalently-closed ends (referred to herein as a “closed-ended DNA vector” or a “ceDNA vector”), where the ceDNA vector comprises a FIX nucleic acid sequence or codon optimized versions thereof. These ceDNA vectors can be used to produce FIX proteins for treatment, monitoring, and diagnosis. The application of ceDNA vectors expressing FIX to the subject for the treatment of hemophilia B is useful to: (i) provide disease modifying levels of FIX enzyme, be minimally invasive in delivery, be repeatable and dosed-to-effect, have rapid onset of therapeutic effect, result in sustained expression of corrective FIX enzyme in the liver, restoring the coagulation cascade, and/or be titratable to achieve the appropriate pharmacologic levels of the defective enzyme.

[0013] In some embodiments, a ceDNA-vector expressing FIX is optionally present in a liposome nanoparticle formulation (LNP) for the treatment of hemophilia B. A ceDNA LNP formulation described herein can provide one or more benefits, including: providing disease modifying levels of FIX protein, being minimally invasive in delivery, being repeatable and dosed-to-effect, having a rapid onset of therapeutic effect that is typically within days of therapeutic intervention, having sustained expression of corrective FIX levels in the circulation, be titratable to achieve the appropriate pharmacologic levels of the defective coagulation factor, and/or provide treatments for other types of hemophilia, including but not limited to Factor VII deficiency.

[0014] Accordingly, the present disclosure relates to a capsid-free (*e.g.*, non-viral) DNA vector with covalently-closed ends (referred to herein as a “closed-ended DNA vector” or a “ceDNA vector”)

comprising a gene encoding FIX, to permit expression of the FIX therapeutic protein in a cell. In one embodiment, the gene encoding FIX is a heterologous gene.

[0015] The ceDNA vectors for expression of FIX protein production as described herein are capsid-free, linear duplex DNA molecules formed from a continuous strand of complementary DNA with covalently-closed ends (linear, continuous and non-encapsidated structure), which comprise a 5' inverted terminal repeat (ITR) sequence and a 3' ITR sequence, where the 5' ITR and the 3' ITR can have the same symmetrical three-dimensional organization with respect to each other, (*i.e.*, symmetrical or substantially symmetrical), or alternatively, the 5' ITR and the 3' ITR can have different three-dimensional organization with respect to each other (*i.e.*, asymmetrical ITRs). In addition, the ITRs can be from the same or different serotypes. In some embodiments, a ceDNA vector can comprise ITR sequences that have a symmetrical three-dimensional spatial organization such that their structure is the same shape in geometrical space, or have the same A, C-C' and B-B' loops in 3D space (*i.e.*, they are the same or are mirror images with respect to each other). In some embodiments, one ITR can be from one AAV serotype, and the other ITR can be from a different AAV serotype.

[0016] Accordingly, some aspects of the technology described herein relate to a ceDNA vector for improved protein expression and/or production of the above described FIX protein that comprise ITR sequences that flank a nucleic acid sequence comprising any FIX nucleic acid sequence disclosed in Table 1 or any open reading frame sequence included in any ceDNA sequence disclosed in Table 12, the ITR sequences being selected from any of: (i) at least one WT ITR and at least one modified AAV inverted terminal repeat (ITR) (*e.g.*, asymmetric modified ITRs); (ii) two modified ITRs where the mod-ITR pair have a different three-dimensional spatial organization with respect to each other (*e.g.*, asymmetric modified ITRs), or (iii) symmetrical or substantially symmetrical WT-WT ITR pair, where each WT-ITR has the same three-dimensional spatial organization, or (iv) symmetrical or substantially symmetrical modified ITR pair, where each mod-ITR has the same three-dimensional spatial organization. The ceDNA vectors disclosed herein can be produced in eukaryotic cells, thus devoid of prokaryotic DNA modifications and bacterial endotoxin contamination in insect cells.

[0017] The methods and compositions described herein relate, in part, to the discovery of a non-viral capsid-free DNA vector with covalently-closed ends (ceDNA vectors) that can be used to express at least one FIX protein, or more than one FIX protein, from a cell, including but not limited to cells of the liver.

[0018] Provided herein in one aspect are DNA vectors (*e.g.*, ceDNA vectors) comprising at least one nucleic acid sequence, wherein the nucleic acid sequence encodes a transgene, operably linked to a promoter positioned between two different AAV inverted terminal repeat sequences (ITRs), one of the ITRs comprising a functional AAV terminal resolution site and a Rep binding site, and one of the ITRs comprising a deletion, insertion, or substitution relative to the other ITR; wherein the transgene encodes an FIX protein; and wherein the DNA when digested with a restriction enzyme having a single recognition site on the DNA vector has the presence of characteristic bands of linear and

continuous DNA as compared to linear and non-continuous DNA controls when analyzed on a non-denaturing gel. Other aspects include delivery of the FIX protein by expressing it *in vivo* from a ceDNA vector as described herein and further, the treatment of hemophilia B using ceDNA vectors encoding the FIX protein. Also contemplated herein are cells comprising a ceDNA vector encoding an FIX protein as described herein.

[0019] Aspects of the disclosure relate to methods to produce the ceDNA vectors useful for FIX protein expression in a cell as described herein. Other embodiments relate to a ceDNA vector produced by the methods provided herein. In one embodiment, the capsid free (*e.g.*, non-viral) DNA vector (ceDNA vector) for FIX protein production is obtained from a plasmid (referred to herein as a “ceDNA-plasmid”) comprising a polynucleotide expression construct template comprising in this order: a first 5' inverted terminal repeat (*e.g.* AAV ITR); a nucleic acid sequence; and a 3' ITR (*e.g.* AAV ITR), where the 5' ITR and 3' ITR can be asymmetric relative to each other, or symmetric (*e.g.*, WT-ITRs or modified symmetric ITRs) as defined herein.

[0020] The ceDNA vector for expression of the FIX protein as disclosed herein is obtainable by a number of means that would be known to the ordinarily skilled artisan after reading this disclosure. For example, a polynucleotide expression construct template used for generating the ceDNA vectors of the present disclosure can be a ceDNA-plasmid, a ceDNA-bacmid, and/or a ceDNA-baculovirus. In one embodiment, the ceDNA-plasmid comprises a restriction cloning site (*e.g.* SEQ ID NO: 123 and/or 124) operably positioned between the ITRs where an expression cassette comprising *e.g.*, a promoter operatively linked to a transgene, *e.g.*, a nucleic acid encoding FIX can be inserted. In some embodiments, ceDNA vectors for expression of FIX protein are produced from a polynucleotide template (*e.g.*, ceDNA-plasmid, ceDNA-bacmid, ceDNA-baculovirus) containing symmetric or asymmetric ITRs (modified or WT ITRs).

[0021] In a permissive host cell, in the presence of *e.g.*, Rep, the polynucleotide template having at least two ITRs replicates to produce ceDNA vectors expressing the FIX protein. ceDNA vector production undergoes two steps: first, excision (“rescue”) of template from the template backbone (*e.g.* ceDNA-plasmid, ceDNA-bacmid, ceDNA-baculovirus genome etc.) via Rep proteins, and second, Rep mediated replication of the excised ceDNA vector. Rep proteins and Rep binding sites of the various AAV serotypes are well known to those of ordinary skill in the art. One of ordinary skill understands to choose a Rep protein from a serotype that binds to and replicates the nucleic acid sequence based upon at least one functional ITR. For example, if the replication competent ITR is from AAV serotype 2, the corresponding Rep would be from an AAV serotype that works with that serotype such as AAV2 ITR with AAV2 or AAV4 Rep but not AAV5 Rep, which does not. Upon replication, the covalently-closed ended ceDNA vector continues to accumulate in permissive cells and ceDNA vector is preferably sufficiently stable over time in the presence of Rep protein under standard replication conditions, *e.g.* to accumulate in an amount that is at least 1 pg/cell, preferably at least 2 pg/cell, preferably at least 3 pg/cell, more preferably at least 4 pg/cell, even more preferably at least 5 pg/cell.

[0022] Accordingly, one aspect of the disclosure relates to a process of producing a ceDNA vector for expression of such FIX proteins comprising the steps of: a) incubating a population of host cells (*e.g.* insect cells) harboring the polynucleotide expression construct template (*e.g.*, a ceDNA-plasmid, a ceDNA-bacmid, and/or a ceDNA-baculovirus), which is devoid of viral capsid coding sequences, in the presence of a Rep protein under conditions effective and for a time sufficient to induce production of the ceDNA vector within the host cells, and wherein the host cells do not comprise viral capsid coding sequences; and b) harvesting and isolating the ceDNA vector from the host cells. The presence of Rep protein induces replication of the vector polynucleotide with a modified ITR to produce the ceDNA vector for expression of FIX protein in a host cell. However, no viral particles (*e.g.* AAV virions) are expressed. Thus, there is no virion-enforced size limitation.

[0023] The presence of the ceDNA vector useful for expression of FIX protein is isolated from the host cells can be confirmed by digesting DNA isolated from the host cell with a restriction enzyme having a single recognition site on the ceDNA vector and analyzing the digested DNA material on denaturing and non-denaturing gels to confirm the presence of characteristic bands of linear and continuous DNA as compared to linear and non-continuous DNA.

[0024] Also provided herein are methods of expressing an FIX protein that has therapeutic uses, in a cell or in a subject, using a ceDNA vector. Such FIX proteins can be used for the treatment of hemophilia B. Accordingly, provided herein are methods for the treatment of hemophilia B comprising administering a ceDNA vector encoding a therapeutic FIX protein to a subject in need thereof.

[0025] In some embodiments, one aspect of the technology described herein relates to a non-viral capsid-free DNA vector with covalently-closed ends (ceDNA vector), wherein the ceDNA vector comprises at least one nucleic acid sequence, operably positioned between two ITR sequences where the ITR sequences can be asymmetric, or symmetric, or substantially symmetrical as these terms are defined herein, wherein at least one of the ITRs comprises a functional terminal resolution site (*trs*) and a Rep binding site, and optionally the nucleic acid sequence encodes a transgene (*e.g.*, FIX protein) and wherein the vector is not in a viral capsid.

[0026] These and other aspects of the disclosure are described in further detail below.

DESCRIPTION OF DRAWINGS

[0027] Embodiments of the present disclosure, briefly summarized above and discussed in greater detail below, can be understood by reference to the illustrative embodiments of the disclosure depicted in the appended drawings. However, the appended drawings illustrate only typical embodiments of the disclosure and are therefore not to be considered limiting of scope, for the disclosure may admit to other equally effective embodiments.

[0028] **FIG. 1A** illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein, comprising asymmetric ITRs. In this embodiment, the exemplary ceDNA vector comprises an expression cassette containing CAG promoter, WPRE, and BGHpA. An open

reading frame (ORF) encoding the FIX transgene can be inserted into the cloning site (R3/R4) between the CAG promoter and WPRE. The expression cassette is flanked by two inverted terminal repeats (ITRs) – the wild-type AAV2 ITR on the upstream (5'-end) and the modified ITR on the downstream (3'-end) of the expression cassette, therefore the two ITRs flanking the expression cassette are asymmetric with respect to each other.

[0029] FIG. 1B illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein comprising asymmetric ITRs with an expression cassette containing CAG promoter, WPRE, and BGHpA. An open reading frame (ORF) encoding the FIX transgene can be inserted into the cloning site between CAG promoter and WPRE. The expression cassette is flanked by two inverted terminal repeats (ITRs) – a modified ITR on the upstream (5'-end) and a wild-type ITR on the downstream (3'-end) of the expression cassette.

[0030] FIG. 1C illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein comprising asymmetric ITRs, with an expression cassette containing an enhancer/promoter, the FIX transgene, a post transcriptional element (WPRE), and a polyA signal. An open reading frame (ORF) allows insertion of the FIX transgene into the cloning site between CAG promoter and WPRE. The expression cassette is flanked by two inverted terminal repeats (ITRs) that are asymmetrical with respect to each other; a modified ITR on the upstream (5'-end) and a modified ITR on the downstream (3'-end) of the expression cassette, where the 5' ITR and the 3' ITR are both modified ITRs but have different modifications (*i.e.*, they do not have the same modifications).

[0031] FIG. 1D illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein, comprising symmetric modified ITRs, or substantially symmetrical modified ITRs as defined herein, with an expression cassette containing CAG promoter, WPRE, and BGHpA. An open reading frame (ORF) encoding the FIX transgene is inserted into the cloning site between CAG promoter and WPRE. The expression cassette is flanked by two modified inverted terminal repeats (ITRs), where the 5' modified ITR and the 3' modified ITR are symmetrical or substantially symmetrical.

[0032] FIG. 1E illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein comprising symmetric modified ITRs, or substantially symmetrical modified ITRs as defined herein, with an expression cassette containing an enhancer/promoter, a transgene, a post transcriptional element (WPRE), and a polyA signal. An open reading frame (ORF) allows insertion of a transgene (*e.g.*, the FIX) into the cloning site between CAG promoter and WPRE. The expression cassette is flanked by two modified inverted terminal repeats (ITRs), where the 5' modified ITR and the 3' modified ITR are symmetrical or substantially symmetrical.

[0033] FIG. 1F illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein, comprising symmetric WT-ITRs, or substantially symmetrical WT-ITRs as defined herein, with an expression cassette containing CAG promoter, WPRE, and BGHpA. An open reading frame (ORF) encoding a transgene (*e.g.*, the FIX) is inserted into the cloning site between

CAG promoter and WPRE. The expression cassette is flanked by two wild type inverted terminal repeats (WT-ITRs), where the 5' WT-ITR and the 3' WT ITR are symmetrical or substantially symmetrical.

[0034] FIG. 1G illustrates an exemplary structure of a ceDNA vector for expression of an FIX protein as disclosed herein, comprising symmetric modified ITRs, or substantially symmetrical modified ITRs as defined herein, with an expression cassette containing an enhancer/promoter, a transgene (*e.g.*, the FIX), a post transcriptional element (WPRE), and a polyA signal. An open reading frame (ORF) allows insertion of a transgene (*e.g.*, the FIX) into the cloning site between CAG promoter and WPRE. The expression cassette is flanked by two wild type inverted terminal repeats (WT-ITRs), where the 5' WT-ITR and the 3' WT ITR are symmetrical or substantially symmetrical.

[0035] FIG. 2A provides the T-shaped stem-loop structure of a wild-type left ITR of AAV2 (SEQ ID NO: 52) with identification of A-A' arm, B-B' arm, C-C' arm, two Rep binding sites (RBE and RBE') and also shows the terminal resolution site (*trs*). The RBE contains a series of 4 duplex tetramers that are believed to interact with either Rep 78 or Rep 68. In addition, the RBE' is also believed to interact with Rep complex assembled on the wild-type ITR or mutated ITR in the construct. The D and D' regions contain transcription factor binding sites and other conserved structure. **FIG. 2B** shows proposed Rep-catalyzed nicking and ligating activities in a wild-type left ITR (SEQ ID NO: 53), including the T-shaped stem-loop structure of the wild-type left ITR of AAV2 with identification of A-A' arm, B-B' arm, C-C' arm, two Rep Binding sites (RBE and RBE') and also shows the terminal resolution site (*trs*), and the D and D' region comprising several transcription factor binding sites and other conserved structure.

[0036] FIG. 3A provides the primary structure (polynucleotide sequence) (left) and the secondary structure (right) of the RBE-containing portions of the A-A' arm, and the C-C' and B-B' arm of the wild type left AAV2 ITR (SEQ ID NO: 54). **FIG. 3B** shows an exemplary mutated ITR (also referred to as a modified ITR) sequence for the left ITR. Shown is the primary structure (left) and the predicted secondary structure (right) of the RBE portion of the A-A' arm, the C arm and B-B' arm of an exemplary mutated left ITR (ITR-1, left) (SEQ ID NO: 113). **FIG. 3C** shows the primary structure (left) and the secondary structure (right) of the RBE-containing portion of the A-A' loop, and the B-B' and C-C' arms of wild type right AAV2 ITR (SEQ ID NO: 55). **FIG. 3D** shows an exemplary right modified ITR. Shown is the primary structure (left) and the predicted secondary structure (right) of the RBE containing portion of the A-A' arm, the B-B' and the C arm of an exemplary mutant right ITR (ITR-1, right) (SEQ ID NO: 114). Any combination of left and right ITR (*e.g.*, AAV2 ITRs or other viral serotype or synthetic ITRs) can be used as taught herein. Each of **FIGS. 3A-3D** polynucleotide sequences refer to the sequence used in the plasmid or bacmid/baculovirus genome used to produce the ceDNA as described herein. Also included in each of **FIGS. 3A-3D** are corresponding ceDNA secondary structures inferred from the ceDNA vector configurations in the plasmid or bacmid/baculovirus genome and the predicted Gibbs free energy values.

[0037] **FIG. 4A** is a schematic illustrating an upstream process for making baculovirus infected insect cells (BIICs) that are useful in the production of a ceDNA vector for expression of the FIX as disclosed herein in the process described in the schematic in FIG. 4B. **FIG. 4B** is a schematic of an exemplary method of ceDNA production and **FIG. 4C** illustrates a biochemical method and process to confirm ceDNA vector production. **FIG. 4D** and **FIG. 4E** are schematic illustrations describing a process for identifying the presence of ceDNA in DNA harvested from cell pellets obtained during the ceDNA production processes in FIG. 4B. **FIG. 4D** shows schematic expected bands for an exemplary ceDNA either left uncut or digested with a restriction endonuclease and then subjected to electrophoresis on either a native gel or a denaturing gel. The leftmost schematic is a native gel, and shows multiple bands suggesting that in its duplex and uncut form ceDNA exists in at least monomeric and dimeric states, visible as a faster-migrating smaller monomer and a slower-migrating dimer that is twice the size of the monomer. The schematic second from the left shows that when ceDNA is cut with a restriction endonuclease, the original bands are gone and faster-migrating (*e.g.*, smaller) bands appear, corresponding to the expected fragment sizes remaining after the cleavage. Under denaturing conditions, the original duplex DNA is single-stranded and migrates as a species twice as large as observed on native gel because the complementary strands are covalently linked. Thus in the second schematic from the right, the digested ceDNA shows a similar banding distribution to that observed on native gel, but the bands migrate as fragments twice the size of their native gel counterparts. The rightmost schematic shows that uncut ceDNA under denaturing conditions migrates as a single-stranded open circle, and thus the observed bands are twice the size of those observed under native conditions where the circle is not open. In this figure “kb” is used to indicate relative size of nucleotide molecules based, depending on context, on either nucleotide chain length (*e.g.*, for the single stranded molecules observed in denaturing conditions) or number of basepairs (*e.g.*, for the double-stranded molecules observed in native conditions). **FIG. 4E** shows DNA having a non-continuous structure. The ceDNA can be cut by a restriction endonuclease, having a single recognition site on the ceDNA vector, and generate two DNA fragments with different sizes (1kb and 2kb) in both neutral and denaturing conditions. **FIG. 4E** also shows a ceDNA having a linear and continuous structure. The ceDNA vector can be cut by the restriction endonuclease, and generate two DNA fragments that migrate as 1kb and 2kb in neutral conditions, but in denaturing conditions, the strands remain connected and produce single strands that migrate as 2kb and 4kb.

[0038] **FIG. 5** is an exemplary picture of a denaturing gel running examples of ceDNA vectors with (+) or without (-) digestion with endonucleases (EcoRI for ceDNA construct 1 and 2; BamHI for ceDNA construct 3 and 4; SpeI for ceDNA construct 5 and 6; and XhoI for ceDNA construct 7 and 8) Constructs 1-8 are described in Example 1 of International Application PCT PCT/US18/49996, which is incorporated herein in its entirety by reference. Sizes of bands highlighted with an asterisk were determined and provided on the bottom of the picture.

[0039] **FIG. 6** depicts the results of the experiments described in Example 7 and specifically shows the IVIS images obtained from mice treated with LNP-polyC control (mouse furthest to the left) and four mice treated with LNP-ceDNA-Luciferase (all but the mouse furthest to the left). The four ceDNA-treated mice show significant fluorescence in the liver-containing region of the mouse.

[0040] **FIG. 7** depicts the results of the experiment described in Example 8. The dark specks indicate the presence of the protein resulting from the expressed ceDNA transgene and demonstrate association of the administered LNP-ceDNA with hepatocytes.

[0041] **FIGS. 8A-8B** depict the results of the ocular studies set forth in Example 9. **FIG. 8A** shows representative IVIS images from JetPEI®-ceDNA-Luciferase-injected rat eyes (upper left) versus uninjected eye in the same rat (upper right) or plasmid-Luciferase DNA-injected rat eye (lower left) and the uninjected eye in that same rat (lower right). **FIG. 8B** shows a graph of the average radiance observed in treated eyes or the corresponding untreated eyes in each of the treatment groups. The ceDNA-treated rats demonstrated prolonged significant fluorescence (and hence luciferase transgene expression) over 99 days, in sharp contrast to rats treated with plasmid-luciferase where minimal relative fluorescence (and hence luciferase transgene expression) was observed.

[0042] **FIGS. 9A and 9B** depict the results of the ceDNA persistence and redosing study in Rag2 mice described in Example 10. **FIG. 9A** shows a graph of total flux over time observed in LNP-ceDNA-Luc-treated wild-type c57bl/6 mice or Rag2 mice. **FIG. 9B** provides a graph showing the impact of redose on expression levels of the luciferase transgene in Rag2 mice, with resulting increased stable expression observed after redose (arrow indicates time of redose administration).

[0043] **FIG. 10** provides data from the ceDNA luciferase expression study in treated mice described in Example 11, showing total flux in each group of mice over the duration of the study. High levels of unmethylated CpG correlated with lower total flux observed in the mice over time, while use of a liver-specific promoter correlated with durable, stable expression of the transgene from the ceDNA vector over at least 77 days.

[0044] **FIGS. 11A and 11B** show hydrodynamic delivery of ceDNA vector expressing FIX. **FIG. 11A** shows FIX expression levels in serum samples at day 3 and 7 from mice after hydrodynamic injection of two different ceDNA vectors expressing FIX (LPS1-FIX-v1; LPS1-FIX-v2), or a control ceDNA vector (a ceDNA expressing luciferase only) (shown as Vehicle). Both these FIX ceDNA vectors showed FIX expression. **FIG. 11B** shows FIX expression levels over a 28 day period in serum samples from mice after hydrodynamic injection of the two different ceDNA vectors expressing FIX (LPS1-FIX-v1; LPS1-FIX-v2), or the vehicle control ceDNA vector (expressing luciferase only).

[0045] **FIG. 12** depicts plasma levels of factor IX in mice injected with an LNP formulated FIX ceDNA construct (2.0 mg/kg) at days 0 and 36 and orally dosed with 300 mg/kg of ruxolitinib at day -2, day -1, day 0, day 1 and day 36.

[0046] **FIG. 13A** and **13B** show FIX expression in male CD-1 mice treated with a ceDNA-FIX construct (ceDNA-FIX v1; ceDNA-FIX 2109; or ceDNA-FIX 2112), each of which contains a codon optimized human FIX sequence. **FIG. 13A** shows human FIX expression levels in CD-1 mice treated with 1 µg of ceDNA-FIX v1; ceDNA-FIX 2109; or ceDNA-FIX 2112 via hydrodynamic delivery, measured at Day 3 and Day 7. **FIG. 13B** shows human FIX expression levels in CD-1 mice treated with 10 µg of ceDNA-FIX v1; ceDNA-FIX 2109; or ceDNA-FIX 2112 *via* hydrodynamic delivery, measured at Day 3 and Day 7.

DETAILED DESCRIPTION

[0047] Provided herein is a method for treating hemophilia B using a ceDNA vector comprising one or more nucleic acids that encode an FIX therapeutic protein or fragment thereof. Also provided herein are ceDNA vectors for expression of FIX protein as described herein comprising one or more nucleic acids that encode for the FIX protein. In some embodiments, the expression of FIX protein can comprise secretion of the therapeutic protein out of the cell in which it is expressed. Alternatively, in some embodiments the expressed FIX protein can act or function (*e.g.*, exert its effect) within the cell in which it is expressed. In some embodiments, the ceDNA vector expresses FIX protein in the liver, a muscle (*e.g.*, skeletal muscle) of a subject, or other body part, which can act as a depot for FIX therapeutic protein production and secretion to many systemic compartments.

I. Definitions

[0048] Unless otherwise defined herein, scientific and technical terms used in connection with the present application shall have the meanings that are commonly understood by those of ordinary skill in the art to which this disclosure belongs. It should be understood that this disclosure is not limited to the particular methodology, protocols, and reagents, etc., described herein and as such can vary. The terminology used herein is for the purpose of describing particular embodiments only and is not intended to limit the scope of the present disclosure, which is defined solely by the claims. Definitions of common terms in immunology and molecular biology can be found in The Merck Manual of Diagnosis and Therapy, 19th Edition, published by Merck Sharp & Dohme Corp., 2011 (ISBN 978-0-911910-19-3); Robert S. Porter *et al.* (eds.), Fields Virology, 6th Edition, published by Lippincott Williams & Wilkins, Philadelphia, PA, USA (2013), Knipe, D.M. and Howley, P.M. (ed.), The Encyclopedia of Molecular Cell Biology and Molecular Medicine, published by Blackwell Science Ltd., 1999-2012 (ISBN 9783527600908); and Robert A. Meyers (ed.), Molecular Biology and Biotechnology: a Comprehensive Desk Reference, published by VCH Publishers, Inc., 1995 (ISBN 1-56081-569-8); Immunology by Werner Luttmann, published by Elsevier, 2006; Janeway's Immunobiology, Kenneth Murphy, Allan Mowat, Casey Weaver (eds.), Taylor & Francis Limited, 2014 (ISBN 0815345305, 9780815345305); Lewin's Genes XI, published by Jones & Bartlett Publishers, 2014 (ISBN-1449659055); Michael Richard Green and Joseph Sambrook, Molecular

Cloning: A Laboratory Manual, 4th ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., USA (2012) (ISBN 1936113414); Davis *et al.*, Basic Methods in Molecular Biology, Elsevier Science Publishing, Inc., New York, USA (2012) (ISBN 044460149X); Laboratory Methods in Enzymology: DNA, Jon Lorsch (ed.) Elsevier, 2013 (ISBN 0124199542); Current Protocols in Molecular Biology (CPMB), Frederick M. Ausubel (ed.), John Wiley and Sons, 2014 (ISBN 047150338X, 9780471503385), Current Protocols in Protein Science (CPPS), John E. Coligan (ed.), John Wiley and Sons, Inc., 2005; and Current Protocols in Immunology (CPI) (John E. Coligan, ADA M Kruisbeek, David H Margulies, Ethan M Shevach, Warren Strobe, (eds.) John Wiley and Sons, Inc., 2003 (ISBN 0471142735, 9780471142737), the contents of which are all incorporated by reference herein in their entireties.

[0049] As used herein, the terms “heterologous nucleic acid sequence” and “transgene” are used interchangeably and refer to a nucleic acid of interest (other than a nucleic acid encoding a capsid polypeptide) that is incorporated into and may be delivered and expressed by a ceDNA vector as disclosed herein. According to some embodiments, the term “heterologous nucleic acid” is meant to refer to a nucleic acid (or transgene) that is not present in, expressed by, or derived from the cell or subject to which it is contacted.

[0050] As used herein, the terms “expression cassette” and “transcription cassette” are used interchangeably and refer to a linear stretch of nucleic acids that includes a transgene that is operably linked to one or more promoters or other regulatory sequences sufficient to direct transcription of the transgene, but which does not comprise capsid-encoding sequences, other vector sequences or inverted terminal repeat regions. An expression cassette may additionally comprise one or more *cis*-acting sequences (*e.g.*, promoters, enhancers, or repressors), one or more introns, and one or more post-transcriptional regulatory elements.

[0051] The terms “polynucleotide” and “nucleic acid,” used interchangeably herein, refer to a polymeric form of nucleotides of any length, either ribonucleotides or deoxyribonucleotides. Thus, this term includes single, double, or multi-stranded DNA or RNA, genomic DNA, cDNA, DNA-RNA hybrids, or a polymer including purine and pyrimidine bases or other natural, chemically or biochemically modified, non-natural, or derivatized nucleotide bases. “Oligonucleotide” generally refers to polynucleotides of between about 5 and about 100 nucleotides of single- or double-stranded DNA. However, for the purposes of this disclosure, there is no upper limit to the length of an oligonucleotide. Oligonucleotides are also known as “oligomers” or “oligos” and may be isolated from genes, or chemically synthesized by methods known in the art. The terms “polynucleotide” and “nucleic acid” should be understood to include, as applicable to the embodiments being described, single-stranded (such as sense or antisense) and double-stranded polynucleotides.

[0052] DNA may be in the form of, *e.g.*, antisense molecules, plasmid DNA, DNA-DNA duplexes, pre-condensed DNA, PCR products, vectors (P1, PAC, BAC, YAC, artificial chromosomes), expression cassettes, chimeric sequences, chromosomal DNA, or derivatives and combinations of

these groups. DNA may be in the form of minicircle, plasmid, bacmid, minigene, ministring DNA (linear covalently closed DNA vector), closed-ended linear duplex DNA (CELiD or ceDNA), doggybone (dbDNA™) DNA, dumbbell shaped DNA, minimalistic immunological-defined gene expression (MIDGE)-vector, viral vector or nonviral vectors. RNA may be in the form of small interfering RNA (siRNA), Dicer-substrate dsRNA, small hairpin RNA (shRNA), asymmetrical interfering RNA (aiRNA), microRNA (miRNA), mRNA, rRNA, tRNA, viral RNA (vRNA), and combinations thereof. Nucleic acids include nucleic acids containing known nucleotide analogs or modified backbone residues or linkages, which are synthetic, naturally occurring, and non-naturally occurring, and which have similar binding properties as the reference nucleic acid. Examples of such analogs and/or modified residues include, without limitation, phosphorothioates, phosphorodiamidate morpholino oligomer (morpholino), phosphoramidates, methyl phosphonates, chiral-methyl phosphonates, 2'-O-methyl ribonucleotides, locked nucleic acid (LNA™), and peptide nucleic acids (PNAs). Unless specifically limited, the term encompasses nucleic acids containing known analogues of natural nucleotides that have similar binding properties as the reference nucleic acid. Unless otherwise indicated, a particular nucleic acid sequence also implicitly encompasses conservatively modified variants thereof (*e.g.*, degenerate codon substitutions), alleles, orthologs, SNPs, and complementary sequences as well as the sequence explicitly indicated.

[0053] “Nucleotides” contain a sugar deoxyribose (DNA) or ribose (RNA), a base, and a phosphate group. Nucleotides are linked together through the phosphate groups.

[0054] “Bases” include purines and pyrimidines, which further include natural compounds adenine, thymine, guanine, cytosine, uracil, inosine, and natural analogs, and synthetic derivatives of purines and pyrimidines, which include, but are not limited to, modifications which place new reactive groups such as, but not limited to, amines, alcohols, thiols, carboxylates, and alkylhalides.

[0055] The term “nucleic acid construct” as used herein refers to a nucleic acid molecule, either single- or double-stranded, which is isolated from a naturally occurring gene or which is modified to contain segments of nucleic acids in a manner that would not otherwise exist in nature or which is synthetic. The term nucleic acid construct is synonymous with the term “expression cassette” when the nucleic acid construct contains the control sequences required for expression of a coding sequence of the present disclosure. An “expression cassette” includes a DNA coding sequence operably linked to a promoter.

[0056] By “hybridizable” or “complementary” or “substantially complementary” it is meant that a nucleic acid (*e.g.*, RNA) includes a sequence of nucleotides that enables it to non-covalently bind, *i.e.* form Watson-Crick base pairs and/or G/U base pairs, “anneal”, or “hybridize,” to another nucleic acid in a sequence-specific, antiparallel, manner (*i.e.*, a nucleic acid specifically binds to a complementary nucleic acid) under the appropriate *in vitro* and/or *in vivo* conditions of temperature and solution ionic strength. As is known in the art, standard Watson-Crick base-pairing includes: adenine (A) pairing

with thymidine (T), adenine (A) pairing with uracil (U), and guanine (G) pairing with cytosine (C). In addition, it is also known in the art that for hybridization between two RNA molecules (*e.g.*, dsRNA), guanine (G) base pairs with uracil (U). For example, G/U base-pairing is partially responsible for the degeneracy (*i.e.*, redundancy) of the genetic code in the context of tRNA anti-codon base-pairing with codons in mRNA. In the context of this disclosure, a guanine (G) of a protein-binding segment (dsRNA duplex) of a subject DNA-targeting RNA molecule is considered complementary to a uracil (U), and vice versa. As such, when a G/U base-pair can be made at a given nucleotide position a protein-binding segment (dsRNA duplex) of a subject DNA-targeting RNA molecule, the position is not considered to be non-complementary, but is instead considered to be complementary.

[0057] The terms “peptide,” “polypeptide,” and “protein” are used interchangeably herein, and refer to a polymeric form of amino acids of any length, which can include coded and non-coded amino acids, chemically or biochemically modified or derivatized amino acids, and polypeptides having modified peptide backbones.

[0058] A DNA sequence that “encodes” a particular FIX protein is a DNA nucleic acid sequence that is transcribed into the particular RNA and/or protein. A DNA polynucleotide may encode an RNA (mRNA) that is translated into protein, or a DNA polynucleotide may encode an RNA that is not translated into protein (*e.g.*, tRNA, rRNA, or a DNA-targeting RNA; also called “non-coding” RNA or “ncRNA”).

[0001] As used herein, the term “fusion protein” refers to a polypeptide which comprises protein domains from at least two different proteins. For example, a fusion protein may comprise (i) FIX or fragment thereof and (ii) at least one non-gene of interest (GOI) protein. Fusion proteins encompassed herein include, but are not limited to, an antibody, or Fc or antigen-binding fragment of an antibody fused to a FIX protein, *e.g.*, an extracellular domain of a receptor, ligand, enzyme or peptide. The FIX protein or fragment thereof that is part of a fusion protein can be a monospecific antibody or a bispecific or multispecific antibody.

[0059] As used herein, the term “genomic safe harbor gene” or “safe harbor gene” refers to a gene or loci that a nucleic acid sequence can be inserted such that the sequence can integrate and function in a predictable manner (*e.g.*, express a protein of interest) without significant negative consequences to endogenous gene activity, or the promotion of cancer. In some embodiments, a safe harbor gene is also a loci or gene where an inserted nucleic acid sequence can be expressed efficiently and at higher levels than a non-safe harbor site.

[0060] As used herein, the term “gene delivery” means a process by which foreign DNA is transferred to host cells for applications of gene therapy.

[0061] As used herein, the term “terminal repeat” or “TR” includes any viral terminal repeat or synthetic sequence that comprises at least one minimal required origin of replication and a region comprising a palindrome hairpin structure. A Rep-binding sequence (“RBS”) (also referred to as RBE (Rep-binding element)) and a terminal resolution site (“TRS”) together constitute a “minimal required

origin of replication” and thus the TR comprises at least one RBS and at least one TRS. TRs that are the inverse complement of one another within a given stretch of polynucleotide sequence are typically each referred to as an “inverted terminal repeat” or “ITR”. In the context of a virus, ITRs mediate replication, virus packaging, integration and provirus rescue. As was unexpectedly found, TRs that are not inverse complements across their full length can still perform the traditional functions of ITRs, and thus the term ITR is used herein to refer to a TR in a ceDNA genome or ceDNA vector that is capable of mediating replication of ceDNA vector. It will be understood by one of ordinary skill in the art that in complex ceDNA vector configurations more than two ITRs or asymmetric ITR pairs may be present. The ITR can be an AAV ITR or a non-AAV ITR, or can be derived from an AAV ITR or a non-AAV ITR. For example, the ITR can be derived from the family *Parvoviridae*, which encompasses *Parvoviruses* and *Dependoviruses* (e.g., canine parvovirus, bovine parvovirus, mouse parvovirus, porcine parvovirus, human parvovirus B-19), or the SV40 hairpin that serves as the origin of SV40 replication can be used as an ITR, which can further be modified by truncation, substitution, deletion, insertion and/or addition. *Parvoviridae* family viruses consist of two subfamilies:

Parvovirinae, which infect vertebrates, and *Densovirinae*, which infect invertebrates.

Dependoparvoviruses include the viral family of the adeno-associated viruses (AAV) which are capable of replication in vertebrate hosts including, but not limited to, human, primate, bovine, canine, equine and ovine species. For convenience herein, an ITR located 5' to (upstream of) an expression cassette in a ceDNA vector is referred to as a “5' ITR” or a “left ITR”, and an ITR located 3' to (downstream of) an expression cassette in a ceDNA vector is referred to as a “3' ITR” or a “right ITR”.

[0062] A “wild-type ITR” or “WT-ITR” refers to the sequence of a naturally occurring ITR sequence in an AAV or other dependovirus that retains, e.g., Rep binding activity and Rep nicking ability. The nucleic acid sequence of a WT-ITR from any AAV serotype may slightly vary from the canonical naturally occurring sequence due to degeneracy of the genetic code or drift, and therefore WT-ITR sequences encompassed for use herein include WT-ITR sequences as result of naturally occurring changes taking place during the production process (e.g., a replication error).

[0063] As used herein, the term “substantially symmetrical WT-ITRs” or a “substantially symmetrical WT-ITR pair” refers to a pair of WT-ITRs within a single ceDNA genome or ceDNA vector that are both wild type ITRs that have an inverse complement sequence across their entire length. For example, an ITR can be considered to be a wild-type sequence, even if it has one or more nucleotides that deviate from the canonical naturally occurring sequence, so long as the changes do not affect the properties and overall three-dimensional structure of the sequence. In some aspects, the deviating nucleotides represent conservative sequence changes. As one non-limiting example, a sequence that has at least 95%, 96%, 97%, 98%, or 99% sequence identity to the canonical sequence (as measured, e.g., using BLAST at default settings), and also has a symmetrical three-dimensional spatial organization to the other WT-ITR such that their 3D structures are the same shape in

geometrical space. The substantially symmetrical WT-ITR has the same A, C-C' and B-B' loops in 3D space. A substantially symmetrical WT-ITR can be functionally confirmed as WT by determining that it has an operable Rep binding site (RBE or RBE') and terminal resolution site (trs) that pairs with the appropriate Rep protein. One can optionally test other functions, including transgene expression under permissive conditions.

[0064] As used herein, the phrases of “modified ITR” or “mod-ITR” or “mutant ITR” are used interchangeably herein and refer to an ITR that has a mutation in at least one or more nucleotides as compared to the WT-ITR from the same serotype. The mutation can result in a change in one or more of A, C, C', B, B' regions in the ITR, and can result in a change in the three-dimensional spatial organization (*i.e.*, its 3D structure in geometric space) as compared to the 3D spatial organization of a WT-ITR of the same serotype.

[0065] As used herein, the term “asymmetric ITRs” also referred to as “asymmetric ITR pairs” refers to a pair of ITRs within a single ceDNA genome or ceDNA vector that are not inverse complements across their full length. As one non-limiting example, an asymmetric ITR pair does not have a symmetrical three-dimensional spatial organization to their cognate ITR such that their 3D structures are different shapes in geometrical space. Stated differently, an asymmetrical ITR pair have the different overall geometric structure, *i.e.*, they have different organization of their A, C-C' and B-B' loops in 3D space (*e.g.*, one ITR may have a short C-C' arm and/or short B-B' arm as compared to the cognate ITR). The difference in sequence between the two ITRs may be due to one or more nucleotide addition, deletion, truncation, or point mutation. In one embodiment, one ITR of the asymmetric ITR pair may be a wild-type AAV ITR sequence and the other ITR a modified ITR as defined herein (*e.g.*, a non-wild-type or synthetic ITR sequence). In another embodiment, neither ITRs of the asymmetric ITR pair is a wild-type AAV sequence and the two ITRs are modified ITRs that have different shapes in geometrical space (*i.e.*, a different overall geometric structure). In some embodiments, one mod-ITRs of an asymmetric ITR pair can have a short C-C' arm and the other ITR can have a different modification (*e.g.*, a single arm, or a short B-B' arm etc.) such that they have different three-dimensional spatial organization as compared to the cognate asymmetric mod-ITR.

[0066] As used herein, the term “symmetric ITRs” refers to a pair of ITRs within a single ceDNA genome or ceDNA vector that are mutated or modified relative to wild-type dependoviral ITR sequences and are inverse complements across their full length. Neither ITRs are wild type ITR AAV2 sequences (*i.e.*, they are a modified ITR, also referred to as a mutant ITR), and can have a difference in sequence from the wild type ITR due to nucleotide addition, deletion, substitution, truncation, or point mutation. For convenience herein, an ITR located 5' to (upstream of) an expression cassette in a ceDNA vector is referred to as a “5' ITR” or a “left ITR”, and an ITR located 3' to (downstream of) an expression cassette in a ceDNA vector is referred to as a “3' ITR” or a “right ITR”.

[0067] As used herein, the terms “substantially symmetrical modified-ITRs” or a “substantially symmetrical mod-ITR pair” refers to a pair of modified-ITRs within a single ceDNA genome or ceDNA vector that are both that have an inverse complement sequence across their entire length. For example, the modified ITR can be considered substantially symmetrical, even if it has some nucleotide sequences that deviate from the inverse complement sequence so long as the changes do not affect the properties and overall shape. As one non-limiting example, a sequence that has at least 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% sequence identity to the canonical sequence (as measured using BLAST at default settings), and also has a symmetrical three-dimensional spatial organization to their cognate modified ITR such that their 3D structures are the same shape in geometrical space. Stated differently, a substantially symmetrical modified-ITR pair have the same A, C-C’ and B-B’ loops organized in 3D space. In some embodiments, the ITRs from a mod-ITR pair may have different reverse complement nucleotide sequences but still have the same symmetrical three-dimensional spatial organization – that is both ITRs have mutations that result in the same overall 3D shape. For example, one ITR (*e.g.*, 5’ ITR) in a mod-ITR pair can be from one serotype, and the other ITR (*e.g.*, 3’ ITR) can be from a different serotype, however, both can have the same corresponding mutation (*e.g.*, if the 5’ ITR has a deletion in the C region, the cognate modified 3’ ITR from a different serotype has a deletion at the corresponding position in the C’ region), such that the modified ITR pair has the same symmetrical three-dimensional spatial organization. In such embodiments, each ITR in a modified ITR pair can be from different serotypes (*e.g.*, AAV1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, and 12) such as the combination of AAV2 and AAV6, with the modification in one ITR reflected in the corresponding position in the cognate ITR from a different serotype. In one embodiment, a substantially symmetrical modified ITR pair refers to a pair of modified ITRs (mod-ITRs) so long as the difference in nucleotide sequences between the ITRs does not affect the properties or overall shape and they have substantially the same shape in 3D space. As a non-limiting example, a mod-ITR that has at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% sequence identity to the canonical mod-ITR as determined by standard means well known in the art such as BLAST (Basic Local Alignment Search Tool), or BLASTN at default settings, and also has a symmetrical three-dimensional spatial organization such that their 3D structure is the same shape in geometric space. A substantially symmetrical mod-ITR pair has the same A, C-C’ and B-B’ loops in 3D space, *e.g.*, if a modified ITR in a substantially symmetrical mod-ITR pair has a deletion of a C-C’ arm, then the cognate mod-ITR has the corresponding deletion of the C-C’ loop and also has a similar 3D structure of the remaining A and B-B’ loops in the same shape in geometric space of its cognate mod-ITR.

[0068] The term “flanking” refers to a relative position of one nucleic acid sequence with respect to another nucleic acid sequence. Generally, in the sequence ABC, B is flanked by A and C. The same is true for the arrangement AxBxC. Thus, a flanking sequence precedes or follows a flanked sequence

but need not be contiguous with, or immediately adjacent to the flanked sequence. In one embodiment, the term flanking refers to terminal repeats at each end of the linear duplex ceDNA vector.

[0069] As used herein, the term “ceDNA genome” refers to an expression cassette that further incorporates at least one inverted terminal repeat region. A ceDNA genome may further comprise one or more spacer regions. In some embodiments the ceDNA genome is incorporated as an intermolecular duplex polynucleotide of DNA into a plasmid or viral genome.

[0070] As used herein, the term “ceDNA spacer region” refers to an intervening sequence that separates functional elements in the ceDNA vector or ceDNA genome. In some embodiments, ceDNA spacer regions keep two functional elements at a desired distance for optimal functionality. In some embodiments, ceDNA spacer regions provide or add to the genetic stability of the ceDNA genome within *e.g.*, a plasmid or baculovirus. In some embodiments, ceDNA spacer regions facilitate ready genetic manipulation of the ceDNA genome by providing a convenient location for cloning sites and the like. For example, in certain aspects, an oligonucleotide “polylinker” containing several restriction endonuclease sites, or a non-open reading frame sequence designed to have no known protein (*e.g.*, transcription factor) binding sites can be positioned in the ceDNA genome to separate the *cis* – acting factors, *e.g.*, inserting a 6mer, 12mer, 18mer, 24mer, 48mer, 86mer, 176mer, etc. between the terminal resolution site and the upstream transcriptional regulatory element. Similarly, the spacer may be incorporated between the polyadenylation signal sequence and the 3’-terminal resolution site.

[0071] As used herein, the terms “Rep binding site,” “Rep binding element,” “RBE” and “RBS” are used interchangeably and refer to a binding site for Rep protein (*e.g.*, AAV Rep 78 or AAV Rep 68) which upon binding by a Rep protein permits the Rep protein to perform its site-specific endonuclease activity on the sequence incorporating the RBS. An RBS sequence and its inverse complement together form a single RBS. RBS sequences are known in the art, and include, for example, 5’-GCGCGCTCGCTCGCTC-3’ (SEQ ID NO: 60), an RBS sequence identified in AAV2. Any known RBS sequence may be used in the embodiments of the disclosure, including other known AAV RBS sequences and other naturally known or synthetic RBS sequences. Without being bound by theory it is thought that the nuclease domain of a Rep protein binds to the duplex nucleic acid sequence GCTC, and thus the two known AAV Rep proteins bind directly to and stably assemble on the duplex oligonucleotide, 5’-(GCGC)(GCTC)(GCTC)(GCTC)-3’ (SEQ ID NO: 60). In addition, soluble aggregated conformers (*i.e.*, undefined number of inter-associated Rep proteins) dissociate and bind to oligonucleotides that contain Rep binding sites. Each Rep protein interacts with both the nitrogenous bases and phosphodiester backbone on each strand. The interactions with the nitrogenous bases provide sequence specificity whereas the interactions with the phosphodiester backbone are non- or less- sequence specific and stabilize the protein-DNA complex.

[0072] As used herein, the terms “terminal resolution site” and “TRS” are used interchangeably herein and refer to a region at which Rep forms a tyrosine-phosphodiester bond with the 5’ thymidine generating a 3’ OH that serves as a substrate for DNA extension via a cellular DNA polymerase, *e.g.*,

DNA pol delta or DNA pol epsilon. Alternatively, the Rep-thymidine complex may participate in a coordinated ligation reaction. In some embodiments, a TRS minimally encompasses a non-base-paired thymidine. In some embodiments, the nicking efficiency of the TRS can be controlled at least in part by its distance within the same molecule from the RBS. When the acceptor substrate is the complementary ITR, then the resulting product is an intramolecular duplex. TRS sequences are known in the art, and include, for example, 5'-GGTTGA-3' (SEQ ID NO: 61), the hexanucleotide sequence identified in AAV2. Any known TRS sequence may be used in the embodiments of the disclosure, including other known AAV TRS sequences and other naturally known or synthetic TRS sequences such as AGTT (SEQ ID NO: 62), GGTTGG (SEQ ID NO: 63), AGTTGG (SEQ ID NO: 64), AGTTGA (SEQ ID NO: 65), and other motifs such as RRTTTR (SEQ ID NO: 66).

[0073] As used herein, the term “ceDNA” refers to capsid-free closed-ended linear double stranded (ds) duplex DNA for non-viral gene transfer, synthetic or otherwise. Detailed description of ceDNA is described in International application of PCT/US2017/020828, filed March 3, 2017, the entire contents of which are expressly incorporated herein by reference. Certain methods for the production of ceDNA comprising various inverted terminal repeat (ITR) sequences and configurations using cell-based methods are described in Example 1 of International applications PCT/US18/49996, filed September 7, 2018, and PCT/US2018/064242, filed December 6, 2018 each of which is incorporated herein in its entirety by reference. Certain methods for the production of synthetic ceDNA vectors comprising various ITR sequences and configurations are described, *e.g.*, in International application PCT/US2019/14122, filed January 18, 2019, the entire content of which is incorporated herein by reference. As used herein, the terms “ceDNA vector” and “ceDNA” are used interchangeably and refer to a closed-ended DNA vector comprising at least one terminal palindrome. In some embodiments, the ceDNA comprises two covalently-closed ends.

[0074] As used herein, the term “ceDNA-plasmid” refers to a plasmid that comprises a ceDNA genome as an intermolecular duplex.

[0075] As used herein, the term “ceDNA-bacmid” refers to an infectious baculovirus genome comprising a ceDNA genome as an intermolecular duplex that is capable of propagating in *E. coli* as a plasmid, and so can operate as a shuttle vector for baculovirus.

[0076] As used herein, the term “ceDNA-baculovirus” refers to a baculovirus that comprises a ceDNA genome as an intermolecular duplex within the baculovirus genome.

[0077] As used herein, the terms “ceDNA-baculovirus infected insect cell” and “ceDNA-BIIC” are used interchangeably, and refer to an invertebrate host cell (including, but not limited to an insect cell (*e.g.*, an Sf9 cell)) infected with a ceDNA-baculovirus.

[0078] As used herein, the term “closed-ended DNA vector” refers to a capsid-free DNA vector with at least one covalently closed end and where at least part of the vector has an intramolecular duplex structure.

[0079] As defined herein, “reporters” refer to proteins that can be used to provide detectable read-outs. Reporters generally produce a measurable signal such as fluorescence, color, or luminescence. Reporter protein coding sequences encode proteins whose presence in the cell or organism is readily observed. For example, fluorescent proteins cause a cell to fluoresce when excited with light of a particular wavelength, luciferases cause a cell to catalyze a reaction that produces light, and enzymes such as β -galactosidase convert a substrate to a colored product. Exemplary reporter polypeptides useful for experimental or diagnostic purposes include, but are not limited to β -lactamase, β -galactosidase (LacZ), alkaline phosphatase (AP), thymidine kinase (TK), green fluorescent protein (GFP) and other fluorescent proteins, chloramphenicol acetyltransferase (CAT), luciferase, and others well known in the art.

[0080] As used herein, the term “effector protein” refers to a polypeptide that provides a detectable read-out, either as, for example, a reporter polypeptide, or more appropriately, as a polypeptide that kills a cell, *e.g.*, a toxin, or an agent that renders a cell susceptible to killing with a chosen agent or lack thereof. Effector proteins include any protein or peptide that directly targets or damages the host cell’s DNA and/or RNA. For example, effector proteins can include, but are not limited to, a restriction endonuclease that targets a host cell DNA sequence (whether genomic or on an extrachromosomal element), a protease that degrades a polypeptide target necessary for cell survival, a DNA gyrase inhibitor, and a ribonuclease-type toxin. In some embodiments, the expression of an effector protein controlled by a synthetic biological circuit as described herein can participate as a factor in another synthetic biological circuit to thereby expand the range and complexity of a biological circuit system’s responsiveness.

[0081] Transcriptional regulators refer to transcriptional activators and repressors that either activate or repress transcription of a gene of interest, such as FIX. Promoters are regions of nucleic acid that initiate transcription of a particular gene. Transcriptional activators typically bind nearby to transcriptional promoters and recruit RNA polymerase to directly initiate transcription. Repressors bind to transcriptional promoters and sterically hinder transcriptional initiation by RNA polymerase. Other transcriptional regulators may serve as either an activator or a repressor depending on where they bind and cellular and environmental conditions. Non-limiting examples of transcriptional regulator classes include, but are not limited to homeodomain proteins, zinc-finger proteins, winged-helix (forkhead) proteins, and leucine-zipper proteins.

[0082] As used herein, a “repressor protein” or “inducer protein” is a protein that binds to a regulatory sequence element and represses or activates, respectively, the transcription of sequences operatively linked to the regulatory sequence element. Preferred repressor and inducer proteins as described herein are sensitive to the presence or absence of at least one input agent or environmental input. Preferred proteins as described herein are modular in form, comprising, for example, separable DNA-binding and input agent-binding or responsive elements or domains.

[0083] As used herein, “carrier” includes any and all solvents, dispersion media, vehicles, coatings, diluents, antibacterial and antifungal agents, isotonic and absorption delaying agents, buffers, carrier solutions, suspensions, colloids, and the like. The use of such media and agents for pharmaceutically active substances is well known in the art. Supplementary active ingredients can also be incorporated into the compositions. The phrase “pharmaceutically-acceptable” refers to molecular entities and compositions that do not produce a toxic, an allergic, or similar untoward reaction when administered to a host.

[0084] As used herein, an “input agent responsive domain” is a domain of a transcription factor that binds to or otherwise responds to a condition or input agent in a manner that renders a linked DNA binding fusion domain responsive to the presence of that condition or input. In one embodiment, the presence of the condition or input results in a conformational change in the input agent responsive domain, or in a protein to which it is fused, that modifies the transcription-modulating activity of the transcription factor.

[0085] The term “*in vivo*” refers to assays or processes that occur in or within an organism, such as a multicellular animal. In some of the aspects described herein, a method or use can be said to occur “*in vivo*” when a unicellular organism, such as a bacterium, is used. The term “*ex vivo*” refers to methods and uses that are performed using a living cell with an intact membrane that is outside of the body of a multicellular animal or plant, *e.g.*, explants, cultured cells, including primary cells and cell lines, transformed cell lines, and extracted tissue or cells, including blood cells, among others. The term “*in vitro*” refers to assays and methods that do not require the presence of a cell with an intact membrane, such as cellular extracts, and can refer to the introducing of a programmable synthetic biological circuit in a non-cellular system, such as a medium not comprising cells or cellular systems, such as cellular extracts.

[0086] The term “promoter,” as used herein, refers to any nucleic acid sequence that regulates the expression of another nucleic acid sequence by driving transcription of the nucleic acid sequence, which can be a heterologous target gene encoding a protein or an RNA. Promoters can be constitutive, inducible, repressible, tissue-specific, or any combination thereof. A promoter is a control region of a nucleic acid sequence at which initiation and rate of transcription of the remainder of a nucleic acid sequence are controlled. A promoter can also contain genetic elements at which regulatory proteins and molecules can bind, such as RNA polymerase and other transcription factors. In some embodiments of the aspects described herein, a promoter can drive the expression of a transcription factor that regulates the expression of the promoter itself. Within the promoter sequence will be found a transcription initiation site, as well as protein binding domains responsible for the binding of RNA polymerase. Eukaryotic promoters will often, but not always, contain “TATA” boxes and “CAT” boxes. Various promoters, including inducible promoters, may be used to drive the expression of transgenes in the ceDNA vectors disclosed herein. A promoter sequence may be bounded at its 3'

terminus by the transcription initiation site and extends upstream (5' direction) to include the minimum number of bases or elements necessary to initiate transcription at levels detectable above background.

[0087] The term “enhancer” as used herein refers to a cis-acting regulatory sequence (*e.g.*, 10-1,500 base pairs) that binds one or more proteins (*e.g.*, activator proteins, or transcription factor) to increase transcriptional activation of a nucleic acid sequence. Enhancers can be positioned up to 1,000,000 base pairs upstream of the gene start site or downstream of the gene start site that they regulate. An enhancer can be positioned within an intronic region, or in the exonic region of an unrelated gene.

[0088] A promoter can be said to drive expression or drive transcription of the nucleic acid sequence that it regulates. The phrases “operably linked,” “operatively positioned,” “operatively linked,” “under control,” and “under transcriptional control” indicate that a promoter is in a correct functional location and/or orientation in relation to a nucleic acid sequence it regulates to control transcriptional initiation and/or expression of that sequence. An “inverted promoter,” as used herein, refers to a promoter in which the nucleic acid sequence is in the reverse orientation, such that what was the coding strand is now the non-coding strand, and vice versa. Inverted promoter sequences can be used in various embodiments to regulate the state of a switch. In addition, in various embodiments, a promoter can be used in conjunction with an enhancer.

[0089] A promoter can be one naturally associated with a gene or sequence, as can be obtained by isolating the 5' non-coding sequences located upstream of the coding segment and/or exon of a given gene or sequence. Such a promoter can be referred to as “endogenous.” Similarly, in some embodiments, an enhancer can be one naturally associated with a nucleic acid sequence, located either downstream or upstream of that sequence.

[0090] In some embodiments, a coding nucleic acid segment is positioned under the control of a “recombinant promoter” or “heterologous promoter,” both of which refer to a promoter that is not normally associated with the encoded nucleic acid sequence it is operably linked to in its natural environment. A recombinant or heterologous enhancer refers to an enhancer not normally associated with a given nucleic acid sequence in its natural environment. Such promoters or enhancers can include promoters or enhancers of other genes; promoters or enhancers isolated from any other prokaryotic, viral, or eukaryotic cell; and synthetic promoters or enhancers that are not “naturally occurring,” *i.e.*, comprise different elements of different transcriptional regulatory regions, and/or mutations that alter expression through methods of genetic engineering that are known in the art. In addition to producing nucleic acid sequences of promoters and enhancers synthetically, promoter sequences can be produced using recombinant cloning and/or nucleic acid amplification technology, including PCR, in connection with the synthetic biological circuits and modules disclosed herein (see, *e.g.*, U.S. Pat. No. 4,683,202, U.S. Pat. No. 5,928,906, each incorporated herein by reference). Furthermore, it is contemplated that control sequences that direct transcription and/or expression of sequences within non-nuclear organelles such as mitochondria, chloroplasts, and the like, can be employed as well.

[0091] As described herein, an “inducible promoter” is one that is characterized by initiating or enhancing transcriptional activity when in the presence of, influenced by, or contacted by an inducer or inducing agent. An “inducer” or “inducing agent,” as defined herein, can be endogenous, or a normally exogenous compound or protein that is administered in such a way as to be active in inducing transcriptional activity from the inducible promoter. In some embodiments, the inducer or inducing agent, *i.e.*, a chemical, a compound or a protein, can itself be the result of transcription or expression of a nucleic acid sequence (*i.e.*, an inducer can be an inducer protein expressed by another component or module), which itself can be under the control of an inducible promoter. In some embodiments, an inducible promoter is induced in the absence of certain agents, such as a repressor. Examples of inducible promoters include but are not limited to, tetracycline, metallothionine, ecdysone, mammalian viruses (*e.g.*, the adenovirus late promoter; and the mouse mammary tumor virus long terminal repeat (MMTV-LTR)) and other steroid-responsive promoters, rapamycin responsive promoters and the like.

[0092] The terms “DNA regulatory sequences,” “control elements,” and “regulatory elements,” used interchangeably herein, refer to transcriptional and translational control sequences, such as promoters, enhancers, polyadenylation signals, terminators, protein degradation signals, and the like, that provide for and/or regulate transcription of a non-coding sequence (*e.g.*, DNA-targeting RNA) or a coding sequence (*e.g.*, site-directed modifying polypeptide, or Cas9/Csn1 polypeptide) and/or regulate translation of an encoded polypeptide.

[0093] The term “open reading frame (ORF)” as used herein is meant to refer to a sequence of several nucleotide triplets which may be translated into a peptide or protein. An open reading frame preferably contains a start codon, *i.e.* a combination of three subsequent nucleotides coding usually for the amino acid methionine (ATG), at its 5'-end and a subsequent region which usually exhibits a length which is a multiple of 3 nucleotides. An ORF is preferably terminated by a stop-codon (*e.g.*, TAA, TAG, TGA). Typically, this is the only stop-codon of the open reading frame. Thus, an open reading frame in the context of the present invention is preferably a nucleotide sequence, consisting of a number of nucleotides that may be divided by three, which starts with a start codon (*e.g.* ATG) and which preferably terminates with a stop codon (*e.g.*, TAA, TGA, or TAG). The open reading frame may be isolated or it may be incorporated in a longer nucleic acid sequence, for example in a ceDNA vector as described herein.

[0094] “Operably linked” refers to a juxtaposition wherein the components so described are in a relationship permitting them to function in their intended manner. For instance, a promoter is operably linked to a coding sequence if the promoter affects its transcription or expression. An “expression cassette” includes a DNA sequence that is operably linked to a promoter or other regulatory sequence sufficient to direct transcription of the transgene in the ceDNA vector. Suitable promoters include, for example, tissue specific promoters. Promoters can also be of AAV origin.

[0095] The term “subject” as used herein refers to a human or animal, to whom treatment, including prophylactic treatment, with the ceDNA vector according to the present disclosure, is

provided. Usually, the animal is a vertebrate such as, but not limited to a primate, rodent, domestic animal or game animal. Primates include but are not limited to, chimpanzees, cynomolgous monkeys, spider monkeys, and macaques, *e.g.*, Rhesus. Rodents include mice, rats, woodchucks, ferrets, rabbits and hamsters. Domestic and game animals include, but are not limited to, cows, horses, pigs, deer, bison, buffalo, feline species, *e.g.*, domestic cat, canine species, *e.g.*, dog, fox, wolf, avian species, *e.g.*, chicken, emu, ostrich, and fish, *e.g.*, trout, catfish and salmon. In certain embodiments of the aspects described herein, the subject is a mammal, *e.g.*, a primate or a human. A subject can be male or female. Additionally, a subject can be an infant or a child. In some embodiments, the subject can be a neonate or an unborn subject, *e.g.*, the subject is *in utero*. Preferably, the subject is a mammal. The mammal can be a human, non-human primate, mouse, rat, dog, cat, horse, or cow, but is not limited to these examples. Mammals other than humans can be advantageously used as subjects that represent animal models of diseases and disorders. In addition, the methods and compositions described herein can be used for domesticated animals and/or pets. A human subject can be of any age, gender, race or ethnic group, *e.g.*, Caucasian (white), Asian, African, black, African American, African European, Hispanic, Mideastern, etc. In some embodiments, the subject can be a patient or other subject in a clinical setting. In some embodiments, the subject is already undergoing treatment. In some embodiments, the subject is an embryo, a fetus, neonate, infant, child, adolescent, or adult. In some embodiments, the subject is a human fetus, human neonate, human infant, human child, human adolescent, or human adult. In some embodiments, the subject is an animal embryo, or non-human embryo or non-human primate embryo. In some embodiments, the subject is a human embryo.

[0096] As used herein, the term “host cell”, includes any cell type that is susceptible to transformation, transfection, transduction, and the like with a nucleic acid construct or ceDNA expression vector of the present disclosure. As non-limiting examples, a host cell can be an isolated primary cell, pluripotent stem cells, CD34⁺ cells), induced pluripotent stem cells, or any of a number of immortalized cell lines (*e.g.*, HepG2 cells). Alternatively, a host cell can be an *in situ* or *in vivo* cell in a tissue, organ or organism.

[0097] The term “exogenous” refers to a substance present in a cell other than its native source. The term “exogenous” when used herein can refer to a nucleic acid (*e.g.*, a nucleic acid encoding a polypeptide) or a polypeptide that has been introduced by a process involving the hand of man into a biological system such as a cell or organism in which it is not normally found and one wishes to introduce the nucleic acid or polypeptide into such a cell or organism. Alternatively, “exogenous” can refer to a nucleic acid or a polypeptide that has been introduced by a process involving the hand of man into a biological system such as a cell or organism in which it is found in relatively low amounts and one wishes to increase the amount of the nucleic acid or polypeptide in the cell or organism, *e.g.*, to create ectopic expression or levels. In contrast, the term “endogenous” refers to a substance that is native to the biological system or cell.

[0098] The term “sequence identity” refers to the relatedness between two nucleotide sequences. For purposes of the present disclosure, the degree of sequence identity between two deoxyribonucleotide sequences is determined using the Needleman-Wunsch algorithm (Needleman and Wunsch, 1970, *supra*) as implemented in the Needle program of the EMBOSS package (EMBOSS: The European Molecular Biology Open Software Suite, Rice et al., 2000, *supra*), preferably version 3.0.0 or later. The optional parameters used are gap open penalty of 10, gap extension penalty of 0.5, and the EDNAFULL (EMBOSS version of NCBI NUC4.4) substitution matrix. The output of Needle labeled “longest identity” (obtained using the -nobrief option) is used as the percent identity and is calculated as follows: (Identical Deoxyribonucleotides.times.100) / (Length of Alignment-Total Number of Gaps in Alignment). The length of the alignment is preferably at least 10 nucleotides, preferably at least 25 nucleotides more preferred at least 50 nucleotides and most preferred at least 100 nucleotides.

[0099] The term “homology” or “homologous” as used herein is defined as the percentage of nucleotide residues that are identical to the nucleotide residues in the corresponding sequence on the target chromosome, after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent sequence identity. Alignment for purposes of determining percent nucleotide sequence homology can be achieved in various ways that are within the skill in the art, for instance, using publicly available computer software such as BLAST, BLAST-2, ALIGN, ClustalW2 or Megalign (DNASTAR) software. Those skilled in the art can determine appropriate parameters for aligning sequences, including any algorithms needed to achieve maximal alignment over the full length of the sequences being compared. In some embodiments, a nucleic acid sequence (*e.g.*, DNA sequence), for example of a homology arm, is considered “homologous” when the sequence is at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or more, identical to the corresponding native or unedited nucleic acid sequence (*e.g.*, genomic sequence) of the host cell.

[00100] The term “heterologous,” as used herein, means a nucleotide or polypeptide sequence that is not found in the native nucleic acid or protein, respectively. A heterologous nucleic acid sequence may be linked to a naturally-occurring nucleic acid sequence (or a variant thereof) (*e.g.*, by genetic engineering) to generate a chimeric nucleotide sequence encoding a chimeric polypeptide. A heterologous nucleic acid sequence may be linked to a variant polypeptide (*e.g.*, by genetic engineering) to generate a nucleic acid sequence encoding a fusion variant polypeptide. Alternatively, the term “heterologous” may refer to a nucleic acid sequence which is not naturally present in a cell or subject.

[00101] A “vector” or “expression vector” is a replicon, such as plasmid, bacmid, phage, virus, virion, or cosmid, to which another DNA segment, *i.e.*, an “insert”, may be attached so as to bring about the replication of the attached segment in a cell. A vector can be a nucleic acid construct designed for delivery to a host cell or for transfer between different host cells. As used herein, a vector can be viral or non-viral in origin and/or in final form, however for the purpose of the present

disclosure, a “vector” generally refers to a cDNA vector, as that term is used herein. The term “vector” encompasses any genetic element that is capable of replication when associated with the proper control elements and that can transfer gene sequences to cells. In some embodiments, a vector can be an expression vector or recombinant vector.

[00102] As used herein, the term “expression vector” refers to a vector that directs expression of an RNA or polypeptide from sequences linked to transcriptional regulatory sequences on the vector. The sequences expressed will often, but not necessarily, be heterologous to the cell. An expression vector may comprise additional elements, for example, the expression vector may have two replication systems, thus allowing it to be maintained in two organisms, for example in human cells for expression and in a prokaryotic host for cloning and amplification. The term “expression” refers to the cellular processes involved in producing RNA and proteins and as appropriate, secreting proteins, including where applicable, but not limited to, for example, transcription, transcript processing, translation and protein folding, modification and processing. “Expression products” include RNA transcribed from a gene, and polypeptides obtained by translation of mRNA transcribed from a gene. The term “gene” means the nucleic acid sequence which is transcribed (DNA) to RNA *in vitro* or *in vivo* when operably linked to appropriate regulatory sequences. The gene may or may not include regions preceding and following the coding region, *e.g.*, 5’ untranslated (5’UTR) or “leader” sequences and 3’ UTR or “trailer” sequences, as well as intervening sequences (introns) between individual coding segments (exons).

[00103] By “recombinant vector” is meant a vector that includes a heterologous nucleic acid sequence, or “transgene” that is capable of expression *in vivo*. It should be understood that the vectors described herein can, in some embodiments, be combined with other suitable compositions and therapies. In some embodiments, the vector is episomal. The use of a suitable episomal vector provides a means of maintaining the nucleotide of interest in the subject in high copy number extra chromosomal DNA thereby eliminating potential effects of chromosomal integration.

[00104] The phrase “genetic disease” as used herein refers to a disease, partially or completely, directly or indirectly, caused by one or more abnormalities in the genome, especially a condition that is present from birth. The abnormality may be a mutation, an insertion or a deletion. The abnormality may affect the coding sequence of the gene or its regulatory sequence. According to some embodiments, the genetic disease is a result of a mutation in an FIX gene. According to some embodiments, the genetic disease is a result of decreased FIX protein expression. According to some embodiments, the genetic disease is hemophilia. According to some embodiments, the hemophilia is hemophilia B.

[00105] As used herein, “Clotting Factor IX (fIX; FIX)” is meant to refer to a vitamin K-dependent protein required for the efficient clotting of blood, which functions in coagulation as an activator of factor X. A concentration of about 1-5 µg/ml of fIX in the blood is considered in the normal range. Deficiency of FIX is associated with hemophilia B, and severe cases result when the concentration of

FIX is less than about 1% of the normal concentration of FIX (*i.e.* less than about 0.01-0.05 µg FIX per ml of blood).

[00106] As used herein, the terms, “administration,” “administering” and variants thereof refers to introducing a composition or agent (*e.g.*, a ceDNA as described herein) into a subject and includes concurrent and sequential introduction of one or more compositions or agents. “Administration” can refer, *e.g.*, to therapeutic, pharmacokinetic, diagnostic, research, placebo, and experimental methods. “Administration” also encompasses *in vitro* and *ex vivo* treatments. The introduction of a composition or agent into a subject is by any suitable route, including orally, pulmonarily, intranasally, parenterally (intravenously, intramuscularly, intraperitoneally, or subcutaneously), rectally, intralymphatically, intratumorally, or topically. Administration includes self-administration and the administration by another. Administration can be carried out by any suitable route. A suitable route of administration allows the composition or the agent to perform its intended function. For example, if a suitable route is intravenous, the composition is administered by introducing the composition or agent into a vein of the subject.

[00107] As used herein, the phrases “nucleic acid therapeutic”, “therapeutic nucleic acid” and “TNA” are used interchangeably and refer to any modality of therapeutic using nucleic acids as an active component of therapeutic agent to treat a disease or disorder. As used herein, these phrases refer to RNA-based therapeutics and DNA-based therapeutics. Non-limiting examples of RNA-based therapeutics include mRNA, antisense RNA and oligonucleotides, ribozymes, aptamers, interfering RNAs (RNAi), Dicer-substrate dsRNA, small hairpin RNA (shRNA), asymmetrical interfering RNA (aiRNA), microRNA (miRNA). Non-limiting examples of DNA-based therapeutics include minicircle DNA, minigene, viral DNA (*e.g.*, Lentiviral or AAV genome) or non-viral synthetic DNA vectors, closed-ended linear duplex DNA (ceDNA / CELiD), plasmids, bacmids, doggybone™ DNA vectors, minimalistic immunological-defined gene expression (MIDGE)-vector, nonviral ministring DNA vector (linear-covalently closed DNA vector), or dumbbell-shaped DNA minimal vector (“dumbbell DNA”). According to some embodiments, the therapeutic nucleic acid is a ceDNA.

[00108] As used herein, the term “immunosuppressant” refers to a group of small molecules, monoclonal antibodies or polypeptide antagonists that inhibits protein kinases, such as tyrosine kinases.

[00109] As used herein the term “therapeutic effect” refers to a consequence of treatment, the results of which are judged to be desirable and beneficial. A therapeutic effect can include, directly or indirectly, the arrest, reduction, or elimination of a disease manifestation. A therapeutic effect can also include, directly or indirectly, the arrest reduction or elimination of the progression of a disease manifestation.

[00110] For any therapeutic agent described herein therapeutically effective amount may be initially determined from preliminary *in vitro* studies and/or animal models. A therapeutically effective dose

may also be determined from human data. The applied dose may be adjusted based on the relative bioavailability and potency of the administered compound. Adjusting the dose to achieve maximal efficacy based on the methods described above and other well-known methods is within the capabilities of the ordinarily skilled artisan. General principles for determining therapeutic effectiveness, which may be found in Chapter 1 of Goodman and Gilman's *The Pharmacological Basis of Therapeutics*, 10th Edition, McGraw-Hill (New York) (2001), incorporated herein by reference, are summarized below.

[00111] Pharmacokinetic principles provide a basis for modifying a dosage regimen to obtain a desired degree of therapeutic efficacy with a minimum of unacceptable adverse effects. In situations where the drug's plasma concentration can be measured and related to therapeutic window, additional guidance for dosage modification can be obtained.

[00112] As used herein, the terms “treat,” “treating,” and/or “treatment” include abrogating, substantially inhibiting, slowing or reversing the progression of a condition, substantially ameliorating clinical symptoms of a condition, or substantially preventing the appearance of clinical symptoms of a condition, obtaining beneficial or desired clinical results. Treating further refers to accomplishing one or more of the following: (a) reducing the severity of the disorder; (b) limiting development of symptoms characteristic of the disorder(s) being treated; (c) limiting worsening of symptoms characteristic of the disorder(s) being treated; (d) limiting recurrence of the disorder(s) in patients that have previously had the disorder(s); and (e) limiting recurrence of symptoms in patients that were previously asymptomatic for the disorder(s).

[00113] Beneficial or desired clinical results, such as pharmacologic and/or physiologic effects include, but are not limited to, preventing the disease, disorder or condition from occurring in a subject that may be predisposed to the disease, disorder or condition but does not yet experience or exhibit symptoms of the disease (prophylactic treatment), alleviation of symptoms of the disease, disorder or condition, diminishment of extent of the disease, disorder or condition, stabilization (*i.e.*, not worsening) of the disease, disorder or condition, preventing spread of the disease, disorder or condition, delaying or slowing of the disease, disorder or condition progression, amelioration or palliation of the disease, disorder or condition, and combinations thereof, as well as prolonging survival as compared to expected survival if not receiving treatment.

[00114] As used herein, the term “increase,” “enhance,” “raise” (and like terms) generally refers to the act of increasing, either directly or indirectly, a concentration, level, function, activity, or behavior relative to the natural, expected, or average, or relative to a control condition.

[00115] As used herein, the term “suppress,” “decrease,” “interfere,” “inhibit” and/or “reduce” (and like terms) generally refers to the act of reducing, either directly or indirectly, a concentration, level, function, activity, or behavior relative to the natural, expected, or average, or relative to a control condition.

[00116] As used herein, a “control” is meant to refer to a reference standard. In some embodiments, the control is a negative control sample obtained from a healthy patient. In other embodiments, the control is a positive control sample obtained from a patient diagnosed with hemophilia. In still other embodiments, the control is a historical control or standard reference value or range of values (such as a previously tested control sample, such as a group of hemophilia A patients with known prognosis or outcome, or group of samples that represent baseline or normal values). A difference between a test sample and a control can be an increase or conversely a decrease. The difference can be a qualitative difference or a quantitative difference, for example a statistically significant difference. In some examples, a difference is an increase or decrease, relative to a control, of at least about 5%, such as at least about 10%, at least about 20%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 100%, at least about 150%, at least about 200%, at least about 250%, at least about 300%, at least about 350%, at least about 400%, at least about 500%, or greater than 500%.

[00117] As used herein the term “comprising” or “comprises” is used in reference to compositions, methods, and respective component(s) thereof, that are essential to the method or composition, yet open to the inclusion of unspecified elements, whether essential or not.

[00118] As used herein the term “consisting essentially of” refers to those elements required for a given embodiment. The term permits the presence of elements that do not materially affect the basic and novel or functional characteristic(s) of that embodiment. The use of “comprising” indicates inclusion rather than limitation.

[00119] The term “consisting of” refers to compositions, methods, and respective components thereof as described herein, which are exclusive of any element not recited in that description of the embodiment.

[00120] As used in this specification and the appended claims, the singular forms “a,” “an,” and “the” include plural references unless the context clearly dictates otherwise. Thus, for example, references to “the method” includes one or more methods, and/or steps of the type described herein and/or which will become apparent to those persons skilled in the art upon reading this disclosure and so forth. Similarly, the word “or” is intended to include “and” unless the context clearly indicates otherwise. Although methods and materials similar or equivalent to those described herein can be used in the practice or testing of this disclosure, suitable methods and materials are described below. The abbreviation, “*e.g.*” is derived from the Latin *exempli gratia*, and is used herein to indicate a non-limiting example. Thus, the abbreviation “*e.g.*” is synonymous with the term “for example.”

[00121] Other than in the operating examples, or where otherwise indicated, all numbers expressing quantities of ingredients or reaction conditions used herein should be understood as modified in all instances by the term “about.” The term “about” when used in connection with percentages can mean $\pm 1\%$. The present disclosure is further explained in detail by the following examples, but the scope of the disclosure should not be limited thereto.

[00122] Groupings of alternative elements or embodiments of the disclosure disclosed herein are not to be construed as limitations. Each group member can be referred to and claimed individually or in any combination with other members of the group or other elements found herein. One or more members of a group can be included in, or deleted from, a group for reasons of convenience and/or patentability. When any such inclusion or deletion occurs, the specification is herein deemed to contain the group as modified thus fulfilling the written description of all Markush groups used in the appended claims.

[00123] In some embodiments of any of the aspects, the disclosure described herein does not concern a process for cloning human beings, processes for modifying the germ line genetic identity of human beings, uses of human embryos for industrial or commercial purposes or processes for modifying the genetic identity of animals which are likely to cause them suffering without any substantial medical benefit to man or animal, and also animals resulting from such processes.

[00124] Other terms are defined herein within the description of the various aspects of the disclosure.

[00125] All patents and other publications; including literature references, issued patents, published patent applications, and co-pending patent applications; cited throughout this application are expressly incorporated herein by reference for the purpose of describing and disclosing, for example, the methodologies described in such publications that might be used in connection with the technology described herein. These publications are provided solely for their disclosure prior to the filing date of the present application. Nothing in this regard should be construed as an admission that the inventors are not entitled to antedate such disclosure by virtue of prior disclosure or for any other reason. All statements as to the date or representation as to the contents of these documents is based on the information available to the applicants and does not constitute any admission as to the correctness of the dates or contents of these documents.

[00126] The description of embodiments of the disclosure is not intended to be exhaustive or to limit the disclosure to the precise form disclosed. While specific embodiments of, and examples for, the disclosure are described herein for illustrative purposes, various equivalent modifications are possible within the scope of the disclosure, as those skilled in the relevant art will recognize. For example, while method steps or functions are presented in a given order, alternative embodiments may perform functions in a different order, or functions may be performed substantially concurrently. The teachings of the disclosure provided herein can be applied to other procedures or methods as appropriate. The various embodiments described herein can be combined to provide further embodiments. Aspects of the disclosure can be modified, if necessary, to employ the compositions, functions and concepts of the above references and application to provide yet further embodiments of the disclosure. Moreover, due to biological functional equivalency considerations, some changes can be made in protein structure without affecting the biological or chemical action in kind or amount.

These and other changes can be made to the disclosure in light of the detailed description. All such modifications are intended to be included within the scope of the appended claims.

[00127] Specific elements of any of the foregoing embodiments can be combined or substituted for elements in other embodiments. Furthermore, while advantages associated with certain embodiments of the disclosure have been described in the context of these embodiments, other embodiments may also exhibit such advantages, and not all embodiments need necessarily exhibit such advantages to fall within the scope of the disclosure.

[00128] The technology described herein is further illustrated by the following examples which in no way should be construed as being further limiting. It should be understood that this disclosure is not limited to the particular methodology, protocols, and reagents, etc., described herein and as such can vary. The terminology used herein is for the purpose of describing particular embodiments only, and is not intended to limit the scope of the present disclosure, which is defined solely by the claims.

II. Expression of a FIX Protein from a ceDNA vector

[00129] The technology described herein is directed in general to the expression and/or production of FIX protein in a cell from a non-viral DNA vector, *e.g.*, a ceDNA vector as described herein. ceDNA vectors for expression of FIX protein are described herein in the section entitled “ceDNA vectors in general”. In particular, ceDNA vectors for expression of FIX protein comprise a pair of ITRs (*e.g.*, symmetric or asymmetric as described herein) and between the ITR pair, a nucleic acid encoding an FIX protein, as described herein, operatively linked to a promoter or regulatory sequence. A distinct advantage of ceDNA vectors for expression of FIX protein over traditional AAV vectors, and even lentiviral vectors, is that there is no size constraint for the nucleic acid sequences encoding a desired protein. Thus, even a full length 6.8kb FIX protein can be expressed from a single ceDNA vector. Thus, the ceDNA vectors described herein can be used to express a therapeutic FIX protein in a subject in need thereof, *e.g.*, a subject with hemophilia B.

[00130] As one will appreciate, the ceDNA vector technologies described herein can be adapted to any level of complexity or can be used in a modular fashion, where expression of different components of a FIX protein can be controlled in an independent manner. For example, it is specifically contemplated that the ceDNA vector technologies designed herein can be as simple as using a single ceDNA vector to express a single gene sequence (*e.g.*, a FIX protein) or can be as complex as using multiple ceDNA vectors, where each vector expresses multiple FIX proteins or associated co-factors or accessory proteins that are each independently controlled by different promoters. The following embodiments are specifically contemplated herein and can be adapted by one of skill in the art as desired.

[00131] In one embodiment, a single ceDNA vector can be used to express a single component of a FIX protein. Alternatively, a single ceDNA vector can be used to express multiple components (*e.g.*, at least 2) of a FIX protein under the control of a single promoter (*e.g.*, a strong promoter), optionally using an IRES sequence(s) to ensure appropriate expression of each of the components, *e.g.*, co-factors or accessory proteins.

[00132] Also contemplated herein, in another embodiment, is a single ceDNA vector comprising at least two inserts (*e.g.*, expressing a heavy chain or light chain), where the expression of each insert is under the control of its own promoter. The promoters can include multiple copies of the same promoter, multiple different promoters, or any combination thereof. As one of skill in the art will appreciate, it is often desirable to express components of a FIX protein at different expression levels, thus controlling the stoichiometry of the individual components expressed to ensure efficient a FIX protein folding and combination in the cell.

[00133] Additional variations of ceDNA vector technologies can be envisioned by one of skill in the art or can be adapted from protein production methods using conventional vectors.

A. Factor IX (FIX) expression

[00134] In some embodiments, a transgene encoding the FIX protein can also encode a secretory sequence so that the a FIX protein is directed to the Golgi Apparatus and Endoplasmic Reticulum whence a FIX protein will be folded into the correct conformation by chaperone molecules as it passes through the ER and out of the cell. Exemplary secretory sequences include, but are not limited to VH-02 (SEQ ID NO: 88) and VK-A26 (SEQ ID NO: 89) and Igk signal sequence (SEQ ID NO: 126), as well as a Gluc secretory signal that allows the tagged protein to be secreted out of the cytosol (SEQ ID NO: 188), TMD-ST secretory sequence, that directs the tagged protein to the golgi (SEQ ID NO: 189).

[00135] Regulatory switches can also be used to fine tune the expression of the FIX protein so that the FIX protein is expressed as desired, including but not limited to expression of the FIX protein at a desired expression level or amount, or alternatively, when there is the presence or absence of particular signal, including a cellular signaling event. For instance, as described herein, expression of the FIX protein from the ceDNA vector can be turned on or turned off when a particular condition occurs, as described herein in the section entitled Regulatory Switches.

[00136] For example, and for illustration purposes only, FIX proteins can be used to turn off undesired reaction, such as too high a level of production of the FIX protein. The FIX gene can contain a signal peptide marker to bring the FIX protein to the desired cell. However, in either situation it can be desirable to regulate the expression of the FIX protein. ceDNA vectors readily accommodate the use of regulatory switches.

[00137] A distinct advantage of ceDNA vectors over traditional AAV vectors, and even lentiviral vectors, is that there is no size constraint for the nucleic acid sequences encoding the FIX protein. Thus, even a full-length FIX, as well as optionally any co-factors or assessor proteins can be expressed from a single ceDNA vector. In addition, depending on the necessary stiochemistry one can express multiple segments of the same FIX protein, and can use same or different promoters, and can also use regulatory switches to fine tune expression of each region. For example, as shown in the Examples, a ceDNA vector that comprises a dual promoter system can be used, so that a different promoter is used for each domain of the FIX protein. Use of a ceDNA plasmid to produce the FIX protein can include a unique combination of promoters for expression of the domains of the FIX protein that results in the

proper ratios of each domain for the formation of functional FIX protein. Accordingly, in some embodiments, a ceDNA vector can be used to express different regions of FIX protein separately (*e.g.*, under control of a different promoter).

[00138] In another embodiment, the FIX protein expressed from the ceDNA vectors further comprises an additional functionality, such as fluorescence, enzyme activity, secretion signal or immune cell activator.

[00139] In some embodiments, the ceDNA encoding the FIX protein can further comprise a linker domain, for example. As used herein “linker domain” refers to an oligo- or polypeptide region from about 2 to 100 amino acids in length, which links together any of the domains/regions of the FIX protein as described herein. In some embodiment, linkers can include or be composed of flexible residues such as glycine and serine so that the adjacent protein domains are free to move relative to one another. Longer linkers may be used when it is desirable to ensure that two adjacent domains do not sterically interfere with one another. Linkers may be cleavable or non-cleavable. Examples of cleavable linkers include 2A linkers (for example T2A), 2A-like linkers or functional equivalents thereof and combinations thereof. The linker can be a linker region is T2A derived from *Thosea asigna* virus.

[00140] It is well within the abilities of one of skill in the art to take a known and/or publically available protein sequence of *e.g.*, the FIX etc., and reverse engineer a cDNA sequence to encode such a protein. The cDNA can then be codon optimized to match the intended host cell and inserted into a ceDNA vector as described herein.

B. ceDNA vectors expressing FIX Protein

[00141] A ceDNA vector for expression of FIX protein having one or more sequences encoding a desired FIX can comprise regulatory sequences such as promoters, secretion signals, polyA regions, and enhancers. At a minimum, a ceDNA vector comprises one or more nucleic acid sequences encoding a FIX protein.

[00142] In order to achieve highly efficient and accurate FIX protein assembly, it is specifically contemplated in some embodiments that the FIX protein comprise an endoplasmic reticulum ER leader sequence to direct it to the ER, where protein folding occurs. For example, a sequence that directs the expressed protein(s) to the ER for folding.

[00143] In some embodiments, a cellular or extracellular localization signal (*e.g.*, secretory signal, nuclear localization signal, mitochondrial localization signal etc.) is comprised in the ceDNA vector to direct the secretion or desired subcellular localization of FIX such that the FIX protein can bind to intracellular target(s) (*e.g.*, an intrabody) or extracellular target(s).

[00144] In some embodiments, a ceDNA vector for expression of FIX protein as described herein permits the assembly and expression of any desired FIX protein in a modular fashion. As used herein, the term “modular” refers to elements in a ceDNA expressing plasmid that can be readily removed from the construct. For example, modular elements in a ceDNA-generating plasmid comprise unique

pairs of restriction sites flanking each element within the construct, enabling the exclusive manipulation of individual elements (see *e.g.*, **FIGs. 1A-1G**). Thus, the ceDNA vector platform can permit the expression and assembly of any desired FIX protein configuration. Provided herein in various embodiments are ceDNA plasmid vectors that can reduce and/or minimize the amount of manipulation required to assemble a desired ceDNA vector encoding FIX protein.

C. Exemplary FIX Proteins expressed by ceDNA vectors

[00145] In particular, a ceDNA vector for expression of FIX protein as disclosed herein can encode, for example, but is not limited to, FIX proteins, as well as variants, and/or active fragments thereof, for use in the treatment, prophylaxis, and/or amelioration of one or more symptoms of hemophilia B. In one aspect, the hemophilia B is a human hemophilia B.

(i) FIX therapeutic proteins and fragments thereof

[00146] Essentially any version of the FIX therapeutic protein or fragment thereof (*e.g.*, functional fragment) can be encoded by and expressed in and from a ceDNA vector as described herein. One of skill in the art will understand that an FIX therapeutic protein includes all splice variants and orthologs of the FIX protein. A FIX therapeutic protein includes intact molecules as well as fragments (*e.g.*, functional fragments) thereof.

Factor IX (FIX)

[00147] Factor IX (or Christmas factor) (EC 3.4.21.22) is one of the serine proteases of the coagulation system; it belongs to peptidase family S1. Deficiency of this protein causes hemophilia B. Factor IX is produced as a zymogen, an inactive precursor. It is processed to remove the signal peptide, glycosylated and then cleaved by factor XIa (of the contact pathway) or factor VIIa (of the tissue factor pathway) to produce a two-chain form where the chains are linked by a disulfide bridge. When activated into factor IXa, in the presence of Ca²⁺, membrane phospholipids, and a Factor VIII cofactor, it hydrolyses one arginine-isoleucine bond in factor X to form factor Xa. Factor IX is Vitamin-K dependent. Factor IX is inhibited by antithrombin.

[00148] The Factor IX gene or protein can also be referred to as F9, Coagulation Factor IX, Plasma Thromboplastin Component, Plasma Thromboplastic Component, Christmas Factor, EC 3.4.21.22, PTC, Christmas Disease, Factor IX F9, Hemophilia B, Factor IX, EC 3.4.21, Factor 9, F9 P22, THPH8, HEMB, FIX, or P19.

[00149] The gene of human FIX lies in the X chromosome, has 8 exons, and spans 33.5 Kb. Factor IX is produced in the liver, and the inactive precursor protein is processed in the endoplasmic reticulum and Golgi, where it undergoes multiple post-translational modifications and is secreted into the bloodstream upon proteolytic cleavage of the propeptide.

[00150] Expression of Factor IX mRNA occurs primarily in the liver. Additional tissues can express Factor IX mRNA, including bone marrow, whole blood, lymph nodes, thymus, brain, cerebral cortex, cerebellum, retina, spinal cord, tibial nerve, heart, artery, smooth muscle, skeletal muscle, small

intestine, colon, adipocytes, kidney, lung, spleen, stomach, esophagus, bladder, pancreas, thyroid, salivary gland, adrenal gland, pituitary gland, breast, skin, ovary, uterus, placenta, prostate, and testis.

[00151] Factor IX protein is predominately expressed in the serum, plasma, and monocytes. Factor IX protein expression can also be detected tissues throughout the body, including but not limited to the tonsil, bone marrow mesenchymal stem cells, spinal cord, heart, colon muscle, oral epithelium, esophagus, stomach, cardia, colon, rectum, liver, fetal liver, kidney, spleen, synovial fluid, vitreous humor, salivary gland, thyroid gland, adrenal gland, breast, pancreas, islet of Langerhans, gallbladder, prostate, urine, urinary bladder, skin, placenta, uterus, cervix, ovary, testis, and seminal vesicle.

[00152] There are at least two known mRNA variants each encoding one protein isoform of Factor IX. Variant 1 represents the longer transcript and encodes the longer isoform 1. Variant 2 lacks an alternate in-frame exon in the 5' coding region, compared to variant 1. It encodes isoform 2, which is shorter than isoform 1. Isoform 2 may undergo proteolytic processing similar to isoform 1.

Representative sequence identifiers for mRNA variants 1 and 2 and protein isoforms 1 and 2 are shown below:

Homo sapiens coagulation factor IX (F9), transcript variant 1, mRNA (NCBI Reference Sequence: NM_000133.3) 1386 bp (SEQ ID NO: 377)

Homo sapiens coagulation factor IX isoform 1 preproprotein (NCBI Reference Sequence: NP_000124.1) 461 amino acids (SEQ ID NO: 378)

Homo sapiens coagulation factor IX (F9), transcript variant 2, mRNA (NCBI Reference Sequence: NM_001313913.1) 2688 bp (SEQ ID NO: 379)

Homo sapiens coagulation factor IX isoform 2 precursor (NCBI Reference Sequence: NP_001300842.1) 423 amino acids (SEQ ID NO: 380)

[00153] A distinct advantage of ceDNA vectors over traditional AAV vectors, and even lentiviral vectors, is that there is no size constraint for the nucleic acid sequences encoding a desired protein. Thus, multiple full length FIX therapeutic proteins can be expressed from a single ceDNA vector.

[00154] Expression of FIX therapeutic protein or fragment thereof from a ceDNA vector can be achieved both spatially and temporally using one or more inducible or repressible promoters, as known in the art or described herein, including regulatory switches as described herein.

[00155] In one embodiment, FIX therapeutic protein is an “therapeutic protein variant,” which refers to the FIX therapeutic protein having an altered amino acid sequence, composition or structure as compared to its corresponding native FIX therapeutic protein. In one embodiment, FIX is a functional version (*e.g.*, wild type). It may also be useful to express a mutant version of FIX protein such as a point mutation or deletion mutation that leads to hemophilia B, *e.g.*, for an animal model of the disease and/or for assessing drugs for hemophilia B. Delivery of mutant or modified FIX proteins to a cell or animal model system can be done in order to generate a disease model. Such a cellular or animal model can be used for research and/or drug screening. FIX therapeutic protein expressed from the ceDNA vectors

may further comprise a sequence/moiety that confers an additional functionality, such as fluorescence, enzyme activity, or secretion signal. In one embodiment, an FIX therapeutic protein variant comprises a non-native tag sequence for identification (*e.g.*, an immunotag) to allow it to be distinguished from endogenous FIX therapeutic protein in a recipient host cell.

[00156] It is well within the abilities of one of skill in the art to take a known and/or publically available protein sequence of *e.g.*, FIX therapeutic protein and reverse engineer a cDNA sequence to encode such a protein. The cDNA can then be codon optimized to match the intended host cell and inserted into a ceDNA vector as described herein.

[00157] In one embodiment, the FIX therapeutic protein encoding sequence can be derived from an existing host cell or cell line, for example, by reverse transcribing mRNA obtained from the host and amplifying the sequence using PCR.

(ii) *FIX therapeutic protein expressing ceDNA vectors*

[00158] A ceDNA vector having one or more sequences encoding a desired FIX therapeutic protein can comprise regulatory sequences such as promoters (*e.g.*, see Table 7), secretion signals, polyA regions (*e.g.*, see Table 10), and enhancers (*e.g.*, see Table 8). At a minimum, a ceDNA vector comprises one or more nucleic acid sequences encoding the FIX therapeutic protein or functional fragment thereof. Exemplary cassette inserts for generating ceDNA vectors encoding the FIX therapeutic proteins are depicted in **Figures 1A-1G**. In one embodiment, the ceDNA vector comprises an FIX sequence listed in **Table 1** herein.

[00159] **Table 1:** Exemplary FIX sequences for treatment of hemophilia B

Description	Length	Reference	SEQ ID NO:
Codon optimized hFIX	1386	Nathwani et al., Blood (2006) 107(7):2653-2661.	381
hFIX Exons and first intron derived from SPK9001	2824	US Patent Publication No. 20160375110A1	382
hFIX Exons only derived from SPK9001	1386	US Patent Publication No. 20160375110A1	383
Endogenous hFIX cDNA	1386	NG_007994.1	384
JCat optimized Padua Factor IX ORF containing the G338L mutation	1386		385
Jcat optimized Wild type, Human Factor IX ORF	1386		386
Jcat optimized Wild type, Canine Factor IX ORF	1359		387
Human FIX ORF padua variant (codon optimized) 1386bp	1386		388
Human FIX ORF (codon optimized) 1386bp	1386		389
Murine Factor IX cDNA	1419		390
Murine Factor IX cDNA with GGGGS linker (SEQ ID NO: 391) and 6xHis Tag (SEQ ID NO: 392)	1452		393
Murine Factor IX cDNA with GGGGS linker (SEQ ID NO: 391) and Myc Tag	1464		394
Murine Factor IX cDNA with Padua Mutation (R338L)	1419		395

Murine Factor IX cDNA with Padua Mutation (R338L) and GGGGS linker (SEQ ID NO: 391) and 6xHis Tag (SEQ ID NO: 392)	1452		396
CpG-free human FIX Padua (G388L) COOL optimized	1386		397
Human Factor IX, WT. Codon optimized.	1194		398
WT Murine Factor IX; mouse codon optimized	1416		399
WT Murine Factor IX with GGGGS linker (SEQ ID NO: 391) and 6x His Tag (SEQ ID NO: 392); mouse codon optimized	1449		400
WT Murine Factor IX with GGGGS linker (SEQ ID NO: 391) and Myc Tag ; mouse codon optimized	1461		401
Padua Variant Murine Factor IX; mouse codon optimized	1416		402
Padua Variant Murine Factor IX with GGGGS linker (SEQ ID NO: 391) and 6x His Tag (SEQ ID NO: 392); mouse codon optimized	1449		403

(iii) *FIX therapeutic proteins and uses thereof for the treatment of hemophilia B*

[00160] The ceDNA vectors described herein can be used to deliver therapeutic FIX proteins for treatment of hemophilia B associated with inappropriate expression of the FIX protein and/or mutations within the FIX proteins.

[00161] ceDNA vectors as described herein can be used to express any desired FIX therapeutic protein. Exemplary therapeutic FIX therapeutic proteins include, but are not limited to any FIX protein expressed by the sequences as set forth in **Table 1** herein.

[00162] In one embodiment, the expressed FIX therapeutic protein is functional for the treatment of a Hemophilia B. In some embodiments, FIX therapeutic protein does not cause an immune system reaction.

[00163] In another embodiment, the ceDNA vectors encoding FIX therapeutic protein or fragment thereof (*e.g.*, functional fragment) can be used to generate a chimeric protein. Thus, it is specifically contemplated herein that a ceDNA vector expressing a chimeric protein can be administered to *e.g.*, to any one or more tissues selected from: liver, kidneys, gallbladder, prostate, adrenal gland. In some embodiments, when a ceDNA vector expressing FIX is administered to an infant, or administered to a subject in utero, one can administer a ceDNA vector expressing FIX to any one or more tissues selected from: liver, adrenal gland, heart, intestine, lung, and stomach, or to a liver stem cell precursor thereof for the *in vivo* or *ex vivo* treatment of hemophilia B.

[00164] Hemophilia B

[00165] Hemophilia B is a blood clotting disorder that causes easy bruising and bleeding due to an inherited mutation of the gene for factor IX, and resulting in a deficiency of factor IX. Hemophilia B is inherited as an X-linked recessive trait. Current treatments to prevent bleeding in people with hemophilia B involves intravenous infusion of factor IX and/or blood transfusions.

[00166] There are many complications related to treatment of hemophilia B. In children, an easily accessible intravenous port can be inserted to minimize frequent traumatic intravenous cannulation. However, these ports are associated with high infection rate and a risk of clots forming at the tip of the catheter, rendering it useless. Viral infections can be common in hemophiliacs due to frequent blood transfusions which put patients at risk of acquiring blood borne infections, such as HIV, hepatitis B and hepatitis C. Prion infections can also be transmitted by blood transfusions.

[00167] In some cases, mutations in the promoter region of the FIX gene result in the less severe hemophilia B Leiden, characterized as a nearly complete absence of FIX in childhood and steady increase in the level of endogenous FIX during puberty to the near-normal values.

[00168] Coagulation Cascade

[00169] Coagulation, also known as clotting, is the process by which blood changes from a liquid to a gel, forming a blood clot. It potentially results in hemostasis, the cessation of blood loss from a damaged vessel, followed by repair. The mechanism of coagulation involves activation, adhesion and aggregation of platelets along with deposition and maturation of fibrin. Disorders of coagulation are disease states which can result in bleeding (hemorrhage or bruising) or obstructive clotting (thrombosis).

[00170] Coagulation begins almost instantly after an injury to the blood vessel has damaged the endothelium lining the blood vessel. Exposure of blood to the subendothelial space initiates two processes: changes in platelets, and the exposure of subendothelial tissue factor to plasma Factor VII, which ultimately leads to fibrin formation. Platelets immediately form a plug at the site of injury; this is called primary hemostasis. Secondary hemostasis occurs simultaneously: additional coagulation factors or clotting factors beyond Factor VII (including Factor VIII) respond in a complex cascade to form fibrin strands, which strengthen the platelet plug.

[00171] The coagulation cascade of secondary hemostasis has two initial pathways which lead to fibrin formation. These are the contact activation pathway (also known as the intrinsic pathway), and the tissue factor pathway (also known as the extrinsic pathway), which both lead to the same fundamental reactions that produce fibrin. The primary pathway for the initiation of blood coagulation is the tissue factor (extrinsic) pathway. The pathways are a series of reactions, in which a zymogen (inactive enzyme precursor) of a serine protease and its glycoprotein co-factor are activated to become active components that then catalyze the next reaction in the cascade, ultimately resulting in cross-linked fibrin. Coagulation factors are generally indicated by Roman numerals, with a lowercase a appended to indicate an active form.

[00172] The coagulation factors are generally serine proteases (enzymes), which act by cleaving downstream proteins. The exceptions are tissue factor, FV, FVIII, FXIII. Tissue factor, FV and FVIII are glycoproteins, and Factor XIII is a transglutaminase. The coagulation factors circulate as inactive zymogens. The coagulation cascade is therefore classically divided into three pathways. The tissue

factor and contact activation pathways both activate the "final common pathway" of factor X, thrombin and fibrin.

[00173] The main role of the tissue factor (extrinsic) pathway is to generate a "thrombin burst", a process by which thrombin, the most important constituent of the coagulation cascade in terms of its feedback activation roles, is released very rapidly. FVIIa circulates in a higher amount than any other activated coagulation factor. The process includes the following steps:

[00174] Step 1: Following damage to the blood vessel, FVII leaves the circulation and comes into contact with tissue factor (TF) expressed on tissue-factor-bearing cells (stromal fibroblasts and leukocytes), forming an activated complex (TF-FVIIa).

[00175] Step 2: TF-FVIIa activates FIX and FX.

[00176] Step 3: FVII is itself activated by thrombin, FXIa, FXII and FXa.

[00177] Step 4: The activation of FX (to form FXa) by TF-FVIIa is almost immediately inhibited by tissue factor pathway inhibitor (TFPI).

[00178] Step 5: FXa and its co-factor FVa form the prothrombinase complex, which activates prothrombin to thrombin.

[00179] Step 6: Thrombin then activates other components of the coagulation cascade, including FV and FVIII (which forms a complex with FIX), and activates and releases FVIII from being bound to von Willebrand factor (vWF).

[00180] Step 7: FVIIIa is the co-factor of FIXa, and together they form the "tenase" complex, which activates FX; and so the cycle continues.

[00181] The contact activation (intrinsic) pathway begins with formation of the primary complex on collagen by high-molecular-weight kininogen (HMWK), prekallikrein, and FXII (Hageman factor). Prekallikrein is converted to kallikrein and FXII becomes FXIIa. FXIIa converts FXI into FXIa. Factor XIa activates FIX, which with its co-factor FVIIIa form the tenase complex, which activates FX to FXa. The minor role that the contact activation pathway has in initiating clot formation can be illustrated by the fact that patients with severe deficiencies of FXII, HMWK, and prekallikrein do not have a bleeding disorder. Instead, contact activation system is more involved in inflammation, and innate immunity.

[00182] The final common pathway shared by the intrinsic and extrinsic coagulation pathways involves the conversion of prothrombin into thrombin and fibrinogen into fibrin. Thrombin has a large array of functions, not only the conversion of fibrinogen to fibrin, the building block of a hemostatic plug. In addition, it is the most important platelet activator and on top of that it activates Factors VIII

and V and their inhibitor protein C (in the presence of thrombomodulin), and it activates Factor XIII, which forms covalent bonds that crosslink the fibrin polymers that form from activated monomers.

[00183] Following activation by the contact factor or tissue factor pathways, the coagulation cascade is maintained in a prothrombotic state by the continued activation of FVIII and FIX to form the tenase complex, until it is down-regulated by the anticoagulant pathways.

[00184] The methods comprise administering to the subject an effective amount of a composition comprising a ceDNA vector encoding the FIX therapeutic protein or fragment thereof (*e.g.*, functional fragment) as described herein. As will be appreciated by a skilled practitioner, the term “effective amount” refers to the amount of the ceDNA composition administered that results in expression of the protein in a “therapeutically effective amount” for the treatment of a disease or disorder.

[00185] The dosage ranges for the composition comprising a ceDNA vector encoding the FIX therapeutic protein or fragment thereof (*e.g.*, functional fragment) depends upon the potency (*e.g.*, efficiency of the promoter), and includes amounts large enough to produce the desired effect, *e.g.*, expression of the desired FIX therapeutic protein, for the treatment of hemophilia B. The dosage should not be so large as to cause unacceptable adverse side effects. Generally, the dosage will vary with the particular characteristics of the ceDNA vector, expression efficiency and with the age, condition, and sex of the patient. The dosage can be determined by one of skill in the art and, unlike traditional AAV vectors, can also be adjusted by the individual physician in the event of any complication because ceDNA vectors do not comprise immune activating capsid proteins that prevent repeat dosing.

[00186] Administration of the ceDNA compositions described herein can be repeated for a limited period of time. In some embodiments, the doses are given periodically or by pulsed administration. In a preferred embodiment, the doses recited above are administered over several months. The duration of treatment depends upon the subject’s clinical progress and responsiveness to therapy. Booster treatments over time are contemplated. Further, the level of expression can be titrated as the subject grows.

[00187] An FIX therapeutic protein can be expressed in a subject for at least 1 week, at least 2 weeks, at least 1 month, at least 2 months, at least 6 months, at least 12 months/one year, at least 2 years, at least 5 years, at least 10 years, at least 15 years, at least 20 years, at least 30 years, at least 40 years, at least 50 years or more. Long-term expression can be achieved by repeated administration of the ceDNA vectors described herein at predetermined or desired intervals.

[00188] As used herein, the term “therapeutically effective amount” is an amount of an expressed FIX therapeutic protein, or functional fragment thereof that is sufficient to produce a statistically significant, measurable change in expression of a disease biomarker or reduction in a given disease symptom (see “Efficacy Measurement” below). Such effective amounts can be gauged in clinical trials as well as animal studies for a given ceDNA composition.

[00189] Precise amounts of the ceDNA vector required to be administered depend on the judgment of the practitioner and are particular to each individual. Suitable regimes for administration are also variable, but are typified by an initial administration followed by repeated doses at one or more intervals by a subsequent injection or other administration. Alternatively, continuous intravenous infusion sufficient to maintain concentrations in the blood in the ranges specified for *in vivo* therapies are contemplated, particularly for the treatment of acute diseases/disorders.

[00190] Agents useful in the methods and compositions described herein can be administered topically, intravenously (by bolus or continuous infusion), intracellular injection, intratissue injection, orally, by inhalation, intraperitoneally, intramuscularly, subcutaneously, intracavity, and can be delivered by peristaltic means, if desired, or by other means known by those skilled in the art. The agent can be administered systemically, if so desired. It can also be administered *in utero*.

[00191] The efficacy of a given treatment for hemophilia B, can be determined by the skilled clinician. However, a treatment is considered "effective treatment," as the term is used herein, if any one or all of the signs or symptoms of the disease or disorder is/are altered in a beneficial manner, or other clinically accepted symptoms or markers of disease are improved, or ameliorated, *e.g.*, by at least 10% following treatment with a ceDNA vector encoding FIX, or a functional fragment thereof. Efficacy can also be measured by failure of an individual to worsen as assessed by stabilization of the disease, or the need for medical interventions (*i.e.*, progression of the disease is halted or at least slowed). Methods of measuring these indicators are known to those of skill in the art and/or described herein. Treatment includes any treatment of a disease in an individual or an animal (some non-limiting examples include a human, or a mammal) and includes: (1) inhibiting the disease, *e.g.*, arresting, or slowing progression of the disease or disorder; or (2) relieving the disease, *e.g.*, causing regression of symptoms; and (3) preventing or reducing the likelihood of the development of the disease, or preventing secondary diseases/disorders associated with the disease, such as liver or kidney failure. An effective amount for the treatment of a disease means that amount which, when administered to a mammal in need thereof, is sufficient to result in effective treatment as that term is defined herein, for that disease.

[00192] Efficacy of an agent can be determined by assessing physical indicators that are particular to hemophilia B. Standard methods of analysis of hemophilia B indicators are known in the art.

[00193] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein can also encode co-factors or other polypeptides, sense or antisense oligonucleotides, or RNAs (coding or non-coding; *e.g.*, siRNAs, shRNAs, micro-RNAs, and their antisense counterparts (*e.g.*, antagoMiR)) that can be used in conjunction with the FIX protein expressed from the ceDNA. Additionally, expression cassettes comprising sequence encoding an FIX protein can also include an exogenous sequence that encodes a reporter protein to be used for experimental or diagnostic purposes, such as β -lactamase, β -galactosidase (LacZ), alkaline phosphatase, thymidine kinase, green fluorescent protein (GFP), chloramphenicol acetyltransferase (CAT), luciferase, and others well known in the art.

[00194] In one embodiment, the ceDNA vector comprises a nucleic acid sequence to express the FIX protein that is functional for the treatment of hemophilia B. In a preferred embodiment, the therapeutic FIX protein does not cause an immune system reaction, unless so desired.

III. ceDNA vector in general for use in production of FIX therapeutic proteins

[00195] Embodiments of the disclosure are based on methods and compositions comprising close ended linear duplexed (ceDNA) vectors that can express the FIX transgene. In some embodiments, the transgene is a sequence encoding an FIX protein. The ceDNA vectors for expression of FIX protein as described herein are not limited by size, thereby permitting, for example, expression of all of the components necessary for expression of a transgene from a single vector. The ceDNA vector for expression of FIX protein is preferably duplex, *e.g.* self-complementary, over at least a portion of the molecule, such as the expression cassette (*e.g.* ceDNA is not a double stranded circular molecule). The ceDNA vector has covalently closed ends, and thus is resistant to exonuclease digestion (*e.g.* exonuclease I or exonuclease III), *e.g.* for over an hour at 37°C.

[00196] In general, a ceDNA vector for expression of FIX protein as disclosed herein, comprises in the 5' to 3' direction: a first adeno-associated virus (AAV) inverted terminal repeat (ITR), a nucleic acid sequence of interest (for example an expression cassette as described herein) and a second AAV ITR. The ITR sequences selected from any of: (i) at least one WT ITR and at least one modified AAV inverted terminal repeat (mod-ITR) (*e.g.*, asymmetric modified ITRs); (ii) two modified ITRs where the mod-ITR pair have a different three-dimensional spatial organization with respect to each other (*e.g.*, asymmetric modified ITRs), or (iii) symmetrical or substantially symmetrical WT-WT ITR pair, where each WT-ITR has the same three-dimensional spatial organization, or (iv) symmetrical or substantially symmetrical modified ITR pair, where each mod-ITR has the same three-dimensional spatial organization.

[00197] Encompassed herein are methods and compositions comprising the ceDNA vector for FIX protein production, which may further include a delivery system, such as but not limited to, a liposome nanoparticle delivery system. Non-limiting exemplary liposome nanoparticle systems encompassed for use are disclosed herein. In some aspects, the disclosure provides for a lipid nanoparticle comprising ceDNA and an ionizable lipid. For example, a lipid nanoparticle formulation that is made and loaded with a ceDNA vector obtained by the process is disclosed in International Application PCT/US2018/050042, filed on September 7, 2018, which is incorporated herein.

[00198] The ceDNA vectors for expression of FIX protein as disclosed herein have no packaging constraints imposed by the limiting space within the viral capsid. ceDNA vectors represent a viable eukaryotically-produced alternative to prokaryote-produced plasmid DNA vectors, as opposed to encapsulated AAV genomes. This permits the insertion of control elements, *e.g.*, regulatory switches as disclosed herein, large transgenes, multiple transgenes etc.

[00199] FIG. 1A-1E show schematics of non-limiting, exemplary ceDNA vectors for expression of FIX protein, or the corresponding sequence of ceDNA plasmids. ceDNA vectors for expression of

FIX protein are capsid-free and can be obtained from a plasmid encoding in this order: a first ITR, an expression cassette comprising a transgene and a second ITR. The expression cassette may include one or more regulatory sequences that allows and/or controls the expression of the transgene, *e.g.*, where the expression cassette can comprise one or more of, in this order: an enhancer/promoter, an ORF reporter (transgene), a post-transcription regulatory element (*e.g.*, WPRE), and a polyadenylation and termination signal (*e.g.*, BGH polyA).

[00200] The expression cassette can also comprise an internal ribosome entry site (IRES) and/or a 2A element. The cis-regulatory elements include, but are not limited to, a promoter, a riboswitch, an insulator, a mir-regulatable element, a post-transcriptional regulatory element, a tissue- and cell type-specific promoter and an enhancer. In some embodiments the ITR can act as the promoter for the transgene, *e.g.*, FIX protein. In some embodiments, the ceDNA vector comprises additional components to regulate expression of the transgene, for example, a regulatory switch, which are described herein in the section entitled “Regulatory Switches” for controlling and regulating the expression of the FIX protein, and can include if desired, a regulatory switch which is a kill switch to enable controlled cell death of a cell comprising a ceDNA vector.

[00201] The expression cassette can comprise more than 4000 nucleotides, 5000 nucleotides, 10,000 nucleotides or 20,000 nucleotides, or 30,000 nucleotides, or 40,000 nucleotides or 50,000 nucleotides, or any range between about 4000-10,000 nucleotides or 10,000-50,000 nucleotides, or more than 50,000 nucleotides. In some embodiments, the expression cassette can comprise a transgene in the range of 500 to 50,000 nucleotides in length. In some embodiments, the expression cassette can comprise a transgene in the range of 500 to 75,000 nucleotides in length. In some embodiments, the expression cassette can comprise a transgene which is in the range of 500 to 10,000 nucleotides in length. In some embodiments, the expression cassette can comprise a transgene which is in the range of 1000 to 10,000 nucleotides in length. In some embodiments, the expression cassette can comprise a transgene which is in the range of 500 to 5,000 nucleotides in length. The ceDNA vectors do not have the size limitations of encapsidated AAV vectors, thus enable delivery of a large-size expression cassette to provide efficient transgene expression. In some embodiments, the ceDNA vector is devoid of prokaryote-specific methylation.

[00202] ceDNA expression cassette can include, for example, an expressible exogenous sequence (*e.g.*, open reading frame) or transgene that encodes a protein that is either absent, inactive, or insufficient activity in the recipient subject or a gene that encodes a protein having a desired biological or a therapeutic effect. The transgene can encode a gene product that can function to correct the expression of a defective gene or transcript. In principle, the expression cassette can include any gene that encodes a protein, polypeptide or RNA that is either reduced or absent due to a mutation or which conveys a therapeutic benefit when overexpressed is considered to be within the scope of the disclosure.

[00203] The expression cassette can comprise any transgene (*e.g.*, encoding FIX protein), for example, FIX protein useful for treating hemophilia B in a subject, *i.e.*, a therapeutic FIX protein. A ceDNA vector can be used to deliver and express any FIX protein of interest in the subject, alone or in combination with nucleic acids encoding polypeptides, or non-coding nucleic acids (*e.g.*, RNAi, miRs etc.), as well as exogenous genes and nucleic acid sequences, including virus sequences in a subjects' genome, *e.g.*, HIV virus sequences and the like. Preferably a ceDNA vector disclosed herein is used for therapeutic purposes (*e.g.*, for medical, diagnostic, or veterinary uses) or immunogenic polypeptides. In certain embodiments, a ceDNA vector is useful to express any gene of interest in the subject, which includes one or more polypeptides, peptides, ribozymes, peptide nucleic acids, siRNAs, RNAis, antisense oligonucleotides, antisense polynucleotides, or RNAs (coding or non-coding; *e.g.*, siRNAs, shRNAs, micro-RNAs, and their antisense counterparts (*e.g.*, antagoMiR)), antibodies, fusion proteins, or any combination thereof.

[00204] The expression cassette can also encode polypeptides, sense or antisense oligonucleotides, or RNAs (coding or non-coding; *e.g.*, siRNAs, shRNAs, micro-RNAs, and their antisense counterparts (*e.g.*, antagoMiR)). Expression cassettes can include an exogenous sequence that encodes a reporter protein to be used for experimental or diagnostic purposes, such as β -lactamase, β -galactosidase (LacZ), alkaline phosphatase, thymidine kinase, green fluorescent protein (GFP), chloramphenicol acetyltransferase (CAT), luciferase, and others well known in the art.

[00205] Sequences provided in the expression cassette, expression construct of a ceDNA vector for expression of FIX protein described herein can be codon optimized for the target host cell. As used herein, the term "codon optimized" or "codon optimization" refers to the process of modifying a nucleic acid sequence for enhanced expression in the cells of the vertebrate of interest, *e.g.*, mouse or human, by replacing at least one, more than one, or a significant number of codons of the native sequence (*e.g.*, a prokaryotic sequence) with codons that are more frequently or most frequently used in the genes of that vertebrate. Various species exhibit particular bias for certain codons of a particular amino acid. Typically, codon optimization does not alter the amino acid sequence of the original translated protein. Optimized codons can be determined using *e.g.*, Aptagen's GENE FORGE® codon optimization and custom gene synthesis platform (Aptagen, Inc., 2190 Fox Mill Rd. Suite 300, Herndon, Va. 20171) or another publicly available database. In some embodiments, the nucleic acid encoding the FIX protein is optimized for human expression, and/or is a human FIX, or functional fragment thereof, as known in the art.

[00206] A transgene expressed by the ceDNA vector for expression of FIX protein as disclosed herein encodes FIX protein. There are many structural features of ceDNA vectors for expression of FIX protein that differ from plasmid-based expression vectors. ceDNA vectors may possess one or more of the following features: the lack of original (*i.e.*, not inserted) bacterial DNA, the lack of a prokaryotic origin of replication, being self-containing, *i.e.*, they do not require any sequences other than the two ITRs, including the Rep binding and terminal resolution sites (RBS and TRS), and an

exogenous sequence between the ITRs, the presence of ITR sequences that form hairpins, and the absence of bacterial-type DNA methylation or indeed any other methylation considered abnormal by a mammalian host. In general, it is preferred for the present vectors not to contain any prokaryotic DNA but it is contemplated that some prokaryotic DNA may be inserted as an exogenous sequence, as a non-limiting example in a promoter or enhancer region. Another important feature distinguishing ceDNA vectors from plasmid expression vectors is that ceDNA vectors are single-strand linear DNA having closed ends, while plasmids are always double-strand DNA.

[00207] ceDNA vectors for expression of FIX protein produced by the methods provided herein preferably have a linear and continuous structure rather than a non-continuous structure, as determined by restriction enzyme digestion assay (**FIG. 4D**). The linear and continuous structure is believed to be more stable from attack by cellular endonucleases, as well as less likely to be recombined and cause mutagenesis. Thus, a ceDNA vector in the linear and continuous structure is a preferred embodiment. The continuous, linear, single strand intramolecular duplex ceDNA vector can have covalently bound terminal ends, without sequences encoding AAV capsid proteins. These ceDNA vectors are structurally distinct from plasmids (including ceDNA plasmids described herein), which are circular duplex nucleic acid molecules of bacterial origin. The complimentary strands of plasmids may be separated following denaturation to produce two nucleic acid molecules, whereas in contrast, ceDNA vectors, while having complimentary strands, are a single DNA molecule and therefore even if denatured, remain a single molecule. In some embodiments, ceDNA vectors as described herein can be produced without DNA base methylation of prokaryotic type, unlike plasmids. Therefore, the ceDNA vectors and ceDNA-plasmids are different both in term of structure (in particular, linear versus circular) and also in view of the methods used for producing and purifying these different objects (see below), and also in view of their DNA methylation which is of prokaryotic type for ceDNA-plasmids and of eukaryotic type for the ceDNA vector.

[00208] There are several advantages of using a ceDNA vector for expression of FIX protein as described herein over plasmid-based expression vectors, such advantages include, but are not limited to: 1) plasmids contain bacterial DNA sequences and are subjected to prokaryotic-specific methylation, *e.g.*, 6-methyl adenosine and 5-methyl cytosine methylation, whereas capsid-free AAV vector sequences are of eukaryotic origin and do not undergo prokaryotic-specific methylation; as a result, capsid-free AAV vectors are less likely to induce inflammatory and immune responses compared to plasmids; 2) while plasmids require the presence of a resistance gene during the production process, ceDNA vectors do not; 3) while a circular plasmid is not delivered to the nucleus upon introduction into a cell and requires overloading to bypass degradation by cellular nucleases, ceDNA vectors contain viral cis-elements, *i.e.*, ITRs, that confer resistance to nucleases and can be designed to be targeted and delivered to the nucleus. It is hypothesized that the minimal defining elements indispensable for ITR function are a Rep-binding site (RBS; 5'-GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60) for AAV2) and a terminal resolution site (TRS; 5'-AGTTGG-3' (SEQ ID NO: 64)

for AAV2) plus a variable palindromic sequence allowing for hairpin formation; and 4) ceDNA vectors do not have the over-representation of CpG dinucleotides often found in prokaryote-derived plasmids that reportedly binds a member of the Toll-like family of receptors, eliciting a T cell-mediated immune response. In contrast, transductions with capsid-free AAV vectors disclosed herein can efficiently target cell and tissue-types that are difficult to transduce with conventional AAV virions using various delivery reagent.

IV. Inverted Terminal Repeats (ITRs)

[00209] As disclosed herein, ceDNA vectors for expression of FIX protein contain a transgene or nucleic acid sequence positioned between two inverted terminal repeat (ITR) sequences, where the ITR sequences can be an asymmetrical ITR pair or a symmetrical- or substantially symmetrical ITR pair, as these terms are defined herein. A ceDNA vector as disclosed herein can comprise ITR sequences that are selected from any of: (i) at least one WT ITR and at least one modified AAV inverted terminal repeat (mod-ITR) (*e.g.*, asymmetric modified ITRs); (ii) two modified ITRs where the mod-ITR pair have a different three-dimensional spatial organization with respect to each other (*e.g.*, asymmetric modified ITRs), or (iii) symmetrical or substantially symmetrical WT-WT ITR pair, where each WT-ITR has the same three-dimensional spatial organization, or (iv) symmetrical or substantially symmetrical modified ITR pair, where each mod-ITR has the same three-dimensional spatial organization, where the methods of the present disclosure may further include a delivery system, such as but not limited to a liposome nanoparticle delivery system.

[00210] In some embodiments, the ITR sequence can be from viruses of the *Parvoviridae* family, which includes two subfamilies: *Parvovirinae*, which infect vertebrates, and *Densovirinae*, which infect insects. The subfamily *Parvovirinae* (referred to as the parvoviruses) includes the genus *Dependovirus*, the members of which, under most conditions, require coinfection with a helper virus such as adenovirus or herpes virus for productive infection. The genus *Dependovirus* includes adeno-associated virus (AAV), which normally infects humans (*e.g.*, serotypes 2, 3A, 3B, 5, and 6) or primates (*e.g.*, serotypes 1 and 4), and related viruses that infect other warm-blooded animals (*e.g.*, bovine, canine, equine, and ovine adeno-associated viruses). The parvoviruses and other members of the *Parvoviridae* family are generally described in Kenneth I. Berns, "Parvoviridae: The Viruses and Their Replication," Chapter 69 in *FIELDS VIROLOGY* (3d Ed. 1996).

[00211] While ITRs exemplified in the specification and Examples herein are AAV2 WT-ITRs, one of ordinary skill in the art is aware that one can as stated above use ITRs from any known parvovirus, for example a dependovirus such as AAV (*e.g.*, AAV1, AAV2, AAV3, AAV4, AAV5, AAV 5, AAV7, AAV8, AAV9, AAV10, AAV 11, AAV12, AAVrh8, AAVrh10, AAV-DJ, and AAV-DJ8 genome. *E.g.*, NCBI: NC 002077; NC 001401; NC001729; NC001829; NC006152; NC 006260; NC 006261), chimeric ITRs, or ITRs from any synthetic AAV. In some embodiments, the AAV can infect warm-blooded animals, *e.g.*, avian (AAAV), bovine (BAAV), canine, equine, and ovine adeno-associated viruses. In some embodiments the ITR is from B19 parvovirus (GenBank Accession No:

NC 000883), Minute Virus from Mouse (MVM) (GenBank Accession No. NC 001510); goose parvovirus (GenBank Accession No. NC 001701); snake parvovirus 1 (GenBank Accession No. NC 006148). In some embodiments, the 5' WT-ITR can be from one serotype and the 3' WT-ITR from a different serotype, as discussed herein.

[00212] An ordinarily skilled artisan is aware that ITR sequences have a common structure of a double-stranded Holliday junction, which typically is a T-shaped or Y-shaped hairpin structure (see *e.g.*, **FIG. 2A** and **FIG. 3A**), where each WT-ITR is formed by two palindromic arms or loops (B-B' and C-C') embedded in a larger palindromic arm (A-A'), and a single stranded D sequence, (where the order of these palindromic sequences defines the flip or flop orientation of the ITR). See, for example, structural analysis and sequence comparison of ITRs from different AAV serotypes (AAV1-AAV6) and described in Grimm *et al.*, J. Virology, 2006; 80(1); 426-439; Yan *et al.*, J. Virology, 2005; 364-379; Duan *et al.*, Virology 1999; 261; 8-14. One of ordinary skill in the art can readily determine WT-ITR sequences from any AAV serotype for use in a ceDNA vector or ceDNA-plasmid based on the exemplary AAV2 ITR sequences provided herein. See, for example, the sequence comparison of ITRs from different AAV serotypes (AAV1-AAV6, and avian AAV (AAAV) and bovine AAV (BAAV)) described in Grimm *et al.*, J. Virology, 2006; 80(1); 426-439; that show the % identity of the left ITR of AAV2 to the left ITR from other serotypes: AAV-1 (84%), AAV-3 (86%), AAV-4 (79%), AAV-5 (58%), AAV-6 (left ITR) (100%) and AAV-6 (right ITR) (82%).

A. Symmetrical ITR pairs

[00213] In some embodiments, a ceDNA vector for expression of FIX protein as described herein comprises, in the 5' to 3' direction: a first adeno-associated virus (AAV) inverted terminal repeat (ITR), a nucleic acid sequence of interest (for example an expression cassette as described herein) and a second AAV ITR, where the first ITR (5' ITR) and the second ITR (3' ITR) are symmetric, or substantially symmetrical with respect to each other – that is, a ceDNA vector can comprise ITR sequences that have a symmetrical three-dimensional spatial organization such that their structure is the same shape in geometrical space, or have the same A, C-C' and B-B' loops in 3D space. In such an embodiment, a symmetrical ITR pair, or substantially symmetrical ITR pair can be modified ITRs (*e.g.*, mod-ITRs) that are not wild-type ITRs. A mod-ITR pair can have the same sequence which has one or more modifications from wild-type ITR and are reverse complements (inverted) of each other. In alternative embodiments, a modified ITR pair are substantially symmetrical as defined herein, that is, the modified ITR pair can have a different sequence but have corresponding or the same symmetrical three-dimensional shape.

[00214] (i) *Wildtype ITRs*

[00215] In some embodiments, the symmetrical ITRs, or substantially symmetrical ITRs are wild type (WT-ITRs) as described herein. That is, both ITRs have a wild-type sequence, but do not necessarily have to be WT-ITRs from the same AAV serotype. That is, in some embodiments, one WT-ITR can be from one AAV serotype, and the other WT-ITR can be from a different AAV

serotype. In such an embodiment, a WT-ITR pair are substantially symmetrical as defined herein, that is, they can have one or more conservative nucleotide modification while still retaining the symmetrical three-dimensional spatial organization.

[00216] Accordingly, as disclosed herein, ceDNA vectors contain a transgene or nucleic acid sequence positioned between two flanking wild-type inverted terminal repeat (WT-ITR) sequences, that are either the reverse complement (inverted) of each other, or alternatively, are substantially symmetrical relative to each other – that is a WT-ITR pair have symmetrical three-dimensional spatial organization. In some embodiments, a wild-type ITR sequence (*e.g.* AAV WT-ITR) comprises a functional Rep binding site (RBS; *e.g.* 5'-GCGCGCTCGCTCGCTC-3' for AAV2, SEQ ID NO: 60) and a functional terminal resolution site (TRS; *e.g.* 5'-AGTT-3', SEQ ID NO: 62).

[00217] In one aspect, ceDNA vectors for expression of FIX protein are obtainable from a vector polynucleotide that encodes a nucleic acid operatively positioned between two WT inverted terminal repeat sequences (WT-ITRs) (*e.g.* AAV WT-ITRs). That is, both ITRs have a wild type sequence, but do not necessarily have to be WT-ITRs from the same AAV serotype. That is, in some embodiments, one WT-ITR can be from one AAV serotype, and the other WT-ITR can be from a different AAV serotype. In such an embodiment, the WT-ITR pair are substantially symmetrical as defined herein, that is, they can have one or more conservative nucleotide modification while still retaining the symmetrical three-dimensional spatial organization. In some embodiments, the 5' WT-ITR is from one AAV serotype, and the 3' WT-ITR is from the same or a different AAV serotype. In some embodiments, the 5' WT-ITR and the 3' WT-ITR are mirror images of each other, that is they are symmetrical. In some embodiments, the 5' WT-ITR and the 3' WT-ITR are from the same AAV serotype.

[00218] WT ITRs are well known. In one embodiment the two ITRs are from the same AAV2 serotype. In certain embodiments one can use WT from other serotypes. There are a number of serotypes that are homologous, *e.g.* AAV2, AAV4, AAV6, AAV8. In one embodiment, closely homologous ITRs (*e.g.* ITRs with a similar loop structure) can be used. In another embodiment, one can use AAV WT ITRs that are more diverse, *e.g.*, AAV2 and AAV5, and still another embodiment, one can use an ITR that is substantially WT - that is, it has the basic loop structure of the WT but some conservative nucleotide changes that do not alter or affect the properties. When using WT-ITRs from the same viral serotype, one or more regulatory sequences may further be used. In certain embodiments, the regulatory sequence is a regulatory switch that permits modulation of the activity of the ceDNA, *e.g.*, the expression of the encoded FIX protein.

[00219] In some embodiments, one aspect of the technology described herein relates to a ceDNA vector for expression of FIX protein, wherein the ceDNA vector comprises at least one nucleic acid sequence encoding the FIX protein, operably positioned between two wild-type inverted terminal repeat sequences (WT-ITRs), wherein the WT-ITRs can be from the same serotype, different serotypes or substantially symmetrical with respect to each other (*i.e.*, have the symmetrical three-dimensional

spatial organization such that their structure is the same shape in geometrical space, or have the same A, C-C' and B-B' loops in 3D space). In some embodiments, the symmetric WT-ITRs comprises a functional terminal resolution site and a Rep binding site. In some embodiments, the nucleic acid sequence encodes a transgene, and wherein the vector is not in a viral capsid.

[00220] In some embodiments, the WT-ITRs are the same but the reverse complement of each other. For example, the sequence AACG in the 5' ITR may be CGTT (*i.e.*, the reverse complement) in the 3' ITR at the corresponding site. In one example, the 5' WT-ITR sense strand comprises the sequence of ATCGATCG and the corresponding 3' WT-ITR sense strand comprises CGATCGAT (*i.e.*, the reverse complement of ATCGATCG). In some embodiments, the WT-ITRs ceDNA further comprises a terminal resolution site and a replication protein binding site (RPS) (sometimes referred to as a replicative protein binding site), *e.g.* a Rep binding site.

[00221] Exemplary WT-ITR sequences for use in the ceDNA vectors for expression of FIX protein comprising WT-ITRs are shown in **Table 2** herein, which shows pairs of WT-ITRs (5' WT-ITR and the 3' WT-ITR).

[00222] As an exemplary example, the present disclosure provides a ceDNA vector for expression of FIX protein comprising a promoter operably linked to a transgene (*e.g.*, nucleic acid sequence), with or without the regulatory switch, where the ceDNA is devoid of capsid proteins and is: (a) produced from a ceDNA-plasmid (*e.g.*, see **FIGS. 1F-1G**) that encodes WT-ITRs, where each WT-ITR has the same number of intramolecularly duplexed base pairs in its hairpin secondary configuration (preferably excluding deletion of any AAA or TTT terminal loop in this configuration compared to these reference sequences), and (b) is identified as ceDNA using the assay for the identification of ceDNA by agarose gel electrophoresis under native gel and denaturing conditions in Example 1.

[00223] In some embodiments, the flanking WT-ITRs are substantially symmetrical to each other. In this embodiment the 5' WT-ITR can be from one serotype of AAV, and the 3' WT-ITR from a different serotype of AAV, such that the WT-ITRs are not identical reverse complements. For example, the 5' WT-ITR can be from AAV2, and the 3' WT-ITR from a different serotype (*e.g.* AAV1, 3, 4, 5, 6, 7, 8, 9, 10, 11, and 12. In some embodiments, WT-ITRs can be selected from two different parvoviruses selected from any to of: AAV1, AAV2, AAV3, AAV4, AAV5, AAV6, AAV7, AAV8, AAV9, AAV10, AAV11, AAV12, AAV13, snake parvovirus (*e.g.*, royal python parvovirus), bovine parvovirus, goat parvovirus, avian parvovirus, canine parvovirus, equine parvovirus, shrimp parvovirus, porcine parvovirus, or insect AAV. In some embodiments, such a combination of WT ITRs is the combination of WT-ITRs from AAV2 and AAV6. In one embodiment, the substantially symmetrical WT-ITRs are when one is inverted relative to the other ITR at least 90% identical, at least 95% identical, at least 96%...97%... 98%... 99%....99.5% and all points in between, and has the same symmetrical three-dimensional spatial organization. In some embodiments, a WT-ITR pair are substantially symmetrical as they have symmetrical three-dimensional spatial organization, *e.g.*, have the same 3D organization of the A, C-C'. B-B' and D arms. In one embodiment, a substantially

symmetrical WT-ITR pair are inverted relative to the other, and are at least 95% identical, at least 96%...97%... 98%... 99%...99.5% and all points in between, to each other, and one WT-ITR retains the Rep-binding site (RBS) of 5'-GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60) and a terminal resolution site (trs). In some embodiments, a substantially symmetrical WT-ITR pair are inverted relative to each other, and are at least 95% identical, at least 96%...97%... 98%... 99%...99.5% and all points in between, to each other, and one WT-ITR retains the Rep-binding site (RBS) of 5'-GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60) and a terminal resolution site (trs) and in addition to a variable palindromic sequence allowing for hairpin secondary structure formation. Homology can be determined by standard means well known in the art such as BLAST (Basic Local Alignment Search Tool), BLASTN at default setting.

[00224] In some embodiments, the structural element of the ITR can be any structural element that is involved in the functional interaction of the ITR with a large Rep protein (*e.g.*, Rep 78 or Rep 68). In certain embodiments, the structural element provides selectivity to the interaction of an ITR with a large Rep protein, *i.e.*, determines at least in part which Rep protein functionally interacts with the ITR. In other embodiments, the structural element physically interacts with a large Rep protein when the Rep protein is bound to the ITR. Each structural element can be, *e.g.*, a secondary structure of the ITR, a nucleic acid sequence of the ITR, a spacing between two or more elements, or a combination of any of the above. In one embodiment, the structural elements are selected from the group consisting of an A and an A' arm, a B and a B' arm, a C and a C' arm, a D arm, a Rep binding site (RBE) and an RBE' (*i.e.*, complementary RBE sequence), and a terminal resolution site (trs).

[00225] By way of example only, **Table 2** indicates exemplary combinations of WT-ITRs.

[00226] **Table 2:** Exemplary combinations of WT-ITRs from the same serotype or different serotypes, or different parvoviruses. The order shown is not indicative of the ITR position, for example, "AAV1, AAV2" demonstrates that the cdDNA can comprise a WT-AAV1 ITR in the 5' position, and a WT-AAV2 ITR in the 3' position, or vice versa, a WT-AAV2 ITR the 5' position, and a WT-AAV1 ITR in the 3' position. Abbreviations: AAV serotype 1 (AAV1), AAV serotype 2 (AAV2), AAV serotype 3 (AAV3), AAV serotype 4 (AAV4), AAV serotype 5 (AAV5), AAV serotype 6 (AAV6), AAV serotype 7 (AAV7), AAV serotype 8 (AAV8), AAV serotype 9 (AAV9), AAV serotype 10 (AAV10), AAV serotype 11 (AAV11), or AAV serotype 12 (AAV12); AAVrh8, AAVrh10, AAV-DJ, and AAV-DJ8 genome (*E.g.*, NCBI: NC 002077; NC 001401; NC001729; NC001829; NC006152; NC 006260; NC 006261), ITRs from warm-blooded animals (avian AAV (AAV), bovine AAV (BAAV), canine, equine, and ovine AAV), ITRs from B19 *Parvovirus* (GenBank Accession No: NC 000883), Minute Virus from Mouse (MVM) (GenBank Accession No. NC 001510); Goose: goose parvovirus (GenBank Accession No. NC 001701); snake: snake parvovirus 1 (GenBank Accession No. NC 006148).

Table 2: Exemplary combinations of WT-ITRs

AAV1,AAV1	AAV2,AAV2	AAV3,AAV3	AAV4,AAV4	AAV5,AAV5
AAV1,AAV2	AAV2,AAV3	AAV3,AAV4	AAV4,AAV5	AAV5,AAV6
AAV1,AAV3	AAV2,AAV4	AAV3,AAV5	AAV4,AAV6	AAV5,AAV7
AAV1,AAV4	AAV2,AAV5	AAV3,AAV6	AAV4,AAV7	AAV5,AAV8
AAV1,AAV5	AAV2,AAV6	AAV3,AAV7	AAV4,AAV8	AAV5,AAV9
AAV1,AAV6	AAV2,AAV7	AAV3,AAV8	AAV4,AAV9	AAV5,AAV10
AAV1,AAV7	AAV2,AAV8	AAV3,AAV9	AAV4,AAV10	AAV5,AAV11
AAV1,AAV8	AAV2,AAV9	AAV3,AAV10	AAV4,AAV11	AAV5,AAV12
AAV1,AAV9	AAV2,AAV10	AAV3,AAV11	AAV4,AAV12	AAV5,AAVRH8
AAV1,AAV10	AAV2,AAV11	AAV3,AAV12	AAV4,AAVRH8	AAV5,AAVRH10
AAV1,AAV11	AAV2,AAV12	AAV3,AAVRH8	AAV4,AAVRH10	AAV5,AAV13
AAV1,AAV12	AAV2,AAVRH8	AAV3,AAVRH10	AAV4,AAV13	AAV5,AAVDJ
AAV1,AAVRH8	AAV2,AAVRH10	AAV3,AAV13	AAV4,AAVDJ	AAV5,AAVDJ8
AAV1,AAVRH10	AAV2,AAV13	AAV3,AAVDJ	AAV4,AAVDJ8	AAV5,AVIAN
AAV1,AAV13	AAV2,AAVDJ	AAV3,AAVDJ8	AAV4,AVIAN	AAV5,BOVINE
AAV1,AAVDJ	AAV2,AAVDJ8	AAV3,AVIAN	AAV4,BOVINE	AAV5,CANINE
AAV1,AAVDJ8	AAV2,AVIAN	AAV3,BOVINE	AAV4,CANINE	AAV5,EQUINE
AAV1,AVIAN	AAV2,BOVINE	AAV3,CANINE	AAV4,EQUINE	AAV5,GOAT
AAV1,BOVINE	AAV2,CANINE	AAV3,EQUINE	AAV4,GOAT	AAV5,SHRIMP
AAV1,CANINE	AAV2,EQUINE	AAV3,GOAT	AAV4,SHRIMP	AAV5,PORCINE
AAV1,EQUINE	AAV2,GOAT	AAV3,SHRIMP	AAV4,PORCINE	AAV5,INSECT
AAV1,GOAT	AAV2,SHRIMP	AAV3,PORCINE	AAV4,INSECT	AAV5,OVINE
AAV1,SHRIMP	AAV2,PORCINE	AAV3,INSECT	AAV4,OVINE	AAV5,B19
AAV1,PORCINE	AAV2,INSECT	AAV3,OVINE	AAV4,B19	AAV5,MVM
AAV1,INSECT	AAV2,OVINE	AAV3,B19	AAV4,MVM	AAV5,GOOSE
AAV1,OVINE	AAV2,B19	AAV3,MVM	AAV4,GOOSE	AAV5,SNAKE
AAV1,B19	AAV2,MVM	AAV3,GOOSE	AAV4,SNAKE	
AAV1,MVM	AAV2,GOOSE	AAV3,SNAKE		
AAV1,GOOSE	AAV2,SNAKE			
AAV1,SNAKE				
AAV6,AAV6	AAV7,AAV7	AAV8,AAV8	AAV9,AAV9	AAV10,AAV10
AAV6,AAV7	AAV7,AAV8	AAV8,AAV9	AAV9,AAV10	AAV10,AAV11
AAV6,AAV8	AAV7,AAV9	AAV8,AAV10	AAV9,AAV11	AAV10,AAV12
AAV6,AAV9	AAV7,AAV10	AAV8,AAV11	AAV9,AAV12	AAV10,AAVRH8
AAV6,AAV10	AAV7,AAV11	AAV8,AAV12	AAV9,AAVRH8	AAV10,AAVRH10
AAV6,AAV11	AAV7,AAV12	AAV8,AAVRH8	AAV9,AAVRH10	AAV10,AAV13

AAV6,AAV12	AAV7,AAVRH8	AAV8,AAVRH10	AAV9,AAV13	AAV10,AAVDJ
AAV6,AAVRH8	AAV7,AAVRH10	AAV8,AAV13	AAV9,AAVDJ	AAV10,AAVDJ8
AAV6,AAVRH10	AAV7,AAV13	AAV8,AAVDJ	AAV9,AAVDJ8	AAV10,AVIAN
AAV6,AAV13	AAV7,AAVDJ	AAV8,AAVDJ8	AAV9,AVIAN	AAV10,BOVINE
AAV6,AAVDJ	AAV7,AAVDJ8	AAV8,AVIAN	AAV9,BOVINE	AAV10,CANINE
AAV6,AAVDJ8	AAV7,AVIAN	AAV8,BOVINE	AAV9,CANINE	AAV10,EQUINE
AAV6,AVIAN	AAV7,BOVINE	AAV8,CANINE	AAV9,EQUINE	AAV10,GOAT
AAV6,BOVINE	AAV7,CANINE	AAV8,EQUINE	AAV9,GOAT	AAV10,SHRIMP
AAV6,CANINE	AAV7,EQUINE	AAV8,GOAT	AAV9,SHRIMP	AAV10,PORCINE
AAV6,EQUINE	AAV7,GOAT	AAV8,SHRIMP	AAV9,PORCINE	AAV10,INSECT
AAV6,GOAT	AAV7,SHRIMP	AAV8,PORCINE	AAV9,INSECT	AAV10,OVINE
AAV6,SHRIMP	AAV7,PORCINE	AAV8,INSECT	AAV9,OVINE	AAV10,B19
AAV6,PORCINE	AAV7,INSECT	AAV8,OVINE	AAV9,B19	AAV10,MVM
AAV6,INSECT	AAV7,OVINE	AAV8,B19	AAV9,MVM	AAV10,GOOSE
AAV6,OVINE	AAV7,B19	AAV8,MVM	AAV9,GOOSE	AAV10,SNAKE
AAV6,B19	AAV7,MVM	AAV8,GOOSE	AAV9,SNAKE	
AAV6,MVM	AAV7,GOOSE	AAV8,SNAKE		
AAV6,GOOSE	AAV7,SNAKE			
AAV6,SNAKE				
AAV11,AAV11	AAV12,AAV12	AAVRH8,AAVRH8	AAVRH10,AAVRH10	AAV13,AAV13
AAV11,AAV12	AAV12,AAVRH8	AAVRH8,AAVRH10	AAVRH10,AAV13	AAV13,AAVDJ
AAV11,AAVRH8	AAV12,AAVRH10	AAVRH8,AAV13	AAVRH10,AAVDJ	AAV13,AAVDJ8
AAV11,AAVRH10	AAV12,AAV13	AAVRH8,AAVDJ	AAVRH10,AAVDJ8	AAV13,AVIAN
AAV11,AAV13	AAV12,AAVDJ	AAVRH8,AAVDJ8	AAVRH10,AVIAN	AAV13,BOVINE
AAV11,AAVDJ	AAV12,AAVDJ8	AAVRH8,AVIAN	AAVRH10,BOVINE	AAV13,CANINE
AAV11,AAVDJ8	AAV12,AVIAN	AAVRH8,BOVINE	AAVRH10,CANINE	AAV13,EQUINE
AAV11,AVIAN	AAV12,BOVINE	AAVRH8,CANINE	AAVRH10,EQUINE	AAV13,GOAT
AAV11,BOVINE	AAV12,CANINE	AAVRH8,EQUINE	AAVRH10,GOAT	AAV13,SHRIMP
AAV11,CANINE	AAV12,EQUINE	AAVRH8,GOAT	AAVRH10,SHRIMP	AAV13,PORCINE
AAV11,EQUINE	AAV12,GOAT	AAVRH8,SHRIMP	AAVRH10,PORCINE	AAV13,INSECT
AAV11,GOAT	AAV12,SHRIMP	AAVRH8,PORCINE	AAVRH10,INSECT	AAV13,OVINE
AAV11,SHRIMP	AAV12,PORCINE	AAVRH8,INSECT	AAVRH10,OVINE	AAV13,B19
AAV11,PORCINE	AAV12,INSECT	AAVRH8,OVINE	AAVRH10,B19	AAV13,MVM
AAV11,INSECT	AAV12,OVINE	AAVRH8,B19	AAVRH10,MVM	AAV13,GOOSE
AAV11,OVINE	AAV12,B19	AAVRH8,MVM	AAVRH10,GOOSE	AAV13,SNAKE
AAV11,B19	AAV12,MVM	AAVRH8,GOOSE	AAVRH10,SNAKE	
AAV11,MVM	AAV12,GOOSE	AAVRH8,SNAKE		
AAV11,GOOSE	AAV12,SNAKE			

AAV11,SNAKE				
AAVDJ,AAVDJ	AAVDJ8,AVVDJ8	AVIAN, AVIAN	BOVINE, BOVINE	CANINE, CANINE
AAVDJ,AAVDJ8	AAVDJ8,AVIAN	AVIAN,BOVINE	BOVINE,CANINE	CANINE,EQUINE
AAVDJ,AVIAN	AAVDJ8,BOVINE	AVIAN,CANINE	BOVINE,EQUINE	CANINE,GOAT
AAVDJ,BOVINE	AAVDJ8,CANINE	AVIAN,EQUINE	BOVINE,GOAT	CANINE,SHRIMP
AAVDJ,CANINE	AAVDJ8,EQUINE	AVIAN,GOAT	BOVINE,SHRIMP	CANINE,PORCIN E
AAVDJ,EQUINE	AAVDJ8,GOAT	AVIAN,SHRIMP	BOVINE,PORCINE	CANINE,INSECT
AAVDJ,GOAT	AAVDJ8,SHRIMP	AVIAN,PORCINE	BOVINE,INSECT	CANINE,OVINE
AAVDJ,SHRIMP	AAVDJ8,PORCINE	AVIAN,INSECT	BOVINE,OVINE	CANINE,B19
AAVDJ,PORCINE	AAVDJ8,INSECT	AVIAN,OVINE	BOVINE,B19	CANINE,MVM
AAVDJ,INSECT	AAVDJ8,OVINE	AVIAN,B19	BOVINE,MVM	CANINE,GOOSE
AAVDJ,OVINE	AAVDJ8,B19	AVIAN,MVM	BOVINE,GOOSE	CANINE,SNAKE
AAVDJ,B19	AAVDJ8,MVM	AVIAN,GOOSE	BOVINE,SNAKE	
AAVDJ,MVM	AAVDJ8,GOOSE	AVIAN,SNAKE		
AAVDJ,GOOSE	AAVDJ8,SNAKE			
AAVDJ,SNAKE				
EQUINE, EQUINE	GOAT, GOAT	SHRIMP, SHRIMP	PORCINE, PORCINE	INSECT, INSECT
EQUINE,GOAT	GOAT,SHRIMP	SHRIMP,PORCINE	PORCINE,INSECT	INSECT,OVINE
EQUINE,SHRIMP	GOAT,PORCINE	SHRIMP,INSECT	PORCINE,OVINE	INSECT,B19
EQUINE,PORCINE	GOAT,INSECT	SHRIMP,OVINE	PORCINE,B19	INSECT,MVM
EQUINE,INSECT	GOAT,OVINE	SHRIMP,B19	PORCINE,MVM	INSECT,GOOSE
EQUINE,OVINE	GOAT,B19	SHRIMP,MVM	PORCINE,GOOSE	INSECT,SNAKE
EQUINE,B19	GOAT,MVM	SHRIMP,GOOSE	PORCINE,SNAKE	
EQUINE,MVM	GOAT,GOOSE	SHRIMP,SNAKE		
EQUINE,GOOSE	GOAT,SNAKE			
EQUINE,SNAKE				
OVINE, OVINE	B19, B19	MVM, MVM	GOOSE, GOOSE	SNAKE, SNAKE
OVINE,B19	B19,MVM	MVM,GOOSE	GOOSE,SNAKE	
OVINE,MVM	B19,GOOSE	MVM,SNAKE		
OVINE,GOOSE	B19,SNAKE			
OVINE,SNAKE				

[00227] By way of example only, **Table 3** shows the sequences of exemplary WT-ITRs from some different AAV serotypes.

Table 3: Exemplary WT-ITRs

AAV serotype	5' WT-ITR (LEFT)	3' WT-ITR (RIGHT)
AAV1	SEQ ID NO: 5	SEQ ID NO: 10
AAV2	SEQ ID NO: 2	SEQ ID NO: 1
AAV3	SEQ ID NO: 6	SEQ ID NO: 11
AAV4	SEQ ID NO: 7	SEQ ID NO: 12
AAV5	SEQ ID NO: 8	SEQ ID NO: 13
AAV6	SEQ ID NO: 9	SEQ ID NO: 14

[00228] In some embodiments, the nucleic acid sequence of the WT-ITR sequence can be modified (*e.g.*, by modifying 1, 2, 3, 4 or 5, or more nucleotides or any range therein), whereby the modification is a substitution for a complementary nucleotide, *e.g.*, G for a C, and vice versa, and T for an A, and vice versa.

[00229] In certain embodiments of the present disclosure, the ceDNA vector for expression of FIX protein does not have a WT-ITR consisting of the nucleic acid sequence selected from any of: SEQ ID NOs: 1, 2, 5-14. In alternative embodiments of the present disclosure, if a ceDNA vector has a WT-ITR comprising the nucleic acid sequence selected from any of: SEQ ID NOs: 1, 2, 5-14, then the flanking ITR is also WT and the ceDNA vector comprises a regulatory switch, *e.g.*, as disclosed herein and in International application PCT/US18/49996 (*e.g.*, see Table 11 of PCT/US18/49996, incorporated by reference in its entirety herein). In some embodiments, the ceDNA vector for expression of FIX protein comprises a regulatory switch as disclosed herein and a WT-ITR selected having the nucleic acid sequence selected from any of the group consisting of: SEQ ID NO: 1, 2, 5-14.

[00230] The ceDNA vector for expression of FIX protein as described herein can include WT-ITR structures that retains an operable RBE, trs and RBE' portion. **FIG. 2A** and **FIG. 2B**, using wild-type ITRs for exemplary purposes, show one possible mechanism for the operation of a trs site within a wild type ITR structure portion of a ceDNA vector. In some embodiments, the ceDNA vector for expression of FIX protein contains one or more functional WT-ITR polynucleotide sequences that comprise a Rep-binding site (RBS; 5'-GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60) for AAV2) and a terminal resolution site (TRS; 5'-AGTT (SEQ ID NO: 62)). In some embodiments, at least one WT-ITR is functional. In alternative embodiments, where a ceDNA vector for expression of FIX protein comprises two WT-ITRs that are substantially symmetrical to each other, at least one WT-ITR is functional and at least one WT-ITR is non-functional.

B. Modified ITRs (mod-ITRs) in general for ceDNA vectors comprising asymmetric ITR pairs or symmetric ITR pairs

[00231] As discussed herein, a ceDNA vector for expression of FIX protein can comprise a symmetrical ITR pair or an asymmetrical ITR pair. In both instances, one or both of the ITRs can be

modified ITRs – the difference being that in the first instance (*i.e.*, symmetric mod-ITRs), the mod-ITRs have the same three-dimensional spatial organization (*i.e.*, have the same A-A', C-C' and B-B' arm configurations), whereas in the second instance (*i.e.*, asymmetric mod-ITRs), the mod-ITRs have a different three-dimensional spatial organization (*i.e.*, have a different configuration of A-A', C-C' and B-B' arms).

[00232] In some embodiments, a modified ITR is an ITRs that is modified by deletion, insertion, and/or substitution as compared to a wild-type ITR sequence (*e.g.* AAV ITR). In some embodiments, at least one of the ITRs in the ceDNA vector comprises a functional Rep binding site (RBS; *e.g.* 5'-GCGCGCTCGCTCGCTC-3' for AAV2, SEQ ID NO: 60) and a functional terminal resolution site (TRS; *e.g.* 5'-AGTT-3', SEQ ID NO: 62.) In one embodiment, at least one of the ITRs is a non-functional ITR. In one embodiment, the different or modified ITRs are not each wild type ITRs from different serotypes.

[00233] Specific alterations and mutations in the ITRs are described in detail herein, but in the context of ITRs, “altered” or “mutated” or “modified”, it indicates that nucleotides have been inserted, deleted, and/or substituted relative to the wild-type, reference, or original ITR sequence. The altered or mutated ITR can be an engineered ITR. As used herein, “engineered” refers to the aspect of having been manipulated by the hand of man. For example, a polypeptide is considered to be “engineered” when at least one aspect of the polypeptide, *e.g.*, its sequence, has been manipulated by the hand of man to differ from the aspect as it exists in nature.

[00234] In some embodiments, a mod-ITR may be synthetic. In one embodiment, a synthetic ITR is based on ITR sequences from more than one AAV serotype. In another embodiment, a synthetic ITR includes no AAV-based sequence. In yet another embodiment, a synthetic ITR preserves the ITR structure described above although having only some or no AAV-sourced sequence. In some aspects, a synthetic ITR may interact preferentially with a wild type Rep or a Rep of a specific serotype, or in some instances will not be recognized by a wild-type Rep and be recognized only by a mutated Rep.

[00235] The skilled artisan can determine the corresponding sequence in other serotypes by known means. For example, determining if the change is in the A, A', B, B', C, C' or D region and determine the corresponding region in another serotype. One can use BLAST® (Basic Local Alignment Search Tool) or other homology alignment programs at default status to determine the corresponding sequence. The disclosure further provides populations and pluralities of ceDNA vectors comprising mod-ITRs from a combination of different AAV serotypes – that is, one mod-ITR can be from one AAV serotype and the other mod-ITR can be from a different serotype. Without wishing to be bound by theory, in one embodiment one ITR can be from or based on an AAV2 ITR sequence and the other ITR of the ceDNA vector can be from or be based on any one or more ITR sequence of AAV serotype 1 (AAV1), AAV serotype 4 (AAV4), AAV serotype 5 (AAV5), AAV serotype 6 (AAV6), AAV serotype 7 (AAV7), AAV serotype 8 (AAV8), AAV serotype 9 (AAV9), AAV serotype 10 (AAV10), AAV serotype 11 (AAV11), or AAV serotype 12 (AAV12).

[00236] Any parvovirus ITR can be used as an ITR or as a base ITR for modification. Preferably, the parvovirus is a dependovirus. More preferably AAV. The serotype chosen can be based upon the tissue tropism of the serotype. AAV2 has a broad tissue tropism, AAV1 preferentially targets to neuronal and skeletal muscle, and AAV5 preferentially targets neuronal, retinal pigmented epithelia, and photoreceptors. AAV6 preferentially targets skeletal muscle and lung. AAV8 preferentially targets liver, skeletal muscle, heart, and pancreatic tissues. AAV9 preferentially targets liver, skeletal and lung tissue. In one embodiment, the modified ITR is based on an AAV2 ITR.

[00237] More specifically, the ability of a structural element to functionally interact with a particular large Rep protein can be altered by modifying the structural element. For example, the nucleic acid sequence of the structural element can be modified as compared to the wild-type sequence of the ITR. In one embodiment, the structural element (*e.g.*, A arm, A' arm, B arm, B' arm, C arm, C' arm, D arm, RBE, RBE', and trs) of an ITR can be removed and replaced with a wild-type structural element from a different parvovirus. For example, the replacement structure can be from AAV1, AAV2, AAV3, AAV4, AAV5, AAV6, AAV7, AAV8, AAV9, AAV10, AAV11, AAV12, AAV13, snake parvovirus (*e.g.*, royal python parvovirus), bovine parvovirus, goat parvovirus, avian parvovirus, canine parvovirus, equine parvovirus, shrimp parvovirus, porcine parvovirus, or insect AAV. For example, the ITR can be an AAV2 ITR and the A or A' arm or RBE can be replaced with a structural element from AAV5. In another example, the ITR can be an AAV5 ITR and the C or C' arms, the RBE, and the trs can be replaced with a structural element from AAV2. In another example, the AAV ITR can be an AAV5 ITR with the B and B' arms replaced with the AAV2 ITR B and B' arms.

[00238] By way of example only, **Table 4** indicates exemplary modifications of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in regions of a modified ITR, where X is indicative of a modification of at least one nucleic acid (*e.g.*, a deletion, insertion and/ or substitution) in that section relative to the corresponding wild-type ITR. In some embodiments, any modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in any of the regions of C and/or C' and/or B and/or B' retains three sequential T nucleotides (*i.e.*, TTT) in at least one terminal loop. For example, if the modification results in any of: a single arm ITR (*e.g.*, single C-C' arm, or a single B-B' arm), or a modified C-B' arm or C'-B arm, or a two arm ITR with at least one truncated arm (*e.g.*, a truncated C-C' arm and/or truncated B-B' arm), at least the single arm, or at least one of the arms of a two arm ITR (where one arm can be truncated) retains three sequential T nucleotides (*i.e.*, TTT) in at least one terminal loop. In some embodiments, a truncated C-C' arm and/or a truncated B-B' arm has three sequential T nucleotides (*i.e.*, TTT) in the terminal loop.

Table 4: Exemplary combinations of modifications of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) to different B-B' and C-C' regions or arms of ITRs (X indicates a nucleotide modification, *e.g.*, addition, deletion or substitution of at least one nucleotide in the region).

B region	B' region	C region	C' region
X			
	X		
X	X		
		X	
			X
		X	X
X		X	
X			X
	X	X	
	X		X
X	X	X	
X	X		X
X		X	X
	X	X	X
X	X	X	X

[00239] In some embodiments, mod-ITR for use in a ceDNA vector for expression of FIX protein comprises an asymmetric ITR pair, or a symmetric mod-ITR pair as disclosed herein, can comprise any one of the combinations of modifications shown in **Table 4**, and also a modification of at least one nucleotide in any one or more of the regions selected from: between A' and C, between C and C', between C' and B, between B and B' and between B' and A. In some embodiments, any modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in the C or C' or B or B' regions, still preserves the terminal loop of the stem-loop. In some embodiments, any modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) between C and C' and/or B and B' retains three sequential T nucleotides (*i.e.*, TTT) in at least one terminal loop. In alternative embodiments, any modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) between C and C' and/or B and B' retains three sequential A nucleotides (*i.e.*, AAA) in at least one terminal loop. In some embodiments, a modified ITR for use herein can comprise any one of the combinations of modifications shown in **Table 4**, and also a modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in any one or more of the regions selected from: A', A and/or D. For example, in some embodiments, a modified ITR for use herein can comprise any one of the combinations of modifications shown in **Table 4**, and also a modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in the A region. In some embodiments, a modified ITR for use herein can comprise any one of the combinations of modifications shown in **Table 4**, and also a modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in the A' region. In some embodiments, a modified ITR for use herein can comprise any one of the combinations of modifications shown in **Table 4**, and also a modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in the A and/or A' region. In some embodiments, a modified ITR for use

herein can comprise any one of the combinations of modifications shown in **Table 4**, and also a modification of at least one nucleotide (*e.g.*, a deletion, insertion and/ or substitution) in the D region.

[00240] In one embodiment, the nucleotide sequence of the structural element can be modified (*e.g.*, by modifying 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 or more nucleotides or any range therein) to produce a modified structural element. In one embodiment, the specific modifications to the ITRs are exemplified herein (*e.g.*, SEQ ID NOS: 3, 4, 15-47, 101-116 or 165-187, or shown in FIG. 7A-7B of International Patent Application No. PCT/US2018/064242, filed on December 6, 2018 (*e.g.*, SEQ ID Nos 97-98, 101-103, 105-108, 111-112, 117-134, 545-54 in PCT/US2018/064242). In some embodiments, an ITR can be modified (*e.g.*, by modifying 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 or more nucleotides or any range therein). In other embodiments, the ITR can have at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or more sequence identity with one of the modified ITRs of SEQ ID NOS: 3, 4, 15-47, 101-116 or 165-187, or the RBE-containing section of the A-A' arm and C-C' and B-B' arms of SEQ ID NO: 3, 4, 15-47, 101-116 or 165-187, or shown in Tables 2-9 (*i.e.*, SEQ ID NO: 110-112, 115-190, 200-468) of International Patent Application No. PCT/US18/49996, which is incorporated herein in its entirety by reference.

[00241] In some embodiments, a modified ITR can for example, comprise removal or deletion of all of a particular arm, *e.g.*, all or part of the A-A' arm, or all or part of the B-B' arm or all or part of the C-C' arm, or alternatively, the removal of 1, 2, 3, 4, 5, 6, 7, 8, 9 or more base pairs forming the stem of the loop so long as the final loop capping the stem (*e.g.*, single arm) is still present (*e.g.*, see ITR-21 in FIG. 7A of PCT/US2018/064242, filed December 6, 2018). In some embodiments, a modified ITR can comprise the removal of 1, 2, 3, 4, 5, 6, 7, 8, 9 or more base pairs from the B-B' arm. In some embodiments, a modified ITR can comprise the removal of 1, 2, 3, 4, 5, 6, 7, 8, 9 or more base pairs from the C-C' arm (see, *e.g.*, ITR-1 in FIG. 3B, or ITR-45 in FIG. 7A of International Patent Application No. PCT/US2018/064242, filed December 6, 2018). In some embodiments, a modified ITR can comprise the removal of 1, 2, 3, 4, 5, 6, 7, 8, 9 or more base pairs from the C-C' arm and the removal of 1, 2, 3, 4, 5, 6, 7, 8, 9 or more base pairs from the B-B' arm. Any combination of removal of base pairs is envisioned, for example, 6 base pairs can be removed in the C-C' arm and 2 base pairs in the B-B' arm. As an illustrative example, **FIG. 3B** shows an exemplary modified ITR with at least 7 base pairs deleted from each of the C portion and the C' portion, a substitution of a nucleotide in the loop between C and C' region, and at least one base pair deletion from each of the B region and B' regions such that the modified ITR comprises two arms where at least one arm (*e.g.*, C-C') is truncated. In some embodiments, the modified ITR also comprises at least one base pair deletion from each of the B region and B' regions, such that the B-B' arm is also truncated relative to WT ITR.

[00242] In some embodiments, a modified ITR can have between 1 and 50 (*e.g.* 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, or 50) nucleotide deletions relative to a full-length

wild-type ITR sequence. In some embodiments, a modified ITR can have between 1 and 30 nucleotide deletions relative to a full-length WT ITR sequence. In some embodiments, a modified ITR has between 2 and 20 nucleotide deletions relative to a full-length wild-type ITR sequence.

[00243] In some embodiments, a modified ITR does not contain any nucleotide deletions in the RBE-containing portion of the A or A' regions, so as not to interfere with DNA replication (*e.g.* binding to an RBE by Rep protein, or nicking at a terminal resolution site). In some embodiments, a modified ITR encompassed for use herein has one or more deletions in the B, B', C, and/or C region as described herein.

[00244] In some embodiments, a ceDNA vector for expression of FIX protein comprising a symmetric ITR pair or asymmetric ITR pair comprises a regulatory switch as disclosed herein and at least one modified ITR selected having the nucleotide sequence selected from any of the group consisting of: SEQ ID NO: 3, 4, 15-47, 101-116 or 165-187.

[00245] In another embodiment, the structure of the structural element can be modified. For example, the structural element a change in the height of the stem and/or the number of nucleotides in the loop. For example, the height of the stem can be about 2, 3, 4, 5, 6, 7, 8, or 9 nucleotides or more or any range therein. In one embodiment, the stem height can be about 5 nucleotides to about 9 nucleotides and functionally interacts with Rep. In another embodiment, the stem height can be about 7 nucleotides and functionally interacts with Rep. In another example, the loop can have 3, 4, 5, 6, 7, 8, 9, or 10 nucleotides or more or any range therein.

[00246] In another embodiment, the number of GAGY binding sites or GAGY-related binding sites within the RBE or extended RBE can be increased or decreased. In one example, the RBE or extended RBE, can comprise 1, 2, 3, 4, 5, or 6 or more GAGY binding sites or any range therein. Each GAGY binding site can independently be an exact GAGY sequence or a sequence similar to GAGY as long as the sequence is sufficient to bind a Rep protein.

[00247] In another embodiment, the spacing between two elements (such as but not limited to the RBE and a hairpin) can be altered (*e.g.*, increased or decreased) to alter functional interaction with a large Rep protein. For example, the spacing can be about 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, or 21 nucleotides or more or any range therein.

[00248] The ceDNA vector for expression of FIX protein as described herein can include an ITR structure that is modified with respect to the wild type AAV2 ITR structure disclosed herein, but still retains an operable RBE, trs and RBE' portion. **FIG. 2A** and **FIG. 2B** show one possible mechanism for the operation of a trs site within a wild type ITR structure portion of a ceDNA vector for expression of FIX protein. In some embodiments, the ceDNA vector for expression of FIX protein contains one or more functional ITR polynucleotide sequences that comprise a Rep-binding site (RBS; 5'-GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60) for AAV2) and a terminal resolution site (TRS; 5'-AGTT (SEQ ID NO: 62)). In some embodiments, at least one ITR (wt or modified ITR) is functional. In alternative embodiments, where a ceDNA vector for expression of FIX protein comprises two

modified ITRs that are different or asymmetrical to each other, at least one modified ITR is functional and at least one modified ITR is non-functional.

[00249] In some embodiments, the modified ITR (*e.g.*, the left or right ITR) of a ceDNA vector for expression of FIX protein as described herein has modifications within the loop arm, the truncated arm, or the spacer. Exemplary sequences of ITRs having modifications within the loop arm, the truncated arm, or the spacer are listed in Table 2 (*i.e.*, SEQ ID NOS: 135-190, 200-233); Table 3 (*e.g.*, SEQ ID Nos: 234-263); Table 4 (*e.g.*, SEQ ID NOS: 264-293); Table 5 (*e.g.*, SEQ ID Nos: 294-318 herein); Table 6 (*e.g.*, SEQ ID NO: 319-468; and Tables 7-9 (*e.g.*, SEQ ID Nos: 101-110, 111-112, 115-134) or Table 10A or 10B (*e.g.*, SEQ ID Nos: 9, 100, 469-483, 484-499) of International Patent Application No. PCT/US18/49996, which is incorporated herein in its entirety by reference.

[00250] In some embodiments, the modified ITR for use in a ceDNA vector for expression of FIX protein comprising an asymmetric ITR pair, or symmetric mod-ITR pair is selected from any or a combination of those shown in Tables 2, 3, 4, 5, 6, 7, 8, 9 and 10A-10B of International Patent Application No. PCT/US18/49996 which is incorporated herein in its entirety by reference.

[00251] Additional exemplary modified ITRs for use in a ceDNA vector for expression of FIX protein comprising an asymmetric ITR pair, or symmetric mod-ITR pair in each of the above classes are provided in **Tables 5A** and **5B**. The predicted secondary structure of the Right modified ITRs in **Table 5A** are shown in FIG. 7A of International Patent Application No. PCT/US2018/064242, filed December 6, 2018, and the predicted secondary structure of the Left modified ITRs in **Table 5B** are shown in FIG. 7B of International Patent Application No. PCT/US2018/064242, filed December 6, 2018, which is incorporated herein in its entirety by reference.

[00252] **Table 5A** and **Table 5B** list the SEQ ID NOs of exemplary right and left modified ITRs.

Table 5A: Exemplary modified right ITRs. These exemplary modified right ITRs can comprise the RBE of GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60), spacer of ACTGAGGC (SEQ ID NO: 69), the spacer complement GCCTCAGT (SEQ ID NO: 70) and RBE' (*i.e.*, complement to RBE) of GAGCGAGCGAGCGCGC (SEQ ID NO: 71).

ITR Construct	SEQ ID NO:
ITR-18 Right	15
ITR-19 Right	16
ITR-20 Right	17
ITR-21 Right	18
ITR-22 Right	19
ITR-23 Right	20
ITR-24 Right	21
ITR-25 Right	22
ITR-26 Right	23
ITR-27 Right	24
ITR-28 Right	25
ITR-29 Right	26

ITR-30 Right	27
ITR-31 Right	28
ITR-32 Right	29
ITR-49 Right	30
ITR-50 right	31

TABLE 5B: Exemplary modified left ITRs. These exemplary modified left ITRs can comprise the RBE of GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60), spacer of ACTGAGGC (SEQ ID NO: 69), the spacer complement GCCTCAGT (SEQ ID NO: 70) and RBE complement (RBE') of GAGCGAGCGAGCGCGC (SEQ ID NO: 71).

ITR Construct	SEQ ID NO:
ITR-33 Left	32
ITR-34 Left	33
ITR-35 Left	34
ITR-36 Left	35
ITR-37 Left	36
ITR-38 Left	37
ITR-39 Left	38
ITR-40 Left	39
ITR-41 Left	40
ITR-42 Left	41
ITR-43 Left	42
ITR-44 Left	43
ITR-45 Left	44
ITR-46 Left	45
ITR-47 Left	46
ITR-48 Left	47

[00253] In one embodiment, a ceDNA vector for expression of FIX protein comprises, in the 5' to 3' direction: a first adeno-associated virus (AAV) inverted terminal repeat (ITR), a nucleic acid sequence of interest (for example an expression cassette as described herein) and a second AAV ITR, where the first ITR (5' ITR) and the second ITR (3' ITR) are asymmetric with respect to each other – that is, they have a different 3D-spatial configuration from one another. As an exemplary embodiment, the first ITR can be a wild-type ITR and the second ITR can be a mutated or modified ITR, or vice versa, where the first ITR can be a mutated or modified ITR and the second ITR a wild-type ITR. In some embodiment, the first ITR and the second ITR are both mod-ITRs, but have different sequences, or have different modifications, and thus are not the same modified ITRs, and have different 3D spatial configurations. Stated differently, a ceDNA vector with asymmetric ITRs comprises ITRs where any changes in one ITR relative to the WT-ITR are not reflected in the other ITR; or alternatively, where the asymmetric ITRs have a modified asymmetric ITR pair can have a different sequence and different three-dimensional shape with respect to each other. Exemplary asymmetric ITRs in the ceDNA vector

for expression of FIX protein and for use to generate a ceDNA-plasmid are shown in **Table 5A and 5B**.

[00254] In an alternative embodiment, a ceDNA vector for expression of FIX protein comprises two symmetrical mod-ITRs - that is, both ITRs have the same sequence, but are reverse complements (inverted) of each other. In some embodiments, a symmetrical mod-ITR pair comprises at least one or any combination of a deletion, insertion, or substitution relative to wild type ITR sequence from the same AAV serotype. The additions, deletions, or substitutions in the symmetrical ITR are the same but the reverse complement of each other. For example, an insertion of 3 nucleotides in the C region of the 5' ITR would be reflected in the insertion of 3 reverse complement nucleotides in the corresponding section in the C' region of the 3' ITR. Solely for illustration purposes only, if the addition is AACG in the 5' ITR, the addition is CGTT in the 3' ITR at the corresponding site. For example, if the 5' ITR sense strand is ATCGATCG with an addition of AACG between the G and A to result in the sequence ATCGAACGATCG (SEQ ID NO: 51). The corresponding 3' ITR sense strand is CGATCGAT (the reverse complement of ATCGATCG) with an addition of CGTT (*i.e.* the reverse complement of AACG) between the T and C to result in the sequence CGATCGTTTCGAT (SEQ ID NO: 49) (the reverse complement of ATCGAACGATCG) (SEQ ID NO: 51).

[00255] In alternative embodiments, the modified ITR pair are substantially symmetrical as defined herein - that is, the modified ITR pair can have a different sequence but have corresponding or the same symmetrical three-dimensional shape. For example, one modified ITR can be from one serotype and the other modified ITR be from a different serotype, but they have the same mutation (*e.g.*, nucleotide insertion, deletion or substitution) in the same region. Stated differently, for illustrative purposes only, a 5' mod-ITR can be from AAV2 and have a deletion in the C region, and the 3' mod-ITR can be from AAV5 and have the corresponding deletion in the C' region, and provided the 5' mod-ITR and the 3' mod-ITR have the same or symmetrical three-dimensional spatial organization, they are encompassed for use herein as a modified ITR pair.

[00256] In some embodiments, a substantially symmetrical mod-ITR pair has the same A, C-C' and B-B' loops in 3D space, *e.g.*, if a modified ITR in a substantially symmetrical mod-ITR pair has a deletion of a C-C' arm, then the cognate mod-ITR has the corresponding deletion of the C-C' loop and also has a similar 3D structure of the remaining A and B-B' loops in the same shape in geometric space of its cognate mod-ITR. By way of example only, substantially symmetrical ITRs can have a symmetrical spatial organization such that their structure is the same shape in geometrical space. This can occur, *e.g.*, when a G-C pair is modified, for example, to a C-G pair or vice versa, or A-T pair is modified to a T-A pair, or vice versa. Therefore, using the exemplary example above of modified 5' ITR as a ATCGAACGATCG (SEQ ID NO: 51), and modified 3' ITR as CGATCGTTTCGAT (SEQ ID NO: 49) (*i.e.*, the reverse complement of ATCGAACGATCG (SEQ ID NO: 51)), these modified ITRs would still be symmetrical if, for example, the 5' ITR had the sequence of ATCGAACCATTCG (SEQ ID NO: 50), where G in the addition is modified to C, and the substantially symmetrical 3' ITR has the

sequence of CGATCGTTTCGAT (SEQ ID NO: 49), without the corresponding modification of the T in the addition to a. In some embodiments, such a modified ITR pair are substantially symmetrical as the modified ITR pair has symmetrical stereochemistry.

[00257] **Table 6** shows exemplary symmetric modified ITR pairs (*i.e.*, a left modified ITRs and the symmetric right modified ITR) for use in a ceDNA vector for expression of FIX protein. The bold (red) portion of the sequences identify partial ITR sequences (*i.e.*, sequences of A-A', C-C' and B-B' loops), also shown in FIGS 31A-46B. These exemplary modified ITRs can comprise the RBE of GCGCGCTCGCTCGCTC-3' (SEQ ID NO: 60), spacer of ACTGAGGC (SEQ ID NO: 69), the spacer complement GCCTCAGT (SEQ ID NO: 70) and RBE' (*i.e.*, complement to RBE) of GAGCGAGCGAGCGCGC (SEQ ID NO: 71).

Table 6: Exemplary symmetric modified ITR pairs in a ceDNA vector for expression of FIX protein

LEFT modified ITR (modified 5' ITR)		Symmetric RIGHT modified ITR (modified 3' ITR)	
ITR-33 left	SEQ ID NO:32	ITR-18, right	SEQ ID NO: 15
ITR-34 left	SEQ ID NO: 33	ITR-51, right	SEQ ID NO: 48
ITR-35 left	SEQ ID NO: 34	ITR-19, right	SEQ ID NO: 16
ITR-36 left	SEQ ID NO: 35	ITR-20, right	SEQ ID NO: 17
ITR-37 left	SEQ ID NO: 36	ITR-21, right	SEQ ID NO: 18
ITR-38 left	SEQ ID NO: 37	ITR-22 right	SEQ ID NO: 19
ITR-39 left	SEQ ID NO: 38	ITR-23, right	SEQ ID NO: 20
ITR-40 left	SEQ ID NO: 39	ITR-24, right	SEQ ID NO: 21
ITR-41 left	SEQ ID NO: 40	ITR-25 right	SEQ ID NO: 22
ITR-42 left	SEQ ID NO: 41	ITR-26 right	SEQ ID NO: 23
ITR-43 left	SEQ ID NO: 42	ITR-27 right	SEQ ID NO: 24
ITR-44 left	SEQ ID NO: 43	ITR-28 right	SEQ ID NO: 25
ITR-45 left	SEQ ID NO: 44	ITR-29, right	SEQ ID NO: 26
ITR-46 left	SEQ ID NO: 45	ITR-30, right)	SEQ ID NO: 27
ITR-47, left	SEQ ID NO: 46	ITR-31, right	SEQ ID NO: 28
ITR-48, left	SEQ ID NO: 47	ITR-32 right	SEQ ID NO: 29

[00258] In some embodiments, a ceDNA vector for expression of FIX protein comprising an asymmetric ITR pair can comprise an ITR with a modification corresponding to any of the modifications in ITR sequences or ITR partial sequences shown in any one or more of **Tables 5A-5B** herein, or the sequences shown in FIG. 7A-7B of International Patent Application No.

PCT/US2018/064242, filed December 6, 2018, which is incorporated herein in its entirety, or disclosed in Tables 2, 3, 4, 5, 6, 7, 8, 9 or 10A-10B of International Patent Application No.

PCT/US18/49996 filed September 7, 2018 which is incorporated herein in its entirety by reference.

V. Exemplary ceDNA vectors

[00259] As described above, the present disclosure relates to recombinant ceDNA expression vectors and ceDNA vectors that encode FIX protein, comprising any one of: an asymmetrical ITR pair, a symmetrical ITR pair, or substantially symmetrical ITR pair as described above. In certain

embodiments, the disclosure relates to recombinant ceDNA vectors for expression of FIX protein having flanking ITR sequences and a transgene, where the ITR sequences are asymmetrical, symmetrical or substantially symmetrical relative to each other as defined herein, and the ceDNA further comprises a nucleic acid sequence of interest (for example an expression cassette comprising the nucleic acid of a transgene) located between the flanking ITRs, wherein said nucleic acid molecule is devoid of viral capsid protein coding sequences.

[00260] The ceDNA expression vector for expression of FIX protein may be any ceDNA vector that can be conveniently subjected to recombinant DNA procedures including nucleic acid sequence(s) as described herein, provided at least one ITR is altered. The ceDNA vectors for expression of FIX protein of the present disclosure are compatible with the host cell into which the ceDNA vector is to be introduced. In certain embodiments, the ceDNA vectors may be linear. In certain embodiments, the ceDNA vectors may exist as an extrachromosomal entity. In certain embodiments, the ceDNA vectors of the present disclosure may contain an element(s) that permits integration of a donor sequence into the host cell's genome. As used herein “transgene”, “nucleic acid sequence” and “heterologous nucleic acid sequence” are synonymous, and encode FIX protein, as described herein.

[00261] Referring now to **FIGS 1A-1G**, schematics of the functional components of two non-limiting plasmids useful in making a ceDNA vector for expression of FIX protein are shown. **FIG. 1A, 1B, 1D, 1F** show the construct of ceDNA vectors or the corresponding sequences of ceDNA plasmids for expression of FIX protein. ceDNA vectors are capsid-free and can be obtained from a plasmid encoding in this order: a first ITR, an expressible transgene cassette and a second ITR, where the first and second ITR sequences are asymmetrical, symmetrical or substantially symmetrical relative to each other as defined herein. ceDNA vectors for expression of FIX protein are capsid-free and can be obtained from a plasmid encoding in this order: a first ITR, an expressible transgene (protein or nucleic acid) and a second ITR, where the first and second ITR sequences are asymmetrical, symmetrical or substantially symmetrical relative to each other as defined herein. In some embodiments, the expressible transgene cassette includes, as needed: an enhancer/promoter, one or more homology arms, a donor sequence, a post-transcription regulatory element (*e.g.*, WPRE, *e.g.*, SEQ ID NO: 67)), and a polyadenylation and termination signal (*e.g.*, BGH polyA, *e.g.*, SEQ ID NO: 68).

[00262] **FIG. 5** is a gel confirming the production of ceDNA from multiple plasmid constructs using the method described in the Examples. The ceDNA is confirmed by a characteristic band pattern in the gel, as discussed with respect to **FIG. 4A** above and in the Examples.

A. Regulatory elements.

[00263] The ceDNA vectors for expression of FIX protein as described herein comprising an asymmetric ITR pair or symmetric ITR pair as defined herein, can further comprise a specific combination of cis-regulatory elements. The cis-regulatory elements include, but are not limited to, a promoter, a riboswitch, an insulator, a mir-regulatable element, a post-transcriptional regulatory

element, a tissue- and cell type-specific promoter and an enhancer. Exemplary promoters are listed in **Table 7**. Exemplary enhancers are listed in **Table 8**. In some embodiments, the ITR can act as the promoter for the transgene, *e.g.*, FIX protein. In some embodiments, the ceDNA vector for expression of FIX protein as described herein comprises additional components to regulate expression of the transgene, for example, regulatory switches as described herein, to regulate the expression of the transgene, or a kill switch, which can kill a cell comprising the ceDNA vector encoding FIX protein thereof. Regulatory elements, including Regulatory Switches that can be used in the present disclosure are more fully discussed in International Patent Application No. PCT/US18/49996, which is incorporated herein in its entirety by reference.

[00264] In embodiments, the second nucleic acid sequence includes a regulatory sequence, and a nucleic acid sequence encoding a nuclease. In certain embodiments the gene regulatory sequence is operably linked to the nucleic acid sequence encoding the nuclease. In certain embodiments, the regulatory sequence is suitable for controlling the expression of the nuclease in a host cell. In certain embodiments, the regulatory sequence includes a suitable promoter sequence, being able to direct transcription of a gene operably linked to the promoter sequence, such as a nucleic acid sequence encoding the nuclease(s) of the present disclosure. In certain embodiments, the second nucleic acid sequence includes an intron sequence linked to the 5' terminus of the nucleic acid sequence encoding the nuclease. In certain embodiments, an enhancer sequence is provided upstream of the promoter to increase the efficacy of the promoter. In certain embodiments, the regulatory sequence includes an enhancer and a promoter, wherein the second nucleic acid sequence includes an intron sequence upstream of the nucleic acid sequence encoding a nuclease, wherein the intron includes one or more nuclease cleavage site(s), and wherein the promoter is operably linked to the nucleic acid sequence encoding the nuclease.

[00265] The ceDNA vectors for expression of FIX protein produced synthetically, or using a cell-based production method as described herein in the Examples, can further comprise a specific combination of cis-regulatory elements such as WHP posttranscriptional regulatory element (WPRE) (*e.g.*, SEQ ID NO: 67) and BGH polyA (SEQ ID NO: 68). Suitable expression cassettes for use in expression constructs are not limited by the packaging constraint imposed by the viral capsid.

(i). Promoters:

[00266] It will be appreciated by one of ordinary skill in the art that promoters used in the ceDNA vectors for expression of FIX protein as disclosed herein should be tailored as appropriate for the specific sequences they are promoting. Sequence identifiers of exemplary promoters operatively linked to a transgene (*e.g.*, FIX) useful in a ceDNA vector are disclosed in **Table 7**, herein.

Table 7: Exemplary promoters

Description	Length	Tissue Specificity	CG Content	SEQ ID NO
chicken B-actin core promoter ; part of constitutive CAG promoter set	278	Constitutive	33	200
hAAT promoter ; part of HAAT promoter Set	348	Liver	12	201
CpG-free human EF1a core promoter (3' sequence AAGCTT may be a spacer/restriction enzyme cut site and was absorbed); part of CET promoter set	226	Constitutive	0	202
murine TTR liver specific promoter (3' CTCCTG may be spacer/restriction enzyme cut site and was absorbed); part of CRM8 VandenDriessche promoter set	225	Liver	5	203
HLP promoter derived from BMN270	143	Liver	5	204
Mutant TTR promoter derived from SPK-8011	222	Liver	4	205
TTR promoter derived from Sangamo CRMSBS2-Intron3	223	Liver	4	206
Endogenous hFIX promoter (-3000 to -1 of 5' flanking genomic sequence)	3000	Endogenous	21	207
hAAT promoter derived from hFIX	205	Liver	10	208
hAAT promoter derived from SPK9001	397	Liver	12	209
Endogenous hG6Pase promoter (-2864 to -1 of 5' Flanking)	2864	Endogenous (Liver)	28	210
Human Rhodopsin kinase (GRK1) promoter (1793-2087 of genbank entry AY327580)	295	Photoreceptors	11	211
Truncated hAAT Core promoter; Part of LP1 promoter set	206	Liver	10	212
Human EF-1a promoter (contains EF-1a intron A)	1179	Constitutive	94	213

hRK promoter- Nearly identical to human rhodopsin kinase (GRK1) promoter (1793-2087 of genbank entry AY327580), but with a few indels of unknown origin.	292	Photoreceptors	11	214
Interphotoreceptor retinoid-binding protein (IRBP) promoter sequence	1325	Photoreceptors	14	215
promoter set containing CpGmin CME Enhancer, SV40_Enhancer_Invivogen, and CpG-free hEF1a core promoter	883	Constitutive	0	216
promoter set containing SV40_Enhancer_Invivogen, CpG-free hEF1a core promoter, and CET Intron	639	Constitutive	0	217
CpGmin hAAT promoter Set; contains CpGmin APOe-CR hAAT enhancer, hAAT core promoter, and CpGmin hAAT-Intron	1272	Liver	24	218
LP1 promoter Set; contains hAAT-HCR_LP1_Enhancer, hAAT_LP1_promoter, and hAAT-Intron	547	Liver	14	219
Synthetic CRM8 TBG promoter set with 5 CpGs; contains 2 copies of HS-CRM8_SERP_Enhancer, TBG promoter, and MVM intron	709	Liver	5	220
TBG core promoter (Thyroxine Binding Globulin; Liver Specific)	460	Liver	1	221
Synthetic CRM8 LP1 promoter set with 18 CpGs; contains 2 copies of HS-CRM8_SERP_Enhancer, hAPO-HCR_LP1_Enhancer, hAAT_LP1_promoter, and hAAT-Intron	699	Liver	18	222
Synthetic mic/bik TBG promoter set; contains 2 copies of mic/bik enhancer, TBG core promoter; does not contain an intron	681	Liver	1	223
Synthetic human CEF1 promoter set; contains human_CMV_Enhancer and hEF1a core promoter	532	Constitutive	0	224
Synthetic human CEF1 promoter set; contains murine_CMV_Enhancer, human_CMV_Enhancer, and hEF1a core promoter (In that order)	955	Constitutive	0	225

Synthetic human CEF1 promoter set; contains human_CMV_Enhancer, murine_CMV_Enhancer, and hEF1a core promoter (In that order)	955	Constitutive	0	226
Constitutive promoter Set containing CMV enhancer, gB-actin_promoter, and CAG-intron	1923	Constitutive	192	227
hAAT promoter Set; contains APOe-CR hAAT enhancer, hAAT core promoter, and hAAT-intron (Composed of hAAT 5' UTR and modSV40 intron)	1272	Liver	26	228
CpG-free CET promoter Set; containing murine_CMV_Enhancer, hEF1a core promoter, and CET synthetic intron	826	Constitutive	0	229
Canonical VandenDriessche promoter set; contains 1 copy of HS-SERP_Enhancer, TTR liver specific promoter, and MVM intron	399	Liver	9	230
Constitutive promoter Set containing CMV enhancer and CMV promoter (no Intron)	654	Constitutive	33	231
Murine Phosphoglycerate Kinase (PGK) promoter	500	Constitutive	39	232
SV40 + Human albumin Invivogen promoter set; containing SV40 enhancer (Invivogen) and huAlb promoter (Invivogen)	450	Liver	3	233
CMV enhancer + Human albumin Invivogen promoter set; contains CMV enhancer and huAlb promoter (Invivogen)	594	Liver	22	234
Human UBC promoter	1210	Constitutive	95	235
Endogenous hGFAP promoter (5' 3kb region)	3000	Muller Cell	44	236
Endogenous hRLBP1 promoter (5' 3kb region)	3000	Muller Cell	32	237
Murine RPE65 promoter	718	RPE Cells	2	238
Rat EF-1a promoter	1313	Constitutive	102	239
Human EF-1a promoter Set composed of SV40_Enhancer_Oz and human_FullLength_EF1a promoter	1420	Constitutive	95	240

Rat EF-1a promoter Set composed of CMV_Enhancer and rat_FullLength_EF1a promoter	1831	Constitutive	124	241
Endogenous hABCB11 promoter (5' 3kb region)	3000	Endogenous (Liver)	21	242
Endogenous hFIX promoter (5' 3kb region)	3095	Endogenous (Liver)	37	243
Murine CD44 Promoter sequence	1807	Muller Cell	34	244
Endogenous hABCB4 promoter (5' 3kb region)	3000	Endogenous (Liver)	91	245
Human RPE65 Promoter (-742 : +15) of NG_008472.1	757	RPE Cells	1	246
tMCK Promoter. Triplet repeat of 2R5S enhancer sequence followed by [-80 : +7] of murine MCK promoter	720	Muscle	16	247
MHCK7 Promoter	772	Muscle	16	248
MCK Promoter derived from rAAVirh74.MCK GALGT2 (Serepta's dystroglycan modifying therapy to promote Utrophin usage). Derived from mouse MCK core enhancer (206bp) fused to the MCK core promoter (351bp)	558	Muscle	12	249
MCK Promoter/5pUTR derived from rAAVirh74.MCK GALGT2 (Serepta's dystroglycan modifying therapy to promote Utrophin usage)	766	Muscle	21	250
Contains MHCK7 Promoter linked to SV40intron	961	Muscle	25	251
Muscle Specific Promoter derived from the human Desmin gene. Contains a ~1.7kb human DES promoter/enhancer region extending from 1.7kb upstream of the transcription start site to 35bp downstream within exon I of DES.	1736	Muscle	39	252
CMV enhancer + CMV Promoter + 5pUTR + Kozak Used in Stargen pONY8.95CMVABCR construct	807	Constitutive	48	253

Endogenous hFIX ORF (-973 to -3)	973	Endogenous (Photoreceptors)	17	254
Muscle Specific CK8 Promoter	450	Muscle	9	255
Muscle Specific human cTnT_Promoter	455	Muscle	4	256
Endogenous hABCB4 promoter (5' 3050bp region)	3050	Endogenous (Liver)	91	257
Endogenous hUSH1b promoter (5' 3kb region)	3000	Endogenous (Photoreceptors)	49	258
Endogenous hUSH2a promoter (5' 3kb region)	3000	Endogenous (Photoreceptors)	21	259
CASI promoter set containing a CMV enhancer, ubiquitin C enhancer elements, and Chicken B-actin core promoter	1053	Constitutive	99	260
Endogenous hABCB4 promoter (5' 3kb region)	3000	Endogenous (Liver)	38	261
Endogenous hABCB4 promoter (5' 3.1kb region)	3102	Liver	33	262
Murine Albumin Promoter (muAlb Enhancer region + core muAlb Promoter)	2337	Liver	15	263
Chimeric Promoter hAPOe Enhancer + TBG core promoter + modSV40intron	1330	Liver	14	264
mCMV enhancer + EF-1a core promoter + SI 126 Intron	937	Constitutive	21	265
LSP Promoter #2 - Synthetic mTTRenh-promoter Shire	367	Liver	11	266
LSP Promoter #4 - HS-CRM8 2x SerpEnh TTRmin MVMIntron	468	Liver	9	267
LSP Promoter #5 - HS-CRM1 AlbEnh TTRmin MVM	426	Liver	7	268
LSP Promoter #6 - HS-CRM2 Apo4Enh TTRmin MVM	396	Liver	7	269
LSP Promoter #7 - HS-CRM10 Enh TTRmin MVM	495	Liver	6	270
LSP Promoter #8 - HS-CRM8 SerpEnh huTBGpro MVM	640	Liver	4	271
LSP Promoter #9 - HS-CRM1 AlbEnh huTBGpro MVM	667	Liver	3	272

LSP Promoter #10 - HS-CRM2 Apo4Enh huTBGpro MVM	637	Liver	3	273
LSP Promoter #11 - HS-CRM10 Enh huTBGpro MVM	736	Liver	2	274
LSP Promoter #12 - HS-CRM8 SerpEnh muAlbpro MVM	515	Liver	6	275
LSP Promoter #13 - HS-CRM1 AlbEnh muAlbpro MVM	542	Liver	5	276
LSP Promoter #14 - HS-CRM2 Apo4Enh muAlbpro MVM	512	Liver	5	277
LSP Promoter #15 - HS-CRM10 Enh muAlbpro MVM	611	Liver	4	278
LSP Promoter #16 - CRM8 SerpEnh huAlbpro MVM	355	Liver	5	279
LSP Promoter #17 - HS-CRM1 AlbEnh huAlbpro MVM	382	Liver	4	280
LSP Promoter #18 - HS-CRM2 Apo4Enh huAlbpro MVM	352	Liver	4	281
LSP Promoter #19 - HS-CRM10 Enh huAlbpro MVM	451	Liver	3	282
LSP Promoter #20 - HS-CRM8 SerpEnh huAATpro MVM	430	Liver	13	283
LSP Promoter #21 - HS-CRM1 AlbEnh huAATpro MVM	457	Liver	12	284
LSP Promoter #22 - HS-CRM2 Apo4Enh huAATpro MVM	427	Liver	12	285
LSP Promoter #23 - HS-CRM10 Enh huAATpro MVM	526	Liver	11	286
LSP Promoter #24 - HS-CRM8 SerpEnh huAATpro SV40in	435	Liver	14	287
LSP Promoter #25 - HS-CRM1 AlbEnh huAATpro SV40in	462	Liver	13	288
LSP Promoter #26 - HS-CRM2 Apo4Enh huAATpro SV40in	448	Liver	16	289
LSP Promoter #27 - HS-CRM10 Enh huAATpro SV40in	531	Liver	12	290
LSP Promoter #28 - HS-CRM8 SerpEnh huTBGpro SV40in	636	Liver	4	291

LSP Promoter #29 - HS-CRM1 AlbEnh huTBGpro SV40in	663	Liver	3	292
LSP Promoter #30 - HS-CRM2 Apo4Enh huTBGpro SV40in	633	Liver	3	293
LSP Promoter #31 - HS-CRM10 Enh huTBGpro SV40in	732	Liver	2	294
LSP Promoter #32 - AMPBenh2x- huTBGpro SV40in	762	Liver	4	295
LSP Promoter #33 - AMPBenh2x- huTBGpro MVM	766	Liver	4	296

[00267] Expression cassettes of the ceDNA vector for expression of FIX protein can include a promoter, *e.g.*, any of the promoter selected from **Table 7**, which can influence overall expression levels as well as cell-specificity. For transgene expression, *e.g.*, expression of FIX protein, they can include a highly active virus-derived immediate early promoter. Expression cassettes can contain tissue-specific eukaryotic promoters to limit transgene expression to specific cell types and reduce toxic effects and immune responses resulting from unregulated, ectopic expression. In some embodiments, an expression cassette can contain a promoter or synthetic regulatory element, such as a CAG promoter (SEQ ID NO: 72). The CAG promoter comprises (i) the cytomegalovirus (CMV) early enhancer element, (ii) the promoter, the first exon and the first intron of chicken beta-actin gene, and (iii) the splice acceptor of the rabbit beta-globin gene. Alternatively, an expression cassette can contain an Alpha-1-antitrypsin (AAT) promoter (SEQ ID NO: 73 or SEQ ID NO: 74), a liver specific (LP1) promoter (SEQ ID NO: 75 or SEQ ID NO: 76), or a Human elongation factor-1 alpha (EF1a) promoter (*e.g.*, SEQ ID NO: 77 or SEQ ID NO: 78). In some embodiments, the expression cassette includes one or more constitutive promoters, for example, a retroviral Rous sarcoma virus (RSV) LTR promoter (optionally with the RSV enhancer), or a cytomegalovirus (CMV) immediate early promoter (optionally with the CMV enhancer, *e.g.*, SEQ ID NO: 79). Alternatively, an inducible promoter, a native promoter for a transgene, a tissue-specific promoter, or various promoters known in the art can be used.

[00268] Suitable promoters, including those described in **Table 7** and above, can be derived from viruses and can therefore be referred to as viral promoters, or they can be derived from any organism, including prokaryotic or eukaryotic organisms. Suitable promoters can be used to drive expression by any RNA polymerase (*e.g.*, pol I, pol II, pol III). Exemplary promoters include, but are not limited to the SV40 early promoter, mouse mammary tumor virus long terminal repeat (LTR) promoter; adenovirus major late promoter (Ad MLP); a herpes simplex virus (HSV) promoter, a cytomegalovirus (CMV) promoter such as the CMV immediate early promoter region (CMVIE), a rous sarcoma virus (RSV) promoter, a human U6 small nuclear promoter (U6, *e.g.*, SEQ ID NO: 80) (Miyagishi *et al.*,

Nature Biotechnology 20, 497-500 (2002)), an enhanced U6 promoter (*e.g.*, Xia *et al.*, *Nucleic Acids Res.* 2003 Sep. 1; 31(17)), a human H1 promoter (H1) (*e.g.*, SEQ ID NO: 81 or SEQ ID NO: 155), a CAG promoter, a human alpha 1-antitrypsin (HAAT) promoter (*e.g.*, SEQ ID NO: 82), and the like. In certain embodiments, these promoters are altered at their downstream intron containing end to include one or more nuclease cleavage sites. In certain embodiments, the DNA containing the nuclease cleavage site(s) is foreign to the promoter DNA.

[00269] In one embodiment, the promoter used is the native promoter of the gene encoding the therapeutic protein. The promoters and other regulatory sequences for the respective genes encoding the therapeutic proteins are known and have been characterized. The promoter region used may further include one or more additional regulatory sequences (*e.g.*, native), *e.g.*, enhancers, (*e.g.* SEQ ID NO: 79 and SEQ ID NO: 83), including a SV40 enhancer (SEQ ID NO: 126).

[00270] In some embodiments, a promoter may also be a promoter from a human gene such as human ubiquitin C (hUbC), human actin, human myosin, human hemoglobin, human muscle creatine, or human metallothionein. The promoter may also be a tissue specific promoter, such as a liver specific promoter, such as human alpha 1-antitrypsin (HAAT), natural or synthetic. In one embodiment, delivery to the liver can be achieved using endogenous ApoE specific targeting of the composition comprising a ceDNA vector to hepatocytes via the low density lipoprotein (LDL) receptor present on the surface of the hepatocyte.

[00271] Non-limiting examples of suitable promoters for use in accordance with the present disclosure include any of the promoters listed in **Table 7**, or any of the following: the CAG promoter of, for example (SEQ ID NO: 72), the HAAT promoter (SEQ ID NO: 82), the human EF1- α promoter (SEQ ID NO: 77) or a fragment of the EF1 α promoter (SEQ ID NO: 78), IE2 promoter (*e.g.*, SEQ ID NO: 84) and the rat EF1- α promoter (SEQ ID NO: 85), mEF1 promoter (SEQ ID NO: 59), or 1E1 promoter fragment (SEQ ID NO: 125).

[00272] (ii) *Enhancers*

[00273] In some embodiments, a ceDNA expressing FIX comprises one or more enhancers. In some embodiments, an enhancer sequence is located 5' of the promoter sequence. In some embodiments, the enhancer sequence is located 3' of the promoter sequence. Sequence identifiers of exemplary enhancers are listed in **Table 8** herein.

Table 8: Exemplary enhancer sequences

Description	Length	Tissue Specificity	CG Content	SEQ ID NO:
cytomegalovirus enhancer	518	Constitutive	22	300
Human apolipoprotein E/C-I liver specific enhancer	777	Liver	13	301
CpG-free Murine CMV enhancer	427	Constitutive	0	302
HS-CRM8 SERP enhancer	83	Liver	4	303

Human apolipoprotein E/C-I liver specific enhancer	777	Liver	12	304
34bp APOe/c-1 Enhancer and 32bp AAT X-region	66	Liver	1	305
Insulting sequence and hAPO-HCR Enhancer	212	Liver	4	306
hAPO-HCR Enhancer derived from SPK9001	330	Liver	4	307
hAPO-HCR Enhancer	194	Liver	3	308
SV40 Enhancer Invivogen	240	Constitutive	0	309
HS-CRM8 SERP enhancer with all spacers/cutsites removed	73	Liver	2	310
Alpha mic/bik Enhancer	100	Liver	0	311
CpG-free Human CMV Enhancer v2	296	Constitutive	0	312
SV40 Enhancer	235	Constitutive	1	313

(iii) 5' UTR sequences and intron sequences

[00274] In some embodiments, a ceDNA vector comprises a 5' UTR sequence and/or an intron sequence that located 3' of the 5' ITR sequence. In some embodiments, the 5' UTR is located 5' of the transgene, *e.g.*, sequence encoding the FIX protein. Sequence identifiers of exemplary 5' UTR sequences listed in **Table 9A**.

Table 9A: Exemplary 5' UTR sequences and intron sequences

Description	Length	Reference	CG Content	SEQ ID NO:
synthetic 5' UTR element composed of chicken B-actin 5'UTR/Intron and rabbit B-globin intron and 1st exon	1127		137	315
modified SV40 Intron	93		0	316
5' UTR of hAAT just upstream of ORF (3' CGGA may be spacer/restriction enzyme cut site, and was absorbed into the sequence)	54		1	317
CET promotor set synthetic intron	173		0	318
Minute Virus Mice (MVM) Intron	91		0	319
5' UTR of hAAT	54		0	320
5' UTR of hAAT combined with modSV40 intron	147		1	321
5' UTR of hAAT (3' TAATTA may be spacer/restriction enzyme cut site, and was	147		0	322

absorbed into the sequence) combined with modSV40 intron				
42bp of 5' UTR of AAT derived from BMN270 - includes Kozak	48	https://www.ncbi.nlm.nih.gov/pubmed/29292164	1	323
Intron/Enhancer from EF1a1	128	US2017/0216408	6	324
Synthetic SBR intron derived from Sangamo CRMSBS2-Intron3 -- includes kozak	98	WO2017074526	2	325
Endogenous hFIX 5' UTR	172	NG_011403.1	0	326
hAAT 5' UTR + modSV40 + kozak	160	http://www.bloodjournal.org/content/early/2005/12/01/blood-2005-10-4035?sso-checked=true	1	327
hFIX 5' UTR and Kozak	29	US20160375110	0	328
Chimeric Intron	133	U47119.2	2	329
Large fragment of Human Alpha-1 Antitrypsin (AAT) 5' UTR	341		9	330
5pUTR	316	US9644216	6	331
Human cDNA ABCB4 5pUTR (Variant A, predominant Isoform)	76	NM_000443	8	332
Human cDNA ABCB11 5pUTR	127	NM_003742	2	333
Human G6Pase 5pUTR	80	NM_000151.3	0	334
MCK 5pUTR derived from rAAVrh74.MCK GALGT2. Contains 53bp of endogenous mouse MCK Exon1 (untranslated), SV40 late 16S/19S splice signals, 5pUTR derived from plasmid pCMVB.	208	https://patentimages.storage.googleapis.com/4f/8a/d6/b915c650f5eeb5/WO2017049031A1.pdf	8	335
CpG Free 5' UTR synthetic (SI 126) Intron	159		0	336
5' UTR of Human Cytochrome b-245 alpha chain (CYBA) gene	36	(NM_000101.4)	5	337
5' UTR of Human 2,4-dienoyl-CoA reductase 1 (DECRL1) gene	141	(NM_001330575.1)	14	338
5' UTR of Human glia maturation factor gamma (GMFG) gene	110	(NM_001301008.1)	4	339
5' UTR of Human late endosomal/lysosomal adaptor, MAPK and MTOR activator 2 (LAMTOR2)	164	(NM_001145264.1)	13	340
5' UTR of Human myosin light chain 6B (MYL6B)	127	(NM_002475.4)	8	341
Large fragment of Human Alpha-1 Antitrypsin (AAT) 5' UTR	341		9	342

[00275] (iv) 3' UTR sequences

[00276] In some embodiments, a cDNA vector comprises a 3' UTR sequence that located 5' of the 3' ITR sequence. In some embodiments, the 3' UTR is located 3' of the transgene, *e.g.*, sequence encoding the FIX protein. Sequence identifiers of exemplary 3' UTR sequences listed in **Table 9B**.

Table 9B: Exemplary 3' UTR sequences and intron sequences

Description	Length	Reference	CG Content	SEQ ID NO:
WHP Posttranscriptional Response Element	581		20	345
Triplet repeat of mir-142 binding site	77		1	346
hFIX 3' UTR and polyA spacer derived from SPK9001	88	US2016/037 5110	0	347
Human hemoglobin beta (HBB) 3pUTR	395		1	348
Interferon Beta S/MAR (Scaffold/matrix-associated Region)	800		0	349
Beta-Globulin MAR (Matrix-associated region)	407		0	350
Human Albumin 3' UTR Sequence	186		1	351
CpG minimized HBB 3pUTR	395		0	352
WHP Posttranscriptional Response Element. Missing 3' Cytosine.	580		20	353
3' UTR of Human Cytochrome b-245 alpha chain (CYBA) gene	64	(NM_00010 1.4)	5	354
Shortened WPRE3 sequence with minimal gamma and alpha elements	247	WPRE3 ref https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3975461/	10	355
Human hemoglobin beta (HBB) 3pUTR	144		1	356
First 62bp of WPRE 3pUTR element	62		1	357

(v). Polyadenylation Sequences:

[00277] A sequence encoding a polyadenylation sequence can be included in the ceDNA vector for expression of FIX protein to stabilize an mRNA expressed from the ceDNA vector, and to aid in nuclear export and translation. In one embodiment, the ceDNA vector does not include a polyadenylation sequence. In other embodiments, the ceDNA vector for expression of FIX protein includes at least 1, at least 2, at least 3, at least 4, at least 5, at least 10, at least 15, at least 20, at least 25, at least 30, at least 40, least 45, at least 50 or more adenine dinucleotides. In some embodiments, the polyadenylation sequence comprises about 43 nucleotides, about 40-50 nucleotides, about 40-55 nucleotides, about 45-50 nucleotides, about 35-50 nucleotides, or any range there between.

[00278] The expression cassettes can include any poly-adenylation sequence known in the art or a variation thereof. In some embodiments, a poly-adenylation (polyA) sequence is selected from any of those listed in **Table 10**. Other polyA sequences commonly known in the art can also be used, *e.g.*, including but not limited to, naturally occurring sequence isolated from bovine BGHpA (*e.g.*, SEQ ID NO: 68) or a virus SV40pA (*e.g.*, SEQ ID NO: 86), or a synthetic sequence (*e.g.*, SEQ ID NO: 87). Some expression cassettes can also include SV40 late polyA signal upstream enhancer (USE)

sequence. In some embodiments, a USE sequence can be used in combination with SV40pA or heterologous poly-A signal. PolyA sequences are located 3' of the transgene encoding the FIX protein.

[00279] The expression cassettes can also include a post-transcriptional element to increase the expression of a transgene. In some embodiments, Woodchuck Hepatitis Virus (WHP) posttranscriptional regulatory element (WPRE) (*e.g.*, SEQ ID NO: 67) is used to increase the expression of a transgene. Other posttranscriptional processing elements such as the post-transcriptional element from the thymidine kinase gene of herpes simplex virus, or hepatitis B virus (HBV) can be used. Secretory sequences can be linked to the transgenes, *e.g.*, VH-02 and VK-A26 sequences, *e.g.*, SEQ ID NO: 88 and SEQ ID NO: 89.

Table 10: Sequence identifiers of exemplary polyA sequences

Description	Length	Reference	CG Content	SEQ ID NO:
bovine growth hormone Terminator and poly-adenylation sequence.	225		3	360
Synthetic polyA derived from BMN270	49	https://www.ncbi.nlm.nih.gov/pubmed/29292164	0	361
Synthetic polyA derived from SPK8011	54	US2017/0216408	2	362
Synthetic polyA and insulating sequence derived from Sangamo_CRMSBS2-Intron3	74	WO2017074526	2	363
SV40 Late polyA and 3' Insulating sequence derived from hFIX	143	http://www.bloodjournal.org/content/early/2005/12/01/blood-2005-10-4035?sso-checked=true	1	364
bGH polyA derived from SPK9001	228	US2016/0375110	0	365
CpGfree SV40 polyA	222		0	366
SV40 late polyA	226		0	367
C60pAC30HSL polyA containing A64 polyA sequence and C30 histone stem loop sequence	129		0	368
polyA used in J. Chou G6Pase constructs containing a SV40 polyA	232	US9644216	4	369
SV40 polyadenylation signal	135		0	370
herpesvirus thymidine kinase polyadenylation signal	49		4	371
SV40 late polyadenylation signal	226		0	372
Human Albumin 3' UTR and Terminator/polyA Sequence	416		2	373
Human Albumin 3' UTR and Terminator/polyA Sequence	415		2	374
CpGfree, Short SV40 polyA	122		0	375

CpGfree, Short SV40 polyA	133		0	376

(vi). *Nuclear Localization Sequences*

[00280] In some embodiments, the ceDNA vector for expression of FIX protein comprises one or more nuclear localization sequences (NLSs), for example, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more NLSs. In some embodiments, the one or more NLSs are located at or near the amino-terminus, at or near the carboxy-terminus, or a combination of these (*e.g.*, one or more NLS at the amino-terminus and/or one or more NLS at the carboxy terminus). When more than one NLS is present, each can be selected independently of the others, such that a single NLS is present in more than one copy and/or in combination with one or more other NLSs present in one or more copies. Sequence identifiers of non-limiting examples of NLSs are shown in **Table 11**.

Table 11: Nuclear Localization Sequence

SOURCE	SEQ ID NO.
SV40 virus large T-antigen	90
nucleoplasmin	92
c-myc	93
	94
hRNPA1 M9	95
IBB domain from importin-alpha	96
myoma T protein	97
	98
human p53	99
mouse c-abl IV	100
influenza virus NS1	117
	118
Hepatitis virus delta antigen	119
mouse Mx1 protein	120
human poly(ADP-ribose) polymerase	121
steroid hormone receptors (human) glucocorticoid	122

B. Additional Components of ceDNA vectors

[00281] The ceDNA vectors for expression of FIX protein of the present disclosure may contain nucleotides that encode other components for gene expression. For example, to select for specific gene targeting events, a protective shRNA may be embedded in a microRNA and inserted into a recombinant ceDNA vector designed to integrate site-specifically into the highly active locus, such as an albumin locus. Such embodiments may provide a system for *in vivo* selection and expansion of gene-modified hepatocytes in any genetic background such as described in Nygaard *et al.*, *A universal system to select gene-modified hepatocytes in vivo*, *Gene Therapy*, June 8, 2016. The ceDNA vectors of the present disclosure may contain one or more selectable markers that permit selection of transformed, transfected, transduced, or the like cells. A selectable marker is a gene the product of

which provides for biocide or viral resistance, resistance to heavy metals, prototrophy to auxotrophs, NeoR, and the like. In certain embodiments, positive selection markers are incorporated into the donor sequences such as NeoR. Negative selection markers may be incorporated downstream the donor sequences, for example a nucleic acid sequence HSV-tk encoding a negative selection marker may be incorporated into a nucleic acid construct downstream the donor sequence.

C. Regulatory Switches

[00282] A molecular regulatory switch is one which generates a measurable change in state in response to a signal. Such regulatory switches can be usefully combined with the ceDNA vectors for expression of FIX protein as described herein to control the output of expression of FIX protein from the ceDNA vector. In some embodiments, the ceDNA vector for expression of FIX protein comprises a regulatory switch that serves to fine tune expression of the FIX protein. For example, it can serve as a biocontainment function of the ceDNA vector. In some embodiments, the switch is an “ON/OFF” switch that is designed to start or stop (*i.e.*, shut down) expression of FIX protein in the ceDNA vector in a controllable and regulatable fashion. In some embodiments, the switch can include a “kill switch” that can instruct the cell comprising the ceDNA vector to undergo cell programmed death once the switch is activated. Exemplary regulatory switches encompassed for use in a ceDNA vector for expression of FIX protein can be used to regulate the expression of a transgene, and are more fully discussed in International application PCT/US18/49996, which is incorporated herein in its entirety by reference

(i) Binary Regulatory Switches

[00283] In some embodiments, the ceDNA vector for expression of FIX protein comprises a regulatory switch that can serve to controllably modulate expression of FIX protein. For example, the expression cassette located between the ITRs of the ceDNA vector may additionally comprise a regulatory region, *e.g.*, a promoter, cis-element, repressor, enhancer etc., that is operatively linked to the nucleic acid sequence encoding FIX protein, where the regulatory region is regulated by one or more cofactors or exogenous agents. By way of example only, regulatory regions can be modulated by small molecule switches or inducible or repressible promoters. Non-limiting examples of inducible promoters are hormone-inducible or metal-inducible promoters. Other exemplary inducible promoters/enhancer elements include, but are not limited to, an RU486-inducible promoter, an ecdysone-inducible promoter, a rapamycin-inducible promoter, and a metallothionein promoter.

(ii) Small molecule Regulatory Switches

[00284] A variety of art-known small-molecule based regulatory switches are known in the art and can be combined with the ceDNA vectors for expression of FIX protein as disclosed herein to form a regulatory-switch controlled ceDNA vector. In some embodiments, the regulatory switch can be selected from any one or a combination of: an orthogonal ligand/nuclear receptor pair, for example retinoid receptor variant/LG335 and GRQCIMFI, along with an artificial promoter controlling expression of the operatively linked transgene, such as that as disclosed in Taylor, et al. BMC

Biotechnology 10 (2010): 15; engineered steroid receptors, *e.g.*, modified progesterone receptor with a C-terminal truncation that cannot bind progesterone but binds RU486 (mifepristone) (US Patent No. 5,364,791); an ecdysone receptor from *Drosophila* and their ecdysteroid ligands (Saez, et al., PNAS, 97(26)(2000), 14512–14517; or a switch controlled by the antibiotic trimethoprim (TMP), as disclosed in Sando R 3rd; Nat Methods. 2013, 10(11):1085-8. In some embodiments, the regulatory switch to control the transgene or expressed by the ceDNA vector is a pro-drug activation switch, such as that disclosed in US patents 8,771,679, and 6,339,070.

(iii) “Passcode” Regulatory Switches

[00285] In some embodiments the regulatory switch can be a “passcode switch” or “passcode circuit”. Passcode switches allow fine tuning of the control of the expression of the transgene from the ceDNA vector when specific conditions occur – that is, a combination of conditions need to be present for transgene expression and/or repression to occur. For example, for expression of a transgene to occur at least conditions A and B must occur. A passcode regulatory switch can be any number of conditions, *e.g.*, at least 2, or at least 3, or at least 4, or at least 5, or at least 6 or at least 7 or more conditions to be present for transgene expression to occur. In some embodiments, at least 2 conditions (*e.g.*, A, B conditions) need to occur, and in some embodiments, at least 3 conditions need to occur (*e.g.*, A, B and C, or A, B and D). By way of an example only, for gene expression from a ceDNA to occur that has a passcode “ABC” regulatory switch, conditions A, B and C must be present.

Conditions A, B and C could be as follows; condition A is the presence of a condition or disease, condition B is a hormonal response, and condition C is a response to the transgene expression. For example, if the transgene edits a defective EPO gene, Condition A is the presence of Chronic Kidney Disease (CKD), Condition B occurs if the subject has hypoxic conditions in the kidney, Condition C is that Erythropoietin-producing cells (EPC) recruitment in the kidney is impaired; or alternatively, HIF-2 activation is impaired. Once the oxygen levels increase or the desired level of EPO is reached, the transgene turns off again until 3 conditions occur, turning it back on.

[00286] In some embodiments, a passcode regulatory switch or “Passcode circuit” encompassed for use in the ceDNA vector comprises hybrid transcription factors (TFs) to expand the range and complexity of environmental signals used to define biocontainment conditions. As opposed to a deadman switch which triggers cell death in the presence of a predetermined condition, the “passcode circuit” allows cell survival or transgene expression in the presence of a particular “passcode”, and can be easily reprogrammed to allow transgene expression and/or cell survival only when the predetermined environmental condition or passcode is present.

[00287] Any and all combinations of regulatory switches disclosed herein, *e.g.*, small molecule switches, nucleic acid-based switches, small molecule-nucleic acid hybrid switches, post-transcriptional transgene regulation switches, post-translational regulation, radiation-controlled switches, hypoxia-mediated switches and other regulatory switches known by persons of ordinary skill in the art as disclosed herein can be used in a passcode regulatory switch as disclosed herein.

Regulatory switches encompassed for use are also discussed in the review article Kis et al., J R Soc Interface. 12: 20141000 (2015), and summarized in Table 1 of Kis. In some embodiments, a regulatory switch for use in a passcode system can be selected from any or a combination of the switches disclosed in Table 11 of International Patent Application PCT/US18/49996, which is incorporated herein in its entirety by reference.

(iv). Nucleic acid-based regulatory switches to control transgene expression

[00288] In some embodiments, the regulatory switch to control the expression of FIX protein by the ceDNA is based on a nucleic-acid based control mechanism. Exemplary nucleic acid control mechanisms are known in the art and are envisioned for use. For example, such mechanisms include riboswitches, such as those disclosed in, *e.g.*, US2009/0305253, US2008/0269258, US2017/0204477, WO2018026762A1, US patent 9,222,093 and EP application EP288071, and also disclosed in the review by Villa JK et al., Microbiol Spectr. 2018 May;6(3). Also included are metabolite-responsive transcription biosensors, such as those disclosed in WO2018/075486 and WO2017/147585. Other art-known mechanisms envisioned for use include silencing of the transgene with an siRNA or RNAi molecule (*e.g.*, miR, shRNA). For example, the ceDNA vector can comprise a regulatory switch that encodes a RNAi molecule that is complementary to the to part of the transgene expressed by the ceDNA vector. When such RNAi is expressed even if the transgene (*e.g.*, FIX protein) is expressed by the ceDNA vector, it will be silenced by the complementary RNAi molecule, and when the RNAi is not expressed when the transgene is expressed by the ceDNA vector the transgene (*e.g.*, FIX protein) is not silenced by the RNAi.

[00289] In some embodiments, the regulatory switch is a tissue-specific self-inactivating regulatory switch, for example as disclosed in US2002/0022018, whereby the regulatory switch deliberately switches transgene (*e.g.*, FIX protein) off at a site where transgene expression might otherwise be disadvantageous. In some embodiments, the regulatory switch is a recombinase reversible gene expression system, for example as disclosed in US2014/0127162 and US Patent 8,324,436.

(v). Post-transcriptional and post-translational regulatory switches.

[00290] In some embodiments, the regulatory switch to control the expression of FIX protein by the ceDNA vector is a post-transcriptional modification system. For example, such a regulatory switch can be an aptazyme riboswitch that is sensitive to tetracycline or theophylline, as disclosed in US2018/0119156, GB201107768, WO2001/064956A3, EP Patent 2707487 and Beilstein et al., ACS Synth. Biol., 2015, 4 (5), pp 526–534; Zhong et al., Elife. 2016 Nov 2;5. pii: e18858. In some embodiments, it is envisioned that a person of ordinary skill in the art could encode both the transgene and an inhibitory siRNA which contains a ligand sensitive (OFF-switch) aptamer, the net result being a ligand sensitive ON-switch.

(vi). Other exemplary regulatory switches

[00291] Any known regulatory switch can be used in the ceDNA vector to control the expression of FIX protein by the ceDNA vector, including those triggered by environmental changes. Additional

examples include, but are not limited to; the BOC method of Suzuki et al., Scientific Reports 8; 10051 (2018); genetic code expansion and a non-physiologic amino acid; radiation-controlled or ultra-sound controlled on/off switches (see, *e.g.*, Scott S *et al.*, Gene Ther. 2000 Jul;7(13):1121-5; US patents 5,612,318; 5,571,797; 5,770,581; 5,817,636; and WO1999/025385A1. In some embodiments, the regulatory switch is controlled by an implantable system, *e.g.*, as disclosed in US patent 7,840,263; US2007/0190028A1 where gene expression is controlled by one or more forms of energy, including electromagnetic energy, that activates promoters operatively linked to the transgene in the ceDNA vector.

[00292] In some embodiments, a regulatory switch envisioned for use in the ceDNA vector is a hypoxia-mediated or stress-activated switch, *e.g.*, such as those disclosed in WO1999060142A2, US patent 5,834,306; 6,218,179; 6,709,858; US2015/0322410; Greco et al., (2004) Targeted Cancer Therapies 9, S368, as well as FROG, TOAD and NRSE elements and conditionally inducible silence elements, including hypoxia response elements (HREs), inflammatory response elements (IREs) and shear-stress activated elements (SSAEs), *e.g.*, as disclosed in U.S. Patent 9,394,526. Such an embodiment is useful for turning on expression of the transgene from the ceDNA vector after ischemia or in ischemic tissues, and/or tumors.

(vii). Kill Switches

[00293] Other embodiments described herein relate to a ceDNA vector for expression of FIX protein as described herein comprising a kill switch. A kill switch as disclosed herein enables a cell comprising the ceDNA vector to be killed or undergo programmed cell death as a means to permanently remove an introduced ceDNA vector from the subject's system. It will be appreciated by one of ordinary skill in the art that use of kill switches in the ceDNA vectors for expression of FIX protein would be typically coupled with targeting of the ceDNA vector to a limited number of cells that the subject can acceptably lose or to a cell type where apoptosis is desirable (*e.g.*, cancer cells). In all aspects, a "kill switch" as disclosed herein is designed to provide rapid and robust cell killing of the cell comprising the ceDNA vector in the absence of an input survival signal or other specified condition. Stated another way, a kill switch encoded by a ceDNA vector for expression of FIX protein as described herein can restrict cell survival of a cell comprising a ceDNA vector to an environment defined by specific input signals. Such kill switches serve as a biological biocontainment function should it be desirable to remove the ceDNA vector e expression of FIX protein in a subject or to ensure that it will not express the encoded FIX protein.

[00294] Other kill switches known to a person of ordinary skill in the art are encompassed for use in the ceDNA vector for expression of FIX protein as disclosed herein, *e.g.*, as disclosed in US2010/0175141; US2013/0009799; US2011/0172826; US2013/0109568, as well as kill switches disclosed in Jusiak et al, Reviews in Cell Biology and molecular Medicine; 2014; 1-56; Kobayashi et al., PNAS, 2004; 101; 8419-9; Marchisio et al., Int. Journal of Biochem and Cell Biol., 2011; 43; 310-319; and in Reinshagen et al., Science Translational Medicine, 2018, 11.

[00295] Accordingly, in some embodiments, the ceDNA vector for expression of FIX protein can comprise a kill switch nucleic acid construct, which comprises the nucleic acid encoding an effector toxin or reporter protein, where the expression of the effector toxin (*e.g.*, a death protein) or reporter protein is controlled by a predetermined condition. For example, a predetermined condition can be the presence of an environmental agent, such as, *e.g.*, an exogenous agent, without which the cell will default to expression of the effector toxin (*e.g.*, a death protein) and be killed. In alternative embodiments, a predetermined condition is the presence of two or more environmental agents, *e.g.*, the cell will only survive when two or more necessary exogenous agents are supplied, and without either of which, the cell comprising the ceDNA vector is killed.

[00296] In some embodiments, the ceDNA vector for expression of FIX protein is modified to incorporate a kill-switch to destroy the cells comprising the ceDNA vector to effectively terminate the *in vivo* expression of the transgene being expressed by the ceDNA vector (*e.g.*, expression of FIX protein). Specifically, the ceDNA vector is further genetically engineered to express a switch-protein that is not functional in mammalian cells under normal physiological conditions. Only upon administration of a drug or environmental condition that specifically targets this switch-protein, the cells expressing the switch-protein will be destroyed thereby terminating the expression of the therapeutic protein or peptide. For instance, it was reported that cells expressing HSV-thymidine kinase can be killed upon administration of drugs, such as ganciclovir and cytosine deaminase. See, for example, Dey and Evans, Suicide Gene Therapy by Herpes Simplex Virus-1 Thymidine Kinase (HSV-TK), in *Targets in Gene Therapy*, edited by You (2011); and Beltinger et al., *Proc. Natl. Acad. Sci. USA* 96(15):8699-8704 (1999). In some embodiments the ceDNA vector can comprise a siRNA kill switch referred to as DISE (Death Induced by Survival gene Elimination) (Murmman et al., *Oncotarget*. 2017; 8:84643-84658. Induction of DISE in ovarian cancer cells *in vivo*).

D. Exemplary ceDNA-FIX vectors

[00297] According to some embodiments, an exemplary ceDNA-FIX vector is selected from a ceDNA-FIX vector shown below in Table 12. According to some embodiments, the disclosure provides a capsid-free close-ended DNA (ceDNA) vector comprising at least one nucleic acid sequence between flanking inverted terminal repeats (ITRs), wherein at least one nucleic acid sequence encodes at least one FIX protein, and wherein the ceDNA vector is selected from a ceDNA-FIX vector shown in **Table 12**.

Table 12: ceDNA-FIX constructs

Construct	Sequence Identifier
ceDNA-FIX v1	SEQ ID NO: 404
ceDNA-FIX 2109	SEQ ID NO: 405
ceDNA-FIX 2112	SEQ ID NO: 406

ceDNA-FIX 2113	SEQ ID NO: 407
ceDNA-FIX 2114	SEQ ID NO: 408
ceDNA-FIX 2115	SEQ ID NO: 409
ceDNA-FIX 2116	SEQ ID NO: 410

[00298] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIXv1, comprising SEQ ID NO: 404. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 404.

[00299] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIX2109, comprising SEQ ID NO: 405. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 405.

[00300] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIX2112, comprising SEQ ID NO: 406. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 406.

[00301] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIX2113, comprising SEQ ID NO: 407. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 407.

[00302] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIX2114, comprising SEQ ID NO: 408. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 408.

[00303] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIX2115, comprising SEQ ID NO: 409. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 409.

[00304] According to some embodiments, the exemplary ceDNA vector is ceDNA-FIX2116, comprising SEQ ID NO: 410. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 410.

[00305] According to some embodiments, the disclosure provides a capsid-free close-ended DNA (ceDNA) vector comprising at least one nucleic acid sequence between flanking inverted terminal

repeats (ITRs), wherein at least one nucleic acid sequence encodes at least one FIX protein, and wherein the ceDNA vector comprises SEQ ID NO: 404. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 404.

[00306] According to some embodiments, the disclosure provides a capsid-free close-ended DNA (ceDNA) vector comprising at least one nucleic acid sequence between flanking inverted terminal repeats (ITRs), wherein at least one nucleic acid sequence encodes at least one FIX protein, and wherein the ceDNA vector comprises SEQ ID NO: 405. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 405.

[00307] According to some embodiments, the disclosure provides a capsid-free close-ended DNA (ceDNA) vector comprising at least one nucleic acid sequence between flanking inverted terminal repeats (ITRs), wherein at least one nucleic acid sequence encodes at least one FIX protein, and wherein the ceDNA vector comprises SEQ ID NO: 406. According to some embodiments, the ceDNA vector is at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98% or at least 99% identical to SEQ ID NO: 406.

[00308] According to some embodiments, SEQ ID NO: 404 comprises the following components, where the numbers indicate nucleic acid residues:

1..141=left-ITR_v1
 142..194 = spacer_left-ITR_v1
 195..524 = Enhancer
 525..921 =Promoter
 922..950 = 5pUTR
 951..1034 = hFIX Signal Peptide
 951..1037 = CDS
 951..3774 = Exon Intron_ORF
 1039..2476 = Intron
 2476..3774 = CDS "Translation 2476-3774"
 3538..3540 = R338L Padua Mutation
 3775..3862 = 3p UTR
 3863..4090 = poly A
 4091..4151 = spacer right-ITR

4152..4281 = right-ITR

[00309] According to some embodiments, SEQ ID NO: 405 comprises the following components, where the numbers indicate nucleic acid residues:

1..141 = left-ITR

142..183 = spacer_left-ITR_v.2.1

184..255 = SerpinEnhancer

184..837 = 3x Serpin-TTRe_PromoterSet

257..328 = Serpin Enhancer

330..401 = Serpin Enhancer

562..745 = Mouse TTR 5pUTR

714..745 = 5pUTR

746..836 = MVM intron

838..845 = PmeI_site

846..854 = Consensus Kozak

855..2240 = FIX-cDNA-691_2

2241..2248 = PacI site

2249..2829 = WPRE_3pUTR

2830..3054 = bGH

3055..3115 = spacer_right-ITR_v1

3116..3245 = right-ITR_v1

[00310] According to some embodiments, SEQ ID NO: 406 comprises the following components, where the numbers indicate nucleic acid residues:

1..141 = left-ITR_v1

142..183 = spacer_left-ITR_v2.1

184..255 = SerpinEnhancer

184..837 = 3x Serpin-TTRe_PromoterSet

257..328 = Serpin Enhancer

330..401 = Serpin Enhancer

562..745 = Mouse TTR 5pUTR (NM_013697.5)

714..745 = 5pUTR

746..836 = MVM Intron

838..845 = PmeI_site

846..854 = Consensus_Kozak

855..2240 =FIX-cDNA-691_16

2241..2248 =PacI_site

2249..2829 = WPRE_3pUTR

2830..3054 = bGH

3055..3115 = spacer_right-ITR_v1

3116..3245 = right-ITR_v1

VI. Detailed method of Production of a ceDNA Vector

A. Production in General

[00311] Certain methods for the production of a ceDNA vector for expression of FIX protein comprising an asymmetrical ITR pair or symmetrical ITR pair as defined herein is described in section IV of International application PCT/US18/49996 filed September 7, 2018, which is incorporated herein in its entirety by reference. In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein can be produced using insect cells, as described herein. In alternative embodiments, a ceDNA vector for expression of FIX protein as disclosed herein can be produced synthetically and in some embodiments, in a cell-free method, as disclosed on International Application PCT/US19/14122, filed January 18, 2019, which is incorporated herein in its entirety by reference.

[00312] As described herein, in one embodiment, a ceDNA vector for expression of FIX protein can be obtained, for example, by the process comprising the steps of: a) incubating a population of host cells (*e.g.* insect cells) harboring the polynucleotide expression construct template (*e.g.*, a ceDNA-plasmid, a ceDNA-Bacmid, and/or a ceDNA-baculovirus), which is devoid of viral capsid coding sequences, in the presence of a Rep protein under conditions effective and for a time sufficient to induce production of the ceDNA vector within the host cells, and wherein the host cells do not comprise viral capsid coding sequences; and b) harvesting and isolating the ceDNA vector from the host cells. The presence of Rep protein induces replication of the vector polynucleotide with a modified ITR to produce the ceDNA vector in a host cell. However, no viral particles (*e.g.* AAV virions) are expressed. Thus, there is no size limitation such as that naturally imposed in AAV or other viral-based vectors.

[00313] The presence of the ceDNA vector isolated from the host cells can be confirmed by digesting DNA isolated from the host cell with a restriction enzyme having a single recognition site on

the ceDNA vector and analyzing the digested DNA material on a non-denaturing gel to confirm the presence of characteristic bands of linear and continuous DNA as compared to linear and non-continuous DNA.

[00314] In yet another aspect, the disclosure provides for use of host cell lines that have stably integrated the DNA vector polynucleotide expression template (ceDNA template) into their own genome in production of the non-viral DNA vector, *e.g.* as described in Lee, L. *et al.* (2013) Plos One 8(8): e69879. Preferably, Rep is added to host cells at an MOI of about 3. When the host cell line is a mammalian cell line, *e.g.*, HEK293 cells, the cell lines can have polynucleotide vector template stably integrated, and a second vector such as herpes virus can be used to introduce Rep protein into cells, allowing for the excision and amplification of ceDNA in the presence of Rep and helper virus.

[00315] In one embodiment, the host cells used to make the ceDNA vectors for expression of FIX protein as described herein are insect cells, and baculovirus is used to deliver both the polynucleotide that encodes Rep protein and the non-viral DNA vector polynucleotide expression construct template for ceDNA, *e.g.*, as described in **FIGS. 4A-4C** and Example 1. In some embodiments, the host cell is engineered to express Rep protein.

[00316] The ceDNA vector is then harvested and isolated from the host cells. The time for harvesting and collecting ceDNA vectors described herein from the cells can be selected and optimized to achieve a high-yield production of the ceDNA vectors. For example, the harvest time can be selected in view of cell viability, cell morphology, cell growth, *etc.* In one embodiment, cells are grown and harvested a sufficient time after baculoviral infection to produce ceDNA vectors but before most cells start to die due to the baculoviral toxicity. The DNA vectors can be isolated using plasmid purification kits such as Qiagen Endo-Free Plasmid kits. Other methods developed for plasmid isolation can be also adapted for DNA vectors. Generally, any nucleic acid purification methods can be adopted.

[00317] The DNA vectors can be purified by any means known to those of skill in the art for purification of DNA. In one embodiment, ceDNA vectors are purified as DNA molecules. In another embodiment, the ceDNA vectors are purified as exosomes or microparticles.

[00318] The presence of the ceDNA vector for expression of FIX protein can be confirmed by digesting the vector DNA isolated from the cells with a restriction enzyme having a single recognition site on the DNA vector and analyzing both digested and undigested DNA material using gel electrophoresis to confirm the presence of characteristic bands of linear and continuous DNA as compared to linear and non-continuous DNA. **FIG. 4C** and **FIG. 4D** illustrate one embodiment for identifying the presence of the closed ended ceDNA vectors produced by the processes herein.

[00319] According to some embodiments, the ceDNA is synthetically produced in a cell-free environment.

B. ceDNA Plasmid

[00320] A ceDNA-plasmid is a plasmid used for later production of a ceDNA vector for expression of FIX protein. In some embodiments, a ceDNA-plasmid can be constructed using known techniques to provide at least the following as operatively linked components in the direction of transcription: (1) a modified 5' ITR sequence; (2) an expression cassette containing a *cis*-regulatory element, for example, a promoter, inducible promoter, regulatory switch, enhancers and the like; and (3) a modified 3' ITR sequence, where the 3' ITR sequence is symmetric relative to the 5' ITR sequence. In some embodiments, the expression cassette flanked by the ITRs comprises a cloning site for introducing an exogenous sequence. The expression cassette replaces the rep and cap coding regions of the AAV genomes.

[00321] In one aspect, a ceDNA vector for expression of FIX protein is obtained from a plasmid, referred to herein as a "ceDNA-plasmid" encoding in this order: a first adeno-associated virus (AAV) inverted terminal repeat (ITR), an expression cassette comprising a transgene, and a mutated or modified AAV ITR, wherein said ceDNA-plasmid is devoid of AAV capsid protein coding sequences. In alternative embodiments, the ceDNA-plasmid encodes in this order: a first (or 5') modified or mutated AAV ITR, an expression cassette comprising a transgene, and a second (or 3') modified AAV ITR, wherein said ceDNA-plasmid is devoid of AAV capsid protein coding sequences, and wherein the 5' and 3' ITRs are symmetric relative to each other. In alternative embodiments, the ceDNA-plasmid encodes in this order: a first (or 5') modified or mutated AAV ITR, an expression cassette comprising a transgene, and a second (or 3') mutated or modified AAV ITR, wherein said ceDNA-plasmid is devoid of AAV capsid protein coding sequences, and wherein the 5' and 3' modified ITRs have the same modifications (*i.e.*, they are inverse complement or symmetric relative to each other).

[00322] In a further embodiment, the ceDNA-plasmid system is devoid of viral capsid protein coding sequences (*i.e.* it is devoid of AAV capsid genes but also of capsid genes of other viruses). In addition, in a particular embodiment, the ceDNA-plasmid is also devoid of AAV Rep protein coding sequences. Accordingly, in a preferred embodiment, ceDNA-plasmid is devoid of functional AAV cap and AAV rep genes GG-3' for AAV2) plus a variable palindromic sequence allowing for hairpin formation.

[00323] A ceDNA-plasmid of the present disclosure can be generated using natural nucleic acid sequences of the genomes of any AAV serotypes well known in the art. In one embodiment, the ceDNA-plasmid backbone is derived from the AAV1, AAV2, AAV3, AAV4, AAV5, AAV 5, AAV7, AAV8, AAV9, AAV10, AAV 11, AAV12, AAVrh8, AAVrh10, AAV-DJ, and AAV-DJ8 genome. *E.g.*, NCBI: NC 002077; NC 001401; NC001729; NC001829; NC006152; NC 006260; NC 006261; Kotin and Smith, *The Springer Index of Viruses*, available at the URL maintained by Springer (at www.oesys.springer.de/viruses/database/mkchapter.asp?virID=42.04)(note -references to a URL or database refer to the contents of the URL or database as of the effective filing date of this application) In a particular embodiment, the ceDNA-plasmid backbone is derived from the AAV2

genome. In another particular embodiment, the ceDNA-plasmid backbone is a synthetic backbone genetically engineered to include at its 5' and 3' ITRs derived from one of these AAV genomes.

[00324] A ceDNA-plasmid can optionally include a selectable or selection marker for use in the establishment of a ceDNA vector-producing cell line. In one embodiment, the selection marker can be inserted downstream (*i.e.*, 3') of the 3' ITR sequence. In another embodiment, the selection marker can be inserted upstream (*i.e.*, 5') of the 5' ITR sequence. Appropriate selection markers include, for example, those that confer drug resistance. Selection markers can be, for example, a blasticidin S-resistance gene, kanamycin, geneticin, and the like. In a preferred embodiment, the drug selection marker is a blasticidin S-resistance gene.

[00325] An exemplary ceDNA (*e.g.*, rAAV0) vector for expression of FIX protein is produced from an rAAV plasmid. A method for the production of a rAAV vector, can comprise: (a) providing a host cell with a rAAV plasmid as described above, wherein both the host cell and the plasmid are devoid of capsid protein encoding genes, (b) culturing the host cell under conditions allowing production of an ceDNA genome, and (c) harvesting the cells and isolating the AAV genome produced from said cells.

C. Exemplary method of making the ceDNA vectors from ceDNA plasmids

[00326] Methods for making capsid-less ceDNA vectors for expression of FIX protein are also provided herein, notably a method with a sufficiently high yield to provide sufficient vector for *in vivo* experiments.

[00327] In some embodiments, a method for the production of a ceDNA vector for expression of FIX protein comprises the steps of: (1) introducing the nucleic acid construct comprising an expression cassette and two symmetric ITR sequences into a host cell (*e.g.*, Sf9 cells), (2) optionally, establishing a clonal cell line, for example, by using a selection marker present on the plasmid, (3) introducing a Rep coding gene (either by transfection or infection with a baculovirus carrying said gene) into said insect cell, and (4) harvesting the cell and purifying the ceDNA vector. The nucleic acid construct comprising an expression cassette and two ITR sequences described above for the production of ceDNA vector can be in the form of a ceDNA plasmid, or Bacmid or Baculovirus generated with the ceDNA plasmid as described below. The nucleic acid construct can be introduced into a host cell by transfection, viral transduction, stable integration, or other methods known in the art.

D. Cell lines

[00328] Host cell lines used in the production of a ceDNA vector for expression of FIX protein can include insect cell lines derived from *Spodoptera frugiperda*, such as Sf9 Sf21, or *Trichoplusia ni* cell, or other invertebrate, vertebrate, or other eukaryotic cell lines including mammalian cells. Other cell lines known to an ordinarily skilled artisan can also be used, such as HEK293, Huh-7, HeLa, HepG2, Hep1A, 911, CHO, COS, MeWo, NIH3T3, A549, HT1 180, monocytes, and mature and immature dendritic cells. Host cell lines can be transfected for stable expression of the ceDNA-plasmid for high yield ceDNA vector production.

[00329] CeDNA-plasmids can be introduced into Sf9 cells by transient transfection using reagents (*e.g.*, liposomal, calcium phosphate) or physical means (*e.g.*, electroporation) known in the art. Alternatively, stable Sf9 cell lines which have stably integrated the ceDNA-plasmid into their genomes can be established. Such stable cell lines can be established by incorporating a selection marker into the ceDNA -plasmid as described above. If the ceDNA -plasmid used to transfect the cell line includes a selection marker, such as an antibiotic, cells that have been transfected with the ceDNA-plasmid and integrated the ceDNA-plasmid DNA into their genome can be selected for by addition of the antibiotic to the cell growth media. Resistant clones of the cells can then be isolated by single-cell dilution or colony transfer techniques and propagated.

E. Isolating and Purifying ceDNA vectors

[00330] Examples of the process for obtaining and isolating ceDNA vectors are described in **FIGS. 4A-4E** and the specific examples below. ceDNA-vectors for expression of FIX protein disclosed herein can be obtained from a producer cell expressing AAV Rep protein(s), further transformed with a ceDNA-plasmid, ceDNA-bacmid, or ceDNA-baculovirus. Plasmids useful for the production of ceDNA vectors include plasmids that encode FIX protein, or plasmids encoding one or more REP proteins.

[00331] In one aspect, a polynucleotide encodes the AAV Rep protein (Rep 78 or 68) delivered to a producer cell in a plasmid (Rep-plasmid), a bacmid (Rep-bacmid), or a baculovirus (Rep-baculovirus). The Rep-plasmid, Rep-bacmid, and Rep-baculovirus can be generated by methods described above.

[00332] Methods to produce a ceDNA vector for expression of FIX protein are described herein. Expression constructs used for generating a ceDNA vector for expression of FIX protein as described herein can be a plasmid (*e.g.*, ceDNA-plasmids), a Bacmid (*e.g.*, ceDNA-bacmid), and/or a baculovirus (*e.g.*, ceDNA-baculovirus). By way of an example only, a ceDNA-vector can be generated from the cells co-infected with ceDNA-baculovirus and Rep-baculovirus. Rep proteins produced from the Rep-baculovirus can replicate the ceDNA-baculovirus to generate ceDNA-vectors. Alternatively, ceDNA vectors for expression of FIX protein can be generated from the cells stably transfected with a construct comprising a sequence encoding the AAV Rep protein (Rep78/52) delivered in Rep-plasmids, Rep-bacmids, or Rep-baculovirus. CeDNA-Baculovirus can be transiently transfected to the cells, be replicated by Rep protein and produce ceDNA vectors.

[00333] The bacmid (*e.g.*, ceDNA-bacmid) can be transfected into permissive insect cells such as Sf9, Sf21, Tni (*Trichoplusia ni*) cell, High Five cell, and generate ceDNA-baculovirus, which is a recombinant baculovirus including the sequences comprising the symmetric ITRs and the expression cassette. ceDNA-baculovirus can be again infected into the insect cells to obtain a next generation of the recombinant baculovirus. Optionally, the step can be repeated once or multiple times to produce the recombinant baculovirus in a larger quantity.

[00334] The time for harvesting and collecting ceDNA vectors for expression of FIX protein as described herein from the cells can be selected and optimized to achieve a high-yield production of the

ceDNA vectors. For example, the harvest time can be selected in view of cell viability, cell morphology, cell growth, etc. Usually, cells can be harvested after sufficient time after baculoviral infection to produce ceDNA vectors (*e.g.*, ceDNA vectors) but before majority of cells start to die because of the viral toxicity. The ceDNA-vectors can be isolated from the Sf9 cells using plasmid purification kits such as Qiagen ENDO-FREE PLASMID® kits. Other methods developed for plasmid isolation can be also adapted for ceDNA vectors. Generally, any art-known nucleic acid purification methods can be adopted, as well as commercially available DNA extraction kits.

[00335] Alternatively, purification can be implemented by subjecting a cell pellet to an alkaline lysis process, centrifuging the resulting lysate and performing chromatographic separation. As one non-limiting example, the process can be performed by loading the supernatant on an ion exchange column (*e.g.* SARTOBIND Q®) which retains nucleic acids, and then eluting (*e.g.* with a 1.2 M NaCl solution) and performing a further chromatographic purification on a gel filtration column (*e.g.* 6 fast flow GE). The capsid-free AAV vector is then recovered by, *e.g.*, precipitation.

[00336] In some embodiments, ceDNA vectors for expression of FIX protein can also be purified in the form of exosomes, or microparticles. It is known in the art that many cell types release not only soluble proteins, but also complex protein/nucleic acid cargoes via membrane microvesicle shedding (Cocucci et al, 2009; EP 10306226.1) Such vesicles include microvesicles (also referred to as microparticles) and exosomes (also referred to as nanovesicles), both of which comprise proteins and RNA as cargo. Microvesicles are generated from the direct budding of the plasma membrane, and exosomes are released into the extracellular environment upon fusion of multivesicular endosomes with the plasma membrane. Thus, ceDNA vector-containing microvesicles and/or exosomes can be isolated from cells that have been transduced with the ceDNA-plasmid or a bacmid or baculovirus generated with the ceDNA-plasmid.

[00337] Microvesicles can be isolated by subjecting culture medium to filtration or ultracentrifugation at 20,000 x g, and exosomes at 100,000 x g. The optimal duration of ultracentrifugation can be experimentally-determined and will depend on the particular cell type from which the vesicles are isolated. Preferably, the culture medium is first cleared by low-speed centrifugation (*e.g.*, at 2000 x g for 5-20 minutes) and subjected to spin concentration using, *e.g.*, an AMICON® spin column (Millipore, Watford, UK). Microvesicles and exosomes can be further purified via FACS or MACS by using specific antibodies that recognize specific surface antigens present on the microvesicles and exosomes. Other microvesicle and exosome purification methods include, but are not limited to, immunoprecipitation, affinity chromatography, filtration, and magnetic beads coated with specific antibodies or aptamers. Upon purification, vesicles are washed with, *e.g.*, phosphate-buffered saline. One advantage of using microvesicles or exosome to deliver ceDNA-containing vesicles is that these vesicles can be targeted to various cell types by including on their membranes proteins recognized by specific receptors on the respective cell types. (See also EP 10306226)

[00338] Another aspect of the disclosure herein relates to methods of purifying ceDNA vectors from host cell lines that have stably integrated a ceDNA construct into their own genome. In one embodiment, ceDNA vectors are purified as DNA molecules. In another embodiment, the ceDNA vectors are purified as exosomes or microparticles.

[00339] FIG. 5 of International application PCT/US18/49996 shows a gel confirming the production of ceDNA from multiple ceDNA-plasmid constructs using the method described in the Examples. The ceDNA is confirmed by a characteristic band pattern in the gel, as discussed with respect to FIG. 4D in the Examples.

VII. Pharmaceutical Compositions

[00340] In another aspect, pharmaceutical compositions are provided. The pharmaceutical composition comprises a ceDNA vector for expression of FIX protein as described herein and a pharmaceutically acceptable carrier or diluent.

[00341] The ceDNA vectors for expression of FIX protein as disclosed herein can be incorporated into pharmaceutical compositions suitable for administration to a subject for *in vivo* delivery to cells, tissues, or organs of the subject. Typically, the pharmaceutical composition comprises a ceDNA-vector as disclosed herein and a pharmaceutically acceptable carrier. For example, the ceDNA vectors for expression of FIX protein as described herein can be incorporated into a pharmaceutical composition suitable for a desired route of therapeutic administration (*e.g.*, parenteral administration). Passive tissue transduction via high pressure intravenous or intra-arterial infusion, as well as intracellular injection, such as intranuclear microinjection or intracytoplasmic injection, are also contemplated. Pharmaceutical compositions for therapeutic purposes can be formulated as a solution, microemulsion, dispersion, liposomes, or other ordered structure suitable to high ceDNA vector concentration. Sterile injectable solutions can be prepared by incorporating the ceDNA vector compound in the required amount in an appropriate buffer with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization including a ceDNA vector can be formulated to deliver a transgene in the nucleic acid to the cells of a recipient, resulting in the therapeutic expression of the transgene or donor sequence therein. The composition can also include a pharmaceutically acceptable carrier.

[00342] Pharmaceutically active compositions comprising a ceDNA vector for expression of FIX protein can be formulated to deliver a transgene for various purposes to the cell, *e.g.*, cells of a subject.

[00343] Pharmaceutical compositions for therapeutic purposes typically must be sterile and stable under the conditions of manufacture and storage. The composition can be formulated as a solution, microemulsion, dispersion, liposomes, or other ordered structure suitable to high ceDNA vector concentration. Sterile injectable solutions can be prepared by incorporating the ceDNA vector compound in the required amount in an appropriate buffer with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization.

[00344] A ceDNA vector for expression of FIX protein as disclosed herein can be incorporated into a pharmaceutical composition suitable for topical, systemic, intra-amniotic, intrathecal, intracranial, intra-arterial, intravenous, intralymphatic, intraperitoneal, subcutaneous, tracheal, intra-tissue (*e.g.*, intramuscular, intracardiac, intrahepatic, intrarenal, intracerebral), intrathecal, intravesical, conjunctival (*e.g.*, extra-orbital, intraorbital, retroorbital, intraretinal, subretinal, choroidal, sub-choroidal, intrastromal, intracameral and intravitreal), intracochlear, and mucosal (*e.g.*, oral, rectal, nasal) administration. Passive tissue transduction via high pressure intravenous or intraarterial infusion, as well as intracellular injection, such as intranuclear microinjection or intracytoplasmic injection, are also contemplated.

[00345] In some aspects, the methods provided herein comprise delivering one or more ceDNA vectors for expression of FIX protein as disclosed herein to a host cell. Also provided herein are cells produced by such methods, and organisms (such as animals, plants, or fungi) comprising or produced from such cells. Methods of delivery of nucleic acids can include lipofection, nucleofection, microinjection, biolistics, liposomes, immunoliposomes, polycation or lipid:nucleic acid conjugates, naked DNA, and agent-enhanced uptake of DNA. Lipofection is described in *e.g.*, U.S. Pat. Nos. 5,049,386, 4,946,787; and 4,897,355) and lipofection reagents are sold commercially (*e.g.*, Transfectam™ and Lipofectin™). Delivery can be to cells (*e.g.*, *in vitro* or *ex vivo* administration) or target tissues (*e.g.*, *in vivo* administration).

[00346] Various techniques and methods are known in the art for delivering nucleic acids to cells. For example, nucleic acids, such as ceDNA for expression of FIX protein can be formulated into lipid nanoparticles (LNPs), lipidoids, liposomes, lipid nanoparticles, lipoplexes, or core-shell nanoparticles. Typically, LNPs are composed of nucleic acid (*e.g.*, ceDNA) molecules, one or more ionizable or cationic lipids (or salts thereof), one or more non-ionic or neutral lipids (*e.g.*, a phospholipid), a molecule that prevents aggregation (*e.g.*, PEG or a PEG-lipid conjugate), and optionally a sterol (*e.g.*, cholesterol).

[00347] Another method for delivering nucleic acids, such as ceDNA for expression of FIX protein to a cell is by conjugating the nucleic acid with a ligand that is internalized by the cell. For example, the ligand can bind a receptor on the cell surface and internalized via endocytosis. The ligand can be covalently linked to a nucleotide in the nucleic acid. Exemplary conjugates for delivering nucleic acids into a cell are described, example, in WO2015/006740, WO2014/025805, WO2012/037254, WO2009/082606, WO2009/073809, WO2009/018332, WO2006/112872, WO2004/090108, WO2004/091515 and WO2017/177326.

[00348] Nucleic acids, such as ceDNA vectors for expression of FIX protein can also be delivered to a cell by transfection. Useful transfection methods include, but are not limited to, lipid-mediated transfection, cationic polymer-mediated transfection, or calcium phosphate precipitation. Transfection reagents are well known in the art and include, but are not limited to, TurboFect Transfection Reagent (Thermo Fisher Scientific), Pro-Ject Reagent (Thermo Fisher Scientific), TRANSPASS™ P Protein

Transfection Reagent (New England Biolabs), CHARIOT™ Protein Delivery Reagent (Active Motif), PROTEOJUICE™ Protein Transfection Reagent (EMD Millipore), 293fectin, LIPOFECTAMINE™ 2000, LIPOFECTAMINE™ 3000 (Thermo Fisher Scientific), LIPOFECTAMINE™ (Thermo Fisher Scientific), LIPOFECTIN™ (Thermo Fisher Scientific), DMRIE-C, CELLFECTIN™ (Thermo Fisher Scientific), OLIGOFECTAMINE™ (Thermo Fisher Scientific), LIPOFECTACE™, FUGENE™ (Roche, Basel, Switzerland), FUGENE™ HD (Roche), TRANSFECTAM™ (Transfectam, Promega, Madison, Wis.), TFX-10™ (Promega), TFX-20™ (Promega), TFX-50™ (Promega), TRANSFECTIN™ (BioRad, Hercules, Calif.), SILENTFECT™ (Bio-Rad), Effectene™ (Qiagen, Valencia, Calif.), DC-chol (Avanti Polar Lipids), GENEPORTER™ (Gene Therapy Systems, San Diego, Calif.), DHARMAFECT 1™ (Dharmacon, Lafayette, Colo.), DHARMAFECT 2™ (Dharmacon), DHARMAFECT 3™ (Dharmacon), DHARMAFECT 4™ (Dharmacon), ESCORT™ III (Sigma, St. Louis, Mo.), and ESCORT™ IV (Sigma Chemical Co.). Nucleic acids, such as ceDNA, can also be delivered to a cell via microfluidics methods known to those of skill in the art.

[00349] ceDNA vectors for expression of FIX protein as described herein can also be administered directly to an organism for transduction of cells *in vivo*. Administration is by any of the routes normally used for introducing a molecule into ultimate contact with blood or tissue cells including, but not limited to, injection, infusion, topical application and electroporation. Suitable methods of administering such nucleic acids are available and well known to those of skill in the art, and, although more than one route can be used to administer a particular composition, a particular route can often provide a more immediate and more effective reaction than another route.

[00350] Methods for introduction of a nucleic acid vector ceDNA vector for expression of FIX protein as disclosed herein can be delivered into hematopoietic stem cells, for example, by the methods as described, for example, in U.S. Pat. No. 5,928,638.

[00351] The ceDNA vectors for expression of FIX protein in accordance with the present disclosure can be added to liposomes for delivery to a cell or target organ in a subject. Liposomes are vesicles that possess at least one lipid bilayer. Liposomes are typically used as carriers for drug/therapeutic delivery in the context of pharmaceutical development. They work by fusing with a cellular membrane and repositioning its lipid structure to deliver a drug or active pharmaceutical ingredient (API). Liposome compositions for such delivery are composed of phospholipids, especially compounds having a phosphatidylcholine group, however these compositions may also include other lipids. Exemplary liposomes and liposome formulations, including but not limited to polyethylene glycol (PEG)-functional group containing compounds are disclosed in International Application PCT/US2018/050042, filed on September 7, 2018 and in International application PCT/US2018/064242, filed on December 6, 2018, *e.g.*, see the section entitled “Pharmaceutical Formulations”.

[00352] Various delivery methods known in the art or modification thereof can be used to deliver ceDNA vectors *in vitro* or *in vivo*. For example, in some embodiments, ceDNA vectors for expression

of FIX protein are delivered by making transient penetration in cell membrane by mechanical, electrical, ultrasonic, hydrodynamic, or laser-based energy so that DNA entrance into the targeted cells is facilitated. For example, a ceDNA vector can be delivered by transiently disrupting cell membrane by squeezing the cell through a size-restricted channel or by other means known in the art. In some cases, a ceDNA vector alone is directly injected as naked DNA into any one of: any one or more tissues selected from: liver, kidneys, gallbladder, prostate, adrenal gland, heart, intestine, lung, and stomach, skin, thymus, cardiac muscle or skeletal muscle. In some cases, a ceDNA vector is delivered by gene gun. Gold or tungsten spherical particles (1–3 μm diameter) coated with capsid-free AAV vectors can be accelerated to high speed by pressurized gas to penetrate into target tissue cells.

[00353] Compositions comprising a ceDNA vector for expression of FIX protein and a pharmaceutically acceptable carrier are specifically contemplated herein. In some embodiments, the ceDNA vector is formulated with a lipid delivery system, for example, liposomes as described herein. In some embodiments, such compositions are administered by any route desired by a skilled practitioner. The compositions may be administered to a subject by different routes including orally, parenterally, sublingually, transdermally, rectally, transmucosally, topically, via inhalation, via buccal administration, intrapleurally, intravenous, intra-arterial, intraperitoneal, subcutaneous, intramuscular, intranasal intrathecal, and intraarticular or combinations thereof. For veterinary use, the composition may be administered as a suitably acceptable formulation in accordance with normal veterinary practice. The veterinarian may readily determine the dosing regimen and route of administration that is most appropriate for a particular animal. The compositions may be administered by traditional syringes, needleless injection devices, “microprojectile bombardment gene guns”, or other physical methods such as electroporation (“EP”), hydrodynamic methods, or ultrasound.

[00354] In some cases, a ceDNA vector for expression of FIX protein is delivered by hydrodynamic injection, which is a simple and highly efficient method for direct intracellular delivery of any water-soluble compounds and particles into internal organs and skeletal muscle in an entire limb.

[00355] In some cases, ceDNA vectors for expression of FIX protein are delivered by ultrasound by making nanoscopic pores in membrane to facilitate intracellular delivery of DNA particles into cells of internal organs or tumors, so the size and concentration of plasmid DNA have great role in efficiency of the system. In some cases, ceDNA vectors are delivered by magnetofection by using magnetic fields to concentrate particles containing nucleic acid into the target cells.

[00356] In some cases, chemical delivery systems can be used, for example, by using nanomeric complexes, which include compaction of negatively charged nucleic acid by polycationic nanomeric particles, belonging to cationic liposome/micelle or cationic polymers. Cationic lipids used for the delivery method includes, but not limited to monovalent cationic lipids, polyvalent cationic lipids, guanidine containing compounds, cholesterol derivative compounds, cationic polymers, (*e.g.*, poly(ethylenimine), poly-L-lysine, protamine, other cationic polymers), and lipid-polymer hybrid.

A. Exosomes:

[00357] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is delivered by being packaged in an exosome. Exosomes are small membrane vesicles of endocytic origin that are released into the extracellular environment following fusion of multivesicular bodies with the plasma membrane. Their surface consists of a lipid bilayer from the donor cell's cell membrane, they contain cytosol from the cell that produced the exosome, and exhibit membrane proteins from the parental cell on the surface. Exosomes are produced by various cell types including epithelial cells, B and T lymphocytes, mast cells (MC) as well as dendritic cells (DC). Some embodiments, exosomes with a diameter between 10nm and 1 μ m, between 20nm and 500nm, between 30nm and 250nm, between 50nm and 100nm are envisioned for use. Exosomes can be isolated for a delivery to target cells using either their donor cells or by introducing specific nucleic acids into them. Various approaches known in the art can be used to produce exosomes containing capsid-free AAV vectors of the present disclosure.

B. Microparticle/Nanoparticles

[00358] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is delivered by a lipid nanoparticle. Generally, lipid nanoparticles comprise an ionizable amino lipid (*e.g.*, heptatriaconta-6,9,28,31-tetraen-19-yl 4-(dimethylamino)butanoate, DLin-MC3-DMA, a phosphatidylcholine (1,2-distearoyl-sn-glycero-3-phosphocholine, DSPC), cholesterol and a coat lipid (polyethylene glycol-dimyristolglycerol, PEG-DMG), for example as disclosed by Tam *et al.* (2013). *Advances in Lipid Nanoparticles for siRNA delivery*. Pharmaceuticals 5(3): 498-507.

[00359] In some embodiments, a lipid nanoparticle has a mean diameter between about 10 and about 1000 nm. In some embodiments, a lipid nanoparticle has a diameter that is less than 300 nm. In some embodiments, a lipid nanoparticle has a diameter between about 10 and about 300 nm. In some embodiments, a lipid nanoparticle has a diameter that is less than 200 nm. In some embodiments, a lipid nanoparticle has a diameter between about 25 and about 200 nm. In some embodiments, a lipid nanoparticle preparation (*e.g.*, composition comprising a plurality of lipid nanoparticles) has a size distribution in which the mean size (*e.g.*, diameter) is about 70 nm to about 200 nm, and more typically the mean size is about 100 nm or less.

[00360] Various lipid nanoparticles known in the art can be used to deliver ceDNA vector for expression of FIX protein as disclosed herein. For example, various delivery methods using lipid nanoparticles are described in U.S. Patent Nos. 9,404,127, 9,006,417 and 9,518,272.

[00361] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is delivered by a gold nanoparticle. Generally, a nucleic acid can be covalently bound to a gold nanoparticle or non-covalently bound to a gold nanoparticle (*e.g.*, bound by a charge-charge interaction), for example as described by Ding *et al.* (2014). *Gold Nanoparticles for Nucleic Acid Delivery*. Mol. Ther. 22(6); 1075-1083. In some embodiments, gold nanoparticle-nucleic acid conjugates are produced using methods described, for example, in U.S. Patent No. 6,812,334.

C. Conjugates

[00362] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is conjugated (*e.g.*, covalently bound to an agent that increases cellular uptake. An “agent that increases cellular uptake” is a molecule that facilitates transport of a nucleic acid across a lipid membrane. For example, a nucleic acid can be conjugated to a lipophilic compound (*e.g.*, cholesterol, tocopherol, *etc.*), a cell penetrating peptide (CPP) (*e.g.*, penetratin, TAT, Syn1B, *etc.*), and polyamines (*e.g.*, spermine). Further examples of agents that increase cellular uptake are disclosed, for example, in Winkler (2013). *Oligonucleotide conjugates for therapeutic applications*. Ther. Deliv. 4(7); 791-809.

[00363] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is conjugated to a polymer (*e.g.*, a polymeric molecule) or a folate molecule (*e.g.*, folic acid molecule). Generally, delivery of nucleic acids conjugated to polymers is known in the art, for example as described in WO2000/34343 and WO2008/022309. In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is conjugated to a poly(amide) polymer, for example as described by U.S. Patent No. 8,987,377. In some embodiments, a nucleic acid described by the disclosure is conjugated to a folic acid molecule as described in U.S. Patent No. 8,507,455.

[00364] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is conjugated to a carbohydrate, for example as described in U.S. Patent No. 8,450,467.

D. Nanocapsule

[00365] Alternatively, nanocapsule formulations of a ceDNA vector for expression of FIX protein as disclosed herein can be used. Nanocapsules can generally entrap substances in a stable and reproducible way. To avoid side effects due to intracellular polymeric overloading, such ultrafine particles (sized around 0.1 μm) should be designed using polymers able to be degraded *in vivo*. Biodegradable polyalkyl-cyanoacrylate nanoparticles that meet these requirements are contemplated for use.

E. Liposomes

[00366] The ceDNA vectors for expression of FIX protein in accordance with the present disclosure can be added to liposomes for delivery to a cell or target organ in a subject. Liposomes are vesicles that possess at least one lipid bilayer. Liposomes are typically used as carriers for drug/therapeutic delivery in the context of pharmaceutical development. They work by fusing with a cellular membrane and repositioning its lipid structure to deliver a drug or active pharmaceutical ingredient (API).

Liposome compositions for such delivery are composed of phospholipids, especially compounds having a phosphatidylcholine group, however these compositions may also include other lipids.

[00367] The formation and use of liposomes is generally known to those of skill in the art. Liposomes have been developed with improved serum stability and circulation half-times (U.S. Pat. No. 5,741,516). Further, various methods of liposome and liposome like preparations as potential drug carriers have been described (U.S. Pat. Nos. 5,567,434; 5,552,157; 5,565,213; 5,738,868 and 5,795,587).

F. Exemplary liposome and Lipid Nanoparticle (LNP) Compositions

[00368] The ceDNA vectors for expression of FIX protein in accordance with the present disclosure can be added to liposomes for delivery to a cell, *e.g.*, a cell in need of expression of the transgene. Liposomes are vesicles that possess at least one lipid bilayer. Liposomes are typically used as carriers for drug/ therapeutic delivery in the context of pharmaceutical development. They work by fusing with a cellular membrane and repositioning its lipid structure to deliver a drug or active pharmaceutical ingredient (API). Liposome compositions for such delivery are composed of phospholipids, especially compounds having a phosphatidylcholine group, however these compositions may also include other lipids.

[00369] Lipid nanoparticles (LNPs) comprising ceDNA vectors are disclosed in International Application PCT/US2018/050042, filed on September 7, 2018, and International Application PCT/US2018/064242, filed on December 6, 2018 which are incorporated herein in their entirety and envisioned for use in the methods and compositions for ceDNA vectors for expression of FIX protein as disclosed herein.

[00370] In some aspects, the disclosure provides for a liposome formulation that includes one or more compounds with a polyethylene glycol (PEG) functional group (so-called “PEG-ylated compounds”) which can reduce the immunogenicity/ antigenicity of, provide hydrophilicity and hydrophobicity to the compound(s) and reduce dosage frequency. Or the liposome formulation simply includes polyethylene glycol (PEG) polymer as an additional component. In such aspects, the molecular weight of the PEG or PEG functional group can be from 62 Da to about 5,000 Da.

[00371] In some aspects, the disclosure provides for a liposome formulation that will deliver an API with extended release or controlled release profile over a period of hours to weeks. In some related aspects, the liposome formulation may comprise aqueous chambers that are bound by lipid bilayers. In other related aspects, the liposome formulation encapsulates an API with components that undergo a physical transition at elevated temperature which releases the API over a period of hours to weeks.

[00372] In some aspects, the liposome formulation comprises sphingomyelin and one or more lipids disclosed herein. In some aspects, the liposome formulation comprises optisomes.

[00373] In some aspects, the disclosure provides for a liposome formulation that includes one or more lipids selected from: N-(carbonyl-methoxypolyethylene glycol 2000)-1,2-distearoyl-sn-glycero-3-phosphoethanolamine sodium salt, (distearoyl-sn-glycero-phosphoethanolamine), MPEG (methoxy polyethylene glycol)-conjugated lipid, HSPC (hydrogenated soy phosphatidylcholine); PEG (polyethylene glycol); DSPE (distearoyl-sn-glycero-phosphoethanolamine); DSPC (distearoylphosphatidylcholine); DOPC (dioleoylphosphatidylcholine); DPPG (dipalmitoylphosphatidylglycerol); EPC (egg phosphatidylcholine); DOPS (dioleoylphosphatidylserine); POPC (palmitoyl-oleoylphosphatidylcholine); SM (sphingomyelin); MPEG (methoxy polyethylene glycol); DMPC (dimyristoyl phosphatidylcholine); DMPG (dimyristoyl phosphatidylglycerol); DSPG (distearoylphosphatidylglycerol); DEPC

(dierucoylphosphatidylcholine); DOPE (dioleoly-sn-glycero-phosphoethanolamine). cholesteryl sulphate (CS), dipalmitoylphosphatidylglycerol (DPPG), DOPC (dioleoly-sn-glycero-phosphatidylcholine) or any combination thereof.

[00374] In some aspects, the disclosure provides for a liposome formulation comprising phospholipid, cholesterol and a PEG-ylated lipid in a molar ratio of 56:38:5. In some aspects, the liposome formulation's overall lipid content is from 2-16 mg/mL. In some aspects, the disclosure provides for a liposome formulation comprising a lipid containing a phosphatidylcholine functional group, a lipid containing an ethanolamine functional group and a PEG-ylated lipid. In some aspects, the disclosure provides for a liposome formulation comprising a lipid containing a phosphatidylcholine functional group, a lipid containing an ethanolamine functional group and a PEG-ylated lipid in a molar ratio of 3:0.015:2 respectively. In some aspects, the disclosure provides for a liposome formulation comprising a lipid containing a phosphatidylcholine functional group, cholesterol and a PEG-ylated lipid. In some aspects, the disclosure provides for a liposome formulation comprising a lipid containing a phosphatidylcholine functional group and cholesterol. In some aspects, the PEG-ylated lipid is PEG-2000-DSPE. In some aspects, the disclosure provides for a liposome formulation comprising DPPG, soy PC, MPEG-DSPE lipid conjugate and cholesterol.

[00375] In some aspects, the disclosure provides for a liposome formulation comprising one or more lipids containing a phosphatidylcholine functional group and one or more lipids containing an ethanolamine functional group. In some aspects, the disclosure provides for a liposome formulation comprising one or more: lipids containing a phosphatidylcholine functional group, lipids containing an ethanolamine functional group, and sterols, *e.g.* cholesterol. In some aspects, the liposome formulation comprises DOPC/ DEPC; and DOPE.

[00376] In some aspects, the disclosure provides for a liposome formulation further comprising one or more pharmaceutical excipients, *e.g.* sucrose and/or glycine.

[00377] In some aspects, the disclosure provides for a liposome formulation that is either unilamellar or multilamellar in structure. In some aspects, the disclosure provides for a liposome formulation that comprises multi-vesicular particles and/or foam-based particles. In some aspects, the disclosure provides for a liposome formulation that are larger in relative size to common nanoparticles and about 150 to 250 nm in size. In some aspects, the liposome formulation is a lyophilized powder. In some aspects, the disclosure provides for a liposome formulation that is made and loaded with ceDNA vectors disclosed or described herein, by adding a weak base to a mixture having the isolated ceDNA outside the liposome. This addition increases the pH outside the liposomes to approximately 7.3 and drives the API into the liposome. In some aspects, the disclosure provides for a liposome formulation having a pH that is acidic on the inside of the liposome. In such cases the inside of the liposome can be at pH 4-6.9, and more preferably pH 6.5. In other aspects, the disclosure provides for a liposome formulation made by using intra-liposomal drug stabilization technology. In such cases,

polymeric or non-polymeric highly charged anions and intra-liposomal trapping agents are utilized, *e.g.* polyphosphate or sucrose octasulfate.

[00378] In some aspects, the disclosure provides for a lipid nanoparticle comprising ceDNA and an ionizable lipid. For example, a lipid nanoparticle formulation that is made and loaded with ceDNA obtained by the process as disclosed in International Application PCT/US2018/050042, filed on September 7, 2018, which is incorporated herein. This can be accomplished by high energy mixing of ethanolic lipids with aqueous ceDNA at low pH which protonates the ionizable lipid and provides favorable energetics for ceDNA/lipid association and nucleation of particles. The particles can be further stabilized through aqueous dilution and removal of the organic solvent. The particles can be concentrated to the desired level.

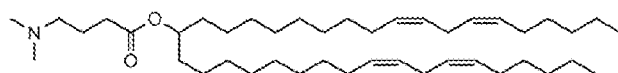
[00379] Generally, the lipid nanoparticles are prepared at a total lipid to ceDNA (mass or weight) ratio of from about 10:1 to 60:1. In some embodiments, the lipid to ceDNA ratio (mass/mass ratio; w/w ratio) can be in the range of from about 1:1 to about 60:1, from about 1:1 to about 55:1, from about 1:1 to about 50:1, from about 1:1 to about 45:1, from about 1:1 to about 40:1, from about 1:1 to about 35:1, from about 1:1 to about 30:1, from about 1:1 to about 25:1, from about 10:1 to about 14:1, from about 3:1 to about 15:1, from about 4:1 to about 10:1, from about 5:1 to about 9:1, about 6:1 to about 9:1; from about 30:1 to about 60:1. According to some embodiments, the lipid particles (*e.g.*, lipid nanoparticles) are prepared at a ceDNA (mass or weight) to total lipid ratio of about 60:1. According to some embodiments, the lipid particles are prepared at a total lipid to ceDNA (mass or weight) ratio of from about 10:1 to 30:1. In some embodiments, the lipid to ceDNA ratio (mass/mass ratio; w/w ratio) can be in the range of from about 1:1 to about 25:1, from about 10:1 to about 14:1, from about 3:1 to about 15:1, from about 4:1 to about 10:1, from about 5:1 to about 9:1, or about 6:1 to about 9:1. The amounts of lipids and ceDNA can be adjusted to provide a desired N/P ratio, for example, N/P ratio of 3, 4, 5, 6, 7, 8, 9, 10 or higher. Generally, the lipid particle formulation's overall lipid content can range from about 5 mg/ml to about 30 mg/mL.

[00380] The ionizable lipid is typically employed to condense the nucleic acid cargo, *e.g.*, ceDNA at low pH and to drive membrane association and fusogenicity. Generally, ionizable lipids are lipids comprising at least one amino group that is positively charged or becomes protonated under acidic conditions, for example at pH of 6.5 or lower. Ionizable lipids are also referred to as cationic lipids herein.

[00381] Exemplary ionizable lipids are described in International PCT patent publications WO2015/095340, WO2015/199952, WO2018/011633, WO2017/049245, WO2015/061467, WO2012/040184, WO2012/000104, WO2015/074085, WO2016/081029, WO2017/004143, WO2017/075531, WO2017/117528, WO2011/022460, WO2013/148541, WO2013/116126, WO2011/153120, WO2012/044638, WO2012/054365, WO2011/090965, WO2013/016058, WO2012/162210, WO2008/042973, WO2010/129709, WO2010/144740, WO2012/099755, WO2013/049328, WO2013/086322, WO2013/086373, WO2011/071860, WO2009/132131,

WO2010/048536, WO2010/088537, WO2010/054401, WO2010/054406, WO2010/054405, WO2010/054384, WO2012/016184, WO2009/086558, WO2010/042877, WO2011/000106, WO2011/000107, WO2005/120152, WO2011/141705, WO2013/126803, WO2006/007712, WO2011/038160, WO2005/121348, WO2011/066651, WO2009/127060, WO2011/141704, WO2006/069782, WO2012/031043, WO2013/006825, WO2013/033563, WO2013/089151, WO2017/099823, WO2015/095346, and WO2013/086354, and US patent publications US2016/0311759, US2015/0376115, US2016/0151284, US2017/0210697, US2015/0140070, US2013/0178541, US2013/0303587, US2015/0141678, US2015/0239926, US2016/0376224, US2017/0119904, US2012/0149894, US2015/0057373, US2013/0090372, US2013/0274523, US2013/0274504, US2013/0274504, US2009/0023673, US2012/0128760, US2010/0324120, US2014/0200257, US2015/0203446, US2018/0005363, US2014/0308304, US2013/0338210, US2012/0101148, US2012/0027796, US2012/0058144, US2013/0323269, US2011/0117125, US2011/0256175, US2012/0202871, US2011/0076335, US2006/0083780, US2013/0123338, US2015/0064242, US2006/0051405, US2013/0065939, US2006/0008910, US2003/0022649, US2010/0130588, US2013/0116307, US2010/0062967, US2013/0202684, US2014/0141070, US2014/0255472, US2014/0039032, US2018/0028664, US2016/0317458, and US2013/0195920, the contents of all of which are incorporated herein by reference in their entirety.

[00382] In some embodiments, the ionizable lipid is MC3 (6Z,9Z,28Z,31Z)-heptatriaconta-6,9,28,31-tetraen-19-yl-4-(dimethylamino) butanoate (DLin-MC3-DMA or MC3) having the following structure:



DLin-M-C3-DMA ("MC3")

[00383] The lipid DLin-MC3-DMA is described in Jayaraman *et al.*, Angew. Chem. Int. Ed Engl. (2012), 51(34): 8529-8533, content of which is incorporated herein by reference in its entirety.

[00384] In some embodiments, the ionizable lipid is the lipid ATX-002 as described in WO2015/074085, content of which is incorporated herein by reference in its entirety.

[00385] In some embodiments, the ionizable lipid is (13Z,16Z)-N,N-dimethyl-3-nonyldocosa-13,16-dien-1-amine (Compound 32), as described in WO2012/040184, content of which is incorporated herein by reference in its entirety.

[00386] In some embodiments, the ionizable lipid is Compound 6 or Compound 22 as described in WO2015/199952, content of which is incorporated herein by reference in its entirety.

[00387] Without limitations, ionizable lipid can comprise 20-90% (mol) of the total lipid present in the lipid nanoparticle. For example, ionizable lipid molar content can be 20-70% (mol), 30-60% (mol) or 40-50% (mol) of the total lipid present in the lipid nanoparticle. In some embodiments, ionizable

lipid comprises from about 50 mol % to about 90 mol % of the total lipid present in the lipid nanoparticle.

[00388] In some aspects, the lipid nanoparticle can further comprise a non-cationic lipid. Non-ionic lipids include amphipathic lipids, neutral lipids and anionic lipids. Accordingly, the non-cationic lipid can be a neutral uncharged, zwitterionic, or anionic lipid. Non-cationic lipids are typically employed to enhance fusogenicity.

[00389] Exemplary non-cationic lipids envisioned for use in the methods and compositions as disclosed herein are described in International Application PCT/US2018/050042, filed on September 7, 2018, and PCT/US2018/064242, filed on December 6, 2018 which is incorporated herein in its entirety. Exemplary non-cationic lipids are described in International Application Publication WO2017/099823 and US patent publication US2018/0028664, the contents of both of which are incorporated herein by reference in their entirety.

[00390] The non-cationic lipid can comprise 0-30% (mol) of the total lipid present in the lipid nanoparticle. For example, the non-cationic lipid content is 5-20% (mol) or 10-15% (mol) of the total lipid present in the lipid nanoparticle. In various embodiments, the molar ratio of ionizable lipid to the neutral lipid ranges from about 2:1 to about 8:1.

[00391] In some embodiments, the lipid nanoparticles do not comprise any phospholipids. In some aspects, the lipid nanoparticle can further comprise a component, such as a sterol, to provide membrane integrity.

[00392] One exemplary sterol that can be used in the lipid nanoparticle is cholesterol and derivatives thereof. Exemplary cholesterol derivatives are described in International application WO2009/127060 and US patent publication US2010/0130588, contents of both of which are incorporated herein by reference in their entirety.

[00393] The component providing membrane integrity, such as a sterol, can comprise 0-50% (mol) of the total lipid present in the lipid nanoparticle. In some embodiments, such a component is 20-50% (mol) 30-40% (mol) of the total lipid content of the lipid nanoparticle.

[00394] In some aspects, the lipid nanoparticle can further comprise a polyethylene glycol (PEG) or a conjugated lipid molecule. Generally, these are used to inhibit aggregation of lipid nanoparticles and/or provide steric stabilization. Exemplary conjugated lipids include, but are not limited to, PEG-lipid conjugates, polyoxazoline (POZ)-lipid conjugates, polyamide-lipid conjugates (such as ATTA-lipid conjugates), cationic-polymer lipid (CPL) conjugates, and mixtures thereof. In some embodiments, the conjugated lipid molecule is a PEG-lipid conjugate, for example, a (methoxy polyethylene glycol)-conjugated lipid. Exemplary PEG-lipid conjugates include, but are not limited to, PEG-diacylglycerol (DAG) (such as 1-(monomethoxy-polyethyleneglycol)-2,3-dimyristoylglycerol (PEG-DMG)), PEG-dialkoxypentyl (DAA), PEG-phospholipid, PEG-ceramide (Cer), a pegylated phosphatidylethanolamine (PEG-PE), PEG succinate diacylglycerol (PEGS-DAG) (such as 4-O-(2',3'-di(tetradecanoyloxy)propyl-1-O-(w-methoxy(polyethoxy)ethyl) butanedioate (PEG-S-DMG)),

PEG dialkoxypolypropylcarbam, N-(carbonyl-methoxypolyethylene glycol 2000)-1,2-distearoyl-sn-glycero-3-phosphoethanolamine sodium salt, or a mixture thereof. Additional exemplary PEG-lipid conjugates are described, for example, in US5,885,613, US6,287,591, US2003/0077829, US2003/0077829, US2005/0175682, US2008/0020058, US2011/0117125, US2010/0130588, US2016/0376224, and US2017/0119904, the contents of all of which are incorporated herein by reference in their entirety.

[00395] In some embodiments, a PEG-lipid is a compound as defined in US2018/0028664, the content of which is incorporated herein by reference in its entirety. In some embodiments, a PEG-lipid is disclosed in US20150376115 or in US2016/0376224, the content of both of which is incorporated herein by reference in its entirety.

[00396] The PEG-DAA conjugate can be, for example, PEG-dilauryloxypropyl, PEG-dimyristyloxypropyl, PEG-dipalmitoyloxypropyl, or PEG-distearoyloxypropyl. The PEG-lipid can be one or more of PEG-DMG, PEG-dilaurylglycerol, PEG-dipalmitoylglycerol, PEG-disteryl glycerol, PEG-dilaurylglycamide, PEG-dimyristylglycamide, PEG-dipalmitoylglycamide, PEG-disteryl glycamide, PEG-cholesterol (1-[8'-(Cholest-5-en-3[beta]-oxy)carboxamido-3',6'-dioxaoctanyl] carbamoyl-[omega]-methyl-poly(ethylene glycol)), PEG-DMB (3,4-Ditetradecoxylbenzyl- [omega]-methyl-poly(ethylene glycol) ether), and 1,2-dimyristoyl-sn-glycero-3-phosphoethanolamine-N-[methoxy(polyethylene glycol)-2000]. In some examples, the PEG-lipid can be selected from the group consisting of PEG-DMG, 1,2-dimyristoyl-sn-glycero-3-phosphoethanolamine-N-[methoxy(polyethylene glycol)-2000],

[00397] Lipids conjugated with a molecule other than a PEG can also be used in place of PEG-lipid. For example, polyoxazoline (POZ)-lipid conjugates, polyamide-lipid conjugates (such as ATTA-lipid conjugates), and cationic-polymer lipid (CPL) conjugates can be used in place of or in addition to the PEG-lipid. Exemplary conjugated lipids, *i.e.*, PEG-lipids, (POZ)-lipid conjugates, ATTA-lipid conjugates and cationic polymer-lipids are described in the International patent application publications WO1996/010392, WO1998/051278, WO2002/087541, WO2005/026372, WO2008/147438, WO2009/086558, WO2012/000104, WO2017/117528, WO2017/099823, WO2015/199952, WO2017/004143, WO2015/095346, WO2012/000104, WO2012/000104, and WO2010/006282, US patent application publications US2003/0077829, US2005/0175682, US2008/0020058, US2011/0117125, US2013/0303587, US2018/0028664, US2015/0376115, US2016/0376224, US2016/0317458, US2013/0303587, US2013/0303587, and US20110123453, and US patents US5,885,613, US6,287,591, US6,320,017, and US6,586,559, the contents of all of which are incorporated herein by reference in their entirety.

[00398] In some embodiments, the one or more additional compound can be a therapeutic agent. The therapeutic agent can be selected from any class suitable for the therapeutic objective. In other words, the therapeutic agent can be selected from any class suitable for the therapeutic objective. In other words, the therapeutic agent can be selected according to the treatment objective

and biological action desired. For example, if the ceDNA within the LNP is useful for treating hemophilia B, the additional compound can be an anti-hemophilia B agent (*e.g.*, a chemotherapeutic agent, or other hemophilia B therapy (including, but not limited to, a small molecule or an antibody). In another example, if the LNP containing the ceDNA is useful for treating an infection, the additional compound can be an antimicrobial agent (*e.g.*, an antibiotic or antiviral compound). In yet another example, if the LNP containing the ceDNA is useful for treating an immune disease or disorder, the additional compound can be a compound that modulates an immune response (*e.g.*, an immunosuppressant, immunostimulatory compound, or compound modulating one or more specific immune pathways). In some embodiments, different cocktails of different lipid nanoparticles containing different compounds, such as a ceDNA encoding a different protein or a different compound, such as a therapeutic may be used in the compositions and methods of the disclosure.

[00399] In some embodiments, the additional compound is an immune modulating agent. For example, the additional compound is an immunosuppressant. Immunosuppressants described herein include protein kinase inhibitors (PKIs), such as tyrosine kinase inhibitors (TKIs), which include but are not limited to small molecule compounds, biologics (such as monoclonal antibodies), and large polypeptide molecules that inhibit the activity of, for example, IFN signaling and production pathways, or any other form of antagonists that can decrease expression of a target protein in the immune response pathway. It is to be understood that the present disclosure contemplates use of any modality of therapeutics that can act as an antagonist of, *e.g.*, the IFN signaling and production pathways that modulate immune responses.

[00400] The immunosuppressants are protein kinase inhibitors belonging to a wide class of compounds that inhibits the activity of protein kinases and can be used in conjunction with any nucleic acid therapeutic that triggers immune responses (innate and/or adaptive) in a host cell or a subject suffering from a genetic disorder. Tyrosine kinases regulate a variety of cellular functions including cell growth (*e.g.*, IFN signaling and production and epidermal growth factor (“EGFR” such as ERBB1, ERBB2/HER2, ERBB3/HER3, ERBB4/HER4)). These are the main signal transducers and activators which act downstream of multiple cytokines, growth factors, and hormones, thereby regulating immune responses. For example, upon binding of a specific ligand to its cognate receptor, conformational changes lead to receptor oligomerization and activation of the receptor-associated JAKs. JAKs auto- and trans-phosphorylate one another and phosphorylate receptor chains, providing the docking sites for STAT molecules. STATs then undergo JAK-mediated phosphorylation, dimerize, and translocate to the nucleus, where they regulate the transcription of target genes involving immune responses (*e.g.*, interferon- α , interferon- β , interferon- γ , TNF α , IL2, IL-6, IL-18, etc.).

[00401] In some embodiments, the immunosuppressant is an antagonist of Jak1, Jak2, Jak3, Stat, Tyk2, c-MET, EGFR, c-KIT, BTK, ALK, ABL, SRC, ROS1, Syk, MEK, ATM, NRF2, Flt3, fms/CSF1R, FDGFRs, RON, IGF1R, EPHA2, EPHA3, VEGF or VEGFR. In some embodiment, the

immunosuppressant is an antagonist of tyrosine kinase. In one embodiment, the immunosuppressant is an antagonist of Jak1. In another embodiment, the immunosuppressant is an antagonist of Jak2. In one embodiment, the immunosuppressant is an antagonist of Jak3. In yet another embodiment, the immunosuppressant is an antagonist of Tyk2. In yet another embodiment, the immunosuppressant is an antagonist of EGFR. In one embodiment, the immunosuppressant is an antagonist of ALK. In yet another embodiment, the immunosuppressant is an antagonist of Syk.

[00402] In one embodiment, the immunosuppressant is a small molecule antagonist. In another embodiment, the immunosuppressant is an antibody that binds to a protein kinase target. In another embodiment, the immunosuppressant is an antibody that binds to a tyrosine kinase. In another embodiment, the immunosuppressant is a monoclonal antibody against a protein kinase. In another embodiment, the immunosuppressant is a monoclonal antibody against tyrosine kinase. In another embodiment, the immunosuppressant is a monoclonal antibody against a target selected from the group consisting of Jak1, Jak2, Jak3, Stat, Tyk2, c-MET, EGFR, c-KIT, BTK, ALK, ABL, SRC, ROS1, Syk, MEK, ATM, NRF2, Flt3, fms/CSF1R, FDGFRs, RON, IGF1R, EPHA2, EPHA3, VEGF and VEGFR. In yet another embodiment, the immunosuppressant is a polypeptide that has binding affinity to a protein kinase. In yet another embodiment, the immunosuppressant is a nucleic acid, such as RNAi or an anti-sense oligonucleotide, that attenuates expression of Jak1, Jak2, Jak3, Stat, Tyk2, c-MET, EGFR, c-KIT, BTK, ALK, ABL, SRC, ROS1, Syk, MEK, ATM, NRF2, Flt3, fms/CSF1R, FDGFRs, RON, IGF1R, EPHA2, EPHA3, VEGF or VEGFR.

[00403] In some embodiments, inhibition of a protein kinase, *e.g.*, a tyrosine kinase, can be achieved by using small molecules that bind to the ATP pocket of a given protein kinase, blocking it from catalyzing the phosphorylation of target proteins. Hence, in some embodiments, the immunosuppressant can be a small molecule antagonist of protein kinase. Non-limiting examples of immunosuppressant protein kinase antagonist include imatinib mesylate (GLEEVEC®), Nilotinib (TASIGNA®), sorafenib (NEXAVAR®), sunitinib (SUTNET®), dasatinib (SPRCEL®), acalabrutinib, alectinib, axitinib, baricitinib, afatinib, bosutinib, brigatinib, cabozantinib, cerdulatinib, ceritinib, cobimetinib, crizotinib, dacomitinib, dasatinib, erlotinib, imatinib, fostamatinib, gefitinib, AG-1478, lapatinib, lorlatinib, TAK-659, ruxolitinib, osimertinib, pazopanib, pegaptanib, ponatinib, regorafenib, saracatinib, tofacitinib, BMS- 986165, vandetinib, vemurafenib, or a pharmaceutically acceptable salt thereof.

[00404] In some embodiments, the immunosuppressant can be a small molecule antagonist of tyrosine kinase and is selected from the group consisting of baricitinib, afatinib, brigatinib, cerdulatinib, ceritinib, cobimetinib, dacomitinib, dasatinib, osimertinib, fostamatinib, saracatinib, TAK-659, ruxolitinib, BMS-986165, tofacitinib, and a pharmaceutically acceptable salt thereof.

[00405] In some embodiments, said TKI is selected from the group consisting of sunitinib, imatinib, sorafenib, dasatinib, entoplectinib, fostamatinib, TAK-659, ruxolitinib, baricitinib, BMS-986165, tofacitinib, and a pharmaceutically acceptable salt thereof.

[00406] In some embodiments, the TKI is selected from the group consisting of fostamatinib, ruxolitinib, BMS-986165, and a pharmaceutically acceptable salt thereof.

[00407] In one embodiment, the TKI is ruxolitinib or ruxolitinib phosphate.

[00408] In some embodiments, the TKI may selectively inhibit one or multiple kinases; or target multiple kinases in the same pathway. For example, ruxolitinib and baricitinib can inhibit Jak1 and Jak2. Lorlatinib can inhibit ROS1 and ALK. Dasatinib can inhibit Alb, Src and c-Kit. Brigatinib, genfinitinib, erlotinib, AG-1478 and lapatinib can inhibit EGFR. Crizitinib can inhibit both ALK and c-Met. Fostamatinib and cerdulitinib can selectively inhibit Syk. Saracatinib can inhibit Src and Abl. In some embodiments, the TKI is an inhibitor of Jak1. In some embodiments, the TKI is an inhibitor of Jak2. In some embodiments, the TKI is an inhibitor of Jak1 and Jak2. In some embodiments, the TKI is an inhibitor of EGFR. In some embodiments, the TKI is an inhibitor of ALK. In some embodiments, the TKI is an inhibitor of Syk.

[00409] Protein kinase activity in immune response pathways may also be inhibited by biologic drugs, such as a monoclonal antibody against a protein kinase. These therapeutics may exert efficacy by preventing receptor protein kinases from activating and are capable of binding cell surface antigens with high specificity. Several monoclonal antibodies target receptor protein kinases that play a role in inhibiting protein kinases involving in DNA sensing immune response signaling pathways.

Trastuzumab and bevacizumab are nonlimiting examples of such monoclonal antibodies.

[00410] In some embodiments, the biologic agent that functions to suppress unwanted immune response to a TNA is a monoclonal antibody selected from the group consisting of ado-trastuzumab emtansine, cetuximab, CetuGEX™, cixutumumab, dalotuzumab, duligotumab, ertumaxomab, futuximab, ganitumab, icrucumab, margetuximab, narnatumab, necitumumab, nimotuzumab (h-R3), olaratumab, onartuzumab, panitumumab, pertuzumab, ranibizumab, ramucirumab, seribantumab, tanibirumab, teprotumumab, trasGEX™, trastuzumab, and zatuximab. In one embodiment, the monoclonal antibody is trastuzumab. In another embodiment, the monoclonal antibody is cetuximab.

[00411] In some embodiments, the protein kinase inhibitor is a peptide. The peptide can be polypeptide having a specific affinity binding to target proteins such as Jak1, Jak2, Jak3, STAT, Tyk2, c-MET, EGFR, c-KIT, BTK, ALK, ABL, SRC, ROS1, Syk, MEK, ATM, NRF2, Flt3, fms/CSF1R, FDGFRs, RON, IGF1r, EPHA2, EPHA3, VEGF or VEGFR. Non-limiting examples of a polypeptide immunosuppressant include aflibercept (VEGF). Binding targets for the peptide can be in a signaling pathway involved in, *e.g.*, IFN response and production pathways.

[00412] In some embodiments, the immunosuppressants of the present disclosure effectively reduce *in vitro* and *in vivo* pro-inflammatory cytokine and chemokine levels when the immunosuppressants are present in combination with the nucleic acid therapeutics. The pro-

inflammatory cytokine can be selected from any or a combination of interferon- α (IFN- α), interferon- γ (IFN- γ), tumor necrosis factor- α (TNF- α), interleukin-1 β (IL-1 β), interleukin-6 (IL-6), interleukin-8 (IL-8), interleukin-10 (IL-10), interleukin-12 (IL-12), interleukin-18 (IL-18), vascular endothelial growth factor (VEGF), leukemia inhibitory factor (LIF), matrixmetalloproteinase 2 (MMP2), monocyte chemoattractant protein-1 (MCP-1), RANTES (CCL5), IP-10 (CXCL10), macrophage inflammatory protein-1 α (MIP-1 α ; CCL3) and/or macrophage inflammatory protein-1 β (MIP-1 β ; CCL4).

[00413] In some embodiments, the additional compound is immune stimulatory agent. Also provided herein is a pharmaceutical composition comprising the lipid nanoparticle-encapsulated insect-cell produced, or a synthetically produced ceDNA vector for expression of FIX protein as described herein and a pharmaceutically acceptable carrier or excipient.

[00414] In some aspects, the disclosure provides for a lipid nanoparticle formulation further comprising one or more pharmaceutical excipients. In some embodiments, the lipid nanoparticle formulation further comprises sucrose, tris, trehalose and/or glycine.

[00415] The ceDNA vector can be complexed with the lipid portion of the particle or encapsulated in the lipid position of the lipid nanoparticle. In some embodiments, the ceDNA can be fully encapsulated in the lipid position of the lipid nanoparticle, thereby protecting it from degradation by a nuclease, *e.g.*, in an aqueous solution. In some embodiments, the ceDNA in the lipid nanoparticle is not substantially degraded after exposure of the lipid nanoparticle to a nuclease at 37°C. for at least about 20, 30, 45, or 60 minutes. In some embodiments, the ceDNA in the lipid nanoparticle is not substantially degraded after incubation of the particle in serum at 37°C. for at least about 30, 45, or 60 minutes or at least about 2, 3, 4, 5, 6, 7, 8, 9, 10, 12, 14, 16, 18, 20, 22, 24, 26, 28, 30, 32, 34, or 36 hours.

[00416] In certain embodiments, the lipid nanoparticles are substantially non-toxic to a subject, *e.g.*, to a mammal such as a human. In some aspects, the lipid nanoparticle formulation is a lyophilized powder.

[00417] In some embodiments, lipid nanoparticles are solid core particles that possess at least one lipid bilayer. In other embodiments, the lipid nanoparticles have a non-bilayer structure, *i.e.*, a non-lamellar (*i.e.*, non-bilayer) morphology. Without limitations, the non-bilayer morphology can include, for example, three dimensional tubes, rods, cubic symmetries, etc. For example, the morphology of the lipid nanoparticles (lamellar vs. non-lamellar) can readily be assessed and characterized using, *e.g.*, Cryo-TEM analysis as described in US2010/0130588, the content of which is incorporated herein by reference in its entirety.

[00418] In some further embodiments, the lipid nanoparticles having a non-lamellar morphology are electron dense. In some aspects, the disclosure provides for a lipid nanoparticle that is either unilamellar or multilamellar in structure. In some aspects, the disclosure provides for a lipid nanoparticle formulation that comprises multi-vesicular particles and/or foam-based particles.

[00419] By controlling the composition and concentration of the lipid components, one can control the rate at which the lipid conjugate exchanges out of the lipid particle and, in turn, the rate at which the lipid nanoparticle becomes fusogenic. In addition, other variables including, *e.g.*, pH, temperature, or ionic strength, can be used to vary and/or control the rate at which the lipid nanoparticle becomes fusogenic. Other methods which can be used to control the rate at which the lipid nanoparticle becomes fusogenic will be apparent to those of ordinary skill in the art based on this disclosure. It will also be apparent that by controlling the composition and concentration of the lipid conjugate, one can control the lipid particle size.

[00420] The pKa of formulated cationic lipids can be correlated with the effectiveness of the LNPs for delivery of nucleic acids (see Jayaraman et al, *Angewandte Chemie, International Edition* (2012), 51(34), 8529-8533; Semple et al, *Nature Biotechnology* 28, 172-176 (2010), both of which are incorporated by reference in their entirety). The preferred range of pKa is ~5 to ~7. The pKa of the cationic lipid can be determined in lipid nanoparticles using an assay based on fluorescence of 2-(p-toluidino)-6-naphthalene sulfonic acid (TNS).

VIII. Methods of Use

[00421] A ceDNA vector for expression of FIX protein as disclosed herein can also be used in a method for the delivery of a nucleic acid sequence of interest (*e.g.*, encoding FIX protein) to a target cell (*e.g.*, a host cell). The method may in particular be a method for delivering FIX protein to a cell of a subject in need thereof and treating hemophilia B. The disclosure allows for the *in vivo* expression of FIX protein encoded in the ceDNA vector in a cell in a subject such that therapeutic effect of the expression of FIX protein occurs. These results are seen with both *in vivo* and *in vitro* modes of ceDNA vector delivery.

[00422] In addition, the disclosure provides a method for the delivery of FIX protein in a cell of a subject in need thereof, comprising multiple administrations of the ceDNA vector of the disclosure encoding said FIX protein. Since the ceDNA vector of the disclosure does not induce an immune response like that typically observed against encapsidated viral vectors, such a multiple administration strategy will likely have greater success in a ceDNA-based system. The ceDNA vector are administered in sufficient amounts to transfect the cells of a desired tissue and to provide sufficient levels of gene transfer and expression of the FIX protein without undue adverse effects. Conventional and pharmaceutically acceptable routes of administration include, but are not limited to, retinal administration (*e.g.*, subretinal injection, suprachoroidal injection or intravitreal injection), intravenous (*e.g.*, in a liposome formulation), direct delivery to the selected organ (*e.g.*, any one or more tissues selected from: liver, kidneys, gallbladder, prostate, adrenal gland, heart, intestine, lung, and stomach), intramuscular, and other parental routes of administration. Routes of administration may be combined, if desired.

[00423] Delivery of a ceDNA vector for expression of FIX protein as described herein is not limited to delivery of the expressed FIX protein. For example, conventionally produced (*e.g.*, using a cell-

based production method (*e.g.*, insect-cell production methods) or synthetically produced ceDNA vectors as described herein may be used with other delivery systems provided to provide a portion of the gene therapy. One non-limiting example of a system that may be combined with the ceDNA vectors in accordance with the present disclosure includes systems which separately deliver one or more co-factors or immune suppressors for effective gene expression of the ceDNA vector expressing the FIX protein.

[00424] The disclosure also provides for a method of treating hemophilia B in a subject comprising introducing into a target cell in need thereof (in particular a muscle cell or tissue) of the subject a therapeutically effective amount of a ceDNA vector, optionally with a pharmaceutically acceptable carrier. While the ceDNA vector can be introduced in the presence of a carrier, such a carrier is not required. The ceDNA vector selected comprises a nucleic acid sequence encoding an FIX protein useful for treating hemophilia B. In particular, the ceDNA vector may comprise a desired FIX protein sequence operably linked to control elements capable of directing transcription of the desired FIX protein encoded by the exogenous DNA sequence when introduced into the subject. The ceDNA vector can be administered via any suitable route as provided above, and elsewhere herein.

[00425] The compositions and vectors provided herein can be used to deliver an FIX protein for various purposes. In some embodiments, the transgene encodes an FIX protein that is intended to be used for research purposes, *e.g.*, to create a somatic transgenic animal model harboring the transgene, *e.g.*, to study the function of the FIX protein product. In another example, the transgene encodes an FIX protein that is intended to be used to create an animal model of hemophilia B. In some embodiments, the encoded FIX protein is useful for the treatment or prevention of hemophilia B states in a mammalian subject. The FIX protein can be transferred (*e.g.*, expressed in) to a patient in a sufficient amount to treat hemophilia B associated with reduced expression, lack of expression or dysfunction of the gene.

[00426] In principle, the expression cassette can include a nucleic acid or any transgene that encodes an FIX protein that is either reduced or absent due to a mutation or which conveys a therapeutic benefit when overexpressed is considered to be within the scope of the disclosure. Preferably, noninserted bacterial DNA is not present and preferably no bacterial DNA is present in the ceDNA compositions provided herein.

[00427] A ceDNA vector is not limited to one species of ceDNA vector. As such, in another aspect, multiple ceDNA vectors expressing different proteins or the same FIX protein but operatively linked to different promoters or cis-regulatory elements can be delivered simultaneously or sequentially to the target cell, tissue, organ, or subject. Therefore, this strategy can allow for the gene therapy or gene delivery of multiple proteins simultaneously. It is also possible to separate different portions of a FIX protein into separate ceDNA vectors (*e.g.*, different domains and/or co-factors required for functionality of a FIX protein) which can be administered simultaneously or at different times, and can be separately regulatable, thereby adding an additional level of control of expression of a FIX protein.

Delivery can also be performed multiple times and, importantly for gene therapy in the clinical setting, in subsequent increasing or decreasing doses, given the lack of an anti-capsid host immune response due to the absence of a viral capsid. It is anticipated that no anti-capsid response will occur as there is no capsid.

[00428] The disclosure also provides for a method of treating hemophilia B in a subject comprising introducing into a target cell in need thereof (in particular a muscle cell or tissue) of the subject a therapeutically effective amount of a ceDNA vector as disclosed herein, optionally with a pharmaceutically acceptable carrier. While the ceDNA vector can be introduced in the presence of a carrier, such a carrier is not required. The ceDNA vector implemented comprises a nucleic acid sequence of interest useful for treating the hemophilia B. In particular, the ceDNA vector may comprise a desired exogenous DNA sequence operably linked to control elements capable of directing transcription of the desired polypeptide, protein, or oligonucleotide encoded by the exogenous DNA sequence when introduced into the subject. The ceDNA vector can be administered via any suitable route as provided above, and elsewhere herein.

IX. Methods of delivering ceDNA vectors for FIX protein production

[00429] In some embodiments, a ceDNA vector for expression of FIX protein can be delivered to a target cell *in vitro* or *in vivo* by various suitable methods. ceDNA vectors alone can be applied or injected. CeDNA vectors can be delivered to a cell without the help of a transfection reagent or other physical means. Alternatively, ceDNA vectors for expression of FIX protein can be delivered using any art-known transfection reagent or other art-known physical means that facilitates entry of DNA into a cell, *e.g.*, liposomes, alcohols, polylysine- rich compounds, arginine-rich compounds, calcium phosphate, microvesicles, microinjection, electroporation and the like.

[00430] The ceDNA vectors for expression of FIX protein as disclosed herein can efficiently target cell and tissue-types that are normally difficult to transduce with conventional AAV virions using various delivery reagent.

[00431] One aspect of the technology described herein relates to a method of delivering an FIX protein to a cell. Typically, for *in vivo* and *in vitro* methods, a ceDNA vector for expression of FIX protein as disclosed herein may be introduced into the cell using the methods as disclosed herein, as well as other methods known in the art. A ceDNA vector for expression of FIX protein as disclosed herein are preferably administered to the cell in a biologically-effective amount. If the ceDNA vector is administered to a cell *in vivo* (*e.g.*, to a subject), a biologically-effective amount of the ceDNA vector is an amount that is sufficient to result in transduction and expression of the FIX protein in a target cell.

[00432] Exemplary modes of administration of a ceDNA vector for expression of FIX protein as disclosed herein includes oral, rectal, transmucosal, intranasal, inhalation (*e.g.*, via an aerosol), buccal (*e.g.*, sublingual), vaginal, intrathecal, intraocular, transdermal, intraendothelial, in utero (or in ovo), parenteral (*e.g.*, intravenous, subcutaneous, intradermal, intracranial, intramuscular [including

administration to skeletal, diaphragm and/or cardiac muscle], intrapleural, intracerebral, and intraarticular). Administration can be systemically or direct delivery to the liver or elsewhere (*e.g.*, any kidneys, gallbladder, prostate, adrenal gland, heart, intestine, lung, and stomach).

[00433] Administration can be topical (*e.g.*, to both skin and mucosal surfaces, including airway surfaces, and transdermal administration), intralymphatic, and the like, as well as direct tissue or organ injection (*e.g.*, but not limited to, liver, but also to eye, muscles, including skeletal muscle, cardiac muscle, diaphragm muscle, or brain).

[00434] Administration of the ceDNA vector can be to any site in a subject, including, without limitation, a site selected from the group consisting of the liver and/or also eyes, brain, a skeletal muscle, a smooth muscle, the heart, the diaphragm, the airway epithelium, the kidney, the spleen, the pancreas, the skin.

[00435] The most suitable route in any given case will depend on the nature and severity of the condition being treated, ameliorated, and/or prevented and on the nature of the particular ceDNA vector that is being used. Additionally, ceDNA permits one to administer more than one FIX protein in a single vector, or multiple ceDNA vectors (*e.g.* a ceDNA cocktail).

A. Intramuscular Administration of a ceDNA vector

[00436] In some embodiments, a method of treating a disease in a subject comprises introducing into a target cell in need thereof (in particular a muscle cell or tissue) of the subject a therapeutically effective amount of a ceDNA vector encoding an FIX protein, optionally with a pharmaceutically acceptable carrier. In some embodiments, the ceDNA vector for expression of FIX protein is administered to a muscle tissue of a subject.

[00437] In some embodiments, administration of the ceDNA vector can be to any site in a subject, including, without limitation, a site selected from the group consisting of a skeletal muscle, a smooth muscle, the heart, the diaphragm, or muscles of the eye.

[00438] Administration of a ceDNA vector for expression of FIX protein as disclosed herein to a skeletal muscle according to the present disclosure includes but is not limited to administration to the skeletal muscle in the limbs (*e.g.*, upper arm, lower arm, upper leg, and/or lower leg), back, neck, head (*e.g.*, tongue), thorax, abdomen, pelvis/perineum, and/or digits. The ceDNA as disclosed herein vector can be delivered to skeletal muscle by intravenous administration, intra-arterial administration, intraperitoneal administration, limb perfusion, (optionally, isolated limb perfusion of a leg and/or arm; see, *e.g.* Arruda et al., (2005) Blood 105: 3458-3464), and/or direct intramuscular injection. In particular embodiments, the ceDNA vector as disclosed herein is administered to the liver, eye, a limb (arm and/or leg) of a subject (*e.g.*, a subject with muscular dystrophy such as DMD) by limb perfusion, optionally isolated limb perfusion (*e.g.*, by intravenous or intra-articular administration. In embodiments, the ceDNA vector as disclosed herein can be administered without employing "hydrodynamic" techniques.

[00439] For instance, tissue delivery (*e.g.*, to retina) of conventional viral vectors is often enhanced by hydrodynamic techniques (*e.g.*, intravenous/intravenous administration in a large volume), which increase pressure in the vasculature and facilitate the ability of the viral vector to cross the endothelial cell barrier. In particular embodiments, the ceDNA vectors described herein can be administered in the absence of hydrodynamic techniques such as high volume infusions and/or elevated intravascular pressure (*e.g.*, greater than normal systolic pressure, for example, less than or equal to a 5%, 10%, 15%, 20%, 25% increase in intravascular pressure over normal systolic pressure). Such methods may reduce or avoid the side effects associated with hydrodynamic techniques such as edema, nerve damage and/or compartment syndrome.

[00440] Furthermore, a composition comprising a ceDNA vector for expression of FIX protein as disclosed herein that is administered to a skeletal muscle can be administered to a skeletal muscle in the limbs (*e.g.*, upper arm, lower arm, upper leg, and/or lower leg), back, neck, head (*e.g.*, tongue), thorax, abdomen, pelvis/perineum, and/or digits. Suitable skeletal muscles include but are not limited to abductor digiti minimi (in the hand), abductor digiti minimi (in the foot), abductor hallucis, abductor ossis metatarsi quinti, abductor pollicis brevis, abductor pollicis longus, adductor brevis, adductor hallucis, adductor longus, adductor magnus, adductor pollicis, anconeus, anterior scalene, articularis genus, biceps brachii, biceps femoris, brachialis, brachioradialis, buccinator, coracobrachialis, corrugator supercilii, deltoid, depressor anguli oris, depressor labii inferioris, digastric, dorsal interossei (in the hand), dorsal interossei (in the foot), extensor carpi radialis brevis, extensor carpi radialis longus, extensor carpi ulnaris, extensor digiti minimi, extensor digitorum, extensor digitorum brevis, extensor digitorum longus, extensor hallucis brevis, extensor hallucis longus, extensor indicis, extensor pollicis brevis, extensor pollicis longus, flexor carpi radialis, flexor carpi ulnaris, flexor digiti minimi brevis (in the hand), flexor digiti minimi brevis (in the foot), flexor digitorum brevis, flexor digitorum longus, flexor digitorum profundus, flexor digitorum superficialis, flexor hallucis brevis, flexor hallucis longus, flexor pollicis brevis, flexor pollicis longus, frontalis, gastrocnemius, geniohyoid, gluteus maximus, gluteus medius, gluteus minimus, gracilis, iliocostalis cervicis, iliocostalis lumborum, iliocostalis thoracis, iliopsoas, inferior gemellus, inferior oblique, inferior rectus, infraspinatus, interspinalis, intertransversi, lateral pterygoid, lateral rectus, latissimus dorsi, levator anguli oris, levator labii superioris, levator labii superioris alaeque nasi, levator palpebrae superioris, levator scapulae, long rotators, longissimus capitis, longissimus cervicis, longissimus thoracis, longus capitis, longus colli, lumbricals (in the hand), lumbricals (in the foot), masseter, medial pterygoid, medial rectus, middle scalene, multifidus, mylohyoid, obliquus capitis inferior, obliquus capitis superior, obturator externus, obturator internus, occipitalis, omohyoid, opponens digiti minimi, opponens pollicis, orbicularis oculi, orbicularis oris, palmar interossei, palmaris brevis, palmaris longus, pectineus, pectoralis major, pectoralis minor, peroneus brevis, peroneus longus, peroneus tertius, piriformis, plantar interossei, plantaris, platysma, popliteus, posterior scalene, pronator quadratus, pronator teres, psoas major, quadratus femoris, quadratus plantae, rectus capitis anterior,

rectus capitis lateralis, rectus capitis posterior major, rectus capitis posterior minor, rectus femoris, rhomboid major, rhomboid minor, risorius, sartorius, scalenus minimus, semimembranosus, semispinalis capitis, semispinalis cervicis, semispinalis thoracis, semitendinosus, serratus anterior, short rotators, soleus, spinalis capitis, spinalis cervicis, spinalis thoracis, splenius capitis, splenius cervicis, sternocleidomastoid, sternohyoid, sternothyroid, stylohyoid, subclavius, subscapularis, superior gemellus, superior oblique, superior rectus, supinator, supraspinatus, temporalis, tensor fascia lata, teres major, teres minor, thoracis, thyrohyoid, tibialis anterior, tibialis posterior, trapezius, triceps brachii, vastus intermedius, vastus lateralis, vastus medialis, zygomaticus major, and zygomaticus minor, and any other suitable skeletal muscle as known in the art.

[00441] Administration of a ceDNA vector for expression of FIX protein as disclosed herein to diaphragm muscle can be by any suitable method including intravenous administration, intra-arterial administration, and/or intra-peritoneal administration. In some embodiments, delivery of an expressed transgene from the ceDNA vector to a target tissue can also be achieved by delivering a synthetic depot comprising the ceDNA vector, where a depot comprising the ceDNA vector is implanted into skeletal, smooth, cardiac and/or diaphragm muscle tissue or the muscle tissue can be contacted with a film or other matrix comprising the ceDNA vector as described herein. Such implantable matrices or substrates are described in U.S. Pat. No. 7,201,898.

[00442] Administration of a ceDNA vector for expression of FIX protein as disclosed herein to cardiac muscle includes administration to the left atrium, right atrium, left ventricle, right ventricle and/or septum. The ceDNA vector as described herein can be delivered to cardiac muscle by intravenous administration, intra-arterial administration such as intra-aortic administration, direct cardiac injection (*e.g.*, into left atrium, right atrium, left ventricle, right ventricle), and/or coronary artery perfusion.

[00443] Administration of a ceDNA vector for expression of FIX protein as disclosed herein to smooth muscle can be by any suitable method including intravenous administration, intra-arterial administration, and/or intra-peritoneal administration. In one embodiment, administration can be to endothelial cells present in, near, and/or on smooth muscle. Non-limiting examples of smooth muscles include the iris of the eye, bronchioles of the lung, laryngeal muscles (vocal cords), muscular layers of the stomach, esophagus, small and large intestine of the gastrointestinal tract, ureter, detrusor muscle of the urinary bladder, uterine myometrium, penis, or prostate gland.

[00444] In some embodiments, of a ceDNA vector for expression of FIX protein as disclosed herein is administered to skeletal muscle, diaphragm muscle and/or cardiac muscle. In representative embodiments, a ceDNA vector according to the present disclosure is used to treat and/or prevent disorders of skeletal, cardiac and/or diaphragm muscle.

[00445] Specifically, it is contemplated that a composition comprising a ceDNA vector for expression of FIX protein as disclosed herein can be delivered to one or more muscles of the eye (*e.g.*, Lateral rectus, Medial rectus, Superior rectus, Inferior rectus, Superior oblique, Inferior oblique), facial

muscles (*e.g.*, Occipitofrontalis muscle, Temporoparietalis muscle, Procerus muscle, Nasalis muscle, Depressor septi nasi muscle, Orbicularis oculi muscle, Corrugator supercilii muscle, Depressor supercilii muscle, Auricular muscles, Orbicularis oris muscle, Depressor anguli oris muscle, Risorius, Zygomaticus major muscle, Zygomaticus minor muscle, Levator labii superioris, Levator labii superioris alaeque nasi muscle, Depressor labii inferioris muscle, Levator anguli oris, Buccinator muscle, Mentalis) or tongue muscles (*e.g.*, genioglossus, hyoglossus, chondroglossus, styloglossus, palatoglossus, superior longitudinal muscle, inferior longitudinal muscle, the vertical muscle, and the transverse muscle).

[00446](i) Intramuscular injection: In some embodiments, a composition comprising a ceDNA vector for expression of FIX protein as disclosed herein can be injected into one or more sites of a given muscle, for example, skeletal muscle (*e.g.*, deltoid, vastus lateralis, ventrogluteal muscle of dorsogluteal muscle, or anterolateral thigh for infants) in a subject using a needle. The composition comprising ceDNA can be introduced to other subtypes of muscle cells. Non-limiting examples of muscle cell subtypes include skeletal muscle cells, cardiac muscle cells, smooth muscle cells and/or diaphragm muscle cells.

[00447] Methods for intramuscular injection are known to those of skill in the art and as such are not described in detail herein. However, when performing an intramuscular injection, an appropriate needle size should be determined based on the age and size of the patient, the viscosity of the composition, as well as the site of injection. **Table 13** provides guidelines for exemplary sites of injection and corresponding needle size:

Table 13: Guidelines for intramuscular injection in human patients

<i>Injection Site</i>	<i>Needle Gauge</i>	<i>Needle Size</i>	<i>Maximum volume of composition</i>
Ventrogluteal site (gluteus medius and gluteus minimus)	<i>Aqueous solutions:</i> 20-25 gauge <i>Viscous or oil-based solution:</i> 18-21 gauge	Thin adult: 15 to 25 mm Average adult: 25 mm Larger adult (over 150 lbs): 25 to 38 mm. Children and infants: will require a smaller needle	3mL
Vastus lateralis	<i>Aqueous solutions:</i> 20-25 gauge <i>Viscous or oil-based solution:</i> 18-21 gauge <u>Children/infants:</u> 22 to 25 gauge	Adult: 25 mm to 38 mm	3mL
Deltoid	22 to 25 gauge	Males:	1mL

		130-260lbs: 25 mm Females: <130 lbs: 16 mm 130-200 lbs: 25mm >200lbs: 38mm	

[00448] In certain embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is formulated in a small volume, for example, an exemplary volume as outlined in **Table 13** for a given subject. In some embodiments, the subject can be administered a general or local anesthetic prior to the injection, if desired. This is particularly desirable if multiple injections are required or if a deeper muscle is injected, rather than the common injection sites noted above.

[00449] In some embodiments, intramuscular injection can be combined with electroporation, delivery pressure or the use of transfection reagents to enhance cellular uptake of the ceDNA vector.

[00450] **(ii) Transfection Reagents:** In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is formulated in compositions comprising one or more transfection reagents to facilitate uptake of the vectors into myotubes or muscle tissue. Thus, in one embodiment, the nucleic acids described herein are administered to a muscle cell, myotube or muscle tissue by transfection using methods described elsewhere herein.

[00451] **(iii) Electroporation:** In certain embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is administered in the absence of a carrier to facilitate entry of ceDNA into the cells, or in a physiologically inert pharmaceutically acceptable carrier (*i.e.*, any carrier that does not improve or enhance uptake of the capsid free, non-viral vectors into the myotubes). In such embodiments, the uptake of the capsid free, non-viral vector can be facilitated by electroporation of the cell or tissue.

[00452] Cell membranes naturally resist the passage of extracellular into the cell cytoplasm. One method for temporarily reducing this resistance is “electroporation”, where electrical fields are used to create pores in cells without causing permanent damage to the cells. These pores are large enough to allow DNA vectors, pharmaceutical drugs, DNA, and other polar compounds to gain access to the interior of the cell. With time, the pores in the cell membrane close and the cell once again becomes impermeable.

[00453] Electroporation can be used in both *in vitro* and *in vivo* applications to introduce *e.g.*, exogenous DNA into living cells. *In vitro* applications typically mix a sample of live cells with the composition comprising *e.g.*, DNA. The cells are then placed between electrodes such as parallel plates and an electrical field is applied to the cell/composition mixture.

[00454] There are a number of methods for *in vivo* electroporation; electrodes can be provided in various configurations such as, for example, a caliper that grips the epidermis overlying a region of cells to be treated. Alternatively, needle-shaped electrodes may be inserted into the tissue, to access more deeply located cells. In either case, after the composition comprising *e.g.*, nucleic acids are

injected into the treatment region, the electrodes apply an electrical field to the region. In some electroporation applications, this electric field comprises a single square wave pulse on the order of 100 to 500 V/cm. of about 10 to 60 ms duration. Such a pulse may be generated, for example, in known applications of the Electro Square Porator T820, made by the BTX Division of Genetronics, Inc.

[00455] Typically, successful uptake of *e.g.*, nucleic acids occurs only if the muscle is electrically stimulated immediately, or shortly after administration of the composition, for example, by injection into the muscle.

[00456] In certain embodiments, electroporation is achieved using pulses of electric fields or using low voltage/long pulse treatment regimens (*e.g.*, using a square wave pulse electroporation system). Exemplary pulse generators capable of generating a pulsed electric field include, for example, the ECM600, which can generate an exponential wave form, and the ElectroSquarePorator (T820), which can generate a square wave form, both of which are available from BTX, a division of Genetronics, Inc. (San Diego, Calif.). Square wave electroporation systems deliver controlled electric pulses that rise quickly to a set voltage, stay at that level for a set length of time (pulse length), and then quickly drop to zero.

[00457] In some embodiments, a local anesthetic is administered, for example, by injection at the site of treatment to reduce pain that may be associated with electroporation of the tissue in the presence of a composition comprising a capsid free, non-viral vector as described herein. In addition, one of skill in the art will appreciate that a dose of the composition should be chosen that minimizes and/or prevents excessive tissue damage resulting in fibrosis, necrosis or inflammation of the muscle.

[00458](iv) *Delivery Pressure:* In some embodiments, delivery of a ceDNA vector for expression of FIX protein as disclosed herein to muscle tissue is facilitated by delivery pressure, which uses a combination of large volumes and rapid injection into an artery supplying a limb (*e.g.*, iliac artery). This mode of administration can be achieved through a variety of methods that involve infusing limb vasculature with a composition comprising a ceDNA vector, typically while the muscle is isolated from the systemic circulation using a tourniquet or vessel clamps. In one method, the composition is circulated through the limb vasculature to permit extravasation into the cells. In another method, the intravascular hydrodynamic pressure is increased to expand vascular beds and increase uptake of the ceDNA vector into the muscle cells or tissue. In one embodiment, the ceDNA composition is administered into an artery.

[00459](v) *Lipid Nanoparticle Compositions:* In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein for intramuscular delivery are formulated in a composition comprising a liposome as described elsewhere herein.

[00460](vi) *Systemic Administration of a ceDNA Vector targeted to Muscle Tissue:* In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is formulated to be targeted to the muscle via indirect delivery administration, where the ceDNA is transported to the

muscle as opposed to the liver. Accordingly, the technology described herein encompasses indirect administration of compositions comprising a ceDNA vector for expression of FIX protein as disclosed herein to muscle tissue, for example, by systemic administration. Such compositions can be administered topically, intravenously (by bolus or continuous infusion), intracellular injection, intratissue injection, orally, by inhalation, intraperitoneally, subcutaneously, intracavity, and can be delivered by peristaltic means, if desired, or by other means known by those skilled in the art. The agent can be administered systemically, for example, by intravenous infusion, if so desired.

[00461] In some embodiments, uptake of a ceDNA vector for expression of FIX protein as disclosed herein into muscle cells/tissue is increased by using a targeting agent or moiety that preferentially directs the vector to muscle tissue. Thus, in some embodiments, a capsid free, ceDNA vector can be concentrated in muscle tissue as compared to the amount of capsid free ceDNA vectors present in other cells or tissues of the body.

[00462] In some embodiments, the composition comprising a ceDNA vector for expression of FIX protein as disclosed herein further comprises a targeting moiety to muscle cells. In other embodiments, the expressed gene product comprises a targeting moiety specific to the tissue in which it is desired to act. The targeting moiety can include any molecule, or complex of molecules, which is/are capable of targeting, interacting with, coupling with, and/or binding to an intracellular, cell surface, or extracellular biomarker of a cell or tissue. The biomarker can include, for example, a cellular protease, a kinase, a protein, a cell surface receptor, a lipid, and/or fatty acid. Other examples of biomarkers that the targeting moieties can target, interact with, couple with, and/or bind to include molecules associated with a particular disease. For example, the biomarkers can include cell surface receptors implicated in cancer development, such as epidermal growth factor receptor and transferrin receptor. The targeting moieties can include, but are not limited to, synthetic compounds, natural compounds or products, macromolecular entities, bioengineered molecules (*e.g.*, polypeptides, lipids, polynucleotides, antibodies, antibody fragments), and small entities (*e.g.*, small molecules, neurotransmitters, substrates, ligands, hormones and elemental compounds) that bind to molecules expressed in the target muscle tissue.

[00463] In certain embodiments, the targeting moiety may further comprise a receptor molecule, including, for example, receptors, which naturally recognize a specific desired molecule of a target cell. Such receptor molecules include receptors that have been modified to increase their specificity of interaction with a target molecule, receptors that have been modified to interact with a desired target molecule not naturally recognized by the receptor, and fragments of such receptors (see, *e.g.*, Skerra, 2000, *J. Molecular Recognition*, 13:167-187). A preferred receptor is a chemokine receptor. Exemplary chemokine receptors have been described in, for example, Lapidot et al, 2002, *Exp Hematol*, 30:973-81 and Onuffer et al, 2002, *Trends Pharmacol Sci*, 23:459-67.

[00464] In other embodiments, the additional targeting moiety may comprise a ligand molecule, including, for example, ligands which naturally recognize a specific desired receptor of a target cell,

such as a Transferrin (Tf) ligand. Such ligand molecules include ligands that have been modified to increase their specificity of interaction with a target receptor, ligands that have been modified to interact with a desired receptor not naturally recognized by the ligand, and fragments of such ligands.

[00465] In still other embodiments, the targeting moiety may comprise an aptamer. Aptamers are oligonucleotides that are selected to bind specifically to a desired molecular structure of the target cell. Aptamers typically are the products of an affinity selection process similar to the affinity selection of phage display (also known as *in vitro* molecular evolution). The process involves performing several tandem iterations of affinity separation, *e.g.*, using a solid support to which the diseased immunogen is bound, followed by polymerase chain reaction (PCR) to amplify nucleic acids that bound to the immunogens. Each round of affinity separation thus enriches the nucleic acid population for molecules that successfully bind the desired immunogen. In this manner, a random pool of nucleic acids may be “educated” to yield aptamers that specifically bind target molecules. Aptamers typically are RNA, but may be DNA or analogs or derivatives thereof, such as, without limitation, peptide nucleic acids (PNAs) and phosphorothioate nucleic acids.

[00466] In some embodiments, the targeting moiety can comprise a photo-degradable ligand (*i.e.*, a ‘caged’ ligand) that is released, for example, from a focused beam of light such that the capsid free, non-viral vectors or the gene product are targeted to a specific tissue.

[00467] It is also contemplated herein that the compositions be delivered to multiple sites in one or more muscles of the subject. That is, injections can be in at least 2, at least 3, at least 4, at least 5, at least 6, at least 7, at least 8, at least 9, at least 10, at least 15, at least 20, at least 25, at least 30, at least 35, at least 40, at least 45, at least 50, at least 55, at least 60, at least 65, at least 70, at least 75, at least 80, at least 85, at least 90, at least 95, at least 100 injection sites. Such sites can be spread over the area of a single muscle or can be distributed among multiple muscles.

B. Administration of the ceDNA vector for expression of FIX protein to non-muscle locations

[00468] In another embodiment, a ceDNA vector for expression of FIX protein is administered to the liver. The ceDNA vector may also be administered to different regions of the eye such as the cornea and/or optic nerve. The ceDNA vector may also be introduced into the spinal cord, brainstem (medulla oblongata, pons), midbrain (hypothalamus, thalamus, epithalamus, pituitary gland, substantia nigra, pineal gland), cerebellum, telencephalon (corpus striatum, cerebrum including the occipital, temporal, parietal and frontal lobes, cortex, basal ganglia, hippocampus and portaamygdala), limbic system, neocortex, corpus striatum, cerebrum, and inferior colliculus.. The ceDNA vector may be delivered into the cerebrospinal fluid (*e.g.*, by lumbar puncture). The ceDNA vector for expression of FIX protein may further be administered intravascularly to the CNS in situations in which the blood-brain barrier has been perturbed (*e.g.*, brain tumor or cerebral infarct).

[00469] In some embodiments, the ceDNA vector for expression of FIX protein can be administered to the desired region(s) of the eye by any route known in the art, including but not limited to, intrathecal, intra-ocular, intracerebral, intraventricular, intravenous (*e.g.*, in the presence of a sugar

such as mannitol), intranasal, intra-aural, intra-ocular (*e.g.*, intra-vitreous, sub-retinal, anterior chamber) and peri-ocular (*e.g.*, sub-Tenon's region) delivery as well as intramuscular delivery with retrograde delivery to motor neurons.

[00470] In some embodiments, the ceDNA vector for expression of FIX protein is administered in a liquid formulation by direct injection (*e.g.*, stereotactic injection) to the desired region or compartment in the CNS. In other embodiments, the ceDNA vector can be provided by topical application to the desired region or by intra-nasal administration of an aerosol formulation. Administration to the eye may be by topical application of liquid droplets. As a further alternative, the ceDNA vector can be administered as a solid, slow-release formulation (see, *e.g.*, U.S. Pat. No. 7,201,898). In yet additional embodiments, the ceDNA vector can be used for retrograde transport to treat, ameliorate, and/or prevent diseases and disorders involving motor neurons (*e.g.*, amyotrophic lateral sclerosis (ALS); spinal muscular atrophy (SMA), etc.). For example, the ceDNA vector can be delivered to muscle tissue from which it can migrate into neurons.

C. *Ex vivo* treatment

[00471] In some embodiments, cells are removed from a subject, a ceDNA vector for expression of FIX protein as disclosed herein is introduced therein, and the cells are then replaced back into the subject. Methods of removing cells from subject for treatment *ex vivo*, followed by introduction back into the subject are known in the art (see, *e.g.*, U.S. Pat. No. 5,399,346; the disclosure of which is incorporated herein in its entirety). Alternatively, a ceDNA vector is introduced into cells from another subject, into cultured cells, or into cells from any other suitable source, and the cells are administered to a subject in need thereof.

[00472] Cells transduced with a ceDNA vector for expression of FIX protein as disclosed herein are preferably administered to the subject in a "therapeutically-effective amount" in combination with a pharmaceutical carrier. Those skilled in the art will appreciate that the therapeutic effects need not be complete or curative, as long as some benefit is provided to the subject.

[00473] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein can encode an FIX protein as described herein (sometimes called a transgene or nucleic acid sequence) that is to be produced in a cell *in vitro*, *ex vivo*, or *in vivo*. For example, in contrast to the use of the ceDNA vectors described herein in a method of treatment as discussed herein, in some embodiments a ceDNA vector for expression of FIX protein may be introduced into cultured cells and the expressed FIX protein isolated from the cells, *e.g.*, for the production of antibodies and fusion proteins. In some embodiments, the cultured cells comprising a ceDNA vector for expression of FIX protein as disclosed herein can be used for commercial production of antibodies or fusion proteins, *e.g.*, serving as a cell source for small or large scale biomanufacturing of antibodies or fusion proteins. In alternative embodiments, a ceDNA vector for expression of FIX protein as disclosed herein is introduced into cells in a host non-human subject, for *in vivo* production of antibodies or fusion proteins, including small scale production as well as for commercial large scale FIX protein production.

[00474] The ceDNA vectors for expression of FIX protein as disclosed herein can be used in both veterinary and medical applications. Suitable subjects for *ex vivo* gene delivery methods as described above include both avians (*e.g.*, chickens, ducks, geese, quail, turkeys and pheasants) and mammals (*e.g.*, humans, bovines, ovines, caprines, equines, felines, canines, and lagomorphs), with mammals being preferred. Human subjects are most preferred. Human subjects include neonates, infants, juveniles, and adults.

D. Dose ranges

[00475] Provided herein are methods of treatment comprising administering to the subject an effective amount of a composition comprising a ceDNA vector encoding an FIX protein as described herein. As will be appreciated by a skilled practitioner, the term “effective amount” refers to the amount of the ceDNA composition administered that results in expression of the FIX protein in a “therapeutically effective amount” for the treatment of hemophilia B.

[00476] *In vivo* and/or *in vitro* assays can optionally be employed to help identify optimal dosage ranges for use. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the condition, and should be decided according to the judgment of the person of ordinary skill in the art and each subject's circumstances. Effective doses can be extrapolated from dose-response curves derived from *in vitro* or animal model test systems

[00477] A ceDNA vector for expression of FIX protein as disclosed herein is administered in sufficient amounts to transfect the cells of a desired tissue and to provide sufficient levels of gene transfer and expression without undue adverse effects. Conventional and pharmaceutically acceptable routes of administration include, but are not limited to, those described above in the “Administration” section, such as direct delivery to the selected organ (*e.g.*, intraportal delivery to the liver), oral, inhalation (including intranasal and intratracheal delivery), intraocular, intravenous, intramuscular, subcutaneous, intradermal, intratumoral, and other parental routes of administration. Routes of administration can be combined, if desired.

[00478] The dose of the amount of a ceDNA vectors for expression of FIX protein as disclosed herein required to achieve a particular “therapeutic effect,” will vary based on several factors including, but not limited to: the route of nucleic acid administration, the level of gene or RNA expression required to achieve a therapeutic effect, the specific disease or disorder being treated, and the stability of the gene(s), RNA product(s), or resulting expressed protein(s). One of skill in the art can readily determine a ceDNA vector dose range to treat a patient having a particular disease or disorder based on the aforementioned factors, as well as other factors that are well known in the art.

[00479] Dosage regime can be adjusted to provide the optimum therapeutic response. For example, the oligonucleotide can be repeatedly administered, *e.g.*, several doses can be administered daily or the dose can be proportionally reduced as indicated by the exigencies of the therapeutic situation. One of ordinary skill in the art will readily be able to determine appropriate doses and schedules of

administration of the subject oligonucleotides, whether the oligonucleotides are to be administered to cells or to subjects.

[00480] A “therapeutically effective dose” will fall in a relatively broad range that can be determined through clinical trials and will depend on the particular application (neural cells will require very small amounts, while systemic injection would require large amounts). For example, for direct *in vivo* injection into skeletal or cardiac muscle of a human subject, a therapeutically effective dose will be on the order of from about 1 µg to 100 g of the ceDNA vector. If exosomes or microparticles are used to deliver the ceDNA vector, then a therapeutically effective dose can be determined experimentally, but is expected to deliver from 1 µg to about 100 g of vector. Moreover, a therapeutically effective dose is an amount ceDNA vector that expresses a sufficient amount of the transgene to have an effect on the subject that results in a reduction in one or more symptoms of the disease, but does not result in significant off-target or significant adverse side effects. In one embodiment, a “therapeutically effective amount” is an amount of an expressed FIX protein that is sufficient to produce a statistically significant, measurable change in expression of hemophilia B biomarker or reduction of a given disease symptom. Such effective amounts can be gauged in clinical trials as well as animal studies for a given ceDNA vector composition.

[00481] Formulation of pharmaceutically-acceptable excipients and carrier solutions is well-known to those of skill in the art, as is the development of suitable dosing and treatment regimens for using the particular compositions described herein in a variety of treatment regimens.

[00482] For *in vitro* transfection, an effective amount of a ceDNA vectors for expression of FIX protein as disclosed herein to be delivered to cells (1×10^6 cells) will be on the order of 0.1 to 100 µg ceDNA vector, preferably 1 to 20 µg, and more preferably 1 to 15 µg or 8 to 10 µg. Larger ceDNA vectors will require higher doses. If exosomes or microparticles are used, an effective *in vitro* dose can be determined experimentally but would be intended to deliver generally the same amount of the ceDNA vector.

[00483] For the treatment of hemophilia B, the appropriate dosage of a ceDNA vector that expresses an FIX protein as disclosed herein will depend on the specific type of disease to be treated, the type of a FIX protein, the severity and course of the hemophilia B disease, previous therapy, the patient's clinical history and response to the antibody, and the discretion of the attending physician. The ceDNA vector encoding a FIX protein is suitably administered to the patient at one time or over a series of treatments. Various dosing schedules including, but not limited to, single or multiple administrations over various time-points, bolus administration, and pulse infusion are contemplated herein.

[00484] Depending on the type and severity of the disease, a ceDNA vector is administered in an amount that the encoded FIX protein is expressed at about 0.3 mg/kg to 100 mg/kg (*e.g.* 15 mg/kg-100 mg/kg, or any dosage within that range), by one or more separate administrations, or by continuous infusion. One typical daily dosage of the ceDNA vector is sufficient to result in the expression of the encoded FIX protein at a range from about 15 mg/kg to 100 mg/kg or more, depending on the factors

mentioned above. One exemplary dose of the ceDNA vector is an amount sufficient to result in the expression of the encoded FIX protein as disclosed herein in a range from about 10 mg/kg to about 50 mg/kg. Thus, one or more doses of a ceDNA vector in an amount sufficient to result in the expression of the encoded FIX protein at about 0.5 mg/kg, 1 mg/kg, 1.5 mg/kg, 2.0 mg/kg, 3 mg/kg, 4.0 mg/kg, 5 mg/kg, 10 mg/kg, 15 mg/kg, 20 mg/kg, 25 mg/kg, 30 mg/kg, 35 mg/kg, 40 mg/kg, 50 mg/kg, 60 mg/kg, 70 mg/kg, 80 mg/kg, 90 mg/kg, or 100 mg/kg (or any combination thereof) may be administered to the patient. In some embodiments, the ceDNA vector is an amount sufficient to result in the expression of the encoded FIX protein for a total dose in the range of 50 mg to 2500 mg. An exemplary dose of a ceDNA vector is an amount sufficient to result in the total expression of the encoded FIX protein at about 50 mg, about 100 mg, 200 mg, 300 mg, 400 mg, about 500 mg, about 600 mg, about 700 mg, about 720 mg, about 1000 mg, about 1050 mg, about 1100 mg, about 1200 mg, about 1300 mg, about 1400 mg, about 1500 mg, about 1600 mg, about 1700 mg, about 1800 mg, about 1900 mg, about 2000 mg, about 2050 mg, about 2100 mg, about 2200 mg, about 2300 mg, about 2400 mg, or about 2500 mg (or any combination thereof). As the expression of the FIX protein from ceDNA vector can be carefully controlled by regulatory switches herein, or alternatively multiple dose of the ceDNA vector administered to the subject, the expression of the FIX protein from the ceDNA vector can be controlled in such a way that the doses of the expressed FIX protein may be administered intermittently, *e.g.* every week, every two weeks, every three weeks, every four weeks, every month, every two months, every three months, or every six months from the ceDNA vector. The progress of this therapy can be monitored by conventional techniques and assays.

[00485] In certain embodiments, a ceDNA vector is administered an amount sufficient to result in the expression of the encoded FIX protein at a dose of 15 mg/kg, 30 mg/kg, 40 mg/kg, 45 mg/kg, 50 mg/kg, 60 mg/kg or a flat dose, *e.g.*, 300 mg, 500 mg, 700 mg, 800 mg, or higher. In some embodiments, the expression of the FIX protein from the ceDNA vector is controlled such that the FIX protein is expressed every day, every other day, every week, every 2 weeks or every 4 weeks for a period of time. In some embodiments, the expression of the FIX protein from the ceDNA vector is controlled such that the FIX protein is expressed every 2 weeks or every 4 weeks for a period of time. In certain embodiments, the period of time is 6 months, one year, eighteen months, two years, five years, ten years, 15 years, 20 years, or the lifetime of the patient.

[00486] Treatment can involve administration of a single dose or multiple doses. In some embodiments, more than one dose can be administered to a subject; in fact, multiple doses can be administered as needed, because the ceDNA vector does not elicit an anti-capsid host immune response due to the absence of a viral capsid. As such, one of skill in the art can readily determine an appropriate number of doses. The number of doses administered can, for example, be on the order of 1-100, preferably 2-20 doses.

[00487] Without wishing to be bound by any particular theory, the lack of typical anti-viral immune response elicited by administration of a ceDNA vector as described by the disclosure (*i.e.*, the absence

of capsid components) allows the ceDNA vector for expression of FIX protein to be administered to a host on multiple occasions. In some embodiments, the number of occasions in which a nucleic acid is delivered to a subject is in a range of 2 to 10 times (*e.g.*, 2, 3, 4, 5, 6, 7, 8, 9, or 10 times). In some embodiments, a ceDNA vector is delivered to a subject more than 10 times.

[00488] In some embodiments, a dose of a ceDNA vector for expression of FIX protein as disclosed herein is administered to a subject no more than once per calendar day (*e.g.*, a 24-hour period). In some embodiments, a dose of a ceDNA vector is administered to a subject no more than once per 2, 3, 4, 5, 6, or 7 calendar days. In some embodiments, a dose of a ceDNA vector for expression of FIX protein as disclosed herein is administered to a subject no more than once per calendar week (*e.g.*, 7 calendar days). In some embodiments, a dose of a ceDNA vector is administered to a subject no more than bi-weekly (*e.g.*, once in a two calendar week period). In some embodiments, a dose of a ceDNA vector is administered to a subject no more than once per calendar month (*e.g.*, once in 30 calendar days). In some embodiments, a dose of a ceDNA vector is administered to a subject no more than once per six calendar months. In some embodiments, a dose of a ceDNA vector is administered to a subject no more than once per calendar year (*e.g.*, 365 days or 366 days in a leap year).

[00489] In some embodiments, a dose of a ceDNA vector is administered on day 0. Following the initial treatment at day 0, a second dosing (re-dose) can be performed in about 1 week, about 2 weeks, about 3 weeks, about 4 weeks, about 5 weeks, about 6 weeks, about 7 weeks, about 8 weeks, or about 3 months, about 4 months, about 5 months, about 6 months, about 7 months, about 8 months, about 9 months, about 10 months, about 11 months, or about 1 year, about 2 years, about 3 years, about 4 years, about 5 years, about 6 years, about 7 years, about 8 years, about 9 years, about 10 years, about 11 years, about 12 years, about 13 years, about 14 years, about 15 years, about 16 years, about 17 years, about 18 years, about 19 years, about 20 years, about 21 years, about 22 years, about 23 years, about 24 years, about 25 years, about 26 years, about 27 years, about 28 years, about 29 years, about 30 years, about 31 years, about 32 years, about 33 years, about 34 years, about 35 years, about 36 years, about 37 years, about 38 years, about 39 years, about 40 years, about 41 years, about 42 years, about 43 years, about 44 years, about 45 years, about 46 years, about 47 years, about 48 years, about 49 years or about 50 years after the initial treatment with the ceDNA vector.

[00490] According to some embodiments, re-dosing of the therapeutic nucleic acid results in an increase in expression of the therapeutic nucleic acid. According to some embodiments, the increase of expression of the therapeutic nucleic acid after re-dosing, compared to the expression of the therapeutic nucleic acid after the first dose is about 0.5-fold to about 10-fold, about 1-fold to about 5-fold, about 1-fold to about 2-fold, or about 0.5-fold, about 1-fold, about 2-fold, about 3-fold, about 4-fold, about 5-fold, about 6-fold, about 7-fold, about 8-fold, about 9-fold or about 10-fold higher after re-dosing of the therapeutic nucleic acid.

[00491] In particular embodiments, more than one administration (*e.g.*, two, three, four or more administrations) of a ceDNA vector for expression of FIX protein as disclosed herein may be employed to achieve the desired level of gene expression over a period of various intervals, *e.g.*, daily, weekly, monthly, yearly, etc.

[00492] In some embodiments, a therapeutic FIX protein encoded by a ceDNA vector as disclosed herein can be regulated by a regulatory switch, inducible or repressible promotor so that it is expressed in a subject for at least 1 hour, at least 2 hours, at least 5 hours, at least 10 hours, at least 12 hours, at least 18 hours, at least 24 hours, at least 36 hours, at least 48 hours, at least 72 hours, at least 1 week, at least 2 weeks, at least 1 month, at least 2 months, at least 6 months, at least 12 months/one year, at least 2 years, at least 5 years, at least 10 years, at least 15 years, at least 20 years, at least 30 years, at least 40 years, at least 50 years or more. In one embodiment, the expression can be achieved by repeated administration of the ceDNA vectors described herein at predetermined or desired intervals. Alternatively, a ceDNA vector for expression of FIX protein as disclosed herein can further comprise components of a gene editing system (*e.g.*, CRISPR/Cas, TALENs, zinc finger endonucleases etc) to permit insertion of the one or more nucleic acid sequences encoding the FIX protein for substantially permanent treatment or “curing” the disease. Such ceDNA vectors comprising gene editing components are disclosed in International Application PCT/US18/64242, and can include the 5' and 3' homology arms (*e.g.*, SEQ ID NO: 151-154, or sequences with at least 40%, 50%, 60%, 70% or 80% homology thereto) for insertion of the nucleic acid encoding aFIX protein into safe harbor regions, such as, but not including albumin gene or CCR5 gene. By way of example, a ceDNA vector expressing a FIX protein can comprise at least one genomic safe harbor (GSH)-specific homology arms for insertion of the FIX transgene into a genomic safe harbor is disclosed in International Patent Application PCT/US2019/020225, filed on March 1, 2019, which is incorporated herein in its entirety by reference.

[00493] As described herein, according to some embodiments, a ceDNA vector expressing a FIX protein can be administered in combination with an additional compound.

[00494] Methods disclosed herein can comprise administering to the subject a combination of an immunosuppressant (*e.g.*, TKI or derivative or salt thereof) and a therapeutic nucleic acid (*e.g.* a ceDNA vector comprising a nucleic acid sequence encoding a FIX protein) in an effective amount to ameliorate a genetic disorder with a sufficient level of reduction in immune responses which allows for the safe administration of the therapeutic nucleic acid. These two agents can be administered at the same time in a co-formulation, at the same time in different formulations, or they can be administered separately at different times.

[00495] In some embodiments, a subject may be administered one or more immunosuppressants (or derivative or salt thereof) and a pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein as described herein concomitantly. For example, the method may comprise administering to a subject an immunosuppressant and a pharmaceutical composition comprising the

ceDNA vector for expression of FIX protein as described herein as two separate formulations but concomitantly. In another example, the method may comprise simultaneously administering to a subject an immunosuppressant and a pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein as described herein in one formulation, thereby the immunosuppressant and the therapeutic nucleic acid can be administered to a subject at the same time.

[00496] In some embodiment, a subject may be administered one or more immunosuppressants (or derivative or salt thereof) and a pharmaceutical composition comprising the ceDNA vector for expression of FIX protein as described herein sequentially. For example, the immunosuppressant may be administered prior to administration of the pharmaceutical composition.

[00497] In cases of sequential administration, there may be a delay period between administration of the one or more immunosuppressants and the pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein as described herein. For example, the immunosuppressant may be administered hours, days, or weeks prior to administration of the pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein as described herein (*e.g.*, at least 30 minutes, at least 1 hour, at least 2 hours, at least 3 hours, at least 4 hours, at least 5 hours, at least 6 hours, at least 7 hours, at least 8 hours, at least 9 hours, at least 10 hours, at least 11 hours, at least 12 hours, at least 13 hours, at least 14 hours, at least 15 hours, at least 16 hours, at least 17 hours, at least 18 hours, at least 19 hours, at least 20 hours, at least 21 hours, at least 22 hours, at least 23 hours, at least 24 hours, at least about 2 days, at least about 3 days, at least about 4 days, at least about 5 days, at least about 6 days, at least about 1 week, at least about 2 weeks, at least about 3 weeks, and at least about 4 weeks prior to the administration of a therapeutic nucleic acid). In some embodiments, an immunosuppressant may be administered about thirty (30) minutes prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant may be administered about one (1) hour prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about two (2) hours prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about three (3) hours prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about four (4) hours prior to the administration of a therapeutic nucleic acid. In some embodiments, an immunosuppressant can be administered about five (5) hours prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about six (6) hours prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about seven (7) hours prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about eight (8) hours prior to the administration of the pharmaceutical composition. In some embodiments, an immunosuppressant can be administered about nine (9) hours prior to the administration of the pharmaceutical composition. In

some embodiments, an immunosuppressant can be administered about ten (10) hours prior to the administration of the pharmaceutical composition.

[00498] In one embodiment, an immunosuppressant is administered no more than about 1 hour, about 2 hours, about 3 hours, about 4 hours, about 5 hours, about 6 hours, about 7 hours, about 8 hours, about 9 hours, about 10 hours, about 11 hours, about 12 hours, about 13 hours, about 14 hours, about 15 hours, about 16 hours, about 17 hours, about 18 hours, about 19 hours, about 20 hours, about 21 hours, about 22 hours, about 23 hours, or 24 hours before the administration of a pharmaceutical composition comprising the ceDNA vector for expression of FIX protein. In some embodiments, an immunosuppressant can be administered no more than about 1 day, about 2 days, about 3 days, about 4 days, about 6 days, or about 7 days before the administration of a pharmaceutical composition comprising the ceDNA vector for expression of FIX protein.

[00499] In some embodiments, an immunosuppressant can be administered about 30 minutes, about 1 hour, about 2 hours, about 3 hours, about 4 hours, about 5 hours, about 6 hours, about 7 hours, about 8 hours, about 9 hours, about 10 hours, about 11 hours, about 12 hours, about 13 hours, about 14 hours, about 15 hours, about 16 hours, about 17 hours, about 18 hours, about 19 hours, about 20 hours, about 21 hours, about 22 hours, about 23 hours, or 24 hours after the administration of a pharmaceutical composition comprising the ceDNA vector for expression of FIX protein. In some embodiments, an immunosuppressant can be administered about 1 day, about 2 days, about 3 days, about 4 days, about 6 days, or about 7 days after the administration of a pharmaceutical composition comprising the ceDNA vector for expression of FIX protein.

[00500] In one embodiment, an immunosuppressant is administered no more than about 1 hour, about 2 hours, about 3 hours, about 4 hours, about 5 hours, about 6 hours, about 7 hours, about 8 hours, about 9 hours, about 10 hours, about 11 hours, about 12 hours, about 13 hours, about 14 hours, about 15 hours, about 16 hours, about 17 hours, about 18 hours, about 19 hours, about 20 hours, about 21 hours, about 22 hours, about 23 hours, or 24 hours after the administration of a pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein. In some embodiments, an immunosuppressant can be administered no more than about 1 day, about 2 days, about 3 days, about 4 days, about 6 days, or about 7 days after the administration of a pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein.

[00501] In some embodiments, one or more immunosuppressants can be administered multiple times before, concurrently with, and/or after the administration of a pharmaceutical composition comprising the ceDNA vectors for expression of FIX protein.

In some embodiments, a ceDNA vector can be administered and re-dosed multiple times in conjunction with one or more immunosuppressant disclosed herein. For example, the ceDNA vector can be administered on day 0 with one or more immunosuppressants that is administered before, after or at the same time with the administration the therapeutic nucleic acid in a first dosing regimen. Following the initial treatment at day 0, a second dosing (re-dose) can be performed in about 1 week,

about 2 weeks, about 3 weeks, about 4 weeks, about 5 weeks, about 6 weeks, about 7 weeks, about 8 weeks, or about 3 months, about 4 months, about 5 months, about 6 months, about 7 months, about 8 months, about 9 months, about 10 months, about 11 months, or about 1 year, about 2 years, about 3 years, about 4 years, about 5 years, about 6 years, about 7 years, about 8 years, about 9 years, about 10 years, about 11 years, about 12 years, about 13 years, about 14 years, about 15 years, about 16 years, about 17 years, about 18 years, about 19 years, about 20 years, about 21 years, about 22 years, about 23 years, about 24 years, about 25 years, about 26 years, about 27 years, about 28 years, about 29 years, about 30 years, about 31 years, about 32 years, about 33 years, about 34 years, about 35 years, about 36 years, about 37 years, about 38 years, about 39 years, about 40 years, about 41 years, about 42 years, about 43 years, about 44 years, about 45 years, about 46 years, about 47 years, about 48 years, about 49 years or about 50 years after the initial treatment with the ceDNA vector, preferably with one or more immunosuppressants disclosed herein.

[00502] The duration of treatment depends upon the subject's clinical progress and responsiveness to therapy. Continuous, relatively low maintenance doses are contemplated after an initial higher therapeutic dose.

E. Unit dosage forms

[00503] In some embodiments, the pharmaceutical compositions comprising a ceDNA vector for expression of FIX protein as disclosed herein can conveniently be presented in unit dosage form. A unit dosage form will typically be adapted to one or more specific routes of administration of the pharmaceutical composition. In some embodiments, the unit dosage form is adapted for droplets to be administered directly to the eye. In some embodiments, the unit dosage form is adapted for administration by inhalation. In some embodiments, the unit dosage form is adapted for administration by a vaporizer. In some embodiments, the unit dosage form is adapted for administration by a nebulizer. In some embodiments, the unit dosage form is adapted for administration by an aerosolizer. In some embodiments, the unit dosage form is adapted for oral administration, for buccal administration, or for sublingual administration. In some embodiments, the unit dosage form is adapted for intravenous, intramuscular, or subcutaneous administration. In some embodiments, the unit dosage form is adapted for subretinal injection, suprachoroidal injection or intravitreal injection.

[00504] In some embodiments, the unit dosage form is adapted for intrathecal or intracerebroventricular administration. In some embodiments, the pharmaceutical composition is formulated for topical administration. The amount of active ingredient which can be combined with a carrier material to produce a single dosage form will generally be that amount of the compound which produces a therapeutic effect.

X. Methods of Treatment

[00505] The technology described herein also demonstrates methods for making, as well as methods of using the disclosed ceDNA vectors for expression of FIX protein in a variety of ways, including, for

example, *ex vivo*, *ex situ*, *in vitro* and *in vivo* applications, methodologies, diagnostic procedures, and/or gene therapy regimens.

[00506] In one embodiment, the expressed therapeutic FIX protein expressed from a ceDNA vector as disclosed herein is functional for the treatment of disease. In a preferred embodiment, the therapeutic FIX protein does not cause an immune system reaction, unless so desired.

[00507] Provided herein is a method of treating hemophilia B in a subject comprising introducing into a target cell in need thereof (for example, a muscle cell or tissue, or other affected cell type) of the subject a therapeutically effective amount of a ceDNA vector for expression of FIX protein as disclosed herein, optionally with a pharmaceutically acceptable carrier. While the ceDNA vector can be introduced in the presence of a carrier, such a carrier is not required. The ceDNA vector implemented comprises a nucleic acid sequence encoding an FIX protein as described herein useful for treating the disease. In particular, a ceDNA vector for expression of FIX protein as disclosed herein may comprise a desired FIX protein DNA sequence operably linked to control elements capable of directing transcription of the desired FIX protein encoded by the exogenous DNA sequence when introduced into the subject. The ceDNA vector for expression of FIX protein as disclosed herein can be administered via any suitable route as provided above, and elsewhere herein.

[00508] Disclosed herein are ceDNA vector compositions and formulations for expression of FIX protein as disclosed herein that include one or more of the ceDNA vectors of the present disclosure together with one or more pharmaceutically-acceptable buffers, diluents, or excipients. Such compositions may be included in one or more diagnostic or therapeutic kits, for diagnosing, preventing, treating or ameliorating one or more symptoms of hemophilia B. In one aspect the disease, injury, disorder, trauma or dysfunction is a human disease, injury, disorder, trauma or dysfunction.

[00509] Another aspect of the technology described herein provides a method for providing a subject in need thereof with a diagnostically- or therapeutically-effective amount of a ceDNA vector for expression of FIX protein as disclosed herein, the method comprising providing to a cell, tissue or organ of a subject in need thereof, an amount of the ceDNA vector as disclosed herein; and for a time effective to enable expression of the FIX protein from the ceDNA vector thereby providing the subject with a diagnostically- or a therapeutically-effective amount of the FIX protein expressed by the ceDNA vector. In a further aspect, the subject is human.

[00510] Another aspect of the technology described herein provides a method for diagnosing, preventing, treating, or ameliorating at least one or more symptoms of hemophilia B, a disorder, a dysfunction, an injury, an abnormal condition, or trauma in a subject. In an overall and general sense, the method includes at least the step of administering to a subject in need thereof one or more of the disclosed ceDNA vector for FIX protein production, in an amount and for a time sufficient to diagnose, prevent, treat or ameliorate the one or more symptoms of the disease, disorder, dysfunction, injury, abnormal condition, or trauma in the subject. In such an embodiment, the subject can be evaluated for efficacy of the FIX protein, or alternatively, detection of the FIX protein or tissue

location (including cellular and subcellular location) of the FIX protein in the subject. As such, the ceDNA vector for expression of FIX protein as disclosed herein can be used as an *in vivo* diagnostic tool, *e.g.*, for the detection of cancer or other indications. In a further aspect, the subject is human.

[00511] Another aspect is use of a ceDNA vector for expression of FIX protein as disclosed herein as a tool for treating or reducing one or more symptoms of hemophilia B or disease states. There are a number of inherited diseases in which defective genes are known, and typically fall into two classes: deficiency states, usually of enzymes, which are generally inherited in a recessive manner, and unbalanced states, which may involve regulatory or structural proteins, and which are typically but not always inherited in a dominant manner. For unbalanced disease states, a ceDNA vector for expression of FIX protein as disclosed herein can be used to create hemophilia B state in a model system, which could then be used in efforts to counteract the disease state. Thus, the ceDNA vector for expression of FIX protein as disclosed herein permit the treatment of genetic diseases. As used herein, hemophilia B state is treated by partially or wholly remedying the deficiency or imbalance that causes the disease or makes it more severe.

A. Host cells

[00512] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein delivers the FIX protein transgene into a subject host cell. In some embodiments, the cells are photoreceptor cells. In some embodiments, the cells are RPE cells. In some embodiments, the subject host cell is a human host cell, including, for example blood cells, stem cells, hematopoietic cells, CD34⁺ cells, liver cells, cancer cells, vascular cells, muscle cells, pancreatic cells, neural cells, ocular or retinal cells, epithelial or endothelial cells, dendritic cells, fibroblasts, or any other cell of mammalian origin, including, without limitation, hepatic (*i.e.*, liver) cells, lung cells, cardiac cells, pancreatic cells, intestinal cells, diaphragmatic cells, renal (*i.e.*, kidney) cells, neural cells, blood cells, bone marrow cells, or any one or more selected tissues of a subject for which gene therapy is contemplated. In one aspect, the subject host cell is a human host cell.

[00513] The present disclosure also relates to recombinant host cells as mentioned above, including a ceDNA vector for expression of FIX protein as disclosed herein. Thus, one can use multiple host cells depending on the purpose as is obvious to the skilled artisan. A construct or a ceDNA vector for expression of FIX protein as disclosed herein including donor sequence is introduced into a host cell so that the donor sequence is maintained as a chromosomal integrant as described earlier. The term host cell encompasses any progeny of a parent cell that is not identical to the parent cell due to mutations that occur during replication. The choice of a host cell will to a large extent depend upon the donor sequence and its source.

[00514] The host cell may also be a eukaryote, such as a mammalian, insect, plant, or fungal cell. In one embodiment, the host cell is a human cell (*e.g.*, a primary cell, a stem cell, or an immortalized cell line). In some embodiments, the host cell can be administered a ceDNA vector for expression of FIX protein as disclosed herein *ex vivo* and then delivered to the subject after the gene

therapy event. A host cell can be any cell type, *e.g.*, a somatic cell or a stem cell, an induced pluripotent stem cell, or a blood cell, *e.g.*, T-cell or B-cell, or bone marrow cell. In certain embodiments, the host cell is an allogenic cell. For example, T-cell genome engineering is useful for cancer immunotherapies, disease modulation such as HIV therapy (*e.g.*, receptor knock out, such as CXCR4 and CCR5) and immunodeficiency therapies. MHC receptors on B-cells can be targeted for immunotherapy. In some embodiments, gene modified host cells, *e.g.*, bone marrow stem cells, *e.g.*, CD34⁺ cells, or induced pluripotent stem cells can be transplanted back into a patient for expression of a therapeutic protein.

B. Additional diseases for gene therapy

[00515] In general, a ceDNA vector for expression of FIX protein as disclosed herein can be used to deliver any FIX protein in accordance with the description above to treat, prevent, or ameliorate the symptoms associated with hemophilia B related to an aborant protein expression or gene expression in a subject.

[00516] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein can be used to deliver an FIX protein to skeletal, cardiac or diaphragm muscle, for production of an FIX protein for secretion and circulation in the blood or for systemic delivery to other tissues to treat, ameliorate, and/or prevent hemophilia B.

[00517] The a ceDNA vector for expression of FIX protein as disclosed herein can be administered to the lungs of a subject by any suitable means, optionally by administering an aerosol suspension of respirable particles comprising the ceDNA vectors, which the subject inhales. The respirable particles can be liquid or solid. Aerosols of liquid particles comprising the ceDNA vectors may be produced by any suitable means, such as with a pressure-driven aerosol nebulizer or an ultrasonic nebulizer, as is known to those of skill in the art. See, *e.g.*, U.S. Pat. No. 4,501,729. Aerosols of solid particles comprising the ceDNA vectors may likewise be produced with any solid particulate medicament aerosol generator, by techniques known in the pharmaceutical art.

[00518] In some embodiments, a ceDNA vector for expression of FIX protein as disclosed herein can be administered to tissues of the CNS (*e.g.*, brain, eye). In some embodiments, a ceDNA vector for expression of FIX protein can be administered to tissues of the CNS, *e.g.*, ocular tissue, for the treatment of ocular hemorrhage associated with hemophilia.

[00519] In some aspects, ceDNA vectors expressing an FIX protein linked to a reporter polypeptide may be used for diagnostic purposes, as well as to determine efficacy or as markers of the ceDNA vector's activity in the subject to which they are administered.

C. Testing for successful gene expression using a ceDNA vector

[00520] Assays well known in the art can be used to test the efficiency of gene delivery of an FIX protein by a ceDNA vector can be performed in both *in vitro* and *in vivo* models. Levels of the expression of the FIX protein by ceDNA can be assessed by one skilled in the art by measuring mRNA and protein levels of the FIX protein (*e.g.*, reverse transcription PCR, western blot analysis, and

enzyme-linked immunosorbent assay (ELISA)). In one embodiment, ceDNA comprises a reporter protein that can be used to assess the expression of the FIX protein, for example by examining the expression of the reporter protein by fluorescence microscopy or a luminescence plate reader. For *in vivo* applications, protein function assays can be used to test the functionality of a given FIX protein to determine if gene expression has successfully occurred. One skilled will be able to determine the best test for measuring functionality of an FIX protein expressed by the ceDNA vector *in vitro* or *in vivo*.

[00521] It is contemplated herein that the effects of gene expression of an FIX protein from the ceDNA vector in a cell or subject can last for at least 1 month, at least 2 months, at least 3 months, at least four months, at least 5 months, at least six months, at least 10 months, at least 12 months, at least 18 months, at least 2 years, at least 5 years, at least 10 years, at least 20 years, or can be permanent.

[00522] In some embodiments, an FIX protein in the expression cassette, expression construct, or ceDNA vector described herein can be codon optimized for the host cell.

D. Determining Efficacy by Assessing FIX protein Expression from the ceDNA vector

[00523] Essentially any method known in the art for determining protein expression can be used to analyze expression of a FIX protein from a ceDNA vector. Non-limiting examples of such methods/assays include enzyme-linked immunoassay (ELISA), affinity ELISA, ELISPOT, serial dilution, flow cytometry, surface plasmon resonance analysis, kinetic exclusion assay, mass spectrometry, Western blot, immunoprecipitation, and PCR.

[00524] For assessing FIX protein expression *in vivo*, a biological sample can be obtained from a subject for analysis. Exemplary biological samples include a biofluid sample, a body fluid sample, blood (including whole blood), serum, plasma, urine, saliva, a biopsy and/or tissue sample etc. A biological sample or tissue sample can also refer to a sample of tissue or fluid isolated from an individual including, but not limited to, tumor biopsy, stool, spinal fluid, pleural fluid, nipple aspirates, lymph fluid, the external sections of the skin, respiratory, intestinal, and genitourinary tracts, tears, saliva, breast milk, cells (including, but not limited to, blood cells), tumors, organs, and also samples of *in vitro* cell culture constituent. The term also includes a mixture of the above-mentioned samples. The term "sample" also includes untreated or pretreated (or pre-processed) biological samples. In some embodiments, the sample used for the assays and methods described herein comprises a serum sample collected from a subject to be tested.

E. Determining Efficacy of the expressed FIX protein by Clinical Parameters

[00525] The efficacy of a given FIX protein expressed by a ceDNA vector for hemophilia B (*i.e.*, functional expression) can be determined by the skilled clinician. However, a treatment is considered "effective treatment," as the term is used herein, if any one or all of the signs or symptoms of hemophilia B is/are altered in a beneficial manner, or other clinically accepted symptoms or markers of disease are improved, or ameliorated, *e.g.*, by at least 10% following treatment with a ceDNA vector encoding a therapeutic FIX protein as described herein. Efficacy can also be measured by failure of an individual to worsen as assessed by stabilization of hemophilia B, or the need for medical

interventions (*i.e.*, progression of the disease is halted or at least slowed). Methods of measuring these indicators are known to those of skill in the art and/or described herein. Treatment includes any treatment of a disease in an individual or an animal (some non-limiting examples include a human, or a mammal) and includes: (1) inhibiting hemophilia B, *e.g.*, arresting, or slowing progression of hemophilia B; or (2) relieving the hemophilia B, *e.g.*, causing regression of a hemophilia B symptom; and (3) preventing or reducing the likelihood of the development of the hemophilia B disease, or preventing secondary diseases/disorders associated with hemophilia B. An effective amount for the treatment of a disease means that amount which, when administered to a mammal in need thereof, is sufficient to result in effective treatment as that term is defined herein, for that disease. Efficacy of an agent can be determined by assessing physical indicators that are particular to hemophilia B disease. A physician can assess for any one or more of clinical symptoms of hemophilia B which include: ******(i) reduced serum Factor IX. Reduction in FIX is a key biomarker in the development of treatments for hemophilia B.

XI. Various applications of ceDNA vectors expressing antibodies or fusion proteins

[00526] As disclosed herein, the compositions and ceDNA vectors for expression of FIX protein as described herein can be used to express an FIX protein for a range of purposes. In one embodiment, the ceDNA vector expressing an FIX protein can be used to create a somatic transgenic animal model harboring the transgene, *e.g.*, to study the function or disease progression of hemophilia B. In some embodiments, a ceDNA vector expressing an FIX protein is useful for the treatment, prevention, or amelioration of hemophilia B states or disorders in a mammalian subject.

[00527] In some embodiments the FIX protein can be expressed from the ceDNA vector in a subject in a sufficient amount to treat a disease associated with increased expression, increased activity of the gene product, or inappropriate upregulation of a gene.

[00528] In some embodiments the FIX protein can be expressed from the ceDNA vector in a subject in a sufficient amount to treat hemophilia B with a reduced expression, lack of expression or dysfunction of a protein.

[00529] It will be appreciated by one of ordinary skill in the art that the transgene may not be an open reading frame of a gene to be transcribed itself; instead it may be a promoter region or repressor region of a target gene, and the ceDNA vector may modify such region with the outcome of so modulating the expression of the FIX gene.

[00530] The compositions and ceDNA vectors for expression of FIX protein as disclosed herein can be used to deliver an FIX protein for various purposes as described above.

[00531] In some embodiments, the transgene encodes one or more FIX proteins which are useful for the treatment, amelioration, or prevention of hemophilia B states in a mammalian subject. The FIX protein expressed by the ceDNA vector is administered to a patient in a sufficient amount to treat hemophilia B associated with an abnormal gene sequence, which can result in any one or more of the

following: increased protein expression, over activity of the protein, reduced expression, lack of expression or dysfunction of the target gene or protein.

[00532] In some embodiments, the ceDNA vectors for expression of FIX protein as disclosed herein are envisioned for use in diagnostic and screening methods, whereby an FIX protein is transiently or stably expressed in a cell culture system, or alternatively, a transgenic animal model.

[00533] Another aspect of the technology described herein provides a method of transducing a population of mammalian cells with a ceDNA vector for expression of FIX protein as disclosed herein. In an overall and general sense, the method includes at least the step of introducing into one or more cells of the population, a composition that comprises an effective amount of one or more of the ceDNA vectors for expression of FIX protein as disclosed herein.

[00534] Additionally, the present disclosure provides compositions, as well as therapeutic and/or diagnostic kits that include one or more of the disclosed ceDNA vectors for expression of FIX protein as disclosed herein or ceDNA compositions, formulated with one or more additional ingredients, or prepared with one or more instructions for their use.

[00535] A cell to be administered a ceDNA vector for expression of FIX protein as disclosed herein may be of any type, including but not limited to neural cells (including cells of the peripheral and central nervous systems, in particular, brain cells), lung cells, retinal cells, epithelial cells (*e.g.*, gut and respiratory epithelial cells), muscle cells, dendritic cells, pancreatic cells (including islet cells), hepatic cells, myocardial cells, bone cells (*e.g.*, bone marrow stem cells), hematopoietic stem cells, spleen cells, keratinocytes, fibroblasts, endothelial cells, prostate cells, germ cells, and the like. Alternatively, the cell may be any progenitor cell. As a further alternative, the cell can be a stem cell (*e.g.*, neural stem cell, liver stem cell). As still a further alternative, the cell may be a cancer or tumor cell. Moreover, the cells can be from any species of origin, as indicated above.

A. Production and Purification of ceDNA vectors expressing FIX

[00536] The ceDNA vectors disclosed herein are to be used to produce FIX protein either *in vitro* or *in vivo*. The FIX proteins produced in this manner can be isolated, tested for a desired function, and purified for further use in research or as a therapeutic treatment. Each system of protein production has its own advantages/disadvantages. While proteins produced *in vitro* can be easily purified and can proteins in a short time, proteins produced *in vivo* can have post-translational modifications, such as glycosylation.

[00537] FIX therapeutic protein produced using ceDNA vectors can be purified using any method known to those of skill in the art, for example, ion exchange chromatography, affinity chromatography, precipitation, or electrophoresis.

[00538] An FIX therapeutic protein produced by the methods and compositions described herein can be tested for binding to the desired target protein.

EXAMPLES

[00539] The following examples are provided by way of illustration not limitation. It will be appreciated by one of ordinary skill in the art that ceDNA vectors can be constructed from any of the wild-type or modified ITRs described herein, and that the following exemplary methods can be used to construct and assess the activity of such ceDNA vectors. While the methods are exemplified with certain ceDNA vectors, they are applicable to any ceDNA vector in keeping with the description.

EXAMPLE 1: Constructing ceDNA Vectors Using an Insect Cell-Based Method

[00540] Production of the ceDNA vectors using a polynucleotide construct template is described in Example 1 of PCT/US18/49996, which is incorporated herein in its entirety by reference. For example, a polynucleotide construct template used for generating the ceDNA vectors of the present disclosure can be a ceDNA-plasmid, a ceDNA-Bacmid, and/or a ceDNA-baculovirus. Without being limited to theory, in a permissive host cell, in the presence of *e.g.*, Rep, the polynucleotide construct template having two symmetric ITRs and an expression construct, where at least one of the ITRs is modified relative to a wild-type ITR sequence, replicates to produce ceDNA vectors. ceDNA vector production undergoes two steps: first, excision (“rescue”) of template from the template backbone (*e.g.* ceDNA-plasmid, ceDNA-bacmid, ceDNA-baculovirus genome etc.) via Rep proteins, and second, Rep mediated replication of the excised ceDNA vector.

[00541] An exemplary method to produce ceDNA vectors is from a ceDNA-plasmid as described herein. Referring to **FIG. 1A and 1B**, the polynucleotide construct template of each of the ceDNA-plasmids includes both a left modified ITR and a right modified ITR with the following between the ITR sequences: (i) an enhancer/promoter; (ii) a cloning site for a transgene; (iii) a posttranscriptional response element (*e.g.* the woodchuck hepatitis virus posttranscriptional regulatory element (WPRE)); and (iv) a poly-adenylation signal (*e.g.* from bovine growth hormone gene (BGHpA). Unique restriction endonuclease recognition sites (R1-R6) (shown in **FIG. 1A** and **FIG. 1B**) were also introduced between each component to facilitate the introduction of new genetic components into the specific sites in the construct. R3 (PmeI) GTTTAAAC (SEQ ID NO: 123) and R4 (PacI) TTAATTAA (SEQ ID NO: 124) enzyme sites are engineered into the cloning site to introduce an open reading frame of a transgene. These sequences were cloned into a pFastBac HT B plasmid obtained from ThermoFisher Scientific.

[00542] Production of ceDNA-bacmids:

[00543] DH10Bac competent cells (MAX EFFICIENCY® DH10Bac™ Competent Cells, Thermo Fisher) were transformed with either test or control plasmids following a protocol according to the manufacturer’s instructions. Recombination between the plasmid and a baculovirus shuttle vector in the DH10Bac cells were induced to generate recombinant ceDNA-bacmids. The recombinant bacmids were selected by screening a positive selection based on blue-white screening in *E. coli* (Φ80dlacZAM15 marker provides α-complementation of the β-galactosidase gene from the bacmid

vector) on a bacterial agar plate containing X-gal and IPTG with antibiotics to select for transformants and maintenance of the bacmid and transposase plasmids. White colonies caused by transposition that disrupts the β -galactoside indicator gene were picked and cultured in 10 ml of media.

[00544] The recombinant ceDNA-bacmids were isolated from the *E. coli* and transfected into Sf9 or Sf21 insect cells using FugeneHD to produce infectious baculovirus. The adherent Sf9 or Sf21 insect cells were cultured in 50 ml of media in T25 flasks at 25°C. Four days later, culture medium (containing the P0 virus) was removed from the cells, filtered through a 0.45 μ m filter, separating the infectious baculovirus particles from cells or cell debris.

[00545] Optionally, the first generation of the baculovirus (P0) was amplified by infecting naïve Sf9 or Sf21 insect cells in 50 to 500 ml of media. Cells were maintained in suspension cultures in an orbital shaker incubator at 130 rpm at 25 °C, monitoring cell diameter and viability, until cells reach a diameter of 18-19 nm (from a naïve diameter of 14-15 nm), and a density of $\sim 4.0 \times 10^6$ cells/mL. Between 3 and 8 days post-infection, the P1 baculovirus particles in the medium were collected following centrifugation to remove cells and debris then filtration through a 0.45 μ m filter.

[00546] The ceDNA-baculovirus comprising the test constructs were collected and the infectious activity, or titer, of the baculovirus was determined. Specifically, four x 20 ml Sf9 cell cultures at 2.5×10^6 cells/ml were treated with P1 baculovirus at the following dilutions: 1/1000, 1/10,000, 1/50,000, 1/100,000, and incubated at 25-27°C. Infectivity was determined by the rate of cell diameter increase and cell cycle arrest and change in cell viability every day for 4 to 5 days.

[00547] A “Rep-plasmid” as disclosed in FIG. 8A of PCT/US18/49996, which is incorporated herein in its entirety by reference, was produced in a pFASTBAC™-Dual expression vector (ThermoFisher) comprising both the Rep78 (SEQ ID NO: 131 or 133) and Rep52 (SEQ ID NO: 132) or Rep68 (SEQ ID NO: 130) and Rep40 (SEQ ID NO: 129). The Rep-plasmid was transformed into the DH10Bac competent cells (MAX EFFICIENCY® DH10Bac™ Competent Cells (Thermo Fisher) following a protocol provided by the manufacturer. Recombination between the Rep-plasmid and a baculovirus shuttle vector in the DH10Bac cells were induced to generate recombinant bacmids (“Rep-bacmids”). The recombinant bacmids were selected by a positive selection that included-blue-white screening in *E. coli* ($\Phi 80$ dlacZ Δ M15 marker provides α -complementation of the β -galactosidase gene from the bacmid vector) on a bacterial agar plate containing X-gal and IPTG. Isolated white colonies were picked and inoculated in 10 ml of selection media (kanamycin, gentamicin, tetracycline in LB broth). The recombinant bacmids (Rep-bacmids) were isolated from the *E. coli* and the Rep-bacmids were transfected into Sf9 or Sf21 insect cells to produce infectious baculovirus.

[00548] The Sf9 or Sf21 insect cells were cultured in 50 ml of media for 4 days, and infectious recombinant baculovirus (“Rep-baculovirus”) were isolated from the culture. Optionally, the first generation Rep-baculovirus (P0) were amplified by infecting naïve Sf9 or Sf21 insect cells and cultured in 50 to 500 ml of media. Between 3 and 8 days post-infection, the P1 baculovirus particles in the medium were collected either by separating cells by centrifugation or filtration or another

fractionation process. The Rep-baculovirus were collected and the infectious activity of the baculovirus was determined. Specifically, four x 20 mL Sf9 cell cultures at 2.5×10^6 cells/mL were treated with P1 baculovirus at the following dilutions, 1/1000, 1/10,000, 1/50,000, 1/100,000, and incubated. Infectivity was determined by the rate of cell diameter increase and cell cycle arrest, and change in cell viability every day for 4 to 5 days.

[00549] ceDNA vector generation and characterization

[00550] With reference to **FIG. 4B**, Sf9 insect cell culture media containing either (1) a sample-containing a ceDNA-bacmid or a ceDNA-baculovirus, and (2) Rep-baculovirus described above were then added to a fresh culture of Sf9 cells (2.5×10^6 cells/ml, 20ml) at a ratio of 1:1000 and 1:10,000, respectively. The cells were then cultured at 130 rpm at 25°C. 4-5 days after the co-infection, cell diameter and viability are detected. When cell diameters reached 18-20nm with a viability of ~70-80%, the cell cultures were centrifuged, the medium was removed, and the cell pellets were collected. The cell pellets are first resuspended in an adequate volume of aqueous medium, either water or buffer. The ceDNA vector was isolated and purified from the cells using Qiagen MIDI PLUS™ purification protocol (Qiagen, 0.2mg of cell pellet mass processed per column).

[00551] Yields of ceDNA vectors produced and purified from the Sf9 insect cells were initially determined based on UV absorbance at 260nm.

[00552] ceDNA vectors can be assessed by identified by agarose gel electrophoresis under native or denaturing conditions as illustrated in **FIG. 4D**, where (a) the presence of characteristic bands migrating at twice the size on denaturing gels versus native gels after restriction endonuclease cleavage and gel electrophoretic analysis and (b) the presence of monomer and dimer (2x) bands on denaturing gels for uncleaved material is characteristic of the presence of ceDNA vector.

[00553] Structures of the isolated ceDNA vectors were further analyzed by digesting the DNA obtained from co-infected Sf9 cells (as described herein) with restriction endonucleases selected for a) the presence of only a single cut site within the ceDNA vectors, and b) resulting fragments that were large enough to be seen clearly when fractionated on a 0.8% denaturing agarose gel (>800 bp). As illustrated in **FIGS. 4D and 4E**, linear DNA vectors with a non-continuous structure and ceDNA vector with the linear and continuous structure can be distinguished by sizes of their reaction products—for example, a DNA vector with a non-continuous structure is expected to produce 1kb and 2kb fragments, while a non-encapsidated vector with the continuous structure is expected to produce 2kb and 4kb fragments.

[00554] Therefore, to demonstrate in a qualitative fashion that isolated ceDNA vectors are covalently closed-ended as is required by definition, the samples were digested with a restriction endonuclease identified in the context of the specific DNA vector sequence as having a single restriction site, preferably resulting in two cleavage products of unequal size (*e.g.*, 1000 bp and 2000 bp). Following digestion and electrophoresis on a denaturing gel (which separates the two complementary DNA strands), a linear, non-covalently closed DNA will resolve at sizes 1000 bp and

2000 bp, while a covalently closed DNA (*i.e.*, a ceDNA vector) will resolve at 2x sizes (2000 bp and 4000 bp), as the two DNA strands are linked and are now unfolded and twice the length (though single stranded). Furthermore, digestion of monomeric, dimeric, and *n*-meric forms of the DNA vectors will all resolve as the same size fragments due to the end-to-end linking of the multimeric DNA vectors (see FIG. 4D).

[00555] As used herein, the phrase “assay for the Identification of DNA vectors by agarose gel electrophoresis under native gel and denaturing conditions” refers to an assay to assess the closedness of the ceDNA by performing restriction endonuclease digestion followed by electrophoretic assessment of the digest products. One such exemplary assay follows, though one of ordinary skill in the art will appreciate that many art-known variations on this example are possible. The restriction endonuclease is selected to be a single cut enzyme for the ceDNA vector of interest that will generate products of approximately 1/3x and 2/3x of the DNA vector length. This resolves the bands on both native and denaturing gels. Before denaturation, it is important to remove the buffer from the sample. The Qiagen PCR clean-up kit or desalting “spin columns,” *e.g.* GE HEALTHCARE ILUSTRA™ MICROSPIN™ G-25 columns are some art-known options for the endonuclease digestion. The assay includes for example, i) digest DNA with appropriate restriction endonuclease(s), 2) apply to *e.g.*, a Qiagen PCR clean-up kit, elute with distilled water, iii) adding 10x denaturing solution (10x = 0.5 M NaOH, 10mM EDTA), add 10X dye, not buffered, and analyzing, together with DNA ladders prepared by adding 10X denaturing solution to 4x, on a 0.8 – 1.0 % gel previously incubated with 1mM EDTA and 200mM NaOH to ensure that the NaOH concentration is uniform in the gel and gel box, and running the gel in the presence of 1x denaturing solution (50 mM NaOH, 1mM EDTA). One of ordinary skill in the art will appreciate what voltage to use to run the electrophoresis based on size and desired timing of results. After electrophoresis, the gels are drained and neutralized in 1x TBE or TAE and transferred to distilled water or 1x TBE/TAE with 1x SYBR Gold. Bands can then be visualized with *e.g.* Thermo Fisher, SYBR® Gold Nucleic Acid Gel Stain (10,000X Concentrate in DMSO) and epifluorescent light (blue) or UV (312nm).

[00556] The purity of the generated ceDNA vector can be assessed using any art-known method. As one exemplary and non-limiting method, contribution of ceDNA-plasmid to the overall UV absorbance of a sample can be estimated by comparing the fluorescent intensity of ceDNA vector to a standard. For example, if based on UV absorbance 4μg of ceDNA vector was loaded on the gel, and the ceDNA vector fluorescent intensity is equivalent to a 2kb band which is known to be 1μg, then there is 1μg of ceDNA vector, and the ceDNA vector is 25% of the total UV absorbing material. Band intensity on the gel is then plotted against the calculated input that band represents – for example, if the total ceDNA vector is 8kb, and the excised comparative band is 2kb, then the band intensity would be plotted as 25% of the total input, which in this case would be .25μg for 1.0μg input. Using the ceDNA vector plasmid titration to plot a standard curve, a regression line equation is then used to

calculate the quantity of the ceDNA vector band, which can then be used to determine the percent of total input represented by the ceDNA vector, or percent purity.

[00557] For comparative purposes, Example 1 describes the production of ceDNA vectors using an insect cell based method and a polynucleotide construct template, and is also described in Example 1 of PCT/US18/49996, which is incorporated herein in its entirety by reference. For example, a polynucleotide construct template used for generating the ceDNA vectors of the present disclosure according to Example 1 can be a ceDNA-plasmid, a ceDNA-Bacmid, and/or a ceDNA-baculovirus. Without being limited to theory, in a permissive host cell, in the presence of *e.g.*, Rep, the polynucleotide construct template having two symmetric ITRs and an expression construct, where at least one of the ITRs is modified relative to a wild-type ITR sequence, replicates to produce ceDNA vectors. ceDNA vector production undergoes two steps: first, excision (“rescue”) of template from the template backbone (*e.g.* ceDNA-plasmid, ceDNA-bacmid, ceDNA-baculovirus genome etc.) via Rep proteins, and second, Rep mediated replication of the excised ceDNA vector.

[00558] An exemplary method to produce ceDNA vectors in a method using insect cell is from a ceDNA-plasmid as described herein. Referring to **FIG. 1A and 1B**, the polynucleotide construct template of each of the ceDNA-plasmids includes both a left modified ITR and a right modified ITR with the following between the ITR sequences: (i) an enhancer/promoter; (ii) a cloning site for a transgene; (iii) a posttranscriptional response element (*e.g.* the woodchuck hepatitis virus posttranscriptional regulatory element (WPRE)); and (iv) a poly-adenylation signal (*e.g.* from bovine growth hormone gene (BGHpA). Unique restriction endonuclease recognition sites (R1-R6) (shown in **FIG. 1A and FIG. 1B**) were also introduced between each component to facilitate the introduction of new genetic components into the specific sites in the construct. R3 (PmeI) GTTTAAAC (SEQ ID NO: 123) and R4 (PacI) TTAATTAA (SEQ ID NO: 124) enzyme sites are engineered into the cloning site to introduce an open reading frame of a transgene. These sequences were cloned into a pFastBac HT B plasmid obtained from ThermoFisher Scientific.

[00559] Production of ceDNA-bacmids:

[00560] DH10Bac competent cells (MAX EFFICIENCY® DH10Bac™ Competent Cells, Thermo Fisher) were transformed with either test or control plasmids following a protocol according to the manufacturer’s instructions. Recombination between the plasmid and a baculovirus shuttle vector in the DH10Bac cells were induced to generate recombinant ceDNA-bacmids. The recombinant bacmids were selected by screening a positive selection based on blue-white screening in *E. coli* (Φ80dlacZΔM15 marker provides α-complementation of the β-galactosidase gene from the bacmid vector) on a bacterial agar plate containing X-gal and IPTG with antibiotics to select for transformants and maintenance of the bacmid and transposase plasmids. White colonies caused by transposition that disrupts the β-galactoside indicator gene were picked and cultured in 10 ml of media.

[00561] The recombinant ceDNA-bacmids were isolated from the *E. coli* and transfected into Sf9 or Sf21 insect cells using FugeneHD to produce infectious baculovirus. The adherent Sf9 or Sf21 insect

cells were cultured in 50 ml of media in T25 flasks at 25°C. Four days later, culture medium (containing the P0 virus) was removed from the cells, filtered through a 0.45 µm filter, separating the infectious baculovirus particles from cells or cell debris.

[00562] Optionally, the first generation of the baculovirus (P0) was amplified by infecting naïve Sf9 or Sf21 insect cells in 50 to 500 ml of media. Cells were maintained in suspension cultures in an orbital shaker incubator at 130 rpm at 25 °C, monitoring cell diameter and viability, until cells reach a diameter of 18-19 nm (from a naïve diameter of 14-15 nm), and a density of ~4.0E+6 cells/mL. Between 3 and 8 days post-infection, the P1 baculovirus particles in the medium were collected following centrifugation to remove cells and debris then filtration through a 0.45 µm filter.

[00563] The ceDNA-baculovirus comprising the test constructs were collected and the infectious activity, or titer, of the baculovirus was determined. Specifically, four x 20 ml Sf9 cell cultures at 2.5E+6 cells/ml were treated with P1 baculovirus at the following dilutions: 1/1000, 1/10,000, 1/50,000, 1/100,000, and incubated at 25-27°C. Infectivity was determined by the rate of cell diameter increase and cell cycle arrest, and change in cell viability every day for 4 to 5 days.

[00564] A “Rep-plasmid” was produced in a pFASTBAC™-Dual expression vector (ThermoFisher) comprising both the Rep78 (SEQ ID NO: 131 or 133) or Rep68 (SEQ ID NO: 130) and Rep52 (SEQ ID NO: 132) or Rep40 (SEQ ID NO: 129). The Rep-plasmid was transformed into the DH10Bac competent cells (MAX EFFICIENCY® DH10Bac™ Competent Cells (Thermo Fisher) following a protocol provided by the manufacturer. Recombination between the Rep-plasmid and a baculovirus shuttle vector in the DH10Bac cells were induced to generate recombinant bacmids (“Rep-bacmids”). The recombinant bacmids were selected by a positive selection that included-blue-white screening in *E. coli* (Φ80dlacZΔM15 marker provides α-complementation of the β-galactosidase gene from the bacmid vector) on a bacterial agar plate containing X-gal and IPTG. Isolated white colonies were picked and inoculated in 10 ml of selection media (kanamycin, gentamicin, tetracycline in LB broth). The recombinant bacmids (Rep-bacmids) were isolated from the *E. coli* and the Rep-bacmids were transfected into Sf9 or Sf21 insect cells to produce infectious baculovirus.

[00565] The Sf9 or Sf21 insect cells were cultured in 50 ml of media for 4 days, and infectious recombinant baculovirus (“Rep-baculovirus”) were isolated from the culture. Optionally, the first generation Rep-baculovirus (P0) were amplified by infecting naïve Sf9 or Sf21 insect cells and cultured in 50 to 500 ml of media. Between 3 and 8 days post-infection, the P1 baculovirus particles in the medium were collected either by separating cells by centrifugation or filtration or another fractionation process. The Rep-baculovirus were collected and the infectious activity of the baculovirus was determined. Specifically, four x 20 mL Sf9 cell cultures at 2.5x10⁶ cells/mL were treated with P1 baculovirus at the following dilutions, 1/1000, 1/10,000, 1/50,000, 1/100,000, and incubated. Infectivity was determined by the rate of cell diameter increase and cell cycle arrest, and change in cell viability every day for 4 to 5 days.

[00566] ceDNA vector generation and characterization

[00567] Sf9 insect cell culture media containing either (1) a sample-containing a ceDNA-bacmid or a ceDNA-baculovirus, and (2) Rep-baculovirus described above were then added to a fresh culture of Sf9 cells (2.5E+6 cells/ml, 20ml) at a ratio of 1:1000 and 1:10,000, respectively. The cells were then cultured at 130 rpm at 25°C. 4-5 days after the co-infection, cell diameter and viability are detected. When cell diameters reached 18-20nm with a viability of ~70-80%, the cell cultures were centrifuged, the medium was removed, and the cell pellets were collected. The cell pellets are first resuspended in an adequate volume of aqueous medium, either water or buffer. The ceDNA vector was isolated and purified from the cells using Qiagen MIDI PLUS™ purification protocol (Qiagen, 0.2mg of cell pellet mass processed per column).

[00568] Yields of ceDNA vectors produced and purified from the Sf9 insect cells were initially determined based on UV absorbance at 260nm. The purified ceDNA vectors can be assessed for proper closed-ended configuration using the electrophoretic methodology described in Example 5.

EXAMPLE 2: Synthetic ceDNA production via excision from a double-stranded DNA molecule

[00569] Synthetic production of the ceDNA vectors is described in Examples 2-6 of International Application PCT/US19/14122, filed January 18, 2019, which is incorporated herein in its entirety by reference. One exemplary method of producing a ceDNA vector using a synthetic method that involves the excision of a double-stranded DNA molecule. In brief, a ceDNA vector can be generated using a double stranded DNA construct, *e.g.*, see FIGS. 7A-8E of PCT/US19/14122. In some embodiments, the double stranded DNA construct is a ceDNA plasmid, *e.g.*, see, *e.g.*, FIG. 6 in International patent application PCT/US2018/064242, filed December 6, 2018).

[00570] In some embodiments, a construct to make a ceDNA vector comprises a regulatory switch as described herein.

[00571] For illustrative purposes, Example 2 describes producing ceDNA vectors as exemplary closed-ended DNA vectors generated using this method. However, while ceDNA vectors are exemplified in this Example to illustrate *in vitro* synthetic production methods to generate a closed-ended DNA vector by excision of a double-stranded polynucleotide comprising the ITRs and expression cassette (*e.g.*, nucleic acid sequence) followed by ligation of the free 3' and 5' ends as described herein, one of ordinary skill in the art is aware that one can, as illustrated above, modify the double stranded DNA polynucleotide molecule such that any desired closed-ended DNA vector is generated, including but not limited to, doggybone DNA, dumbbell DNA and the like. Exemplary ceDNA vectors for production of antibodies or fusion proteins that can be produced by the synthetic production method described in Example 2 are discussed in the sections entitled "III ceDNA vectors in general". Exemplary antibodies and fusion proteins expressed by the ceDNA vectors are described in the section entitled "IIC Exemplary antibodies and fusion proteins expressed by the ceDNA vectors".

[00572] The method involves (i) excising a sequence encoding the expression cassette from a double-stranded DNA construct and (ii) forming hairpin structures at one or more of the ITRs and (iii) joining the free 5' and 3' ends by ligation, *e.g.*, by T4 DNA ligase.

[00573] The double-stranded DNA construct comprises, in 5' to 3' order: a first restriction endonuclease site; an upstream ITR; an expression cassette; a downstream ITR; and a second restriction endonuclease site. The double-stranded DNA construct is then contacted with one or more restriction endonucleases to generate double-stranded breaks at both of the restriction endonuclease sites. One endonuclease can target both sites, or each site can be targeted by a different endonuclease as long as the restriction sites are not present in the ceDNA vector template. This excises the sequence between the restriction endonuclease sites from the rest of the double-stranded DNA construct (see Fig. 9 of PCT/US19/14122). Upon ligation a closed-ended DNA vector is formed.

[00574] One or both of the ITRs used in the method may be wild-type ITRs. Modified ITRs may also be used, where the modification can include deletion, insertion, or substitution of one or more nucleotides from the wild-type ITR in the sequences forming B and B' arm and/or C and C' arm (see, *e.g.*, Figs. 6-8 and 10 FIG. 11B of PCT/US19/14122), and may have two or more hairpin loops (see, *e.g.*, Figs. 6-8 FIG. 11B of PCT/US19/14122) or a single hairpin loop (see, *e.g.*, Fig. 10A-10B FIG. 11B of PCT/US19/14122). The hairpin loop modified ITR can be generated by genetic modification of an existing oligo or by de novo biological and/or chemical synthesis.

[00575] In a non-limiting example, ITR-6 Left and Right (SEQ ID NOS: 111 and 112), include 40 nucleotide deletions in the B-B' and C-C' arms from the wild-type ITR of AAV2. Nucleotides remaining in the modified ITR are predicted to form a single hairpin structure. Gibbs free energy of unfolding the structure is about -54.4 kcal/mol. Other modifications to the ITR may also be made, including optional deletion of a functional Rep binding site or a Trs site.

EXAMPLE 3: ceDNA production via oligonucleotide construction

[00576] Another exemplary method of producing a ceDNA vector using a synthetic method that involves assembly of various oligonucleotides, is provided in Example 3 of PCT/US19/14122, where a ceDNA vector is produced by synthesizing a 5' oligonucleotide and a 3' ITR oligonucleotide and ligating the ITR oligonucleotides to a double-stranded polynucleotide comprising an expression cassette. FIG. 11B of PCT/US19/14122 shows an exemplary method of ligating a 5' ITR oligonucleotide and a 3' ITR oligonucleotide to a double stranded polynucleotide comprising an expression cassette.

[00577] As disclosed herein, the ITR oligonucleotides can comprise WT-ITRs (*e.g.*, see **FIG. 3A**, **FIG. 3C**), or modified ITRs (*e.g.*, see, **FIG. 3B** and **FIG. 3D**). (See also, *e.g.*, FIGS. 6A, 6B, 7A and 7B of PCT/US19/14122, which is incorporated herein in its entirety). Exemplary ITR oligonucleotides include, but are not limited to SEQ ID NOS: 134-145 (*e.g.*, see Table 7 in of PCT/US19/14122). Modified ITRs can include deletion, insertion, or substitution of one or more nucleotides from the

wild-type ITR in the sequences forming B and B' arm and/or C and C' arm. ITR oligonucleotides, comprising WT-ITRs or mod-ITRs as described herein, to be used in the cell-free synthesis, can be generated by genetic modification or biological and/or chemical synthesis. As discussed herein, the ITR oligonucleotides in Examples 2 and 3 can comprise WT-ITRs, or modified ITRs (mod-ITRs) in symmetrical or asymmetrical configurations, as discussed herein.

EXAMPLE 4: ceDNA production via a single-stranded DNA molecule

[00578] Another exemplary method of producing a ceDNA vector using a synthetic method is provided in Example 4 of PCT/US19/14122, and uses a single-stranded linear DNA comprising two sense ITRs which flank a sense expression cassette sequence and are attached covalently to two antisense ITRs which flank an antisense expression cassette, the ends of which single stranded linear DNA are then ligated to form a closed-ended single-stranded molecule. One non-limiting example comprises synthesizing and/or producing a single-stranded DNA molecule, annealing portions of the molecule to form a single linear DNA molecule which has one or more base-paired regions of secondary structure, and then ligating the free 5' and 3' ends to each other to form a closed single-stranded molecule.

[00579] An exemplary single-stranded DNA molecule for production of a ceDNA vector comprises, from 5' to 3':

- a sense first ITR;
- a sense expression cassette sequence;
- a sense second ITR;
- an antisense second ITR;
- an antisense expression cassette sequence; and
- an antisense first ITR.

[00580] A single-stranded DNA molecule for use in the exemplary method of Example 4 can be formed by any DNA synthesis methodology described herein, *e.g.*, *in vitro* DNA synthesis, or provided by cleaving a DNA construct (*e.g.*, a plasmid) with nucleases and melting the resulting dsDNA fragments to provide ssDNA fragments.

[00581] Annealing can be accomplished by lowering the temperature below the calculated melting temperatures of the sense and antisense sequence pairs. The melting temperature is dependent upon the specific nucleotide base content and the characteristics of the solution being used, *e.g.*, the salt concentration. Melting temperatures for any given sequence and solution combination are readily calculated by one of ordinary skill in the art.

[00582] The free 5' and 3' ends of the annealed molecule can be ligated to each other, or ligated to a hairpin molecule to form the ceDNA vector. Suitable exemplary ligation methodologies and hairpin molecules are described in Examples 2 and 3.

EXAMPLE 5: Purifying and/or confirming production of ceDNA

[00583] Any of the DNA vector products produced by the methods described herein, *e.g.*, including the insect cell based production methods described in Example 1, or synthetic production methods described in Examples 2-4 can be purified, *e.g.*, to remove impurities, unused components, or byproducts using methods commonly known by a skilled artisan; and/or can be analyzed to confirm that DNA vector produced, (in this instance, a ceDNA vector) is the desired molecule. An exemplary method for purification of the DNA vector, *e.g.*, ceDNA is using Qiagen Midi Plus purification protocol (Qiagen) and/or by gel purification,

[00584] The following is an exemplary method for confirming the identity of ceDNA vectors.

[00585] ceDNA vectors can be assessed by identified by agarose gel electrophoresis under native or denaturing conditions as illustrated in **FIG. 4D**, where (a) the presence of characteristic bands migrating at twice the size on denaturing gels versus native gels after restriction endonuclease cleavage and gel electrophoretic analysis and (b) the presence of monomer and dimer (2x) bands on denaturing gels for uncleaved material is characteristic of the presence of ceDNA vector.

[00586] Structures of the isolated ceDNA vectors were further analyzed by digesting the purified DNA with restriction endonucleases selected for a) the presence of only a single cut site within the ceDNA vectors, and b) resulting fragments that were large enough to be seen clearly when fractionated on a 0.8% denaturing agarose gel (>800 bp). As illustrated in **FIGS. 4C and 4D**, linear DNA vectors with a non-continuous structure and ceDNA vector with the linear and continuous structure can be distinguished by sizes of their reaction products—for example, a DNA vector with a non-continuous structure is expected to produce 1kb and 2kb fragments, while a ceDNA vector with the continuous structure is expected to produce 2kb and 4kb fragments.

[00587] Therefore, to demonstrate in a qualitative fashion that isolated ceDNA vectors are covalently closed-ended as is required by definition, the samples were digested with a restriction endonuclease identified in the context of the specific DNA vector sequence as having a single restriction site, preferably resulting in two cleavage products of unequal size (*e.g.*, 1000 bp and 2000 bp). Following digestion and electrophoresis on a denaturing gel (which separates the two complementary DNA strands), a linear, non-covalently closed DNA will resolve at sizes 1000 bp and 2000 bp, while a covalently closed DNA (*i.e.*, a ceDNA vector) will resolve at 2x sizes (2000 bp and 4000 bp), as the two DNA strands are linked and are now unfolded and twice the length (though single stranded). Furthermore, digestion of monomeric, dimeric, and *n*-meric forms of the DNA vectors will all resolve as the same size fragments due to the end-to-end linking of the multimeric DNA vectors (see **FIG. 4E**).

[00588] As used herein, the phrase “assay for the Identification of DNA vectors by agarose gel electrophoresis under native gel and denaturing conditions” refers to an assay to assess the close-endedness of the ceDNA by performing restriction endonuclease digestion followed by electrophoretic assessment of the digest products. One such exemplary assay follows, though one of ordinary skill in

the art will appreciate that many art-known variations on this example are possible. The restriction endonuclease is selected to be a single cut enzyme for the ceDNA vector of interest that will generate products of approximately 1/3x and 2/3x of the DNA vector length. This resolves the bands on both native and denaturing gels. Before denaturation, it is important to remove the buffer from the sample. The Qiagen PCR clean-up kit or desalting “spin columns,” *e.g.* GE HEALTHCARE ILUSTRA™ MICROSPIN™ G-25 columns are some art-known options for the endonuclease digestion. The assay includes for example, i) digest DNA with appropriate restriction endonuclease(s), 2) apply to *e.g.*, a Qiagen PCR clean-up kit, elute with distilled water, iii) adding 10x denaturing solution (10x = 0.5 M NaOH, 10mM EDTA), add 10X dye, not buffered, and analyzing, together with DNA ladders prepared by adding 10X denaturing solution to 4x, on a 0.8 – 1.0 % gel previously incubated with 1mM EDTA and 200mM NaOH to ensure that the NaOH concentration is uniform in the gel and gel box, and running the gel in the presence of 1x denaturing solution (50 mM NaOH, 1mM EDTA). One of ordinary skill in the art will appreciate what voltage to use to run the electrophoresis based on size and desired timing of results. After electrophoresis, the gels are drained and neutralized in 1x TBE or TAE and transferred to distilled water or 1x TBE/TAE with 1x SYBR Gold. Bands can then be visualized with *e.g.* Thermo Fisher, SYBR® Gold Nucleic Acid Gel Stain (10,000X Concentrate in DMSO) and epifluorescent light (blue) or UV (312nm). The foregoing gel-based method can be adapted to purification purposes by isolating the ceDNA vector from the gel band and permitting it to renature.

[00589] The purity of the generated ceDNA vector can be assessed using any art-known method. As one exemplary and non-limiting method, contribution of ceDNA-plasmid to the overall UV absorbance of a sample can be estimated by comparing the fluorescent intensity of ceDNA vector to a standard. For example, if based on UV absorbance 4μg of ceDNA vector was loaded on the gel, and the ceDNA vector fluorescent intensity is equivalent to a 2kb band which is known to be 1μg, then there is 1μg of ceDNA vector, and the ceDNA vector is 25% of the total UV absorbing material. Band intensity on the gel is then plotted against the calculated input that band represents – for example, if the total ceDNA vector is 8kb, and the excised comparative band is 2kb, then the band intensity would be plotted as 25% of the total input, which in this case would be .25μg for 1.0μg input. Using the ceDNA vector plasmid titration to plot a standard curve, a regression line equation is then used to calculate the quantity of the ceDNA vector band, which can then be used to determine the percent of total input represented by the ceDNA vector, or percent purity.

EXAMPLE 6: Controlled transgene expression from ceDNA: transgene expression from the ceDNA vector *in vivo* can be sustained and/or increased by re-dose administration.

[00590] A ceDNA vector was produced according to the methods described in Example 1 above, using a ceDNA plasmid comprising a CAG promoter (SEQ ID NO: 72) and a luciferase transgene (SEQ ID NO: 56) as an exemplary FIX, flanked between asymmetric ITRs (*e.g.*, a 5' WT-ITR (SEQ

ID NO: 2) and a 3' mod-ITR (SEQ ID NO: 3) and was assessed in different treatment paradigms *in vivo*. This ceDNA vector was used in all subsequent experiments described in Examples 6-10. In Example 6, the ceDNA vector was purified and formulated with a lipid nanoparticle (LNP ceDNA) and injected into the tail vein of each CD-1® IGS mice. Liposomes were formulated with a suitable lipid blend comprising four components to form lipid nanoparticles (LNP) liposomes, including cationic lipids, helper lipids, cholesterol and PEG-lipids.

[00591] To assess the sustained expression of the transgene *in vivo* from the ceDNA vector over a long time period, the LNP-ceDNA was administered in sterile PBS by tail vein intravenous injection to CD-1® IGS mice of approximately 5-7 weeks of age. Three different dosage groups were assessed: 0.1mg/kg, 0.5 mg/kg, and 1.0 mg/kg, ten mice per group (except 1.0 mg/kg which had 15 mice per group). Injections were administered on day 0. Five mice from each of the groups were injected with an additional identical dose on day 28. Luciferase expression was measured by IVIS imaging following intravenous administration into CD-1® IGS mice (Charles River Laboratories; WT mice). Luciferase expression was assessed by IVIS imaging following intraperitoneal injection of 150 mg/kg luciferin substrate on days 3, 4, 7, 14, 21, 28, 31, 35, and 42, and routinely (*e.g.*, weekly, biweekly or every 10-days or every 2 weeks), between days 42-110 days. Luciferase transgene expression as the exemplary FIX as measured by IVIS imaging for at least 132 days after 3 different administration protocols (data not shown).

[00592] An extension study was performed to investigate the effect of a re-dose, *e.g.*, a re-administration of LNP-ceDNA expressing luciferase of the LNP-ceDNA treated subjects. In particular, it was assessed to determine if expression levels can be increased by one or more additional administrations of the ceDNA vector.

[00593] In this study, the biodistribution of luciferase expression from a ceDNA vector was assessed by IVIS in CD-1® IGS mice after an initial intravenous administration of 1.0mg/kg (*i.e.*, a priming dose) at days 0 and 28 (Group A). A second administration of a ceDNA vector was administered via tail vein injection of 3mg/kg (Group B) or 10mg/kg (Group C) in 1.2 mL in the tail vein at day 84. In this study, five (5) CD-1® mice were used in each of Groups A, B and C. IVIS imaging of the mice for luciferase expression was performed prior to the additional dosing at days 49, 56, 63, and 70 as described above, as well as post-redose on day 84 and on days 91, 98, 105, 112, and 132. Luciferase expression was assessed and detected in all three Groups A, B and C until at least 110 days (the longest time period assessed).

[00594] The level of expression of luciferase was shown to be increased by a re-dose (*i.e.*, re-administration of the ceDNA composition) of the LNP-ceDNA-Luc, as determined by assessment of luciferase activity in the presence of luciferin. Luciferase transgene expression as an exemplary FIX as measured by IVIS imaging for at least 110 days after 3 different administration protocols (Groups A, B and C). The mice that had not been given any additional redose (1mg/kg priming dose (*i.e.*, Group A) treatment had stable luciferase expression observed over the duration of the study. The mice

in Group B that had been administered a re-dose of 3mg/kg of the ceDNA vector showed an approximately seven-fold increase in observed radiance relative to the mice in Group C. Surprisingly, the mice re-dosed with 10 mg/kg of the ceDNA vector had a 17-fold increase in observed luciferase radiance over the mice not receiving any redose (Group A).

[00595] Group A shows luciferase expression in CD-1[®] IGS mice after intravenous administration of 1mg/kg of a ceDNA vector into the tail vein at days 0 and 28. Group B and C show luciferase expression in CD-1[®] IGS mice administered 1mg/kg of a ceDNA vector at a first time point (day 0) and re-dosed with administration of a ceDNA vector at a second time point of 84 days. The second administration (*i.e.*, re-dose) of the ceDNA vector increased expression by at least 7-fold, even up to 17-fold.

[00596] A 3-fold increase in the dose (*i.e.*, the amount) of ceDNA vector in a re-dose administration in Group B (*i.e.*, 3mg/kg administered at re-dose) resulted in a 7-fold increase in expression of the luciferase. Also unexpectedly, a 10-fold increase in the amount of ceDNA vector in a re-dose administration (*i.e.*, 10mg/kg re-dose administered) in Group C resulted in a 17-fold increase in expression of the luciferase. Thus, the second administration (*i.e.*, re-dose) of the ceDNA increased expression by at least 7-fold, even up to 17-fold. This shows that the increase in transgene expression from the re-dose is greater than expected and dependent on the dose or amount of the ceDNA vector in the re-dose administration, and appears to be synergistic to the initial transgene expression from the initial priming administration at day 0. That is, the dose-dependent increase in transgene expression is not additive, rather, the expression level of the transgene is dose-dependent and greater than the sum of the amount of the ceDNA vector administered at each time point.

[00597] Both Groups B and C showed significant dose-dependent increase in expression of luciferase as compared to control mice (Group A) that were not re-dosed with a ceDNA vector at the second time point. Taken together, these data show that the expression of a transgene from ceDNA vector can be increased in a dose-dependent manner by re-dose (*i.e.*, re-administration) of the ceDNA vector at least a second time point.

[00598] Taken together, these data demonstrate that the expression level of a transgene, *e.g.*, FIX from ceDNA vectors can be maintained at a sustained level for at least 84 days and can be increased *in vivo* after a redose of the ceDNA vector administered at least at a second time point.

EXAMPLE 7: Sustained transgene expression *in vivo* of LNP-Formulated ceDNA vectors

[00599] The reproducibility of the results in Example 6 with a different lipid nanoparticle was assessed *in vivo* in mice. Mice were dosed on day 0 with either ceDNA vector comprising a luciferase transgene driven by a CAG promoter that was encapsulated in an LNP different from that used in Example 6 or with that same LNP comprising polyC but lacking ceDNA or a luciferase gene. Specifically, male CD-1[®] mice of approximately 4 weeks of age were treated with a single injection of 0.5 mg/kg LNP-TTX-luciferase or control LNP-polyC, administered intravenously via lateral tail

vein on day 0. At day 14 animals were dosed systemically with luciferin at 150 mg/kg via intraperitoneal injection at 2.5 mL/kg. At approximately 15 minutes after luciferin administration each animal was imaged using an *In vivo* Imaging System (“IVIS”).

[00600] As shown in **FIG. 7**, significant fluorescence in the liver was observed in all four ceDNA-treated mice, and very little other fluorescence was observed in the animals other than at the injection site, indicating that the LNP mediated liver-specific delivery of the ceDNA construct and that the delivered ceDNA vector was capable of controlled sustained expression of its transgene for at least two weeks after administration.

EXAMPLE 8: Sustained transgene expression in the liver *in vivo* from ceDNA vector administration

[00601] In a separate experiment, the localization of LNP-delivered ceDNA within the liver of treated animals was assessed. A ceDNA vector comprising a functional transgene of interest was encapsulated in the same LNP as used in Example 7 and administered to mice *in vivo* at a dose level of 0.5 mg/kg by intravenous injection. After 6 hours the mice were terminated and liver samples taken, formalin fixed and paraffin-embedded using standard protocols. RNAscope® in situ hybridization assays were performed to visualize the ceDNA vectors within the tissue using a probe specific for the ceDNA transgene and detecting using chromogenic reaction and hematoxylin staining (Advanced Cell Diagnostics). **FIG. 8** shows the results, which indicate that ceDNA is present in hepatocytes.

EXAMPLE 9: Sustained Ocular transgene Expression of ceDNA *in vivo*

[00602] The sustainability of ceDNA vector transgene expression in tissues other than the liver was assessed to determine tolerability and expression of a ceDNA vector after ocular administration *in vivo*. While luciferase was used as an exemplary transgene in Example 9, one of ordinary skill can readily substitute the luciferase transgene with an FIX sequence from any of those listed in **Table 1** or included in **Table 12**.

[00603] On day 0, male Sprague Dawley rats of approximately 9 weeks of age were injected sub-retinally with 5 µL of either ceDNA vector comprising a luciferase transgene formulated with jetPEI® transfection reagent (Polyplus) or plasmid DNA encoding luciferase formulated with jetPEI®, both at a concentration of 0.25 µg/µL. Four rats were tested in each group. Animals were sedated and injected sub-retinally in the right eye with the test article using a 33 gauge needle. The left eye of each animal was untreated. Immediately after injection eyes were checked with optical coherence tomography or fundus imaging in order to confirm the presence of a subretinal bleb. Rats were treated with buprenorphine and topical antibiotic ointment according to standard procedures.

[00604] At days 7, 14, 21, 28, and 35, the animals in both groups were dosed systemically with freshly made luciferin at 150 mg/kg via intraperitoneal injection at 2.5mL/kg. at 5-15 minutes post luciferin administration, all animals were imaged using IVIS while under isoflurane anesthesia. Total

Flux [p/s] and average Flux (p/s/sr/cm²) in a region of interest encompassing the eye were obtained over 5 minutes of exposure. The results were graphed as average radiance of each treatment group in the treated eye (“injected”) relative to the average radiance of each treatment group in the untreated eye (“uninjected”) (**FIG. 9B**). Significant fluorescence was readily detectable in the ceDNA vector-treated eyes but much weaker in the plasmid-treated eyes (**FIG. 9A**). After 35 days, the plasmid-injected rats were terminated, while the study continued for the ceDNA-treated rats, with luciferin injection and IVIS imaging at days 42, 49, 56, 63, 70, and 99. The results demonstrate that ceDNA vector introduced in a single injection to rat eye mediated transgene expression *in vivo* and that that expression was sustained at a high level at least through 99 days after injection.

EXAMPLE 10: Sustained dosing and redosing of ceDNA vector in Rag2 mice.

[00605] In situations where one or more of the transgenes encoded in the gene expression cassette of the ceDNA vector is expressed in a host environment (*e.g.*, cell or subject) where the expressed protein is recognized as foreign, the possibility exists that the host will mount an adaptive immune response that may result in undesired depletion of the expression product, which could potentially be confused for lack of expression. In some cases, this may occur with a reporter molecule that is heterologous to the normal host environment. Accordingly, ceDNA vector transgene expression was assessed *in vivo* in the Rag2 mouse model which lacks B and T cells and therefore does not mount an adaptive immune response to non-native murine proteins such as luciferase. Briefly, c57bl/6 and Rag2 knockout mice were dosed intravenously via tail vein injection with 0.5 mg/kg of LNP-encapsulated ceDNA vector expressing luciferase or a polyC control at day 0, and at day 21 certain mice were redosed with the same LNP-encapsulated ceDNA vector at the same dose level. All testing groups consisted of 4 mice each. IVIS imaging was performed after luciferin injection as described in Example 9 at weekly intervals.

[00606] Comparing the total flux observed from the IVIS analyses, the fluorescence observed in the wild-type mice (an indirect measure of the presence of expressed luciferase) dosed with LNP-ceDNA vector-Luc decreased gradually after day 21 whereas the Rag2 mice administered the same treatment displayed relatively constant sustained expression of luciferase over the 42-day experiment (**FIG. 10A**). The approximately 21-day time point of the observed decrease in the wild-type mice corresponds to the timeframe in which an adaptive immune response might expect to be produced. Re-administration of the LNP-ceDNA vector in the Rag2 mice resulted in a marked increase in expression which was sustained over the at least 21 days it was tracked in this study (**FIG. 10B**). The results suggest that adaptive immunity may play a role when a non-native protein is expressed from a ceDNA vector in a host, and that observed decreases in expression in the 20+ day timeframe from initial administration may signal a confounding adaptive immune response to the expressed molecule rather than (or in addition to) a decline in expression. Of note, this response is expected to be low

when expressing native proteins in a host where it is anticipated that the host will properly recognize the expressed molecules as self and will not develop such an immune response.

EXAMPLE 11: Impact of liver-specific expression and CpG modulation on sustained expression

[00607] As described in Example 10, undesired host immune response may in some cases artificially dampen what would otherwise be sustained expression of one or more desired transgenes from an introduced ceDNA vector. Two approaches were taken to assess the impact of avoiding and/or dampening potential host immune response on sustained expression from a ceDNA vector. First, since the ceDNA-Luc vector used in the preceding examples was under the control of a constitutive CAG promoter, a similar construct was made using a liver-specific promoter (hAAT) or a different constitutive promoter (hEF-1) to see whether avoiding prolonged exposure to myeloid cells or non-liver tissue reduced any observed immune effects. Second, certain of the ceDNA-luciferase constructs were engineered to be reduced in CpG content, a known trigger for host immune reaction. ceDNA-encoded luciferase gene expression upon administration of such engineered and promoter-switched ceDNA vectors to mice was measured.

[00608] Three different ceDNA vectors were used, each encoding luciferase as the transgene. The first ceDNA vector had a high number of unmethylated CpG (~350) and comprised the constitutive CAG promoter ("ceDNA CAG"); the second had a moderate number of unmethylated CpG (~60) and comprised the liver-specific hAAT promoter ("ceDNA hAAT low CpG"); and the third was a methylated form of the second, such that it contained no unmethylated CpG and also comprised the hAAT promoter ("ceDNA hAAT No CpG"). The ceDNA vectors were otherwise identical. The vectors were prepared as described above.

[00609] Four groups of four male CD-1® mice, approximately 4 weeks old, were treated with one of the ceDNA vectors encapsulated in an LNP or a polyC control. On day 0 each mouse was administered a single intravenous tail vein injection of 0.5 mg/kg ceDNA vector in a volume of 5 mL/kg. Body weights were recorded on days -1, -, 1, 2, 3, 7, and weekly thereafter until the mice were terminated. Whole blood and serum samples were taken on days 0, 1, and 35. In-life imaging was performed on days 7, 14, 21, 28, and 35, and weekly thereafter using an *in vivo* imaging system (IVIS). For the imaging, each mouse was injected with luciferin at 150 mg/kg via intraperitoneal injection at 2.5 mL/kg. After 15 minutes, each mouse was anaesthetized and imaged. The mice were terminated at day 93 and terminal tissues collected, including liver and spleen. Cytokine measurements were taken 6 hours after dosing on day 0.

[00610] While all of the ceDNA-treated mice displayed significant fluorescence at days 7 and 14, the fluorescence decreased rapidly in the ceDNA CAG mice after day 14 and more gradually decreased for the remainder of the study. In contrast, the total flux for the ceDNA hAAT low CpG and No CpG-treated mice remained at a steady high level (**Fig. 11**). This suggested that directing the ceDNA vector delivery specifically to the liver resulted in sustained, durable transgene expression

from the vector over at least 77 days after a single injection. Constructs that were CpG minimized or completely absent of CpG content (No CpG) had similar durable sustained expression profiles, while the high CpG constitutive promoter construct exhibited a decline in expression over time, suggesting that host immune activation by the ceDNA vector introduction may play a role in any decreased expression observed from such vector in a subject. These results provide alternative methods of tailoring the duration of the response to the desired level by selecting a tissue-restricted promoter and/or altering the CpG content of the ceDNA vector in the event that a host immune response is observed – a potentially transgene-specific response.

EXAMPLE 12: *In vivo* Effects of Selected Tyrosine Kinase Inhibitors

[00611] In an extended study, a therapeutic ceDNA carrying human Factor IX (FIX) was dosed in mice (n=4) to evaluate the effect of the combination of an immunosuppressant TKI (*e.g.*, ruxolitinib) and a therapeutic FIX nucleic acid on *in vivo* expression of FIX over a period of 56 days. The study design and details were carried out as set forth below.

[00612] Study Design

[00613] **Table 14** sets forth the design of the kinase inhibitor administration component of the study. As shown in **Table 14**, two groups of male CD-1 mice (Group 1, n=4; Group 2, n=4) were orally administered either vehicle or ruxolitinib (300 mg/kg) at a dose volume of 10 mL/kg. For both Groups 1 and 2 dosing was carried out at days -2, -1, 1, 0 and 36.

Table 14: Study Design of Kinase Inhibitor Administration

Group No.	Animals per Group	Inhibitor ^a	Dose Level (mg/kg)	Dose Volume (mL/kg)	Treatment Regimen, via <u>PO</u>
1	4	Vehicle	NA	10	Days -2, -1 & 1 Day 0: 90 min. pre-dose
2	4	Ruxolitinib	300		Day 36: 30 min. pre-dose & 5 hours post dose

No. = Number; PO = oral gavage; ROA = route of administration; min = minutes; hrs = hours. ^a

Vehicle for dosing and inhibitor preparation = 0.5% methylcellulose

[00614] **Table 15** sets forth the design of the test material administration component of the study. As shown in **Table 15**, one group of male CD-1 mice (Group 1, n=4) was intravenously administered either LNP:Empty on day 0 or LNP:Empty on day 36 at a dose level of 1 mg/kg and a dose volume of 5 mL/kg. The second group of male CD-1 mice (Group 2, n=4) was intravenously administered LNP:ceDNA-FIX on day 0 and re-dosed with LNP:ceDNA-FIX on day 36 at a dose level of 2 mg/kg and a dose volume of 5 mL/kg. Day 56 was the terminal time point of the study.

Table 15: Study Design of LNP:ceDNA-FIX Administration

Group No.	Animals per Group	Treatment	Dose Level (mg/kg)	Dose Volume (mL/kg)	Treatment Regimen, IV	Terminal Time Point
1	4	LNP:Empty (Day 0) or LNP:Empty (Day 36)	1.0	5	Once on Day 0 & 36	Day 56
2	4	LNP:ceDNA-FIX LNP:ceDNA-FIX (Day 36)	2.0			

No. = Number; IV = intravenous; ROA = route of administration

[00615] Sample Collection

[00616] Blood collection or plasma collection (interim) was carried out as follows. For mice in both Groups 1 and 2, a sample of 120 μ L of whole blood was collected orbitally on days 7, 14, 21, 28, 35, 42, 49, and 56. A plasma sample was collected from the blood on days 7, 14, 21, 28, 35, 42, 49, and 56. To process and store the blood samples, 120 μ L of whole blood was added to a tube pre-coated with 13.33 μ L of 3.2% sodium citrate and kept ambient until processed. To process and store the plasma samples, one aliquot of plasma was frozen at nominally -70°C.

[00617] Study Details

[00618] Body Weights: Body weights for all animals were be recorded on Days -2, -1, 0, 1, 2, 3, 7, 14, 21, 28, 35, 36, 37, 38, 39, 42, 49, and 56 (prior to euthanasia). Additional body weights were recorded as needed.

[00619] Interim Blood Collection: All animals in Groups 1 – 2 had interim blood collected on Day 0 at 6 hours post Test Material dose ($\pm 5\%$); then on Day 7, 14, 21, 28, 35, 42, 49, and 56 as indicated above.

[00620] Inhibitor Administration: Inhibitor or vehicle was dosed on Days -2, -1, 0 & 1 and again on Day 36 by PO administration (oral gavage) at 10 mL/kg. On Day 0, inhibitor or vehicle was dosed 1.5 hours (± 10 minutes) prior to the Day 0 ceDNA administration. On Day 36, inhibitor or vehicle was be dosed 0.5 hours (± 10 minutes) prior to the Day 0 ceDNA administration and 5 hours (± 20 minutes) post administration. Inhibitors were administered at approximately the same time each day (± 1 hour).

[00621] Dose Administration: Test articles were be dosed at 5 mL/kg on Day 0 and Day 36 for Groups 1 – 2 by intravenous (IV) administration via lateral tail vein.

[00622] Euthanasia & Terminal Collection: On Day 56, after bleed for plasma, animals were euthanized by CO₂ asphyxiation followed by thoracotomy or cervical dislocation. No tissues will be collected.

[00623] As shown in **FIG. 12**, mice treated with ruxolitinib (300 mg/kg) at days -2, -1, 1, 0 and 36 and LNP:ceDNA-FIX (2.0 mg/kg) at day 0 and day 36 expressed factor IX (FIX) protein (IU/mL) that was detected *in vivo* beginning at day 7 through the end of the study (day 56). Notably, re-dosing with ceDNA-FIX at day 36 resulted in a dramatic increase in FIX expression beyond day 42 to the end of

the study. In contrast, mice treated with vehicle control at -48h, -24h, -1.5h, and 24hr and LNP:empty (1.0 mg/kg) at day 0 or day 36 did not express FIX protein.

[00624] These results also demonstrated that a therapeutic nucleic acid (*e.g.*, ceDNA-FIX) can be administered and re-dosed multiple times in conjunction with one or more immunosuppressant TKIs (*e.g.* the JAK inhibitor ruxolitinib) in a therapeutic model. As shown in **FIG. 12**, the combination approach allowed for a re-dosing of ceDNA-FIX, which led to a considerable increase in FIX expression.

EXAMPLE 13: Hydrodynamic Delivery of ceDNA Expressing FIX

[00625] A well-known method of introducing nucleic acid to the liver in rodents is by hydrodynamic tail vein injection. In this system, the pressurized injection in a large volume of non-encapsulated nucleic acid results in a transient increase in cell permeability and delivery directly into tissues and cells. This provides an experimental mechanism to bypass many of the host immune systems, such as macrophage delivery, providing the opportunity to observe delivery and expression in the absence of such activity.

[00626] Two different ceDNA vectors, each with a wild-type left ITR and a truncation mutant right ITR and having a transgene region encoding FIX, were prepared and purified as described above in **Examples 1 and 5**. ceDNA FIX vectors under the control of a liver-specific promoter or PBS without vector were administered to male C57bl/6J mice of approximately 6 weeks of age. The naked ceDNA vectors were dosed at 0.005 mg per animal (4 animals per group) by hydrodynamic intravenous injection via lateral tail vein in a volume of 100mL/kg administered over 5-8 seconds. Body weights were measured on days 0, 1, 2, 3 and 7 and weekly thereafter. Blood samples were collected from each treated animal on days 3, 7, 14, 21, and at terminal day 28. The presence of expressed FIX in the plasma samples was measured using the Factor IX (F9, FIX) ELISA kit (Affinity Biologicals).

[00627] As shown in **FIG. 11A**, FIX was readily detected in day 3 and 7 plasma samples from mice treated with each of the ceDNA FIX vectors, but was not observed in mice treated with PBS. **FIG. 11B** shows that FIX expression persisted over the duration of the 28-day study, plateauing at the 21 day time point. This experiment demonstrated that ceDNA vectors were able to express FIX from the liver after hydrodynamic injection, and that FIX was rapidly and readily detectable in the plasma after ceDNA administration.

EXAMPLE 14: Identification of FIX Constructs via Hydrodynamic Delivery in male CD-1 mice

[00628] The objective of this study was to determine FIX protein expression after hydrodynamic injection of recombinant DNA.

[00629] STUDY DESIGN**Table 16: Study Design of ceDNA-FIX Hydrodynamic Administration**

Group No.	No. of Animals	Test Material	Dose Levels (µg/an)	Dose Volume (mL/kg)	Dosing Regimen ROA	Terminal Time Point
1	5	PBS	NA	90 – 100 ML/KG (SET VOLUME)	ONCE ON DAY 0 BY IV HYDRODYNA MIC	DAY 7
2	5	CEDNA-FIX V1	1.0			
3	5	CEDNA-FIX V1	10.0			
4	5	CEDNA-FIX 2109	1.0			
5	5	CEDNA-FIX 2109	10.0			
6	5	CEDNA-FIX 2112	1.0			
7	5	CEDNA-FIX 2112	10.0			

No. = Number; IV = intravenous; ROA = route of administration; an = animal

Species: *Mus Musculus*

Strain: CD-1

Number of Males: 35 plus 3 spares

Age: 4 weeks of age at arrival

[00630] CD-1 mice were group housed in clear polycarbonate cages with contact bedding and provided ad libitum Mouse Diet 5058 and filtered tap water acidified with 1N HCl to a targeted pH of 2.5-3.0.

[00631] Dose Formulation, Administration and Observation

[00632] ceDNA-FIX constructs (*i.e.*, ceDNA-FIX v1, ceDNA-FIX 2109 and ceDNA-FIX 2112) were warmed to room temperature and diluted with the provided PBS immediately prior to use. Sequence information of exemplary ceDNA-FIX v1 vectors is set forth herein in **Table 12**. ceDNA-FIX v1, 2109 and 2112 were dosed on Day 0 by hydrodynamic IV administration, at a set volume per animal, 90 – 100 mL / kg (dependent on the lightest animal in the group) via lateral tail vein (dosed within 5 seconds). Doses were rounded to the nearest 0.1 mL. Cage side animal health checks were performed at least once daily to check for general health, mortality and moribundity. Clinical observations were performed ~1-hour post dose, by the end of the dosing day (3-6 hours) and then ~24 hours post the Day 0 Test Material dose. Additional observations were made per exception. Body weights for all animals were recorded on Days 0, 1, 2, 3, 7. Additional body weights were recorded as needed.

[00633] Blood Collection

[00634] All animals in Groups 1 – 7, had interim blood collected on Day 3. After collection animals received 0.5 – 1.0 mL lactated Ringer's; subcutaneously. For plasma collections, whole blood were collected into non-coated Eppendorf style tubes via orbital sinus puncture under anesthesia. 120µL were withdrawn and placed into tubes containing 13.33 µL of 3.2% sodium citrate. Blood samples were gently mixed and maintained ambient until processed. Whole blood samples were centrifuged at

2,000g for 15 minutes under ambient conditions (20-25°C). Plasma samples were withdrawn avoiding the cell pack. Terminal whole blood were collected by syringe and 500 µL placed immediately tubes containing 55.6 µL of 3.2% sodium citrate.

[00635] Results

[00636] As shown in **FIG. 13A**, at 1 µg per animal, expression is equivalent for all three constructs. At a higher dose (10 µg per animal), ceDNA-FIX 2109 showed superior and increased level of FIX expression at Day 7 (**FIG. 13B**) as compared to the other constructs tested.

EXAMPLE 15: A Study to Evaluate ceDNA-FIX Formulations via IV Delivery in Male C57Bl/6 Mice

[00637] The following study was carried out to determine protein expression after IV injection of LNP formulated ceDNA. ceDNA-FIX was formulated in two different LNPs compositions (LNP formulation1: Ionizable lipid: DSPC : Cholesterol : PEG-Lipid + DSPE-PEG-GalNAc4 (47.5 : 10.0 : 39.2 : 3.3) (designated “DP No.1”); and LNP formulation 2: Ionizable lipid : DSPC : Cholesterol : PEG-Lipid + DSPE-PEG2000-GalNAc4 (47.3 : 10.0 : 40.5 : 2.3) (designated “DP No.2”). Doses of test material were administered on Day 0 by intravenous dosing into the lateral tail vein. LNPs containing ceDNA-FIX were administered at a dose volume of 5 mL/kg (2 mg/kg). Test materials for the study are shown in **Table 17** below. ceDNA expressing Factor IX (ceDNA-FIX) was used as an independent control.

Table 17

Group No.	No. of Animals	Test Material	Dose Levels (mg/kg)	Dose Volume (mL/kg)	Dosing Regimen ROA	Terminal Time Point
1	5	PBS	NA	5	Once on Day 0 by IV	Day 14
4	5	ceDNA-FIX v1	2.0			
5	5	ceDNA-FIX v1	2.0			
6	5	ceDNA-FIX v1	2.0			

[00638] Mice treated with ceDNA-FIX v1 LNP formulations (DP No. 1 or DP No. 2) exhibited the presence of human FIX in its plasma as compared to mice treated with vehicles that showed no hFIX, indicating that ceDNA-FIX v1 LNP formulation could successfully target and be integrated into cells which lead to expression of FIX protein. Overall, the ceDNA-FIX LNP formulations were well tolerated.

EXAMPLE 16: A 14-Day Single Dose Intravenous Infusion Toxicity Study of a Lipid Nano Particle Formulation in Cynomolgus Monkeys

[00639] The objective of this study was to determine the toxicity effects of a single intravenous (IV) dose of a lipid nanoparticle ceDNA transgene expression after IV administration of LNP formulated

ceDNA to male *Cynomolgus* monkeys. Dosing was by IV infusion (70 minutes $\pm$ 10 minutes) to the saphenous vein (cephalic or tail vein was used, if necessary) dosed at 0.42 mL/kg/hr for 15 min and then escalating to 4.59 mL/kg/hr for 55 min. Prolonged infusion with escalating dosing rate design was necessary to prevent/mitigate infusion reactions. The first day of dosing was designated as day 1. Dosing was performed once on day 1 and was carried out for 15 days. Study details are shown in

Table 18.

Table 18: Test material administration in ce-DNA-FIX formulation toxicity study

Group No.	Test Material	Dose Level (mg/kg/dose)	Dose Volume ^a (mL/kg)	Dose Concentration (mg/mL)	No. of Animals
					Males ^b
1	Vehicle	0	4.31	0	2
2	ceDNA-FIX v1	2.0	4.31	0.46	2
3	ceDNA-FIX v1	2.0	4.31	0.46	2
4	ceDNA-FIX v1	2.0	4.31	0.46	2

Based on the most recent body weight measurement. The first day of dosing was based on Day 1 body weights.

[00640] Prior to the start of infusion, the catheters were flushed with approximately 2 mL of sterile saline. Next the dosing formulations were administered at 0.42 mL/kg/hr for the first 15 minutes (target time). The infusion pump was stopped, reprogrammed to infuse the remaining dose for an infusion rate of 4.59 mL/kg/hr, for the remaining 55 minutes (target time) of the infusion. An approximate 1.0 mL flush of sterile saline was administered *via* the catheter after dose administration.

[00641] To mitigate potential infusion reactions, all animals were pretreated approximately 30 $\pm$ 5 minutes prior to start of infusion with diphenhydramine and dexamethasone. In addition, all animals received a second dose of diphenhydramine and dexamethasone approximately 4 hours $\pm$ 10 minutes post infusion. Diphenhydramine was administered as an intramuscular injection at a dose volume of 0.1 mL/kg to achieve a dose level of 5 mg/kg/dose. Dexamethasone was administered as an intramuscular injection at a dose volume of 0.25 mL/kg to achieve a dose level of 1 mg/kg/dose.

Table 19: Factor IX Sample Collection, Processing and Analysis

Group Nos.	Factor IX Blood Sample Collection		
	Pretreatment	Day 5	Day 14
Groups 1 – 4	X	X	X

X = sample to be collected

[00642] Samples were mixed gently and centrifuged as soon as practical. The resultant plasma was separated and split into two aliquots. All aliquots were made in uniquely labeled polypropylene tubes, and frozen immediately over dry ice or in a freezer set to maintain -70°C or colder until sample analysis.

[00643] Results

[00644] *Cynomolgus* monkeys treated with ceDNA-FIX v1 showed an elevated plasma concentration of human FIX (IU/mL) as compared to *Cynomolgus* monkeys treated only with vehicle

that showed no expression. No adverse event was observed, and the test article appeared to be well tolerated, suggesting that the ceDNA constructs disclosed herein can be used to increase plasma FIX protein to promote blood clotting in primates and potentially human patients.

REFERENCES

[00645] All publications and references, including but not limited to patents and patent applications, cited in this specification and Examples herein are incorporated by reference in their entirety as if each individual publication or reference were specifically and individually indicated to be incorporated by reference herein as being fully set forth. Any patent application to which this application claims priority is also incorporated by reference herein in the manner described above for publications and references.

CLAIMS

1. A capsid-free closed-ended DNA (ceDNA) vector comprising:
at least one nucleic acid sequence between flanking inverted terminal repeats (ITRs), wherein the at least one nucleic acid sequence encodes at least one FIX protein.
2. The ceDNA vector of claim 1, wherein the least one nucleic acid sequence that encodes at least one FIX protein is selected from any of the sequences set forth in Table 1 or any open reading frame (ORF) sequence included in any ceDNA sequence listed in Table 12.
3. The ceDNA vector of claims 1 or 2, wherein the ceDNA vector comprises a promoter sequence selected from any of those in Table 7 operatively linked to the least one nucleic acid sequence that encodes at least one FIX protein.
4. The ceDNA vector of any of claims 1 to 3, wherein the ceDNA vector comprises an enhancer sequence selected from any of those in Table 8.
5. The ceDNA vector of any of claims 1 to 4, wherein the ceDNA vector comprises a 5' UTR and/or intron sequence selected from any of those in Table 9A.
6. The ceDNA vector of any of claims 1 to 5, wherein the ceDNA vector comprises a 3' UTR sequence selected from any of those in Table 9B.
7. The ceDNA vector of any of claims 1 to 6, wherein the ceDNA vector comprises at least one poly A sequence selected from any of those in Table 10.
8. The ceDNA vector of any one of claims 1-7, wherein the ceDNA vector comprises at least one promoter sequence operably linked to at least one nucleic acid sequence.
9. The ceDNA vector of any one of claims 1-8, wherein the at least one nucleic acid sequence is cDNA.
10. The ceDNA vector of any one of claims 1-9, wherein at least one ITR comprises a functional terminal resolution site and a Rep binding site.
11. The ceDNA vector of any one of claims 1-10, wherein one or both of the ITRs are from a virus selected from a *Parvovirus*, a *Dependovirus*, and an adeno-associated virus (AAV).
12. The ceDNA vector of any one of claims 1-11, wherein the flanking ITRs are symmetric or asymmetric with respect to one another.
13. The ceDNA vector of claim 12, wherein the flanking ITRs are symmetrical or substantially symmetrical.
14. The ceDNA vector of claim 12, wherein the flanking ITRs are asymmetric.
15. The ceDNA vector of any one of claims 1-14, wherein one or both of the ITRs are wild type, or wherein both of the ITRs are wild-type ITRs.
16. The ceDNA vector of any one of claims 1-15, wherein the flanking ITRs are from different viral serotypes.

17. The ceDNA vector of any one of claims 1-16, wherein the flanking ITRs are selected from any pair of viral serotypes shown in Table 2.
18. The ceDNA vector of any one of claims 1-17, wherein one or both of the ITRs comprises a sequence selected from one or more of the sequences in Table 3.
19. The ceDNA vector of any one of claims 1-18, wherein at least one of the ITRs is altered from a wild-type AAV ITR sequence by a deletion, addition, or substitution that affects the overall three-dimensional conformation of the ITR.
20. The ceDNA vector of any one of claims 1-19, wherein one or both of the ITRs are derived from an AAV serotype selected from AAV1, AAV2, AAV3, AAV4, AAV5, AAV6, AAV7, AAV8, AAV9, AAV10, AAV11, and AAV12.
21. The ceDNA vector of any one of claims 1-20, wherein one or both of the ITRs are synthetic.
22. The ceDNA vector of any one of claims 1-21, wherein one or both of the ITRs are not a wild type ITR, or wherein both of the ITRs are not wild-type ITRs.
23. The ceDNA vector of any one of claims 1-22, wherein one or both of the ITRs are modified by a deletion, insertion, and/or substitution in at least one of the ITR regions selected from A, A', B, B', C, C', D, and D'.
24. The ceDNA vector of claim 23, wherein the deletion, insertion, and/or substitution results in the deletion of all or part of a stem-loop structure normally formed by the A, A', B, B', C, or C' regions.
25. The ceDNA vector of any one of claims 1-24, wherein one or both of the ITRs are modified by a deletion, insertion, and/or substitution that results in the deletion of all or part of a stem-loop structure normally formed by the B and B' regions.
26. The ceDNA vector of any one of claims 1-24, wherein one or both of the ITRs are modified by a deletion, insertion, and/or substitution that results in the deletion of all or part of a stem-loop structure normally formed by the C and C' regions.
27. The ceDNA vector of any one of claims 1-24, wherein one or both of the ITRs are modified by a deletion, insertion, and/or substitution that results in the deletion of part of a stem-loop structure normally formed by the B and B' regions and/or part of a stem-loop structure normally formed by the C and C' regions.
28. The ceDNA vector of any one of claims 1-27, wherein one or both of the ITRs comprise a single stem-loop structure in the region that normally comprises a first stem-loop structure formed by the B and B' regions and a second stem-loop structure formed by the C and C' regions.
29. The ceDNA vector of any one of claims 1-28, wherein one or both of the ITRs comprise a single stem and two loops in the region that normally comprises a first stem-loop structure formed by the B and B' regions and a second stem-loop structure formed by the C and C' regions.
30. The ceDNA vector of any one of claims 1-29, wherein one or both of the ITRs comprise a single stem and a single loop in the region that normally comprises a first stem-loop structure formed

by the B and B' regions and a second stem-loop structure formed by the C and C' regions.

31. The ceDNA vector of any one of claims 1-30, wherein both ITRs are altered in a manner that results in an overall three-dimensional symmetry when the ITRs are inverted relative to each other.
32. The ceDNA vector of any one of claims 1-31, wherein one or both of the ITRs comprises a nucleic acid sequence selected from the sequences set forth in Tables 3, 5A, 5B, and 6.
33. The ceDNA vector of any one of claims 1-32, wherein at least one nucleic acid sequence is under the control of at least one regulatory switch.
34. The ceDNA vector of claim 33, wherein the at least one regulatory switch is selected from the group consisting of: a binary regulatory switch, a small molecule regulatory switch, a passcode regulatory switch, a nucleic acid-based regulatory switch, a post-transcriptional regulatory switch, a radiation-controlled or ultrasound controlled regulatory switch, a hypoxia-mediated regulatory switch, an inflammatory response regulatory switch, a shear-activated regulatory switch, and a kill switch.
35. A capsid-free close-ended DNA (ceDNA) vector comprising a nucleic acid sequence selected from Table 12.
36. A capsid-free close-ended DNA (ceDNA) vector comprising a nucleic acid sequence at least 85% identical to SEQ ID NO: 404, SEQ ID NO: 405 or SEQ ID NO: 406.
37. A capsid-free close-ended DNA (ceDNA) vector consisting of a nucleic acid sequence selected from the group consisting of SEQ ID NO: 404, SEQ ID NO: 405 and SEQ ID NO: 406.
38. A method of expressing an FIX protein in a cell comprising contacting the cell with the ceDNA vector of any one of claims 1-37.
39. The method of claim 38, wherein the cell is a hepatocyte.
40. The method of claim 38 or 39, wherein the cell is *in vitro* or *in vivo*.
41. The method of any one of claims 38-40, wherein the at least one nucleic acid sequence is codon optimized for expression in the eukaryotic cell.
42. The method of any one of claims 38-41, wherein the at least one nucleic acid sequence that is codon optimized is selected from any one of the sequences set forth in Table 1 or any open reading frame (ORF) sequence included in any ceDNA sequence listed in Table 12.
43. A method of treating a subject with hemophilia B, comprising administering to the subject a ceDNA vector of any one of claims 1-37, wherein at least one nucleic acid sequence encodes at least one FIX protein.
44. The method of claim 43, wherein the least one nucleic acid sequence that encodes the at least one FIX protein is selected from any one of the sequences set forth in Table 1.
45. The method of claim 43 or 44, wherein the ceDNA vector is administered to a hepatocyte.
46. The method of any of claims 44 to 45, wherein the ceDNA vector expresses the FIX protein in a hepatocyte.
47. The method of any of claims 44-46, wherein the ceDNA vector is administered by intravenous or intramuscular injection.

48. The method of any one of claims 44-47, further comprising administering to the subject an immune modulating agent.
49. The method of claim 48, wherein the immune modulating agent is an immunosuppressant.
50. The method of claim 49, wherein the immunosuppressant is a tyrosine kinase inhibitor (TKI).
51. The method of claim 50, wherein the TKI is administered to the subject at a dosage of about 0.5 mg/kg to about 700 mg/kg.
52. A pharmaceutical composition comprising the ceDNA vector of any one of claims 1-37.
53. The pharmaceutical composition of claim 52, further comprising an additional compound.
54. The pharmaceutical composition of claim 52, wherein the additional compound is an immune modulating agent.
55. The pharmaceutical composition of claim 54, wherein the immune modulating agent is an immunosuppressant.
56. The pharmaceutical composition of claim 55, wherein the immunosuppressant is a tyrosine kinase inhibitor (TKI).
57. The pharmaceutical composition of claim 56, wherein the composition further comprises an excipient or carrier.
58. A cell containing a ceDNA vector of any of claims 1-37.
59. The cell of claim 58, wherein the cell is a hepatocyte.
60. A composition comprising a ceDNA vector of any of claims 1-37 and a lipid.
61. The composition of claim 60, wherein the lipid is a lipid nanoparticle (LNP).
62. A kit comprising the ceDNA vector of any one of claims 1-37 or the composition of any one of claims 52-57, or the cell of claim 58.

FIG. 1A

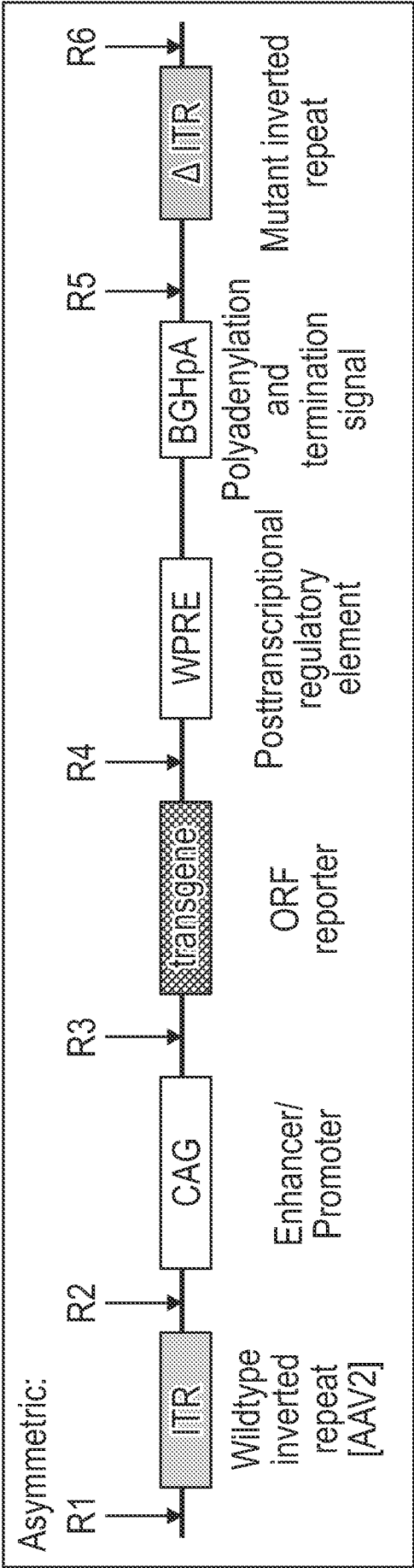


FIG. 1B

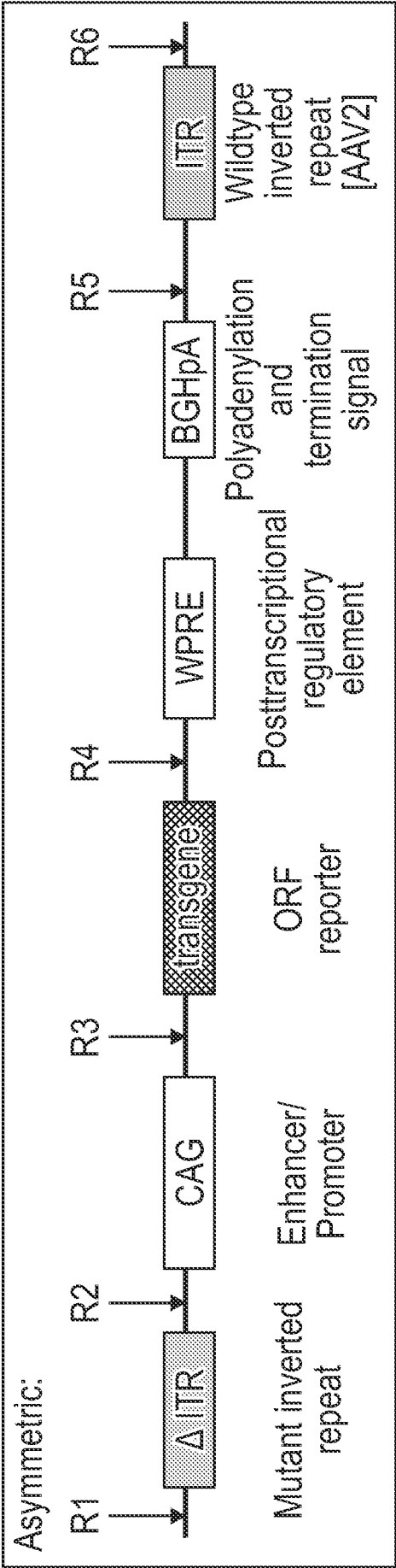


FIG. 1C

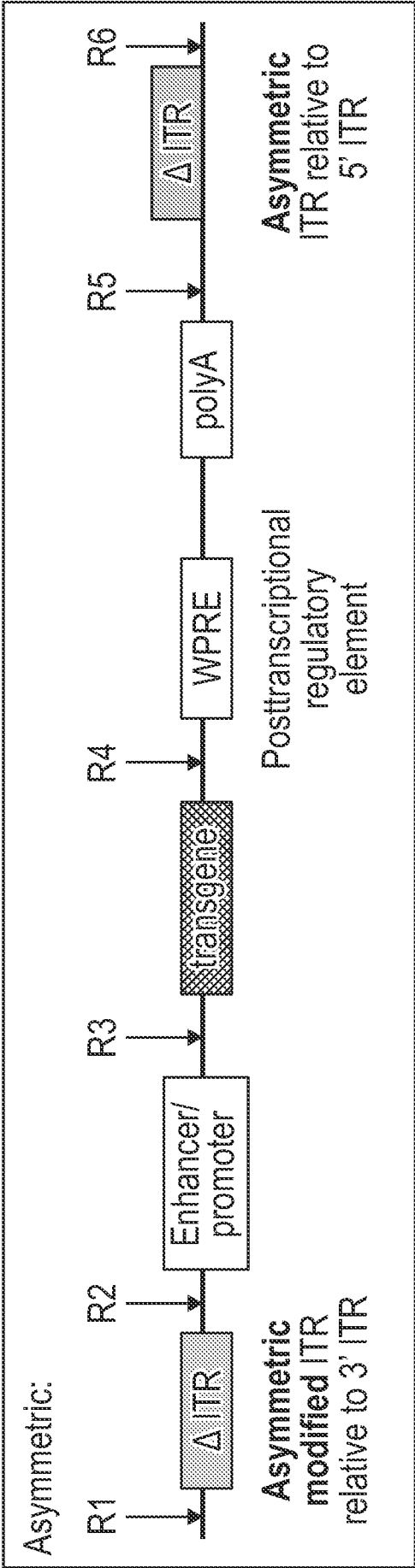


FIG. 1D

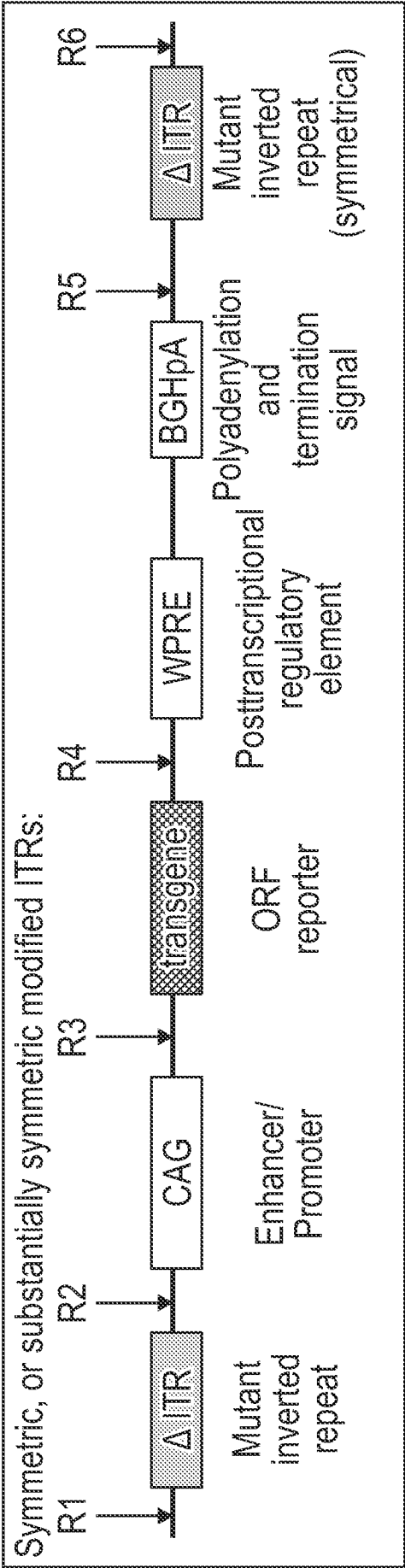


FIG. 1E

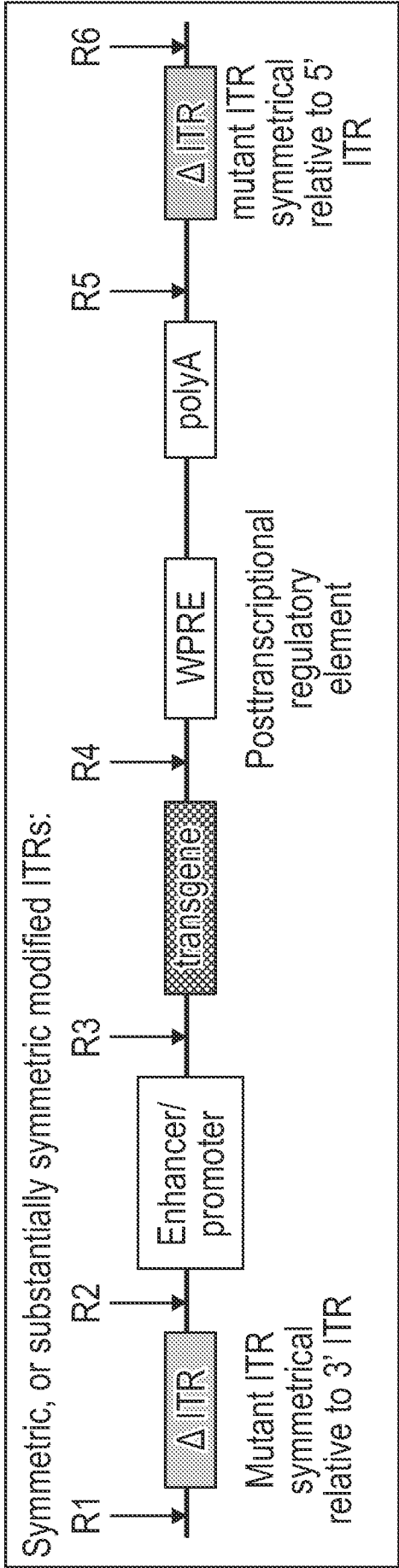


FIG. 1F

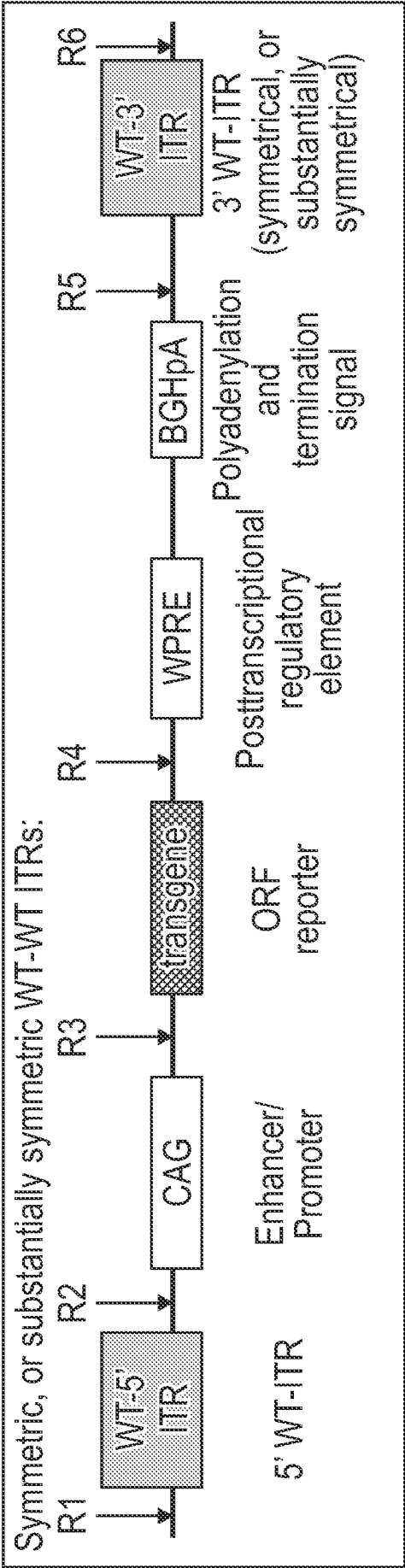
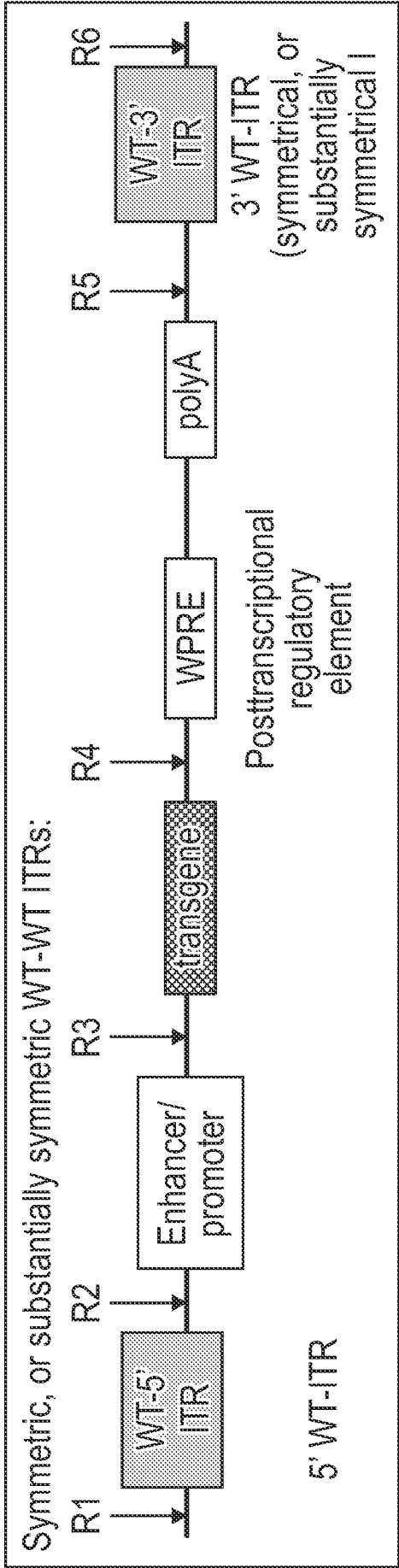


FIG. 1G



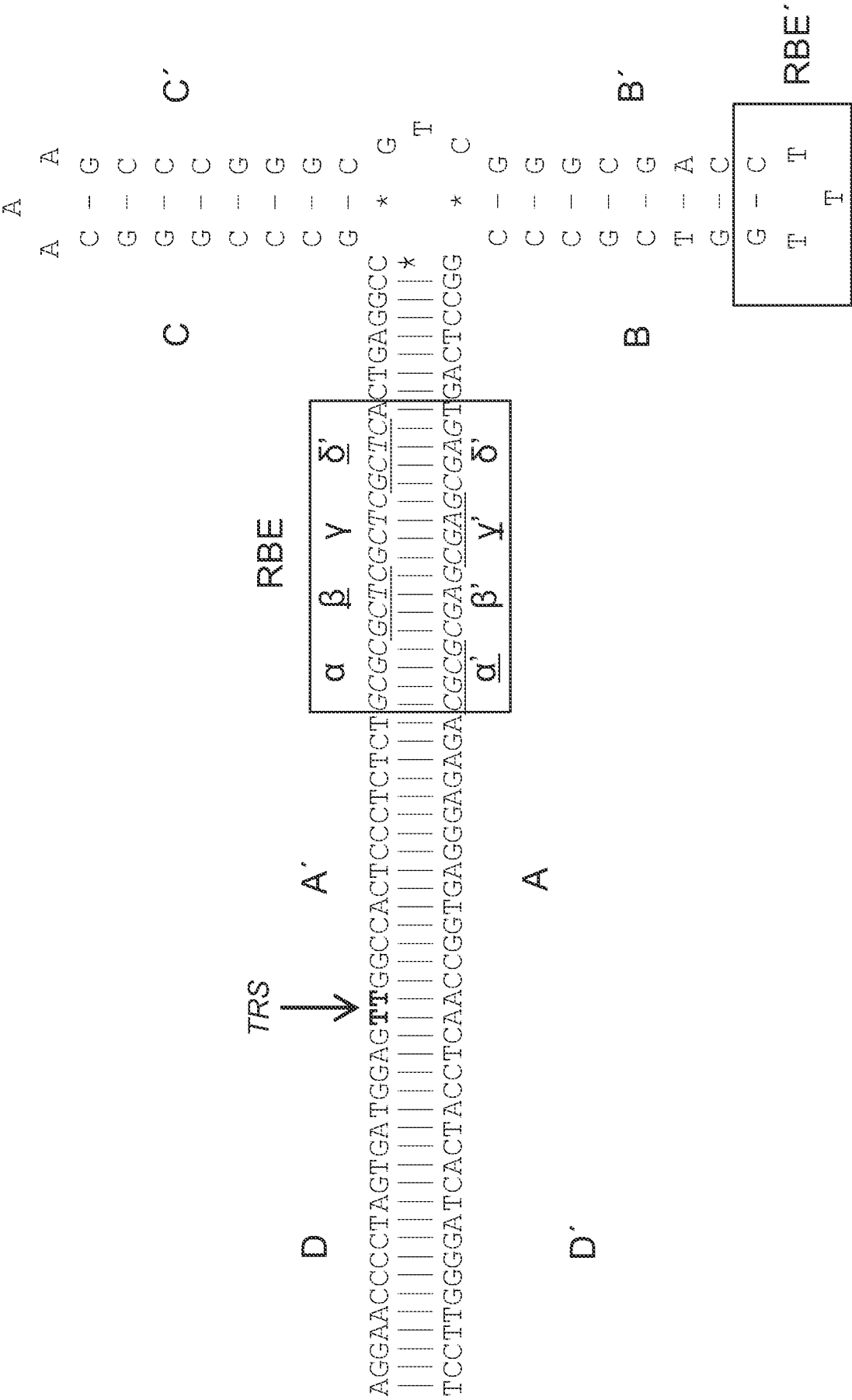


FIG. 2A

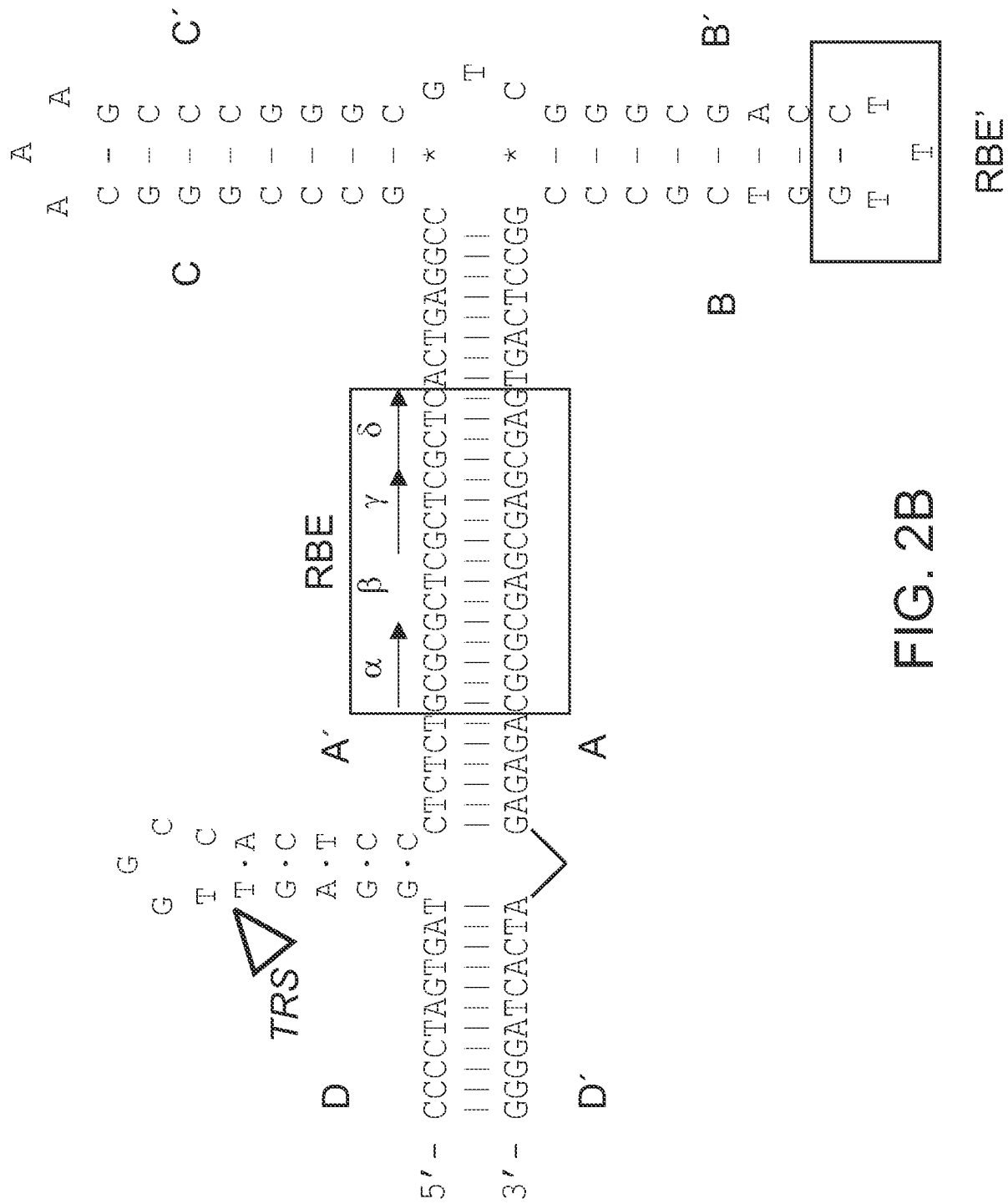
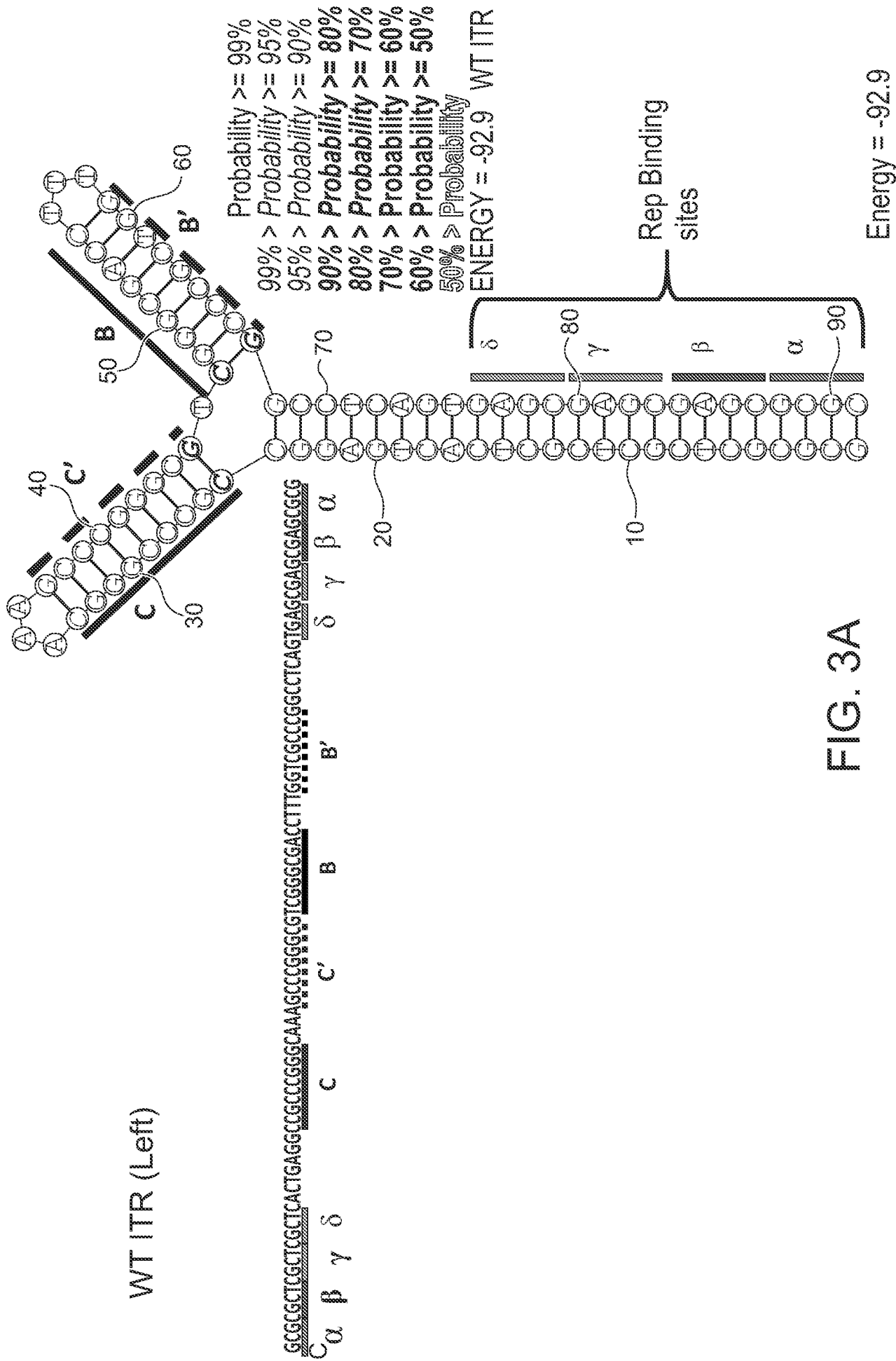
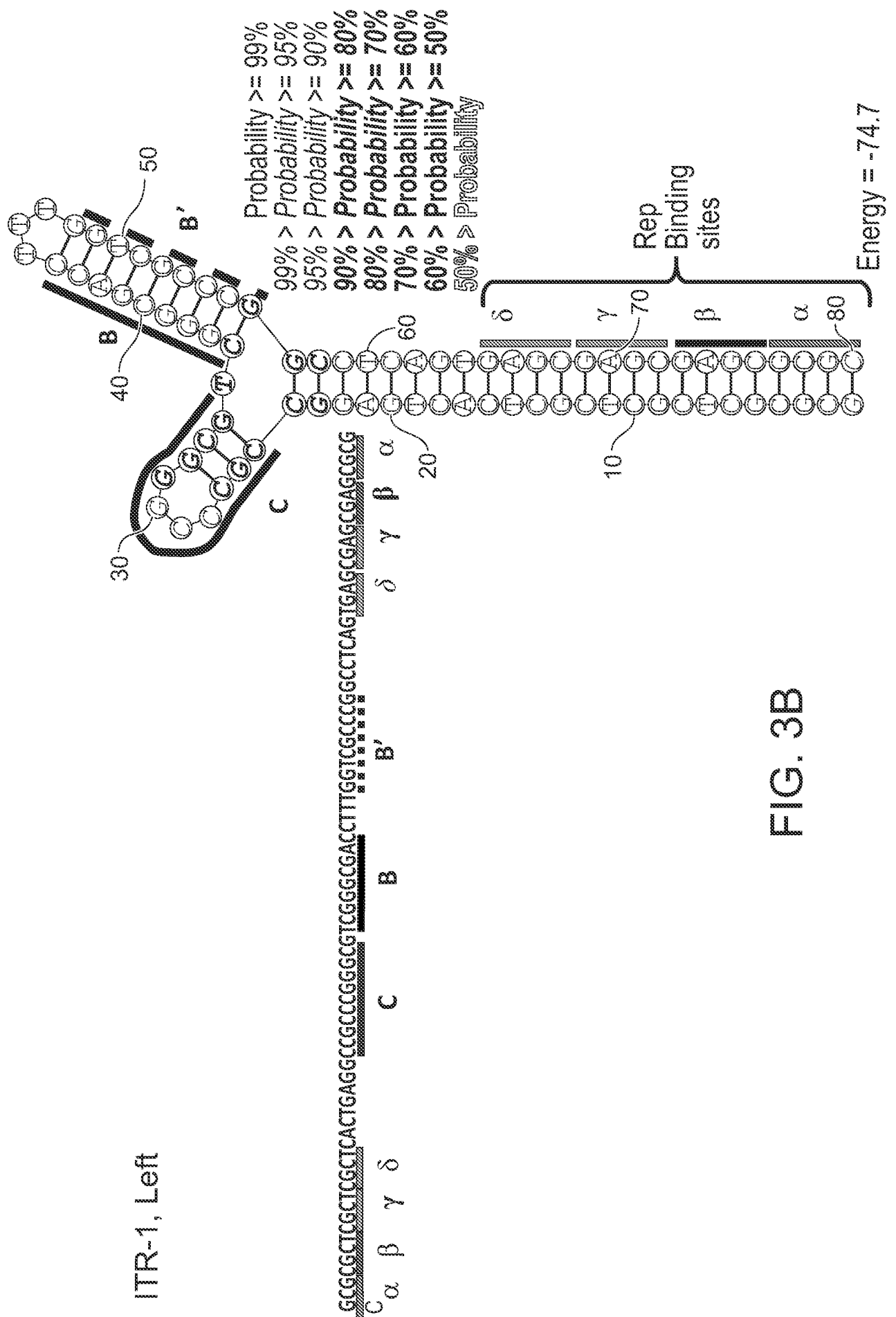


FIG. 2B





336

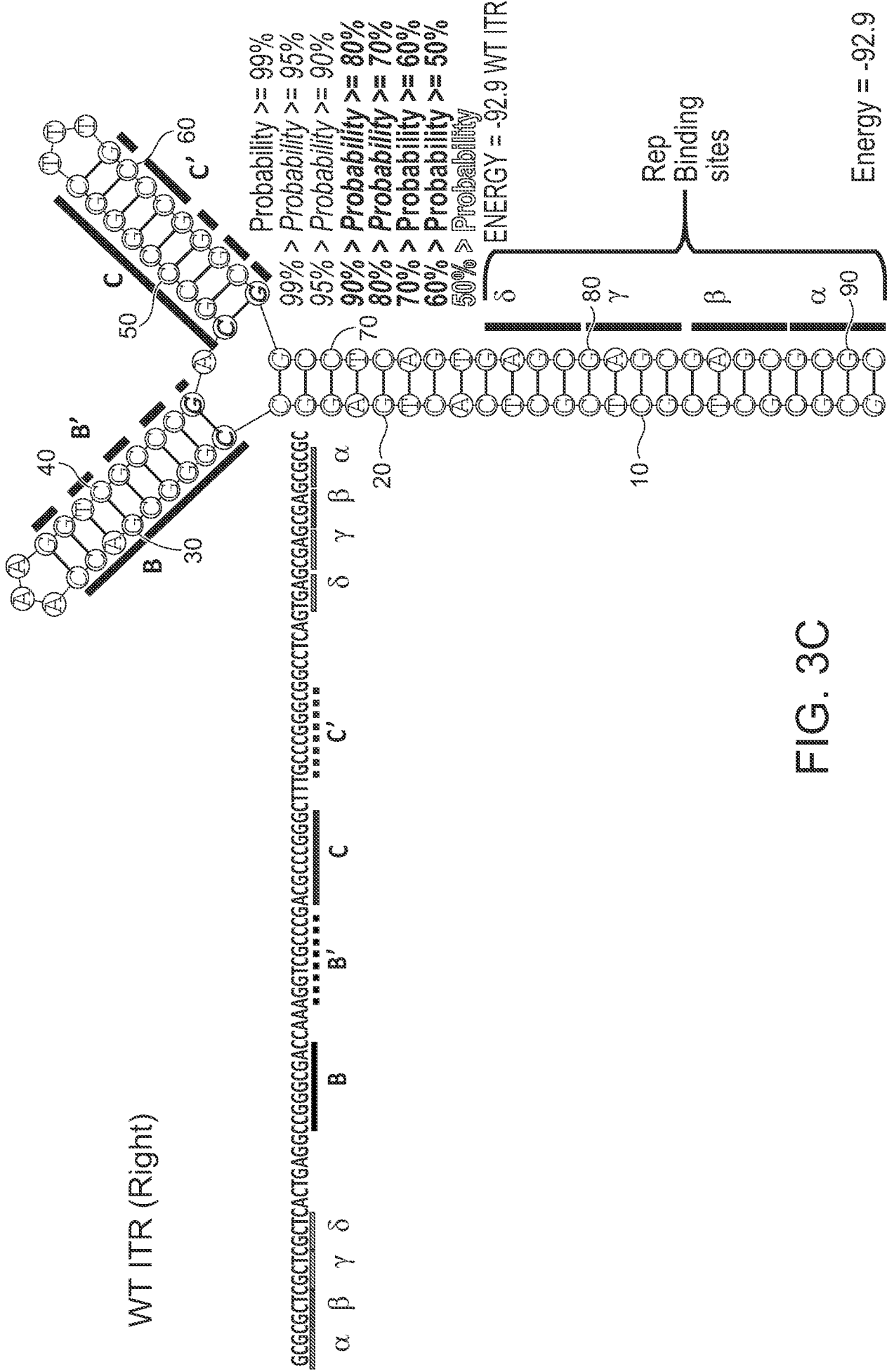


FIG. 3C

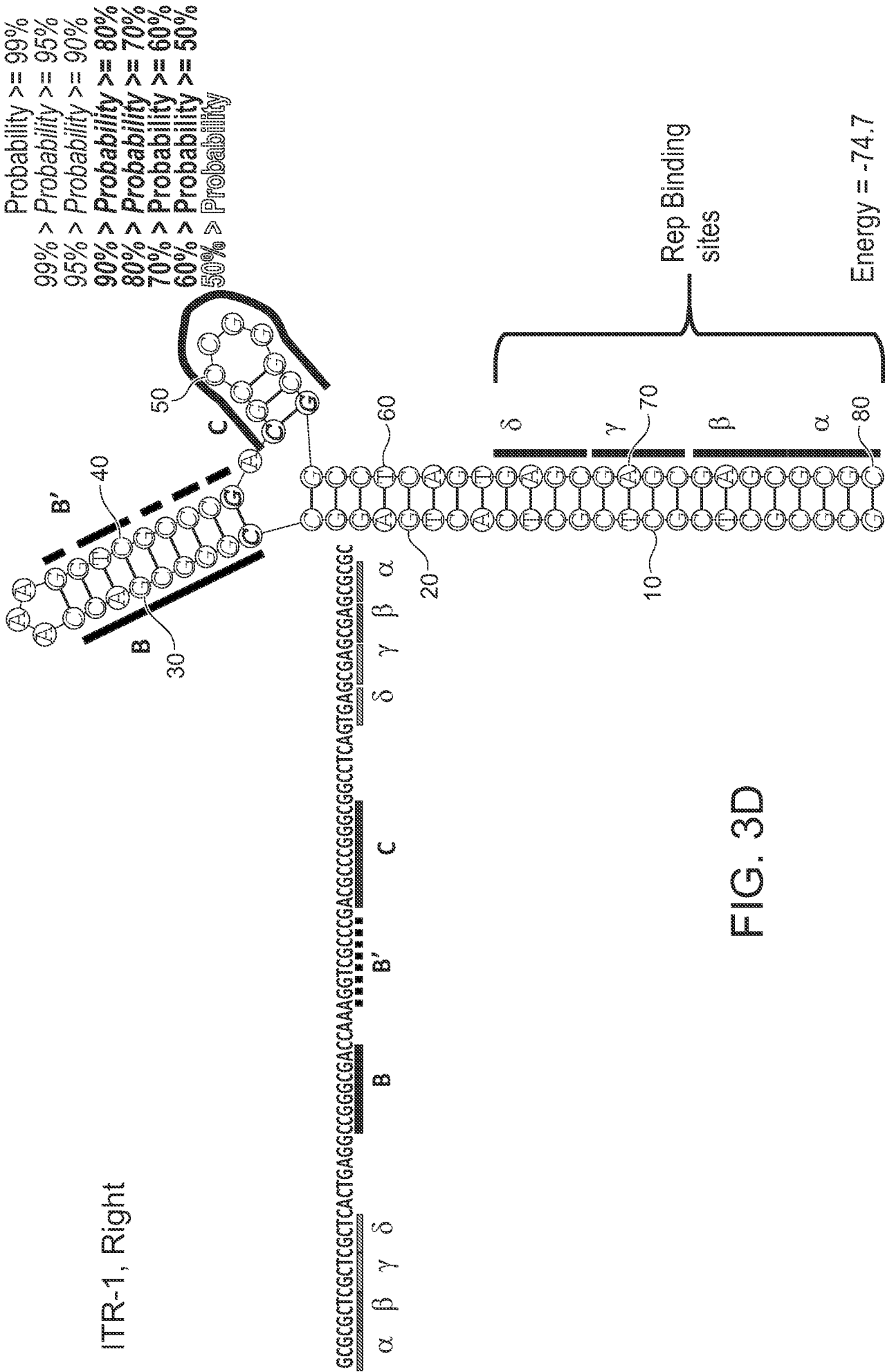


FIG. 3D

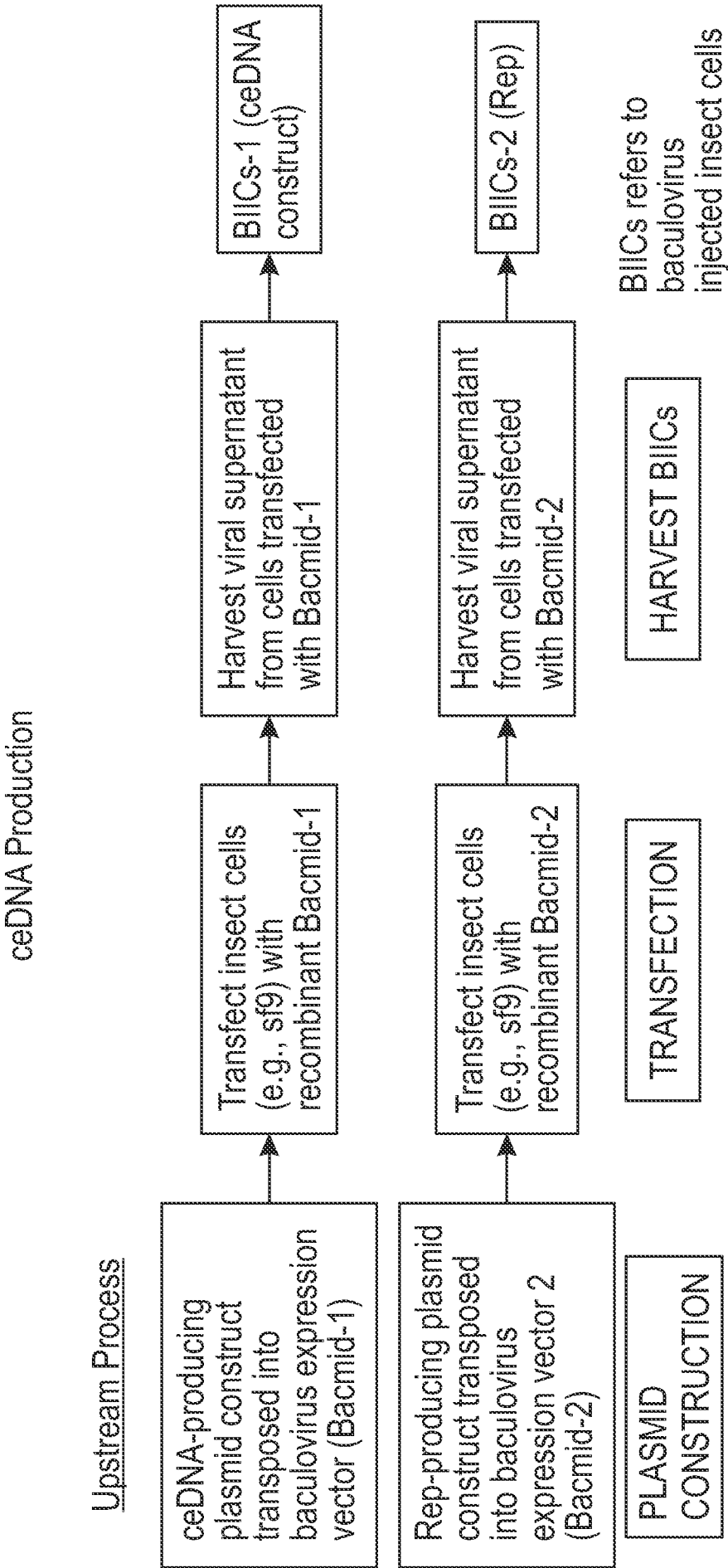


FIG. 4A

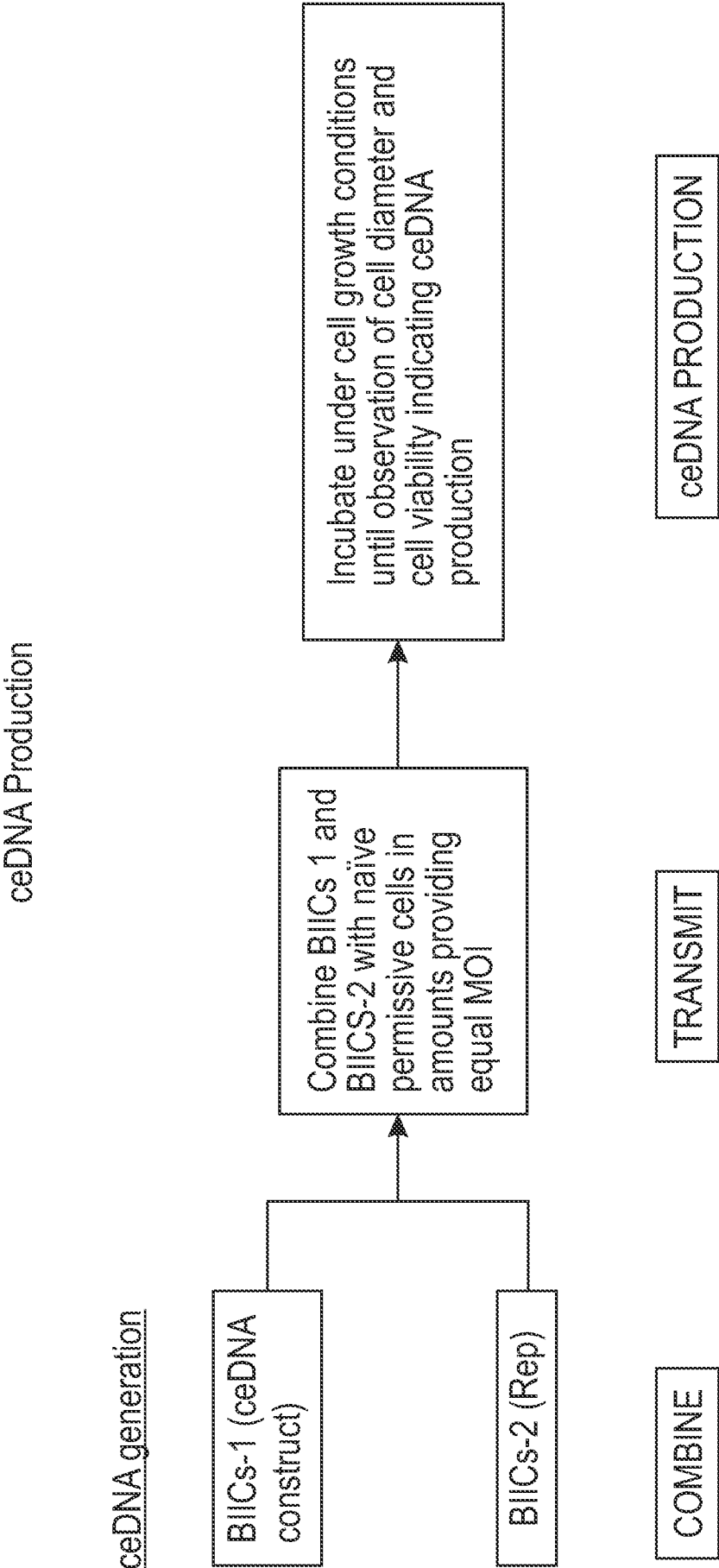


FIG. 4B

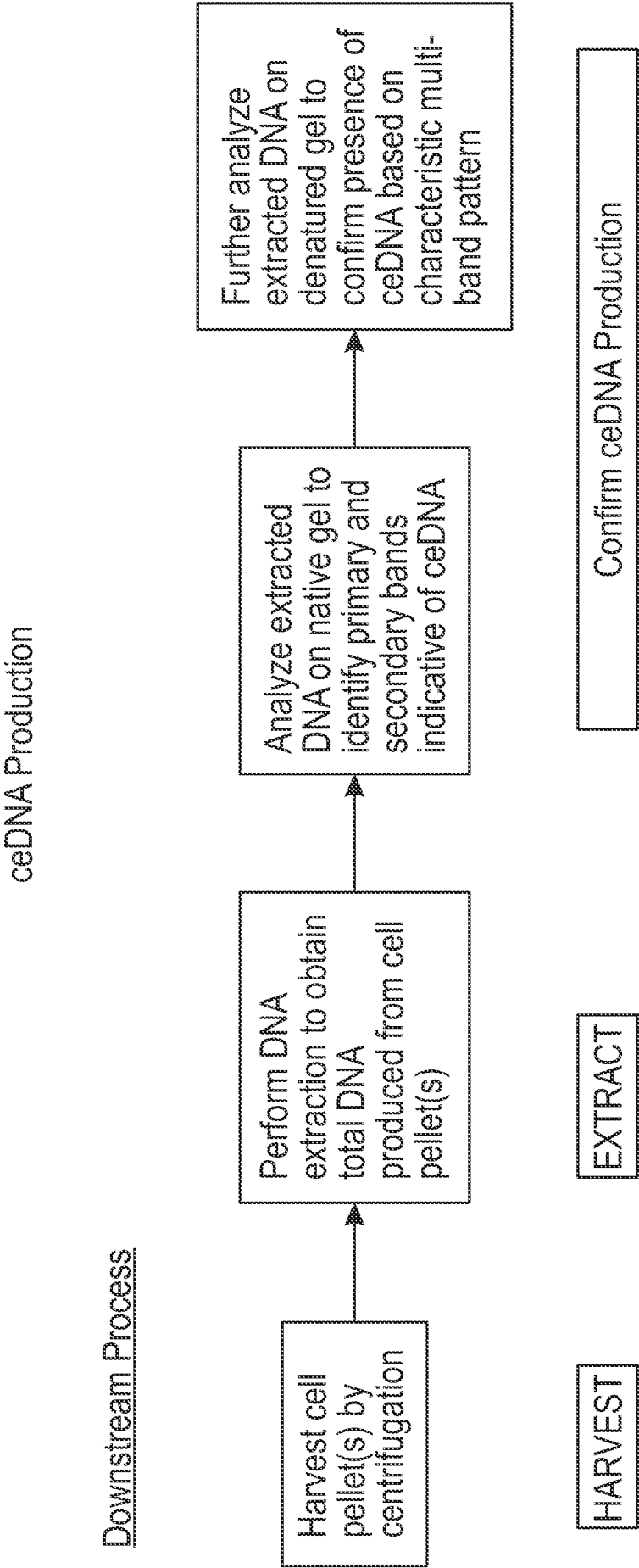


FIG. 4C

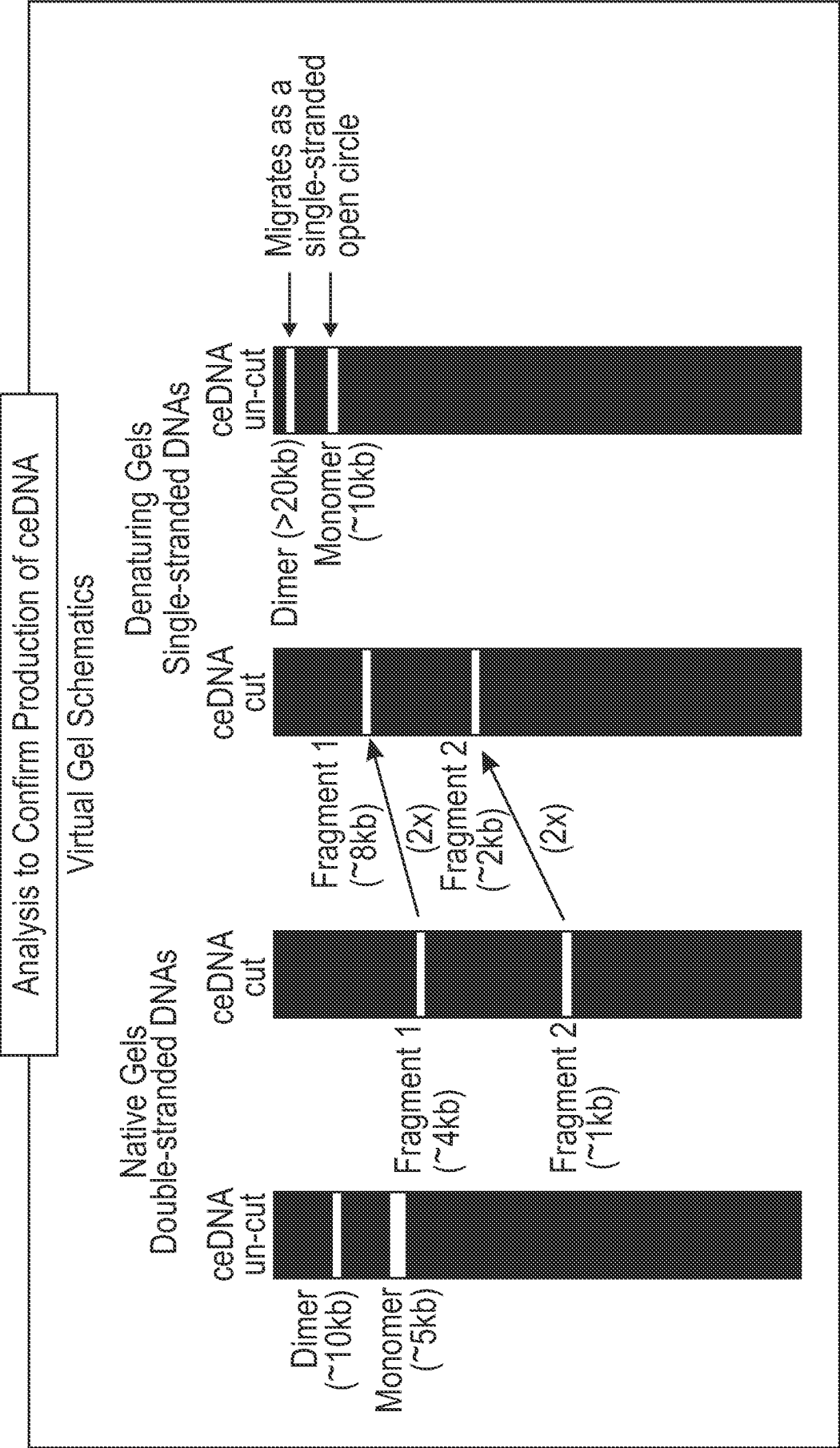
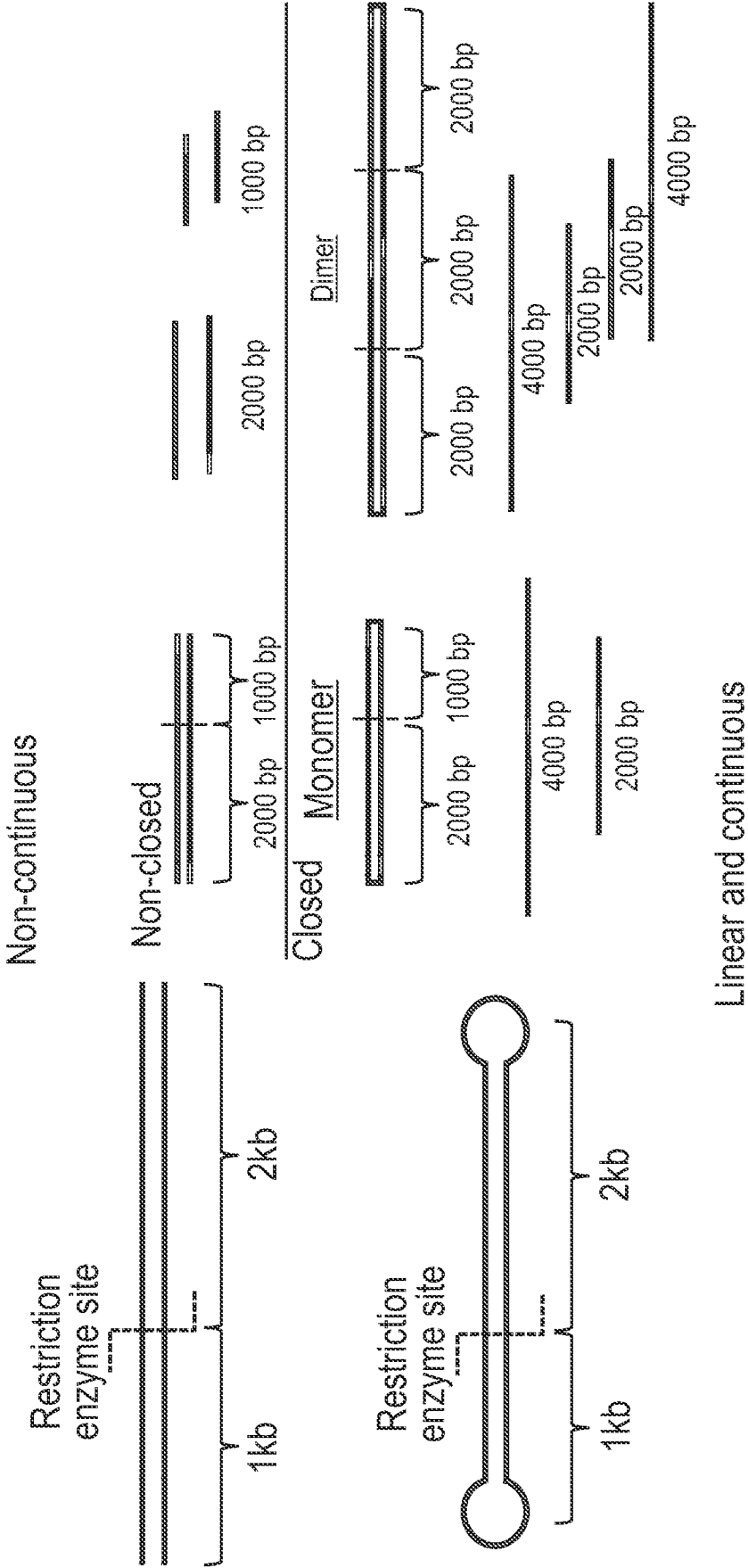


FIG. 4D

FIG. 4E



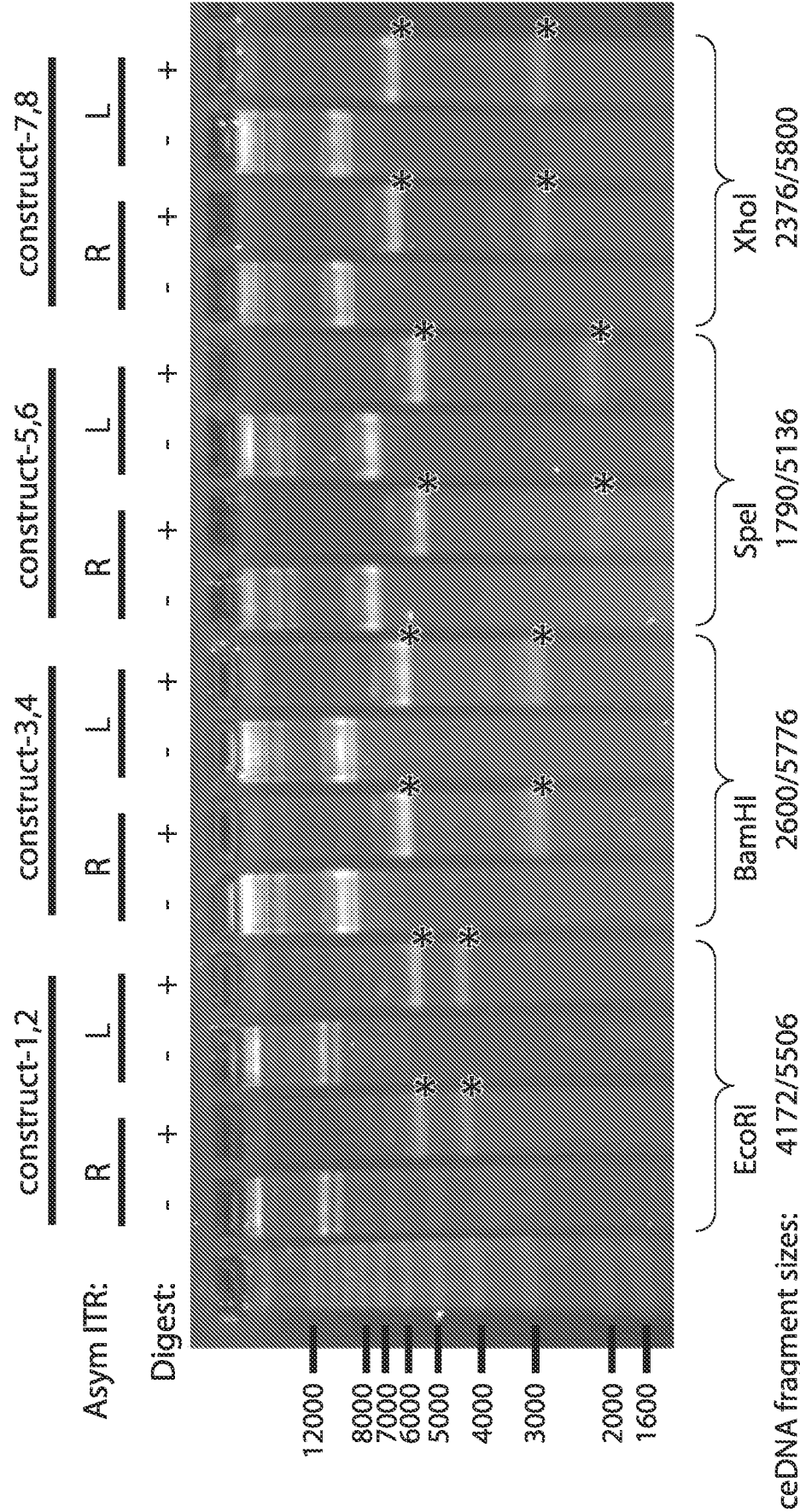
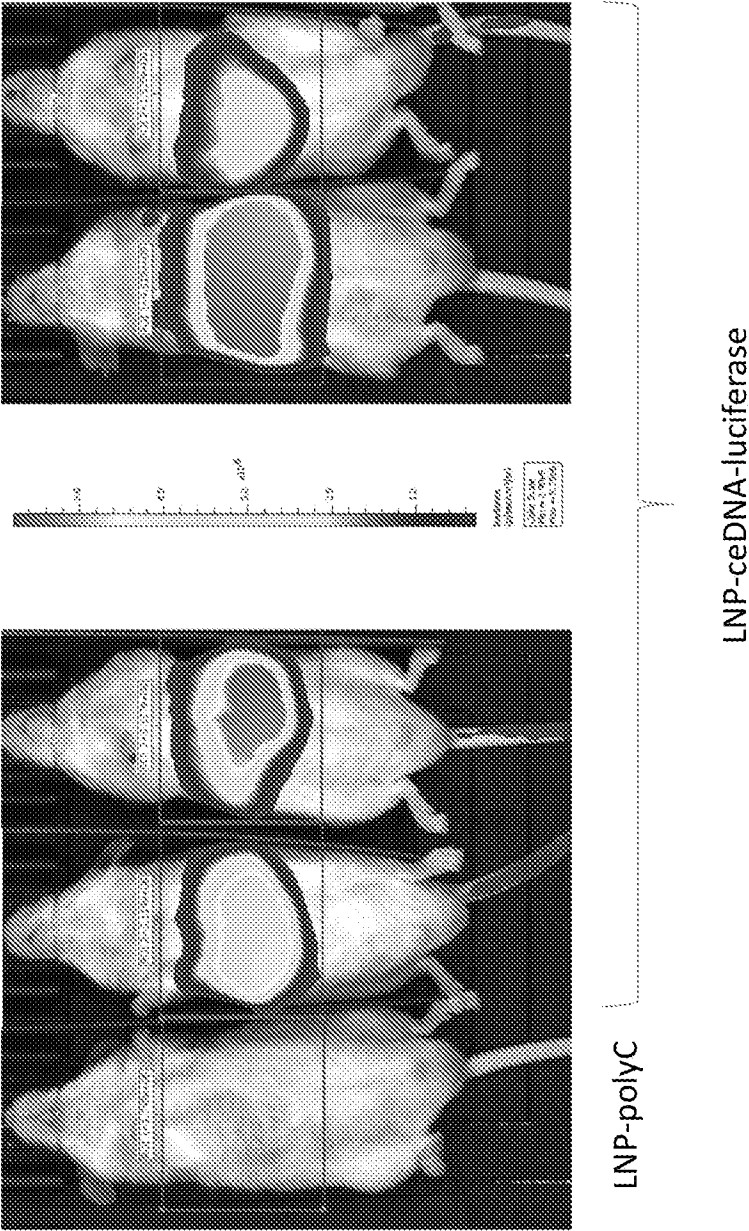


FIG. 5

FIG. 6



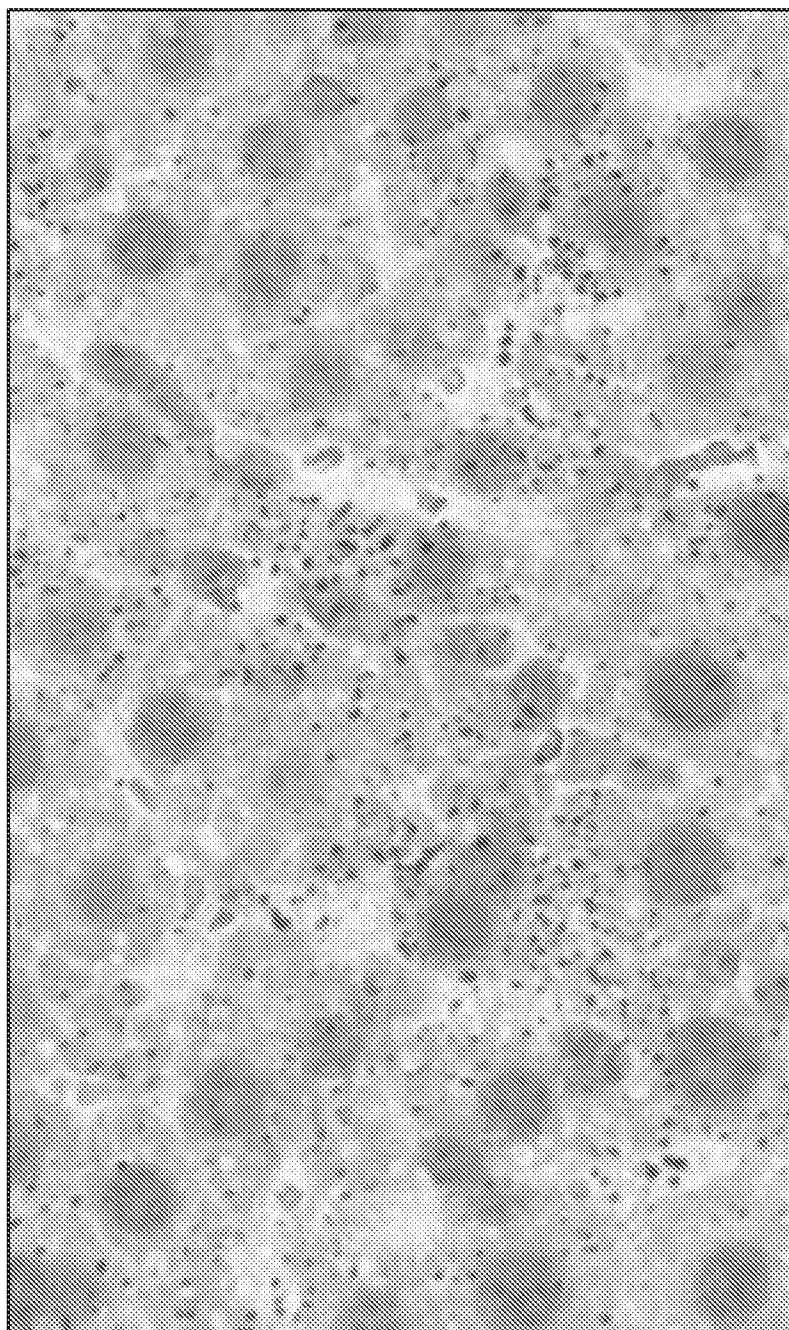


FIG. 7

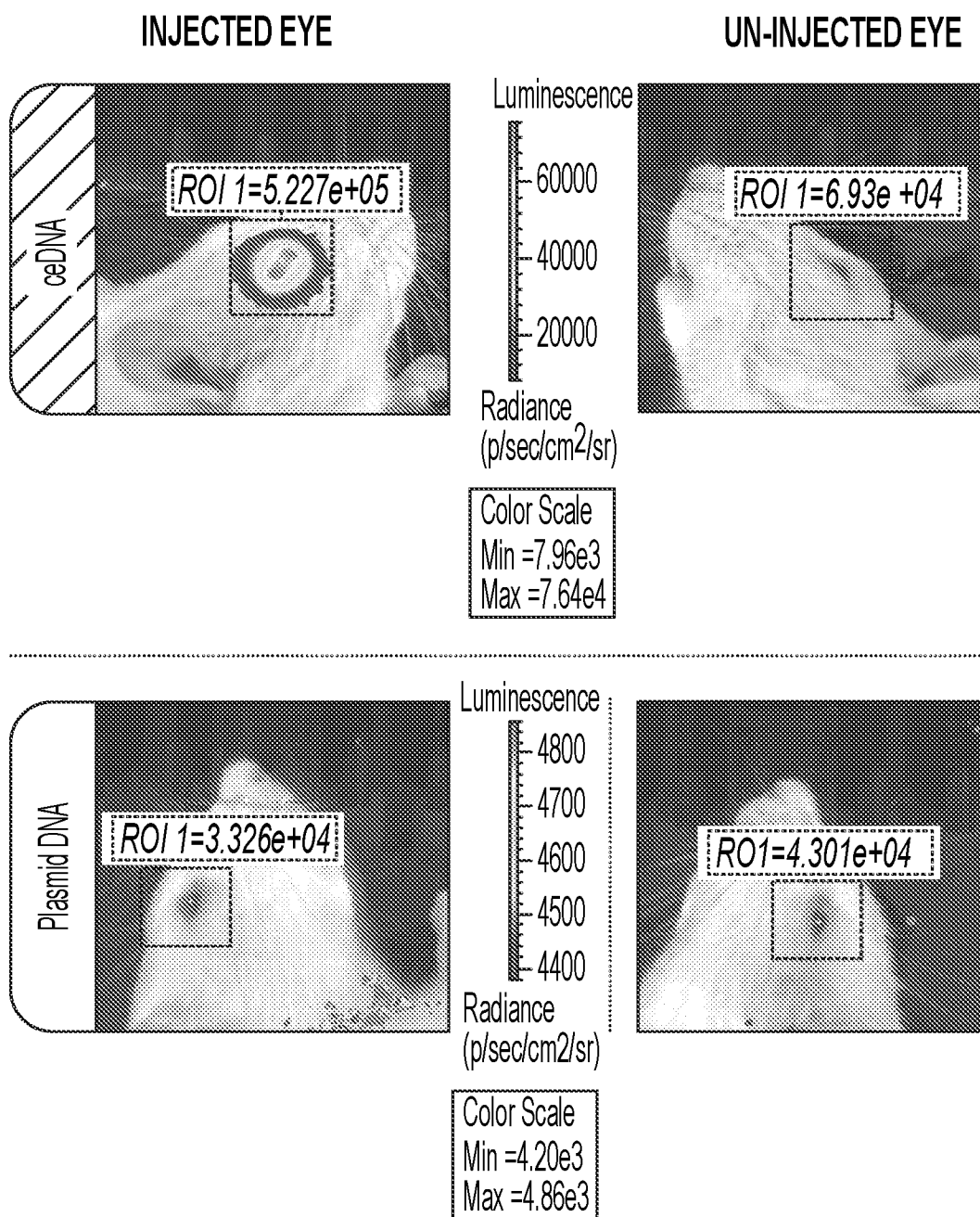


FIG. 8A

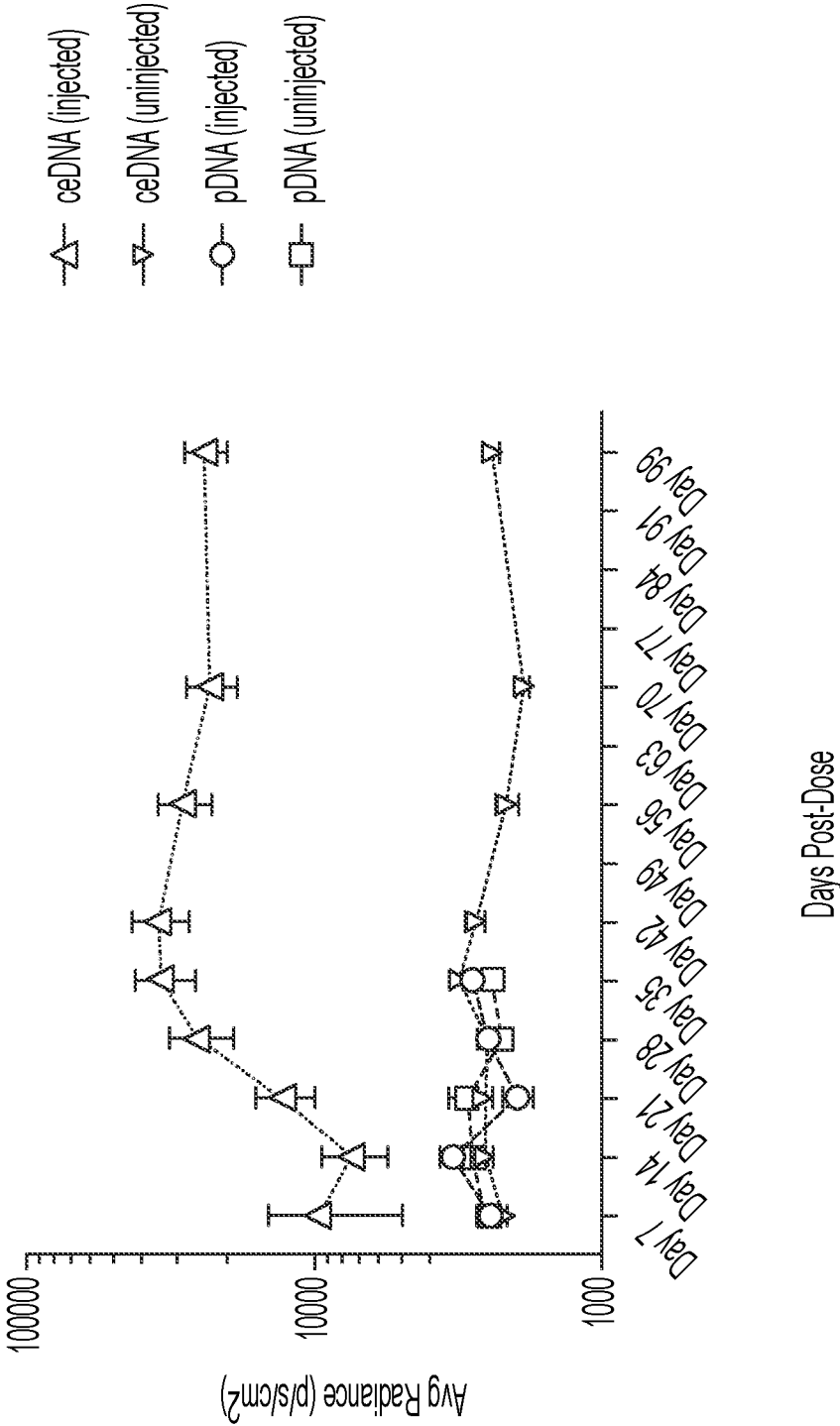


FIG. 8B

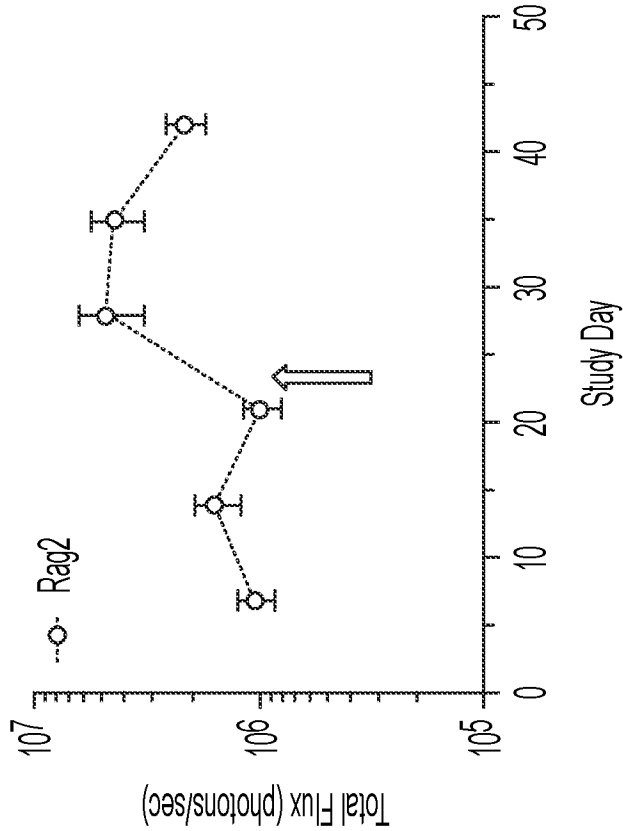


FIG. 9B

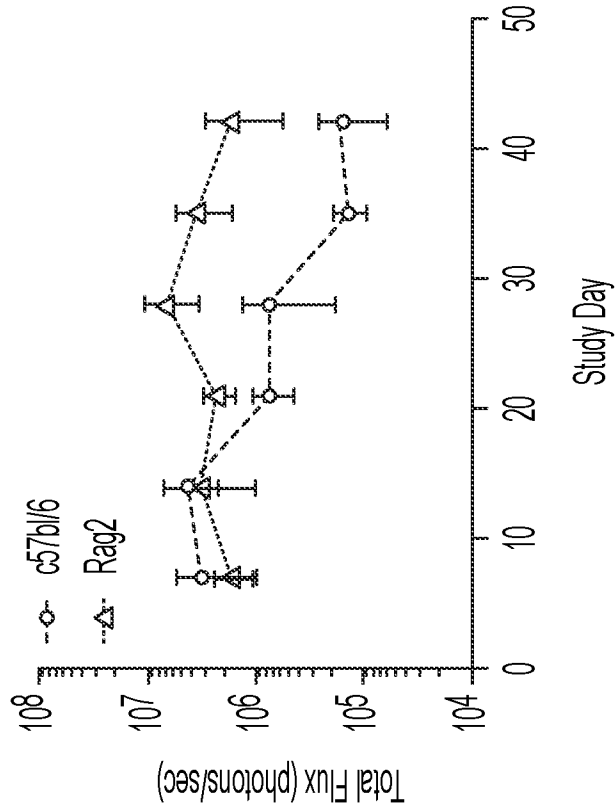


FIG. 9A

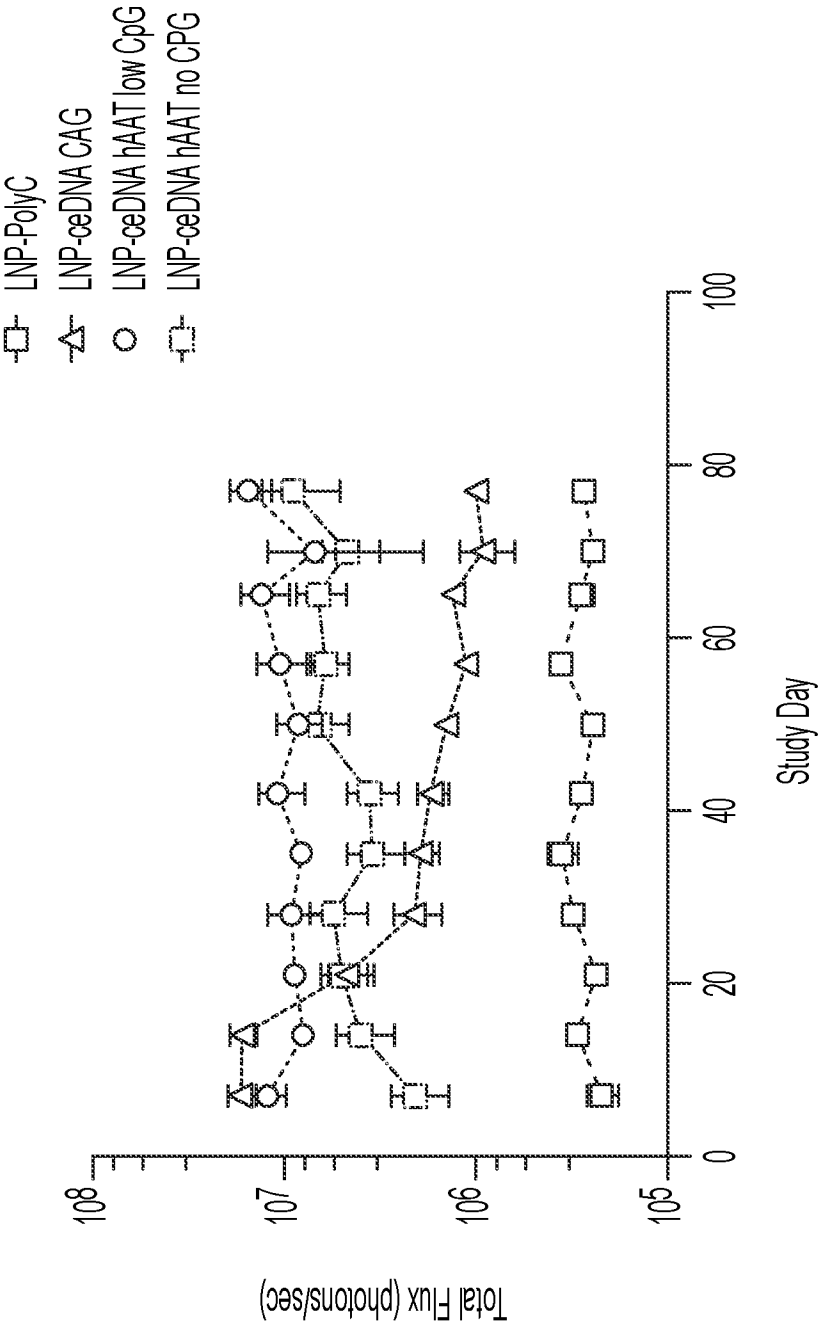


FIG. 10

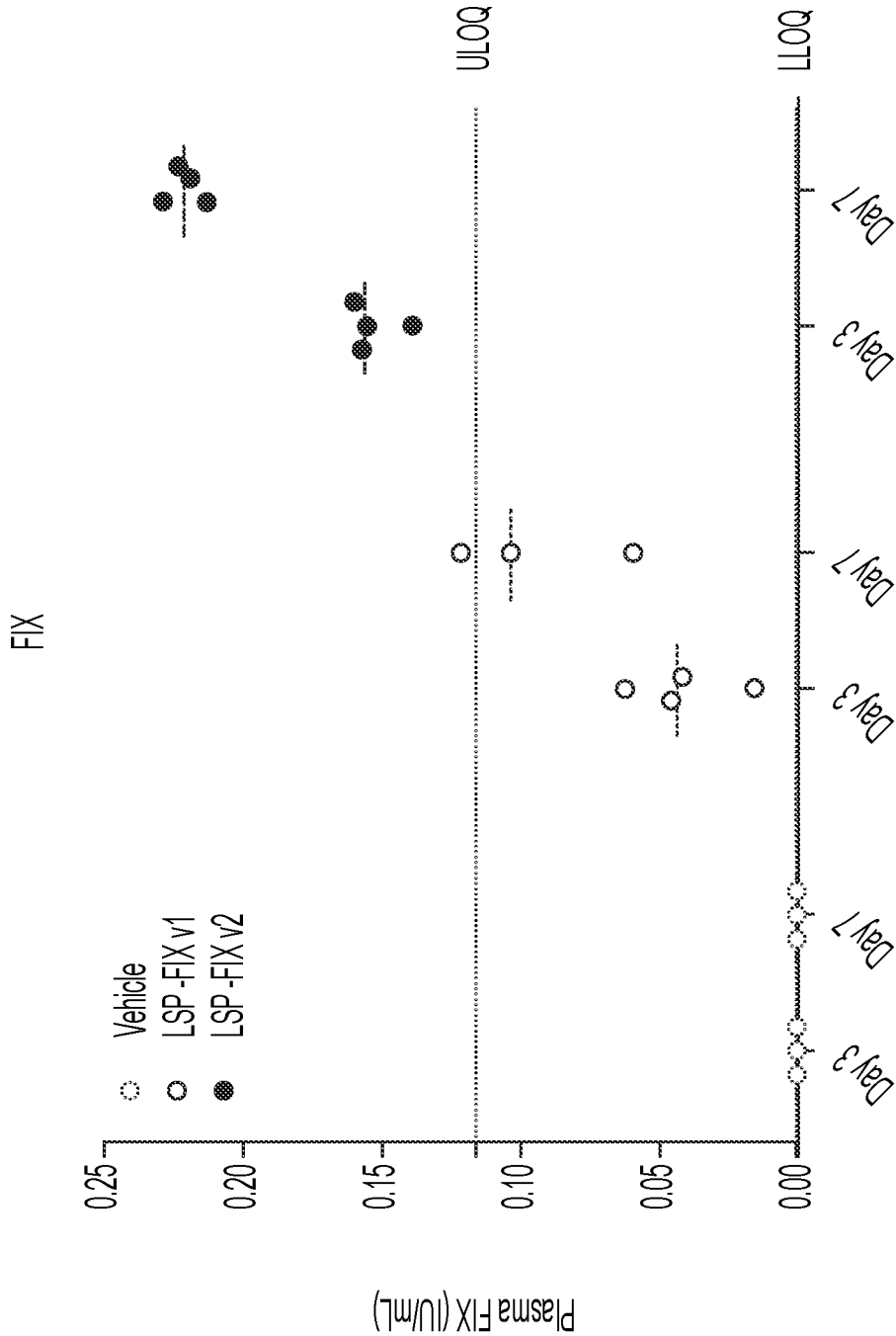


FIG. 11A

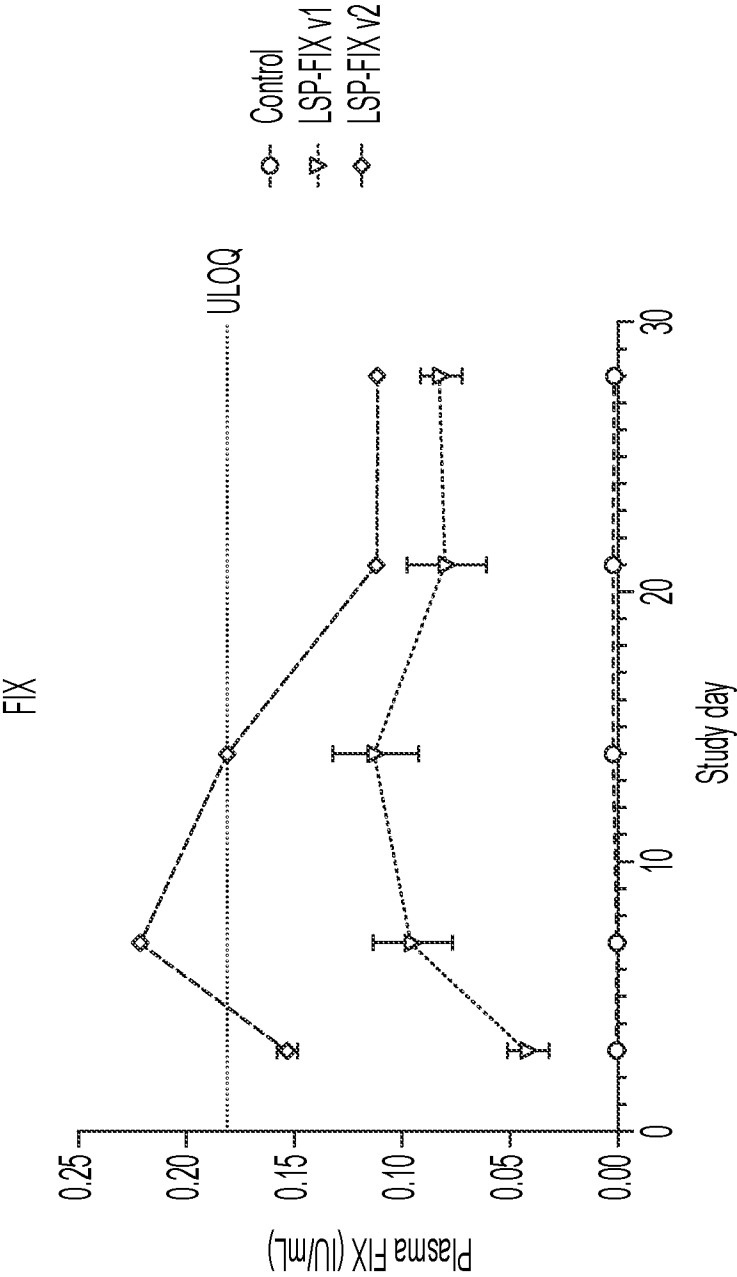


FIG. 11B

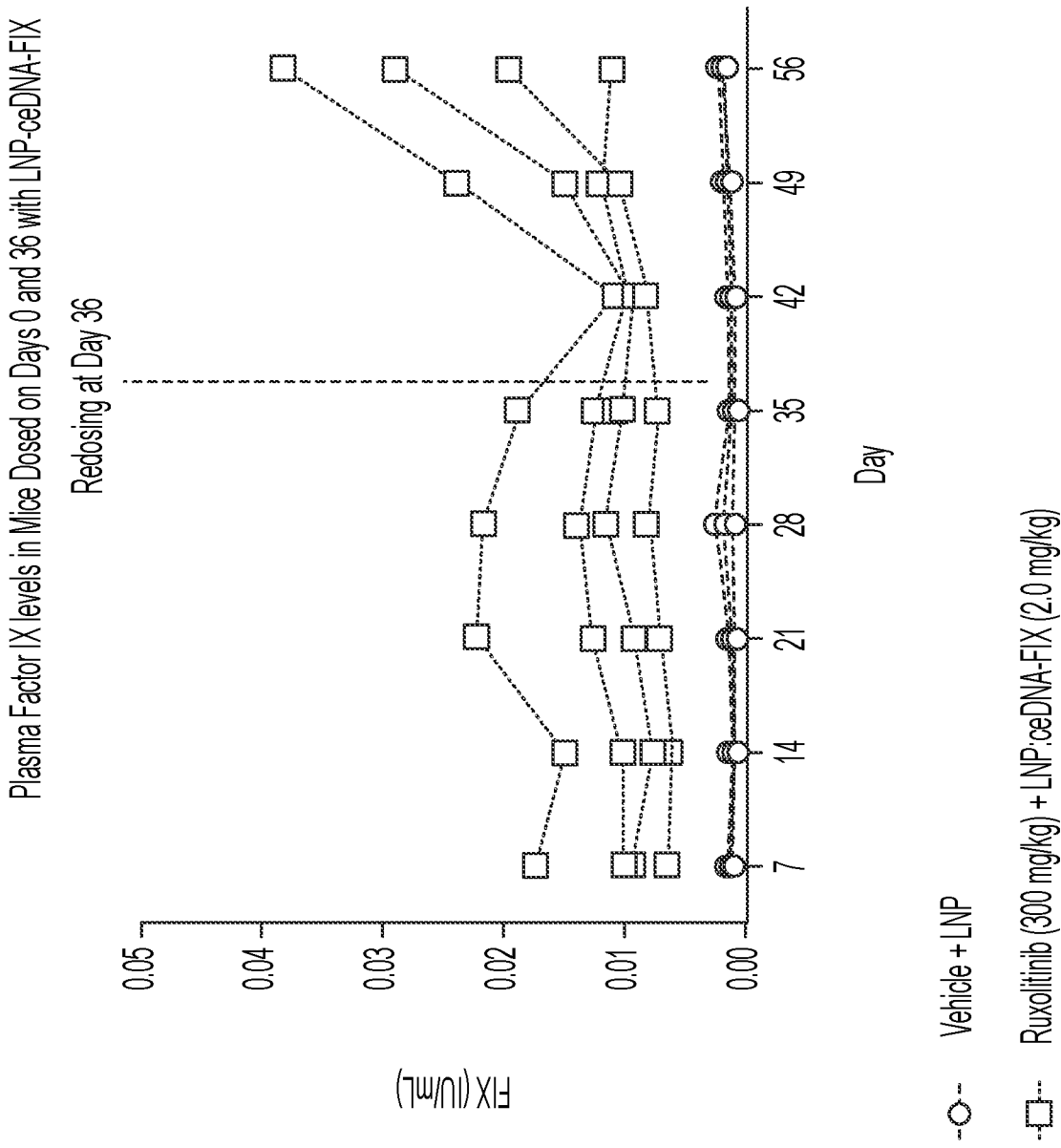


FIG. 12

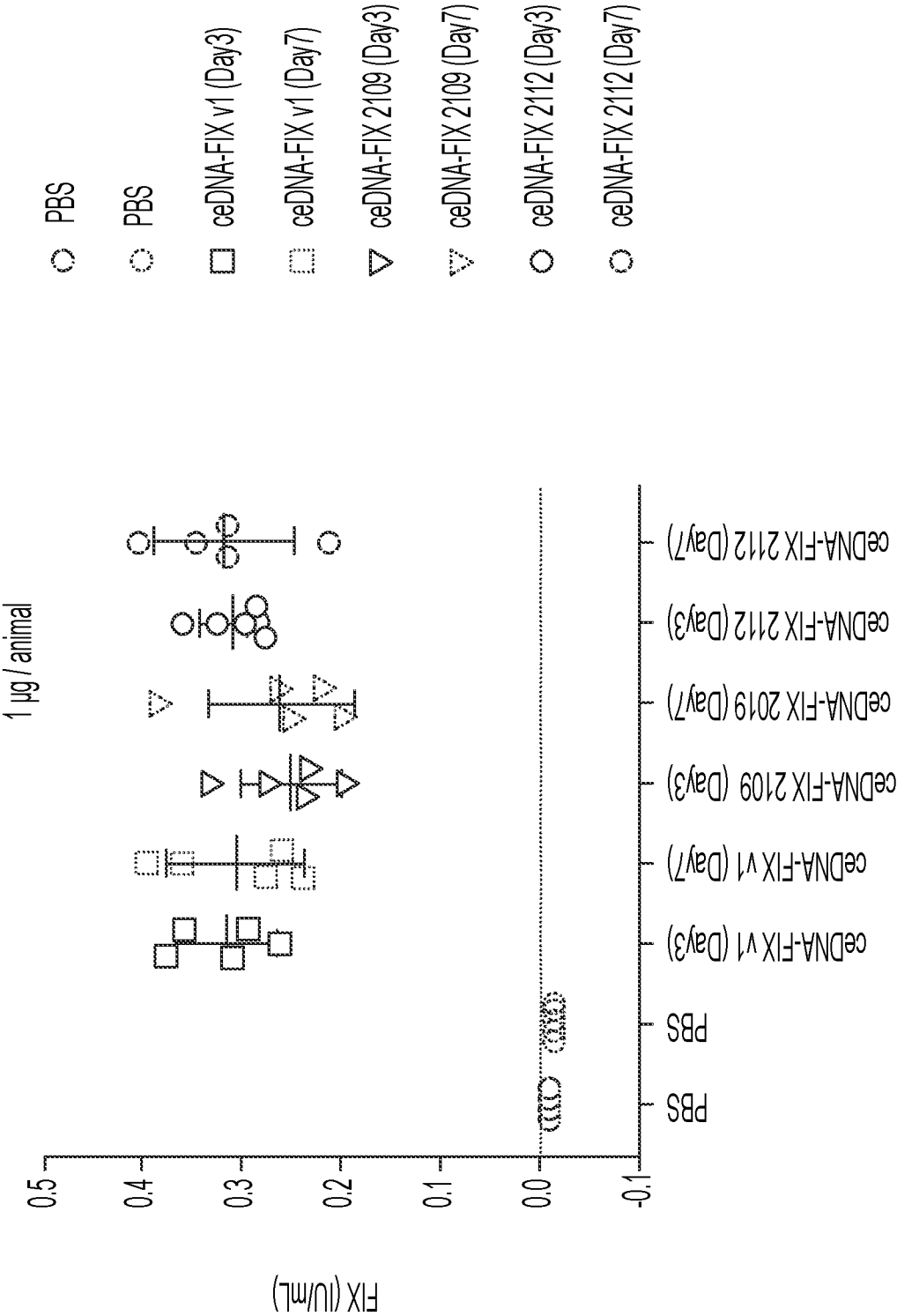


FIG. 13A

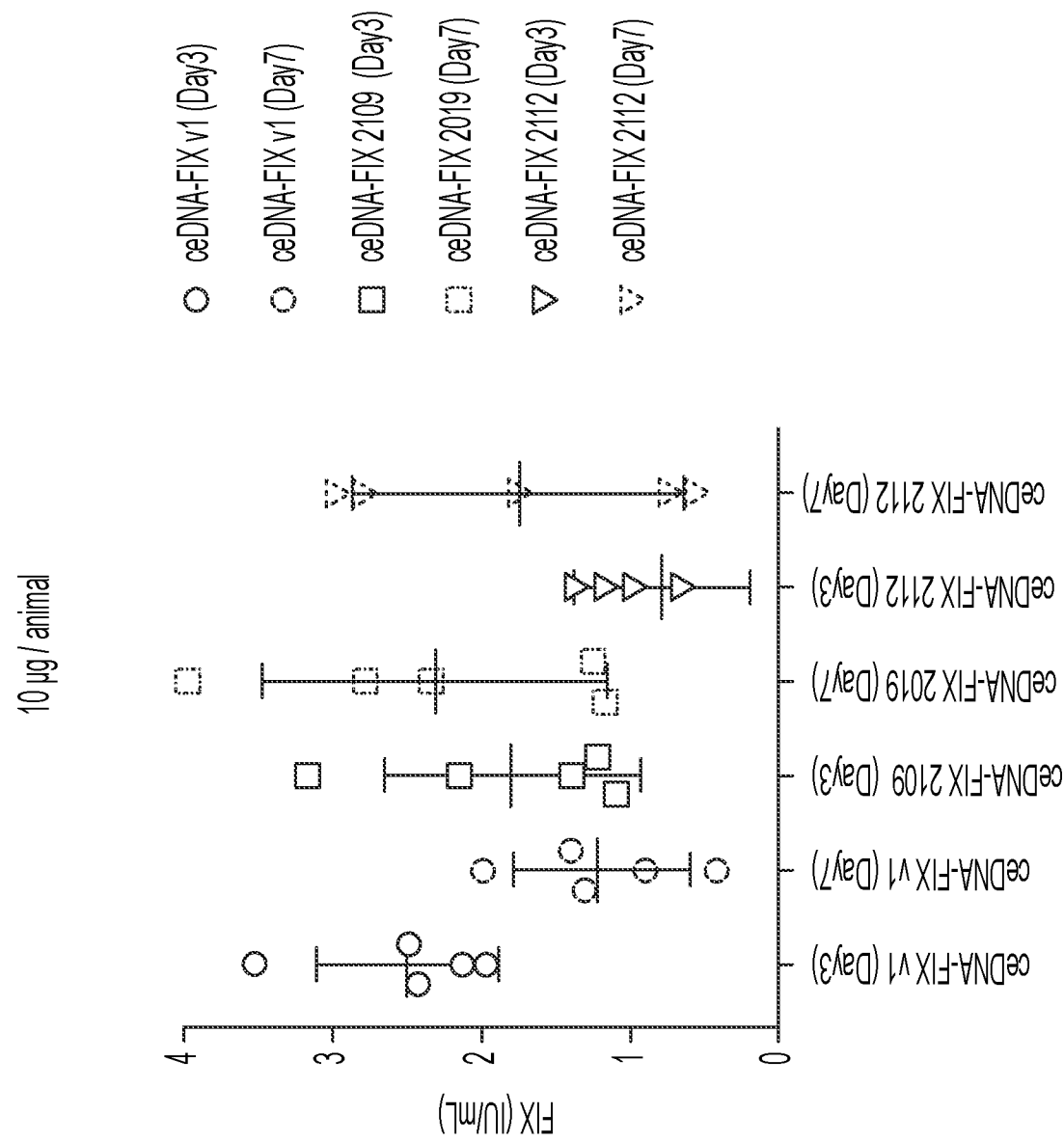


FIG. 13B

SEQUENCE LISTING

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PASACKOW, Eli Hertzal

<120> NON-VIRAL DNA VECTORS AND USES THEREOF FOR EXPRESSING FACTOR IX
THERAPEUTICS

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 cagacggcag aggtctcctc tgccggcccc accgagcgag cgagcgcgca tagagggagt 120
 gggcaactcc atcactaggg gtat 144

<210> 10
 <211> 143
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 10
 ttaccctagt gatggagttg cccactccct ctctgcgcgc gtcgctcgct cgggtggggcc 60
 ggcagaggag acctctgccg tctgcggacc tttggtccgc aggccccacc gagcgagcga 120
 gcgcgcagag agggagtggg caa 143

<210> 11
 <211> 144
 <212> DNA
 <213> Artificial Sequence

<220>

<221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 11
 atacctctag tgatggagtt ggccactccc tctatgcgca ctgctcgct cgggtggggcc 60
 ggacgtggaa acccacgtcc gtctggcgac ctttggtcgc caggccccac cgagcgagcg 120
 agtgcgcata gagggagtgg ccaa 144

<210> 12
 <211> 127
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 12
 agttggccac attagctatg gcgctcgct cactcactcg gccctggaga ccaaaggtct 60
 ccagactgcc ggctcttggc cggcagggcc gagtgagtga gcgagcgcg atagaggag 120
 tggccaa 127

<210> 13
 <211> 166
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 13
 cttacaaac ccccttgctt gagagtgtgg cactctcccc cctgtcgcgt tcgctcgctc 60
 gctggctcgt ttgggggggt ggcagctcaa agagctgcca gacgacggcc ctctggccgt 120
 cgcccccca aacgagccag cgagcgagcg aacgcgacag ggggga 166

<210> 14
 <211> 144
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 14
 atacccttag tgatggagtt gccactccc tctatgcgca ctgctcgct cgggtggggcc 60
 ggcagaggag acctctgccg tctgcggacc tttggtccgc aggccccacc gagcgagcga 120
 gcgcgcatta gagggagtgg gcaa 144

<210> 15
 <211> 120
 <212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 15

aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60

cgcacgcccg gggtttcccg gcggcctcag tgagcgagcg agcgcgagc tgcctgcagg 120

<210> 16

<211> 122

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 16

aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60

ccgacgcccg ggctttgccc gggcggcctc agtgagcgag cgagcgcgca gctgcctgca 120

gg 122

<210> 17

<211> 129

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 17

aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60

ccggcgacc aaaggtcgcc cgacgcccgg gcgcctcagt gagcgagcga gcgcgcagct 120

gcctgcagg 129

<210> 18

<211> 101

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 18

aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60

ctttgcctca gtgagcgagc gagcgcgag ctgcctgcag g 101

<210> 19

<211> 139

<212> DNA

<213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 19
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgaca aagtcgcccg acgcccgggc ttgcccggg cggcctcagt gagcgagcga 120
 gcgcgagct gcctgcagg 139

<210> 20
 <211> 137
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 20
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcaaaa atcgcccgac gcccgggctt tgcccgggcg gcctcagtga gcgagcgagc 120
 gcgagctgc ctgcagg 137

<210> 21
 <211> 135
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 21
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcaaaa cgcccgacgc ccgggctttg cccgggcggc ctactgagc gagcgagcgc 120
 gcagctgcct gcagg 135

<210> 22
 <211> 133
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 22
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcaaag cccgacgccc gggctttgcc cgggcggcct cagtgagcga gcgagcgcgc 120
 agctgcctgc agg 133

<210> 23

<211> 139
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 23
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgcccgg gtttcccggg cggcctcagt gagcgagcga 120
 gcgcgcagct gcctgcagg 139

<210> 24
 <211> 137
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 24
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgcccgg tttccgggcg gcctcagtga gcgagcgagc 120
 gcgcagctgc ctgcagg 137

<210> 25
 <211> 135
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 25
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgcccgt ttcgggcggc ctactgagc gagcgagcgc 120
 gcagctgcct gcagg 135

<210> 26
 <211> 133
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 26
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgcccct tgggcggcct cagtgagcga gcgagcgcg 120
 agctgcctgc agg 133

<210> 27
 <211> 131
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

 <400> 27
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgccttt ggccgacctca gtgagcgagc gagcgcgag 120
 ctgcctgcag g 131

<210> 28
 <211> 129
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

 <400> 28
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgccttt ggccctcagt gagcgagcga gcgcgcagct 120
 gcctgcagg 129

<210> 29
 <211> 127
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

 <400> 29
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgggcgacc aaaggtcgcc cgacgttttc gcctcagtga gcgagcgagc gcgcagctgc 120
 ctgcagg 127

<210> 30
 <211> 122
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

 <400> 30
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60

ccgggcgacc aaaggtcgcc cgacggcctc agtgagcgag cgagcgcgca gctgcctgca 120

gg 122

<210> 31

<211> 130

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 31

aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60

ccgggcgacc aaaggtcgcc cgacgcccgg gcggcctcag tgagcgagcg agcgcgagc 120

tgcctgcagg 130

<210> 32

<211> 120

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 32

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggaaacc cgggcgtgcg 60

cctcagttag cgagcgagcg cgagagagg gaggggccaa ctccatcact aggggttcct 120

<210> 33

<211> 122

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 33

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgt cgggcgacct ttggtcgccc 60

ggcctcagtg agcgagcgag cgcgagaga gggagtggcc aactccatca ctaggggttc 120

ct 122

<210> 34

<211> 122

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 34

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggcaaag cccgggcgtc 60

ggcctcagtg agcgagcgag cgcgagaga gggagtggcc aactccatca ctaggggttc 120

ct 122

<210> 35
 <211> 129
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 35
 cctgcaggca gctgcgcgct cgctcgctca ctgaggcgcc cgggcgctcg gcgacctttg 60
 gtcgcccggc ctcaagtgcg gagcgagcgc gcagagaggg agtggccaac tccatcacta 120
 ggggttcct 129

<210> 36
 <211> 101
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 36
 cctgcaggca gctgcgcgct cgctcgctca ctgaggcaaa gcctcagtga gcgagcgagc 60
 gcgcagagag ggagtggcca actccatcac taggggttc t 101

<210> 37
 <211> 139
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 37
 cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggcaaag cccgggcgctc 60
 gggcgacttt gtcgcccggc ctcaagtgcg gagcgagcgc gcagagaggg agtggccaac 120
 tccatcacta ggggttcct 139

<210> 38
 <211> 137
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 38

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggcaaag cccgggcgtc	60
gggcgatttt cgcccggcct cagtgagcga gcgagcgcgc agagaggag tggccaactc	120
catcactagg ggttcct	137

<210> 39
 <211> 135
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 39	
cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggcaaag cccgggcgtc	60
gggcgtttcg cccggcctca gtgagcgagc gagcgcgcag agaggagtg gccaaactcca	120
tcactagggg ttcct	135

<210> 40
 <211> 133
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 40	
cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggcaaag cccgggcgtc	60
gggctttgcc cggcctcagt gagcgagcga gcgcgagag agggagtggc caactccatc	120
actagggggtt cct	133

<210> 41
 <211> 139
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 41	
cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgggaaacc cgggcgtcgg	60
gcgacctttg gtcgccggc ctcaagtgagc gagcgagcgc gcagagagg agtggccaac	120
tccatcacta ggggttcct	139

<210> 42
 <211> 137
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic

polynucleotide"

<400> 42

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccggaaccg ggcgtcgggc 60
gacctttggt cgcccgccct cagtgcgcga ggcgagcgcg agagagggag tggccaactc 120
catcactagg ggttcct 137

<210> 43

<211> 135

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 43

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccgaaacggg cgtcgggcga 60
cctttggtcg cccggcctca gtgagcgagc gagcgcgag agagggagtg gccaaactcca 120
tcactagggg ttcct 135

<210> 44

<211> 133

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 44

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc ccaaagggcg tcgggcgacc 60
tttggtcgcc cggcctcagt gagcgagcga ggcgagagag agggagtggc caactccatc 120
actaggggtt cct 133

<210> 45

<211> 131

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 45

cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc caaaggcgtc gggcgacatt 60
tggtcgcccg gcctcagtga ggcgagcagc ggcgagagag ggagtggcca actccatcac 120
taggggttcc t 131

<210> 46

<211> 129

<212> DNA

<213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 46
 cctgcaggca gctgcgcgct cgctcgctca ctgaggccgc aaagcgtcgg gcgaccttg 60
 gtcgcccggc ctcaagtgagc gagcgagcgc gcagagaggg agtggccaac tccatcacta 120
 ggggttcct 129

<210> 47
 <211> 127
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 47
 cctgcaggca gctgcgcgct cgctcgctca ctgaggccga aacgtcgggc gacctttggt 60
 cgcccggcct cagtgagcga gcgagcgcgc agagagggag tggccaactc catcactagg 120
 ggttcct 127

<210> 48
 <211> 122
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 48
 aggaacccct agtgatggag ttggccactc cctctctgcg cgctcgctcg ctcaactgagg 60
 ccgggcgacc aaaggtcgcc cgacggcctc agtgagcgag cgagcgcgca gctgcctgca 120
 gg 122

<210> 49
 <211> 12
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 49
 cgatcgttcg at 12

<210> 50
 <211> 12
 <212> DNA
 <213> Artificial Sequence

<220>

<221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 50
 atcgaaccat cg 12

<210> 51
 <211> 12
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 51
 atcgaacgat cg 12

<210> 52
 <211> 165
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 52
 aggaaccct agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagg 60
 ccgcccgggc aaagcccggg cgtcgggcga cctttggtcg cccggcctca gtgagcgagc 120
 gagcgcgag agagggagtg gccaaactcca tcactagggg ttctt 165

<210> 53
 <211> 140
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 53
 cccctagtga tggagttggc cactccctct ctgcgcgctc gctcgctcac tgaggccgcc 60
 cgggcaaagc ccgggcgtcg ggcgaccttt ggctgcccgg cctcagttag cgagcgagcg 120
 cgagagaga tcactagggg 140

<210> 54
 <211> 91
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 54

gcgcgctcgc tcgctcactg aggccgccc ggcaaagccc gggcgtcggg cgacctttgg 60

tcgcccggcc tcagtgagcg agcgagcgcg c 91

<210> 55
 <211> 91
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 55
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgacgcc cgggctttgc 60

ccgggaggcc tcagtgagcg agcgagcgcg c 91

<210> 56
 <211> 1662
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 56
 gccgccacca tggagacgc caaaaacata aagaaaggcc cggcgccatt ctatccgctg 60

gaagatggaa ccgctggaga gcaactgcat aaggctatga agagatacgc cctggttcct 120

ggaacaattg cttttacaga tgcacatatc gaggtggaca tcacttacgc tgagtacttc 180

gaaatgtccg ttcggttggc agaagctatg aaacgatatg ggctgaatac aaatcacaga 240

atcgtcgtat gcagtgaaaa ctctcttcaa ttctttatgc cgggtgttggg cgcgttattt 300

atcggagttg cagttgcgcc cgcgaacgac atttataatg aacgtgaatt gctcaacagt 360

atgggcattt cgagcctac cgtgggtgtc gtttcacaaa aggggttgca aaaaattttg 420

aacgtgcaaa aaaagctccc aatcatccaa aaaattatta tcatggattc taaaacggat 480

taccagggat ttcagtcgat gtacacgttc gtcacatctc atctacctcc cggttttaat 540

gaatacgatt ttgtgccaga gtccttcgat agggacaaga caattgcact gatcatgaac 600

tcctctggat ctactggtct gcctaaaggc gtcgctctgc ctcatagaac tgcctgcgtg 660

agattctcgc atgccagaga tcctatTTTT ggcaatcaaa tcattccgga tactgagatt 720

ttaagtgttg ttccattcca tcacggtttt ggaatgttta ctacactcgg atatttgata 780

tgtggatttc gagtcgtctt aatgtataga tttgaagaag agctgtttct gaggagcctt 840

caggattaca agattcaaag tgcgtgctg gtgccaaccc tattctcctt cttcgccaaa 900

agcactctga ttgacaaata cgatttatct aatttacacg aaattgcttc tgggtggcgt 960

cccctctcta aggaagtcgg ggaagcgggt gccaaagggt tccatctgcc aggtatcagg 1020

caaggatatg ggctcactga gactacatca gctattctga ttacaccga gggggatgat 1080

aaaccgggcg cggtcggtaa agttgttcca ttttttgaag cgaaggttgt ggatctggat 1140

accgggaaaa cgctgggcgt taatcaaaga ggcgaactgt gtgtgagagg tcctatgatt 1200
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ctacattctg gagacatagc ttactgggac gaagacgaac acttcttcat cgttgaccgc 1320
ctgaagtctc tgattaagta caaaggctat caggtggctc ccgctgaatt ggaatccatc 1380
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atcgtggatt acgtcgccag tcaagtaaca accgcgaaaa agttgcgcgg aggagttgtg 1560
tttgtggacg aagtaccgaa aggtcttacc ggaaaactcg acgcaagaaa aatcagagag 1620
atcctcataa aggccaagaa gggcggaaag atcgccgtgt aa 1662

<210> 57
<211> 453
<212> PRT
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polypeptide"

<220>
<221> MOD_RES
<222> (1)..(1)
<223> Any amino acid

<400> 57
Xaa Val Gln Leu Val Glu Ser Gly Gly Gly Val Val Gln Pro Gly Arg
1 5 10 15

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Ala Phe Ser Ser Tyr
20 25 30

Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45

Ala Val Ile Trp Phe Asp Gly Thr Lys Lys Tyr Tyr Thr Asp Ser Val
50 55 60

Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65 70 75 80

Leu Gln Met Asn Thr Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
85 90 95

Ala Arg Asp Arg Gly Ile Gly Ala Arg Arg Gly Pro Tyr Tyr Met Asp
100 105 110

Val Trp Gly Lys Gly Thr Thr Val Thr Val Ser Ser Ala Ser Thr Lys
115 120 125

Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly
130 135 140

Gly Thr Ala Ala Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro
145 150 155 160

Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr
165 170 175

Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val
180 185 190

Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn
195 200 205

Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro
210 215 220

Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu
225 230 235 240

Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp
245 250 255

Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp
260 265 270

Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly
275 280 285

Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn
290 295 300

Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp
305 310 315 320

Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro
325 330 335

Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
340 345 350

Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn
355 360 365

Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile
370 375 380

Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr
385 390 395 400

Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys

405

410

415

Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys
 420 425 430

Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu
 435 440 445

Ser Leu Ser Pro Gly
 450

<210> 58

<211> 214

<212> PRT

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
 polypeptide"

<400> 58

Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly
 1 5 10 15

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser Tyr
 20 25 30

Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly
 50 55 60

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
 65 70 75 80

Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ser Tyr Ser Thr Pro Leu
 85 90 95

Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 100 105 110

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
 115 120 125

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
 130 135 140

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
 145 150 155 160

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
 165 170 175

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
180 185 190

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
195 200 205

Phe Asn Arg Gly Glu Cys
210

<210> 59
<211> 1310
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 59
ggagccgaga gtaattcata caaaaggagg gatcgcttc gcaaggggag agcccaggga 60
ccgtccctaa attctcacag acccaaatcc ctgtagccgc cccacgacag cgcgaggagc 120
atgcgctcag ggctgagcgc ggggagagca gagcacacaa gtcctagac cctggctctg 180
ggggggagga ccggggagct ggcgcggggc aaactgggaa agcgggtgtcg tgtgctggct 240
ccgccctctt ccgaggggtg ggggagaacg gtatataagt gcggcagtcg ccttgacgt 300
tctttttcgc aacgggtttg ccgtcagaac gcagggtgagg ggcggggtgtg gcttccgcgg 360
gccgccgagc tggaggtcct gctccgagcg ggccggggccc cgctgtcgtc ggcggggatt 420
agctgcgagc attcccgctt cgagttgcgg gcggcgcggg aggcagagtg cgaggcctag 480
cggcaacccc gtagcctcgc ctctgtgtccg gcttgaggcc tagcgtggtg tccgcgccgc 540
cgccgcgtgc tactccggcc gcactctggt cttttttttt tttgtttgtt ttgccctgct 600
gccttcgatt gccgttcagc aataggggct aacaaaggga gggcgcgggg cttgctcgcc 660
cggagcccgg agaggtcatg gttggggagg aatggaggga caggagtggc ggctggggcc 720
cgccgcctt cggagcacat gtccgacgcc acctggatgg gcgaggcct ggggtttttc 780
ccgaagcaac caggctgggg ttagcgtgcc gaggccatgt ggccccagca cccggcacga 840
tctggcttgg cggcgccgcg ttgccctgcc tccctaacta gggtagggcc atccgtccg 900
gcaccagtgt cgtgcgtgga aagatggccg ctcccgggcc ctgttgcaag gagctcaaaa 960
tggaggacgc ggcagcccgg tggagcgggc gggtagagtca cccacacaaa ggaagagggc 1020
ctggtccctc accggctgct gttcctgtg acccgtggt cctatcgcc gcaatagtca 1080
cctcgggctt ttgagcacgg ctagtcgcgg cggggggagg ggatgtaatg gcgttgagtg 1140
ttgttcacat ttggtgggtg gagactagtc aggccagcct ggcgctggaa gtcatttttg 1200
gaatttgtcc cttgagttt tgagcggagc taattctcgg gcttcttagc ggttcaaagg 1260
tatcttttaa accctttttt aggtgttgtg aaaaccaccg ctaattcaaa 1310

<210> 60
<211> 16
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 60
gcgcgctcgc tcgctc

16

<210> 61
<211> 6
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 61
ggttga

6

<210> 62
<211> 4
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 62
agtt

4

<210> 63
<211> 6
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 63
ggttgg

6

<210> 64
<211> 6
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 64
agttgg

6

<210> 65
 <211> 6
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 65
 agttga 6

<210> 66
 <211> 6
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 66
 rrtrrr 6

<210> 67
 <211> 581
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

 <400> 67
 gagcatctta ccgccattta ttcccatatt tgttctgttt ttcttgattt gggatatacat 60

 ttaaatgtta ataaaacaaa atgggtggggc aatcatttac atttttaggg atatgtaatt 120

 actagttcag gtgtattgcc acaagacaaa catgttaaga aactttcccg ttatttacgc 180

 tctgttcctg ttaatcaacc tctggattac aaaatttgtg aaagattgac tgatattctt 240

 aactatgttg ctctttttac gctgtgtgga tatgctgctt tatagcctct gtatctagct 300

 attgcttccc gtacggcttt cgttttctcc tccttgtata aatcctgggt gctgtctctt 360

 ttagaggagt tgtggcccg tgtccgtcaa cgtggcgtgg tgtgctctgt gtttgctgac 420

 gcaaccccca ctggctgggg cattgccacc acctgtcaac tcctttctgg gactttcgct 480

 ttccccctcc cgatcgccac ggacagaactc atcgccgcct gccttgcccg ctgctggaca 540

 ggggctaggt tgctgggcac tgataattcc gtgggtgtgt c 581

<210> 68
 <211> 225
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic

polynucleotide"

<400> 68
tgtgccttct agttgccagc catctgttgt ttgccctcc cccgtgcctt ccttgaccct 60
ggaagggtgcc actcccactg tcctttccta ataaaatgag gaaattgcat cgcattgtct 120
gagtaggtgt cattctattc tggggggtgg ggtggggcag gacagcaagg gggaggattg 180
ggaagacaat agcaggcatg ctggggatgc ggtgggctct atggc 225

<210> 69
<211> 8
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 69
actgaggc 8

<210> 70
<211> 8
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 70
gcctcagt 8

<210> 71
<211> 16
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 71
gagcgagcga gcgcgc 16

<210> 72
<211> 1923
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 72
tcaatattgg ccattagcca tattattcat tggttatata gcataaatca atattggcta 60
ttggccattg catacgttgt atctatatca taatatgtac atttatattg gctcatgtcc 120

aatatgaccg ccatgttggc attgattatt gactagttat taatagtaat caattacggg	180
gtcattagtt catagcccat atatggagtt ccgcgttaca taacttacgg taaatggccc	240
gcctggctga ccgccaacg acccccggcc attgacgtca ataatgacgt atgttcccat	300
agtaacgcca atagggactt tccattgacg tcaatgggtg gagtatttac ggtaaactgc	360
ccacttggca gtacatcaag tgtatcatat gccaaagtccg cccctattg acgtcaatga	420
cggtaaatgg cccgcctggc attatgccc gtacatgacc ttacgggact ttcctacttg	480
gcagtacatc tacgtattag tcatcgctat taccatggtc gaggtgagcc ccacgttctg	540
cttcactctc cccatctccc cccctcccc accccaatt ttgtatttat ttatTTTTTA	600
attatTTTgt gcagcgatgg gggcgggggg gggggggggg cgcgcgccag gcggggcggg	660
gcggggcgag gggcggggcg gggcgaggcg gagagggtcg gcggcagcca atcagagcgg	720
cgcgctccga aagtttcctt ttatggcgag gcggcggcgg gcggcgccct ataaaaagcg	780
aagcgcgcgg cggcggggag tcgctgcgac gctgccttcg ccccggtccc cgctccgccg	840
ccgcctcgcg ccgcccggc cggctctgac tgaccgcgtt actcccacag gtgagcgggc	900
gggacggccc ttctcctccg ggctgtaatt agcgttgggt ttaatgacgg cttgtttctt	960
ttctgtggct gcgtgaaagc cttgaggggc tccgggaggg ccctttgtgc gggggggagc	1020
ggctcggggg gtgcgtgcgt gtgtgtgtgc gtggggagcg ccgcgtgcgg cccgcgtgc	1080
ccggcggctg tgagcgtgc gggcgcggcg cggggctttg tgcgtccgc agtgtgcgcg	1140
aggggagcgc ggccgggggc ggtgccccgc ggtgcggggg gggctgcgag gggaacaaag	1200
gctgcgtgcg ggggtgtgtc gtgggggggt gagcaggggg tgtgggcgcg gcggtcgggc	1260
tgtaaccccc ccctgcaccc ccctccccga gttgctgagc acggcccggc ttcgggtgcg	1320
gggctccgta cggggcgtgg cgcggggctc gccgtgccgg gcgggggggtg gcggcaggtg	1380
ggggtgccgg gcggggcggg gccgcctcgg gccggggagg gctcggggga ggggcgcggc	1440
ggccccgga gcgccggcgg ctgtcgaggc gcggcgagcc gcagccattg ctttttatgg	1500
taatcgtgcg agagggcgca gggacttcct ttgtccaaa tctgtgcgga gccgaaatct	1560
gggaggcgcc gccgcacccc ctctagcggg cgcggggcga agcgggtcgg cgccggcagg	1620
aaggaaatgg gcggggaggg ccttcgtgcg tcgccgcgcc gccgtcccct tctccctctc	1680
cagcctcggg gctgtccgcg gggggacggc tgccttcggg ggggacgggg cagggcgggg	1740
ttcggcttct ggctgtgac ggcggtctt agagcctctg ctaaccatgt tttagccttc	1800
ttctttttcc tacagctcct gggcaacgtg ctggttattg tgctgtctca tcatttgtcg	1860
acagaattcc tcgaagatcc gaagggttc aagcttggca ttccggtact gttggtaaag	1920
cca	1923

<210> 73

<211> 1272

<212> DNA

<213> Artificial Sequence

<220>

<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 73
aggctcagag gcacacagga gtttctgggc tcaccctgcc cccttccaac ccctcagttc 60
ccatcctcca gcagctgttt gtgtgctgcc tctgaagtcc aactgaaca aacttcagcc 120
tactcatgtc cctaaaatgg gcaaacattg caagcagcaa acagcaaaca cacagccctc 180
cctgcctgct gaccttggag ctggggcaga ggtcagagac ctctctgggc ccatgccacc 240
tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagagggt gtcctggcgt 300
ggtttaggta gtgtgagagg gtccgggttc aaaaccactt gctgggtggg gaggcgtcag 360
taagtggcta tgccccgacc ccgaagcctg tttccccatc tgtacaatgg aaatgataaa 420
gacgcccac tcataggggtt tttgtggcaa ataaacattt ggtttttttg ttttgttttg 480
ttttgttttt tgagatggag gtttgctctg tcgcccaggc tggagtgcag tgacacaatc 540
tcatctcacc acaaccttcc cctgcctcag cctcccaagt agctgggatt acaagcatgt 600
gccaccacac ctggctaatt ttctattttt agtagagacg ggtttctcca tgttggtcag 660
cctcagcctc ccaagtaact gggattacag gcctgtgcc cccacccgg ctaatttttt 720
ctatttttga caggacggg gtttcacat gttggtcagg ctgggtctaga ggtaccggat 780
cttgctacca gtggaacagc cactaaggat tctgcagtga gagcagagg ccagctaagt 840
ggtactctcc cagagactgt ctgactcac ccacccctc caccttggac acaggacgct 900
gtggtttctg agccaggta aatgactcct ttcggtaagt gcagtggaag ctgtacactg 960
cccaggcaaa gcgtccgggc agcgtaggc ggcgactcag atcccagcca gtggacttag 1020
cccctgtttg ctctccgat aactgggggtg accttggtta atattcacca gcagcctccc 1080
ccgttgcccc tctggatcca ctgcttaaat acggacgagg acagggccct gtctcctcag 1140
cttcaggcac caccactgac ctgggacagt gaatccggac tctaaggtaa atataaaatt 1200
tttaagtgtg taatgtgtta aactactgat tctaattgtt tctctctttt agattccaac 1260
ctttggaact ga 1272

<210> 74
<211> 1177
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 74
ggctcagagg ctgagaggca cacaggagtt tctgggctca ccctgcccc ttccaacccc 60
tcagttccca tcctccagca gctgtttgtg tgctgcctct gaagtccaca ctgaacaaac 120
ttcagcctac tcatgtccct aaaatgggca aacattgcaa gcagcaaaca gcaaacacac 180
agccctccct gcctgctgac cttggagctg gggcagaggc cagagacctc tctgggcca 240

tgccacctcc aacatccact cgaccccttg gaatttcggt ggagaggagc agaggttgtc	300
ctggcgtggt ttaggtagtg tgagagggtc cgggttcaaa accacttgct ggggtggggag	360
tcgtcagtaa gtggctatgc ccgcaccccg aagcctgttt ccccatctgt acaatggaaa	420
tgataaagac gcccatctga tagggttttt gtggcaaata aacatttggt ttttttgttt	480
tgttttgttt tgttttttga gatggaggtt tgctctgtcg cccaggctgg agtgcagtga	540
cacaatctca tctcaccaca accttcccct gcctcagcct cccaagtagc tgggattaca	600
agcatgtgcc accacacctg gctaattttc tatttttagt agagacgggt ttctccatgt	660
tggtcagcct cagcctccca agtaactggg attacaggcc tgtgccacca caccggcta	720
attttttcta tttttgacag ggacgggggtt tcaccatgtt ggtcaggctg gtctagaggt	780
accggatctt gctaccagtg gaacagccac taaggattct gcagtgagag cagagggcca	840
gctaagtggg actctcccag agactgtctg actcacgcca cccctccac cttggacaca	900
ggacgctgtg gtttctgagc caggtacaat gactcctttc ggtaagtga gtggaagctg	960
tacactgccc aggcaaagcg tccgggcagc gtaggcgggc gactcagatc ccagccagt	1020
gacttagccc ctgtttgctc ctccgataac tggggtgacc ttggttaata ttcaccagca	1080
gcctcccccg ttgcccctct ggatccactg cttaaatacg gacgaggaca gggcctgtc	1140
tcctcagctt caggcaccac cactgacctg ggacagt	1177

<210> 75

<211> 547

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 75

ccctaaaatg ggcaaacatt gcaagcagca aacagcaaac acacagccct ccctgcctgc	60
tgaccttgga gctggggcag aggtcagaga cctctctggg cccatgccac ctccaacatc	120
cactcgaccc cttggaattt ttcggtggag aggagcagag gttgtcctgg cgtggtttag	180
gtagtgtgag aggggaatga ctcttttcgg taagtgcagt ggaagctgta cactgcccag	240
gcaaagcgtc cgggcagcgt aggcgggcga ctcagatccc agccagtgga cttagcccct	300
gtttgtctct ccgataactg gggtgacctt ggtaaataatt caccagcagc ctccccgtt	360
gcccctctgg atccactgct taaatacggg cgaggacagg gccctgtctc ctgagcttca	420
ggcaccacca ctgacctggg acagtgaatc cggactctaa ggtaaatata aaatttttaa	480
gtgtataatg tgtaaacta ctgattctaa ttgtttctct ctttttagatt ccaacctttg	540
gaactga	547

<210> 76

<211> 556

<212> DNA

<213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 76
 ccctaaaatg ggcaaacatt gcaagcagca aacagcaaac acacagccct ccctgcctgc 60
 tgaccttgga gctggggcag aggtcagaga cctctctggg cccatgccac ctccaacatc 120
 cactcgaccc cttggaatth cgggtggagag gagcagaggt tgtcctggcg tggtttaggt 180
 agtgtgagag gggaaatgact cctttcggtg agtgcagtgg aagctgtaca ctgcccaggc 240
 aaagcgtccg ggacagcgtg gcggggcgact cagatcccag ccagtggact tagcccctgt 300
 ttgctcctcc gataactggg gtgaccttgg ttaatatcca ccagcagcct cccccgttgc 360
 ccctctggat ccactgctta aatacggacg aggacactcg agggccctgt ctcctcagct 420
 tcaggcacca ccactgacct gggacagtga atccggacat cgattctaag gtaaataata 480
 aatthtttaag tgtataatth gttaaactac tgattctaat tgtttctctc ttttagattc 540
 caacctttgg aactga 556

<210> 77
 <211> 1179
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 77
 ggctccggtg cccgtcagtg ggacagagcg acatcgccca cagtccccga gaagttgggg 60
 ggagggggtcg gcaattgaac cgggtgcctag agaaggtggc gcggggtaaa ctgggaaagt 120
 gatgtcgtgt actggctccg cctthtttccc gaggggtggg gagaaccgta tataagtgca 180
 gtagtcgccg tgaacgttct ttttcgcaac gggtttgccg ccagaacaca ggtaagtgcc 240
 gtgtgtggtt cccgcgggcc tggcctcttt acgggttatg gcccttgctg gccttgaatt 300
 acttccacct ggctgcagta cgtgattctt gatcccgagc ttcgggttgg aagtgggtgg 360
 gagagttcga ggccttgccg ttaaggagcc ctttcgcctc gtgcttgagt tgaggcctgg 420
 cctgggcgct gggggccgcc cgtgcgaatc tgggtggcacc ttcgcgcctg tctcgtgct 480
 ttcgataagt ctctagccat ttaaaattht tgatgacctg ctgcgacgct tthtttctgg 540
 caagatagtc ttgtaaatgc gggccaagat ctgcacactg gtatttcggt ttttggggcc 600
 gcgggcggcg acggggcccg tgcgtcccag cgcacatgth cggcgaggcg gggcctgcga 660
 gcgcggccac cgagaatcgg acgggggtag tctcaagctg gccggcctgc tctggtgcct 720
 ggtctcgcgc cgccgtgtat cggcccgccc tgggcggcaa ggctggcccg gtcggcacca 780
 gttgcgtgag cggaaagatg gccgcttccc ggccttgctg caggagctc aaaatggagg 840
 acgcggcgct cgggagagcg ggcgggtgag tcaccacac aaagggaaaag ggcctthtcg 900

tcctcagccg tcgcttcacg tgactccacg gaggaccggg cgccgtccag gcacctcgat	960
tagttctcga gcttttgag tacgtcgtct ttaggttggg gggaggggtt ttatgcatg	1020
gagtttccc acactgagtg ggtggagact gaagttaggc cagcttgga cttgatgtaa	1080
ttctccttg aatttgccct ttttgagttt ggatcttggt tcattctcaa gcctcagaca	1140
gtggttcaaa gtttttttct tccatttcag gtgtcgtga	1179

<210> 78
 <211> 141
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 78 aataaacgat aacgccgttg gtggcgtgag gcatgtaaaa ggttacatca ttatcttgtt	60
cgccatccgg ttggtataaa tagacgttca tgttggtttt tgtttcagtt gcaagttggc	120
tgcggcgcgc gcagcacctt t	141

<210> 79
 <211> 317
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 79 ggtgtggaaa gtccccaggc tccccagcag gcagaagtat gcaaagcatg catctcaatt	60
agtcagcaac caggtgtgga aagttcccag gctccccagc aggcagaagt atgcaaagca	120
tgcattctaa ttagtcagca accatagtcc cgcccctaac tccgcccata ccgcccctaa	180
ctccgcccag ttccgcccata tctccgcccc atggctgact aatttttttt atttatgcag	240
aggccgaggc cgcttcggcc tctgagctat tccagaagta gtgaggaggc ttttttgag	300
gcctaggctt ttgcaaa	317

<210> 80
 <211> 241
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 80 gagggcctat ttccatgat tccttcatat ttgcatatac gatacaaggc tgtagagag	60
ataattggaa ttaatttgac tgtaaacaca aagatattag taaaaatac gtgacgtaga	120
aagtaataat ttcttggtga gtttcagtt taaaattat gttttaaaat ggactatcat	180

atgcttaccg taacttgaaa gtatttcgat ttcttggctt tatatatctt gtggaaagga 240

c 241

<210> 81

<211> 215

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 81

gaacgctgac gtcacatcaacc cgctccaagg aatcgcgggc ccagtgtcac taggcgggaa 60

cacccagcgc gcgtgcgccc tggcaggaag atggctgtga gggacagggg agtggcgccc 120

tgcaatatatt gcatgtcgct atgtgttctg ggaaatcacc ataaacgtga aatgtctttg 180

gatttgggaa tcgtataaga actgtatgag accac 215

<210> 82

<211> 546

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 82

ccctaaaatg ggcaaacatt gcaagcagca aacagcaaac acacagccct ccctgcctgc 60

tgaccttgga gctggggcag aggtcagaga cctctctggg cccatgccac ctccaacatc 120

cactcgaccc cttggaattt ttcggtggag aggagcagag gttgtcctgg cgtgggttag 180

gtagtgtgag aggggaatga ctcttttcgg taagtgcagt ggaagctgta cactgcccag 240

gcaaagcgtc cgggcagcgt aggcgggcga ctcatatccc agccagtgga cttagcccct 300

gtttgctcct ccgataactg gggtgacctt ggtaaatatt caccagcagc ctccccggtt 360

gcccccttgg atccactgct taaatacggg cgaggacagg gccctgtctc ctgagcttca 420

ggcaccacca ctgacctggg acagtgaatc cggactctaa ggtaaatata aaatttttaa 480

gtgtataatg tgtaaaacta ctgattctaa ttgtttctct ctttttagatt ccaacctttg 540

gaactg 546

<210> 83

<211> 576

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 83

tagtaatcaa ttacggggtc attagttcat agcccatata tggagttccg cgttacataa	60
cttacggtaa atggcccgcc tggctgaccg cccaacgacc cccgcccatt gacgtcaata	120
atgacgtatg ttcccatagt aacgccaata gggactttcc attgacgtca atgggtggag	180
tatttacggt aaactgcccc cttggcagta catcaagtgt atcatatgcc aagtacgccc	240
cctattgacg tcaatgacgg taaatggccc gcctggcatt atgcccagta catgacctta	300
tgggactttc ctacttgga gtacatctac gtattagtca tcgctattac catggtgatg	360
cggttttggc agtacatcaa tgggcgtgga tagcggtttg actcacgggg atttccaagt	420
ctccacccca ttgacgtcaa tgggagtttg ttttggcacc aaaatcaacg ggactttcca	480
aaatgtcgta acaactccgc ccattgacg caaatgggcg gtaggcgtgt acggtgggag	540
gtctatataa gcagagctgg tttagtgaac cgtcag	576

<210> 84
 <211> 150
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 84	
ataaacgata acgccgttgg tggcgtgagg catgtaaaag gttacatcat tatcttggtc	60
gccatccggt tggataaat agacgttcat gttgggtttt gtttcagttg caagttggct	120
gcggcgcgcg cagcaccttt gcggccatct	150

<210> 85
 <211> 1313
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 85	
ggagccgaga gtaattcata caaaaggagg gatcgcttc gcaaggggag agcccaggga	60
ccgtccctaa attctcacag acccaaattc ctgtagccgc cccacgacag cgcgaggagc	120
atgcgcccag ggctgagcgc gggtagatca gagcacacaa gctcacagtc cccggcggtg	180
gggggagggg cgcgctgagc gggggccagg gagctggcgc ggggcaaact gggaaagtgg	240
tgtcgtgtgc tggctccgcc ctcttcccga gggtagggga gaacggtata taagtgcggt	300
agtgccttg gacgttcttt ttcgcaacgg gtttgccgtc agaacgcagg tgagtggcgg	360
gtgtggcttc cgcgggcccc ggagctggag ccctgctctg agcgggcccgg gctgatatgc	420
gagtgtcgtc cgagggttt agctgtgagc attcccactt cgagtggcgg gcggtgcggg	480
ggtgagagtg cgaggcctag cggcaacccc gtagcctcgc ctctgtgtccg gcttgaggcc	540
tagcgtggtg tccgcgccg cgtgccactc cggccgcact atgcgttttt tgtccttgct	600

gccctcgatt gccttccagc agcatgggct aacaaaggga ggggtgtgggg ctcaacttta	660
aggagcccat gaagcttacg ttggatagga atggaagggc aggaggggcg actggggccc	720
gcccgccttc ggagcacatg tccgacgcca cctggatggg gcgaggcctg tggctttccg	780
aagcaatcgg gcgtgagttt agcctacctg ggccatgtgg ccctagcact gggcacggtc	840
tggcctggcg gtgccgcgtt cccttgcttc ccaacaaggg tgaggccgtc ccgcccggca	900
ccagttgctt gcgcggaaag atggccgctc ccggggccct gttgcaagga gctcaaaatg	960
gaggacgcgg cagcccgggtg gagcgggcgg gtgagtcacc cacacaaagg aagagggcct	1020
tgcccctcgc cggccgctgc ttcctgtgac cccgtggtct atcggccgca tagtcacctc	1080
gggcttctct tgagcaccgc tcgtcgcggc ggggggaggg gatctaattg cgttggagtt	1140
tgttcacatt tggtaggttg agactagtca ggccagcctg gcgctggaag tcattcttgg	1200
aatttgcccc tttgagtttg gagcaggct aatttcaag cctcttagcg gttcaaaggt	1260
attttctaaa cccgtttcca ggtgttgtga aagccaccgc taattcaaag caa	1313

<210> 86
 <211> 213
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 86	
taagatacat tgatgagttt ggacaaacca caactagaat gcagtgaaaa aaatgcttta	60
tttgtgaaat ttgtgatgct attgctttat ttgtaacct tataagctgc aataaacaag	120
ttaacaacaa caattgcatt ctttttatgt ttcaggttca gggggaggtg tgggaggttt	180
tttaaagcaa gtaaaacctc tacaaatgtg gta	213

<210> 87
 <211> 7
 <212> PRT
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic peptide"

<400> 87	
Pro Lys Lys Lys Arg Lys Val	
1	5

<210> 88
 <211> 19
 <212> PRT
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic"

peptide"

<400> 88

Met Asp Trp Thr Trp Arg Ile Leu Phe Leu Val Ala Ala Ala Thr Gly
1 5 10 15

Ala His Ser

<210> 89

<211> 19

<212> PRT

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
peptide"

<400> 89

Met Leu Pro Ser Gln Leu Ile Gly Phe Leu Leu Leu Trp Val Pro Ala
1 5 10 15

Ser Arg Gly

<210> 90

<211> 7

<212> PRT

<213> Simian virus 40

<400> 90

Pro Lys Lys Lys Arg Lys Val
1 5

<210> 91

<211> 21

<212> DNA

<213> Simian virus 40

<400> 91

cccaagaaga agaggaaggt g

21

<210> 92

<211> 16

<212> PRT

<213> Unknown

<220>

<221> source

<223> /note="Description of Unknown:
Nucleoplasmin bipartite NLS sequence"

<400> 92

Lys Arg Pro Ala Ala Thr Lys Lys Ala Gly Gln Ala Lys Lys Lys Lys
1 5 10 15

<210> 93

<211> 9

<212> PRT

<213> Unknown

<220>
<221> source
<223> /note="Description of Unknown:
C-myc NLS sequence"

<400> 93
Pro Ala Ala Lys Arg Val Lys Leu Asp
1 5

<210> 94
<211> 11
<212> PRT
<213> Unknown

<220>
<221> source
<223> /note="Description of Unknown:
C-myc NLS sequence"

<400> 94
Arg Gln Arg Arg Asn Glu Leu Lys Arg Ser Pro
1 5 10

<210> 95
<211> 38
<212> PRT
<213> Homo sapiens

<400> 95
Asn Gln Ser Ser Asn Phe Gly Pro Met Lys Gly Gly Asn Phe Gly Gly
1 5 10 15

Arg Ser Ser Gly Pro Tyr Gly Gly Gly Gly Gln Tyr Phe Ala Lys Pro
20 25 30

Arg Asn Gln Gly Gly Tyr
35

<210> 96
<211> 42
<212> PRT
<213> Unknown

<220>
<221> source
<223> /note="Description of Unknown:
IBB domain from importin-alpha sequence"

<400> 96
Arg Met Arg Ile Glx Phe Lys Asn Lys Gly Lys Asp Thr Ala Glu Leu
1 5 10 15

Arg Arg Arg Arg Val Glu Val Ser Val Glu Leu Arg Lys Ala Lys Lys
20 25 30

Asp Glu Gln Ile Leu Lys Arg Arg Asn Val
35 40

<210> 97
<211> 8
<212> PRT

<213> Unknown

<220>

<221> source

<223> /note="Description of Unknown:
Myoma T protein sequence"

<400> 97

Val Ser Arg Lys Arg Pro Arg Pro
1 5

<210> 98

<211> 8

<212> PRT

<213> Unknown

<220>

<221> source

<223> /note="Description of Unknown:
Myoma T protein sequence"

<400> 98

Pro Pro Lys Lys Ala Arg Glu Asp
1 5

<210> 99

<211> 8

<212> PRT

<213> Homo sapiens

<400> 99

Pro Gln Pro Lys Lys Lys Pro Leu
1 5

<210> 100

<211> 12

<212> PRT

<213> Mus musculus

<400> 100

Ser Ala Leu Ile Lys Lys Lys Lys Met Ala Pro
1 5 10

<210> 101

<211> 70

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 101

gcgcgctcgc tcgctcactg aggccgcccc ggaaacccgg gcgtg'gcct cagtgcgcga 60

gcgagcgcg 70

<210> 102

<211> 70

<212> DNA

<213> Artificial Sequence

<220>

<221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 102
 gcgcgctcgc tcgctcactg aggcgcacgc ccgggtttcc cgggcggcct cagtgagcga 60
 gcgagcgcg 70

<210> 103
 <211> 72
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 103
 gcgcgctcgc tcgctcactg aggccgtcgg gcgacctttg gtcgcccggc ctcaagtgagc 60
 gagcgagcgc gc 72

<210> 104
 <211> 72
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 104
 gcgcgctcgc tcgctcactg aggccggggcg accaaaggtc gcccgcgggc ctcaagtgagc 60
 gagcgagcgc gc 72

<210> 105
 <211> 72
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 105
 gcgcgctcgc tcgctcactg aggccgcccg ggcaaagccc gggcgtcggc ctcaagtgagc 60
 gagcgagcgc gc 72

<210> 106
 <211> 72
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 106

gcgcgctcgc tcgctcactg aggccgacgc ccgggctttg cccgggcggc ctcaagtgc 60
gagcgagcgc gc 72

<210> 107
<211> 83
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 107
gcgcgctcgc tcgctcactg aggccgcccg ggcaaagccc gggcgtcggg ctttgcccgg 60
cctcaagtgc cgagcgagcg cgc 83

<210> 108
<211> 83
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 108
gcgcgctcgc tcgctcactg aggccgggca aagcccgcgc cccgggcttt gcccgggcgg 60
cctcaagtgc cgagcgagcg cgc 83

<210> 109
<211> 77
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 109
gcgcgctcgc tcgctcactg aggccgaaac gtcgggcgc ctttggtcgc ccggcctcag 60
tgagcgagcg agcgcgc 77

<210> 110
<211> 77
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 110
gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgacgtt tcggcctcag 60
tgagcgagcg agcgcgc 77

<210> 111
 <211> 51
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 111
 gcgcgctcgc tcgctcactg aggcaaagcc tcagtgagcg agcgagcgcg c 51

 <210> 112
 <211> 51
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 112
 gcgcgctcgc tcgctcactg aggctttgcc tcagtgagcg agcgagcgcg c 51

 <210> 113
 <211> 80
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 113
 gcgcgctcgc tcgctcactg aggccgcccg ggcgtcgggc gacctttggt cgcccggcct 60
 cagtgagcga gcgagcgcg 80

 <210> 114
 <211> 80
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 114
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgacgcc cgggcggcct 60
 cagtgagcga gcgagcgcg 80

 <210> 115
 <211> 79
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 115
gcgcgctcgc tcgctcactg aggcgcccgg gcgtcgggcg acctttggtc gcccggcctc 60
agtgagcgag cgagcgcgc 79

<210> 116
<211> 79
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 116
gcgcgctcgc tcgctcactg aggcggggcg accaaaggct gcccgacgcc cgggcgcctc 60
agtgagcgag cgagcgcgc 79

<210> 117
<211> 5
<212> PRT
<213> Influenza virus

<400> 117
Asp Arg Leu Arg Arg
1 5

<210> 118
<211> 7
<212> PRT
<213> Influenza virus

<400> 118
Pro Lys Gln Lys Lys Arg Lys
1 5

<210> 119
<211> 10
<212> PRT
<213> Hepatitis delta virus

<400> 119
Arg Lys Leu Lys Lys Lys Ile Lys Lys Leu
1 5 10

<210> 120
<211> 10
<212> PRT
<213> Mus musculus

<400> 120
Arg Glu Lys Lys Lys Phe Leu Lys Arg Arg
1 5 10

<210> 121
<211> 20
<212> PRT
<213> Homo sapiens

<400> 121

Lys Arg Lys Gly Asp Glu Val Asp Gly Val Asp Glu Val Ala Lys Lys
1 5 10 15

Lys Ser Lys Lys
20

<210> 122
<211> 17
<212> PRT
<213> Homo sapiens

<400> 122
Arg Lys Cys Leu Gln Ala Gly Met Asn Leu Glu Ala Arg Lys Thr Lys
1 5 10 15

Lys

<210> 123
<211> 8
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 123
gtttaaac 8

<210> 124
<211> 8
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 124
ttaattaa 8

<210> 125
<211> 141
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 125
aataaacgat aacgccgttg gtggcgtgag gcatgtaaaa ggttacatca ttatcttgtt 60
cgccatccgg ttggtataaa tagacgttca tgttggtttt tgtttcagtt gcaagttggc 120
tgcggcgcgc gcagcacctt t 141

<210> 126

<211> 317
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 126
ggtgtggaaa gtccccaggc tccccagcag gcagaagtat gcaaagcatg catctcaatt 60
agtcagcaac cagggtgtgga aagtccccag gctccccagc aggcagaagt atgcaaagca 120
tgcatctcaa ttagtcagca accatagtcc cgcccctaac tccgcccatac ccgcccctaa 180
ctccgcccag ttccgcccatac tctccgcccc atggctgact aattttttttt atttatgcag 240
aggccgaggc cgccctcgcc tctgagctat tccagaagta gtgaggaggc ttttttggag 300
gcctaggctt ttgcaaa 317

<210> 127
<211> 72
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 127
gcgcgctcgc tcgctcactg aggccgggag accaaaggct gcccgacggc ctcaagtgcg 60
gagcgagcgc gc 72

<210> 128
<211> 60
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 128
gagacagaca cactcctgct atgggtactg ctgctctggg ttccaggctc cactggtgac 60

<210> 129
<211> 1260
<212> DNA
<213> Adeno-associated virus - 2

<400> 129
atggagctgg tcgggtggct cgtggacaag gggattacct cggagaagca gtggatccag 60
gaggaccagg cctcatacat ctctttcaat gcggcctcca actcgcggtc ccaaatcaag 120
gctgccttgg acaatgcggg aaagattatg agcctgacta aaaccgcccc cgactacctg 180
gtgggccagc agcccgtgga ggacatttcc agcaatcgga ttataaaaat tttggaacta 240
aacgggtacg atccccaata tgcggcttcc gtctttctgg gatgggccac gaaaaagttc 300

ggcaagagga acaccatctg gctgtttggg cctgcaacta ccgggaagac caacatcgcg	360
gaggccatag cccacactgt gcccttctac ggggtgcgtaa actggaccaa tgagaacttt	420
cccttcaacg actgtgtcga caagatgggt atctgggtggg aggaggggaa gatgaccgcc	480
aaggtcgtgg agtcggccaa agccattctc ggaggaagca aggtgcgcgt ggaccagaaa	540
tgcaagtcct cggcccagat agacccgact cccgtgatcg tcacctcaa caccaacatg	600
tgcgccgtga ttgacgggaa ctcaacgacc ttcgaacacc agcagccgtt gcaagaccgg	660
atgttcaaat ttgaactcac ccgccgtctg gatcatgact ttgggaaggt caccaagcag	720
gaagtcaaag actttttccg gtgggcaaag gatcacgtgg ttgaggtgga gcatgaattc	780
tacgtcaaaa aggggtggagc caagaaaaga cccgccccca gtgacgcaga tataagtgag	840
cccaaacggg tgcgcgagtc agttgcgcag ccatcgacgt cagacgcgga agcttcgatc	900
aactacgcag acaggtacca aaacaaatgt tctcgtcacg tgggcatgaa tctgatgctg	960
tttccctgca gacaatgcga gagaatgaat cagaattcaa atatctgctt cactcacgga	1020
cagaaagact gtttagagtg ctttcccgtg tcagaatctc aaccgtttc tgtcgtcaaa	1080
aaggcgtatc agaaactgtg ctacattcat catatcatgg gaaaggtgcc agacgcttgc	1140
actgcctgcg atctgggtcaa tgtggatttg gatgactgca tctttgaaca ataaatgatt	1200
taaatcaggt atggctgccg atggttatct tccagattgg ctcgaggaca ctctctctga	1260

<210> 130

<211> 1932

<212> DNA

<213> Adeno-associated virus - 2

<400> 130

atgccggggt ttacgagat tgtgattaag gtccccagcg accttgacga gcatctgccc	60
ggcatttctg acagctttgt gaactgggtg gccgagaagg aatgggagtt gccgccagat	120
tctgacatgg atctgaatct gattgagcag gcacccctga ccgtggccga gaagctgcag	180
cgcgactttc tgacggaatg gcgccgtgtg agtaaggccc cggaggccct tttctttgtg	240
caatttgaga agggagagag ctacttcac atgcacgtgc tcgtggaaac caccggggtg	300
aaatccatgg ttttgggacg tttcctgagt cagattcgcg aaaaactgat tcagagaatt	360
taccgcggga tcgagccgac ttgccaac tggttcgcgg tcacaaagac cagaaatggc	420
gccggaggcg ggaacaaggt ggtggatgag tgctacatcc ccaattactt gctcccaaa	480
accagcctg agctccagtg ggcgtggact aatatggaac agtatttaag cgcctgtttg	540
aatctcacgg agcgtaaacg gttggtggcg cagcatctga cgcacgtgtc gcagacgcag	600
gagcagaaca aagagaatca gaatcccaat tctgatgcgc cggtgatcag atcaaaaact	660
tcagccaggt acatggagct ggtcgggtgg ctcgtggaca aggggattac ctcggagaag	720
cagtggatcc aggaggacca ggcctcatac atctccttca atgcggcctc caactcgcgg	780
tcccaaatca aggctgcctt ggacaatgcg ggaaagatta tgagcctgac taaaaccgcc	840
cccgactacc tggtagggcca gcagcccgtg gaggacattt ccagcaatcg gatttataaa	900

atthttggaac taaacgggta cgatcccaa tatgaggctt ccgtctttct gggatgggcc	960
acgaaaaagt tcggcaagag gaacaccatc tggctgtttg ggcttgcaac taccgggaag	1020
accaacatcg cggaggccat agcccacact gtgcccttct acgggtgcgt aaactggacc	1080
aatgagaact ttcccttcaa cgactgtgtc gacaagatgg tgatctgggtg ggaggagggg	1140
aagatgaccg ccaaggtcgt ggagtcggcc aaagccattc tcggaggaag caaggtgcgc	1200
gtggaccaga aatgcaagtc ctcggcccag atagaccga ctcccgtgat cgtcacctcc	1260
aacaccaaca tgtgcgccgt gattgacggg aactcaacga ctttgaaca ccagcagccg	1320
ttgcaagacc ggatgttcaa atttgaactc acccgccgtc tggatcatga ctttgggaag	1380
gtcaccaagc aggaagtcaa agactttttc cggatgggcaa aggatcacgt ggttgaggtg	1440
gagcatgaat tctacgtcaa aaagggtgga gccaagaaa gaccgcccc cagtgcgca	1500
gatataagtg agcccaaacg ggtgcgcgag tcagttgcgc agccatcgac gtcagacgcg	1560
gaagcttcga tcaactacgc agacaggtac caaacaat gttctcgtca cgtgggcatg	1620
aatctgatgc tgtttcctg cagacaatgc gagagaatga atcagaattc aaatatctgc	1680
ttcactcacg gacagaaaga ctgtttagag tgctttcccg tgtcagaatc tcaaccgtt	1740
tctgtcgtca aaaaggcgta tcagaaactg tgctacattc atcatatcat gggaaagggtg	1800
ccagacgctt gactgcctg cgatctggtc aatgtggatt tggatgactg catctttgaa	1860
caataaatga tttaaatcag gtatggctgc cgatggttat cttccagatt ggctcgagga	1920
cactctctct ga	1932

<210> 131

<211> 1876

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 131

cgagccacc atggcggggt ttacgagat tgtgattaag gtcccagcg acctgacgg	60
gcatctgccc ggcatcttctg acagctttgt gaactgggtg gccgagaagg aatgggagtt	120
gccgccagat tctgacatgg atctgaatct gattgagcag gcaccctga ccgtggccga	180
gaagctgcag cgcgactttc tgacggaatg gcgccgtgtg agtaaggccc cggaggccct	240
tttctttgtg caatttgaga agggagagag ctacttcac atgcacgtgc tcgtggaac	300
caccggggtg aaatccatgg ttttgggacg tttcctgagt cagattcgcg aaaaactgat	360
tcagagaatt taccgcggga tcgagccgac ttgccaaac tggttcgcg tcacaaagac	420
cagaaatggc gccggaggcg ggaacaaggt ggtggatgag tgctacatcc ccaattactt	480
gctcccaaa acccagcctg agctccagtg ggcgtggact aatatggaac agtatttaag	540
cgctgtttg aatctcacgg agcgtaaacg gttggtggcg cagcatctga cgcacgtgtc	600
gcagacgcag gagcagaaca aagagaatca gaatcccaat tctgatgcgc cggatgatcag	660

atcaaaaact tcagccaggt acatggagct ggtcgggtgg ctcgtggaca aggggattac	720
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caactcgcgg tcccaaatca aggctgcctt ggacaatgcg ggaaagatta tgagcctgac	840
taaaaccgcc ccgcactacc tgggtgggcca gcagcccgtg gaggacattt ccagcaatcg	900
gatttataaa attttgaac taaacgggta cgatcccaa tatgcggctt ccgtctttct	960
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taccggaag accaacatcg cggaggccat agcccacact gtgcccttct acgggtgcgt	1080
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ggaggagggg aagatgaccg ccaaggctgt ggagtcggcc aaagccattc tcggaggaag	1200
caaggtgcgc gtggaccaga aatgcaagtc ctcggcccag atagaccga ctcccgtgat	1260
cgtcacctcc aacaccaaca tgtgcgccgt gattgacggg aactcaacga ctttcgaaca	1320
ccagcagccg ttgcaagacc ggatgttcaa atttgaactc acccgccgtc tggatcatga	1380
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cagtgcgcga gatataagtg agcccaaacg ggtgcgcgag tcagttgcgc agccatcgac	1560
gtcagacgcg gaagcttcga tcaactacgc agacaggtac caaaacaaat gttctcgtca	1620
cgtgggcatg aatctgatgc tgtttccctg cagacaatgc gagagaatga atcagaattc	1680
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tcaaccggtt tctgtcgtca aaaaggcgta tcagaaactg tgctacattc atcatatcat	1800
gggaaaggtg ccagacgctt gcactgcctg cgatctggtc aatgtggatt tggatgactg	1860
catctttgaa caataa	1876

<210> 132

<211> 1194

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 132

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gaggaccagg cctcatacat ctcttcaat gcggcctcca actcgcggtc ccaaatcaag	120
gctgccttgg acaatgcggg aaagattatg agcctgacta aaaccgcccc cgactacctg	180
gtgggccagc agcccgtgga ggacatttcc agcaatcgga tttataaaat tttggaacta	240
aacgggtacg atccccaata tgcggcttcc gtctttctgg gatgggccac gaaaaagttc	300
ggcaagagga acaccatctg gctgtttggg cctgcaacta ccgggaagac caacatcgcg	360
gaggccatag ccacactgt gcccttctac gggcgcgtaa actggaccaa tgagaacttt	420

cccttcaacg actgtgtcga caagatggtg atctggtggg aggaggggaa gatgaccgcc	480
aaggctcgtgg agtcggccaa agccattctc ggaggaagca aggtgcgcgt ggaccagaaa	540
tgcaagtcct cggcccagat agacccgact cccgtgatcg tcacctcaa caccaacatg	600
tgcgccgtga ttgacgggaa ctcaacgacc ttcgaacacc agcagccgtt gcaagaccgg	660
atgttcaa at ttgaactcac ccgccgtctg gatcatgact ttgggaaggt caccaagcag	720
gaagtcaaag actttttccg gtggggcaaag gatcacgtgg ttgaggtgga gcatgaattc	780
tacgtcaaaa aggggtggagc caagaaaaga cccgccccca gtgacgcaga tataagtgag	840
cccaaacggg tgcgcgagtc agttgcgcag ccatcgacgt cagacgcgga agcttcgatc	900
aactacgcag accgctacca aaacaaatgt tctcgtcacg tgggcatgaa tctgatgctg	960
tttccttgca gacaatgcga gagaatgaat cagaattcaa atatctgctt cactcacgga	1020
cagaaagact gtttagagtg ctttcccgtg tcagaatctc aaccgtttc tgcgtcaaa	1080
aaggcgtatc agaaactgtg ctacattcat catatcatgg gaaaggtgcc agacgcttgc	1140
actgcctgcg atctggtcaa tgtggatttg gatgactgca tctttgaaca ataa	1194

<210> 133

<211> 1876

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 133

cgcagccacc atggcggggt tttacgagat tgtgattaag gtccccagcg accttgacgg	60
gcatctgccc ggcatcttctg acagctttgt gaactgggtg gccgagaagg aatgggagtt	120
gccgccagat tctgacatgg atctgaatct gattgagcag gcacccctga ccgtggccga	180
gaagctgcag cgcgactttc tgacggaatg gcgccgtgtg agtaaggccc cggaggccct	240
tttctttgtg caatttgaga agggagagag ctacttcac atgcacgtgc tcgtggaaac	300
caccggggtg aaatccatgg ttttgggacg tttcctgagt cagattcgcg aaaaactgat	360
tcagagaatt taccgcggga tcgagccgac tttgcaaac tggttcgcgg tcacaaagac	420
cagaaatggc gccggaggcg ggaacaaggt ggtggatgag tgctacatcc ccaattactt	480
gctcccaaaa acccagcctg agctccagt ggctgggact aatatggaac agtatttaag	540
cgcctgtttg aatctcacgg agcgtaaacg gttggtggcg cagcatctga cgcacgtgtc	600
gcagacgcag gagcagaaca aagagaatca gaatcccaat tctgatgcgc cggatgatcag	660
atcaaaaaact tcagccaggt acatggagct ggtcgggtgg ctctgggaca aggggattac	720
ctcggagaag cagtggatcc aggaggacca ggccctatac atctccttca atgcggcctc	780
caactcgcgg tcccaaatca aggctgcctt ggacaatgcg ggaaagatta tgagcctgac	840
taaaaccgcc cccgactacc tgggtgggcca gcagcccgtg gaggacattt ccagcaatcg	900
gatttataaa attttgaac taaacgggta cgatcccaa tatgcggctt ccgtctttct	960

gggatgggcc acgaaaaagt tcggcaagag gaacaccatc tggctgtttg ggcctgcaac	1020
taccgggaag accaacaatcg cggaggccat agcccacact gtgcccttct acgggtgcgt	1080
aaactggacc aatgagaact ttcccttcaa cgactgtgtc gacaagatgg tgatctggtg	1140
ggaggagggg aagatgaccg ccaaggtcgt ggagtcggcc aaagccattc tcggaggaag	1200
caaggtgcgc gtggaccaga aatgcaagtc ctcggcccag atagaccga ctcccgtgat	1260
cgtcacctcc aacaccaaca tgtgcgccgt gattgacggg aactcaacga ccttcgaaca	1320
ccagcagccg ttgcaagacc ggatgttcaa atttgaactc acccgccgtc tggatcatga	1380
ctttgggaag gtcaccaagc aggaagtcaa agactttttc cggtagggcaa aggatcacgt	1440
ggttgagggtg gagcatgaat tctacgtcaa aaagggtgga gccaagaaaa gaccgcgcc	1500
cagtgcgca gatataagtg agcccaaacg ggtgcgcgag tcagttgcgc agccatcgac	1560
gtcagacgcg gaagcttcga tcaactacgc agacaggtac caaaacaaat gttctcgtca	1620
cgtgggcatg aatctgatgc tgtttccctg cagacaatgc gagagaatga atcagaattc	1680
aaatatctgc ttactcacg gacagaaaga ctgttttagag tgctttcccg tgtcagaatc	1740
tcaaccggtt tctgtcgtca aaaaggcgta tcagaaactg tgctacattc atcatatcat	1800
gggaaagggtg ccagacgctt gcactgcctg cgatctggtc aatgtggatt tggatgactg	1860
catctttgaa caataa	1876

<210> 134
 <211> 51
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 134	
ctaggccgcc cgggcaaagc ccgggcgtcg ggcgaccttt ggtagcccg c	51

<210> 135
 <211> 65
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 135	
ctaggactga ggccgcccgg gcaaagcccg ggcgtcgggc gacctttggt cgcccgccct	60
cagtc	65

<210> 136
 <211> 67
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 136
 ggactgaggc cgcccgggca aagcccgggc gtcgggcgac ctttggtcgc ccggcctcag 60
 tcctgca 67

<210> 137
 <211> 41
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 137
 gtgcgggcga ccaaaggctc gccgacgccc gggcgccactc a 41

<210> 138
 <211> 56
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 138
 ggactgaggc cgggcgacca aaggctgccc gacgcccggg cggcctcagt cctgca 56

<210> 139
 <211> 54
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 139
 ctaggactga ggccgcccgg gcgtcgggcg acctttggtc gcccggcctc agtc 54

<210> 140
 <211> 48
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 140
 ggactgaggc cgggcgacca aaggctgccc gacggcctca gtcctgca 48

<210> 141
 <211> 46

<212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 141
 ctaggactga ggccgtcggg cgacctttgg tcgcccggcc tcagtc 46

 <210> 142
 <211> 67
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 142
 ggactgaggc ccgggcgacc aaaggtcgcc cgacgcccgg gctttgcccg ggcgcctcag 60
 tcctgca 67

 <210> 143
 <211> 47
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 143
 atacctaggc acgcgtgtta ctagttatta atagtaatca attacgg 47

 <210> 144
 <211> 29
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 144
 atacctaggg gccgcacgcg tgttactag 29

 <210> 145
 <211> 42
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 145
 atacactcag tgcttcgagg cacgtggtcc ggagatccag ac 42

<210> 146
<211> 3754
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 146
cctaggtgag cgagcgagcg cgcagagagg gaggggccaa ctccatcact aggggttcct 60
tgtagttaat gattaacccg ccatgctact tatcgcgcc gctcaatatt ggccattagc 120
catattattc attggttata tagcataaat caatattggc tattggccat tgcatacgtt 180
gtatctatat cataatatgt acatttatat tggctcatgt ccaatatgac cgccatgttg 240
gcattgatta ttgactagtt attaatagta atcaattacg gggtcattag ttcataagccc 300
atatatggag ttccgcgtta cataacttac ggtaaattggc ccgcctggct gaccgcccac 360
cgacccccgc ccattgacgt caataatgac gtatgttccc atagtaacgc caatagggac 420
tttccattga cgtcaatggg tggagtattt acggtaaact gcccacttgg cagtacatca 480
agtgtatcat atgccaagtc cgccccctat tgacgtcaat gacggtaaatt ggcccgctg 540
gcattatgcc cagtacatga ccttacggga ctttctact tggcagtaca tctacgtatt 600
agtcacgct attaccatgg tcgaggtgag cccacgctt tgcttcactc tccccatctc 660
ccccccctcc ccacccccaa ttttgtattt atttattttt taattatttt gtgcagcgat 720
gggggcgggg gggggggggg ggcgcgcgcc aggcggggcg gggcggggcg aggggcgggg 780
cggggcgagg cggagaggtg cggcggcagc caatcagagc ggcgcgctcc gaaagtttcc 840
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agtcgctgag acgctgcctt cgccccgtgc cccgctccgc cgccgctcg cgccggccgc 960
cccggctctg actgaccgag ttactccac aggtgagcgg gcgggacggc ctttctctc 1020
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gccttgaggg gctccgggag ggccctttgt gcggggggga gcggctcggg ggggtgcgtgc 1140
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gcggtgcccc gcggtgcggg gggggctgag aggggaacaa aggtgcgtg cggggtgtgt 1320
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ccccctccc gagttgctga gcacggccc gcttcgggtg cggggctccg tacggggcgt 1440
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ggctgtcgag gcgcggcgag ccgcagccat tgccttttat ggtaatcgt cgagagggcg 1620
cagggacttc ctttgtccca aatctgtgc gagccgaaat ctgggaggcg ccgccgacc 1680
ccctctagcg ggcgcggggc gaagcggtgc ggcgcggga ggaaggaaat gggcggggag 1740

ggccttcgtg cgtcgccgcg ccgccgtccc cttctccctc tccagcctcg gggctgtccg	1800
cgggggggacg gctgccttcg ggggggacgg ggcagggcgg ggttcggctt ctggcgtgtg	1860
accggcggct ctagagcctc tgctaaccat gttttagcct tcttcttttt cctacagctc	1920
ctgggcaacg tgtctggttat tgtctgtctc catcatttgt cgacagaatt cctcgaagat	1980
ccgaaggggt tcaagcttgg cattccggtc ctgttggtta agccagtta aacgccgcca	2040
ccatggtgag caagggcgag gagctgttca ccgggggtgg gcccattcctg gtcgagctgg	2100
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acaagctgga gtacaactac aacagccaca acgtctatat catggccgac aagcagaaga	2520
acggcatcaa ggtgaacttc aagatccgcc acaacatcga ggacggcagc gtgcagctcg	2580
ccgaccacta ccagcagaac acccccattc gcgacggccc cgtgctgctg cccgacaacc	2640
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aattaattaa gagcatctta ccgccattta ttcccatatt tgttctgttt ttcttgattt	2820
gggtatacat ttaaatgtta ataaaacaaa atgggtggggc aatcatttac atttttaggg	2880
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gactttcgct tccccctcc cgatgccac ggcagaactc atcgccgcct gccttgcccg	3300
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tcgcccgcgc cccgggcggc ctgagtcctg cagg	3754

<210> 147
<211> 8418
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 147
ggcagctgcg cgctcgctcg ctcacctagg ccgcccgggc aaagcccggg cgtcgggcga 60
cctttggtcg cccggcctag gtgagcgagc gagcgcgag agaggagtg gccaaactcca 120
tcactagggg ttccttgtag ttaatgatta acccgccatg ctacttatcg cggccgctca 180
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gccattgcat acgttgatc tatatcataa tatgtacatt tatattggct catgtccaat 300
atgaccgcca tgttggcatt gattattgac tagttattaa tagtaatcaa ttacggggtc 360
attagttcat agcccatata tggagtccg cgttacataa cttacggtaa atggcccgcc 420
tggctgaccg cccaacgacc cccgcccatt gacgtcaata atgacgatg ttcccatagt 480
aacgccaata gggactttcc attgacgtca atgggtggag tatttacggg aaactgcca 540
cttggcagta catcaagtgt atcatatgcc aagtcgccc cctattgacg tcaatgacgg 600
taaatggccc gcctggcatt atgcccagta catgacctta cgggactttc ctacttggca 660
gtacatctac gtattagtca tcgctattac catggtcgag gtgagcccca cgttctgctt 720
cactctcccc atctcccccc cctccccacc cccaattttg tatttattta ttttttaatt 780
attttgtgca gcgatggggg cggggggggg gggggggcgc gcgccaggcg gggcggggcg 840
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cgcgcgggcg gcgggagtcg ctgcgacgct gccttcgccc cgtgccccgc tccgccgccg 1020
cctcgcgccg cccgccccgg ctctgactga ccgcgttact cccacaggag agcgggcggg 1080
acggcccttc tcctccgggc tgtaattagc gcttggttta atgacggctt gtttcttttc 1140
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gcgtgcgggg tgtgtgcgtg ggggggtgag cagggggtgt gggcgcggcg gtcgggctgt 1440
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gtgccgggcg gggcggggcc gcctcgggcc ggggagggct cgggggaggg gcgcggcggc 1620
ccccggagcg ccggcggtg tcgaggcgcg gcgagccgca gccattgcct tttatggtaa 1680
tcgtgcgaga gggcgaggg acttcctttg tcccaaatct gtgcggagcc gaaatctggg 1740

aggcgccgcc gcacccccctc tagcggggcgc ggggcgaagc ggtgcggcgc cggcaggaag	1800
gaaatgggcg gggagggcct tcgtgcgtcg ccgcgccgcc gtccccttct ccctctccag	1860
cctcggggct gtccgcgggg ggacggctgc cttcgggggg gacggggcag ggcggggttc	1920
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gaattcctcg aagatccgaa ggggttcaag cttaggcattc cggctactgtt ggtaaagcca	2100
gtttaaacgc cgccaccatg gtgagcaagg gcgaggagct gttcaccggg gtggtgcca	2160
tcctggtcga gctggacggc gacgtaaacg gccacaagtt cagcgtgtcc ggcgagggcg	2220
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ccgtgccctg gcccaccctc gtgaccaccc tgacctacgg cgtgcagtgc ttcagccgct	2340
accccgacca catgaagcag cacgacttct tcaagtccgc catgcccga ggctacgtcc	2400
aggagcgcac catcttcttc aaggacgacg gcaactacaa gaccgcgcc gaggtgaagt	2460
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 <213> Artificial Sequence

<220>
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gagtaggtgt cattctattc tgggggggtgg ggtggggcag gacagcaagg gggaggattg	180
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<210> 149
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 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 149	
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<210> 150

<211> 1326

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 150

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<210> 151
 <211> 573
 <212> DNA
 <213> Mus musculus

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<210> 152
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<400> 152 acgatcggga actggcatct tcaggagta gcttaggtca gtgaagagaa gaacaaaaag	60
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<210> 153
 <211> 1350
 <212> DNA
 <213> Homo sapiens

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<210> 154
 <211> 1223
 <212> DNA
 <213> Homo sapiens

<400> 154	
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gatccactgg ggagcaggaa atatctgtgg gcttgtgaca cggactcaag tgggctggtg	240
accagtcag agttgtgcac atggcttagt tttcatacac agcctgggct gggggtgggg	300
tgggagaggt cttttttaa aggaagttac tgttatagag ggtctaagat tcatccattt	360
at ttggcatc tgtttaaagt agattagatc ttttaagccc atcaattata gaaagccaaa	420
tcaaaatatg ttgatgaaaa atagcaacct ttttatctcc cttcacatg catcaagtta	480
ttgacaaact ctcccttcac tccgaaagtt cttatgtat atttaaaaga aagcctcaga	540
gaattgtga ttcttgagtt tagtgatctg aacagaaata ccaaattat ttcagaaatg	600
tacaactttt tacctagtac aaggcaacat ataggttgta aatgtgttta aaacaggtct	660
ttgtcttgct atggggagaa aagacatgaa tatgattagt aaagaaatga cacttttcat	720
gtgtgatttc ccctccaagg tatggttaat aagtttact gacttagaac caggcgagag	780
acttgtggcc tgggagagct ggggaagctt cttaaagag aaggaatttg agttggatca	840
tctattgctg gcaaagacag aagcctcact gcaagcactg catgggcaag cttggctgta	900
gaaggagaca gagctggttg ggaagacatg gggaggaagg acaaggctag atcatgaaga	960
accttgacgg cattgctccg tctaagtcac gagctgagca gggagatcct ggttggtgtt	1020
gcagaaggtt tactctgtgg ccaaaggagg gtcaggaagg atgagcattt agggcaagga	1080

gaccaccaac agccctcagg tcagggtgag gatggcctct gctaagctca aggcgtgagg	1140
atgggaagga gggaggtatt cgtaaggatg ggaaggaggg aggtattcgt gcagcatatg	1200
aggatgcaga gtcagcagaa ctg	1223

<210> 155
 <211> 215
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 155	
gaacgctgac gtcacatcaacc cgctccaagg aatcgcgggc ccagtgtcac taggcgggaa	60
cacccagcgc gctgcgccc tggcaggaag atggctgtga gggacagggg agtggcgccc	120
tgcaatatatt gcatgtcgct atgtgttctg ggaaatcacc ataaacgtga aatgtctttg	180
gatttgggaa tcttataagt tctgtatgag accac	215

<210> 156
 <211> 141
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 156	
cctgcaggca gctgcgcgct cgctcgctca cctaggccgc ccgggcaaag cccgggcgtc	60
gggcgacctt tggtcgcccg gcctaggtga gcgagcgagc gcgcagagag ggagtggcca	120
actccatcac taggggttcc t	141

<210> 157
 <211> 19
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 157	
gcgcgctcgc tcgctcacc	19

<210> 158
 <211> 22
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 158
 ctaggtgagc gagcgagcgc gc 22

<210> 159
 <211> 75
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 159
 cctgcaggac tgaggccgcc cgggcaaagc ccgggcgtcg ggcgacctt ggtcgcccgg 60
 cctcagtcct gcagg 75

<210> 160
 <211> 130
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 160
 aggaaccctt agtgatggag ttggccactc cctctctgcg cgctcgctcg ctactgagt 60
 gcgggcgacc aaaggtcgcc cgacgcccgg gcgcactcag tgagcgagcg agcgcgagc 120
 tgcctgcagg 130

<210> 161
 <211> 142
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 161
 cctgcaggca gctgcgcgt cgctcgctcc ctaggactga ggccgcccgg gcgtcgggcg 60
 acctttggtc gcccggcctc agtcctaggg agcgagcgag cgcgagaga gggagtggcc 120
 aactccatca ctaggggttc ct 142

<210> 162
 <211> 80
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 162
 gcgcgctcg tcgctcactg agtgcgggcg accaaaggtc gcccgacgcc cgggcgcact 60

cagtgagcga gcgagcgcgc 80

<210> 163
<211> 21
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 163
gcgcgctcgc tcgctcactg a 21

<210> 164
<211> 18
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 164
gtgagcgagc gagcgcgc 18

<210> 165
<211> 89
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 165
gcgcgctcgc tcgctcactg aggccgcccg ggcaaagccc gggcgtcggg cgactttgtc 60

gcccggcctc agtgagcgag cgagcgcgc 89

<210> 166
<211> 89
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 166
gcgcgctcgc tcgctcactg aggccgggcg acaaagtcgc ccgacgcccg ggctttgccc 60

gggcggcctc agtgagcgag cgagcgcgc 89

<210> 167
<211> 87
<212> DNA
<213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 167
 gcgcgctcgc tcgctcactg aggccgccc ggcaaagccc gggcgtcggg cgattttcgc 60
 ccggcctcag tgagcgagcg agcgcgc 87

<210> 168
 <211> 87
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 168
 gcgcgctcgc tcgctcactg aggccgggcg aaaatcgccc gacgcccggg ctttgcccgg 60
 gcggcctcag tgagcgagcg agcgcgc 87

<210> 169
 <211> 85
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 169
 gcgcgctcgc tcgctcactg aggccgccc ggcaaagccc gggcgtcggg cgtttcgccc 60
 ggcctcagtg agcgagcgag cgcgc 85

<210> 170
 <211> 85
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 170
 gcgcgctcgc tcgctcactg aggccgggcg aaacgcccga cgcccgggct ttgcccgggc 60
 ggcctcagtg agcgagcgag cgcgc 85

<210> 171
 <211> 89
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 171
 gcgcgctcgc tcgctcactg aggccgcccg ggaaacccgg gcgtcgggcg acctttggtc 60
 gcccggcctc agtgagcgag cgagcgcgc 89

<210> 172
 <211> 89
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 172
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgacgcc cgggtttccc 60
 gggcggcctc agtgagcgag cgagcgcgc 89

<210> 173
 <211> 87
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 173
 gcgcgctcgc tcgctcactg aggccgcccg gaaaccgggc gtcgggcgac ctttggtcgc 60
 ccggcctcag tgagcgagcg agcgcgc 87

<210> 174
 <211> 87
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 174
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgacgcc cggtttcgg 60
 gcggcctcag tgagcgagcg agcgcgc 87

<210> 175
 <211> 85
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 175
 gcgcgctcgc tcgctcactg aggccgcccg aaacgggcgt cgggcgacct ttggtcgccc 60
 ggcctcagtg agcgagcgag cgcgc 85

<210> 176
 <211> 85
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 176
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgcgcc cgtttcgggc 60
 ggctcagtgc agcgagcgag cgcgc 85

 <210> 177
 <211> 83
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 177
 gcgcgctcgc tcgctcactg aggccgccc aaggcgctcg ggcgacctt ggctgcccgg 60
 cctcagtgcg cgagcgagcg cgc 83

 <210> 178
 <211> 83
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 178
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgcgcc ctttgggcgg 60
 cctcagtgcg cgagcgagcg cgc 83

 <210> 179
 <211> 81
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 179
 gcgcgctcgc tcgctcactg aggccgccc aaggcgctgg cgacctttgg tcgcccggcc 60
 tcagtgcgagc agcgagcgcg c 81

 <210> 180
 <211> 81
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 180
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgcgccc ttggcgggcc 60
 tcagtgcgagc agcgagcgcg c 81

<210> 181
 <211> 79
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 181
 gcgcgctcgc tcgctcactg aggccgcaaa gcgtcgggcg acctttggtc gcccggcctc 60
 agtgagcgag cgagcgcgcg 79

<210> 182
 <211> 79
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 182
 gcgcgctcgc tcgctcactg aggccgggcg accaaaggtc gcccgcgctc ttgcggcctc 60
 agtgagcgag cgagcgcgcg 79

<210> 183
 <211> 81
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 183
 ctgcgcgctc gctcgctcac tgaggccgaa acgtcgggcg acctttggtc gcccggcctc 60
 agtgagcgag cgagcgcgca g 81

<210> 184
 <211> 81
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 184
 ctgcgctc gctcgctcac tgaggccggg cgaccaaagg tcgcccgcg tttcggcctc 60
 agtgagcgag cgagcgcgca g 81

<210> 185
 <211> 72
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 185
 gcgcgctcgc tcgctcactg aggccggggc accaaaggct gcccgacggc ctcaagtgc 60
 gagcgagcgc gc 72

<210> 186
 <211> 80
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 186
 gcgcgctcgc tcgctcactg aggccggggc accaaaggct gcccgacgcc cgggcggcct 60
 cagtgcgca gcgagcgcg 80

<210> 187
 <211> 79
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 187
 gcgcgctcgc tcgctcactg aggcgcccgg gcgtcggggc acctttggct gcccggcctc 60
 agtgagcgag cgagcgcg 79

<210> 188
 <211> 48
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 188
 ggagtcaaag ttctgtttgc cctgatctgc atcgtgtgg ccgaggcc 48

<210> 189
<211> 99
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 189
attcatacca acttgaagaa aaagttcagc ctcttcatcc tggcttttct cctgttcgca 60
gtcatctgtg ttggaagaa agggagcgac tatgaggcc 99

<210> 190
<211> 588
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 190
gccctctcc cccccccc cctaacttta ctggccgaag ccgcttgga taaggccggt 60
gtgcgtttgt ctatatgtta tttccacca tattgccgtc ttttggaat gtgagggcc 120
ggaaacctgg ccctgtcttc ttgacgagca ttcctagggg tctttcccct ctcgcaaag 180
gaatgcaagg tctgttgaat gtcgtgaagg aagcagttcc tctggaagct tcttgaagac 240
aaacaacgtc ttagcgcacc ctttgcaggc agcggaaacc cccacctggc gacaggtgcc 300
tctgcggcca aaagccacgt gtataagata cacctgcaaa ggcggcacaa cccagtgcc 360
acgttgtgag ttggatagtt gtggaaagag tcaaatggct ctctcaagc gtattcaaca 420
aggggctgaa ggatgccag aaggtagccc attgtatggg atctgatctg gggcctcggt 480
gcacatgctt tacatgtgtt tagtcgaggt taaaaaacg tctaggcccc ccgaaccacg 540
gggacgtggt tttcctttga aaaacacgat gataatatgg ccacaacc 588

<210> 191

<400> 191
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<210> 192

<400> 192
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<210> 193

<400> 193
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<210> 194

<400> 194

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<210> 195

<400> 195
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<210> 196

<400> 196
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<210> 197

<400> 197
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<210> 198

<400> 198
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<210> 199

<400> 199
000

<210> 200

<211> 278

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 200

tcgaggtgag cccacgttc tgcttcactc tccccatctc cccccctcc ccaccccaa 60

ttttgtattt atttattttt taattatttt gtgcagcgat gggggcgggg gggggggggg 120

ggcgcgcgcc aggcggggcg gggcggggcg aggggcgggg cggggcgagg cggagaggtg 180

cggcggcagc caatcagagc ggcgcgctcc gaaagtttcc ttttatggcg aggcggcggc 240

ggcggcggcc ctataaaaag cgaagcgcgc ggcgggcg 278

<210> 201

<211> 348

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 201

gatcttgcta ccagtgaac agccactaag gattctgcag tgagagcaga gggccagcta 60

agtgttactc tccagagac tgtctgactc acgccacccc ctccaccttg gacacaggac 120

gctgtggttt ctgagccagg tacaatgact cctttcggta agtgcagtgg aagctgtaca	180
ctgcccaggc aaagcgtccg ggacagcgtag gcgggcgact cagatcccag ccagtggact	240
tagccccgtg ttgctcctcc gataactggg gtgaccttgg ttaatatcca ccagcagcct	300
cccccgttgc ccctctggat ccactgctta aatacggacg aggacagg	348

<210> 202
 <211> 226
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 202 gtggagaaga gcatgcttga gggctgagtg cccctcagtg ggcagagagc acatggccca	60
cagtccttga gaagttgggg ggaggggtgg gcaattgaac tggcgcctag agaaggtggg	120
gcttgggtaa actgggaaag tgatgtggtg tactggctcc acctttttcc ccagggtggg	180
ggagaacat atataagtgc agtagtctct gtgaacattc aagctt	226

<210> 203
 <211> 225
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 203 ccgtctgtct gcacatttcg tagagcgagt gttccgatac tctaattctcc ctaggcaagg	60
ttcatatttg ttaggttac ttattctcct tttgttgact aagtcaataa tcagaatcag	120
caggtttgga gtcagcttgg cagggatcag cagcctgggt tggaaggagg gggataaaaa	180
gccccctcac caggagaagc cgtcacacag atccacaagc tcctg	225

<210> 204
 <211> 143
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 204 ggcgactcag atcccagcca gtggacttag cccctgtttg ctctccgat aactggggtg	60
accttggtta atattcacca gcagcctccc ccgttgcccc tctggatcca ctgcttaaat	120
acggacgagg acagggccct gtc	143

<210> 205

<211> 222
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 205
gtctgtctgc acatttcgta gagcgagtgt tccgatactc taatctccct aggcaagggt 60
catattgact taggttactt attctccttt tgttgactaa gtcaataatc agaatcagca 120
ggtttgaggt cagcttgga gggatcagca gcctgggttg gaaggagggg gtataaaagc 180
cccttcacca ggagaagccg tcacacagat ccacaagctc ct 222

<210> 206
<211> 223
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 206
gtctgtctgc acatttcgta gagcgagtgt tccgatactc taatctccct aggcaagggt 60
catatttgtg taggttactt attctccttt tgttgactaa gtcaataatc agaatcagca 120
ggtttgaggt cagcttgga gggatcagca gcctgggttg gaaggagggg gtataaaagc 180
cccttcacca ggagaagccg tcacacagat ccacaagctc ctg 223

<210> 207
<211> 3000
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 207
gttcaagcga ttctcctgcc tcagcctccc aagtagctgg gactacaggc acgtgccacc 60
atgcccggct aattttttgt atttttagta gaggaggagt ttcatcttgt tagctaggat 120
ggtctagatc tcctgacctc gtgatctgcc cgcctcagcc tcccaaagt ctgggattac 180
aggtgtgagc caccgtgccc ggccatattt tgatttaaaa ttagcaata atagataaaa 240
ttttcaatca actaagccct tgggccaggg aatgctattc cttaaaaagt gcttctatca 300
atatagcctc tgactcatta ctttgtaaat ttttaaattg tatttcattc ctgattaaca 360
ttcccacca gattattaat tatacaatct gttactgta gaacctcaa catgttggat 420
tgtactgtat ttgtctggaa gacacatttt taaaacattg taatcgctat aagagaagca 480
ctgggaaaga aaggagcttc tatgcctgca gtgcctgagg agccctttaa cagtgtgccc 540
cgcccctaag ctactcatgc agtcatcccc atcccagtta gtcaacttta ttccaaaaaa 600

cttgggtgttc caaatTTTTc cttctcaaag cccacagatc caaaattcat cagcagttcc	660
cacaaacgtt accctcacia tgaatccagc cttttttcac cctctccagt ggtaccatca	720
tagcccaagc cgccaccatt tctcaccccc ggtaaacagg ccaccctcct tctaccctta	780
tcctgctaga gtttgtttta tctacagtga tcagaaagat cagcctaata gataattctg	840
atcaccaccc tcctctactc acaaccggc cgtgtctccc cattgccctc agtgtagaag	900
tcaatgtccc tttgctgaaa tgcaacctta gtgaaacttt ccatgactaa cctcctttta	960
aattgcaacc tgggtccacc ttactcccc ttaccctact tctctttttt gcacagcact	1020
tattttacct tctaacatac tgtataatgt actcatgtat tgtaattatt gcttatcatc	1080
cctctttcag ttgcttatat ttttcatcaa tgtgtaccca gtgcctagga caatatctgt	1140
ctaggacaaa tgggtagtta tgtggctgta ggcaagccat ttaacctctc tgtacctcag	1200
ttactttatc tgtatccact ttgcgggtgt gtcatgagga ttaaatcaga tagcctatgt	1260
gtagcacctg gcagtgaatt tatcacctg tactgttaact gtctactttt ctgtctcctc	1320
cattggactg tcattcccag ggggttggga actgggattt cttcatttct gaggcataga	1380
agtatagcat agtggtagg agcatgactt ctggagccag agtacatggg tttgaatgct	1440
accactcaca agctgtgtgg ccatggagaa gttgcctaac ctctccgtgc ttcagtttca	1500
tcacccataa aatgaaggta agaatagtac ctgtatttaa aagcacctag aacagttcct	1560
ggcatatagt gtcagctgtc atctctgcat ctttgtacct gtcagagagg agtgtttatc	1620
aaaggggctt cttgtgcct gtttccaaac cagtcgacaa tataccaatt gtcacctaac	1680
acattcttgt ttgtgcagaa ctgagctcaa tgataacatt tttatagcaa ccctgatcaa	1740
gtttcttctc ataactcttt acactttgag gccctgcag gggccctcac tctccctaat	1800
aaacattaac ctgagtaggg tgtttgagct caccatggct acattctgat gtaaagagat	1860
atatcctata cctgggcaa atgtaaagag cctggaaaag tgtaggtta aaaacaaaac	1920
aaaataaata aatgaataaa tgccaggtgg ttatgagtgc tattgagaaa aatgaagcca	1980
agagggatat cagtgatgca ggtgggggta aagagcttac aacataaatg tgggtgtcca	2040
tatttaaacc tcattcaaca gggaagattg gagctgaaat gtgaaggagt tgtgggagt	2100
gaactacgtg gaaatctggg ggaaagggtgt tttgggtaaa agaaatagca agtggtgagg	2160
tccaggggca tgagtgtgct tgatatTTTA gggaagagta aggagaccag tataaccaga	2220
gtgagatgag actacagagg tcaggagaaa gggcatgcag accatgtggg atgctctagg	2280
acctaggcca tggtaaagat gtagggTTTT accctgatgg aggtcagaag ccattggagg	2340
attctgagaa gaggagtgc aggactcgct ttatagtttt aaattataac tataaattat	2400
agtttttaaa acaatagttg cctaacctca tgttatatgt aaaactacag ttttaaaaac	2460
tataaattcc tcatactggc agcagtgtga ggggcaaggg caaaagcaga gagactaaca	2520
ggttgctggg tactcttgct agtgcaagtg aattctagaa tcttcgacaa catccagaac	2580
ttctcttgct gctgccactc aggaagaggg ttggagtagg ctaggaatag gagcacaat	2640

taaagctcct gttcactttg acttctccat ccctctcctc ctttccttaa aggttctgat	2700
taaagcagac ttatgccctt actgctctca gaagtgaatg ggtaaagttt agcagcctcc	2760
cttttgctac ttcagttctt cctgtggctg cttccctactg ataaaaagga agcaatccta	2820
tcggttactg cttagtgtg agcacatcca gtgggtaaag ttccttaaaa tgctctgcaa	2880
agaaattggg acttttcatt aaatcagaaa ttttactttt ttcccctcct gggagctaaa	2940
gatattttag agaagaatta accttttgct tctccagttg aacatttgta gcaataagtc	3000

<210> 208
 <211> 205
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 208	
aatgactcct ttcgtaagt gcagtgaag ctgtacactg cccaggcaaa gcgtccgggc	60
agcgtaggcg ggcgactcag atcccagcca gtggacttag cccctgtttg ctcctccgat	120
aactggggtg accttggtta atattcacca gcagcctccc ccgttgcccc tctggatcca	180
ctgcttaaata acggacgagg acagg	205

<210> 209
 <211> 397
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 209	
gatcttgcta ccagtgaac agccactaag gattctgcag tgagagcaga gggccagcta	60
agtggacttc tcccagagac tgtctgactc acgccacccc ctccacctg gacacaggac	120
gctgtggttt ctgagccagg tacaatgact cttttcggtg agtgcagtgg aagctgtaca	180
ctgcccaggc aaagcgtccg ggcagcgtag gcgggcgact cagatcccag ccagtggact	240
tagccccgtg ttgctcctcc gataactggg gtgaccttgg ttaatatcca ccagcagcct	300
cccccgttgc ccctctggat ccaactgctta aatacggacg aggacagggc cctgtctcct	360
cagcttcagg caccaccact gacctgggac agtgaat	397

<210> 210
 <211> 2864
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 210

cctttgagaa tccacggtgt ctcgatgcag tcagctttct aacaagctgg ggcctcacct	60
gttttccac ggataaaaac gtgctggagg aagcagaaag gggctggcag gtggaaagat	120
gaggaccagc tcatcgtctc atgactatga ggttgctctg atccagaggg tccccctgcc	180
tggtggccca ccgccaggaa gactcccact gtccctggat gccagagtg ggatgtcaac	240
tccatcactt atcaactcct tatccatagg ggtattcttc ctgaggcgtc tcagaaaaca	300
gggccctccc catatgctga ccacataata gaaccctcc caactcagag accctggctg	360
ctagctgccc tggcatgacc cagacagtgg ctttgtata tgtttttaga ctcaccttga	420
ctcacctctg accatagaaa ctctcatccc agaggctact gcaatagtta ctccacaaca	480
gaggcttatac tgggtagagg gaggctccct acctatggcc cagcagccct gacagtgcag	540
atcacatata cccacgccc cagcactgcc tgccacgcat gggcttactt tacaccacc	600
cacagtcacc aacacattac ctgctctcca aggttaggcg tggcaggaga agtttgcttg	660
gaccagcaga aaccatgcag tcaaggacaa ctggagtcag catgggctgg gtgcgagccc	720
ttggtgggggt ggggaggaga ctccaggcca tacctcctgg aggatgtttt aatcatttcc	780
agcatggaat gctgtcaact tttgccacag attcattagc tctgagtttc tttttctgt	840
ccccagctac cccttacatg tcaatatgga cttaatgatg ggaaattcag gcaagttttt	900
aaacatttta tccccctgg ctcttatcct caaaaaatgc atgaatttgg aggcagtggc	960
tcatgcctgt aatcccaatg ctttgctagg ttgaggcggg aggatcactt gaagccagga	1020
atttgagacc agcctgggcc gcatagttag accccgtttc taaaaaata aataaataaa	1080
taataaataa tagtgatatg aagcatgatt aaatagccct attttttaaa atgcatgagt	1140
tcgttacctg attcattccc tggttccttt cacagtcctc cgtgacccaa gtgttagggt	1200
tttggctctc ctactatttg taggctgata tatagtatac acacacacac acacacacat	1260
atacacacac acagtgtatc ttgagctttc ttttgtatat ctacacacat atgtataaga	1320
aagctcaaga tatagaagcc ctttttcaaa aataactgaa agtttcaaac tctttaagtc	1380
tccagttacc attttgctgg tattcttatt tggaaccata cattcatcat attgttgcac	1440
agtaagacta tacattcatt attttgctta aacgtatgag ttaaaacact tggccaggca	1500
tggtggttca cacctgtaat ccagagctt tgggaagcca agactggcag atctcttgag	1560
ctcaggaatt caagaccagc ctgggcaaca tggaaaaacc ccatctctac aaaagataga	1620
aaaattagcc aggcattggtg gcgtgtgcct gtgggtccag ctactcagga ggctgaggtg	1680
ggaggatcac attagcccag gaggttgagg ctgcagttag ccgtgattat gccactgcac	1740
tccagcctgg gagacagagt gagaccctgt ttcaaaaaaa agagagagaa aatttaaaaa	1800
agaaaacaac accaagggtc gtaactttaa ggatcattaaa tgaattaatc actgcattca	1860
aaaacgatta ctttctggcc ctaagagaca tgaggccaat accaggaagg gggttgatct	1920
cccaaaccag aggcagaccc tagactctaa tacagttaag gaaagaccag caagatgata	1980
gtccccaata caatagaagt tactatatat tatttgttgt ttttcttttg ttttgttttg	2040

ttttgttttg ttttgtttta gagactgggg tcttgctcga ttgcccaggc tgtagtcag	2100
cggtagggaca atagctcact gcagactcca actcctgggc tcaagcaatc ctcctgcctc	2160
agcctcctga atagctggga ctacaagggt acaccatcac acacaccaa acaatttttt	2220
aaatttttgt gtgaaacga ggggtcttgct ttgttgccca ggctgggtctc caactcctgg	2280
cttcaaggga tcctcccacc tcagcctccc aaattgctgg gattacaggt gtgagccacc	2340
acaaccagcc agaactttac taattttaaa attaagaact taaaacttga atagctagag	2400
caccaagatt tttctttgtc cccaataag tgcagttgca ggcatagaaa atctgacatc	2460
tttgcaagaa tcatcgtgga ttagactct gtctgtgtc tctggcctgg tttcggggac	2520
caggagggca gacccttgca ctgccaagaa gcatgcaaaa gttaatcatt ggccctgctg	2580
agtacatggc cgatcaggct gtttttgtgt gcctgttttt ctattttacg taaatcacc	2640
tgaacatgtt tgcacaaacc tactgggtgat gcacctttga tcaatacatt ttagacaaac	2700
gtggtttttg agtccaaaga tcagggtggt gttgacctga atactggata cagggcata	2760
aaaacagggg caaggcacag actcatagca gagcaatcac caccaagcct ggaataactg	2820
caagggtctt gctgacatct tcctgaggtg ccaaggaaat gagg	2864

<210> 211

<211> 295

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 211

gggccccaga agcctggtgg ttgtttgtcc ttctcagggg aaaagtgagg cggccccttg	60

gaggaagggg ccgggcagaa tgatctaatac ggattccaag cagctcaggg gattgtcttt	120

ttctagcacc ttcttgccac tcctaagcgt cctccgtgac cccggctggg atttagcctg	180

gtgctgtgtc agccccgggc tcccaggggc ttcccagtgg tccccaggga accctcgaca	240

gggccagggc gtctctctcg tccagcaagg gcagggacgg gccacaggca agggc	295

<210> 212

<211> 206

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 212

gaatgactcc tttcggttaag tgcagtggaa gctgtacact gcccaggcaa agcgtccggg	60

cagcgtaggc gggcgactca gatcccagcc agtggactta gcccctgttt gtcctccga	120

taactggggg gaccttggtt aatattcacc agcagcctcc cccgttgccc ctctggatcc	180

actgcttaaa tacggacgag gacagg	206

<210> 213
<211> 1179
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 213
ggctccggtg cccgtcagtg ggcagagcgc acatcgccca cagtccccga gaagttgggg 60
ggaggggtcg gcaattgaac cggcgcctag agaaggtggc gcggggtaaa ctgggaaagt 120
gatgtcgtgt actggctccg cttttttccc gaggggtggg gagaaccgta tataagtgca 180
gtagtcgccg tgaacgttct ttttcgcaac gggtttgccg ccagaacaca ggtaagtgcc 240
gtgtgtgggt cccgcgggcc tggcctcttt acgggttatg gcccttgctg gccttgaatt 300
acttcacact ggtgcagta cgtgattctt gatcccgagc ttcgggttgg aagtgggtgg 360
gagagttcga ggccttgccg ttaaggagcc ctttcgcctc gtgcttgagt tgaggcctgg 420
cctgggcgct ggggccgccg cgtgcgaatc tgggtggcacc ttcgcgcctg tctcgctgct 480
ttcgataagt ctctagccat ttaaaatctt tgatgacctg ctgcgacgct ttttttctgg 540
caagatagtc ttgtaaagtc gggccaagat ctgcacactg gtatttcggt ttttggggcc 600
gcgggcggcg acggggcccc tgcgtcccag cgcacatgtt cggcgaggcg gggcctgcga 660
gcgcggccac cgagaatcgg acgggggtag tctcaagctg gccggcctgc tctggtgcct 720
ggtctcgccg cgccgtgtat cgccccgccc tgggcggcaa ggctggcccc gtcggcacca 780
gttgcgtgag cggaaagatg gccgcttccc ggccctgctg caggagagctc aaaatggagg 840
acgcggcgct cgggagagcg ggcgggtgag tcaccacac aaaggaaaag ggcctttccg 900
tcctcagccg tcgcttcatt tgactccacg gagtaccggg cgccgtccag gcacctcgat 960
tagttctcga gcttttgag tacgtcgtct ttaggttggg gggaggggtt ttatgcgatg 1020
gagtttccc acactgagtg ggtggagact gaagttaggc cagcttgga cttgatgtaa 1080
ttctccttgg aatttgccct ttttgagttt ggatcttggc tcatttcaa gcctcagaca 1140
gtggttcaaa gtttttttct tccatttcag gtgtcgtga 1179

<210> 214
<211> 292
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 214
gggccccaga agcctggtgg ttgtttgtcc ttctcagggg aaaagtgagg cggccccttg 60
gaggaagggg ccgggcagaa tgatctaata ggattccaag cagctcaggg gattgtcttt 120

ttctagcacc ttcttgccac tcctaagcgt cctccgtgac cccggctggg atttagcctg	180
gtgctgtgtc agccccggtc tcccaggggc ttcccagtgg tccccaggaa ccctcgacag	240
ggccccgtct ctctcgtcca gcaagggcag ggacggggcca caggccaagg gc	292

<210> 215
 <211> 1325
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 215 gctgcctact gaggcacaca ggggcgcctg cctgctgccc gctcagccaa ggcggtgttg	60
ctggagccag ctggggacag ctctcccaac gctctgccct ggccttgca cccactctct	120
gggccgtagt tgtctgtctg ttaagtgagg aaagtgccca tctccagagg cattcagcgg	180
caaagcaggg cttccagggtt ccgaccccat agcaggactt cttggatttc tacagccagt	240
cagttgcaag cagcacccaa attatttcta taagaagtgg caggagctgg atctgaagag	300
tcagcagtct acctttccct gtttcttgct ctttatgcag tcaggaggaa tgatctggat	360
tccatgtgaa gcctgggacc acggagacct aagacttcct gcttgattct ccctgcgaac	420
tgcaggctgt gggctgagcc ttcaagaagc aggagtcccc tctagccatt aactctcaga	480
gctaacctca tttgaatggg aacactagtc ctgtgatgtc tggaagggtg gggcctctac	540
actccacacc ctacatggtg gtccagacac atcattccca gcattagaaa gctctagggg	600
gaccgtttct gttccctgag gcattaaagg gacatagaaa taaatctcaa gctctgaggc	660
tgatgccagc ctgagactca gcctctgcac tgtatgggcc aattgtagcc ccaaggactt	720
cttcttgctg cacccttat ctgtccacac ctaaacgat gggcttctat tagttacaga	780
actctctggc ctgttttggt ttgctttgct ttgttttggt ttgttttttt gtttttttgt	840
tttttagcta tgaacagag gtaatatcta atacagataa cttaccagta atgagtgctt	900
cctacttact gggctactggg aagaagtgtt ttacacatat tttctcattt aatctacaca	960
ataagtaatt aagacatttc cctgaggcca cgggagagac agtggcagaa cagttctcca	1020
aggaggactt gcaagttaat aactggactt tgcaaggctc tggaggaaac tgctagcttg	1080
taaaggatgg agcacagtgt ctggcatgta gcaggaacta aaataatggc agtgattaat	1140
gttatgatat gcagacacaa cacagcaaga taagatgcaa tgtaccttct gggtaaacc	1200
accctggcca ctctccccg ataccaggg ttgatgtgct tgaattagac aggattaaag	1260
gcttactgga gctggaagcc ttgccccaac tcaggagttt agccccagac cttctgtcca	1320
ccagc	1325

<210> 216
 <211> 883
 <212> DNA
 <213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 216

gagtcaatgg gaaaaaccca ttggagccaa gtacactgac tcaataggga ctttccattg	60
ggttttgccc agtacataag gtcaataggg ggtgagtcaa caggaaagtc ccattggagc	120
caagtacatt gagtcaatag ggacttttcca atgggttttg cccagtacat aaggtcaatg	180
ggaggtaagc caatgggttt ttcccattac tgacatgtat actgagtcac tagggacttt	240
ccaatgggtt ttgcccagta cataaggtca ataggggtga atcaacagga aagtcccatt	300
ggagccaagt aactgagtc aatagggact ttccattggg ttttgcccag taaaaaggt	360
caataggggg tgagtcaatg ggtttttccc attattggca catacataag gtcaataggg	420
gtggggcctg aaataacctc tgaaagagga acttggttag gtaccttctg aggctgaaag	480
aaccagctgt ggaatgtgtg tcagttaggg tgtggaaagt cccaggctc cccagcaggc	540
agaagtatgc aaagcatgca tctcaattag tcagcaacca ggtgtggaaa gtcccaggc	600
tccccagcag gcagaagtat gcaaagcatg catctcaatt agtcagcaac catagtccca	660
ctagtggaga agagcatgct tgagggtga gtgcccctca gtgggcagag agcacatggc	720
ccacagtccc tgagaagttg gggggagggg tgggcaattg aactggtgcc tagagaaggt	780
ggggcttggg taaactggga aagtgtgtg gtgtactggc tccacctttt tccccagggt	840
gggggagaac catatataag tgcagtagtc tctgtgaaca ttc	883

<210> 217

<211> 639

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 217

gggcctgaaa taacctctga aagaggaact tggtaggta cttctgagg ctgaaagaac	60
cagctgtgga atgtgtgtca gtaggggtgt ggaaagtccc caggctcccc agcaggcaga	120
agtatgcaaa gcatgcatct caattagtca gcaaccaggt gtggaaagtc cccaggctcc	180
ccagcaggca gaagtatgca aagcatgcat ctcaattagt cagcaacat agtcccacta	240
gtggagaaga gcatgcttga gggctgagtg cccctcagtg ggcagagagc acatggccca	300
cagtccctga gaagttgggg ggaggggtgg gcaattgaac tggcgcctag agaaggtggg	360
gcttgggtaa actgggaaag tgatgtggtg tactggctcc acctttttcc ccagggtggg	420
ggagaacat atataagtgc agtagtctct gtgaacattc aagcttctgc cttctccctc	480
ctgtgagttt ggtaagtcac tgactgtcta tgcctgggaa aggggtgggca ggagatgggg	540
cagtgcagga aaagtggcac tatgaaccct gcagccctag acaattgtac taaccttctt	600

ctcttttcctc tcctgacagg ttggtgtaca gtagcttcc

639

<210> 218

<211> 1272

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 218

aggctcagag gcacacagga gtttctgggc tcaccctgcc cccttccaac ccctcagttc 60

ccatcctcca gcagctgttt gtgtgctgcc tctgaagtcc aactgaaca aacttcagcc 120

tactcatgtc ctaaaatgg gcaaacattg caagcagcaa acagcaaaca cacagccctc 180

cctgcctgct gaccttggag ctggggcaga ggtcagagac ctctctgggc ccatgccacc 240

tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagaggtt gtcctggcgt 300

ggttaggta gtgtgagagg gtccgggttc aaaaccactt gctgggtggg gagtcgtcag 360

taagtggcta tgccccgacc ccgaagcctg tttcccatc tgtacaatgg aaatgataaa 420

gacgcccatac tgatagggtt tttgtggcaa ataaacattt ggtttttttg ttttgttttg 480

ttttgttttg tgagatggag gtttgctctg tcgcccaggc tggagtgcag tgacacaatc 540

tcatctcacc acaaccttcc cctgcctcag cctcccaagt agctgggatt acaagcatgt 600

gccaccacac ctggctaatt ttctattttt agtagagacg ggtttctcca tgttggtcag 660

cctcagcctc ccaagtaact gggattacag gcctgtgcc aacacccgg ctaatttttt 720

ctatttttga cagggacggg gtttcacatc gttggtcagg ctggtctaga ggtactggat 780

cttgctacca gtggaacagc cactaaggat tctgcagtga gagcagaggg ccagctaagt 840

ggtactctcc cagagactgt ctgactcacg ccaccccctc caccttggac acaggacgct 900

gtggtttctg agccaggtac aatgactcct ttcggtaagt gcagtggaag ctgtacactg 960

cccaggcaaa gcgtccgggc agcgtaggcg ggcgactcag atcccagcca gtggacttag 1020

cccctgtttg ctctccgat aactggggtg accttggtta atattcacca gcagcctccc 1080

ccgttgcccc tctggatcca ctgcttaaat acggacgagg acagggccct gtctcctcag 1140

cttcaggcac caccactgac ctgggacagt gaataattac tctaaggtaa atataaaatt 1200

tttaagtgtg taatgtgtta aactactgat tctaattgtt tctctctttt agattccaac 1260

ctttggaact ga 1272

<210> 219

<211> 547

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 219
ccctaaaatg ggcaaacatt gcaagcagca aacagcaaac acacagccct ccctgcctgc 60
tgaccttgga gctggggcag aggtcagaga cctctctggg cccatgccac ctccaacatc 120
cactcgaccc cttggaattt ttcggtggag aggagcagag gttgtcctgg cgtgggttag 180
gtagtgtgag aggggaatga ctcttttcgg taagtgcagt ggaagctgta cactgcccag 240
gcaaagcgtc cgggcagcgt aggcgggcga ctcagatccc agccagtgga cttagcccct 300
gtttgctcct ccgataactg gggtgacctt ggttaatat caccagcagc ctccccggtt 360
gccccctctgg atccactgct taaatacggg cgaggacagg gccctgtctc ctgagcttca 420
ggcaccacca ctgacctggg acagtgaatc cggactctaa ggtaaatata aaatTTTTaa 480
gtgtataatg tgtaaacta ctgattctaa ttgtttctct cttttagatt ccaacctttg 540
gaactga 547

<210> 220
<211> 709
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 220
cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag 60
gggctaagtc cacatacggg ggaggctgct ggtgaatatt aaccaaggct accccagtta 120
tcggaggagc aaacaggggc taagtccaca tagggctgga agctaccttt gacatcattt 180
cctctgcgaa tgcatgtata atttctacag aacctattag aaaggatcac ccagcctctg 240
ctttgtaca actttccctt aaaaaactgc caattccact gctgtttggc ccaatagtga 300
gaactttttc ctgctgcctc ttggtgcttt tgcctatggc ccctattctg cctgctgaag 360
acactcttgc cagcatggac ttaaaccctt ccagctctga caatcctctt tctcttttgt 420
ttacatgaa gggctctggc gccaaagcaa tcaactaaag ttcaaaccct atcatttttt 480
gctttgttcc tcttggcctt ggttttgtac atcagctttg aaaataccat cccagggtta 540
atgctggggg taatttataa ctaagagtgc tctagttttg caatacagga catgctataa 600
aaatggaaag atctcctgaa gaggtaaggg tttaagggat gggttggttg tggggtatta 660
atgtttaatt acctggagca cctgcctgaa atcacttttt ttcaggttg 709

<210> 221
<211> 460
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 221

gggctggaag ctacctttga catcatttcc tctgcgaatg catgtataat ttctacagaa	60
cctattagaa aggatcaccc agcctctgct tttgtacaac tttcccttaa aaaactgcca	120
attccactgc tgtttggccc aatagtgaga actttttcct gctgcctctt ggtgcttttg	180
cctatggccc ctattctgcc tgctgaagac actcttgcca gcatggactt aaaccctcc	240
agctctgaca atcctctttc tcttttgttt tacatgaagg gtctggcagc caaagcaatc	300
actcaaagtt caaaccttat catTTTTTgc tttgttcctc ttggccttgg tttgtacat	360
cagctttgaa aataccatcc cagggttaat gctggggtta atttataact aagagtgctc	420
tagttttgca atacaggaca tgctataaaa atggaaagat	460

<210> 222

<211> 699

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 222

cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag	60
gggctaagtc cacatacggg ggaggctgct ggtgaatatt aaccaaggtc accccagtta	120
tcggaggagc aaacaggggc taagtccaca taccctaaaa tgggcaaaca ttgcaagcag	180
caaacagcaa acacacagcc ctccctgcct gctgaccttg gagctggggc agaggtcaga	240
gacctctctg ggcccatgcc acctccaaca tccactcgac cccttggaat ttttcggtgg	300
agaggagcag aggttgctct ggcgtgggtt aggtagtgtg agaggggaat gactcctttc	360
ggtaagtgca gtggaagctg tacactgccc aggcaaagcg tccgggcagc gtaggcgggc	420
gactcagatc ccagccagtg gacttagccc ctgtttgctc ctccgataac tggggtgacc	480
ttggttaata ttcaccagca gcctcccccg ttgcccctct ggatccactg cttaaatacg	540
gacgaggaca gggccctgtc tcctcagctt caggcaccac cactgacctg ggacagtgaa	600
tccggactct aaggtaaata taaaatTTTT aagtgtataa tgtgttaaac tactgattct	660
aattgtttct ctcttttaga ttccaacctt tggaactga	699

<210> 223

<211> 681

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 223

aggtaattt taaaaagca gtcaaaagtc caagtggccc ttggcagcat ttactctctc	60
tgtttgctct ggtaataat ctcaggagca caaacattcc agatccaggt taatttttaa	120
aaagcagtca aaagtccaag tggcccttgg cagcatttac tctctctgtt tgctctggtt	180

aataatctca ggagcacaaa cattccagat cctgctctcc agggctggaa gctacctttg	240
acatcatttc ctctgcgaat gcatgtataa tttctacaga acctattaga aaggatcacc	300
cagcctctgc ttttgtaaa ctttccctta aaaaactgcc aattccactg ctgtttggcc	360
caatagttag aactttttcc tgctgcctct tgggtgcttt gcctatggcc cctattctgc	420
ctgctgaaga cactcttgcc agcatggact taaaccctc cagctctgac aatcctcttt	480
ctcttttggt ttacatgaag ggtctggcag ccaaagcaat cactcaaagt tcaaacctta	540
tcattttttg ctttgcttct cttggccttg gttttgtaca tcagctttga aaataccatc	600
ccagggttaa tgctgggggt aatttataac taagagtgtc ctagttttgc aatacaggac	660
atgctataaa aatggaaaga t	681

<210> 224

<211> 532

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 224

gttacataac ttatggtaaa tggcctgcct ggctgactgc ccaatgaccc ctgcccaatg	60
atgtcaataa tgatgtatgt tcccatgtaa tgccaatagg gactttccat tgatgtcaat	120
gggtggagta tttatggtaa ctgccactt ggcagtacat caagtgtatc atatgccaaag	180
tatgccccct attgatgtca atgatggtaa atggcctgcc tggcattatg ccagttacat	240
gaccttatgg gactttccta cttggcagta catctatgta ttagtcattg ctattaccat	300
gggaattcac tagtggagaa gagcatgctt gagggctgag tgcccctcag tgggcagaga	360
gcacatggcc cacagtcctt gagaagttag ggggaggggt gggcaattga actggtgcct	420
agagaagggt gggcttgggt aaactgggaa agtgatgtgg tgtactggct ccaccttttt	480
ccccagggtg ggggagaacc atatataagt gcagtagtct ctgtgaacat tc	532

<210> 225

<211> 955

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 225

gagtcaatgg gaaaaaccca ttggagccaa gtacactgac tcaataggga ctttccattg	60
ggttttgccc agtacataag gtcaataggg ggtgagtcaa caggaaagtc ccattggagc	120
caagtacatt gagtcaatag ggactttcca atgggttttg ccagttacat aagggtcaatg	180
ggaggtaagc caatgggttt ttcccattac tgacatgtat actgagtcac tagggacttt	240

ccaatgggtt ttgcccagta cataaggtca ataggggtga atcaacagga aagtcccatt	300
ggagccaagt aactgagtc aatagggact ttccattggg ttttggccag tacaaaaggt	360
caataggggg tgagtcaatg ggtttttccc attattggca catacataag gtcaataggg	420
gtggttacat aacttatggt aaatggcctg cctggctgac tgcccaatga cccctgcca	480
atgatgtcaa taatgatgta tgttcccatg taatgccaat agggactttc cattgatgtc	540
aatgggtgga gtatttatgg taactgccc cttggcagta catcaagtg atcatatgcc	600
aagtatgccc cctattgatg tcaatgatgg taaatggcct gcctggcatt atgcccagta	660
catgacctta tgggactttc ctacttggca gtacatctat gtattagtca ttgctattac	720
catgggaatt cactagtgga gaagagcatg cttgagggtc gagggtcccct cagtgggcag	780
agagcacatg gccacagtc cctgagaagt tggggggagg ggtgggcaat tgaactggtg	840
cctagagaag gtggggcttg ggtaaactgg gaaagtgatg tgggtgtactg gctccacctt	900
tttccccagg gtgggggaga accatatata agtgcagtag tctctgtgaa cattc	955

<210> 226

<211> 955

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 226

gttacataac ttatggtaaa tggcctgcct ggctgactgc ccaatgacct ctgcccaatg	60
atgtcaataa tgatgtatgt tcccatgtaa tgccaatagg gactttccat tgatgtcaat	120
gggtggagta tttatggtaa ctgcccactt ggcagtacat caagtgtatc atatgccaa	180
tatgccccct attgatgtca atgatggtaa atggcctgcc tggcattatg cccagtacat	240
gaccttatgg gactttccta cttggcagta catctatgta ttagtcattg ctattacat	300
gggagtcaat gggaaaaacc cattggagcc aagtacactg actcaatagg gactttccat	360
tgggttttgc ccagtacata aggtcaatag ggggtgagtc aacaggaaag tcccattgga	420
gccaagtaca ttgagtcaat agggactttc caatgggttt tgcccagtac ataagggtcaa	480
tgggaggtaa gccaatgggt ttttcccatt actgacatgt atactgagtc attagggact	540
ttccaatggg ttttggccag tacataaggt caataggggt gaatcaacag gaaagtcca	600
ttggagccaa gtacactgag tcaataggga ctttccattg ggttttgccc agtacaaaag	660
gtcaataggg ggtgagtcaa tgggtttttc ccattattgg cacatacata aggtcaatag	720
gggtggaatt cactagtgga gaagagcatg cttgagggtc gagggtcccct cagtgggcag	780
agagcacatg gccacagtc cctgagaagt tggggggagg ggtgggcaat tgaactggtg	840
cctagagaag gtggggcttg ggtaaactgg gaaagtgatg tgggtgtactg gctccacctt	900
tttccccagg gtgggggaga accatatata agtgcagtag tctctgtgaa cattc	955

<210> 227
 <211> 1923
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 227
 tcaatattgg ccattagcca tattattcat tggttatata gcataaatca atattggcta 60
 ttggccattg catacgttgt atctatatca taatatgtac atttatattg gtcattgtcc 120
 aatatgaccg ccatgttggc attgattatt gactagttat taatagtaat caattacggg 180
 gtcattagtt catagcccat atatggagtt ccgcgttaca taacttacgg taaatggccc 240
 gcctggctga ccgccaacg acccccggcc attgacgtca ataatgacgt atgttcccat 300
 agtaacgcca atagggactt tccattgacg tcaatgggtg gagtatttac ggtaaactgc 360
 ccacttggca gtacatcaag tgtatcatat gccaaagtcc cccctattg acgtcaatga 420
 cggtaaatgg cccgcctggc attatgcccc gtacatgacc ttacgggact ttcctacttg 480
 gcagtacatc tacgtattag tcatcgctat taccatggtc gaggtgagcc ccacgttctg 540
 cttcactctc cccatctccc cccctcccc acccccaatt ttgtatttat ttatTTTTTA 600
 attatTTTgt gcagcgatgg gggcgggggg gggggggggg cgcgcgccag gcggggcggg 660
 gcggggcgag gggcggggcg gggcgaggcg gagagggtgc gcggcagcca atcagagcgg 720
 cgcgctccga aagtttcctt ttatggcgag gcggcggcgg cggcggccct ataaaaagcg 780
 aagcgcgcgg cgggcgggag tcgctgcgac gctgccttcg ccccggtccc cgctccgccg 840
 ccgcctcgcg ccgcccggc ggctctgac tgaccgcgtt actccacag gtgagcgggc 900
 gggacggccc ttctcctccg ggctgtaatt agcgcttggg ttaatgacgg cttgtttctt 960
 ttctgtggct gcgtgaaagc cttgaggggc tccgggaggg ccctttgtgc gggggggagc 1020
 ggctcggggg gtgcgtgcgt gtgtgtgtgc gtggggagcg ccgcgtgcgg ccgcgctgc 1080
 ccggcggctg tgagcgtgc gggcgcggcg cggggctttg tgcgtccgc agtgtgcgcg 1140
 aggggagcgc gcccgggggc ggtgccccgc ggtgcggggg gggctgcgag gggaacaaag 1200
 gctgcgtgcg ggggtgtgtc gtgggggggt gagcaggggg tgtgggcgcg gcggtcgggc 1260
 tgtaaccccc ccctgcaccc ccctccccga gttgctgagc acggcccggc ttcgggtgcg 1320
 gggctccgta cggggcggtg cgcggggctc gccgtgccgg gcgggggggt gcggcaggtg 1380
 ggggtgccgg gcggggcggg gccgcctcgg gccggggagg gctcggggga ggggcgcggc 1440
 ggccccgga gcgccggcgg ctgtcgaggc gcggcgagcc gcagccattg ctttttatgg 1500
 taatcgtgcg agagggcgca gggacttcct ttgtccaaa tctgtgcgga gccgaaatct 1560
 gggaggcgcc gccgcacccc ctctagcggg cgcggggcga agcgggtcgg cgccggcagg 1620
 aaggaaatgg gcggggaggg ccttcgtgcg tcgccgcgc gccgtcccct tctccctctc 1680
 cagcctcggg gctgtccgcg gggggacggc tgccttcggg ggggacgggg cagggcgggg 1740

ttcggcttct ggcgtgtgac cggcggctct agagcctctg ctaacatgt tttagccttc	1800
ttctttttcc tacagctcct gggcaacgtg ctggttattg tgctgtctca tcatttgtcg	1860
acagaattcc tcgaagatcc gaagggggtc aagcttggca ttccggtact gttggtaaag	1920
cca	1923

<210> 228
 <211> 1272
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 228	
aggctcagag gcacacagga gtttctgggc tcaccctgcc cccttccaac ccctcagttc	60
ccatcctcca gcagctgttt gtgtgctgcc tctgaagtcc aactgaaca aacttcagcc	120
tactcatgtc cctaaaatgg gcaaacattg caagcagcaa acagcaaaca cacagccctc	180
cctgcctgct gaccttggag ctggggcaga ggtcagagac ctctctgggc ccatgccacc	240
tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagagggt gtcctggcgt	300
ggtttaggta gtgtgagagg gtccgggttc aaaaccactt gctgggtggg gagtcgtcag	360
taagtggcta tgccccgacc ccgaagcctg tttccccatc tgtacaatgg aaatgataaa	420
gacgccccatc tgatagggtt tttgtggcaa ataaacattt ggtttttttg ttttgttttg	480
ttttgttttt tgagatggag gtttgctctg tcgcccaggc tggagtgcag tgacacaatc	540
tcatctcacc acaaccttcc cctgcctcag cctcccaagt agctgggatt acaagcatgt	600
gccaccacac ctggctaatt ttctattttt agtagagacg ggtttctcca tgttggtcag	660
cctcagcctc ccaagtaact gggattacag gcctgtgcca ccacaccgg ctaatttttt	720
ctatttttga caggacggg gtttcacat gttggtcagg ctggtctaga ggtaccggat	780
cttgctacca gtggaacagc cactaaggat tctgcagtga gagcagaggg ccagctaagt	840
ggtactctcc cagagactgt ctgactcacg ccaccccctc caccttggac acaggacgct	900
gtggtttctg agccaggtac aatgactcct ttcggtaagt gcagtggaag ctgtacactg	960
cccaggcaaa gcgtccgggc agcgtaggcg ggcgactcag atcccagcca gtggacttag	1020
cccctgtttg ctctccgat aactgggggtg accttggtta atattcacca gcagcctccc	1080
ccgttgcccc tctggatcca ctgcttaaat acggacgagg acagggccct gtctcctcag	1140
cttcaggcac caccactgac ctgggacagt gaatccggac tctaaggtaa atataaaatt	1200
tttaagtgtg taatgtgtta aactactgat tctaattgtt tctctctttt agattccaac	1260
ctttggaact ga	1272

<210> 229
 <211> 826
 <212> DNA
 <213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 229

gagtcaatgg gaaaaaccca ttggagccaa gtacactgac tcaataggga ctttccattg	60
ggttttgccc agtacataag gtcaataggg ggtgagtcaa caggaaagtc ccattggagc	120
caagtacatt gagtcaatag ggacttttcca atggggtttg cccagtacat aagggtcaatg	180
ggaggtaagc caatgggttt ttcccattac tgacatgtat actgagtcac tagggacttt	240
ccaatgggtt ttgcccagta cataaggtca ataggggtga atcaacagga aagtcccatt	300
ggagccaagt aactgagtc aatagggtact ttccattggg ttttgcccag tacaaaaggt	360
caataggggg tgagtcaatg ggtttttccc attattggca catacataag gtcaataggg	420
gtgactagtg gagaagagca tgcttgaggg ctgagtgccc ctgagtgagg agagagcaca	480
tggcccacag tccctgagaa gttgggggga ggggtgggca attgaactgg tgcctagaga	540
aggtggggct tgggtaaact gggaaagtga tgtggtgtac tggctccacc tttttcccca	600
gggtggggga gaaccatata taagtgcagt agtctctgtg aacattcaag cttctgcctt	660
ctccctcctg tgagtttggg aagtcactga ctgtctatgc ctgggaaagg gtgggcagga	720
gatggggcag tgcaggaaaa gtggcactat gaaccctgca gccctagaca attgtactaa	780
ccttcttctc tttcctctcc tgacaggttg gtgtacagta gcttcc	826

<210> 230

<211> 399

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 230

cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag	60
gggctaagtc cacacgcgtg gtaccgtctg tctgcacatt tcgtagagcg agtgttccga	120
tactctaadc tccctaggca aggttcatat ttgtgtaggt tacttattct cttttgttg	180
actaagtcaa taatcagaat cagcaggttt ggagtcagct tggcagggat cagcagcctg	240
ggttgggaagg agggggtata aaagcccctt caccaggaga agccgtcaca cagatccaca	300
agctcctgaa gaggtgaagg tttaagggat ggttggttgg tggggtatta atgtttaatt	360
acctggagca cctgcctgaa atcacttttt ttcaggttg	399

<210> 231

<211> 654

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 231

gacattgatt attgactagt tattaatagt aatcaattac ggggtcatta gttcatagcc	60
catatatgga gttccgcgtt acataactta cggtaaatgg cccgcctggc tgaccgcca	120
acgacccccg cccattgacg tcaataatga cgtatgttcc catagtaacg ccaataggga	180
ctttccattg acgtcaatgg gtggactatt tacggtaaac tgcccacttg gcagtacatc	240
aagtgtatca tatgccaagt acgcccccta ttgacgtcaa tgacggtaaa tggccgcct	300
ggcattatgc ccagtacatg accttatggg actttcctac ttggcagtac atctacgtat	360
tagtcatcgc tattaccatg gtgatgcggt tttggcagta catcaatggg cgtggatagc	420
ggtttgactc acggggattt ccaagtctcc accccattga cgtcaatggg agtttgttt	480
ggcaccaaaa tcaacgggac tttccaaaat gtcgtaacaa ctccgcccc ttgacgcaaa	540
tgggcggtag gcgtgtacgg tgggagggtct atataagcag agctctctgg ctaactagag	600
aaccactgc ttactggctt atcgaaatta atacgactca ctataggag accc	654

<210> 232

<211> 500

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 232

gggtaggggga ggcgcttttc ccaaggcagt ctggagcatg cgcttttagca gccccgctgg	60
gcacttggcg ctacacaagt ggcctctggc ctgcacaca ttccacatcc accggtaggc	120
gccaaccggc tccgttcttt ggtggcccct tcgcgccacc ttctactcct cccctagtca	180
ggaagttccc cccgccccg cagctcgcgt cgtgcaggac gtgacaaatg gaagtagcac	240
gtctcactag tctcgtgcag atggacagca ccgctgagca atggaagcgg gtaggccttt	300
ggggcagcgg ccaatagcag ctttgctcct tcgctttctg ggctcagagg ctgggaaggg	360
gtgggtccgg gggcgggctc aggggcgggc tcaggggcgg ggcgggcgcc cgaaggtcct	420
ccggaggccc ggcattctgc acgcttcaaa agcgcacgtc tgccgcgctg ttctcctctt	480
cctcatctcc gggcctttcg	500

<210> 233

<211> 450

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 233

gggcctgaaa taacctctga aagaggaact tggttaggta ctttctgagg ctgaaagaac	60

cagctgtgga atgtgtgtca gttagggtgt ggaaagtccc caggctcccc agcaggcaga	120
agtatgcaaa gcatgcatct caattagtca gcaaccaggt gtggaaagtc cccaggctcc	180
ccagcaggca gaagtatgca aagcatgcat ctcaattagt cagcaaccat agtcccacta	240
gttccagatg gtaaatatac acaagggatt tagtcaaaca attttttggc aagaatatta	300
tgaattttgt aatcggttgg cagccaatga aatacaaaga tgagtctagt taataatcta	360
caattattgg ttaaagaagt atattagtgc taatttcctt ccgtttgtcc tagcttttct	420
cttctgtcaa cccacacgc ctttggcacc	450

<210> 234

<211> 594

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 234

gacattgatt attgactagt tattaatagt aatcaattac ggggtcatta gttcatagcc	60
catatatgga gttccgcgtt acataactta cggtaaattgg cccgcctggc tgaccgcca	120
acgacccccg cccattgacg tcaataatga cgtatgttcc catagtaacg ccaataggga	180
ctttccattg acgtcaatgg gtggactatt tacggtaaac tgcccacttg gcagtacatc	240
aagtgtatca tatgccaagt acgcccccta ttgacgtcaa tgacggtaaa tggccgcct	300
ggcattatgc ccagtacatg accttatggg actttcctac ttggcagtac atctacgtat	360
tagtcatcgc tattaccatg actagttcca gatggtaaata atacacaagg gatttagtca	420
aacaatTTTT tggcaagaat attatgaatt ttgtaatcgg ttggcagcca atgaaataca	480
aagatgagtc tagttaataa tctacaatta ttggttaaag aagtatatta gtgctaattt	540
ccctccgttt gtcctagctt ttcttttctg tcaacccac acgccttttg cacc	594

<210> 235

<211> 1210

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 235

ggcctccgcg ccgggttttg ggcctcccg cgggcgcccc cctcctcacg gcgagcgctg	60
ccacgtcaga cgaaggcgcg aggagcgtcc tgatccttcc gcccgacgc tcaggacagc	120
ggcccgtgc tcataagact cggccttaga accccagtat cagcagaagg acattttagg	180
acgggacttg ggtgactcta gggcactggt tttctttcca gagagcggaa caggcgagga	240
aaagtagtcc cttctcgcg attctgcgga gggatctccg tggggcggtg aacgccgatg	300

attatataag gacgcgccgg gtgtggcaca gctagttccg tcgcagccgg gatttgggtc	360
gcggttcttg tttgtggatc gctgtgatcg tcacttgggtg agtagcgggc tgctgggctg	420
gccggggcctt tcgtggccgc cgggccgctc ggtgggacgg aagcgtgtgg agagaccgcc	480
aagggctgta gtcgtgggtcc gcgagcaagg ttgccctgaa ctggggggttgg gggggagcgc	540
agcaaaatgg cggctgttcc cgagtcttga atggaagacg cttgtgaggc gggctgtgag	600
gtcgttgaaa caaggtgggg ggcatgggtgg gcggcaagaa cccaaggtct tgaggccttc	660
gctaatacgcg gaaagctctt attcgggtga gatgggctgg ggcaccatct ggggaccctg	720
acgtgaagtt tgtcactgac tggagaactc ggtttgtcgt ctgttgccgg gcgggcagtt	780
atgcggtgcc gttgggcagt gcacccgtac ctttgggagc gcgcgccctc gtcgtgtcgt	840
gacgtcacc gttctgttgg cttataatgc aggggtggggc cacctgccgg taggtgtgcg	900
gtaggcctttt ctccgtcgca ggacgcaggg ttccgggccta gggtaggctc tcctgaatcg	960
acaggcgccg gacctctggt gaggggaggg ataagtgagg cgtcagtttc tttggtcgggt	1020
tttatgtacc tatcttctta agtagctgaa gctccggttt tgaactatgc gctcggggtt	1080
ggcgagtgtg ttttgtgaag ttttttaggc accttttgaa atgtaatcat ttgggtcaat	1140
atgtaatttt cagtgttaga ctagtaaatt gtccgctaaa ttctggccgt ttttggcttt	1200
tttgtagac	1210

<210> 236

<211> 3000

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 236

ttaagggttg agtgtgagga aaggtctgag ggttgagaag gggtaggagga tgcacctggg	60
cctatgacag ggtccacgg aggtggctga tggcaaaagc tgggggactc caactgctga	120
tgctgaaaca agcttgtgtc tcacatacac agggacagtt cactgagctt caatgacagg	180
cacctcctgc tcatcacatc ttttctctct aggacagctt tgcccttatt ttaactagac	240
ttcccttgaa ccaaaagga aggctacatg ctgtgacttg ctgggcagcc tggaaaggcg	300
ggccactcct agccacagag atgagacaga gttcagacaa gagcttatcc ccagtcttcc	360
ttttctattt tgtttatttt attttatttt tttatttatt gagacagagt ctctgtcacc	420
caggctgggg tgtagtgatg cgacattggc ttactgcagt ctccacctcc tgggctcagg	480
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gaactcctgg actcaagcga tccgccacc tcagcctccc aaagtgctcg gattacaggc	660
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gcctctgggc acagtgacct cagtggggtg aggggagctc tccccatagc tgggctgcgg	2880
cccaacccca cccctcagg ctatgccagg ggggtgttgcc aggggcaccc gggcatcgcc	2940
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<210> 237

<211> 3000

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 237

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tggcccagat cccagcccca cagccgggct tccacatggc cgagcaggaa ctccagagca	180
ggagcacaca aaggagggct ttgatgcgcc tccagccagg cccaggcctc tcccctctcc	240
cctttctctc tgggtcttcc tttgccccac tgagggcctc ctgtgagccc gatttaacgg	300
aaactgtggg cggtgagaag ttccttatga cactactaat ccaacctgct gaccggacca	360
cgctccagc ggagggaacc tctagagctc caggacattc aggtaccagg tagccccaag	420
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aggactgggg tgaggactca ggggagcagg gagaccacgt cccaagatgc ctgtaaaact	540
gaaaccacct ggccattctc caggttgagc cagaccaatt tgatggcaga tttagcaaat	600
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ggtttttcta acccctgcag ccctgacttc ctatcctagg gaaggggccg gctggagagg	2940
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<210> 238

<211> 718

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 238

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cctggatact gaactgtgcc gaagaaggat agcagggtta aaacatgcaa agacagcacc	180
tcataatacct ctaatgttgt taacaatagc taacttttat caaacagtgt cctgtcacca	240

tgacagttac aacataatga taatgactgt acttttctcta accaggtcta gatcacttat	300
aataaatata tcttttagta attgagtaaa tgaattacag tgaggataac agcaaagaaa	360
tggtggacag atgtttacac caagaaagta tgatgactga ggtcagctca ggactgcatg	420
gcaggccac atggctcttt tttatccaac tcactactcc ctctcccttg aaaggatcca	480
agtctggaat atagccaaaa cactgttatg taaacaccaa gtccaaataa tgtgcaagca	540
tctaaagtat tgaaagccac ttttgttacc ttccatcagc tgaggggtgg agagggttcc	600
cagagccgca ggctcctcca ataaggatta gattgcatac aaaaaagccc tggctaagaa	660
cttgcttcct catcctacag ctggtaccag aactctctct aatcttcact ggaagaaa	718

<210> 239

<211> 1313

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 239

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atgcgcccag ggctgagcgc gggtagatca gagcacacaa gctcacagtc cccggcggtg	180
gggggagggg cgcgtgagc gggggccagg gagctggcgc ggggcaaact gggaaagtgg	240
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agtcgccttg gacgttcttt ttgcgaacgg gtttgccgtc agaacgcagg tgagtggcgg	360
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gagtgtcgtc cgcagggttt agctgtgagc attccactt cgagtggcgg gcggtgcggg	480
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gccctcgatt gccttcagc agcatgggct aacaaaggga ggggtgtggg ctactctta	660
aggagcccat gaagcttacg ttggatagga atggaagggc aggaggggag actggggccc	720
gcccgccttc ggagcacatg tccgacgcca cctggatggg gcgaggcctg tggctttccg	780
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aatttgcccc tttgagtttg gagcagggtc aattctcaag cctcttagcg gttcaaaggt	1260

attttctaaa cccgtttcca ggtgttgtga aagccaccgc taattcaaag caa 1313

<210> 240

<211> 1420

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 240

ggcctgaaat aacctctgaa agaggaactt ggtaggtac cttctgaggc ggaaagaacc 60
agctgtggaa tgtgtgtcag ttaggggtgtg gaaagtcccc aggctcccca gcaggcagaa 120
gtatgcaaag catgcatctc aattagtcag caaccagggt tggaaagtcc ccaggctccc 180
cagcaggcag aagtatgcaa agcatgcatc tcaattagtc agcaaccata gtcccactag 240
tggctccggt gcccgtcagt gggcagagcg cacatcgccc acagtccccg agaagttggg 300
gggagggggtc ggcaattgaa ccggtgccta gagaaggtgg cgcggggtaa actgggaaag 360
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cgtgtgtggt tcccgcgggc ctggcctctt tacgggttat ggcccttgcg tgccttgaat 540
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agtggttcaa agtttttttc ttccatttca ggtgtcgtga 1420

<210> 241

<211> 1831

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 241

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aatatgaccg ccatgttggc attgattatt gactagttat taatagtaat caattacggg	180
gtcattagtt catagcccat atatggagtt ccgcgttaca taacttacgg taaatggccc	240
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<210> 242
<211> 3000
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 242
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caattaggtg aatttagaag attttcataa ctcccccttt ctctctttta tttgtacatg	1920
tgtgtatata tgtatgtaat acatatcacg gtctcctcct gtgacctcca tgggtctgca	1980
tttcatcata aggaatagat gcttcaatgg tggccagcag tttcctcagg gtcttctcag	2040
cagtgcattg ggccacatt agctcctctg gctccaagcg aagagatggg ctctagcccc	2100
ctgtttgatt tggggcactt acagtcctct cgccagctaa actctcacac tcgtcagcat	2160
ccagacgctg aggggaaaat accagctgct tctgtgctct gcttactctt cggtaacttct	2220
ctgccatttc tggttcctga agatgtttat ttttatttat ttgagtctga ctgtatctct	2280
ttttaaaaac atgttatcca ccattgctat atatttgaag cagagaaagt tagtgaagca	2340
taaacttcat gctgaatcga gtgtctatat cctggaattc tcagcctgta ccctctataa	2400
actaatTTTT cactgtgaa taagactaat catgactctg tcgacattta cattttattt	2460
agaaaatgtc ttccttctgt tcctttgatc caagcttgac tcaccttacc ttgaggttgc	2520
atttacaag gaacactgaa ggttaccaa cagtatgtgg gtgtcgttca tcaactacag	2580
tgactcaaga atatcaccag ttggtttgcc tttctcatgg ttttaatgtt ttctcattaa	2640
aaataaataa agcacagata agcagaaaga ataaccatcc atccaacaac tagaggaaaa	2700
tttatcaatg gttttgcttt atctttccta taattaagct ataaaaaaca accatccatg	2760
taacaactag agaaaacctt tatcaatgac tgtggcttat ctttcctgat aattaggctc	2820
tttcaggag ttattaaccg attttaaac ttttgtctga gattgattag taaagattat	2880
ttctgaacc aaattgttct ttcgtttggc tactttgatt aaagaagaaa gaagagataa	2940
taattgcaat gattctttta ttttatttta tagggtcgtt ggctgtgggt tgcaattacc	3000

<210> 243

<211> 3095

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 243

tatggcacaa gcaatctctt atttttatct tagtgcataa ataaattttt cttttttgcc	60
agaataattt tttttaaaga agcgattagt ttttcttctc tcagatagca atgatgtgct	120
ttcctctcaa cctagattta gggcattttt atgtgagata ggattaaaaa ttccattttt	180
gtacaaccac tatggagaac agtttggcag ttccccaaaa aactaaaaat agagctacta	240
tatgatctag tgatcccact gctgggtata tacctataag aaaggaaatc agtatatcaa	300
agagatgtct gttcttttat gtttgttgca gcactgttca caatagccaa gatctggaag	360
caaccaagt ctccatcaac atgggtttta aaaaaatgtg gtactttaat acacaatgga	420
gtactattca gcaataaaaa agaattagat cctgttattt gcaataacat ggacagaact	480

ggaggtcatt atgtcaaatg aaataagcca ggcacagaaa gccaaacatc acatattctc	540
actcatatgt ggggtctaaa aatcaaaaca atctgattca tggagctaga gagtagagag	600
ctaattacca gaggtgggga agggtagtag gggcctggag gggaggtgga gatggttaat	660
aggtacaaaa aaatatagaa agaatgaata agacctagta tttgatagta caacagggcg	720
aatatagtca aaataattta attatacatt taaaaatacc tgaaagagta taattggctt	780
gtttgcaaca caaaagataa atgcttgagg ggatggatgc cccattttca atgatgtgat	840
tattacacat tgcatgcctg tatcaaaata ttgcacatac tccatgaatg catacatcta	900
ctatgttccc acaaaaatta aacattagaa aaaagagttg cattttcagc tgttatgggg	960
agaagaaaga aaagctatca ttttgttgtc ctaaaaatta tgttgtcctc atttcaaaca	1020
ggaaagcaaa agtatttgag agccagtga gtccttgggt gttgggtgaa acatagattg	1080
aatttgggcc atttgtttta acttcctagg cctcagtttc ttgcctatta aaaggagtg	1140
catagtccat gggattgtta agaggaagaa gtgaaacat gcacgtggag agcgtggcac	1200
agtgtctaag acagagtgtg catgcaaata agtagataat attccttgct tttctttatt	1260
gcatgcctgt aatatttttg gagttgtcac attcattgcc ctcaagtagc atcaagggat	1320
gaaattatgt ttgtaagaaa atcctgaggc tgaggaatac aacatgtttt atgtctacta	1380
cactgaaaaa tgccggagtc agataaagaa tacagattct cctgaggatg gaaatcaaga	1440
tcttcgcctt caatatttaa caacattgag cttccaactt actatgggaa atattcatca	1500
ggcccctaaa ggttcctttt ggacagaaat tgcacttggt atatctgtat tcttagcaga	1560
cagtagacag cctggcacat cataaaggct taaggaatcc taaatatccc ttaaaattct	1620
cattttaag acaaaaacaa aacaaaaaaa aaaaacaaaa aaaaactgag gcatgggctt	1680
gaccaaata gtggtagaac caagagttaa accacttggt ttgaatccta aacctgagtt	1740
ttattttact tatttattta tttatttggt tatttatttt cagatgcttg gtcaaagaac	1800
agtgggagga gagggatggg cttccagcaa cttttattat tggcttattt tcttacagcc	1860
cattactttc tcttgggaaa atattaagca ggcactcaag gcttgaggcc cctgagtttt	1920
cacatccttt ctgaacctct gaacctgctt tccagcattc ttttatactt tgttttacct	1980
cctggtcagt aatgcctcac cctcagtctt ctctaaaagt gtggttaatg gcatcttcct	2040
gactatttga agaccactgg ccaaatccca ccagctcact catagaccat cccctactt	2100
tactttcttc aaaagactta gccctaccta aacttattta tatgtttatt ttctgcccac	2160
cagaatggca gcatagctgg ggaggcagag tctgttttgt tcattgctgt attcccaaag	2220
actagaacac caccaagcac acggtacagg tctcagtaat tattgtcaaa tttatgtgga	2280
tttgctttta aacaatatct tccatttact gagtgtttat gtggaagaac tgtactaaat	2340
tttaatgcat ttctttattc ctattcttaa aaccttcag caaggtggct ctaccacct	2400
cttttccgag cttcaggagc agttgtgcga atagctggag aacaccaggc tggatttaaa	2460
cccagatcgc tcttacattt gctctttacc tgctgtgctc agcggttcacg tgccctctag	2520

ctgtagtttt ctgaagtcag cgcacagcaa ggcagtgtgc ttagaggtta acagaaggga	2580
aaacaacaac aacaaaaatc taaatgagaa tcctgactgt ttcagctggg ggtaaggggg	2640
gcggattatt catataattg ttataccaga cggtcgcagg cttagtccaa ttgcagagaa	2700
ctcgcttccc aggcttctga gaggccccga agtgcctaaa cctgtccta cgcaggggct	2760
tgggtggccc gtcgctccct ggcttcttcc ctttaccag ggcgggcagc gaagtgtgc	2820
ctcctgcgtc cccacacccc tccctcagcc cctcccctcc ggcccgtcct gggcaggtga	2880
cctggagcat ccggcaggct gccctggcct cctgcgtcag gacaacgccc acgaggggcg	2940
ttactgtgcg gagatgcacc acgcaagaga caccctttgt aactctcttc tcctccctag	3000
tgcgaggtta aaaccttcag cccacgtgc tgtttgcaaa cctgcctgta cctgaggccc	3060
taaaaagcca gagacctcac tcccggggag ccagc	3095

<210> 244

<211> 1807

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 244

agcttgtaga tactcggaac aaatgcaatt cttacgaata cttttagtct atacacagaa	60
aaagctggct gaaaaataaa atgattatct ttaatatctt aacagttatt aattgtgtgt	120
atgtggcagg cctgtgacag gtagaggaca acttgccata ggcacccatgt gggttccgaa	180
ggatctaact tgtcccatgc ttggcagcaa gcatttatca ctggccatct tcccagtcct	240
agctgtagtt tgcagtatat ttatactgc agcagccact ggcttgtgtg ggagctagt	300
cctagaccaa accaggattg cttctcttga aaccctctgg cactcattac gtgcttgatg	360
aataaatgga tggacagggt gctgtgtaca tttctctcac ttctcagttt ctttcagtaa	420
atcccaaat atcattttcc ttcagaaatt ctggcatgat tcattccggg tcctgccctg	480
gcatgcctt ctgtgtttct cattcagtaa gaagtcact cagatttagt tcacattaaa	540
aaataaacag agctttgata tccaaatgtc aacttgcagg gtattagaga agatagggaa	600
ttgcaatctt acatacgatt ttccccgatt ttcagccttg agatttcgtc cttgaaagca	660
tatggcaaat gtgcatccct ctttgaaatg tactaagatg taaaggggaa tttgaatgta	720
ttaaagtttg cagcaaagag aatataaatg taaacaagaa agaacagtta aatgtgtgag	780
tggatatggg gatgggtaga atgagagacg ggaacccatgt atgtgcgtcg ggatggatag	840
gaaatatgat gaacagatat agctgaggag ggggtgtgaaa aggattgaaa agttgtgcag	900
gtgggcgaat acaagaattg gtgggcagggt gtagtatggc tagattagt ctttgcaga	960
aggaagatgg gtggacagag gaatggatgg gtggattgtg agtcgagaag gatttaagaa	1020
attgtagat attttgagag catgaatgaa atgtgttgag cacccttggg ttttccccgg	1080
atcaaagatc agatgagcgg tttggacttc tctcagaggg aaagaggaaa gaacactccc	1140

acaagttccc cacttttcag tccccaccct ggccaggaaa gcactctcca ctaggatgga	1200
tctctctagt ctctctctct cccttcagcc tctttctttc ttcagttcct ccctaagata	1260
agtccagctt cctcagcttc ctgggaaaac cagtctttcc ctagccaggt tcccaagttt	1320
agtgggaaag gagaaactgg aagatttaac tgagaggggc gaggtcttag aactcagtca	1380
ttctccttgt cccaggcagc gcttctcata ggctggtagg ctgggccagg gtaggaagcc	1440
tgtggagtgg ccctggagaa cgtggggcgg cacgggggct ggggggggag gggggcggcc	1500
attctcttct gtccaagaga gcagggcagg agtgcagggg cagtagcgaa agcaggctgg	1560
tgtgtcttta aacttccgtt ggctgcttag tcacagcccc ctgcttttgg gtgtgtcctt	1620
cgcgcgctcc ctccctctta ggtcactcac tctttcaaag cctggaataa aaaccacagc	1680
caacttccga agcgggtctca ttgcccagca gccccagcc agtgacaggt tccattcacc	1740
ctcgttgccc ttctccccac gaccttttcc cagaggcgac tagatccctc cgtttcatcc	1800
agcacgc	1807

<210> 245

<211> 3000

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 245

gaaaatttgt cacaactaa agaaaacaag aaagagacag tagatgaaag agtgctcatt	60
aggtgaaagg aaaatgatcc aagagggtag ctttgagatg taggaagaaa caaaaagcaa	120
gaaaatgata aatgttttga taaagctaaa taagtatcaa ctcataaaga aataatattc	180
ccagaagagt catgaatata cagagaaaat taaagtacat gacaatggca atgtaaaagt	240
taggggtgaa taaaaaagag acttaagagt tctaaaatca ttgcattgtc ctggaagagg	300
aaaaagtaca atgattagtc aaagatacat gtcataatcc ctagaaagga gatcattatt	360
aaatagaaaa taaaagaata catcttatag aaaggaaatc taaatgataa tattaacag	420
atctaaaata aggcaaaagt gaggataaaa aagaaagatg gaaccaatgg ggcaaataga	480
aaaagtaaga tagcgtggta gggcattaat tccagcctta catcaatgca taagtatctc	540
aatattctac tgtaaaggga aagtaaagat ttcttacagc ctgagtgtaa tggagaaatc	600
tagtttatca tagtgcttta aatattgtaa gtcttcaact tctagttgat gaataaatga	660
tggaattctc agtgatactg cactgttatc aaataaatat aaaaggagct cctggaattg	720
gatgtaatac aggtaaagaa gtaaacacag ccatataggc atggcttctt gcagggacaa	780
ctttgtgaat cggctcagac agacagacag gcaaatacac ctcatcgcct catacatgtt	840
atttgcttta gtttttgttc tgaaccttcc tactccttca agtatctgca ttacttttat	900
caaattctct tttagtagag actgaagaaa ctgtcatctc cttatgtgct aatgagttta	960

ataatgtcct ccagtcacca caagccttct ttcaaactac acaattccaa ctgcttccgt	1020
ctcagagtat cttgaaataa tgatctgacc gcctgttaga ccagtgaagg gaaggaattt	1080
gggttgattt aagaagagaa tcctcatggt catggtagac tgatatggag agaaaacatt	1140
ttgaggaaaa atactcaact aaattcattt ctactccagc atgcagtttc aagtcaagtt	1200
ccaccttagc tccaggtggc aggcagagca ggatgcagag gcacagcaca agtaaggggt	1260
gagtgccgaa gctgctggct cctgttccag tctttcttcc ttggcctcgc ctgaactttt	1320
actataataa tagtcaccat ttattaggtg tctcctacgt gcaggacact ttacacacag	1380
tatccctaata cctaataaca cccttatttt atagatccaa tgactgagtc aagaattaca	1440
taacctggcc agacagctgg tacatgggaa aggtgagatt cacaccaggg tccaccagc	1500
atctctactt ataccatgct ctgctttaag gttctctgag aactcagaca agccttgggc	1560
taacaattgt gttacagga catagcaggt gcaaggacct actggtcatc ctgctacctg	1620
atcagaagga aggaaagttg tatttgttg tcacctacta tgttttaggc atagtactag	1680
gtgcttttac ctagtactta attcccttat cctcaactca tttattcctc gcaataacct	1740
gataagggag atgtttttat cctcatttta catataagga aacaggccta gagaaatgag	1800
cacagtgtcc aaagtcacat agttaataag atgtgaagct ctgagtttga aagtctccgg	1860
tttcaaagcc atgaaactta tggctccccg ttttagacac ttccttttgg gaagagtgtg	1920
gaggaattaa tcagaaagaa gaaagtcata ctcaaatagg tggtaggagc agagacaatt	1980
caatacagac agaagtctta gatgagagca gtgagccagg gcaactggact gggactcagg	2040
aggcttcccc tagactctgg ttccaccgat gcagcctcag gcaggacttc acctctctgg	2100
gcataccgttt cttcatatgt taaacatacg gggttttaat tagatgatcg ctgaagacct	2160
ctctagccct aaaactctgt gtctcttaag tgctaagagg gcaccaacag cgttcctcct	2220
ccccaaggag cataatgtga tggttcctgc cggccctggc tgactctcgc cgtccttggg	2280
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tattgctctc tctcggctct gggccgccct gtgattgttg ggcgtccgga aactgtctcc	2400
cctatgggtt taaaaacaaa actgagcgcc catgggggtgt gacagtcac tgcaggggct	2460
tgggtggccc atcaggcgag gctttctcgg caccagaggc tccagcctga tctcggctct	2520
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ccccggcggc aagagcggca ggctgcgccc ctggcccgcg cctagcctgg ggagagagct	2640
gggcgggagg cgggagctgc tctcgcgggc cgcggccctc gccctggctg caacggtagg	2700
cgtttcccgg gccggacgcg cgtggggggc gggggcgggg gcgggggcga ggccgcggcg	2760
agcaaagtcc agggccctct gctgcagcgc ccgcgcgtcc agaggccctg ccagacacgc	2820
gcgaggttcg aggtgagaga ggtccgggcg cgtctggcct cgaagggaga cccgggacgt	2880
ggggcgcggg gcgggagtg ccggacctcc acccagtgcc cccgggcccc gcgactcgtg	2940
cgccgggccc ccggagaggg tgtacttggt tctgaggctg tggtttctcc tcaggctgag	3000

<210> 246
<211> 757
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 246
tgaattgatg ctgtatactc tcagagtgcc aaacatatac caatggacaa gaaggtgagg 60
cagagagcag acaggcatta gtgacaagca aagatatgca gaatttcatt ctcagcaaat 120
caaaagtcct caacctgggtt ggaagaatat tggcactgaa tggatatcaat aagggttgcta 180
gagagggtta gaggtgcaca atgtgcttcc ataacatttt atacttctcc aatcttagca 240
ctaatacaac atggttgaat actttgttta ctataactct tacagagtta taagatctgt 300
gaagacaggg acagggacaa taccatctc tgtctggttc ataggtggta tgtaatagat 360
atttttaaaa ataagtgagt taatgaatga gggtgagaat gaaggcacag aggtattagg 420
gggaggtggg cccagagaa tggtgccaag gtccagtggg gtgactggga tcagctcagg 480
cctgacgctg gccactccca cctagctcct ttctttctaa tctgttctca ttctccttgg 540
gaaggattga ggtctctgga aaacagccaa acaactgtta tgggaacagc aagcccaaat 600
aaagccaagc atcaggggga tctgagagct gaaagcaact tctgttcccc ctccctcagc 660
tgaaggggtg gggaagggtc ccaaagcca taactccttt taagggattt agaaggcata 720
aaaaggcccc tggctgagaa cttccttctt cattctg 757

<210> 247
<211> 720
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 247
ccactacggg tctaggctgc ccatgtaagg aggcaaggcc tggggacacc cgagatgcct 60
ggttataatt aacccaaca cctgtgccc ccccccccc aacacctgct gcctgagcct 120
gagcggttac cccaccccg tgctgggtc ttaggctctg tacaccatgg aggagaagct 180
cgctctaaaa ataaccctgt ccctgggtgg cccactacgg gtctaggctg cccatgtaag 240
gaggcaaggc ctggggacac ccgagatgcc tggttataat taacccaac acctgctgcc 300
cccccccc caacacctgc tgcctgagcc tgagcggtta cccaccccg gtgcctgggt 360
cttaggtctt gtacaccatg gaggagaagc tcgctctaaa aataaccctg tccctggtgg 420
gccactacgg gtctaggctg cccatgtaag gaggcaaggc ctggggacac ccgagatgcc 480
tggttataat taacccaac acctgctgcc ccccccccc caacacctgc tgcctgagcc 540
tgagcggtta cccaccccg gtgcctgggt ctaggctct gtacaccatg gaggagaagc 600

tcgctctaaa aataaccctg tccctggtgg gccctccct ggggacagcc ctcctggct	660
agtcacaccc ttaggctcc tctatataac ccaggggcac aggggctgcc cccgggtcac	720

<210> 248
 <211> 772
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 248 acccttcaga ttaaaaataa ctgaggtaag ggcctgggta ggggaggtgg tgtgagacgc	60
tcctgtctct cctctatctg cccatcggcc ctttggggag gaggaatgtg cccaaggact	120
aaaaaaaggc catggagcca gaggggagag ggcaacagac ctttcatggg caaaccttgg	180
ggccctgctg tctagcatgc ccactacgg gtctaggctg cccatgtaag gaggcaaggc	240
ctggggacac ccgagatgcc tggttataat taaccagac atgtggctgc ccccccccc	300
ccaacacctg ctgcctctaa aaataaccct gtccctggtg gatcccctgc atgcgaagat	360
cttcgaacaa ggctgtgggg gactgagggc aggctgtaac aggcttgggg gccagggctt	420
atactgcct gggactccca aagtattact gttccatgtt cccggcgaag ggccagctgt	480
ccccgccag ctagactcag cacttagttt aggaaccagt gagcaagtca gcccttgggg	540
cagcccatac aaggccatgg ggctgggcaa gctgcacgcc tgggtccggg gtgggcacgg	600
tgcccgggca acgagctgaa agctcatctg ctctcagggg cccctccctg gggacagccc	660
ctcctggcta gtcacaccct gtaggctcct ctatataacc caggggcaca ggggctgccc	720
tcattctacc accacctcca cagcacagac agacactcag gagccagcca gc	772

<210> 249
 <211> 558
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 249 cagccactat ggtcttaggc tgcccatgta aggaggcaag gcctggggac acccgagatg	60
cctggttata attaaccag acatgtggct gctcccccc cccaacacc tgctgcctga	120
gcctcacccc cacccggtg cctgggtctt aggctctgta caccatggag gagaagctcg	180
ctctaaaaat aaccctgtcc ctggtgggct gtgggggact gagggcaggc tgtaacaggc	240
ttgggggcca gggcttatac gtgcctggga ctcccaaagt attactgttc catgttccc	300
gcgaagggcc agctgtcccc cgccagctag actcagcact tagtttagga accagtgagc	360
aagtcagccc ttggggcagc ccatacaagg ccatggggct gggcaagctg cagcctggg	420
tccggggtgg gcacggtgcc cgggcaacga gctgaaagct catctgctct caggggcccc	480

tccctgggga cagccccctcc tggctagtca caccctgtag gctcctctat ataaccagg	540
ggcacagggg ctgcccc	558

<210> 250
 <211> 766
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 250	
cagccactat gggcttaggc tgcccatgta aggaggcaag gcctggggac acccgagatg	60
cctggttata attaaccag acatgtggct gtccccccc cccaacacc tgctgcctga	120
gcctcacccc caccggtg cctgggtctt aggtctgtga caccatggag gagaagctcg	180
ctctaaaaat aacctgtcc ctgggtggct gtgggggact gagggcaggc tgtaacaggc	240
ttgggggcca gggcttatac gtgcctggga ctcccaaagt attactgttc catgttccc	300
gcgaagggcc agctgtcccc cgccagctag actcagcact tagtttagga accagtgagc	360
aagtcagccc ttggggcagc ccatacaagg ccatggggct gggcaagctg cacgcctggg	420
tccggggtgg gcacggtgcc cgggcaacga gctgaaagct catctgctct caggggcccc	480
tccctgggga cagccccctcc tggctagtca caccctgtag gctcctctat ataaccagg	540
ggcacagggg ctgcccccg gtcaccacca cctccacagc acagacagac actcaggagc	600
cagccagcca ggtaagttta gtctttttgt cttttatttc aggtcccga tccggtgggtg	660
gtgcaaatca aagaactgct cctcagtgga tgttgctttt acttctaggc ctgtacggaa	720
gtgttacttc tgctctaaaa gctgcggaat tgtacccgcg gccgcg	766

<210> 251
 <211> 961
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 251	
gtttaacaa gcttgcattg ctaagctaga cccttcagat taaaaataac tgaggtaagg	60
gcctgggtag gggaggtggg gtgagacgct cctgtctctc ctctatctgc ccatcgggcc	120
tttggggagg aggaatgtgc ccaaggacta aaaaaaggcc atggagccag aggggcgagg	180
gcaacagacc tttcatgggc aaaccttggg gccctgctgt ctagcatgcc ccactacggg	240
tctaggctgc ccatgtaagg aggcaaggcc tggggacacc cgagatgcct gggtataatt	300
aaccagaca tgtggctgcc ccccccccc caacacctgc tgcctctaaa aataaccctg	360
tccctggtgg atcccctgca tgcgaagatc ttcgaacaag gctgtggggg actgagggca	420

ggctgtaaca ggcttggggg ccagggttta tacgtgcctg ggactcccaa agtattactg	480
ttccatgttc ccggcgaagg gccagctgtc ccccgccagc tagactcagc acttagttta	540
ggaaccagtg agcaagtcag cccttggggc agcccataca aggccatggg gctgggcaag	600
ctgcacgcct gggtccgggg tgggcacggt gcccgggcaa cgagctgaaa gctcatctgc	660
tctcaggggc ccctccctgg ggacagcccc tcctggctag tcacaccctg taggctcctc	720
tatataaccc aggggcacag gggctgcctt cattctacca ccacctcac agcacagaca	780
gacactcagg agccagccag cggcgcgccc aggtaagttt agtctttttg tcttttattt	840
cagggtcccgg atccggtggt ggtgcaaatc aaagaactgc tcctcagtgg atgttgcctt	900
tacttctagg cctgtacgga agtgttactt ctgctctaaa agctgcggaa ttgtaccgc	960
g	961

<210> 252

<211> 1736

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 252

aaaagagtgc agtaacaaag ccccttttac aatttaccgc gcacattcac acccatcctg	60
aggccaaagc cacaggctgt gaggtctcac tgtctcagct tcctgagcta taaaatggga	120
atgatgctag tgtctacctc ctagggttgg agaattgggg gtcattgggtg tgaagtgctc	180
agcagcttgg cccacactag gtggctcagta catgtaaggt attattgttg ctacatacat	240
tagtagggcc tgggcctctt taaaccttta tagggtagca tggcaaggct aaccatcctc	300
actttatatc tgacaagctg gggctcagag aggacgtgcc tgagctgggg ctcagacaag	360
gacacaccta ctagtaaccc ctccagctgg tgatggcagg tctagggtag gaccagtgc	420
tggctcctaa tcgagcactc tattttcagg gtttgcattc caaaagggtc aggtccaaga	480
gggacctgga gtgccaagtg gaggtgtaga ggcacggcca gtacccatgg agaattggtg	540
atgtccttag gggtagcaa gtgccgtgtg ctaaggaggg ggctttggag gttgggcagg	600
ccctctgtgg ggctccattt ttgtgggggt gggggctgga gcattatagg gggtaggaag	660
tgattggggc tgtcaccta gccttcctta tctgacgcc acccatgcct cctcaggtac	720
cccctgcccc ccacagctcc tctcctgtgc cttgtttccc agccatgcgt tctcctctat	780
aaataccgc tctggtattt ggggttggca gctgttgctg ccaggagat ggttgggttg	840
acatgcggct cctgacaaaa cacaaccccc tgggtgtgtg gggcgtgggt ggtgtgagta	900
gggggatgaa tcaggagggg ggcgggggac ccagggggca ggagccacac aaagtctgtg	960
cgggggtggg agcgcacata gcaattggaa actgaaagct tatcagacct tttctggaaa	1020
tcagcccact gtttataaac ttgaggcccc accctcgaca gtaccgggga ggaagagggc	1080
ctgcactagt ccagagggaa actgaggctc agggctagct cgcccataga catacatggc	1140

aggcaggctt tggccaggat ccctccgcct gccaggcgtc tccctgccct cccttcctgc	1200
ctagagaccc ccaccctcaa gcctggctgg tctttgcctg agacccaaac ctcttcgact	1260
tcaagagaat atttaggaac aagggtggtt agggcctttc ctgggaacag gccttgaccc	1320
tttaagaaat gacccaaagt ctctccttga ccaaaaaggg gaccctcaaa ctaaagggaa	1380
gcctctcttc tgctgtctcc cctgacccca ctcccccca cccaggacg aggagataac	1440
cagggtgaa agaggcccg ctgggggctg cagacatgct tgctgcctgc cctggcgaag	1500
gattggcagg cttgccgctc acaggacccc cgctggctga ctcagggcg caggcctctt	1560
gcgggggagc tggcctcccc gccccacgg ccacgggccc ccctttcctg gcaggacagc	1620
gggatcttgc agctgtcagg ggaggggagg cgggggctga tgctaggagg gatacaaata	1680
gtgccgacgg ctggggggccc tgtctcccct cgccgcatcc actctccggc cggccg	1736

<210> 253

<211> 807

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 253

gacattgatt attgactagt tattaatagt aatcaattac ggggtcatta gttcatagcc	60
catatatgga gttccgcgtt acataactta cggtaaatgg cccgcctggc tgaccgcca	120
acgacccccg ccattgacg tcaataatga cgtatgttcc catagtaacg ccaataggga	180
ctttccattg acgtcaatgg gtggagtatt tacggtaaac tgcccacttg gcagtacatc	240
aagtgtatca tatgccaagt acgcccccta ttgacgtcaa tgacggtaaa tggccgcct	300
ggcattatgc ccagtacatg accttatggg actttcctac ttggcagtac atctacgtat	360
tagtcatcgc tattaccatg gtgatgcggt tttggcagta catcaatggg cgtggatagc	420
ggtttgactc acggggattt ccaagtctcc acccattga cgtcaatggg agtttgtttt	480
ggcaccaaaa tcaacgggac tttccaaaat gtcgtaaaa ctccgcccc ttgacgaaa	540
tgggcggtag gcatgtacgg tgggaggtct atataagcag agctcgttta gtgaaccgtc	600
agatcgcttg gagacgcat ccacgtgtt ttgacctca tagaagacac cgggaccgat	660
ccagcctccg cggccccaag cttcagctgc tcgaggcgcg gcctctagag ctagcgttgc	720
ggccgcctgg ctcttaacgg cgtttatgtc ctttgctgtc tgaggggcct cagctctgac	780
caatctggtc ttcgtgtggt cattagc	807

<210> 254

<211> 973

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 254

aagtcagcat ccattcctct ctgtggttct ccctccgccc catccaggtc tcaagggtct	60
agagtctttc aaagagaaca cattctgaga tttagaggagg cagagacaaa aagttccact	120
gcgaagtgcc agggaggcctt ctgtttgggg tgtcccttgg gatcacagat cccccacctg	180
gtgatgagtc aaccagcac caccctattg cagggtctgga atgacagtaa tgggcccacc	240
tgctgcctct cctcataccc gcaccccagt cagacattgc aagtcagtca cggctctgtc	300
ctgctgggcc tggagtgttc cagtgccttt tccatcacag caccaagcag ccactactag	360
tcgatcaatt tcagcacaag agataaacat cattaccctc tgctaagctc agagataacc	420
caactagctg accataatga cttcagtcac tacggagcaa gataaaagac taaaagaggg	480
agggatcact tcagatctgc cgagtgagtc gattggactt aaagggccag tcaaaccctg	540
actgccggct catggcaggc tcttgccgag gacaaatgcc cagcctatat ttatgcaaag	600
agattttgtt ccaaacttaa ggtcaaagat acctaaagac atccccctca ggaaccctc	660
tcatggagga gagtgcctga gggctcttgg ttcccattgc atccccacc tcaatttccc	720
tggtgcccag ccacttgtgt ctttagggtt ctctttctct ccataaaagg gagccaacac	780
agtgtcggcc tcctctcccc aactaagggc ttatgtgtaa ttaaaaggga ttatgctttg	840
aaggggaaaa gtagccttta atcaccagga gaaggacaca gcgtccggag ccagaggcgc	900
tcttaacggc gtttatgtcc ttgtctgtct gaggggcctc agctctgacc aatctggtct	960
tcgtgtggtc att	973

<210> 255

<211> 450

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 255

ctagactagc atgctgcccc tgtaaggagg caaggcctgg ggacaccga gatgcctggt	60
tataattaac ccagacatgt ggctgcccc cccccccaa cacctgctgc ctctaaaaat	120
aaccctgcat gccatgttcc ggcggaagg ccagctgtcc cccgccagct agactcagca	180
cttagtttag gaaccagtga gcaagtcagc ctttggggca gccatacaa ggccatgggg	240
ctgggcaagc tgcacgcctg ggtccggggg gggcacggtg cccgggcaac gagctgaaag	300
ctcatctgct ctcaggggcc cctccctggg gacagcccct cctggctagt cacaccctgt	360
aggctcctct atataacca ggggcacagg ggctgccctc attctaccac cacctccaca	420
gcacagacag aactcagga gccagccagc	450

<210> 256

<211> 455

<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 256
ctgctccag ctggccctcc caggcctggg ttgctggcct ctgctttatc aggattctca 60
agagggacag ctggtttatg ttgcatgact gttccctgca tatctgctct ggttttaaat 120
agcttatctg ctagcctgct cccagctggc cctcccaggc ctgggttgct ggcctctgct 180
ttatcaggat tctcaagagg gacagctggt ttatgttgca tgactgttcc ctgcatactt 240
gctctggttt taaatagctt atctgagcag ctggaggacc acatgggctt atatggggca 300
cctgcaaaaa tagcagccaa ccccccccc tgctgcacat tcctccctgg ctcaccaggc 360
cccagccac atgcctgctt aaagccctct ccatcctctg cctcaccag tccccgctga 420
gactgagcag acgcctccag gatctgtcgg cagct 455

<210> 257
<211> 3050
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 257
atttttcaag ataaaagtga aataaat tttt caggaaaaaa aagctgagaa aatttgtcac 60
aaactaaaga aaacaagaaa gagacagtag atgaaagagt gctcattagg tgaaaggaaa 120
atgatccaag agggtagctt tgagatgtag gaagaacaa aaagcaagaa aatgataaat 180
gttttgataa agctaaataa gtatcaactc ataaagaaat aatattccca gaagagtcac 240
gaatatacag agaaaattaa agtacatgac aatggcaatg taaaagttag gggatgaataa 300
aaaagagact taagagttct aaaatcattg cattgtcctg gaagaggaaa aagtacaatg 360
attagtcaaa gatcatgtc ataatcccta gaaaggagat cattattaaa tagaaaataa 420
aagaatacat cttatagaaa ggaatcttaa atgataatat taaacagatc taaaataagg 480
caaaagttag gataaaaaag aaagatggaa ccaatggggc aaatagaaaa agtaagatag 540
cgtggtaggg cattaattcc agccttacat caatgcataa gtatctcaat attctactgt 600
aaagggaag taaagatttc ttacagcctg agtgtaatgg agaaatctag tttatcatag 660
tgctttaaat attgtaagtc ttcaacttct agttgatgaa taaatgatgg aattctcagt 720
gatactgcac tgttatcaaa taaatataaa aggagctcct ggaattggat gtaatacagg 780
taaagaagta aacacagcca tataggcatg gcttcttgca gggacaactt tgtgaatcgg 840
ctcagacaga cagacaggca ggcaaataca cctcattgcc tcatacatgt tatttgcttt 900
agtttttggt ctgaaccttc ctactccttc aagtatctgc atttacttta tcaaattctc 960

ttttattaga gactgaagaa actgtcatct ccttatgtgc taatgagttt aataatgtcc	1020
tccagtcacc acaagccttc tttcaaacta cacaattcca actgcttccg tctcagagta	1080
tcttgaaata atgatctgac cgcctgttag accagtgaag ggaaggaatt tgggttgatt	1140
taagaagaga atcctcatgg tcatggtaga ctgatatgga gagaaaacat tttgaggaaa	1200
aataactcaac taaattcatt tctactccag catgcagttt caagtcaagt tccaccttag	1260
ctccaggtgg caggcagagc aggatgcaga ggcacagcac aagtaagggg tgagtgccga	1320
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atagtcacca tttattaggt gtctcctacg tgcaggacac tttacacaca gtatccctaa	1440
tcctaataca cccttatttt atagatccaa tgactgagtc aagaattaca taacctggcc	1500
agacagctgg tacatgggaa aggtgagatt cacaccaggg tccaccagc atctctactt	1560
ataccatgct ctgctttaag gttctctgag aactcagaca agccttgggc taacaattgt	1620
gttaacagga catagcaggt gcaaggaccc actggtcatc ctgctacctg atcagaagga	1680
aggaaagttg tatttgttgc tcacctacta tgtttttaggc atagtactag gtgcttttac	1740
ctagtactta attcccttat cctcaactca tttattcctc gcaataacct gataaggag	1800
atgtttttat cctcatttta catataagga aacaggccta gagaaatgag cacagtgtcc	1860
aaagtcacat agttaataag atgtgaagct ctgagtttga aagtctccgg tttcaaagcc	1920
atgaaactta tggctccccg ttttagacac ttccttttgg gaagagtgtg gaggaattaa	1980
tcagaaagaa gaaagtcata ctcaaatagg tggtaggagc agagacaatt caatacagac	2040
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tagactctgg ttccaccgat gcagcctcag gcaggacttc acctctctgg gcatccgttt	2160
cttcatatgt taaacatacg gggttttaat tagatgatcg ctgaagaccc ctctagccct	2220
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taaaaacaaa actgagcgcc catgggggtgt gacagtcacg tgcaggggct tgggtggccc	2520
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aagagcggca ggctgcgccc ctggcccgcg cctagcctgg ggagagagct gggcgggcgg	2700
cgggagctgc tctcgcgggc cgcggccctc gccctggctg caacggtagg cgtttcccg	2760
gccggacgcg cgtggggggc gggggcgggg gcgggggcga ggccgcggcg agcaaagtcc	2820
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aggtgagaga ggtccgggcg cgtctggcct cgaagggaga cccgggacgt ggggcgcggg	2940
gcgggagtg ccggacctc acccagtgcc cccgggcccc gcgactcgtg cgccgggccc	3000
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<210> 258
<211> 3000
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 258
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ccccaacccg cagagcaggt tgtgggggca atgttgcatc gaccagcgtc agagaacaca 120
catcagaggc aaggggtgggt gtgcaggagg gagaaggcgc agaaggcagg gctttagctc 180
agcactctcc ctctgcatc gctctgcctg accgtttcct ctctgagtcc caaacagcca 240
ggtagaggag gaagaaatgg ggctgagacc ccagcacatc agtgattaag tcaggatcag 300
gtgcggtttc ctgctcaggt gctgagacag caggcggtgt cctgcaaaca acaggaggca 360
cctgaagcta gcctggggggg ccacgcccc ggtgcggtgc attcagcagc acagccagag 420
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cccagacctt ggaggcccc tccggctccg cctcctggga gaaggctctg gagtgaggag 600
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agacacacac agagagacgg cgaggaagg aaagaccag agggacgcct agaacgagac 720
ttggagccag acagaggaag aggggacgtg tgtttgcaga ctggctgggc ccgtgacca 780
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ttgtccatcg aatcttctct gagctggggc tggggtagt ggcatcctgg ggccagggga 1140
atagacatgc tgtggtggca gagagaagag tccgttctct ctgtctcctt tgctttctct 1200
ctgacactct ttatctccgt ttttggataa gtcacttctc tcctctatgc cccaaatatc 1260
ccatctgtga aatgggagta tgaagcccc acagccaggg ttgtagtggg gaagaggtaa 1320
aatcaggatat agacatagaa atacaatac agtctatgcc ccctgttgtc agttggaaaa 1380
gaaattaact tgaagggtgt ctagttctca tttttagaaa tgaaatgtct gtctggtcat 1440
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cgctgtctcc acagcctggg caccatcttc agacaagtgc ccggtggcaa ctgcctgctg 1620
gccctgttga atccacatct ccaccaggca tccagactag ttcagggtctc tggaaggact 1680

gtgggtttgc tgtgtcccag agctccaggg caggggtcag ggctcggatg tcgggcagtg	1740
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cgcattctag gagacaggtt gggagccaga ggggcctcat accagggtct gtaggctggg	1860
gctgcctttt aagctccttc ctgaggccgt ctctgggtct ggccctgtgc tggacaaggc	1920
tggagacaag gcaatgtctc agaccctctc ccattggcca catcctgccc tggatcaact	1980
cgccaacttt gggggcagag gtgggactga cccttaccct gacaacataa tgcataatag	2040
caaaatggga taaaggggaa tatagaggct cttggcagct tgggagtggg cagggaaggc	2100
ttcctggagg aggtatcatc tgaactgagc catgaacat aagtggaaat tcactagtca	2160
aaatttcagg tagaagggcc agtgtgtgaa ggccaggaga tggcaagagc tggcgtat	2220
caggaacagt gagtactga ggatgtcaa gtataagggt aggaaaggga gtgagcagtg	2280
agagaaaaga ccgaggcatc agcaggggcc agatttgtct gggcctagcg gggcgggccc	2340
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tgaccctgcc tccccgtcag cagccagtgc tctgcacaga gccatcctga gggcagtggg	2460
tgctcttgag aggtttcagg cagggtgtgc tgtgagcagg tcatgcccag cccttgacct	2520
tctgctcagt caggcttgct cttgtcacc acattcctgg ggcagtcctt aagctgagtg	2580
ccggagatta agtcctagtc ctaaatttgc tctggctagc tgtgtgacct tgggcaagtc	2640
ttggtccctc tctgggcccc tttgccgtag gtccctgggt gggccagact tgctactttc	2700
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gcagtgggtgc tggtgacagt ggctgaggct caggtcacac aggctgggca gtggtcagag	2820
ggagagaagc caaggagggt tcccttgagg gaggaggagc tggggctttg ggaggagccc	2880
aggtgacccc agccaggctc aaggcttcca gggctggcct gccagaagc atgacatggt	2940
ctctctccct gcagaactgt gcctggccca gtgggcagca ggagctcctg acttgggacc	3000

<210> 259

<211> 3000

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 259

taataggcag agtttcttaa tgtggactag agttgcta cttagattat ccatttgagt	60
catgatttcc tactatacaa agcaggagtt gttatggggg agaagaattt ttatcccagg	120
aatgacaaag ataagttgaa gcactacagt aaaaaattag agttagacat ggacacgtag	180
aagggaacaa cagactctac agactctagg acctacttga ggctgaaggg tgggaggagg	240
tggaagattg aaaaactacc tatcaggtag tgtgcttatt acctggatga tgacataatc	300
tgtacatcta acccccatga cacacaattt acctatataa caaacctcca aatgtacccc	360
tgaacctaaa ataaaagttt agaaaaaatg agaattagtt cttggattca caagatataa	420

agagaagcca gccattgaat accttgtttg aaagtaggtt gacttcatgt tttgtagcag	480
gtctgaataa tccatttgtc taattcactg tgctctataa tacctatttt caaagatagt	540
ttcccaagtt ctgagaagtc cttacatatt agctgacttt atactaaaat ttgggtttaa	600
aaaaattttt ttttagagac atgggtctcac tctgtcatcc aggttaaagt gcagtgggtg	660
tgtgataata gtttactgca gcctcgaaat cctgggctca acaaccctcc cacctcagca	720
tcctaagtag ctgggactac gagtgtgtgc caccatgcct ggcttaaatt tttttatttt	780
tatttttatt tttatttttt ttttggagac gtgggatttc actatgttgc acagcatggt	840
cttgaactcc tggcttcaag caatcctccc accttggcct cccaaatccc taggaggcac	900
aagcatgagc catttgtctt tgccctaaaa tttgttttaa attaaagttt ttctggtaag	960
aatgtaatag cgtattttga caaagggtga gaaaggcttc ttctggaagc aactaatgct	1020
aattgataaa attgatatat aaatgggttg tggtttcag ctctcttctg ggagagaaat	1080
aaaagggaat ctaataaaga acaatgttgg tttttctctg gctgctttac taacaagaaa	1140
caccatgaaa catttctctc atttctaaac atttctataa aaaagataac ttatagagaa	1200
caaaatcaca atcgaccagt tatttcccaa acaaattttc catttttaca atacaaagg	1260
aaagctacaa gtattagctg atttagaata tttctcatct aggatgagat gtcccagatg	1320
gcagagtaga gagagttttg gatataattg aaactctata gaattgggtg caaatgtgca	1380
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agatgctact cataatgtaa ggaagcatgt attatggaga ggttatcctg ggtgaaaggt	1560
acagcaacaa ctgaatagtc aaccgaaact tctatcaatg ggccaagctt tgggagcatc	1620
aatatataaa agtttagaat tccattttgt atcctcttct cccccaaaaa gaaagagcac	1680
tggaaattat tccttggtg gtgtttaata gtggtagatc attttgatta aggaattaaa	1740
tggattgagg tgcatgagag caagaaagag gaggggcaag aggggggatt ataggataag	1800
gtgtactgct actttaaaat tatgtatgca tgatcccatc caggtcctc cactgcttg	1860
aggtaccagc ggaaagcttg ggcagctcag ttccaagagg gccaccaagc agaccacgct	1920
ctgagcttca ggtaaccaag tgtttgtctt gcagaatact ttacctgggc acccaagtct	1980
tccttcagc attcctgctg ctacagccta tttgctgagt aaccaggggt tacagcagcg	2040
ttgccaggca acgagggaca gcggtcctgt tgaagagcca tttgtcacac tgaggggact	2100
ggttgaaatg caataaagaa atggtaactc agcttattta tcaatacaat tacttgca	2160
gtattagga tccatgtgta acctacaaat tcatagtcac atgaggaaac acagaaacat	2220
tttgctaaat attaaagcat aggacagaca gatggtgttg ggtttctaata cagctttact	2280
ctgagcttaa agttgctgca catgctggga taaggggaaa ggcccaaagt cctttgccag	2340
ctttattttg ggcatctgta agttagctct gggttacaat gtacagtgca tgtgtaaaga	2400
aatctacaa gattcttttc cctgttaagt agagctggta atgccattgc taattccctg	2460

gggtgaagta acaacacaaa attattgtat gtgtaatata ttattaataa ttatatatat	2520
ataaaacaca cacatatatt atataaatat ttatgtataa ctggttataa atattactgg	2580
ttgtcctgtg gacttataaa gtgcttgatt tgcccaatgc aatcaagaga tttaccaaaa	2640
ggatgagtat tttactctga gcaactgtgct tcaaaatggt ttttgagaag ttcagtagtg	2700
ttgcttctag gagctcaaag tcctcaggcc tgggatgagc ttcagtttta aagggtgcagc	2760
agctttccct tgacgcccta cgTTTTTgat tcccagatac cagcagctac tcatgtcttc	2820
gccattgcta agaacgtcgt tggattacc ttactctgag aacgtgtctg cagtttccag	2880
aaaatggagt atcgcaacat cacttaaagt accctgcttc aaagtattgc tggcaagtgg	2940
cgtgggcctg attattttatt tagaaatgct ttatcaggag gagaatgctt ttttgtaaac	3000

<210> 260

<211> 1053

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 260

cgttacataa cttacggtaa atggcccgcc tggctgaccg cccaacgacc cccgcccatt	60
gacgtcaata atgacgtatg ttcccatagt aacgccaata gggactttcc attgacgtca	120
atgggtggag tattttacggt aaactgcccc cttggcagta catcaagtgt atcatatgcc	180
aagtacgcc cctattgacg tcaatgacgg taaatggccc gcctggcatt atgccagta	240
catgacctta tgggactttc ctacttgga gtacatctac gtattagtca tcgctattac	300
catggtcgag gtgagcccca cgttctgctt cactctcccc atctcccccc cctccccacc	360
cccaatTTTg tattttattta ttttttaatt attttTgtga gcgatggggg cggggggggg	420
ggggggggcgc gcgccaggcg gggcggggcg gggcgagggg cggggcgggg cgaggcggag	480
aggtgcggcg gcagccaatc agagcggcgc gctccgaaag tttcctttta tggcgaggcg	540
gcggcggcgg cggccctata aaaagcgaag cgcgcggcgg gcgggagtcg ctgcgcgctg	600
ccttcgcccc gtgccccgct ccgccgccgc ctgcgcgccgc ccgccccggc tctgactgac	660
cgcgttacta aaacaggtaa gtccggcctc cgcgccgggt tttggcgcct cccgcgggcg	720
ccccctcct cacggcgagc gctgccacgt cagacgaagg gcgcagcgag cgtcctgac	780
cttcgcccc gacgctcagg acagcggccc gctgctcata agactcggcc ttagaacccc	840
agtatcagca gaaggacatt ttaggacggg acttgggtga ctctagggca ctggttttct	900
ttccagagag cggaacaggc gaggaaaagt agtcccttct cggcgattct gcggagggat	960
ctccgtgggg cggtgaacgc cgatgatgcc tctactaacc atgttcatgt tttctttttt	1020
tttctacagg tcctgggtga cgaacaggct agc	1053

<210> 261

<211> 3000

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 261

gcttgctact gaaaagctaa ggccagaggt aaagactatg gatttgggga atgaatattc	60
tgtgaagcca taagataatg gcctgaggtg ctgaggacca gtagtgctag gaactttgca	120
tccatgacta tagggctctt tagaactgtg ccacagtaca gcatcatgca gtagaatcta	180
agttgttctt tgtaataatg aatgccagca atattttaaa ataataataa taccattaaa	240
aagtgggcaa aggacatgaa tagacatttt tcaaaaggaa acatacaaat cgccaagaag	300
tatatgaaaa attaacagtt aatgttcatt gaatacttat tgcaggctag gtactgagtt	360
gagcattttg catgcatcat ctcaacttaa ataatgtatg tcccagcctg gccaacatgg	420
tgaaccccca tctctactaa aaatacaaaa attagccaga catggtggta catgcctgta	480
atcccagcta ctcaggaggc tgaggcagga gaattgcttg aatctgggag gcagaggttg	540
cagtgagccg agattgcacc actgcactct agcctgggtg acagagcgat actctgtctc	600
aaaagataat aataataaaa taatgtatgt caattgttga aattttggaa aatgaacaag	660
tgtgtgtgtg aataactggg tgtattctat acatatggct ttataactta cctattaact	720
taaggtcatt aatgcaatgt catcaaatac tctttggatc atctagattg ttgcacatta	780
tcctataata tgagatgcca caatttattt acacagtcga caattgtaac ccagcttgct	840
tttggctttt actgttttac ataatacttg gtaaaaatcc tcatataaat atttgaaaat	900
ttcctaagtg tccatttgtg aatgtaaaaa ttattttaga gatctaagat ttggtgcaaa	960
acttgcaatc agctacatag ttctacttga ggcaattttc actcaaaata tatcataaac	1020
catagtacaa aaatagagca tagacctctc cttgtgaagc agttgttttt gccttacatt	1080
tttttttttt tttttttttt ttgagatgga gtctcgctct gtcgccggg ctggagtgc	1140
gtggcgcaat ctgagctcac tgcaagctcc gcctcccggg ttcacgcat tctcctgcct	1200
cagcctcccg agcagctggg actacaggtg cccgctacca cgtctggcta attttttata	1260
tttttagtag aaacgggggt tcaactgtgt agccaggatg gtctcgatct cctgacctcg	1320
tgatccgccc acctggcct cccaaagtgc tgggattaca ggtgtgagcc accgtgcgtg	1380
gctgccttaa atttttaata atcattgtgc aaattattta gcactccagt gttttgattt	1440
ttctcctctg ctgggtagga ataacaataa tactgttatt caccatggtg gtgtgggaag	1500
tttcaaagag cacatgtcta taaagtgctt agtgcaaggc ttggcatgca gttaacacaa	1560
aataaatgcg agctgctgtc attaacaata ctgactacac ggcaactgtga tgcttatgta	1620
aatgccaggc tgtgtgtctg taacctgagg tatttgtgta aatattttcc taaaataaat	1680
ctaactaagg ttgttcttct cacttgtatg gggatcatct atgcggtaga tgctcaaaca	1740
caaattccag atacagagtg ggcagtggta gttaggaaga tagaaaggct agggagtgtt	1800

cctgggaagt cagtaaactt ggaagatcta aggttatatt aaaaatgttg tatcagaaca	1860
aaggctcagg acgttagtgt tagcagaaac cagatatctt agagcagtgg tttgtcaact	1920
ttgccagcaa tccacagtaa gaaattcaac tccggccggg cgcgggcctg taatcccagc	1980
actttgggaa gccgaggcgg gtggatgact tgaggtcagg agttcgagac catcctggct	2040
aacacagtga aaccccgctc ctactaaaaa tacaaaaatt agccgggcgt ggtggtgtgt	2100
gcctgtaatc ccagctactt gggaggttga ggcaggagaa tcacttgaac acaggaggcg	2160
gaggtgacag tgagccgaga tcgtgccatt gcactccagc ctgggtgaca gagggagact	2220
ctatctcaaa aaaagaaaaa aaagaaattc aactccacta acaccacaa tgcaaataaa	2280
tgtgtgaatg tgtacaacta ttttatcaag cagtacttat tatatgtgct gtaatctgat	2340
attttatagc ctgtttcatt ttattttaat gttgattgtt acccactaaa tttatttcat	2400
tgagaccccc taatttgaaa tattgccttg aatatatata tacatatata tacacatata	2460
tacatatata tacacacata tatacacata tatacacaca tatatacaca tatatatata	2520
tatatacaca tatatacata tatacacata tatacatata tacatatata tacacatata	2580
tacatatata cacatatata catatatata catatatata tatatacaca tatatacata	2640
tatacatata tatacatata tacacatata tacatatata catatatata catatatata	2700
tatatatata cacatacata tatatatata cccttgttta aaaataaaaag gtttgcagct	2760
ccatattttt taaaaaatc ttaccaagc atttaatcag tactgaatgg ttttgttctt	2820
gtcttcatgt caagtngaatt ttgggggtac tattccagaa tattttacatg ttagacaatg	2880
ttctgtaaaa ggggcattgt agcagcatgc aggcagtatt caacaaaaa ctgggcaaga	2940
gtcataattc actctggttt ctctttcctt ttaagcaggt agttccaatt tgccagcaga	3000

<210> 262

<211> 3102

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 262

cgggagtcct gagggtagca gaagggtgcg gatttaaagt tactgttaga gtggctggaa	60
aatgggagac cggttcagag acattttatc tacttaaaaa ctgtgccttt tgtatcacgt	120
caaagtgaat gcaaaacaaa gaacaaaagg gttaaaggct caggtttaaa tcccaggtat	180
atgtacattt caattgaggt attttttttt tcttttctaa atgatcagta cacttattct	240
ttctaaagaa aatacttttc ttaactactc tctattttta aacttctccc acaaagatga	300
gaaaacattt aaaaatcatt ggggctattt ttctgtttac cgagtaaaga gaatctctaa	360
accatattta taactcttac tctaaatatt tgcatctacc ctcatgccag agcccgttga	420
tgactgacta aacagagttt caaagtttga agaacaggaa atttagaaat gactaacaat	480
tatgtagggtt tatttctctc agtatagaat gttcatatag aattaatgcc agaggttttc	540

agagaaaaat gcagaaat	ttactttgca aatccagaag	atgcaattgt tcaagtattt	600
gttaagaaac attaat	ttta agtatgcaga tatcattgag	aattaaatat tttaatttct	660
aaactattaa tcttttagta	ggatgcacat atgcaaaatg	cctcattagt actgtaagaa	720
aagattcttg gccgggcgcg	gtggctcatg actgtaatcc	cagcacttag ggaggccgag	780
gtgggcggat gacgagg	tca ggagatcgag accaccctgg	cacacgg	840
ctactaaaga taaaaaa	at tagccgggcg tgatggcggg	cgctgtagt cccagctact	900
cgggaggctg aggcaga	aga atggcgtgaa ctcgggaggc	ggagcttgca agtgagccga	960
gatagtgcc	ctgcactcca gtctgggcga	aagagcgaga ctccatctca	1020
aaaaaaaaaaga	aaagattc	tttaggtttca tcaat	1080
tttagatata	gaaaatcaat tcaaag	tttc tattcag	1140
taggtttc	ttt tgtatt	taac aatatattac	1200
ctaatattat	tcttttcatt	tgtaa	1260
ataaagcct	gaattt	gtca attag	1320
gggtaggcct	ttcccagga	cttctacaag	1380
ataaataatg	cctacaattt	ctaggaagtt	1440
ttattttc	ttt aacttaga	at agttt	1500
agaaaaatg	cttg	ttttt	1560
ctgtagtcag	aaagacaagt	ctgg	1620
aatcttccta	tctatctatg	tatctatcat	1680
gctttattag	ttaattagta	cccatttctg	1740
aatttctttc	attgtaagac	tgaatagaaa	1800
aaaaataata	ctttgttagt	ctctgtgcct	1860
ttttgactga	ctgagttata	taattaagta	1920
tgtagtatca	gagaaagttg	ttcttcctac	1980
agcaaaagg	gagaggtagg	atagcttttc	2040
aggaagattt	atgaaatatt	taaaaaatta	2100
gttaaattta	ttttgaaatt	taaagcaaca	2160
aaagagagaa	aggaaaaaaa	tatgaaacag	2220
tcaagggttc	tcataaccta	cagagaat	2280
caaagataat	catcatctca	tttgggtcca	2340
tcccatgaga	tataactcaca	ctgaatctaa	2400
gggccacagc	aggaatggga	acatggaatt	2460
tatgtctttt	tcttagtttg	catctgaggc	2520
ggaatagtaa	aaaagacaca	gaagccctaa	2580

tgcgtagta cttgtgtgta aatttttcat tatctatagg taaaagcaca cttggaatta	2640
gcaatagatg caatttggga cttactctt tcagtatgtc ttatttctaa gcaaagtatt	2700
tagtttgggt agtaattact aaacactgag aactaaattg caaacaccaa gaactaaaat	2760
gttcaagtgg gaaattacag ttaaatacca tggtaatgaa taaaaggtac aaatcgtttt	2820
aactcttatg taaaatttga taagatgttt tacacaactt taatacattg acaaggtctt	2880
gtggagaaaa cagttccaga tggtaaatat acacaaggga tttagtcaaa caattttttg	2940
gcaagaatat tatgaatttt gtaatcggtt ggagccaat gaaatacaaa gatgagtcta	3000
gttaataatc tacaattatt ggttaaagaa gtatattagt gctaatttcc ctccgtttgt	3060
cctagctttt ctcttctgtc aacccacac gcctttggca ca	3102

<210> 263

<211> 2337

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 263

tctagcttcc ttagcatgac gttccacttt tttctaaggt ggagcttact tctttgattt	60
gatcttttgt gaaacttttg gaaattacc atcttcctaa gcttctgctt ctctcagttt	120
tctgcttgct cattccactt ttccagctga ccctgcccc taccaacatt gctccacaag	180
cacaaattca tccagagaaa ataaattcta agttttatag ttgtttggat cgcataggta	240
gctaaagagg tggcaaccca cacatcctta ggcatgagct tgattttttt tgatttagaa	300
ccttcccctc tctgttccta gactacacta cacattctgc aagcatagca cagagcaatg	360
ttctacttta attactttca ttttcttgta tcctcacagc ctagaaaata acctgcgtta	420
cagcatccac tcagtatccc ttgagcatga ggtgacacta cttacatag ggacgagatg	480
gtactttgtg tctcctgctc tgtcagcagg gcactgtact tgctgatacc agggaatgtt	540
tgttcttaaa taccatcatt ccggacgtgt ttgccttggc cagttttcca tgtacatgca	600
gaaagaagtt tggactgatc aatacagtcc tctgccttta aagcaatagg aaaaggccaa	660
cttgtctacg tttagtatgt ggctgtagaa agggatataga tataaaaatt aaaactaatg	720
aaatggcagt cttacacatt tttggcagct tatttaaagt cttggtgtta agtacgtgg	780
agctgtcaca gctaccaatc aggcattgtc gggaatgagt acacggggac cataagttac	840
tgacattcgt ttcccattcc atttgaatac acacttttgt catggtattg cttgctgaaa	900
ttgttttgca aaaaaaacc cttcaaattc atatatatta ttttaataaa tgaattttaa	960
tttatctcaa tgttataaaa aagtcaattt taataattag gtacttatat acccaataat	1020
atctaacaat catttttaaa catttgttta ttgagcttat tatggatgaa tctatctcta	1080
tatactctat atactctaaa aaagaagaaa gaccatagac aatcatctat ttgatattgt	1140
taaagtttac atgtgagtag acatcagatg ctccatttct cactgtaata ccatttatag	1200

ttacttgcaa aactaactgg aattctagga cttaaataatt ttaagtttta gctgggtgac	1260
tggttgga aaatttaggta agtactgaaa ccaagagatt ataaaacaat aaattctaaa	1320
gttttagaag tgatcataat caaatattac cctctaataa aaatattcca aagttgagct	1380
acagaaaatt caacataaga taatttttagc tgtaacaatg taatttggtg tctattttct	1440
tttgagatac agttttttct gtctagcttt ggctgtcctg gaccttgctc tgtagaccag	1500
gttggtcttg aactcagaga tctgcttgcc tctgccttgc aagtgcagg attaaaagca	1560
tgtgccacca ctgcctggct acaatctatg ttttataaga gattataaag ctctggcttt	1620
gtgacattaa tctttcagat aataagtctt ttggattgtg tctggagaac atacagactg	1680
tgagcagatg ttcagaggta tatttgctta ggggtgaatt caatctgcag caataattat	1740
gagcagaatt actgacactt ccattttata cattctactt gctgatctat gaaacataga	1800
taagcatgca ggcatctatc atagttttct ttatctggaa aaacattaaa tatgaaagaa	1860
gcactttatt aatacagttt agatgtgttt tgccatcttt taatttctta agaaatacta	1920
agctgatgca gagtgaagag tgtgtgaaaa gcagtgggtg agcttggtt gaactcgttc	1980
tccagcttgg gatcgacctg caggcatgct tccatgccaa ggcccacact gaaatgctca	2040
aatgggagac aaagagatta agctcttatg taaaatttgc tgttttacat aactttaatg	2100
aatggacaaa gtcttgtgca tgggggtggg ggtgggttga gagggaaca gctccagatg	2160
gcaaacatac gcaagggtt tagtcaaaca actttttggc aaagatggta tgattttgta	2220
atggggtagg aaccaatgaa atgcgaggta agtatggtta atgatctaca gttattggtt	2280
aaagaagtat attagagcga gtctttctgc acacagatca cttttctat caacccc	2337

<210> 264

<211> 1330

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 264

aggctcagag gcacacagga gtttctgggc tcaccctgcc cccttccaac ccctcagttc	60
ccatcctcca cgagctgttt gtgtgctgcc tctgaagtcc aactgaaca aacttcagcc	120
tactcatgtc cctaaaatgg gcaaacattg caagcagcaa acagcaaaca cacagccctc	180
cctgcctgct gaccttgag ctggggcaga ggtcagagac ctctctgggc ccatgccacc	240
tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagaggtt gtcctggcgt	300
ggtttaggta gtgtgagagg gtccgggttc aaaaccactt gctgggtggg gaggcgtcag	360
taagtggcta tgccccgacc ccgaagcctg tttcccatc tgtacaatgg aaatgataaa	420
gacgcccatac tgatagggtt tttgtggcaa ataaacattt ggtttttttg ttttgttttg	480
ttttgttttt tgagatggag gtttgctctg tcgcccaggc tggagtgcag tgacacaatc	540

tcattctcacc acaaccttcc cctgcctcag cctcccaagt agctgggatt acaagcatgt	600
gccaccacac ctggctaatt ttctatTTTT agtagagacg ggTTTTctca tgttggtcag	660
cctcagcctc ccaagtaact gggattacag gcctgtgccca ccacacccgg ctaatTTTT	720
ctatTTTTga cagggacggg gtttcaccat gttggtcagg ctgggtctaga ggtaccgggg	780
ctggaagcta cctttgacat catttcctct gcgaatgcat gtataatttc tacagaacct	840
attagaaagg atcaccacagc ctctgctttt gtacaacttt cccttaaaaa actgccaatt	900
ccactgctgt ttggcccaat agtgagaact ttttcctgct gcctcttggt gcttttgcct	960
atggccccta ttctgcctgc tgaagacact cttgccagca tggacttaaa cccctccagc	1020
tctgacaatc ctctttctct tttgttttac atgaagggtc tggcagccaa agcaatcact	1080
caaagttcaa accttatcat tttttgcttt gttcctcttg gccttggttt tgtacatcag	1140
ctttgaaaat accatcccag ggtaaatgct ggggttaatt tataactaag agtgctctag	1200
ttttgcaata caggacatgc tataaaaaatg gaaagatctc taaggtaaata ataaaatttt	1260
taagtgtata atgtgttaaa ctactgattc taattgtttc tctcttttag attccaacct	1320
ttggaactga	1330

<210> 265

<211> 937

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 265

agattgtacc tgcccgtaaca taagggtcaat aggggggtgaa tcaacaggaa agtcccattg	60
gagccaagta cactgcgtca atagggactt tccattgggt tttgcccgtt acataagggtc	120
aataggggat gagtcaatgg gaaaaaccca ttggagccaa gtacactgac tcaataggga	180
ctttccattg ggTTTTgccc agtacataag gtcaataggg ggtgagtcaa caggaaagtc	240
ccattggagc caagtacatt gagtcaatag ggactttcca atgggttttg cccagtacat	300
aagggtcaatg ggaggtaagc caatgggttt ttcccattac tggcacgtat actgagtcac	360
tagggacttt ccaatgggtt ttgccagta cataagggtc atagggtga atcaacagga	420
aagtcccatt ggagccaagt aactgagtc aatagggtt ttccattggg ttttgcccag	480
tacaaaaggt caataggggg tgagtcaatg ggTTTTtccc attattggca cgtacataag	540
gtcaataggg gtgactagtc agtgggcaga gcgcacatcg cccacagtcc ccgagaagtt	600
ggggggaggg gtcggcaatt gaaccggtgc ctagagaagg tggcgcgagg taaactggga	660
aagtgatgtc gtgtactggc tccgcctttt tcccagggtt gggggagaa cgtatataag	720
tgtagtagtt gccgtgaacg ttctttttcg caacgggttt gccgccagaa cacagctgaa	780
gcttctgcct tctccctcct gtgagtttgg taagtactg actgtctatg cctgggaaag	840
ggtgggcagg agatggggca gtgcaggaaa agtggcacta tgaaccctgc agccctagac	900

aattgtacta accttcttct ctttcctctc ctgacag

937

<210> 266

<211> 367

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 266

gagcttgggc tgcaggtcga gggcactggg aggatgttga gtaagatgga aaactactga	60
tgacccttgc agagacagag tattaggaca tgtttgaaca ggggccgggc gatcagcagg	120
tagctctaga ggatccccgt ctgtctgcac atttcgtaga gcgagtgttc cgatactcta	180
atctccctag gcaaggttca tatttgtgta ggttacttat tctccttttg ttgactaagt	240
caataatcag aatcagcagg tttggagtca gcttggcagg gatcagcagc ctgggttgga	300
aggaggggggt ataaaagccc cttcaccagg agaagccgtc acacagacta ggcgcgccac	360
cgccacc	367

<210> 267

<211> 468

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 267

cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag	60
gggctaagtc cacatacggg ggaggctgct ggtgaatatt aaccaaggtc accccagtta	120
tcggaggagc aaacaggggc taagtccaca taccgtctgt ctgcacattt cgtagagcga	180
gtgttccgat actctaattc ccctaggcaa ggttcatatt tgtgtaggtt acttattctc	240
cttttgttga ctaagtcaat aatcagaatc agcaggtttg gagtcagctt ggcagggatc	300
agcagcctgg gttggaagga gggggtataa aagccccttc accaggagaa gccgtcacac	360
agatccacaa gtcctgaag aggtgaagggt ttaagggatg gttggttggt ggggtattaa	420
tgtttaatta cctggagcac ctgcctgaaa tcactttttt tcaggttg	468

<210> 268

<211> 426

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 268

agccaatgaa atacaaagat gagtctagtt aataatctac aattattggt taaagaagta	60
tattagtgtc aatttcctc cgtttgtcct agcttttctc atgcgtgtta ccgtctgtct	120
gcacatttcg tagagcgagt gttccgatac tctaattctcc ctaggcaagg ttcataattg	180
tgtaggttac ttattctcct tttgttgact aagtcaataa tcagaatcag caggtttgga	240
gtcagcttgg cagggatcag cagcctgggt tggaaggagg gggataaaaa gcccttcac	300
caggagaagc cgtcacacag atccacaagc tcctgaagag gtaagggttt aagggatggt	360
tggttggtgg ggtattaatg ttttaattacc tggagcacct gcctgaaatc actttttttc	420
aggttg	426

<210> 269

<211> 396

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 269

gaatgacctt cagcctgttc ccgtccctga tatgggcaaa cattgcaagc agcaaacagc	60
aaacacatag atgcgtgtta ccgtctgtct gcacatttcg tagagcgagt gttccgatac	120
tctaattctcc ctaggcaagg ttcataattg tgtaggttac ttattctcct tttgttgact	180
aagtcaataa tcagaatcag caggtttgga gtcagcttgg cagggatcag cagcctgggt	240
tggaaggagg gggataaaaa gcccttcac caggagaagc cgtcacacag atccacaagc	300
tcctgaagag gtaagggttt aagggatggt tggttggtgg ggtattaatg ttttaattacc	360
tggagcacct gcctgaaatc actttttttc aggttg	396

<210> 270

<211> 495

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 270

gatgctctaa tctctctaga caaggttcat atttgatgg gttacttatt ctctctttgt	60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc	120
tagctcagga gaagtgaagta taaaagcccc aggctgggag cagccatcaa tgcgtgttac	180
cgtctgtctg cacatttcgt agagcgagtg ttccgatact ctaatctccc taggcaaggt	240
tcataattgt gtaggttact tattctcctt ttgttgacta agtcaataat cagaatcagc	300
aggtttggag tcagcttggc agggatcagc agcctgggtt ggaaggaggg ggtataaaag	360
ccccttcacc aggagaagcc gtcacacaga tccacaagct cctgaagagg taagggttta	420
agggatggtt ggttggtggg gtattaatgt ttaattacct ggagcacctg cctgaaatca	480

<210> 271

<211> 640

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 271

cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag 60
gggctaagtc cacatgcgtg ttagggctgg aagctacctt tgacatcatt tcctctgcga 120
atgcatgtat aatttctaca gaacctatta gaaaggatca cccagcctct gcttttgtac 180
aactttccct taaaaaactg ccaattccac tgctgtttgg cccaatagtg agaacttttt 240
cctgctgcct ctgggtgctt ttgcctatgg ccctatttct gcctgctgaa gacactcttg 300
ccagcatgga cttaaaccct tccagctctg acaatcctct ttctcttttg ttttacatga 360
agggtctggc agccaaagca atcactcaaa gttcaaacct tatcattttt tgctttgttc 420
ctcttgccct tggttttgta catcagcttt gaaaatacca tcccagggtt aatgctgggg 480
ttaatttata actaagagtg ctctagtttt gcaatacagg acatgctata aaaatggaaa 540
gatctcctga agaggtaagg gttaagga tggttggttg gtgggggtatt aatgtttaat 600
tacctggagc acctgcctga aatcactttt tttcaggttg 640

<210> 272

<211> 667

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 272

agccaatgaa atacaagat gagtctagtt aataatctac aattattggt taaagaagta 60
tattagtgtt aatttccctc cgtttgcct agcttttctc atgcgtgtta gggctggaag 120
ctacctttga catcatttcc tctgcgaatg catgtataat ttctacagaa cctattagaa 180
aggatcacc agcctctgct tttgtacaac tttcccttaa aaaactgcca attccactgc 240
tgtttgccc aatagtgaga actttttcct gctgcctctt ggtgcttttg cctatggccc 300
ctattctgcc tgctgaagac actcttgcca gcatggactt aaaccctcc agctctgaca 360
atcctctttc tcttttgttt tacatgaagg gtctggcagc caaagcaatc actcaaagtt 420
caaaccttat cattttttgc tttgttctc ttggccttgg tttgtacat cagctttgaa 480
aataccatcc cagggttaat gctggggtta atttataact aagagtgtc tagttttgca 540
atacaggaca tgctataaaa atggaaagat ctctgaaga ggtaagggtt taagggatgg 600

ttggttggtg gggattaat gtttaattac ctggagcacc tgcctgaaat cacttttttt 660
caggttg 667

<210> 273
<211> 637
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 273
gaatgacctt cagcctgttc ccgtccctga tatgggcaaa cattgcaagc agcaaacagc 60
aaacacatag atgcgtgtta gggctggaag ctacctttga catcatttcc tctgcgaatg 120
catgtataat ttctacagaa cctattagaa aggatcacc agcctctgct tttgtacaac 180
tttcccttaa aaaactgcc aattccactgc tgtttgcccc aatagtgaga actttttcct 240
gctgcctctt ggtgcttttg cctatggccc ctattctgcc tgctgaagac actcttgcca 300
gcatggactt aaacccctcc agctctgaca atcctctttc tcttttgttt tacatgaagg 360
gtctggcagc caaagcaatc actcaaagtt caaaccttat cttttttgc tttgttcctc 420
ttggccttgg tttgtacat cagctttgaa aataccatcc cagggttaat gctggggtta 480
atttataact aagagtgtc tagttttgca atacaggaca tgctataaaa atggaaagat 540
ctcctgaaga ggtaagggtt taagggatgg ttggttggtg gggattaat gtttaattac 600
ctggagcacc tgcctgaaat cacttttttt caggttg 637

<210> 274
<211> 736
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 274
gatgctctaa tctctctaga caaggttcat atttgtatgg gttacttatt ctctctttgt 60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc 120
tagctcagga gaagttagta taaaagcccc aggctgggag cagccatcaa tgcgtgtag 180
ggctggaagc tacctttgac atcatttcct ctgcgaatgc atgtataatt tctacagaac 240
ctattagaaa ggatcaccca gcctctgctt ttgtacaact ttcccttaaa aaactgccaa 300
ttccactgct gtttgccca atagtgagaa ctttttcctg ctgcctcttg gtgcttttgc 360
ctatggcccc tattctgcct gctgaagaca ctcttgccag catggactta aaccctcca 420
gctctgacaa tcctctttct cttttgtttt acatgaaggg tctggcagcc aaagcaatca 480
ctcaaagttc aaaccttatc attttttgct ttgttcctct tggccttggt tttgtacatc 540
agctttgaaa ataccatccc aggggttaat ctgggggttaa ttataacta agagtgtctc 600

agttttgcaa tacaggacat gctataaaaa tggaaagatc tcctgaagag gtaagggttt	660
aagggatggt tggttggtgg ggtattaatg tttaattacc tggagcacct gcctgaaatc	720
actttttttc aggttg	736

<210> 275
 <211> 515
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 275 cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag	60
gggctaagtc cacatgcgtg ttaggcatgc ttccatgccca aggcccacac tgaaatgctc	120
aaatgggaga caaagagatt aagctcttat gtaaaatttg ctgtttttaca taactttaat	180
gaatggacaa agtcttgtgc atgggggtgg ggggtggggtt agaggggaac agctccagat	240
ggcaaacata cgcaagggat ttagtcaaac aacttttttg caaagatggt atgattttgt	300
aatggggtag gaaccaatga aatgcgaggt aagtatggtt aatgatctac agttattggt	360
taaagaagta tattagagcg agtctttctg cacacagatc acctttccta tcaacccct	420
cctgaagagg taagggttta agggatggtt ggttggtggg gtattaatgt ttaattacct	480
ggagcacctg cctgaaatca ctttttttca ggttg	515

<210> 276
 <211> 542
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 276 agccaatgaa atacaagat gagtctagtt aataatctac aattattggt taaagaagta	60
tattagtgtc aatttccctc cgtttgcct agcttttctc atgcgtgtta ggcatgcttc	120
catgccaagg cccacactga aatgctcaaa tgggagacaa agagattaag ctcttatgta	180
aaatttgctg ttttacataa cttaatgaa tggacaaagt cttgtgcatg ggggtggggg	240
tggggttaga ggggaacagc tccagatggc aaacatacgc aagggttta gtcaaacaac	300
tttttgcaa agatggtatg attttgtaat ggggtaggaa ccaatgaaat gcgaggtaag	360
tatggttaat gatctacagt tattggttaa agaagtatat tagagcgagt ctttctgcac	420
acagatcacc tttcctatca acccctcct gaagaggtaa gggtttaagg gatggttggt	480
tgggtgggta ttaatgttta attacctgga gcacctgcct gaaatcactt tttttcaggt	540
tg	542

<210> 277
<211> 512
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 277
gaatgacctt cagcctgttc ccgtccctga tatgggcaaa cattgcaagc agcaaacagc 60
aaacacatag atgcgtgtta ggcatgcttc catgccaagg cccacactga aatgctcaaa 120
tgggagacaa agagattaag ctcttatgta aaatttgctg ttttacataa ctttaatgaa 180
tggacaaagt cttgtgcatg ggggtggggg tggggttaga ggggaacagc tccagatggc 240
aaacatacgc aagggtatga gtcaaacaac tttttggcaa agatgggtatg attttgtaat 300
ggggtaggaa ccaatgaaat gcgaggttaag tatgggttaat gatctacagt tattgggttaa 360
agaagtatat tagagcgagt ctttctgcac acagatcacc tttcctatca acccctcct 420
gaagaggtaa ggggttaagg gatggttggt tggtagggta ttaatgttta attacctgga 480
gcacctgcct gaaatcactt tttttcaggt tg 512

<210> 278
<211> 611
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 278
gatgctctaa tctctctaga caaggttcat atttgtatgg gttacttatt ctctctttgt 60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc 120
tagctcagga gaagtgaagta taaaagcccc aggctgggag cagccatcaa tgcgtgttag 180
gcatgcttcc atgccaaggc ccacactgaa atgctcaaat gggagacaaa gagattaagc 240
tcttatgtaa aatttgctgt tttacataac tttaatgaat ggacaaagtc ttgtgcatgg 300
gggtgggggt ggggttagag gggaacagct ccagatggca aacatacgca agggatttag 360
tcaaacaact ttttgcaaaa gatggtatga ttttgtaatg gggtaggaac caatgaaatg 420
cgaggtaagt atggttaatg atctacagtt attggttaaa gaagtatatt agagcgagtc 480
tttctgcaca cagatcacct ttcctatcaa cccctcctg aagaggtaag ggtttaaggg 540
atggttggtt ggtgggggat taatgtttta ttacctggag cacctgcctg aaatcacttt 600
ttttcaggtt g 611

<210> 279
<211> 355
<212> DNA
<213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 279
 cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag 60
 gggctaagtc cacatgcgtg ttaaacagtt ccagatggta aatatacaca agggatttag 120
 tcaaacaatt ttttggcaag aatattatga attttghta cggttggcag ccaatgaaat 180
 acaaagatga gtctagttaa taatctacaa ttattggta aagaagtata ttagtgctaa 240
 tttccctccg tttgtcctct cctgaagagg taagggttta agggatgggt ggttggtggg 300
 gtattaatgt ttaattacct ggagcacctg cctgaaatca ctttttttca ggttg 355

<210> 280
 <211> 382
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 280
 agccaatgaa atacaaagat gagtctagtt aataatctac aattattggt taaagaagta 60
 tattagtgtt aatttccttc cgtttgtcct agcttttctc atgcgtgtta aacagttcca 120
 gatggtaa atacacaagg gatttagtca aacaattttt tggcaagaat attatgaatt 180
 ttgtaatcgg ttggcagcca atgaaatata aagatgagtc tagttaataa tctacaatta 240
 ttggttaaag aagtatatta gtgctaattt ccctccgttt gtcctctcct gaagaggtaa 300
 gggtttaagg gatggttggt tgggtgggta ttaatgttta attacctgga gcacctgcct 360
 gaaatcactt tttttcaggt tg 382

<210> 281
 <211> 352
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 281
 gaatgacctt cagcctgttc ccgtccctga tatgggcaaa cattgcaagc agcaaacagc 60
 aaacacatag atgcgtgtta aacagttcca gatggtaa atacacaagg gatttagtca 120
 aacaattttt tggcaagaat attatgaatt ttgtaatcgg ttggcagcca atgaaatata 180
 aagatgagtc tagttaataa tctacaatta ttggttaaag aagtatatta gtgctaattt 240
 ccctccgttt gtcctctcct gaagaggtaa gggtttaagg gatggttggt tgggtgggta 300
 ttaatgttta attacctgga gcacctgcct gaaatcactt tttttcaggt tg 352

<210> 282
<211> 451
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 282
gatgctctaa tctctctaga caaggttcat atttgtatgg gttacttatt ctctctttgt 60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc 120
tagctcagga gaagtgahta taaaagcccc aggctgggag cagccatcaa tgcgtgttaa 180
acagttccag atggtaaata tacacaaggg atttagtcaa acaatttttt ggcaagaata 240
ttatgaattt tgtaatcggt tggcagccaa tgaaatacaa agatgagtct agttaataat 300
ctacaattat tggttaaaga agtatattag tgctaatttc cctccgtttg tcctctcctg 360
aagaggttaag ggtttaaggg atggttggtt ggtggggtat taatgtttta ttacctggag 420
cacctgcctg aaatcacttt ttttcaggtt g 451

<210> 283
<211> 430
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 283
cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag 60
gggctaagtc cacatgcgtg ttaaatagact cttttcggtg agtgcagtgg aagctgtaca 120
ctgcccaggc aaagcgtccg ggcagcgtag gcgggcgact cagatcccag ccagtggact 180
tagcccctgt ttgctcctcc gataactggg gtgaccttgg ttaatattca ccagcagcct 240
ccccgttgc ccctctggat cactgctta aatacggacg aggacagggc cctgtctcct 300
cagcttcagg caccaccact gacctgggac agtctcctga agaggtaagg gttaaggga 360
tggttggttg gtggggtatt aatgtttaat tacctggagc acctgcctga aatcactttt 420
tttcaggttg 430

<210> 284
<211> 457
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 284
agccaatgaa atacaaagat gagtctagtt aataatctac aattattggt taaagaagta 60

tattagtgtc aatttcctc cgtttgtcct agcttttctc atgcgtgtta aatgactcct	120
ttcggtaagt gcagtggaag ctgtacactg cccaggcaaa gcgtccgggc agcgtaggcg	180
ggcgactcag atcccagcca gtggacttag cccctgtttg ctctccgat aactgggggtg	240
accttggtta atattcacca gcagcctccc ccgttgcccc tctggatcca ctgcttaaat	300
acggacgagg acagggccct gtctcctcag cttcaggcac caccactgac ctgggacagt	360
ctcctgaaga ggtaagggtt taagggatgg ttggttggtg gggatattaat gtttaattac	420
ctggagcacc tgcctgaaat cacttttttt caggttg	457

<210> 285

<211> 427

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 285

gaatgacctt cagcctgttc ccgtccctga tatgggcaaa cattgcaagc agcaaacagc	60
aaacacatag atgcgtgtta aatgactcct ttcggtaagt gcagtggaag ctgtacactg	120
cccaggcaaa gcgtccgggc agcgtaggcg ggcgactcag atcccagcca gtggacttag	180
cccctgtttg ctctccgat aactgggggtg accttggtta atattcacca gcagcctccc	240
ccgttgcccc tctggatcca ctgcttaaat acggacgagg acagggccct gtctcctcag	300
cttcaggcac caccactgac ctgggacagt ctcctgaaga ggtaagggtt taagggatgg	360
ttggttggtg gggatattaat gtttaattac ctggagcacc tgcctgaaat cacttttttt	420
caggttg	427

<210> 286

<211> 526

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 286

gatgctctaa tctctctaga caaggttcat atttgtatgg gttacttatt ctctctttgt	60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc	120
tagctcagga gaagtgagta taaaagcccc aggctgggag cagccatcaa tgcgtgttaa	180
atgactcctt tcggtaagtg cagtggaagc tgtacactgc ccaggcaaag cgtccgggca	240
gcgtaggcgg gcgactcaga tcccagccag tggacttagc ccctgtttgc tcctccgata	300
actgggggtga ctttggttaa tattcaccag cagcctcccc cgttgcccct ctggatccac	360
tgcttaaata cggacgagga cagggccctg tctcctcagc ttcaggcacc accactgacc	420

tgggacagtc tcctgaagag gtaagggttt aagggatggt tggttggtgg ggtattaatg 480

tttaattacc tggagcacct gcctgaaatc actttttttc aggttg 526

<210> 287

<211> 435

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 287

cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag 60

gggctaagtc cacatgcgtg ttaaattgact cttttcggtg agtgcagtgg aagctgtaca 120

ctgcccaggc aaagcgtccg ggtagcgtg gcgggtagc cagatcccag ccagtggact 180

tagcccctgt ttgctcctcc gataactggg gtgaccttgg ttaatatcca ccagcagcct 240

ccccgttgc ccctctggat ccactgctta aatacggacg aggacagggc cctgtctcct 300

cagcttcagg caccaccact gacctgggac agtgaatccg gactctaagg taaatataaa 360

atttttaagt gtataatgtg ttaaactact gattctaatt gtttctctct ttagattcc 420

aacctttgga actga 435

<210> 288

<211> 462

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 288

agccaatgaa atacaaagat gagtctagtt aataatctac aattattggt taaagaagta 60

tattagtgtc aatttccttc cgtttgcct agcttttctc atgcgtgtta aatgactcct 120

ttcggtaagt gcagtggaag ctgtacactg cccaggcaaa gcgtccgggc agcgtaggcg 180

ggcgactcag atcccagcca gtggacttag cccctgtttg ctctccgat aactgggggtg 240

accttggtta atattcacca gcagcctccc ccgttgcccc tctggatcca ctgcttaaat 300

acggacgagg acagggccct gtctcctcag cttcaggcac caccactgac ctgggacagt 360

gaatccggac tctaaggtaa atataaaatt tttaagtgtg taatgtgtta aactactgat 420

tctaattggt tctctctttt agattccaac ctttgaact ga 462

<210> 289

<211> 448

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic

polynucleotide"

<400> 289

gcggccgcga atgaccttca gcctgttccc gtccctgata tgggcaaaca ttgcaagcag	60
caaacagcaa acacatagat gcgtgttaaa tgactccttt cggttaagtgc agtggaagct	120
gtacactgcc caggcaaagc gtccggggcag cgtaggcggg cgactcagat cccagccagt	180
ggacttagcc cctgtttgct cctccgataa ctgggggtgac cttgggttaat attcaccagc	240
agcctcccc gttgccctc tggatccact gcttaaatac ggacgaggac agggccctgt	300
ctcctcagct tcaggcacca ccactgacct gggacagtga atccggactc taaggtaaata	360
ataaaatatt taagtgtata atgtgttaaa ctactgattc taattgtttc tctcttttag	420
attccaacct ttggaactga gtttaaac	448

<210> 290

<211> 531

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 290

gatgctctaa tctctctaga caaggttcat atttgtatgg gttacttatt ctctctttgt	60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc	120
tagctcagga gaagttagta taaaagcccc aggctgggag cagccatcaa tgcgtgttaa	180
atgactcctt tcggtaagtg cagtgggaagc tgtacactgc ccaggcaaag cgtccgggca	240
gcgtaggcgg gcgactcaga tcccagccag tggacttagc ccctgtttgc tcctccgata	300
actgggggtga cttgggttaa tattcaccag cagcctcccc cgttgcccct ctggatccac	360
tgcttaaata cggacgagga cagggccctg tctcctcagc ttcaggcacc accactgacc	420
tgggacagtg aatccggact ctaaggtaaa tataaaatatt ttaagtgtat aatgtgttaa	480
actactgatt ctaattgttt ctctctttta gattccaacc tttggaactg a	531

<210> 291

<211> 636

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 291

cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag	60
gggctaagtc cacatgcgtg ttagggctgg aagctacctt tgacatcatt tcctctgcga	120
atgcatgtat aatttctaca gaacctatta gaaaggatca cccagcctct gcttttgtac	180
aactttccct taaaaaactg ccaattccac tgctgtttgg cccaatagtg agaacttttt	240

cctgctgcct cttggtgctt ttgcctatgg cccctattct gcctgctgaa gacactcttg	300
ccagcatgga cttaaacccc tccagctctg acaatcctct ttctcttttg ttttacatga	360
agggtctggc agccaaagca atcactcaaa gttcaaacct tatcattttt tgctttgttc	420
ctcttggcct tggttttgta catcagcttt gaaaatacca tcccagggtt aatgctgggg	480
ttaatttata actaagagt ctctagtttt gcaatacagg acatgctata aaaatggaaa	540
gatctctaag gtaaataata aatttttaag tgtataatgt gttaaactac tgattctaata	600
tgtttctctc ttttagattc caacctttgg aactga	636

<210> 292

<211> 663

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 292

agccaatgaa atacaaagat gagtctagtt aataatctac aattattggt taaagaagta	60
tattagtgtc aatttcctc cgtttgtcct agcttttctc atgcgtgtta gggctggaag	120
ctacctttga catcatttcc tctgcgaatg catgtataat ttctacagaa cctattagaa	180
aggatcacc agcctctgct tttgtacaac tttcccttaa aaaactgcca attccactgc	240
tgtttggccc aatagtgaga actttttcct gctgcctctt ggtgcttttg cctatggccc	300
ctattctgcc tgctgaagac actcttgcca gcatggactt aaaccctcc agctctgaca	360
atcctctttc tcttttgtt tacatgaagg gtctggcagc caaagcaatc actcaaagtt	420
caaaccttat cattttttgc tttgttcctc ttggccttgg ttttgtacat cagctttgaa	480
aataccatcc cagggttaat gctgggggtta atttataact aagagtgtc tagttttgca	540
atacaggaca tgctataaaa atggaaagat ctctaaggta aatataaaat ttttaagtgt	600
ataatgtgtt aaactactga ttctaattgt ttctctctt tagattccaa cctttggaac	660
tga	663

<210> 293

<211> 633

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 293

gaatgacctt cagcctgttc ccgtccctga tatgggcaaa cattgcaagc agcaaacagc	60
aaacacatag atgcgtgtta gggctggaag ctacctttga catcatttcc tctgcgaatg	120
catgtataat ttctacagaa cctattagaa aggatcacc agcctctgct tttgtacaac	180
tttcccttaa aaaactgcca attccactgc tgtttggccc aatagtgaga actttttcct	240

gctgcctctt ggtgcttttg cctatggccc ctattctgcc tgctgaagac actcttgcca	300
gcatggactt aaacccctcc agctctgaca atcctctttc tcttttggtt tacatgaagg	360
gtctggcagc caaagcaatc actcaaagtt caaaccttat catTTTTTgc tttgttcctc	420
ttggccttgg ttttgtacat cagctttgaa aataccatcc cagggttaat gctggggtta	480
atttataact aagagtgctc tagttttgca atacaggaca tgctataaaa atggaaagat	540
ctctaaggta aatataaaat ttttaagtgt ataatgtgtt aaactactga ttctaattgt	600
ttctctcttt tagattccaa cctttggaac tga	633

<210> 294

<211> 732

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 294

gatgctctaa tctctctaga caaggttcat atttgtatgg gttacttatt ctctctttgt	60
tgactaagtc aataatcaga atcagcaggt ttgcagtcag attggcaggg ataagcagcc	120
tagctcagga gaagtgahta taaaagcccc aggctgggag cagccatcaa tgcgtgtag	180
ggctggaagc tacctttgac atcatttcct ctgcgaatgc atgtataatt tctacagaac	240
ctattagaaa ggatcaccca gcctctgctt ttgtacaact ttcccttaaa aaactgcaa	300
ttccactgct gtttgccca atagttagaa ctttttcctg ctgcctcttg gtgcttttgc	360
ctatggcccc tattctgcct gctgaagaca ctcttgccag catggactta aacccctcca	420
gctctgacaa tcctctttct cttttgtttt acatgaaggg tctggcagcc aaagcaatca	480
ctcaaagttc aaaccttatc attttttgct ttgttcctct tggccttggt tttgtacatc	540
agctttgaaa ataccatccc agggttaatg ctggggttaa ttataacta agagtgtctt	600
agttttgcaa tacaggacat gctataaaaa tggaaagatc tctaaggtaa atataaaatt	660
tttaagtgtg taatgtgtta aactactgat tctaattgtt tctctctttt agattccaac	720
ccttggaact ga	732

<210> 295

<211> 762

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 295

aggttaattt ttaaaaagca gtcaaaagtc caagtggccc ttggcagcat ttactctctc	60
tgtttgctct ggtaataat ctcaggagca caaacattcc agatccaggt taatttttaa	120

aaagcagtca aaagtccaag tggcccttgg cagcatttac tctctctggt tgctctgggt	180
aataatctca ggagcacaaa cattccagat ccggcgcgcc agggctggaa gctacctttg	240
acatcatttc ctctgcgaat gcatgtataa tttctacaga acctattaga aaggatcacc	300
cagcctctgc ttttgtacaa ctttccctta aaaaactgcc aattccactg ctgtttggcc	360
caatagttag aactttttcc tgctgcctct tgggtgcttt gcctatggcc cctattctgc	420
ctgctgaaga cactcttgcc agcatggact taaaccctc cagctctgac aatcctcttt	480
ctcttttggt ttacatgaag ggtctggcag ccaaagcaat cactcaaagt tcaaacctta	540
tcattttttg ctttgttcct cttggccttg gttttgtaca tcagctttga aaataccatc	600
ccagggttaa tgctgggggt aatttataac taagagtgtc ctagttttgc aatacaggac	660
atgctataac tctaaggtaa atataaaatt tttaagtgt taatgtgtta aactactgat	720
tctaattggt tctctctttt agattccaac ctttggaact ga	762

<210> 296

<211> 766

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 296

aggttaattt ttaaaaagca gtcaaaagtc caagtggccc ttggcagcat ttactctctc	60
tgtttgctct ggtaataat ctccaggagca caaacattcc agatccaggt taatttttaa	120
aaagcagtca aaagtccaag tggcccttgg cagcatttac tctctctggt tgctctgggt	180
aataatctca ggagcacaaa cattccagat ccggcgcgcc agggctggaa gctacctttg	240
acatcatttc ctctgcgaat gcatgtataa tttctacaga acctattaga aaggatcacc	300
cagcctctgc ttttgtacaa ctttccctta aaaaactgcc aattccactg ctgtttggcc	360
caatagttag aactttttcc tgctgcctct tgggtgcttt gcctatggcc cctattctgc	420
ctgctgaaga cactcttgcc agcatggact taaaccctc cagctctgac aatcctcttt	480
ctcttttggt ttacatgaag ggtctggcag ccaaagcaat cactcaaagt tcaaacctta	540
tcattttttg ctttgttcct cttggccttg gttttgtaca tcagctttga aaataccatc	600
ccagggttaa tgctgggggt aatttataac taagagtgtc ctagttttgc aatacaggac	660
atgctataac tcctgaagag gtaagggttt aagggatggt tggttggtgg ggtattaatg	720
tttaattacc tggagcacct gcctgaaatc actttttttc aggttg	766

<210> 297

<400> 297

000

<210> 298

<400> 298
000

<210> 299

<400> 299
000

<210> 300
<211> 518
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 300
tcaatattgg ccattagcca tattattcat tggttatata gcataaatca atattggcta 60
ttggccattg catacgttgt atctatatca taatatgtac atttatattg gctcatgtcc 120
aatatgaccg ccatgttggc attgattatt gactagttat taatagtaat caattacggg 180
gtcattagtt catagcccat atatggagtt ccgcgttaca taacttacgg taaatggccc 240
gcctggctga ccgccaacg acccccgccc attgacgtca ataatgacgt atgttcccat 300
agtaacgcca atagggactt tccattgacg tcaatgggtg gagtatttac ggtaaactgc 360
ccacttgga gtacatcaag tgtatcatat gccaaagtcc cccctattg acgtcaatga 420
cggtaaatgg ccgcctggc attatgccca gtacatgacc ttacgggact ttcctacttg 480
gcagtacatc tacgtattag tcatcgctat taccatgg 518

<210> 301
<211> 777
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 301
aggctcagag gcacacagga gtttctgggc tcaccctgcc cccttccaac ccctcagttc 60
ccatcctcca gcagctgttt gtgtgctgcc tctgaagtcc aactgaaca aacttcagcc 120
tactcatgtc ctaaaatgg gcaaacattg caagcagcaa acagcaaaca cacagccctc 180
cctgcctgct gaccttgag ctggggcaga ggtcagagac ctctctgggc ccatgccacc 240
tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagagggt gtcctggcgt 300
ggtttaggta gtgtgagagg gtccgggttc aaaaccactt gctgggtggg gagtgcgcag 360
taagtggcta tgccccgacc ccgaagcctg tttccccatc tgtacaatgg aaatgataaa 420
gacgcccac tcataggggt tttgtggcaa ataaacattt ggtttttttg ttttgttttg 480
ttttgttttt tgagatggag gtttgctctg tcgcccaggc tggagtgcag tgacacaatc 540

tcattctcacc acaaccttcc cctgcctcag cctcccaagt agctgggatt acaagcatgt	600
gccaccacac ctggctaatt ttctatTTTT agtagagacg ggTTTctcca tGttggTcag	660
cctcagcctc ccaagtaact gggattacag gcctgtgccca ccacaccgga ctaatTTTT	720
ctatTTTTga cagggacggg gtttcacCAT gttggTcagg ctggTctaga ggtaccg	777

<210> 302
 <211> 427
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 302 gagtcaatgg gaaaaaccca ttggagccaa gtacactgac tcaataggga ctttccattg	60
ggTTTTgcc agtacataag gtcaataggg ggtgagtcaa caggaaagtc ccattggagc	120
caagtacatt gagtcaatag ggactttcca atgggTTTTg ccagTacat aaggTcaatg	180
ggaggTaaGc caatgggTtt ttcccattac tgacatgtat actgagTcat tagggacttt	240
ccaatgggTt ttgccagta cataaggTca ataggggTga atcaacagga aagTcccatt	300
ggagccaagt aactgagTc aatagggact ttccattggg ttttgccag taaaaaggt	360
caataggggg tgagtcaatg ggtTTTTccc attattggca catacataag gtcaataggg	420
gtgacta	427

<210> 303
 <211> 83
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 303 cgggggaggc tgctggTgaa tattaaccaa ggtcaccCCA gttatcggag gagcaaacag	60
gggctaagTc cacacgcgtg gta	83

<210> 304
 <211> 777
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 304 aggTcagag gcacacagga gtttctgggc tcaccTgcc ccTtccaac ccTcagTtc	60
ccatcctcca gcagTgttt gTgtgctgcc tctgaagTcc aactgaaca aactTcagcc	120
tactcatgtc ctaaaatgg gaaacattg caagcagcaa acagcaaaca cacagccctc	180

cctgcctgct gaccttggag ctggggcaga ggtcagagac ctctctgggc ccatgccacc	240
tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagagggt gtcctggcgt	300
ggtttaggta gtgtgagagg gtccgggttc aaaaccactt gctgggtggg gagtgcgcag	360
taagtggcta tgccccgacc ccgaagcctg tttccccatc tgtacaatgg aaatgataaa	420
gacgccccatc tgatagggtt tttgtggcaa ataaacattt ggtttttttg ttttgttttg	480
ttttgttttt tgagatggag gtttgctctg tcgcccaggc tggagtgcag tgacacaatc	540
tcatctcacc acaaccttcc cctgcctcag cctcccaagt agctgggatt acaagcatgt	600
gccaccacac ctggctaatt ttctatTTTT agtagagacg ggtttctcca tgttggtcag	660
cctcagcctc ccaagtaact gggattacag gcctgtgcc cccaccccgg ctaatttttt	720
ctatTTTTga cagggacggg gtttcacat gttggtcagg ctggtctaga ggtactg	777

<210> 305
 <211> 66
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 305	
gtttgctgct tgcaatgttt gccattttta ggggtggacac aggacgctgt ggtttctgag	60
ccaggg	66

<210> 306
 <211> 212
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 306	
ggaggggtgg agtcgtgacc ctaaaatgg gcaaacattg caagcagcaa acagcaaaca	60
cacagccctc cctgcctgct gaccttggag ctggggcaga ggtcagagac ctctctgggc	120
ccatgccacc tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagagggt	180
gtcctggcgt ggtttaggta gtgtgagagg gg	212

<210> 307
 <211> 330
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 307

aggctcagag gcacacagga gtttctgggc tcaccctgcc cccttccaac ccctcagttc	60
ccatcctcca gcagctgttt gtgtgctgcc tctgaagtcc aactgaaca aacttcagcc	120
tactcatgtc cctaaaatgg gcaaacattg caagcagcaa acagcaaaca cacagccctc	180
cctgcctgct gaccttggag ctggggcaga ggtagagac ctctctgggc ccatgccacc	240
tccaacatcc actcgacccc ttggaatttc ggtggagagg agcagaggtt gtcctggcgt	300
ggttaggta gtgtgagagg ggtaccggg	330

<210> 308
 <211> 194
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 308 ccctaaaatg ggcaaacatt gcaagcagca aacagcaaac acacagccct ccctgcctgc	60
tgaccttga gctggggcag aggtcagaga cctctctggg cccatgccac ctccaacatc	120
cactcgaccc cttggaattt ttcggtggag aggagcagag gttgtcctgg cgtggttag	180
gtagtgtgag aggg	194

<210> 309
 <211> 240
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 309 gggcctgaaa taacctctga aagaggaact tggttaggta cttctgagg ctgaaagaac	60
cagctgtgga atgtgtgtca gtaggggtgt ggaaagtccc caggctccc agcaggcaga	120
agtatgcaaa gcatgcatct caattagtca gcaaccaggt gtggaaagtc cccaggctcc	180
ccagcaggca gaagtatgca aagcatgcat ctcaattagt cagcaaccat agtcccacta	240

<210> 310
 <211> 73
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 310 cgggggaggc tgctggtgaa tattaaccaa ggtcacccca gttatcggag gagcaaacag	60
gggctaagtc cac	73

<210> 311
<211> 100
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 311
aggttaattt ttaaaaagca gtcaaaagtc caagtggccc ttggcagcat ttactctctc 60
tgtttgctct ggtaataat ctcaggagca caaacattcc 100

<210> 312
<211> 296
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 312
gttacataac ttatggtaaa tggcctgcct ggctgactgc ccaatgaccc ctgcccattg 60
atgtcaataa tgatgtatgt tcccatgtaa tgccaatagg gactttccat tgatgtcaat 120
gggtggagta tttatggtaa ctgcccactt ggcaagtacat caagtgtatc atatgccaag 180
tatgccccct attgatgtca atgatggtaa atggcctgcc tggcattatg cccagtacat 240
gaccttatgg gactttccta cttggcagta catctatgta ttagtcattg ctatta 296

<210> 313
<211> 235
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 313
ggcctgaaat aacctctgaa agaggaactt ggtaggtac cttctgaggc ggaaagaacc 60
agctgtggaa tgtgtgtcag ttaggggtgtg gaaagtcccc aggctcccca gcaggcagaa 120
gtatgcaaag catgcatctc aattagtcag caaccaggtg tggaaagtcc ccaggctccc 180
cagcaggcag aagtatgcaa agcatgcatc tcaattagtc agcaaccata gtccc 235

<210> 314

<400> 314
000

<210> 315
<211> 1127
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 315
ggagtcgctg cgacgtgcc ttcgccccgt gccccgctcc gccgccgcct cgcgccgccc 60
gccccggctc tgactgaccg cgttactccc acaggtgagc gggcgggacg gcccttctcc 120
tccgggctgt aattagcgct tggtttaatg acggcttggt tcttttctgt ggctgcgtga 180
aagccttgag gggctccggg agggcccttt gtgcgggggg gagcggctcg gggggtgcgt 240
gcgtgtgtgt gtgcgtgggg agcgcgcgt gcggcccgcg ctgcccggcg gctgtgagcg 300
ctgcgggcgc ggcgcggggc tttgtgcgt ccgcagtgtg cgcgagggga gcgcggccgg 360
gggcggtgcc ccgcggtgcg gggggggctg cgaggggaac aaaggctgcg tgcggggtgt 420
gtgcgtgggg gggtagcag ggggtgtggg cgcggcggtc gggctgtaac cccccctgc 480
acccccctcc ccgagttgct gagcacggcc cggcttcggg tgcggggctc cgtacggggc 540
gtggcgcggg gctcgccgtg ccgggcgggg ggtggcggca ggtgggggtg ccgggcgggg 600
cggggccgcc tcgggccggg gagggctcgg gggaggggcg cggcggcccc cggagcgccg 660
gcggctgtcg aggcgcggcg agccgcagcc attgcctttt atggtaatcg tgcgagaggg 720
cgcagggact tcctttgtcc caaatctgtg cggagccgaa atctgggagg cgcgccgca 780
ccccctctag cgggcgcggg gcgaagcggg gcggcgccgg caggaaggaa atgggcgggg 840
agggccttcg tgcgtcgccg gcgcgccgtc cccttctccc tctccagcct cggggctgtc 900
cgcgggggga cggctgcctt cgggggggac ggggcagggc ggggttcggc ttctggcgtg 960
tgaccggcgg ctctagagcc tctgctaacc atgttttagc cttcttcttt ttcttacagc 1020
tcctgggcaa cgtgctgggt attgtgctgt ctcattcatt gtcgacagaa ttctcgaag 1080
atccgaaggg gttcaagctt ggcattccgg tactgttggg aaagcca 1127

<210> 316
<211> 93
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 316
ctctaaggta aatataaaat ttttaagtgt ataatgtgtt aaactactga ttctaattgt 60
ttctctcttt tagattccaa cctttggaac tga 93

<210> 317
<211> 54
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic

oligonucleotide"

<400> 317

gccctgtctc ctcagcttca ggcaccacca ctgacctggg acagtgaatc cgga 54

<210> 318

<211> 173

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 318

ctgccttctc cctcctgtga gtttggtgaag tctactgactg tctatgcctg ggaaagggtg 60

ggcaggagat ggggcagtgc aggaaaagtg gcactatgaa ccctgcagcc ctagacaatt 120

gtactaacct tcttctcttt cctctcctga caggttgggtg tacagtagct tcc 173

<210> 319

<211> 91

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 319

aagaggtaag ggtttaaggg atggttggtt ggtggggtat taatgtttaa ttacctggag 60

cacctgcctg aaatcacttt ttttcaggtt g 91

<210> 320

<211> 54

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 320

gccctgtctc ctcagcttca ggcaccacca ctgacctggg acagtgaata atta 54

<210> 321

<211> 147

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 321

gccctgtctc ctcagcttca ggcaccacca ctgacctggg acagtgaatc cggactctaa 60

ggtaaataata aaatttttaa gtgtataatg gtgtaaacta ctgattctaa ttgtttctct 120

cttttagatt ccaacctttg gaactga 147

<210> 322

<211> 147

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 322

gccctgtctc ctcagcttca ggcaccacca ctgacctggg acagtgaata attactctaa 60

ggtaaataata aaatttttaa gtgtataatg tgttaaacta ctgattctaa ttgtttctct 120

cttttagatt ccaacctttg gaactga 147

<210> 323

<211> 48

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 323

tcctcagctt caggcaccac cactgacctg ggacagtga tgcacc 48

<210> 324

<211> 128

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 324

gctagcaggt aagtgccgtg tgtgggtccc gcgggcctgg cctctttacg ggttatggcc 60

cttgctgcc ttgaattact gacactgaca tccacttttt ctttttctcc acaggtttaa 120

acgccacc 128

<210> 325

<211> 98

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 325

aagaggtaag ggtttaagtt atcgtagtt cgtgcacat taatgtttaa ttacctggag 60

cacctgcctg aaatcatttt tttttcaggt tggctagt 98

<210> 326
<211> 172
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 326
gcttagtgct gagcacatcc agtgggtaaa gttccttaaa atgctctgca aagaaattgg 60
gacttttcat taaatcagaa attttacttt tttcccctcc tgggagctaa agatatttta 120
gagaagaatt aaccttttgc ttctccagtt gaacatttgt agcaataagt ca 172

<210> 327
<211> 160
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 327
gccctgtctc ctcagcttca ggcaccacca ctgacctggg acagtgaatc cggactctaa 60
ggtaaataata aaatttttta gtgtataatg tgtaaacta ctgattctaa ttgtttctct 120
cttttagatt ccaacctttg gaactgaatt ctagaccacc 160

<210> 328
<211> 29
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 328
accactttca caatctgcta gcaaaggtt 29

<210> 329
<211> 133
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 329
gtaagtatca aggttacaag acaggtttta ggagaccaat agaaactggg cttgtcgaga 60
cagagaagac tcttgcgttt ctgataggca cctattggtc ttactgacat ccactttgcc 120
tttctctcca cag 133

<210> 330

<211> 341
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 330
tgggcaggaa ctgggcactg tgcccagggc atgcactgcc tccacgcagc aaccctcaga 60
gtcctgagct gaaccaagaa ggaggagggg gtcgggcctc cgaggaaggc ctagccgctg 120
ctgctgccag gaattccagg ttggaggggc ggcaacctcc tgccagcctt caggccactc 180
tcctgtgcct gccagaagag acagagcttg aggagagctt gaggagagca ggaaagcctc 240
ccccgttgcc cctctggatc cactgcttaa atacggacga ggacagggcc ctgtctcctc 300
agcttcaggc accaccactg acctgggaca gtgaatcgac a 341

<210> 331
<211> 316
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 331
tctagagaag ctttattgcg gtagtttatc acagttaaatt tgctaacgca gtcagtgcctt 60
ctgacacaac agtctcgaac ttaagctgca gtgactctct taaggtagcc ttgcagaagt 120
tggtcgtgag gactcgggca ggtaagtatc aaggttacaa gacaggttta aggagaccaa 180
tagaaactgg gcttgtcgag acagagaaga ctcttgcgtt tctgataggc acctattggt 240
cttactgaca tccactttgc ctttctctcc acaggtgtcc actcccagtt caattacagc 300
tcttaaggcc ctgcag 316

<210> 332
<211> 76
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 332
caaagtccag gccctctgc tgcagcgccc gcgcgtccag aggccctgcc agacacgcgc 60
gaggttcgag gctgag 76

<210> 333
<211> 127
<212> DNA
<213> Artificial Sequence

<220>

<221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 333
 agaatgatga aaaccgaggt tggaaaaggt tgtgaaacct tttaactctc cacagtggag 60
 tccattatatt cctctggctt cctcaaattc atattcacag ggtcgttggc tgtgggttgc 120
 aattacc 127

<210> 334
 <211> 80
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 334
 atagcagagc aatcaccacc aagcctggaa taactgcaag ggctctgctg acatcttcct 60
 gaggtgccaa ggaaatgagg 80

<210> 335
 <211> 208
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 335
 gggtcaccac cacctccaca gcacagacag acactcagga gccagccagc caggtaagtt 60
 tagtcttttt gtcttttatt tcaggtcccg gatccggtgg tggtgcaaat caaagaactg 120
 ctcttcagtg gatgttgctt ttacttctag gcctgtacgg aagtgttact tctgctctaa 180
 aagctgcgga attgtacccg cggccgcg 208

<210> 336
 <211> 159
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 336
 aagcttctgc cttctccctc ctgtgagttt ggtaagtcac tgactgtcta tgcctgggaa 60
 aggggtgggca ggagatgggg cagtcagga aaagtggcac tatgaaccct gcagccctag 120
 acaattgtac taaccttctt ctctttcctc tcctgacag 159

<210> 337
 <211> 36
 <212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 337

cgcgccatgc agtgtcccag ccgggttcgt gtcgcc

36

<210> 338

<211> 141

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 338

acgccgcctg ggtcccagtc cccgtcccat cccccggcgg cctaggcagc gtttccagcc

60

ccgagaactt tgttcttttt gtcccgcccc ctgcgcccac ccgcctgcgc cgccttcggg

120

cccagattct ggagactcaa c

141

<210> 339

<211> 110

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 339

gttgatgaa accttctctc tactgcacag cccgcccccc tacagccccg gtccccacgc

60

ctagaagaca gcggaactaa gaaaagaaga ggcctgtgga cagaacaatc

110

<210> 340

<211> 164

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 340

ggtggggcgg ggttgagtcg gaaccacaat agccaggcga agaaactaca actcccaggg

60

cgtcccggag caggccaacg ggactacggg aagcagcggg cagcggcccc cgggaggcac

120

ctcggagatc tgggtgcaaa agcccagggt taggaaccgt aggc

164

<210> 341

<211> 127

<212> DNA

<213> Artificial Sequence

<220>

<221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 341
 ggccaccgga attaacctt cagggtggg ggccgcgcta tgccccgcc cctccccagc 60
 cccagacacg gacccgcag gagatgggtg ccccatccg cacactgtcc ttggccacc 120
 ggacatc 127

<210> 342
 <211> 341
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 342
 tgggcaggaa ctgggcactg tgcccagggc atgcactgcc tccacgcagc aaccctcaga 60
 gtcctgagct gaaccaagaa ggaggagggg gtcgggcctc cgaggaaggc ctagccgctg 120
 ctgctgccag gaattccagg ttggaggggc ggcaacctcc tgccagcctt caggccactc 180
 tcctgtgcct gccagaagag acagagcttg aggagagctt gaggagagca ggaaagcctc 240
 ccccgttgcc cctctggatt cactgcttaa atacggacga ggacagggcc ctgtctcctc 300
 agcttcaggc accaccactg acctgggaca gtgaatcgac a 341

<210> 343

<400> 343
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<210> 344

<400> 344
 000

<210> 345
 <211> 581
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 345
 gagcatctta ccgccattta ttcccatatt tgttctgttt ttcttgattt gggatatacat 60
 ttaaatgtta ataaaacaaa atgggtgggc aatcatttac attttaggg atatgtaatt 120
 actagttcag gtgtattgcc acaagacaaa catgttaaga aactttccg ttatttacgc 180
 tctgttcctg ttaatcaacc tctggattac aaaatttggtg aaagattgac tgatattctt 240
 aactatgttg ctctttttac gctgtgtgga tatgtgctt tatagcctct gtatctagct 300

attgcttccc gtacggcttt cgttttctcc tccttgtata aatcctggtt gctgtctctt	360
ttagaggagt tgtggcccgt tgtccgtcaa cgtggcgtgg tgtgctctgt gtttgctgac	420
gcaacccccca ctggctgggg cattgccacc acctgtcaac tcctttctgg gactttcgct	480
ttccccctcc cgatcgccac ggcagaactc atcgccgcct gccttgcccc ctgctggaca	540
ggggctaggt tgctgggcac tgataattcc gtggtgttgt c	581

<210> 346
 <211> 77
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 346	
tccataaagt aggaacact acacgattcc ataaagtagg aaacactaca tcactccata	60
aagtaggaaa cactaca	77

<210> 347
 <211> 88
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

<400> 347	
tgaaagatgg atttccaagg ttaattcatt ggaattgaaa attaacagag atctagagct	60
gaattcctgc agccaggggg atcagcct	88

<210> 348
 <211> 395
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

<400> 348	
taaaatacag catagcaaaa cttaacctc caaatcaagc ctctacttga atccttttct	60
gagggatgaa taaggcatag gcatcagggg ctgttgccaa tgtgcattag ctgtttgcag	120
cctcaccttc tttcatggag tttaagatat agtgtatttt cccaaggttt gaactagctc	180
ttcatttctt tatgttttaa atgcactgac ctccacatt cccttttttag taaaatattc	240
agaaataatt taaatacatc attgcaatga aaataaatgt tttttattag gcagaatcca	300
gatgctcaag gcccttcata atatccccc gtttagtagt tggacttagg gaacaaagga	360
acctttaata gaaattggac agcaagaaag cgagc	395

<210> 349
<211> 800
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 349
agtcaatatg ttcaccccaa aaaagctggt tggttaacttg ccaacctcat tctaaaatgt 60
atatagaagc ccaaaagaca ataacaaaaa tattcttgta gaacaaaatg ggaaagaatg 120
ttccactaaa tatcaagatt tagagcaaag catgagatgt gtggggatag acagtgaggc 180
tgataaaata gagtagagct cagaaacaga ccatttgata tatgtaagt acctatgaaa 240
aaaatatggc attttacaat gggaaaatga tgggtctttt cttttttaga aaaacaggga 300
aatatattta tatgtaaaaa ataaaaggga acccatatgt cataccatac acacaaaaaa 360
attccagtga attataagtc taaatggaga aggcaaaact ttaaattctt tagaaaataa 420
tatagaagca tgccatcaag acttcagtgt agagaaaaat ttcttatgac tcaaagtcct 480
aaccacaaag aaaagattgt taattagatt gcatgaatat taagacttat ttttaaaatt 540
aaaaaaccat taagaaaagt caggccatag aatgacagaa aatatttgca acacccagt 600
aaagagaatt gtaatatgca gattataaaa agaagtctta caaatcagta aaaaataaaa 660
ctagacaaaa atttgaacag atgaaagaga aactctaaat aatcattaca catgagaaac 720
tcaatctcag aaatcagaga actatcattg catatacact aaattagaga aatattaaaa 780
ggctaagtaa catctgtggc 800

<210> 350
<211> 407
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 350
aattatctct aaggcatgtg aactggctgt ctgggttttc atctgtactt catctgctac 60
ctctgtgacc tgaacatat ttataattcc attaagctgt gcatatgata gatttatcat 120
atgtattttc cttaaaggat ttttgtaaga actaattgaa ttgatacctg taaagtcttt 180
atcacactac ccaataaata ataaatctct ttgttcagct ctctgtttct ataaatatgt 240
accagtttta ttgttttttag tggtagtgat tttattctct ttctatatat atacacacac 300
atgtgtgcat tcataaatat atacaatttt tatgaataaa aaattattag caatcaatat 360
tgaaaaccac tgatttttgt ttatgtgagc aaacagcaga ttaaaag 407

<210> 351
<211> 186
<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 351

catcacattt aaaagcatct cagcctacca tgagaataag agaaagaaaa tgaagatcaa	60
aagcttattc atctgttttt ctttttcggt ggtgtaaagc caacaccctg tctaaaaaac	120
ataaatttct ttaatcattt tgcctctttt ctctgtgctt caattaataa aaaatggaaa	180
gaatct	186

<210> 352

<211> 395

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 352

taaaaatacag catagcaaaa cttaacctc caaatcaagc ctctacttga atccttttct	60
gagggatgaa taaggcatag gcatcagggg ctgttgccaa tgtgcattag ctgtttgcag	120
cctcaccttc tttcatggag ttaagatat agtgtatttt cccaaggttt gaactagctc	180
ttcatttctt tatgttttaa atgcactgac ctcccacatt cccttttttag taaaatattc	240
agaaataatt taaatacatc attgcaatga aaataaatgt tttttattag gcagaatcca	300
gatgctcaag gcccttcata atatcccca gtttagtagt tggacttagg gaacaaagga	360
acctttaata gaaattggac agcaagaaag ccagc	395

<210> 353

<211> 580

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 353

gagcatctta cgcatttta ttcccatatt tgttctgttt ttcttgattt gggtatacat	60
ttaaatgtta ataaaacaaa atggtggggc aatcatttac atttttaggg atatgtaatt	120
actagttcag gtgtattgcc acaagacaaa catgttaaga aactttcccg ttatttacgc	180
tctgttcctg ttaatcaacc tctggattac aaaatttgtg aaagattgac tgatattctt	240
aactatgttg ctctttttac gctgtgtgga tatgctgctt tatagcctct gtatctagct	300
attgcttccc gtacggcttt cgttttctcc tccttgtata aatcctgggt gctgtctctt	360
ttagaggagt tgtggccgtg tgtccgtcaa cgtggcgtgg tgtgctctgt gtttgctgac	420
gcaaccccca ctggctgggg cattgccacc acctgtcaac tcctttctgg gactttcgct	480

ttccccctcc cgatcgccac ggcagaactc atcgccgcct gccttgcccg ctgctggaca	540
ggggctaggt tgctgggcac tgataattcc gtggtgttgt	580
<210> 354	
<211> 64	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	
<223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"	
<400> 354	
cctcgccccg gacctgccct cccgccaggt gcacccacct gcaataaatg cagcgaagcc	60
ggga	64
<210> 355	
<211> 247	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"	
<400> 355	
gataatcaac ctctggatta caaaatttgt gaaagattga ctggtattct taactatgtt	60
gctcctttta cgctatgttg atacgctgct ttaatgcctt tgtatcatgc tattgcttcc	120
cgtatggctt tcattttctc ctccctgtat aaatcctggt tagttcttgc cacggcggaa	180
ctcatcgccg cctgccttgc ccgctgctgg acaggggctc ggctgttggg cactgacaat	240
tccgtgg	247
<210> 356	
<211> 144	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"	
<400> 356	
aaatacatca ttgcaatgaa aataaatgtt ttttattagg cagaatccag atgtcaagg	60
cccttcataa tatccccag tttagtagtt ggacttaggg aacaaaggaa cctttaatag	120
aaattggaca gcaagaaagc gagc	144
<210> 357	
<211> 62	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	

<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 357

gagcatctta ccgccattta ttcccatatt tgttctgttt ttcttgattt gggatacat 60

tt 62

<210> 358

<400> 358

000

<210> 359

<400> 359

000

<210> 360

<211> 225

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 360

tgtgccttct agttgccagc catctgttgt ttgccctcc cccgtgcctt ccttgaccct 60

ggaagggtgcc actcccactg tcctttccta ataaaatgag gaaattgcat cgcattgtct 120

gagtaggtgt cattctattc tggggggtgg ggtggggcag gacagcaagg gggaggattg 180

ggaagacaat agcaggcatg ctggggatgc ggtgggctct atggc 225

<210> 361

<211> 49

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 361

aataaaagat ctttattttc attagatctg tgtgttggtt ttttgttg 49

<210> 362

<211> 54

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic
oligonucleotide"

<400> 362

gcggccgcaa taaaagatca gagctctaga gatctgtgtg ttggtttttt gtgt 54

<210> 363
 <211> 74
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 oligonucleotide"

 <400> 363
 ggatccaata aaatatcttt attttcatta catctgtgtg ttggtttttt gtgtgttttc 60
 ctgtaacgat cggg 74

 <210> 364
 <211> 143
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

 <400> 364
 ctcgatgctt tatttgtgaa atttgtgatg ctattgcttt atttgtaacc attataagct 60
 gcaataaaca agttaacaac aacaattgca ttcattttat gtttcagggt cagggggagg 120
 tgtgggaggt tttttaaaact agt 143

 <210> 365
 <211> 228
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

 <400> 365
 ctactgtgcc ttctagttagc cagccatctg ttgtttgccc ctcccccttg ccttccttga 60
 ccctggaagg tgccactccc actgtccttt cctaataaaa tgaggaaatt gcatcacatt 120
 gtctgagtag gtgtcattct attctggggg gtgggggtggg gcaggacagc aagggggagg 180
 attgggaaga caatagcagg catgctgggg atgcagtggg ctctatgg 228

 <210> 366
 <211> 222
 <212> DNA
 <213> Artificial Sequence

 <220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic
 polynucleotide"

 <400> 366
 cagacatgat aagatacatt gatgagtttg gacaaaccac aactagaatg cagtgaaaaa 60
 aatgctttat ttgtgaaatt tgtgatgcta ttgctttatt tgtaaccatt ataagctgca 120

ataaacaagt taacaacaac aattgcattc attttatgtt tcaggttcag ggggagatgt	180
gggagggtttt ttaaagcaag taaaacctct acaaatgtgg ta	222
<210> 367	
<211> 226	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"	
<400> 367	
ccagacatga taagatacat tgatgagttt ggacaaacca caactagaat gcagtgaaaa	60
aaatgcttta tttgtgaaat ttgtgatgct attgctttat ttgtaaccat tataagctgc	120
aataaacaag ttaacaacaa caattgcatt cattttatgt ttcaggttca gggggagggtg	180
tgggagggttt tttaaagcaa gtaaacctc tacaatgtg gtatgg	226
<210> 368	
<211> 129	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"	
<400> 368	
gttaacaaaa aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa	60
aaaaaaaaaa tgcattcccc ccccccccc ccccccccc ccccccaaag gctcttttca	120
gagccacca	129
<210> 369	
<211> 232	
<212> DNA	
<213> Artificial Sequence	
<220>	
<221> source	
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"	
<400> 369	
gcggccgcgg ggatccagac atgataagat acattgatga gtttgacaa accacaacta	60
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ccattataag ctgcaataaa caagttaaca acaacaattg cattcatttt atgtttcagg	180
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<210> 370	
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<220>	

<221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

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 <213> Artificial Sequence

<220>
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 <223> /note="Description of Artificial Sequence: Synthetic oligonucleotide"

<400> 371
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<210> 372
 <211> 226
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 372
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 ttacaaataa agcaatagca tcacaaattt cacaaataaa gcattttttt cactgcattc 180
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<210> 373
 <211> 416
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 373
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 gaatctaata gagggtgaca gcactgttat ttttcaaaga tgtgttgcta tcctgaaaat 240
 tctgtaggtt ctgtggaagt tccagtgttc tctcttattc cacttcggta gaggatttct 300
 agtttcttgt gggctaatta aataaatcat taatactctt ctaagttatg gattataaac 360

attcaaaata atattttgac attatgataa ttctgaataa aagaacaaaa accatg 416

<210> 374

<211> 415

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 374

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agcttattca tctgtttttc tttttcgttg gtgtaaagcc aacaccctgt ctaaaaaaca 120

taaatttctt taatcathtt gcctcttttc tctgtgcttc aattaataaa aaatggaaag 180

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<210> 375

<211> 122

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 375

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tt 122

<210> 376

<211> 133

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 376

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gaggtttttt aaa 133

<210> 377

<211> 1386

<212> DNA

<213> Homo sapiens

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acttaa 1386

<210> 378
<211> 461
<212> PRT
<213> Homo sapiens

<400> 378
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Ile Cys Leu Leu Gly Tyr Leu Leu Ser Ala Glu Cys Thr Val Phe Leu
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35 40 45

Ser Gly Lys Leu Glu Glu Phe Val Gln Gly Asn Leu Glu Arg Glu Cys
50 55 60

Met Glu Glu Lys Cys Ser Phe Glu Glu Ala Arg Glu Val Phe Glu Asn
65 70 75 80

Thr Glu Arg Thr Thr Glu Phe Trp Lys Gln Tyr Val Asp Gly Asp Gln
85 90 95

Cys Glu Ser Asn Pro Cys Leu Asn Gly Gly Ser Cys Lys Asp Asp Ile
100 105 110

Asn Ser Tyr Glu Cys Trp Cys Pro Phe Gly Phe Glu Gly Lys Asn Cys
115 120 125

Glu Leu Asp Val Thr Cys Asn Ile Lys Asn Gly Arg Cys Glu Gln Phe
130 135 140

Cys Lys Asn Ser Ala Asp Asn Lys Val Val Cys Ser Cys Thr Glu Gly
145 150 155 160

Tyr Arg Leu Ala Glu Asn Gln Lys Ser Cys Glu Pro Ala Val Pro Phe
165 170 175

Pro Cys Gly Arg Val Ser Val Ser Gln Thr Ser Lys Leu Thr Arg Ala
180 185 190

Glu Thr Val Phe Pro Asp Val Asp Tyr Val Asn Ser Thr Glu Ala Glu
195 200 205

Thr Ile Leu Asp Asn Ile Thr Gln Ser Thr Gln Ser Phe Asn Asp Phe
210 215 220

Thr Arg Val Val Gly Gly Glu Asp Ala Lys Pro Gly Gln Phe Pro Trp
225 230 235 240

Gln Val Val Leu Asn Gly Lys Val Asp Ala Phe Cys Gly Gly Ser Ile
245 250 255

Val Asn Glu Lys Trp Ile Val Thr Ala Ala His Cys Val Glu Thr Gly
260 265 270

Val Lys Ile Thr Val Val Ala Gly Glu His Asn Ile Glu Glu Thr Glu
275 280 285

His Thr Glu Gln Lys Arg Asn Val Ile Arg Ile Ile Pro His His Asn
290 295 300

Tyr Asn Ala Ala Ile Asn Lys Tyr Asn His Asp Ile Ala Leu Leu Glu
305 310 315 320

Leu Asp Glu Pro Leu Val Leu Asn Ser Tyr Val Thr Pro Ile Cys Ile

325

330

335

Ala Asp Lys Glu Tyr Thr Asn Ile Phe Leu Lys Phe Gly Ser Gly Tyr
340 345 350

Val Ser Gly Trp Gly Arg Val Phe His Lys Gly Arg Ser Ala Leu Val
355 360 365

Leu Gln Tyr Leu Arg Val Pro Leu Val Asp Arg Ala Thr Cys Leu Arg
370 375 380

Ser Thr Lys Phe Thr Ile Tyr Asn Asn Met Phe Cys Ala Gly Phe His
385 390 395 400

Glu Gly Gly Arg Asp Ser Cys Gln Gly Asp Ser Gly Gly Pro His Val
405 410 415

Thr Glu Val Glu Gly Thr Ser Phe Leu Thr Gly Ile Ile Ser Trp Gly
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435 440 445

Arg Tyr Val Asn Trp Ile Lys Glu Lys Thr Lys Leu Thr
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<210> 379

<211> 2688

<212> DNA

<213> Homo sapiens

<400> 379

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<210> 380

<211> 423

<212> PRT

<213> Homo sapiens

<400> 380

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Asp His Glu Asn Ala Asn Lys Ile Leu Asn Arg Pro Lys Arg Tyr Asn
35 40 45

Ser Gly Lys Leu Glu Glu Phe Val Gln Gly Asn Leu Glu Arg Glu Cys
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Met Glu Glu Lys Cys Ser Phe Glu Glu Ala Arg Glu Val Phe Glu Asn
65 70 75 80

Thr Glu Arg Thr Thr Glu Phe Trp Lys Gln Tyr Val Asp Val Thr Cys
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Asn Ile Lys Asn Gly Arg Cys Glu Gln Phe Cys Lys Asn Ser Ala Asp
100 105 110

Asn Lys Val Val Cys Ser Cys Thr Glu Gly Tyr Arg Leu Ala Glu Asn
115 120 125

Gln Lys Ser Cys Glu Pro Ala Val Pro Phe Pro Cys Gly Arg Val Ser
130 135 140

Val Ser Gln Thr Ser Lys Leu Thr Arg Ala Glu Thr Val Phe Pro Asp
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Val Asp Tyr Val Asn Ser Thr Glu Ala Glu Thr Ile Leu Asp Asn Ile
165 170 175

Thr Gln Ser Thr Gln Ser Phe Asn Asp Phe Thr Arg Val Val Gly Gly
180 185 190

Glu Asp Ala Lys Pro Gly Gln Phe Pro Trp Gln Val Val Leu Asn Gly
195 200 205

Lys Val Asp Ala Phe Cys Gly Gly Ser Ile Val Asn Glu Lys Trp Ile
210 215 220

Val Thr Ala Ala His Cys Val Glu Thr Gly Val Lys Ile Thr Val Val
225 230 235 240

Ala Gly Glu His Asn Ile Glu Glu Thr Glu His Thr Glu Gln Lys Arg
245 250 255

Asn Val Ile Arg Ile Ile Pro His His Asn Tyr Asn Ala Ala Ile Asn
260 265 270

Lys Tyr Asn His Asp Ile Ala Leu Leu Glu Leu Asp Glu Pro Leu Val

275

280

285

Leu Asn Ser Tyr Val Thr Pro Ile Cys Ile Ala Asp Lys Glu Tyr Thr
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Asn Ile Phe Leu Lys Phe Gly Ser Gly Tyr Val Ser Gly Trp Gly Arg
 305 310 315 320

Val Phe His Lys Gly Arg Ser Ala Leu Val Leu Gln Tyr Leu Arg Val
 325 330 335

Pro Leu Val Asp Arg Ala Thr Cys Leu Arg Ser Thr Lys Phe Thr Ile
 340 345 350

Tyr Asn Asn Met Phe Cys Ala Gly Phe His Glu Gly Gly Arg Asp Ser
 355 360 365

Cys Gln Gly Asp Ser Gly Gly Pro His Val Thr Glu Val Glu Gly Thr
 370 375 380

Ser Phe Leu Thr Gly Ile Ile Ser Trp Gly Glu Glu Cys Ala Met Lys
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Gly Lys Tyr Gly Ile Tyr Thr Lys Val Ser Arg Tyr Val Asn Trp Ile
 405 410 415

Lys Glu Lys Thr Lys Leu Thr
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<210> 381

<211> 1386

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 381

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<210> 382

<211> 2824

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 382

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tgggatctac acaaaagtat ccagatatgt gaactggatt aaggagaaaa ccaagctgac	2820
ttga	2824

<210> 383
<211> 1386
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 383
atgcagaggg tgaacatgat catggctgag agccctggcc tgatcaccat ctgcctgctg 60
ggctacctgc tgtctgctga atgtacagtt tttcttgatc atgaaaatgc caacaaaatt 120
ctgaatagac caaagaggta taactctggc aagcttgaag agtttgtaca ggggaatctg 180
gagagagagt gtatggaaga gaagtgcagc tttgaggaag ccagagaagt gtttgaaaat 240
acagagagaa caactgaatt ttggaagcag tatgtggatg gtgatcaatg tgagagcaat 300
ccctgcttga atggggggag ctgtaaagat gatatcaaca gctatgaatg ttggtgtccc 360
tttgatttg aggggaaaaa ctgtgagctt gatgtgacct gtaatatcaa gaatggcagg 420
tgtgagcaat tttgcaagaa ttctgctgat aacaaagtgg tctgtagctg cactgaggga 480
tataggctgg ctgaaaacca gaagagctgt gaacctgcag tgccttttcc ctgtgggaga 540
gtgtctgtga gccaaaccag caagctgact agggctgaag cagtctttcc tgatgtagat 600
tatgtgaata gactgaggc tgagacaatc cttgacaata tcactcagag cacacagagc 660
ttcaatgact tcaccagggt ggtaggaggg gaggatgcca agcctgggca gttcccctgg 720
caggtagtgc tcaatggaaa agtggatgcc ttttgtggag gttcaattgt aaatgagaag 780
tggattgtga ctgcagccca ctgtgtggaa actggagtca agattactgt ggtggctgga 840
gagcacaata ttgaggaaac tgagcacact gagcagaaga ggaatgtgat caggattatc 900
ccccaccaca actacaatgc tgctatcaac aagtacaacc atgacattgc cctcctggaa 960
ctggatgaac ccctggctct gaacagctat gtgacacca tctgtattgc tgataaagag 1020
tacaccaaca tcttcttgaa atttgggtct ggatatgtgt ctggctgggg cagggtgttc 1080
cataaaggca ggtctgccct ggtattgcag tatttgaggg tgcctctggt ggatagagca 1140
acctgcttgc tgagcaccaa gtttacaatc tacaacaata tgttctgtgc agggttccat 1200
gaaggtggta gagacagctg ccaggagat tctgggggtc cccatgtgac tgaggtggag 1260
ggaaccagct tcctgactgg gattatcagc tggggtgagg agtgtgctat gaagggaag 1320
tatgggatct acacaaaagt atccagatat gtgaactgga ttaaggagaa aaccaagctg 1380
acttga 1386

<210> 384
<211> 1386
<212> DNA
<213> Homo sapiens

<400> 384
atgcagcgcg tgaacatgat catggcagaa tcaccaggcc tcatcaccat ctgcctttta 60
ggatatctac tcagtgtga atgtacagtt tttcttgatc atgaaaacgc caacaaaatt 120

ctgaatcggc caaagaggta taattcaggt aaattggaag agtttgttca agggaacctt	180
gagagagaat gtatggaaga aaagtgtagt tttgaagaag cacgagaagt ttttgaaaac	240
actgaaagaa caactgaatt ttggaagcag tatgttgatg gagatcagtg tgagtccaat	300
cctatgtttaa atggcggcag ttgcaaggat gacattaatt cctatgaatg ttggtgtccc	360
tttggttttg aaggaaagaa ctgtgaatta gatgtaacat gtaacattaa gaatggcaga	420
tgcgagcagt tttgtaaaaa tagtgctgat aacaaggtgg tttgctcctg tactgaggga	480
tatcgacttg cagaaaacca gaagtcctgt gaaccagcag tgccatttcc atgtggaaga	540
gtttctgttt cacaaacttc taagctcacc cgtgctgaga ctgtttttcc tgatgtggac	600
tatgtaaatt ctactgaagc tgaaccatt ttggataaca tcaactaaag cacccaatca	660
tttaatgact tcactcgggt tgttggtgga gaagatgcca aaccaggtca attcccttgg	720
caggttggtt tgaatggtaa agttgatgca ttctgtggag gctctatcgt taatgaaaaa	780
tggattgtaa ctgctgcccc ctgtgttgaa actggtgtta aaattacagt tgtcgaggt	840
gaacataata ttgaggagac agaacataca gagcaaaagc gaaatgtgat tcgaattatt	900
cctcaccaca actacaatgc agctattaat aagtacaacc atgacattgc ctttctggaa	960
ctggacgaac ccttagtgct aaacagctac gttacaccta ttgcatcgc tgacaaggaa	1020
tacacgaaca tcttcctcaa atttgatctt ggctatgtaa gtggctgggg aagagtcttc	1080
cacaaaggga gatcagcttt agttcttcag taccttagag ttccacttgt tgaccgagcc	1140
acatgtcttc gatctacaaa gttcaccatc tataacaaca tgttctgtgc tggcttccat	1200
gaaggaggta gagattcatg tcaaggagat agtgggggac cccatgttac tgaagtggaa	1260
gggaccagtt tcttaactgg aattattagc tggggtgaag agtgtgcaat gaaaggcaaa	1320
tatggaatat ataccaaggt atcccggat gtcaactgga ttaaggaaaa aacaaagctc	1380
acttaa	1386

<210> 385

<211> 1386

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 385

atgcagcgcg tgaacatgat catggccgag agccccggcc tgatcaccat ctgcctgctg	60
ggctacctgc tgagcgccga gtgcaccgtg ttcttgacc acgagaacgc caacaagatc	120
ctgaaccgcc ccaagcgcta caacagcggc aagctggagg agttcgtgca gggcaacctg	180
gagcgcgagt gcatggagga gaagtgcagc ttcgaggagg cccgcgaggt gttcgagaac	240
accgagcgca ccaccaggtt ctggaagcag tacgtggacg gcgaccagtg cgagagcaac	300
ccctgcctga acggcggcag ctgcaaggac gacatcaaca gctacgagtg ctggtgcccc	360

ttcggcttcg agggcaagaa ctgcgagctg gacgtgacct gcaacatcaa gaacggccgc	420
tgcgagcagt tctgcaagaa cagcgccgac aacaaggtgg tgtgcagctg caccgagggc	480
taccgcctgg ccgagaacca gaagagctgc gagcccgccg tgcccttccc ctgcggcagg	540
gtgagcgtga gccagaccag caagctgacc cgcgccgagg ccgtgttccc cgacgtggac	600
tacgtgaaca gcaccgaggc cgagaccatc ctggacaaca tcacccagag caccagagc	660
ttcaacgact tcacccgcgt ggtgggaggc gaggacgcca agcccggcca gttcccctgg	720
caggtggtgc tgaacggcaa ggtggacgcc ttctgcggcg gcagcatcgt gaacgagaag	780
tggatcgtga ccgccgccc ctgcgtggag accggcgtga agatcaccgt ggtggccggc	840
gagcacaaca tcgaggagac cgagcacacc gagcagaagc gcaacgtgat ccgcatcatc	900
ccccaccaca actacaacgc cgccatcaac aagtacaacc acgacatcgc cctgctggag	960
ctggacgagc ccctggtgct gaacagctac gtgaccccc tctgcatcgc cgacaaggag	1020
tacaccaaca tcttctgaa gttcggcagc ggctacgtga gcggctgggg ccgcgtgttc	1080
cacaagggcc gcagcgccct ggtgctgcag tacctgcgcg tgcccctggt ggaccgcgcc	1140
acctgcctgc tgagcaccaa gttcaccatc tacaacaaca tgttctgcgc cggcttcac	1200
gagggcggca gggacagctg ccaggcgac agcggcggcc cccacgtgac cgaggtggag	1260
ggcaccagct tcctgaccgg catcatcagc tggggcgagg agtgcgccat gaagggaag	1320
tacggcatct acaccaaggt gagccgtac gtgaactgga tcaaggagaa gaccaagctg	1380
acctaa	1386

<210> 386

<211> 1386

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 386

atgcagcgcg tgaacatgat catggccgag agccccggcc tgatcaccat ctgcctgctg	60
ggctacctgc tgagcgccga gtgcaccgtg ttcttgacc acgagaacgc caacaagatc	120
ctgaaccgcc ccaagcgcta caacagcggc aagctggagg agttcgtgca gggcaacctg	180
gagcgcgagt gcatggagga gaagtgcagc ttcgaggagg cccgcgaggt gttcgagaac	240
accgagcgca ccaccgagtt ctggaagcag tacgtggacg gcgaccagtg cgagagcaac	300
ccctgcctga acggcggcag ctgcaaggac gacatcaaca gctacgagtg ctggtgcccc	360
ttcggcttcg agggcaagaa ctgcgagctg gacgtgacct gcaacatcaa gaacggccgc	420
tgcgagcagt tctgcaagaa cagcgccgac aacaaggtgg tgtgcagctg caccgagggc	480
taccgcctgg ccgagaacca gaagagctgc gagcccgccg tgcccttccc ctgcggcagg	540
gtgagcgtga gccagaccag caagctgacc cgcgccgagg ccgtgttccc cgacgtggac	600
tacgtgaaca gcaccgaggc cgagaccatc ctggacaaca tcacccagag caccagagc	660

ttcaacgact tcacccgcgt ggtgggcggc gaggacgcca agcccggcca gttcccctgg	720
caggtggtgc tgaacggcaa ggtggacgcc ttctgcggcg gcagcatcgt gaacgagaag	780
tggatcgtga ccgccgcccc ctgcgtggag accggcgtga agatcaccgt ggtggccggc	840
gagcacaaca tcgaggagac cgagcacacc gagcagaagc gcaacgtgat ccgcatcatc	900
ccccaccaca actacaacgc cgccatcaac aagtacaacc acgacatcgc cctgctggag	960
ctggacgagc ccctggtgct gaacagctac gtgaccccca tctgcatcgc cgacaaggag	1020
tacaccaaca tcttcctgaa gttcggcagc ggctacgtga gcggctgggg ccgctgttc	1080
cacaagggcc gcagcgccct ggtgctgcag tacctgcgcg tgcccctggt ggaccgcgcc	1140
acctgcctgc gcagcaccaa gttcaccatc tacaacaaca tgttctgcgc cggcttcac	1200
gagggcggca gggacagctg ccaggcgac agcggcggcc cccacgtgac cgaggtggag	1260
ggcaccagct tcctgaccgg catcatcagc tggggcgagg agtgcgccat gaagggaag	1320
tacggcatct acaccaaggt gagccgtac gtgaactgga tcaaggagaa gaccaagctg	1380
acctaa	1386

<210> 387

<211> 1359

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 387

atggccgagg ccagcggcct ggtgaccgtg tgcctgctgg gctacctgct gagcgccgag	60
tgcgccgtgt tcctggaccg cgagaacgcc accaagatcc tgagccgccc caagcgctac	120
aacagcggca agctggagga gttcgtgcgc ggcaacctgg agcgcgagtg catcgaggag	180
aagtgcagct tcgaggaggc ccgcgaggtg ttcgagaaca ccgagaagac caccgagttc	240
tggaagcagt acgtggacgg cgaccagtgc gagagcaacc cctgcctgaa cgacggcgtg	300
tgcaaggacg acatcaacag ctacgagtgc tggtgccgcg ccggcttcga gggcaagaac	360
tgcgagctgg acgtgacctg caacatcaag aacggccgct gcaagcagtt ctgcaagctg	420
ggccccgaca acaaggtggt gtgcagctgc accaccggct accagctggc cgaggaccag	480
cgcagctgcg agcccgccgt gcccttcccc tgcggcaggg tgagcgtgcc ccacatcagc	540
atgacccgca cccgcgccga gaccctgttc agcaacatgg actacgagaa cagcaccgag	600
gtggagaaga tcctggacaa cgtgaccag cccctgaacg acttcacccg cgtggtgggc	660
ggcaaggacg ccaagcccgg ccagttcccc tggcaggtgc tgctgaacgg caaggtggac	720
gccttctgcg gcggcagcat catcaacgag aagtgggtgg tgaccgccgc cactgcatc	780
gagcccgacg tgaagatcac catcgtggcc ggcgagcaca acaccgagaa gcgcgagcac	840
accgagcaga agcgaacgt gatccgcacc atcctgcacc acagctacaa cgccaccatc	900

aacaagtaca accacgacat cgccctgctg gagctggacg agcccctgac cctgaacagc	960
tacgtgaccc ccatctgcat cgccgaccgc gagtacagca acatcttcct gaagttcggc	1020
agcggctacg tgagcggctg gggccgcgtg ttcaacaagg gccgcagcgc cagcatcctg	1080
cagtacctga aggtgcccct ggtggaccgc gccacctgcc tgcgcagcac caagttcacc	1140
atctacaaca acatgtttctg cgccggcttc cacgagggcg gcaaggacag ctgccagggc	1200
gacagcggcg gccccacgt gaccgaggtg gagggcatca gcttcctgac cggcatcatc	1260
agctggggcg aggagtgcgc catgaaggcg aagtacggca tctacaccaa ggtgagccgc	1320
tacgtgaact ggatcaagga gaagaccaag ctgacctaa	1359

<210> 388

<211> 1386

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 388

atgcagcgcg tgaacatgat tatggccgag tctcccgcc tgatcaccat ctgtctgctg	60
ggctatctgc tgagcgccga gtgcaccgtg tttctggatc acgagaacgc caacaagatc	120
ctgaacagac ccaagcggta caacagcggc aagctggaag agttcgtgca gggcaacctg	180
gaacgcgagt gcatggaaga gaagtgcagc ttcgaagagg ccagagaggt gttcgagaac	240
accgagagaa ccaccgagtt ctggaagcag tacgtggacg gcgatcagtg cgagagcaac	300
ccttgcttga atggcggcag ctgcaaggac gacatcaaca gctacgagtg ctggtgcccc	360
ttcggcttcg agggcaagaa ttgcgagctg gacgtgacct gcaacatcaa gaacggcaga	420
tgcgagcagt tctgcaagaa cagcgccgac aacaaggctg tgtgctcctg cacagagggc	480
tacagactgg ccgagaacca gaagtcttgc gagcccgctg tgccctttcc atgtggcaga	540
gtgtctgtgt cccagaccag caagctgacc agagccgaag ccgtgtttcc cgacgtggac	600
tacgtgaaca gcaccgaggc cgagacaatc ctggacaaca tcaccagag caccagttcc	660
ttcaacgact tcaccagagt cgtcggcggc gaggatgcta agcctggaca gtttccttgg	720
caagtgggtg tgaacggcaa ggtggacgct ttttgtggcg gctccatcgt gaacgagaag	780
tggatcgtga ccgccgctca ctgtgtggaa accggcgtga agattacagt ggtggccggc	840
gagcacaaca tcgaggaaac agagcacacc gagcagaaac ggaacgtgat cagaatcatc	900
cctcaccaca actacaacgc cgccatcaac aagtacaacc acgatatcgc cctgctggaa	960
ctggacgagc ccctggctct gaactcttac gtgacccta tctgtatcgc cgacaaagag	1020
tacaccaaca tctttctgaa gttcggcagc ggctacgtgt ccggctgggg aagagttttc	1080
cacaagggca gatcagccct ggtgctgcag tacctgagag tgcccctggt ggatagagcc	1140
acatgcctgc tgagcaccaa gttcaccatc tacaacaaca tgttctgcgc cggcttccac	1200
gaaggcggca gagattcttg tcaaggcgat tctggcggcc ctcacgtgac agaggttgag	1260

ggcacaagct ttctgaccgg catcatcagc tggggcgaag agtgtgccat gaaggggaag	1320
tacggcatct acaccaaggt gtccagatac gtgaactgga tcaaagaaaa gaccaagctc	1380
acctag	1386

<210> 389
 <211> 1386
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 389	
atgcagcgcg tgaacatgat tatggccgag tctcccgcc tgatcaccat ctgtctgctg	60
ggctatctgc tgagcgccga gtgcaccgtg tttctggatc acgagaacgc caacaagatc	120
ctgaacagac ccaagcggt aacagcggc aagctggaag agttcgtgca gggcaacctg	180
gaacgcgagt gcatggaaga gaagtgcagc ttcgaagagg ccagagaggt gttcgagaac	240
accgagagaa ccaccgagtt ctggaagcag tacgtggacg gcgatcagtg cgagagcaac	300
ccttgctga atggcggcag ctgcaaggac gacatcaaca gctacgagtg ctggtgcccc	360
ttcggcttcg agggcaagaa ttgcgagctg gacgtgacct gcaacatcaa gaacggcaga	420
tgcgagcagt tctgaagaa cagcgccgac aacaaggctg tgtgctcctg cacagagggc	480
tacagactgg ccgagaacca gaagtcttgc gagcccgctg tgccctttcc atgtggcaga	540
gtgtctgtgt cccagaccag caagctgacc agagccgaag ccgtgtttcc cgacgtggac	600
tacgtgaaca gcaccgaggc cgagacaatc ctggacaaca tcaccagag caccagtcc	660
ttcaacgact tcaccagagt cgtcggcggc gaggatgcta agcctggaca gtttccttgg	720
caagtgggtc tgaacggcaa ggtggacgct ttttgtggcg gctccatcgt gaacgagaag	780
tggatcgtga ccgccgtca ctgtgtggaa accggcgtga agattacagt ggtggccggc	840
gagcacaaca tcgaggaaac agagcacacc gagcagaaac ggaacgtgat cagaatcatc	900
cctcaccaca actacaacgc cgccatcaac aagtacaacc acgatatcgc cctgctggaa	960
ctggacgagc ccctggctct gaactcttac gtgacccta tctgtatcgc cgacaaagag	1020
tacaccaaca tctttctgaa gttcggcagc ggctacgtgt ccggctgggg aagagttttc	1080
cacaagggca gatcagccct ggtgctgcag tacctgagag tgcccctggt ggatagagcc	1140
acctgtctga gaagcaccaa gttcaccatc tacaacaaca tgttctgcgc cggcttcac	1200
gaaggcggca gagattcttg tcaaggcgat tctggcggcc ctacgtgac agaggttgag	1260
ggcacaagct ttctgaccgg catcatcagc tggggcgaag agtgtgccat gaaggggaag	1320
tacggcatct acaccaaggt gtccagatac gtgaactgga tcaaagaaaa gaccaagctc	1380
acctag	1386

<210> 390

<211> 1419
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 390
atgaagcacc tgaacaccgt catggcagaa tccccggctc tcatcaccat cttcctttta 60
ggatatctac tcagtaccga atgtgcagtt ttccttgatc gtgaaaatgc caccaaaatt 120
cttaccgctc caaagagata taattcagga aaactagaag agtttgttcg aggaaacctt 180
gaaagagagt gtatagaaga aagatgtagt tttgaagaag cacgagaagt ttttgaaaac 240
actgaaaaaa ctactgaatt ttggaagcag tatgttgatg gagatcagtg tgaatcaaatt 300
ccttgtttta atggtggaat atgcaaggat gatattagtt cctatgaatg ctggtgccaa 360
gttggttttg aaggaaggaa ctgtgaatta gatgcaacgt gtaacattaa aaatggcagg 420
tgcaagcagt tttgtaaaaa cagtcctgat aacaaggtaa tttgttcctg cactgaggga 480
taccaacttg cagaagacca gaagtcctgt gaaccaacag ttccatttcc atgtgggaga 540
gcttctatct catacagttc taaaaagatc acgagagctg agactgtttt ctctaataatg 600
gactatgaaa attctactga agctgtattc attcaagatg acatcactga tggtgccatt 660
cttaataacg tcactgaaag tagtgaatca cttaatgact tcactcgagt tggttggtgga 720
gaaaacgcaa aaccgggtca aatcccttgg cagggtcattt taaatggtga aattgaggca 780
ttctgtggag gtgccatcat taatgaaaaa tggattgtaa ctgctgcca ctgtcttaaa 840
cctggtgata aaattgaggt tgttgctggt gaataataca ttgataagaa ggaagacaca 900
gaacaaagga gaaatgtgat tcgaactatc cctcatcacc agtacaatgc aactattaat 960
aagtatagtc atgacattgc cttgctggaa ctggataaac ctttaataact aaacagctat 1020
gtaacaccta tctgtgttgc caataggga tatacaaata tcttcctcaa gtttggttct 1080
ggctatgtca gtggctgggg aaaagtcttc aacaaaggga gacaggcttc cattcttcag 1140
taccttagag ttccactggt ggatagagcc acatgcctta ggtccacaac attcactatc 1200
tataacaaca tgttctgtgc aggctaccgt gaaggaggca aagattcgtg tgaaggagat 1260
agtgggggac cccatgttac tgaagtagaa gggacaagtt tcttaactgg cattattagc 1320
tggggtgaag aatgtgcaat gaaaggcaaa tatggaatat atactaaggt ttcccgttac 1380
gtcaactgga ttaaggaaaa aacaaagcta acttaatga 1419

<210> 391
<211> 5
<212> PRT
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic peptide"

<400> 391
Gly Gly Gly Gly Ser
1 5

<210> 392
<211> 6
<212> PRT
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
6xHis tag"

<400> 392
His His His His His His
1 5

<210> 393
<211> 1452
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic
polynucleotide"

<400> 393
atgaagcacc tgaacaccgt catggcagaa tccccggctc tcatcaccat cttcctttta 60
ggatatctac tcagtaccga atgtgcagtt ttccttgatc gtgaaaatgc caccaaaatt 120
cttaccctgc caaagagata taattcagga aaactagaag agtttgttcg aggaaacctt 180
gaaagagagt gtatagaaga aagatgtagt tttgaagaag cagagaagt ttttgaaaac 240
actgaaaaaa ctactgaatt ttggaagcag tatgttgatg gagatcagtg tgaatcaaat 300
ccttgtttaa atgggtggaat atgcaaggat gatattagtt cctatgaatg ctggtgccaa 360
gttggaattg aaggaaggaa ctgtgaatta gatgcaacgt gtaacattaa aaatggcagg 420
tgcaagcagt tttgtaaaaa cagtcctgat aacaaggtaa tttgttcctg cactgaggga 480
taccaacttg cagaagacca gaagtcctgt gaaccaacag ttccatttcc atgtgggaga 540
gcttctattt catacagttc taaaaagatc acgagagctg agactgtttt ctctaatatg 600
gactatgaaa attctactga agctgtattc attcaagatg acatcactga tggtgccatt 660
cttaataacg tcaactgaaag tagtgaatca cttaatgact tcactcgagt tgttggtgga 720
gaaaacgcaa aaccgggtca aatcccttgg caggctcattt taaatggtga aattgaggca 780
ttctgtggag gtgccatcat taatgaaaaa tggattgtaa ctgctgccca ctgtcttaaa 840
cctggtgata aaattgaggt tgttgctggt gaatataaca ttgataagaa ggaagacaca 900
gaacaaagga gaaatgtgat tcgaactatc cctcatcacc agtacaatgc aactattaat 960
aagtatagtc atgacattgc cttgctggaa ctggataaac ctttaatact aaacagctat 1020
gtaacaccta tctgtgttgc caatagggaa tatacaaata tcttcctcaa gtttggttct 1080
ggctatgtca gtggctgggg aaaagtcttc aacaaaggga gacaggcttc cattcttcag 1140

taccttagag ttccactggt ggatagagcc acatgcctta ggtccacaac attcactatc	1200
tataacaaca tgttctgtgc aggctaccgt gaaggaggca aagattcgtg tgaaggagat	1260
agtgggggac cccatgttac tgaagtagaa gggacaagtt tcttaactgg cattattagc	1320
tgggggtgaag aatgtgcaat gaaaggcaaa tatggaatat atactaaggt ttcccgttac	1380
gtcaactgga ttaaggaaaa aacaaagcta actgggggtg gaggtctca tcaccatcac	1440
catcactaat ga	1452

<210> 394

<211> 1464

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 394

atgaagcacc tgaacaccgt catggcagaa tccccggctc tcatcaccat cttcctttta	60
ggatatctac tcagtaccga atgtgcagtt ttccttgatc gtgaaaatgc caccaaaatt	120
cttaccgctc caaagagata taattcagga aaactagaag agtttgttcg aggaaacctt	180
gaaagagagt gtatagaaga aagatgtagt tttgaagaag cacgagaagt ttttgaaaac	240
actgaaaaaa ctactgaatt ttggaagcag tatgttgatg gagatcagtg tgaatcaaat	300
ccttgtttaa atggtggaat atgcaaggat gatattagtt cctatgaatg ctggtgccaa	360
gttggttttg aaggaaggaa ctgtgaatta gatgcaacgt gtaacattaa aaatggcagg	420
tgcaagcagt tttgtaaaaa cagtcctgat aacaaggtaa tttgttcctg cactgaggga	480
taccaacttg cagaagacca gaagtcctgt gaaccaacag ttccatttcc atgtgggaga	540
gcttctatct catcacgttc taaaaagatc acgagagctg agactgtttt ctctaatatg	600
gactatgaaa attctactga agctgtattc attcaagatg acatcactga tggtgccatt	660
cttaataacg tcactgaaag tagtgaatca cttaatgact tcactcgagt tgttggtgga	720
gaaaacgcaa aaccgggtca aatcccttgg caggctcattt taaatggtga aattgaggca	780
ttctgtggag gtgccatcat taatgaaaaa tggattgtaa ctgctgcca ctgtcttaaa	840
cctggtgata aaattgaggt tgttgctggt gaataataca ttgataagaa ggaagacaca	900
gaacaaagga gaaatgtgat tcgaactatc cctcatcacc agtacaatgc aactattaat	960
aagtatagtc atgacattgc cttgctggaa ctggataaac ctttaatact aaacagctat	1020
gtaacaccta tctgtgttgc caataggga tatacaaata tcttcctcaa gtttggttct	1080
ggctatgtca gtggctgggg aaaagtcttc aacaaaggga gacaggcttc cattcttcag	1140
taccttagag ttccactggt ggatagagcc acatgcctta ggtccacaac attcactatc	1200
tataacaaca tgttctgtgc aggctaccgt gaaggaggca aagattcgtg tgaaggagat	1260
agtgggggac cccatgttac tgaagtagaa gggacaagtt tcttaactgg cattattagc	1320
tgggggtgaag aatgtgcaat gaaaggcaaa tatggaatat atactaaggt ttcccgttac	1380

gtcaactgga ttaaggaaaa aacaaagcta actgggggtg gaggctctga gcagaaactc 1440

atctctgaac aggatctgta atga 1464

<210> 395

<211> 1419

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 395

atgaagcacc tgaacaccgt catggcagaa tccccggctc tcatcaccat cttcctttta 60

ggatatctac tcagtaccga atgtgcagtt ttccttgatc gtgaaaatgc caccaaaatt 120

cttaccgctc caaagagata taattcagga aaactagaag agtttgttcg aggaaacctt 180

gaaagagagt gtatagaaga aagatgtagt tttgaagaag cacgagaagt ttttgaaaac 240

actgaaaaaa ctactgaatt ttggaagcag tatgttgatg gagatcagtg tgaatcaaatt 300

ccttgtttaa atggtggaat atgcaaggat gatattagtt cctatgaatg ctggtgccaa 360

gttggttttg aaggaaggaa ctgtgaatta gatgcaacgt gtaacattaa aaatggcagg 420

tgcaagcagt tttgtaaaaa cagtcctgat aacaaggtaa tttgttcctg cactgaggga 480

taccaacttg cagaagacca gaagtcctgt gaaccaacag ttccatttcc atgtgggaga 540

gcttctatatt catcacgttc taaaaagatc acgagagctg agactgtttt ctctaatatg 600

gactatgaaa attctactga agctgtattc attcaagatg acatcactga tggtgccatt 660

cttaataacg tcactgaaag tagtgaatca cttaatgact tcactcgagt tggttggtgga 720

gaaaacgcaa aaccgggtca aatcccttgg caggctcattt taaatggtga aattgaggca 780

ttctgtggag gtgccatcat taatgaaaaa tggattgtaa ctgctgccca ctgtcttaaa 840

cctggtgata aaattgaggt tgttgctggt gaatataaca ttgataagaa ggaagacaca 900

gaacaaagga gaaatgtgat tcgaactatc cctcatcacc agtacaatgc aactattaat 960

aagtatagtc atgacattgc cttgctggaa ctggataaac ctttaatact aaacagctat 1020

gtaacaccta tctgtgttgc caataggga tatacaaata tcttcctcaa gtttggttct 1080

ggctatgtca gtggctgggg aaaagtcttc aacaaaggga gacaggcttc cattcttcag 1140

taccttagag ttccactggt ggatagagcc acatgccttc tgtccacaac attcactatc 1200

tataacaaca tgttctgtgc aggctaccgt gaaggaggca aagattcgtg tgaaggagat 1260

agtgggggac cccatgttac tgaagtagaa gggacaagtt tcttaactgg cattattagc 1320

tgggggtgaag aatgtgcaat gaaaggcaaa tatggaatat atactaaggt ttcccgtac 1380

gtcaactgga ttaaggaaaa aacaaagcta acttaatga 1419

<210> 396

<211> 1452

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 396

atgaagcacc tgaacaccgt catggcagaa tccccggctc tcatcaccat cttcctttta	60
ggatatctac tcagtaccga atgtgcagtt ttccttgatc gtgaaaatgc caccaaaatt	120
cttaccgctc caaagagata taattcagga aaactagaag agtttgttcg aggaaacctt	180
gaaagagagt gtatagaaga aagatgtagt tttgaagaag cacgagaagt ttttgaaaac	240
actgaaaaaa ctactgaatt ttggaagcag tatgttgatg gagatcagtg tgaatcaaat	300
ccttgtttaa atgggtggaat atgcaaggat gatattagtt cctatgaatg ctggtgccaa	360
gttggttttg aaggaaggaa ctgtgaatta gatgcaacgt gtaacattaa aaatggcagg	420
tgcaagcagt tttgtaaaaa cagtcctgat aacaaggtaa tttgttcctg cactgaggga	480
taccaacttg cagaagacca gaagtcctgt gaaccaacag ttccatttcc atgtgggaga	540
gcttctatth catacagttc taaaaagatc acgagagctg agactgtttt ctctaatatg	600
gactatgaaa attctactga agctgtattc attcaagatg acatcactga tggtgccatt	660
cttaataacg tcactgaaag tagtgaatca cttaatgact tcactcgagt tgttggtgga	720
gaaaacgcaa aaccgggtca aatcccttgg caggctcattt taaatggtga aattgaggca	780
ttctgtggag gtgccatcat taatgaaaaa tggattgtaa ctgctgcca ctgtcttaaa	840
cctggtgata aaattgaggt tgttgctggt gaatataaca ttgataagaa ggaagacaca	900
gaacaaagga gaaatgtgat tcgaactatc cctcatcacc agtacaatgc aactattaat	960
aagtatagtc atgacattgc cttgctggaa ctggataaac ctttaatact aaacagctat	1020
gtaacaccta tctgtgttgc caatagggaa tatacaaata tcttcctcaa gtttggttct	1080
ggctatgtca gtggctgggg aaaagtcttc aacaaaggga gacaggcttc cattcttcag	1140
taccttagag ttccactggt ggatagagcc acatgccttc tgtccacaac attcactatc	1200
tataacaaca tgttctgtgc aggctaccgt gaaggaggca aagattcgtg tgaaggagat	1260
agtgggggac cccatgttac tgaagtagaa gggacaagtt tcttaactgg cattattagc	1320
tgggggtgaag aatgtgcaat gaaaggcaaa tatggaatat atactaaggt ttcccgttac	1380
gtcaactgga ttaaggaaaa aacaaagcta actgggggtg gaggctctca tcaccatcac	1440
catcactaat ga	1452

<210> 397

<211> 1386

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 397
atgcagagag tcaacatgat catggcagag tccccagggt tgattactat ctgtttgctg 60
gggtacctgt tgtctgcaga atgtacagt ttcttagacc atgagaatgc caacaagatt 120
ctcaacaggc ccaagaggta caacagtgga aagctggagg aatttgtgca gggtaacctg 180
gaaagggagt gcatggagga gaaatgtagt tttgaggagg caagagaagt atttgaaaac 240
actgagagga ccactgagtt ctggaagcag tatgttgatg gagaccagt tgagagcaac 300
ccctgcttga atgggggctc atgtaaagat gatattaatt catatgaatg ctggtgcccc 360
tttggctttg aaggtaagaa ctgtgagctg gatgtcacat gcaacattaa aaatggaaga 420
tgtgaacagt tctgcaaaaa ctgagctgac aacaaagttg tctgttcctg cactgaaggc 480
tacagacttg cagagaatca gaagagttgt gaacctgcag tcccatttcc atgtgggagg 540
gtttcagtct ctcaaacttc taaattaacc agggctgaag ctgtgtttcc tgatgtggac 600
tatgtaaatt ctacagaggc tgagacaata ttggataata tcaccagag tacacagagc 660
ttcaatgact tcacaagagt ggtgggaggg gaagatgcc agcctggcca gtttccctgg 720
caggtagtcc tcaatgggaa ggtagatgcc ttctgtggtg gaagcattgt taatgaaaag 780
tggattgtga cagctgcccc ctgtgtggag actggtgtga agatcactgt ggtggctggt 840
gaacataata ttgaggaaac agagcacact gagcagaaaa gaaatgtgat aagaatcatc 900
cctcaccaca actacaatgc tgccatcaac aaatataatc atgacatagc actcctagaa 960
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gaaggaggaa gagatagctg ccaggagat tcaggaggcc cccatgtaac agaggtagag 1260
gggacctctt tcctcacagg gatcatttcc tggggagaag agtgtgcat gaagggaag 1320
tatggaatct acacaaaggt gtctaggtat gtgaactgga tcaaggagaa gaccaagctg 1380
acctga 1386

<210> 398

<211> 1194

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 398

atggaggaga agtgctcatt tgaagaggcc agggagggtt ttgagaacac agagaggacc 60
acagagttct ggaagcagta tgtggatggg gaccagtgtg agagcaaccc ctgccttaat 120
gggggcagct gtaaagatga tattaatagc tatgaatgct ggtgcccctt tggatttgag 180
gggaaaaact gtgaattgga tgttacttgc aacatcaaaa atggtagatg tgagcagttc 240

tgcaagaact ctgcagacaa taaagtggtc tgctcctgca ctgaagggtg cagactggca	300
gaaaaccaga agagtgtgta gccagctgtg cccttccctt gtggcagagt ttctgtgagc	360
cagaccagca aactcaccag agctgaggct gtctttccag atgtggacta tgtgaactcc	420
acagaagctg agactatcct ggacaacatt actcagagca cccagtcctt caatgacttc	480
acaagggtgg ttggaggaga agatgccaaag ccagggcagt ttccctggca ggtggtactg	540
aatggaaaag ttgatgcttt ctgtggaggg agcattgtga atgaaaaatg gattgtcact	600
gctgcccact gtgtggaaac tgggggtgaag atcactgtgg tggctgggga gcataatatt	660
gaagaaacag agcacactga acagaaaaga aatgtgatca ggatcatccc ccaccacaac	720
tacaatgcag ccatcaacaa atacaacat gacattgccc tgctggagct ggatgagccc	780
ctggtgctga acagctatgt gacccccatc tgtattgctg acaaggagta cacaaatatt	840
ttcctgaagt ttggctctgg ctatgtgagt ggctggggca gagtgttcca caagggaaga	900
tctgccctgg tgctgcagta cctgagggtg ccactggtgg acagggccac ctgcctgagg	960
agcacaaagt tcaccattta taacaacatg ttttgtgctg gcttccatga gggaggcagg	1020
gacagctgcc agggagattc tggaggggcc catgtgactg aggtggaggg cacctccttt	1080
ctgacaggca ttatcagctg gggagaggag tgtgccatga agggcaagta tggcatctac	1140
accaaggtgt ccagatatgt caactggatc aaggaaaaga ccaaactgac ctaa	1194

<210> 399

<211> 1416

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 399

atgaagcacc tgaacacagt gatggccgag agccctgctc tgatcaccat cttcctgctg	60
ggctacctgc tgtccacaga gtgcgccgtg tttctggatc gggagaacgc taccaagatc	120
ctgacacggc caaagaggta caacagcggc aagctggagg agttcgtgag gggaaacctg	180
gagagagagt gcatcgagga gaggtgttcc ttcgaggagg ccagagaggt gtttgagaac	240
accgagaaga ccacagagtt ttggaagcag tacgtggacg gcgatcagtg cgagagcaac	300
ccatgtctga acggcggaat ctgtaaggac gatatcagct cctacgagtg ctggtgtcaa	360
gtgggcttcg agggacggaa ctgcgagctg gacgccacct gtaacatcaa gaacggacgc	420
tgcaagcagt tttgtaagaa ctccccgac aacaaggtga tctgctcttg taccgagggc	480
taccagctgg ctgaggatca gaagtcttgc gagcctacag tgcccttccc ttgtggaagg	540
gccagcatct cctactctag caagaagatc accagagctg agacagtgtt cagcaacatg	600
gactacgaga actccaccga ggccgtgttt atccaggacg atatcaccga tggcgctatc	660
ctgaacaacg tgacagagtc ctctgagtcc ctgaacgact tcacaagagt ggtgggcgga	720
gagaacgcca agccaggcca gatcccctgg cagggtgatcc tgaacggaga gatcgaggcc	780

ttttgcggcg gagctatcat caacgagaag tggatcgtga ccgccgctca ctgtctgaag	840
ccaggcgata agatcgaggt ggtggctgga gagtacaaca tcgacaagaa ggaggatacc	900
gagcagagga gaaacgtgat caggacaatc cccaccacc agtacaacgc caccatcaac	960
aagtactctc acgacatcgc tctgctggag ctggataagc ctctgaccta gaacagctac	1020
gtgaccccaa tctgcgtggc caacagagag tacacaaaca tcttcctgaa gtttggctct	1080
ggatacgtga gcggctgggg aaaggtgttc aacaagggcc ggcaggcctc catcctgcag	1140
tacctgaggg tgccactggg ggacagggtt acctgtctgc gctctaccac attcacaatc	1200
tacaacaaca tgttttgcgc tggatacagg gaggaggagaa aggactcctg tgaggagat	1260
tctggaggac ctcacgtgac cgaggtggag ggcacctctt ttctgacagg catcatcagc	1320
tggggagagg agtgcgctat gaagggaag tacggaatct acacaaaggt gtcccgtac	1380
gtgaactgga tcaaggagaa gaccaagctg acatga	1416

<210> 400

<211> 1449

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 400

atgaagcacc tgaacacagt gatggccgag agccctgctc tgatcaccat cttcctgctg	60
ggctacctgc tgtccacaga gtgcgccgtg tttctggatc gggagaacgc taccaagatc	120
ctgacacggc caaagaggta caactctggc aagctggagg agttcgtgag gggaaacctg	180
gagagagagt gcatcgagga gaggtgtagc ttcgaggagg ccagagaggt gtttgagaac	240
accgagaaga ccacagagtt ttggaagcag tacgtggacg gcgatcagtg cgagagcaac	300
ccatgtctga acggcggaat ctgtaaggac gatatcagct cctacgagtg ctggtgtcaa	360
gtgggcttcg agggacggaa ctgcgagctg gacgccacct gtaacatcaa gaacggacgc	420
tgcaagcagt tttgtaagaa ctccccgac aacaaggtga tctgctcttg taccgagggc	480
taccagctgg ctgaggatca gaagtcctgc gaggctacag tgcccttccc ttgtggaagg	540
gccagcatct cctactctag caagaagatc accagagctg agacagtgtt cagcaacatg	600
gactacgaga actccaccga ggccgtgttt atccaggacg atatcaccga tggcgctatc	660
ctgaacaacg tgacagagtc ctctgagtct ctgaacgact tcacaagagt ggtgggcgga	720
gagaacgcca agccaggcca gatcccctgg cagggtgatcc tgaacggaga gatcgaggcc	780
ttttgcggcg gagctatcat caacgagaag tggatcgtga ccgccgctca ctgtctgaag	840
ccaggcgata agatcgaggt ggtggctgga gagtacaaca tcgacaagaa ggaggatacc	900
gagcagagga gaaacgtgat caggacaatc cccaccacc agtacaacgc caccatcaac	960
aagtactctc acgacatcgc tctgctggag ctggataagc ctctgaccta gaacagctac	1020

gtgaccccaa tctgcgtggc caacagagag tacacaaaca tcttcctgaa gtttggtctt	1080
ggatactga gcggtgggg aaaggtgttc aacaagggcc ggcaggcctc catcctgcag	1140
tacctgaggg tgccactggt ggacagggct acctgtctgc gctctaccac attcacaatc	1200
tacaacaaca tgttttgcgc tggatacagg gaggaggaa aggactcctg tgaggagat	1260
tctggaggac ctcacgtgac cgaggtggag ggcacctctt ttctgacagg catcatcagc	1320
tggggagagg agtgcgtat gaagggaag tacggaatct acacaaaagt gagccgtac	1380
gtgaactgga tcaaggagaa gaccaagctg acaggaggag gaggcagcca ccaccaccac	1440
caccactga	1449

<210> 401

<211> 1461

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 401

atgaagcacc tgaacacagt gatggccgag tctcctgctc tgatcaccat cttcctgctg	60
ggctacctgc tgagcacaga gtgcgccgtg tttctggatc gggagaacgc taccaagatc	120
ctgacacggc caaagaggta caactctggc aagctggagg agttcgtgag gggaaacctg	180
gagagagagt gcatcgagga gaggtgtagc ttcgaggagg ccagagaggt gtttgagaac	240
accgagaaga ccacagagtt ttggaagcag tacgtggacg gcgatcagtg cgagagcaac	300
ccatgtctga acggcggaat ctgtaaggac gatatcagct cctacgagtg ctggtgtcaa	360
gtgggcttcg agggacggaa ctgcgagctg gacgccacct gtaacatcaa gaacggacgc	420
tgcaagcagt tttgtaagaa cagccccgac aacaaggtga tctgctcctg taccgagggc	480
taccagctgg ctgaggatca gaagtcctgc gagcctacag tgcccttccc ttgtggaagg	540
gccagcatct cctactctag caagaagatc accagagctg agacagtgtt ctctaacatg	600
gactacgaga acagcaccga ggccgtgttt atccaggacg atatcaccga tggcgctatc	660
ctgaacaacg tgacagagtc ctctgagtct ctgaacgact tcacaagagt ggtgggcgga	720
gagaacgcca agccaggcca gatcccctgg cagggtgatcc tgaacggaga gatcgaggcc	780
ttttgcggcg gagctatcat caacgagaag tggatcgtga ccgccgtca ctgtctgaag	840
ccaggcgaca agatcgaggt ggtggctgga gagtacaaca tcgacaagaa ggaggatacc	900
gagcagagga gaaacgtgat caggacaatc cccaccacc agtacaacgc caccatcaac	960
aagtactccc acgacatcgc tctgctggag ctggataagc ctctgacatc gaactcttac	1020
gtgaccccaa tctgcgtggc caacagagag tacacaaaca tcttcctgaa gtttggtctt	1080
ggatactga gcggtgggg aaaggtgttc aacaagggcc ggcaggccag catcctgcag	1140
tacctgaggg tgccactggt ggatagggt acctgtctgc gctccaccac attcacaatc	1200
tacaacaaca tgttttgcgc tggatacagg gaggaggaa aggactcctg tgaggagat	1260

tctggaggac ctcacgtgac cgaggtggag ggcacctcct ttctgacagg catcatctct	1320
tggggagagg agtgcgctat gaagggaag tacggaatct acacaaaggt gtcccgtac	1380
gtgaactgga tcaaggagaa gaccaagctg acaggcggag gcggaagcga gcagaagctg	1440
atctccgagc aggacctgtg a	1461

<210> 402
 <211> 1416
 <212> DNA
 <213> Artificial Sequence

<220>
 <221> source
 <223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 402	
atgaagcacc tgaacacagt gatggccgag agccctgctc tgatcaccat cttcctgctg	60
ggctacctgc tgtccacaga gtgcgccgtg tttctggatc gggagaacgc taccaagatc	120
ctgacacggc caaagaggta caacagcggc aagctggagg agttcgtgag gggaaacctg	180
gagagagagt gcatcgagga gaggtgttcc ttcgaggagg ccagagaggt gtttgagaac	240
accgagaaga ccacagagtt ttggaagcag tacgtggacg gcgatcagtg cgagagcaac	300
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gtgggcttcg aggggaaggaa ctgcgagctg gacgccacct gtaacatcaa gaacggaaga	420
tgcaagcagt tttgtaagaa ctccccgcac aacaaggatga tctgctcttg taccgagggc	480
taccagctgg ctgaggatca gaagtcttgc gaggctacag tgcccttccc ttgtggacgg	540
gccagcatct cctactctag caagaagatc acccgcgctg agacagtgtt cagcaacatg	600
gactacgaga actccaccga ggccgtgttt atccaggacg atatcaccga tggcgctatc	660
ctgaacaacg tgacagagtc ctctgagtcc ctgaacgact tcacacgggt ggtgggcgga	720
gagaacgcca agccaggcca gatccccctgg cagggtgatcc tgaacggaga gatcgaggcc	780
ttttgcggcg gagctatcat caacgagaag tggatcgtga ccgccgctca ctgtctgaag	840
ccaggcgata agatcgagggt ggtggctgga gagtacaaca tcgacaagaa ggaggatacc	900
gagcagagga gaaacgtgat ccgcacaatc cccaccacc agtacaacgc caccatcaac	960
aagtactctc acgacatcgc tctgctggag ctggataagc ctctgacatc gaacagctac	1020
gtgaccccaa tctgcgtggc caaccgggag tacacaaaca tcttcctgaa gtttggctct	1080
ggatacgtga gcggctgggg aaaggtgttc aacaagggcc gccaggcctc catcctgcag	1140
tacctgaggg tgccactggg ggacagggct acctgtctgc tgtctaccac attcacaatc	1200
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<210> 403
<211> 1449
<212> DNA
<213> Artificial Sequence

<220>
<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 403
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<211> 4281
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<220>

<221> source
<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 404

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<210> 405

<211> 3245

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 405

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<210> 406

<211> 3245

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 406

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<210> 407

<211> 3245

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 407

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<210> 408

<211> 3245

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 408

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<210> 409

<211> 3245

<212> DNA

<213> Artificial Sequence

<220>

<221> source

<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

<400> 409

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<220>
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<223> /note="Description of Artificial Sequence: Synthetic polynucleotide"

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