



US 20060257912A1

(19) **United States**

(12) **Patent Application Publication**
Kaemmerer et al.

(10) **Pub. No.: US 2006/0257912 A1**

(43) **Pub. Date: Nov. 16, 2006**

(54) **METHODS AND SEQUENCES TO SUPPRESS
PRIMATE HUNTINGTON GENE
EXPRESSION**

Related U.S. Application Data

(60) Provisional application No. 60/678,729, filed on May 6, 2005.

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Publication Classification

(51) **Int. Cl.**
A61K 48/00 (2006.01)
C12Q 1/68 (2006.01)
C07H 21/04 (2006.01)
C12P 21/06 (2006.01)
C07K 14/705 (2006.01)
(52) **U.S. Cl.** **435/6**; 514/44; 435/69.1; 435/320.1;
435/362; 530/350; 536/23.5

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(57) **ABSTRACT**

Disclosed herein are sequences, molecules and methods used to suppress the expression of HD genes encoding for huntingtin protein in primates including *Macaca mulatta* and *Homo sapiens*. These sequences, molecules and methods aid in the study of the pathogenesis of HD and can also provide a treatment for this disease.

(21) Appl. No.: **11/429,491**

(22) Filed: **May 4, 2006**

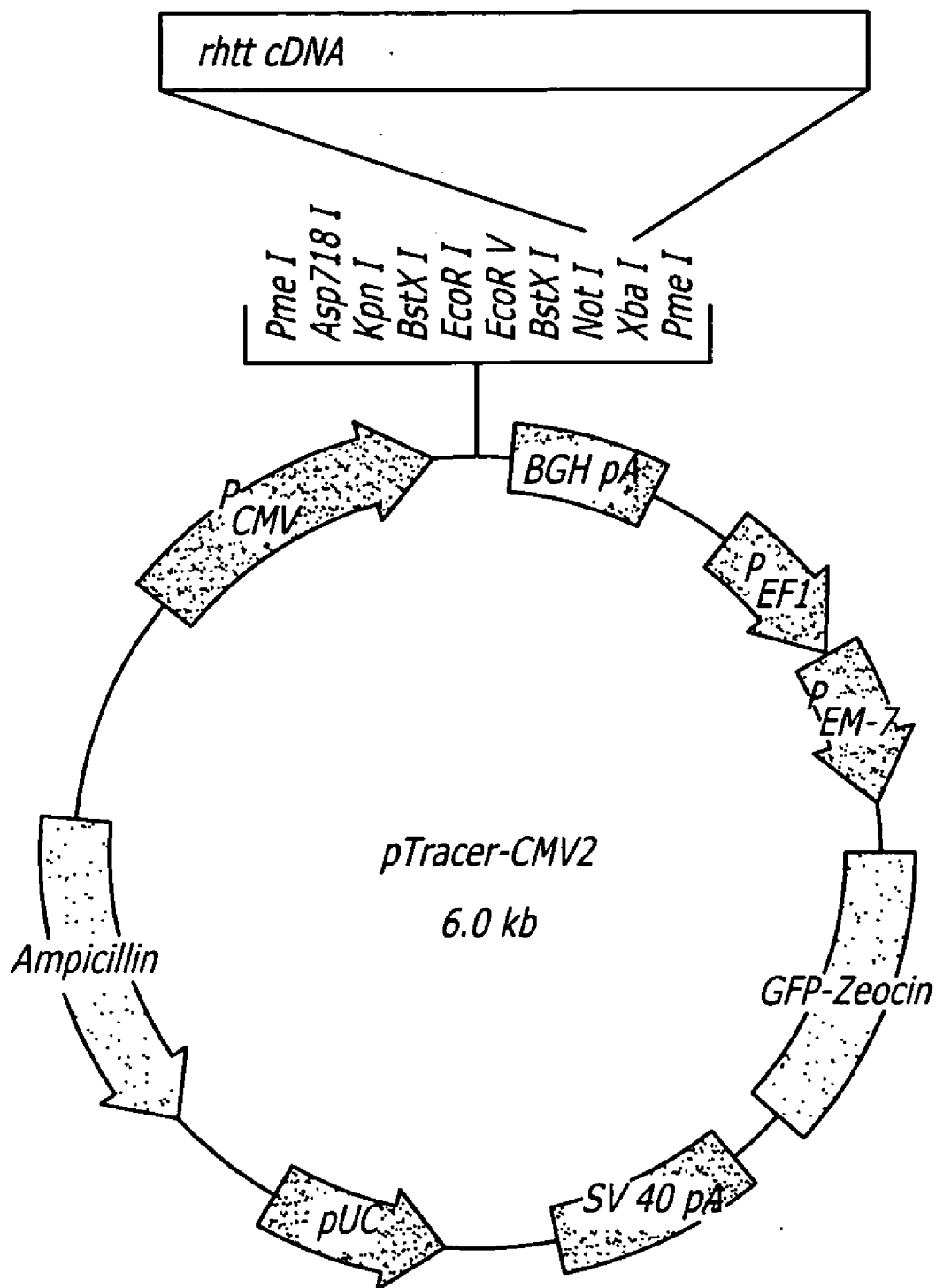


FIG. 1

FIG. 2

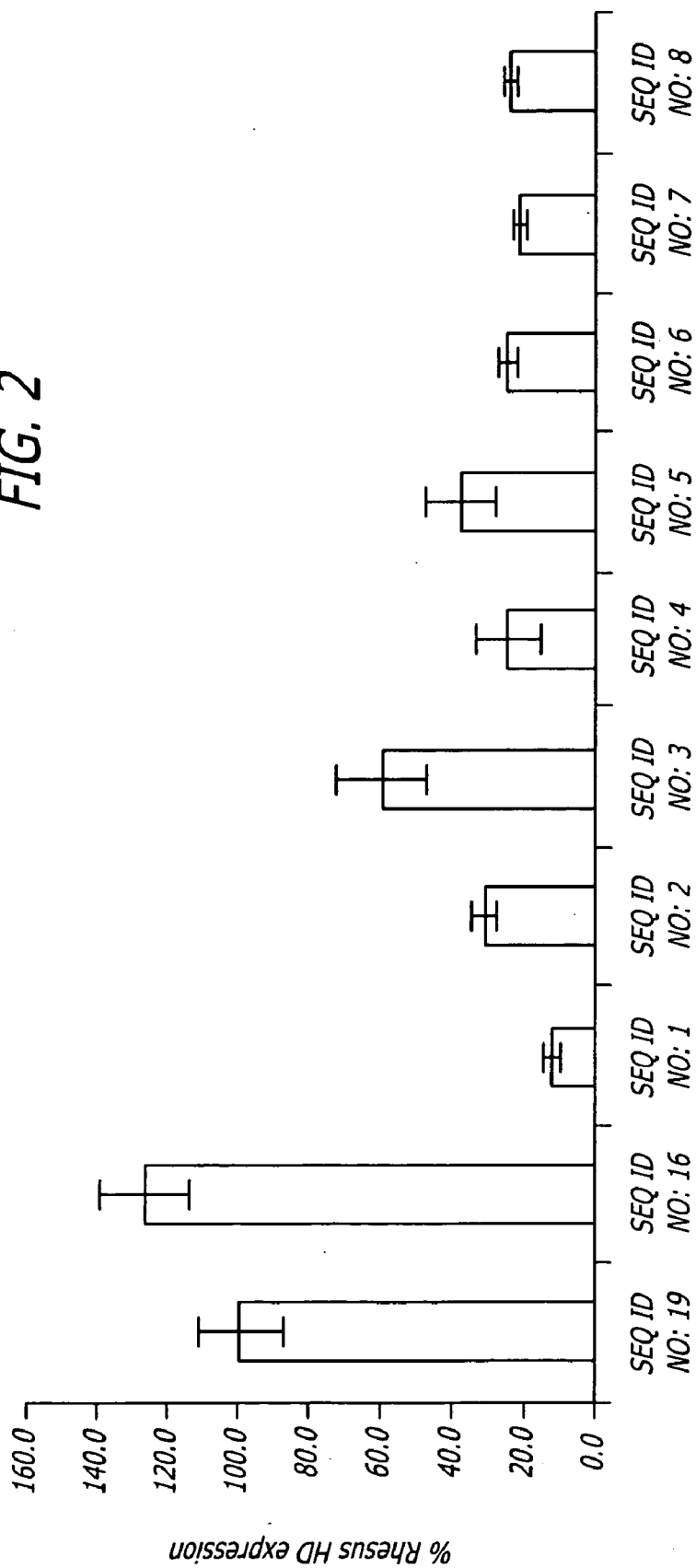


FIG. 3

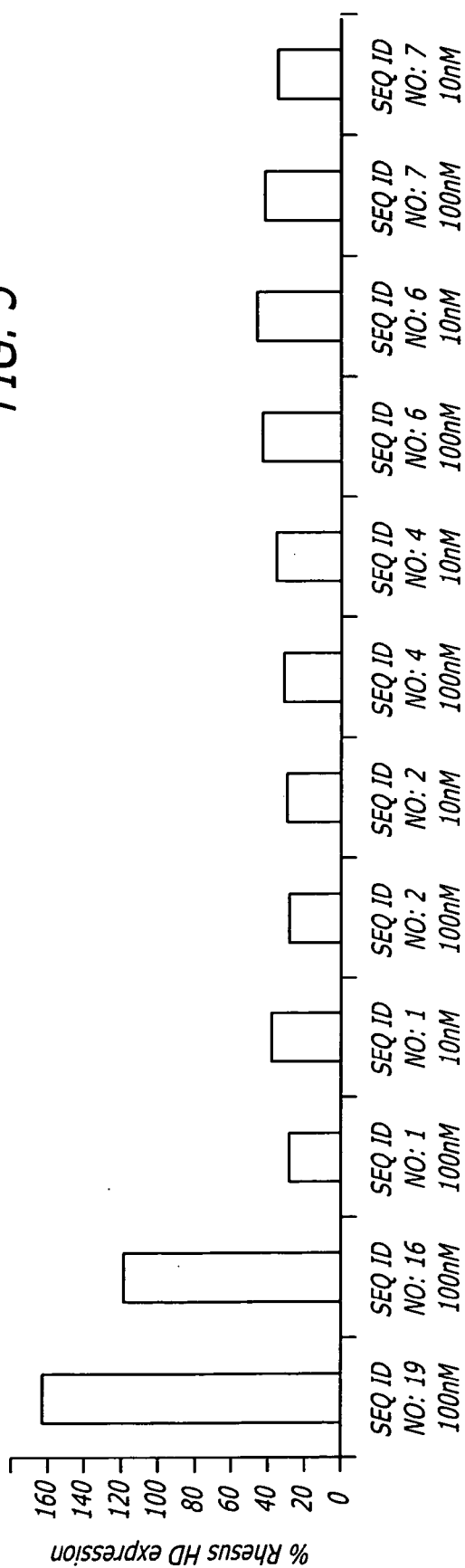


FIG. 4

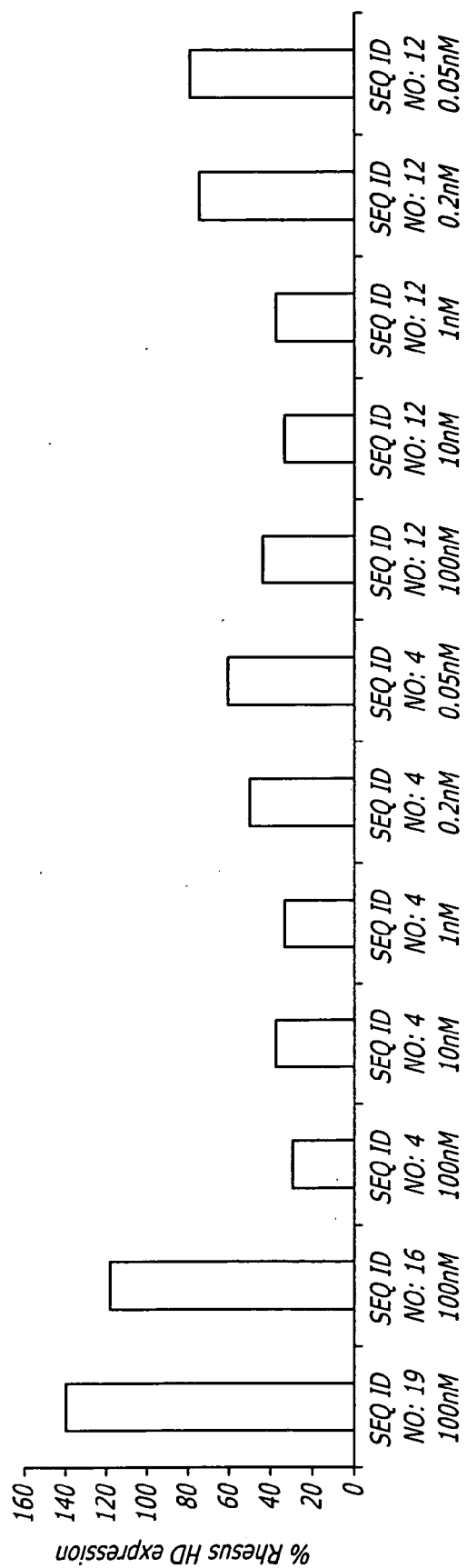


FIG. 5



FIG. 6

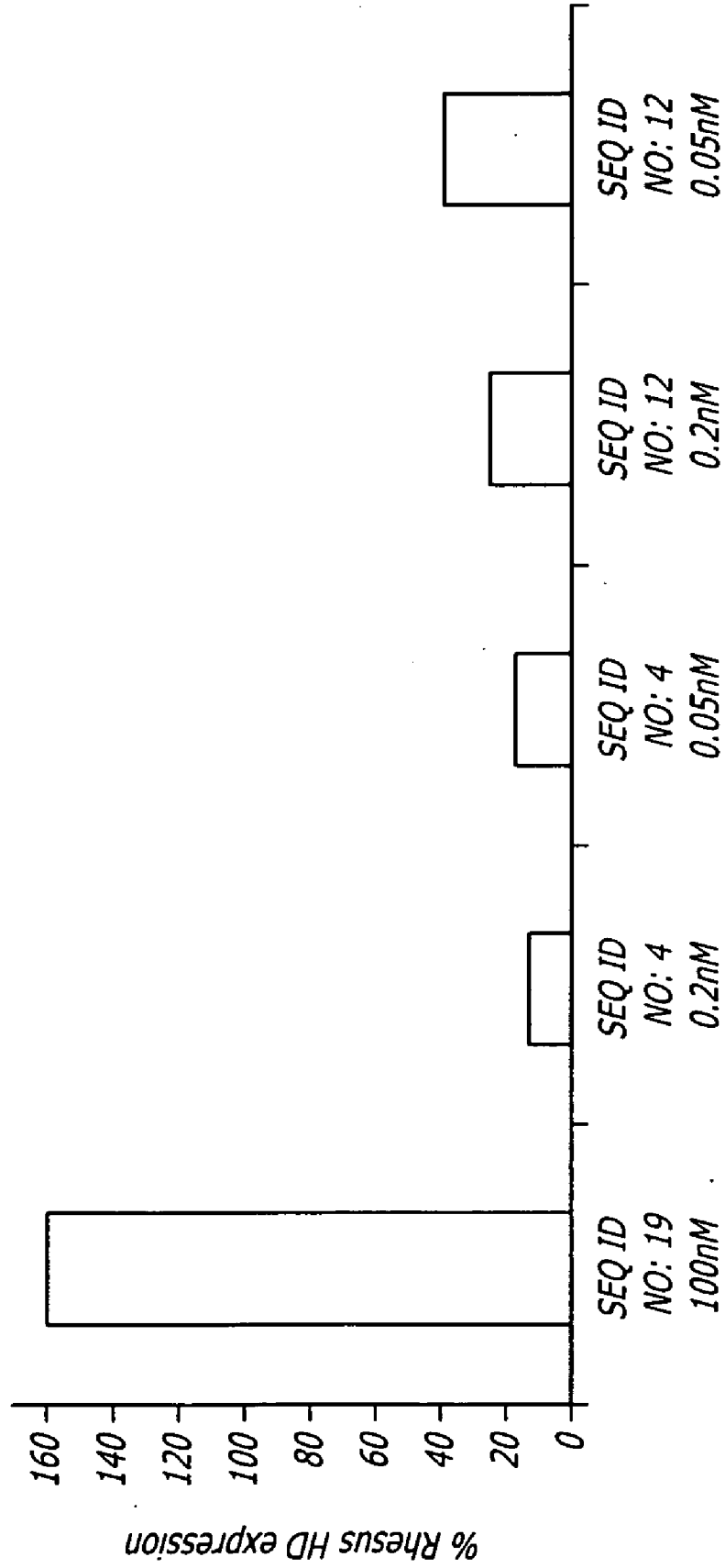
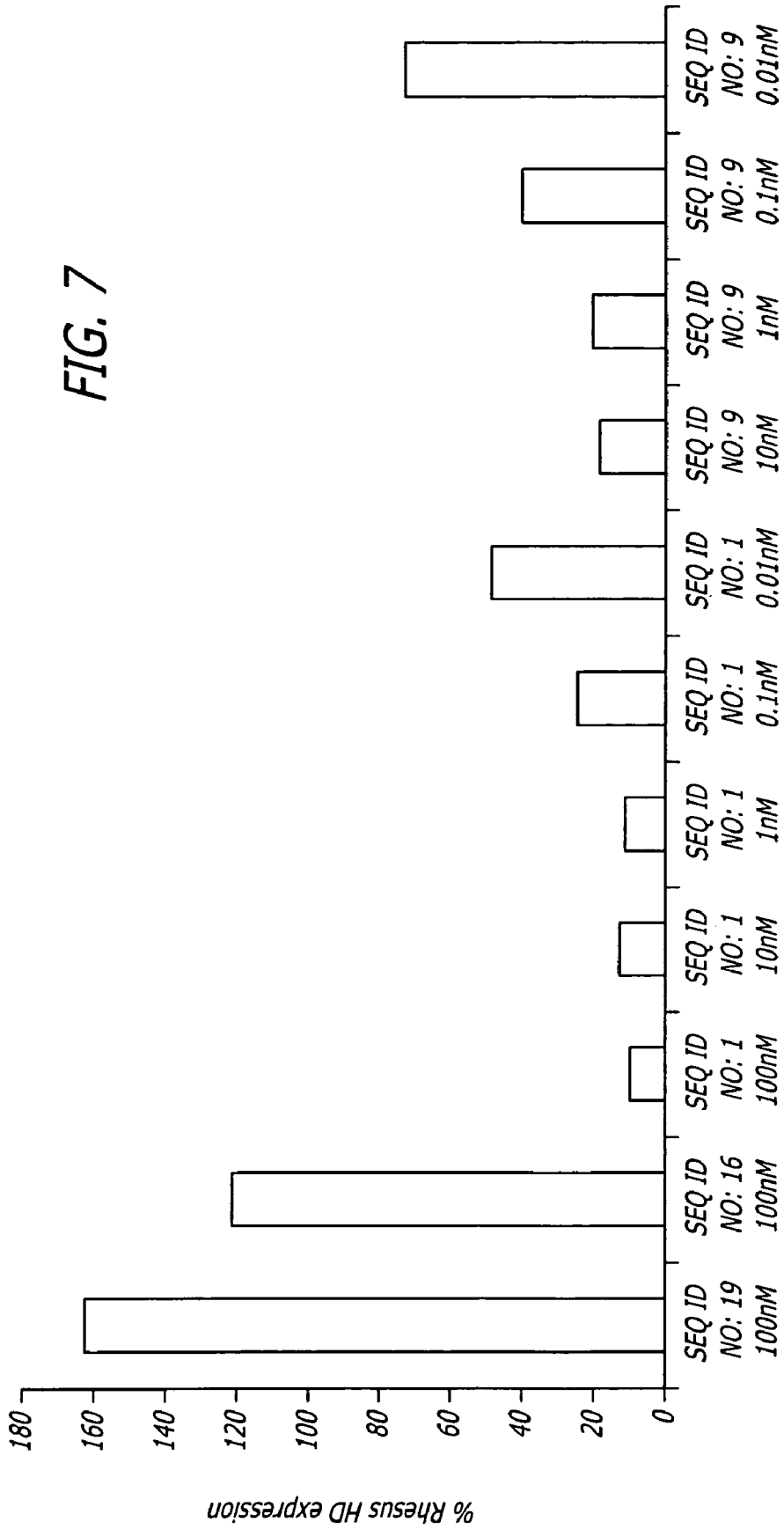


FIG. 7



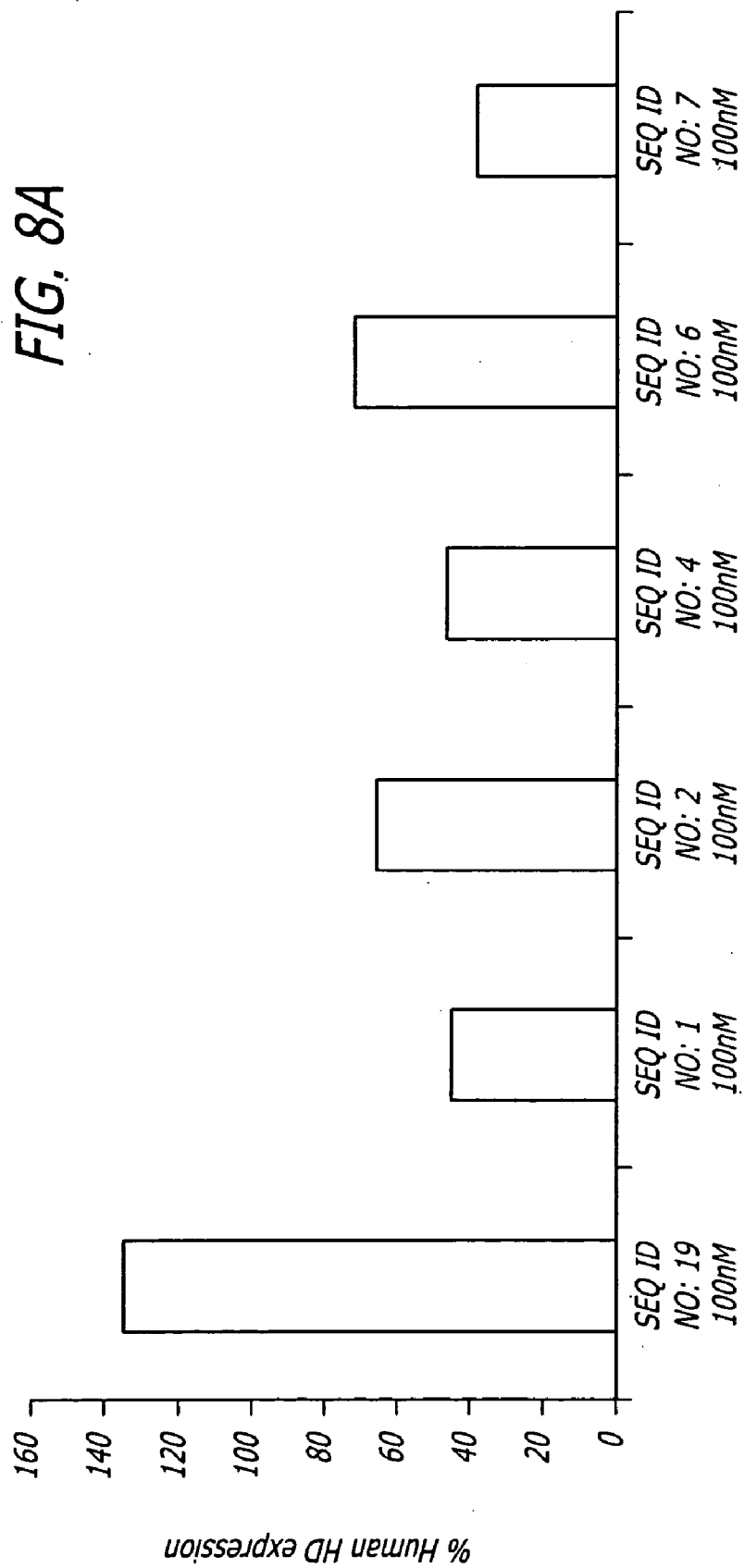


FIG. 8B



FIG. 8C

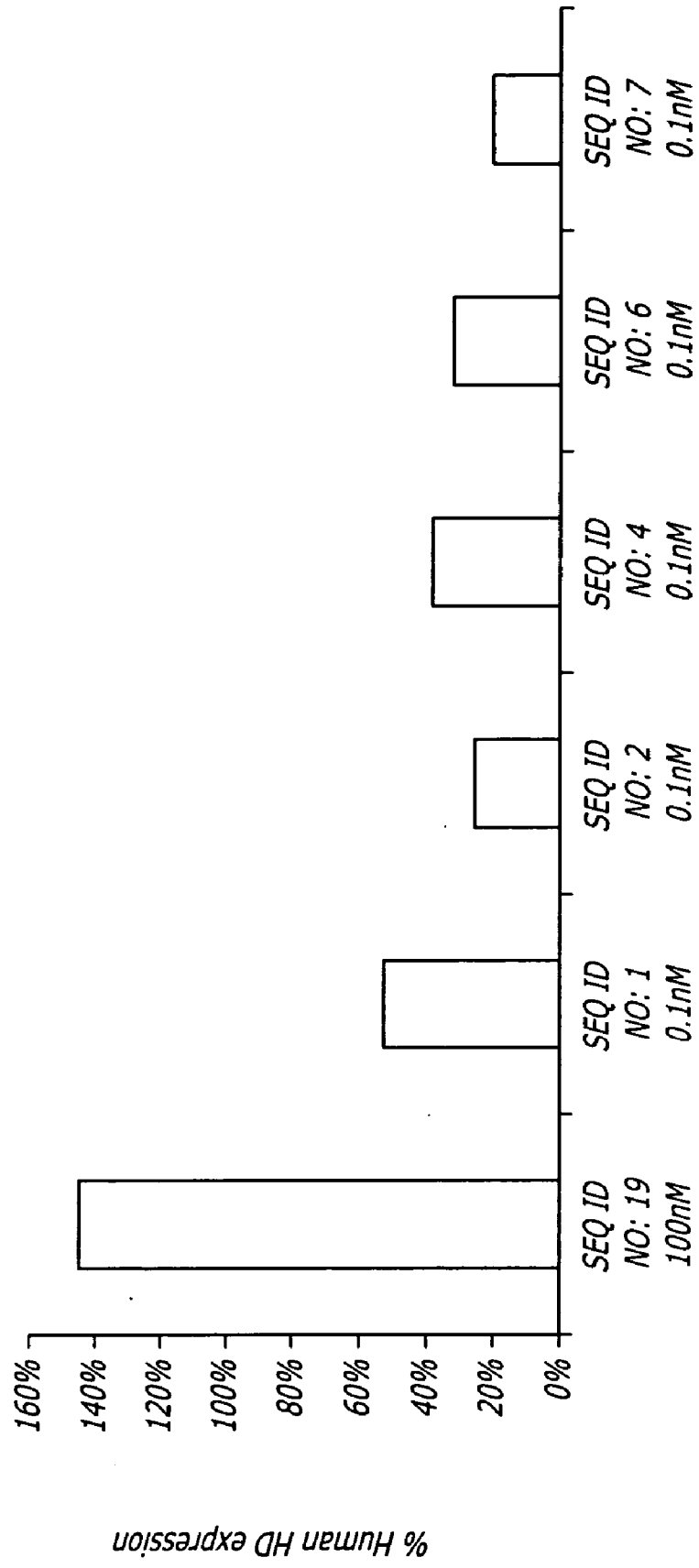


FIG. 8D

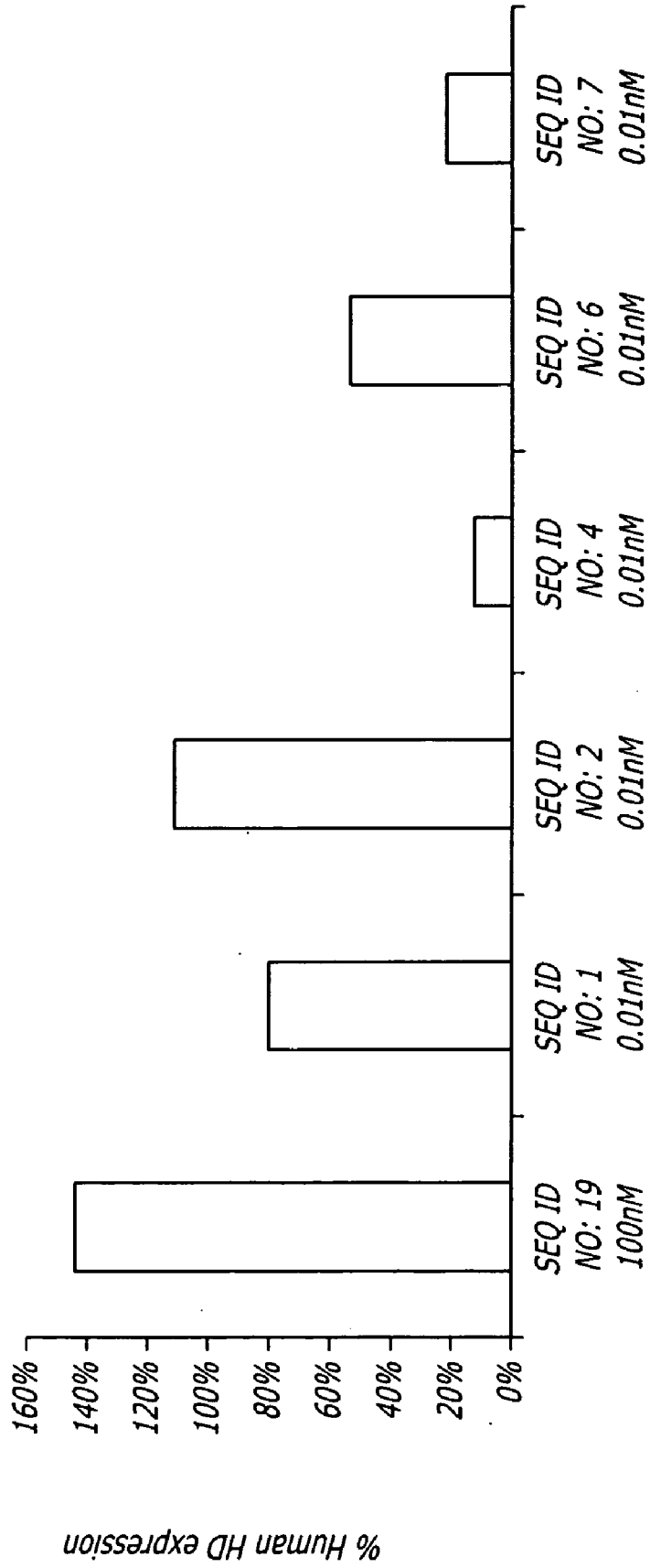


FIG. 9

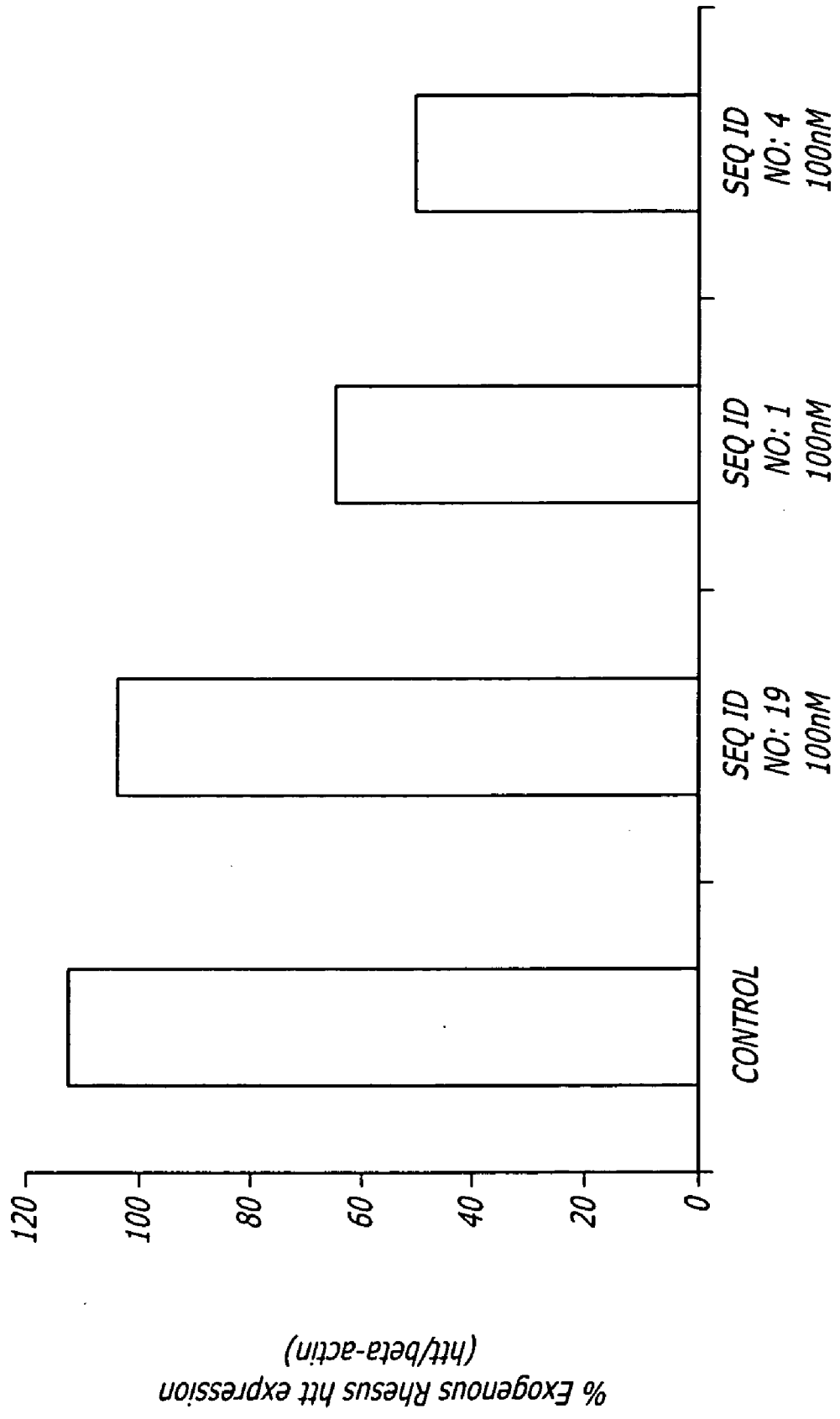


FIG. 10

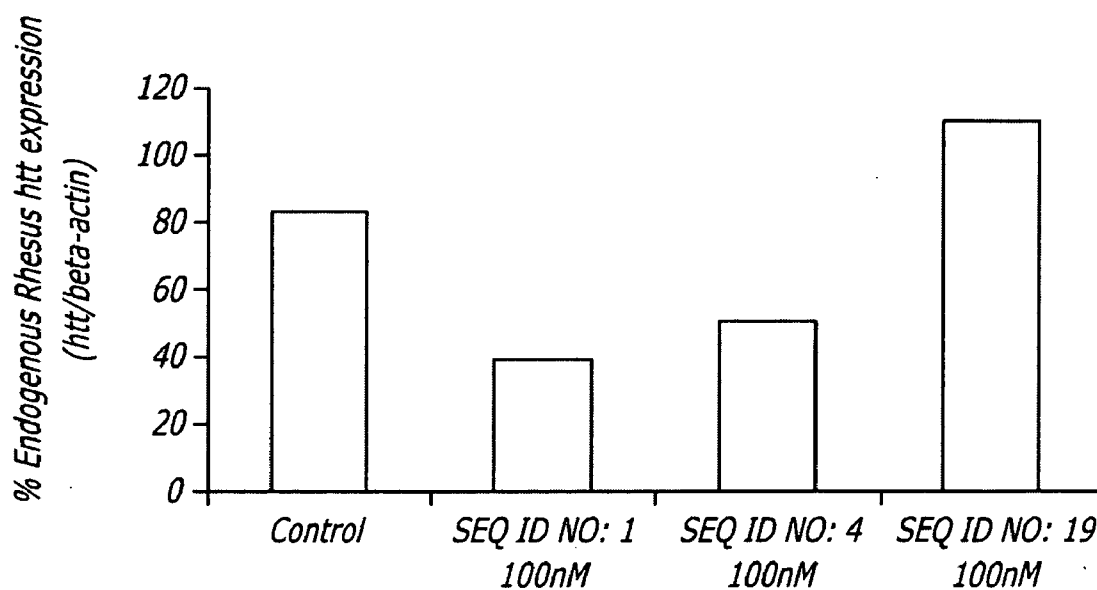
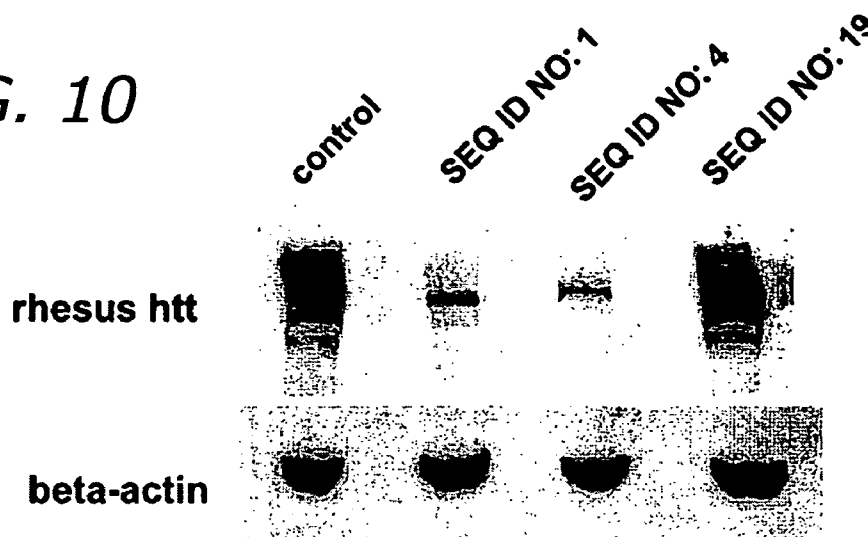


FIG. 11

FIG. 12A

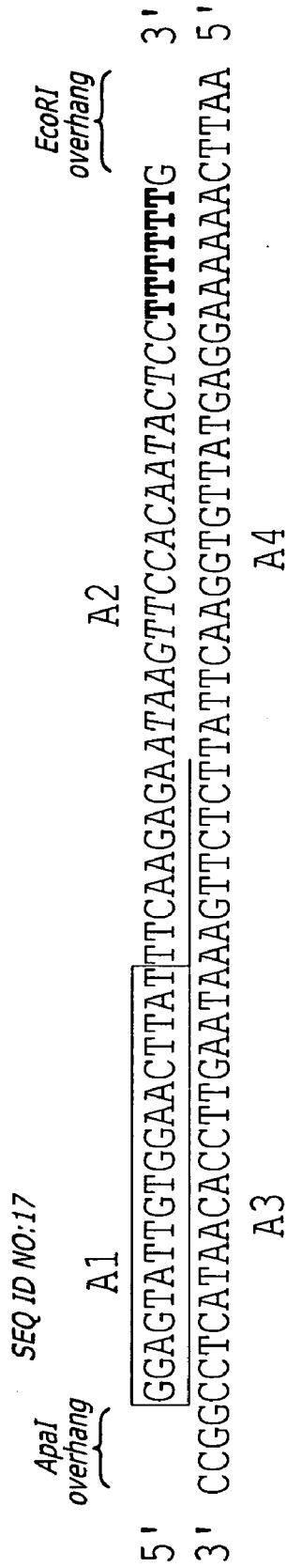


FIG. 12B

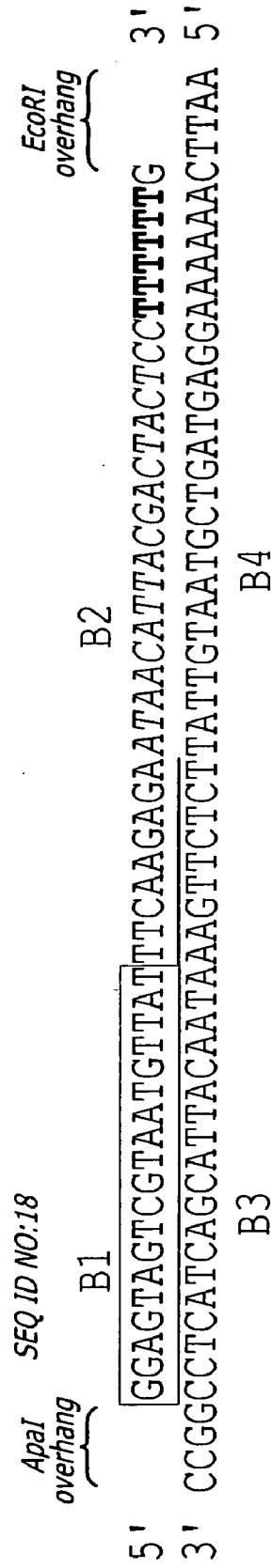


FIG. 13A

SEQ ID NO:20

5' ACCGGGAGTAGTCGTAATGTTATGCGAAGTGTTCAGAGAGACACTTCGCATAACATTACGACTACTCC**TTTTTTC** 3'
 3' CCTCATCAGCATTACAATACCGCTTCACAAGTTCCTCTGTGAAGCGTATTGTAATGCTGATGAGGAAAAAAGACCGT 5'

FIG. 13B

SEQ ID NO:21

5' ACCGGGAGTAGTCGTAATGTTAT**CTTCCTGTCAATAACATTACGACTACTCCTTTTTTC** 3'
 3' CCTCATCAGCATTACAATAGAAGGACAGTATTGTAATGCTGATGAGGAAAAAAGACCGT 5'

FIG. 13C

SEQ ID NO:22

5' ACCGGGAGTATTGTGGAAC**TTATTTC**AGAGAAATAAGTTCACAAATACTCC**TTTTTTC** 3'
 3' CCTCATAACACCC**TTGAATAAAGTTCCTT**ATTCAAGGTGTTATGAGGAAAAAAGACCGT 5'

FIG. 13D

SEQ ID NO:23

5' ACCGGGAGTATTGTGGAACTTATCTTCCCTGTCAATAAGTTCACACAATACTCCTTTTTTTC 3'
 3' CCTCATAACACACCTTGAATAGAAGGACAGTTATTCAAGGTGTTATGAGGAAAAAAGACCGT 5'

FIG. 13E

SEQ ID NO:24

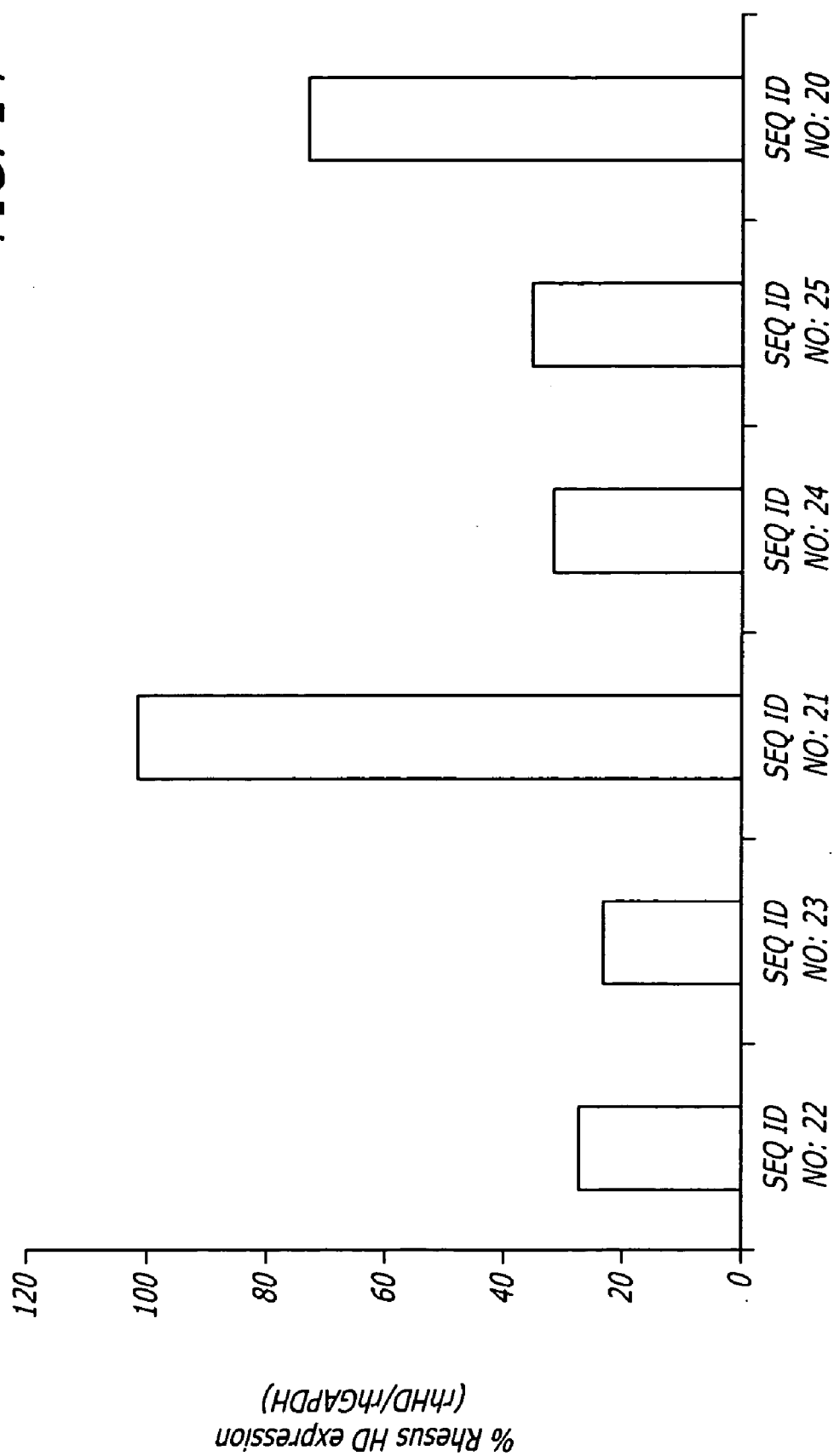
5' ACCGGGAGTATTGTGGAACTTATAGCTGGAGTTCAAGAGACTCCAGCTATAAGTTCACACAATACTCCTTTTTTTC 3'
 3' CCTCATAACACACCTTGAATATCGACCTCAAGTTCCTGAGGTCGATATTC AAGGTGTTATGAGGAAAAAAGACCGT 5'

FIG. 13F

SEQ ID NO:25

5' ACCGGGAGTATTGTGGAACTTATAGCTGGAGCTTCCCTGTCACTCCAGCTATAAGTTCACACAATACTCCTTTTTTTC 3'
 3' CCTCATAACACACCTTGAATATCGACCTCGAAGGACAGTGAGGTCGATATTC AAGGTGTTATGAGGAAAAAAGACCGT 5'

FIG. 14



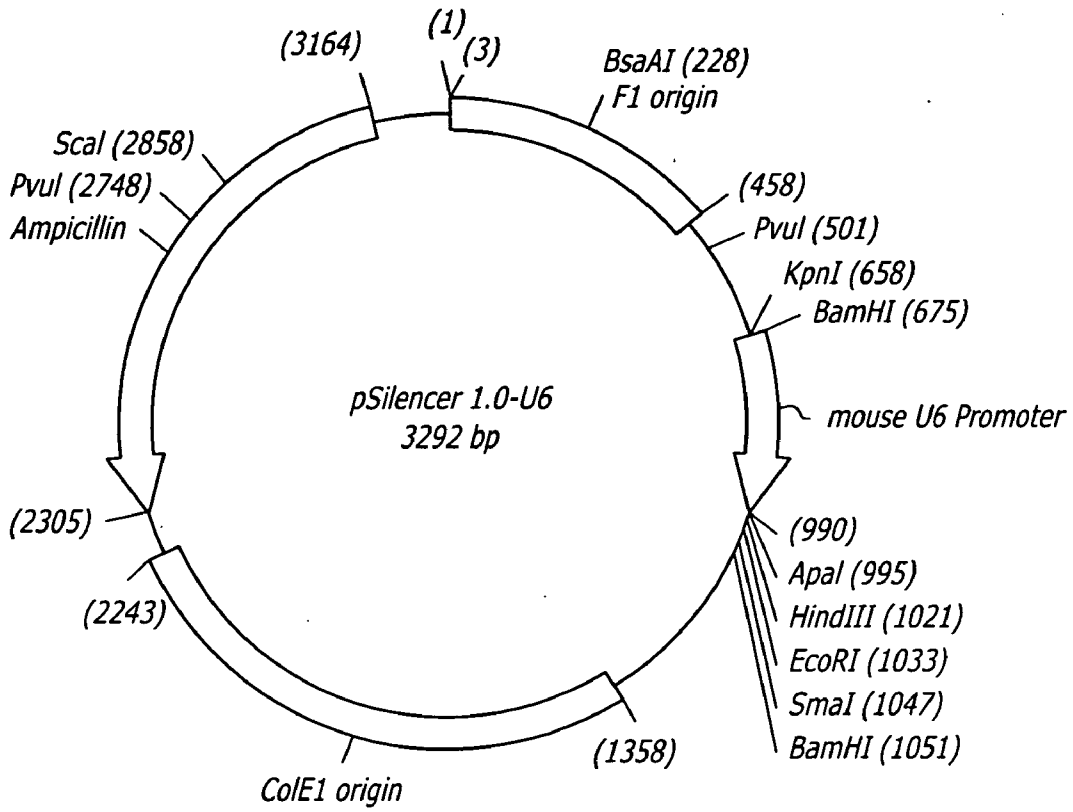


FIG. 15

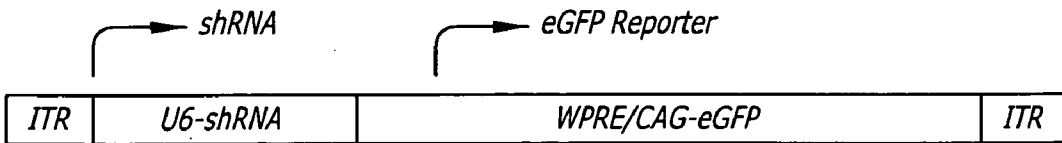


FIG. 16

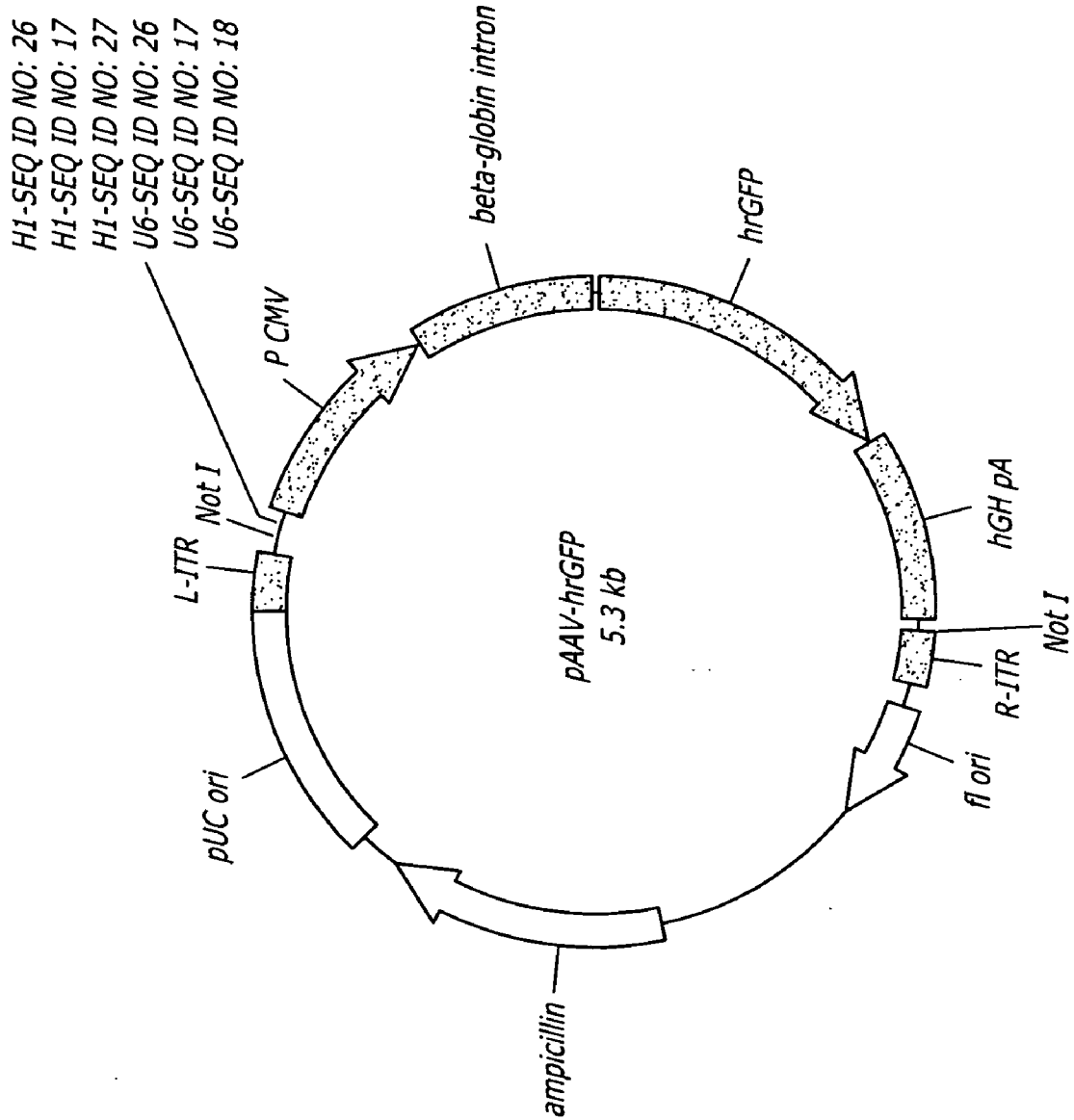


FIG. 17

FIG. 18A

SEQ ID NO: 26

5' ACCG**TGACAGCAGTGTGATAAA**TTCAAGAGATTTATCAACACTGCTGTCA**TTTTTTC** 3'
3' ACTGTCGCACAAC**TATTTAAGTTCTCTAAATAGTTGTGACGACAGTAAAAAAGACCGT** 5'

FIG. 18B

SEQ ID NO: 27

5' ACCG**TGACGAAGTCGTGATAAA**TTCAAGAGATTTAATCACGACTTCGTCAT**TTTTTTC** 3'
3' ACTGCTTCAGC**ACTAATTTAAGTTCTCTAAATTAGTCTGAAGCAGTAAAAAAGACCGT** 5'

FIG. 19

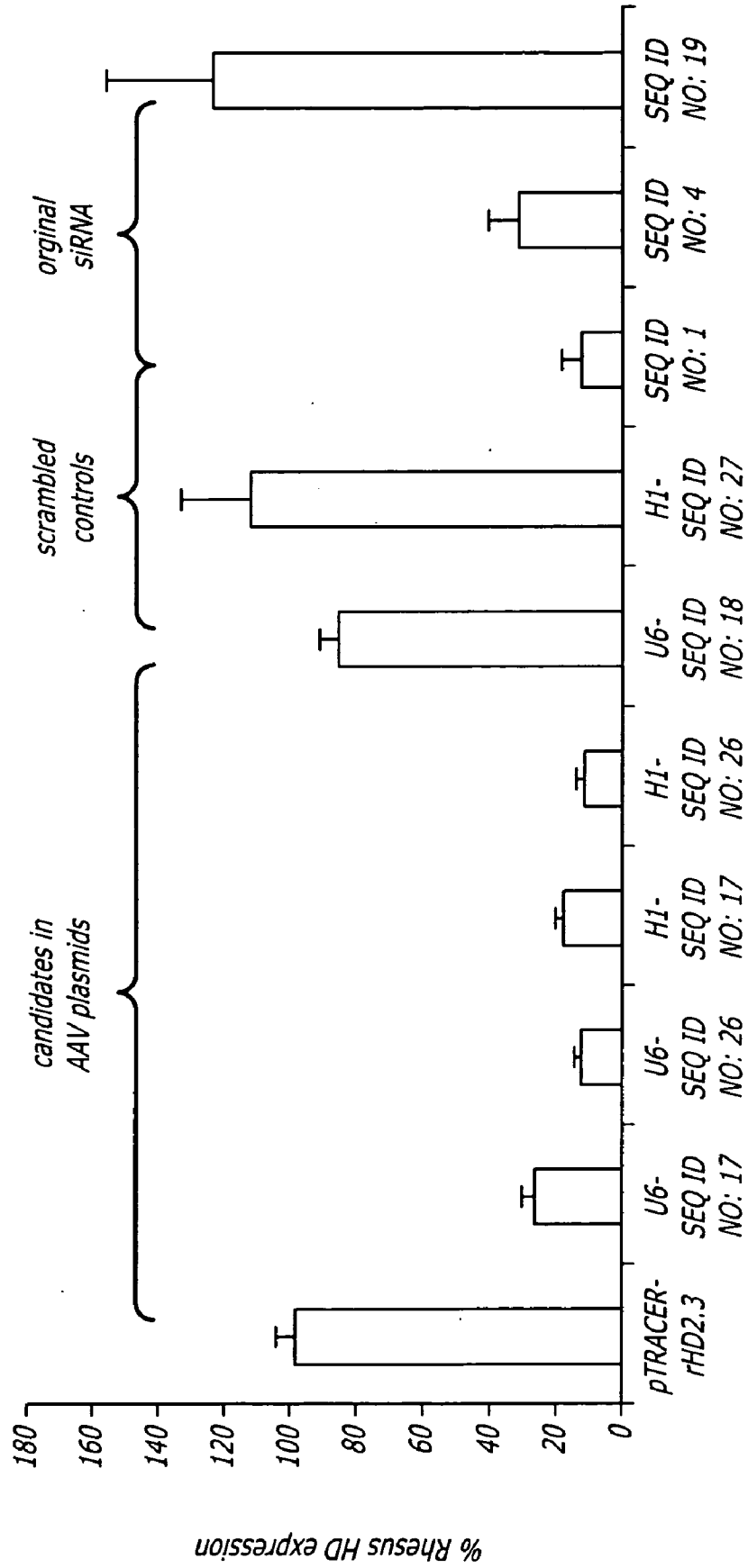


FIG. 20

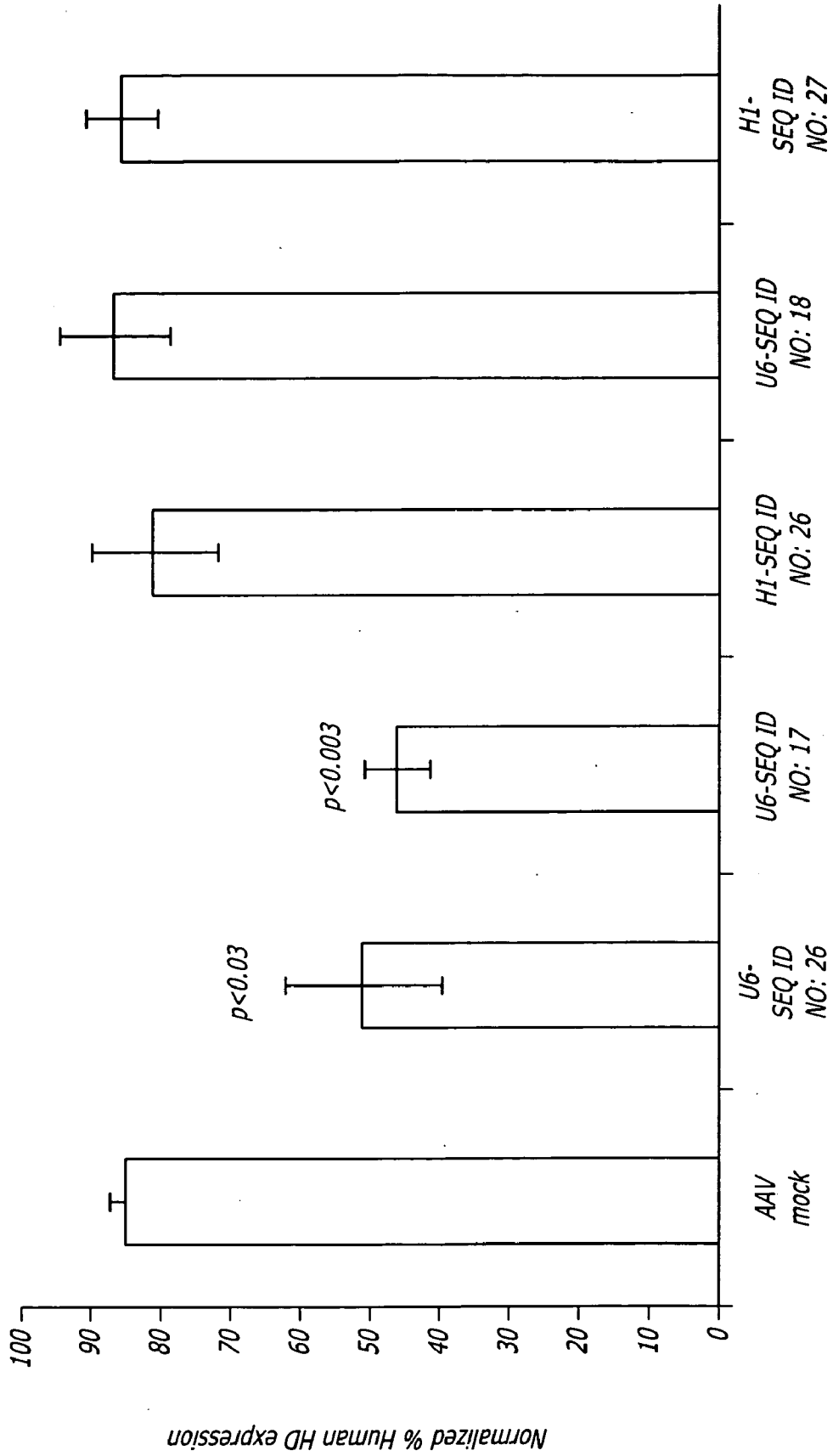


FIG. 21

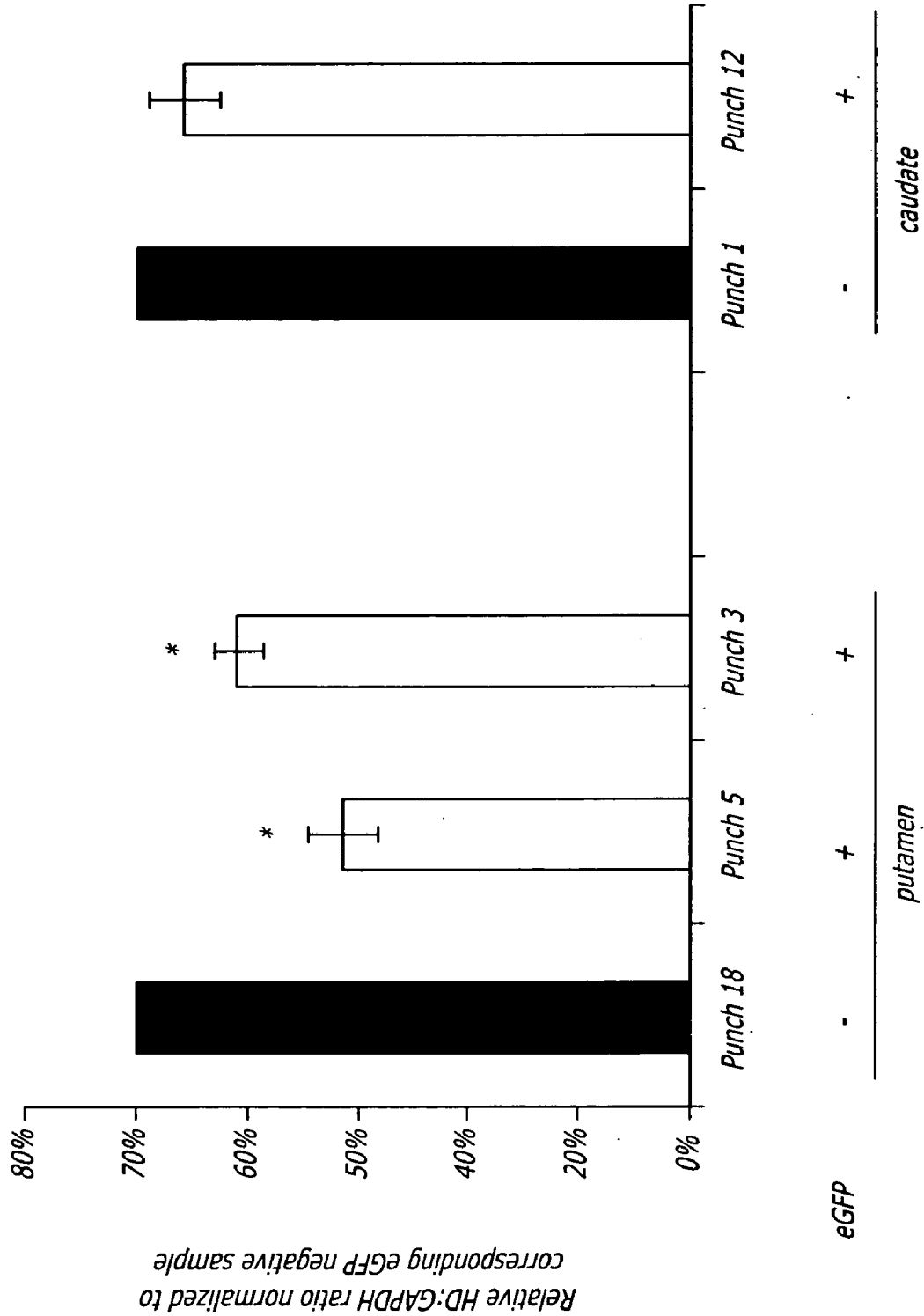
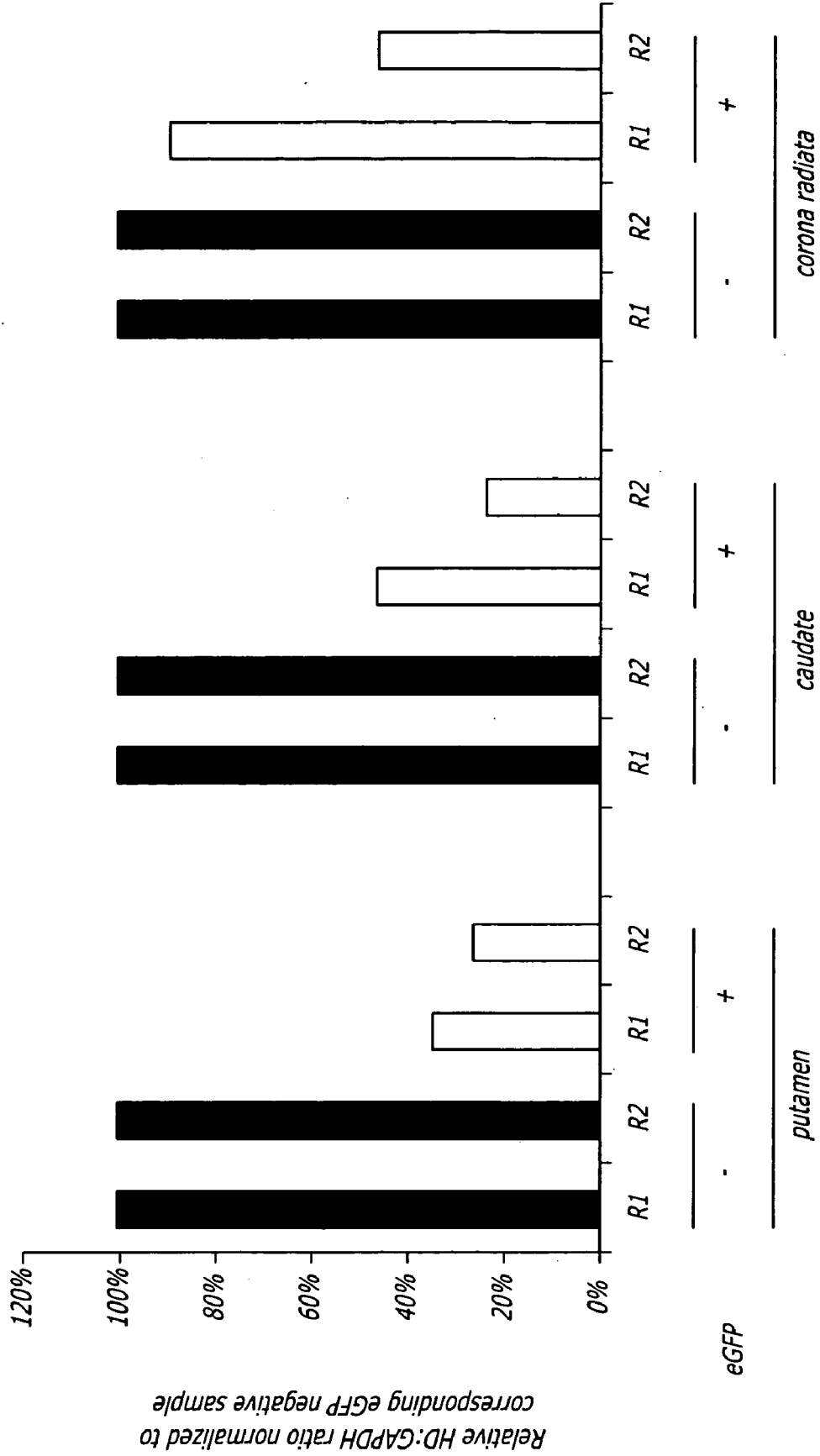


FIG. 22



METHODS AND SEQUENCES TO SUPPRESS PRIMATE HUNTINGTON GENE EXPRESSION

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority under 35 U.S.C. §119(e) to U.S. Provisional Patent Application No. 60/678,729 filed May 6, 2005.

FIELD OF THE INVENTION

[0002] The present invention relates to inhibitory nucleic acid molecules that suppress the expression of the Huntington's disease gene in primates, including rhesus monkeys (*Macaca mulatta*) and humans (*Homo sapiens*).

BACKGROUND OF THE INVENTION

[0003] Huntington's disease ("HD") is a neurodegenerative brain disorder with a juvenile or adult onset. It slowly destroys an affected individual's ability to walk, think, talk and reason. Symptoms include changes in cognitive ability, such as impaired short-term memory and a decreased ability to concentrate; changes in mood, such as the development of mood swings, depression and irritability; and changes in coordination and physical movement such as clumsiness, involuntary movements and twitching. These symptoms gradually worsen until HD patients die, approximately 15-20 years after the onset of the disease.

[0004] While the biochemical cause of HD is not yet fully understood, it is now known that HD is inherited as an autosomal dominant trait. This inheritance feature means that every individual who inherits a mutated (expanded) HD gene from either parent will develop the disease.

[0005] One breakthrough in research regarding HD has been the identification of the mutated gene that causes HD. Based on this breakthrough, researchers and physicians now can predict which individuals will develop HD. Specifically, researchers and physicians can predict which individuals will develop HD by counting the number of "CAG repeats" that exist within a given individual's HD gene. If a person has 35 or fewer CAG repeats in both of their HD genes, that person will not develop HD. If a person has more than 35 CAG repeats in either of their HD genes, that person will develop the disease. The more CAG repeats a person has over 35, the earlier the person will develop the symptoms of HD.

[0006] The HD gene encodes for a protein called "huntingtin" (also known as "htt"). The exact function of huntingtin is not known. The expression of a mutant, expanded huntingtin protein is known to be the cause of HD, however. Some of the evidence that has led scientists to this conclusion includes mouse studies showing that the introduction of an expanded HD transgene in the mouse leads to the pathological and behavioral features of HD and its removal can resolve these effects. Thus, suppressing the production of huntingtin in brain cells may prevent or alleviate the symptoms or occurrence of HD.

[0007] Recent developments in genetic technologies have made the selective suppression of certain proteins, such as huntingtin, possible. Some background in the art is required to understand the potential impact of these technologies. Generally, for a protein to exert an effect, the cell that will

use the protein must create it. To create a protein the cell first makes a copy of the protein's gene sequence in the nucleus of the cell. This copy of the gene sequence that encodes for the protein (called messenger RNA ("mRNA")) leaves the nucleus and is trafficked to a region of the cell containing ribosomes. Ribosomes read the sequence of the mRNA and create the protein for which it encodes. This process of new protein synthesis is known as translation. A variety of factors affect the rate and efficiency of protein translation. Among the most significant of these factors is the intrinsic stability of the mRNA itself. If the mRNA is degraded quickly within the cell (such as before it reaches a ribosome), it is unable to serve as a template for new protein translation, thus reducing the cell's ability to create the protein for which it encoded.

[0008] Based on the foregoing, the technology of RNA interference ("RNAi") has emerged. RNA interference is, in fact, a naturally-occurring mechanism for suppressing gene expression and subsequent protein translation. RNA interference suppresses protein translation by either degrading the mRNA before it can be translated or by binding the mRNA and directly preventing its translation. This naturally-occurring mechanism of RNA interference can also be artificially induced to occur in cells. For example, RNA interference can be achieved by introducing into cells short, double-stranded nucleic acid oligonucleotides corresponding to the mRNA for the gene to be suppressed, or by introducing into cells a sequence of DNA that encodes for a short, hairpin transcript of nucleic acids that folds back upon itself and forms a short, double-stranded nucleic acid oligonucleotide following further processing in the cell. This technology provides a means to suppress the expression of huntingtin in cells. The suppression of huntingtin in cells can be useful in the study of HD pathogenesis. Suppressing huntingtin in a patient also could prevent or alleviate the symptoms of HD.

SUMMARY OF THE INVENTION

[0009] The present invention describes methods, nucleic acid sequences and molecules, expression cassettes, and vectors for using RNA interference ("RNAi") to suppress expression of the HD gene. Suppressing expression of the HD gene can reduce levels of huntingtin within cells. This suppression and reduction can be useful in the study of HD pathogenesis. This suppression and reduction also can be useful in the prevention and treatment of the symptoms of HD. Specifically, RNAi is mediated by double stranded RNA ("dsRNA"), short hairpin RNA ("shRNA") or other nucleic acid molecules with similar characteristics. These nucleic acid molecules are processed or cut into smaller pieces by cellular enzymes including Dicer and Drosha. The smaller fragments of the nucleic acid molecules can then be taken up by a protein complex called the RNA-induced silencing complex ("RISC complex") that mediates degradation of mRNAs. The RISC complex will degrade mRNA that complementarily base pairs with the nucleic acid molecules it has taken up. In this manner, the mRNA is specifically destroyed, thus preventing the protein for which the mRNA encoded from being made.

[0010] The understanding of the mechanism of RNAi now allows geneticists to create nucleic acid molecules with sequences that are homologous to known gene sequences in order to suppress the expression or formation of certain

is complementary to at least 23 contiguous nucleotides of the first strand. In another embodiment of the present invention, the invention includes an isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 27 contiguous nucleotides encoded by SEQ ID NO: 11 and wherein the second strand is complementary to at least 23 contiguous nucleotides of the first strand. In another embodiment of the present invention, the invention includes an isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 27 contiguous nucleotides encoded by SEQ ID NO: 12 and wherein the second strand is complementary to at least 23 contiguous nucleotides of the first strand. In another embodiment of the present invention, the invention includes an isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 27 contiguous nucleotides encoded by SEQ ID NO: 13 and wherein the second strand is complementary to at least 23 contiguous nucleotides of the first strand. In another embodiment of the present invention, the invention includes an isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 27 contiguous nucleotides encoded by SEQ ID NO: 14 and wherein the second strand is complementary to at least 23 contiguous nucleotides of the first strand. In another embodiment of the present invention, the invention includes an isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 27 contiguous nucleotides encoded by SEQ ID NO: 15 and wherein the second strand is complementary to at least 23 contiguous nucleotides of the first strand.

[0014] In one embodiment of the present invention, the nucleic acid duplex is between 19 and 30 base pairs in length.

[0015] In another embodiment of the present invention, the first and/or second strand of the nucleic acid duplex comprises an overhang region. In another embodiment of the present invention, the first and/or second strand of the nucleic acid duplex comprises a 3' overhang region, a 5' overhang region, or both 3' and 5' overhang regions. In another embodiment of the present invention, the first and/or second strand of the nucleic acid duplex comprises an overhang region that is from approximately 1 to approximately 10 nucleotides in length.

[0016] In another embodiment of the present invention, the first and second strand of the nucleic acid duplex are operably linked by means of a nucleic acid loop strand that forms a hairpin structure comprising a duplex structure and a loop structure. In another embodiment of the present invention, the first and second strand of the nucleic acid duplex are operably linked by means of a nucleic acid loop that contains from 4 to 13 nucleotides.

[0017] In another embodiment of the present invention, the invention includes an expression cassette. In one embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 1. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 2. In another embodiment, the expression cassette comprises a nucleic acid sequence

encoding SEQ ID NO: 3. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 4. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 5. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 6. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 7. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 8. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 9. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 10. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 11. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 12. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 13. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 14. In another embodiment, the expression cassette comprises a nucleic acid sequence encoding SEQ ID NO: 15.

[0018] In one embodiment of the present invention, the expression cassette also comprises a promoter. In another embodiment of the present invention, the expression cassette comprises a regulatable promoter. In another embodiment of the present invention, the expression cassette comprises a constitutive promoter. In another embodiment of the present invention, the expression cassette comprises a promoter that is a cytomegalovirus ("CMV") promoter. In another embodiment of the present invention, the expression cassette comprises a promoter that is a Rous sarcoma virus ("RSV") promoter. In another embodiment of the present invention, the expression cassette comprises a promoter utilized by RNA polymerase II. In another embodiment of the present invention, the expression cassette comprises a promoter utilized by RNA polymerase III.

[0019] In one embodiment of the present invention, the expression cassette comprises a polyadenylation signal. In another embodiment of the present invention, the expression cassette comprises a polyadenylation signal that is a synthetic minimal polyadenylation signal. In another embodiment of the present invention, the expression cassette comprises a marker gene.

[0020] In one embodiment of the present invention, the invention includes a vector comprising one or more of the expression cassettes previously described. In another embodiment of the present invention, the vector comprises a first and a second expression cassette.

[0021] In one embodiment, the first expression cassette of the vector encodes a nucleotide sequence encoding for SEQ ID NO. 1 and the second expression cassette of the vector encodes for a nucleotide sequence that is complementary to at least 15 contiguous nucleotides of SEQ ID. NO. 1. In another embodiment, the first expression cassette of the vector encodes a nucleotide sequence encoding for SEQ ID NO. 2 and the second expression cassette of the vector encodes for a nucleotide sequence that is complementary to at least 15 contiguous nucleotides of SEQ ID. NO. 2. In

includes a method of suppressing the accumulation of huntingtin in a cell comprising introducing a previously-described nucleic acid duplex into the cell in an amount sufficient to suppress accumulation of huntingtin in the cell. In another method of the present invention, accumulation of huntingtin is suppressed by at least 10%.

[0027] Another method of the present invention includes a method of preventing cytotoxic effects of mutant huntingtin in a cell comprising introducing a previously-described nucleic acid duplex into the cell in an amount sufficient to suppress accumulation of the mutant huntingtin so that the nucleic acid duplex prevents cytotoxic effects of mutant huntingtin in the cell.

[0028] Another method of the present invention includes a method of inhibiting expression of a huntingtin gene in a cell comprising introducing a previously-described nucleic acid duplex into the cell in an amount sufficient to inhibit expression of huntingtin so that the nucleic acid duplex inhibits expression of huntingtin.

[0029] Another method of the present invention includes a method of inhibiting expression of huntingtin in *Macaca mulatta* and *Homo sapiens* comprising providing a *Macaca mulatta* or *Homo sapiens* containing a neuronal cell that contains and expresses the huntingtin gene and is susceptible to nucleic acid interference and contacting the *Macaca mulatta* or *Homo sapiens* with a previously-described nucleic acid duplex thereby inhibiting expression of the huntingtin gene. In another method of the present invention, expression of huntingtin is inhibited by at least 10%.

[0030] Another method of the present invention includes a method of preventing cytotoxic effects of Huntington's disease ("HD") in a *Macaca mulatta* or *Homo sapiens* comprising introducing a previously-described vector into a cell in an amount sufficient to suppress accumulation of a protein associated with HD, so that the resulting nucleic acid duplex prevents the cytotoxic effects of HD.

[0031] Another method of the present invention includes a method of inhibiting expression of the huntingtin gene in a *Macaca mulatta* or *Homo sapiens* comprising introducing a previously-described vector into a cell in an amount sufficient to inhibit expression of the huntingtin gene so that the resulting nucleic acid duplex inhibits expression of the huntingtin protein.

[0032] Another method of the present invention includes a method of inhibiting expression of huntingtin in a *Macaca mulatta* or *Homo sapiens* comprising providing a *Macaca mulatta* or *Homo sapiens* containing a neuronal cell, wherein the neuronal cell contains and expresses the huntingtin gene and is susceptible to nucleic acid interference, and contacting the *Macaca mulatta* or *Homo sapiens* with a previously-described vector, thereby inhibiting expression of the huntingtin gene.

[0033] In one embodiment of the present invention, the nucleic acid duplexes or vectors are administered to the intrathecal space of the spinal cord. In another embodiment of the present invention, the nucleic acid duplexes or vectors are administered to the cerebrospinal fluid in one or more of the cerebral ventricles of the brain. In another embodiment of the present invention, the nucleic acid duplexes or vectors are administered directly into the brain tissue of the cerebral cortex. In another embodiment of the methods of the present

invention, the nucleic acid duplexes or vectors are administered locally to the basal ganglia. In another embodiment of the present invention, the nucleic acid duplexes or vectors are administered specifically to the caudate nucleus and the putamen.

BRIEF DESCRIPTION OF THE FIGURES

[0034] FIG. 1 shows a target plasmid for nucleic acid sequence characterization in HEK293 cells.

[0035] FIG. 2 shows rhesus monkey HD mRNA suppression by siNA sequences in vitro in HEK293 cells.

[0036] FIG. 3 shows suppression of the endogenous rhesus HD gene at two different siNA doses in 4MBR5 cells.

[0037] FIG. 4 shows suppression of the endogenous rhesus HD gene at five different siNA doses using both 19 and 27 nucleotide length siNAs in 4MBR5 cells.

[0038] FIGS. 5 and 6 show suppression of the endogenous rhesus HD gene using both 19 and 27 nucleotide length siNAs in LLC-MK2 cells.

[0039] FIG. 7 also shows suppression of the endogenous rhesus HD gene, using different doses of both 19 and 27 nucleotide length siNAs in LLC-MK2 cells.

[0040] FIGS. 8A-8D show suppression of the endogenous human HD gene at 4 different siNA doses in HeLa cells.

[0041] FIG. 9 shows suppression of exogenous rhesus huntingtin protein expression in LLC-MK2 cells.

[0042] FIGS. 10 and 11 show suppression of endogenous rhesus huntingtin protein expression in LLC-MK2 cells.

[0043] FIG. 12 depicts a structure and construction of anti-HD (FIG. 12A) and control (FIG. 12B) shNA sequences.

[0044] FIGS. 13A-F show additional shNA sequences used in embodiments according to the present invention.

[0045] FIG. 14 shows suppression of the endogenous rhesus HD gene in LLC-MK2 cells transfected with a plasmid expressing 19 or 27 nucleotide length shNAs with EB4 or mir23 loop structures.

[0046] FIG. 15 shows a schematic description of a pSilencer 1.0-U6 plasmid that can be used in the preparation of anti-HD and control shNA sequences.

[0047] FIG. 16 shows an exemplary format of AAV viral constructs.

[0048] FIG. 17 shows a plasmid used to construct pAAV vectors and to generate AAV in accordance with the present invention.

[0049] FIGS. 18A-B show additional shNA sequences used in embodiments according to the present invention.

[0050] FIG. 19 shows suppression of exogenous rhesus HD by pAAV plasmids.

[0051] FIG. 20 shows suppression of endogenous human HD in HeLa cells and HEK293T cells by recombinant adeno-associated virus (AAV) expressing anti-HD shNA.

[0052] FIGS. 21 and 22 show in vivo suppression of rhesus HD gene expression by recombinant adeno-associated virus (AAV) expressing anti-HD shNA.

DETAILED DESCRIPTION OF THE
INVENTION

[0053] The term “SEQ ID NO: X” (where X is any number from 1 to 15) refers to, in one embodiment, each number’s sequence as defined in Table 1 (identified sequence). SEQ ID NO: X must also be read to encompass sequences that would hybridize with the complementary strand of a sequence set forth in SEQ ID NOS: 1-15 and reduce the particular SEQ ID NO:’s target mRNA in a cell type selected from, without limitation, HEK293 cells, 4MBR5 cells, LLC-MK2 cells, HeLA cells or any other *Macaca mulatta* or *Homo sapien* cell type. Under this definition, claimed sequences can include at least 99% sequence homology with the identified sequence; at least 98% sequence homology with the identified sequence; at least 95% sequence homology with the identified sequence; at least 90% sequence homology with the identified sequence; or at least 85% sequence homology with the identified sequence.

[0054] The terms “nucleic acid” or “nucleic acid molecules” refer to deoxyribonucleotides or ribonucleotides and polymers thereof in either single- or double-stranded form, composed of monomers (nucleotides) containing a sugar, phosphate and a base that is either a purine or pyrimidine. Unless specifically limited, the term encompasses nucleic acids containing known analogs of natural nucleotides that have similar binding properties as the reference nucleic acid and are metabolized in a manner similar to naturally occurring nucleotides. Unless otherwise indicated, a particular nucleic acid sequence also encompasses conservatively modified variants thereof. The nucleic acid molecules of the present invention can include any type of nucleic acid molecule capable of mediating RNA interference, such as, without limitation, short interfering nucleic acid (siNA), short hairpin nucleic acid (shNA), short interfering RNA (siRNA), short hairpin RNA (shRNA), micro-RNA (miRNA), and double-stranded RNA (dsRNA). The nucleic acid molecules of the present invention also include similar DNA sequences. Further, the nucleic acid and nucleic acid molecules of the present invention can contain unmodified or modified nucleotides. Modified nucleotides refer to nucleotides which contain a modification in the chemical structure of a nucleotide base, sugar and/or phosphate. Such modifications can be made to improve the stability and/or efficacy of nucleic acid molecules and are described in patents and publications such as U.S. Pat. No. 6,617,438, U.S. Pat. No. 5,334,711; U.S. Pat. No. 5,716,824; U.S. Pat. No. 5,627,053; U.S. Patent Application No. 60/082,404, International Patent Cooperation Treaty Publication Number (“PCTPN”) WO 98/13526; PCTPN WO 92/07065; PCTPN WO 03/070897; PCTPN WO 97/26270; PCTPN WO 93/15187; Beigelman et al., 1995, *J. Biol. Chem.*, 270, 25702; Usman and Cedergren, 1992, *TIBS*, 17, 34; Usman et al., 1994, *Nucleic Acids Symp. Ser.* 31, 163; Burgin et al., 1996, *Biochemistry*, 35, 14090; Perrault et al. *Nature*, 1990, 344, 565-568; Pieken et al. *Science*, 1991, 253, 314-317; Usman and Cedergren, *Trends in Biochem. Sci.*, 1992, 17, 334-339; Karpeisky et al., 1998, *Tetrahedron Lett*, 39, 1131; Earnshaw and Gait, 1998, *Biopolymers (Nucleic Acid Sciences)*, 48, 39-55; Verma and Eckstein, 1998, *Annu. Rev. Biochem.*, 67, 99-134; Burlina et al., 1997, *Bioorg. Med. Chem.*, 5, 1999-2010; Limbach et al., 1994, *Nucleic Acids Res.* 22, 2183; and Burgin et al., 1996, *Biochemistry*, 35, 14090. Such

patents and publications describe general methods and strategies to modify nucleic acid molecules and are incorporated by reference herein.

[0055] The phrase “expression cassette” as used herein means a nucleic acid sequence capable of directing expression of a particular nucleotide sequence in an appropriate host cell, with additional sequences that facilitate appropriate transcription of the nucleic acid sequence of interest. In addition to the nucleotide sequence of interest, the expression cassette can include a promoter operably linked to the nucleotide sequence of interest that also can be operably linked to termination signals. The expression cassette also can include expression enhancers. The expression cassette including the nucleotide sequence of interest can be chimeric. The expression cassette also can be one that is naturally occurring but has been obtained in a recombinant form useful for heterologous expression. The expression of the nucleotide sequence in the expression cassette can be under the control of a constitutive promoter or of a regulatable promoter that initiates transcription only when the host cell is exposed to some particular stimulus. In the case of a multicellular organism, the promoter also can be specific to a particular tissue or organ or stage of development.

[0056] The term “promoter” refers to a nucleotide sequence, usually upstream (5 prime) of the nucleotide sequence of interest, which directs and/or controls expression of the nucleotide sequence of interest by providing for recognition by RNA polymerase and other factors required for proper transcription. As used herein, the term “promoter” includes (but is not limited to) a minimal promoter that is a short DNA sequence comprised of a TATA-box and other sequences that serve to specify the site of transcription initiation, to which regulatory elements are added for control of expression. The term “promoter” also refers to a nucleotide sequence that includes a minimal promoter plus regulatory elements that is capable of controlling the expression of a coding sequence or functional RNA. This type of promoter sequence consists of proximal and more distal upstream elements, the latter elements often referred to as enhancers. The term “enhancer” refers to a DNA sequence that can stimulate promoter activity and can be an innate element of the promoter or a heterologous element inserted to enhance the level or tissue specificity of a promoter. Enhancers are capable of operating in both orientations (normal or flipped), and are capable of functioning even when moved either upstream or downstream of the promoter. Both enhancers and other upstream promoter elements bind sequence-specific DNA-binding proteins that mediate their effects. Promoters can be derived in their entirety from a native gene, or be composed of different elements derived from different promoters found in nature, or even be comprised of synthetic DNA segments. A promoter also can contain DNA sequences that are involved in the binding of protein factors that control the effectiveness of transcription initiation in response to physiological or developmental conditions. Specific promoters used in accordance with the present invention can include, for example and without limitation pol II promoters (including, without limitation cytomegalovirus (“CMV”) promoters, chicken β -actin (“CBA”) promoters, Rous sarcoma virus (“RSV”) promoters and neuron-specific enolase (“NSE”) promoters). Furthermore, specific promoters used in accordance with the present invention can include, for example and without limitation, pol III promoters (including, without limitation,

human H1 and human or murine U6 promoters, as well as H1 and U6 promoters engineered to be expressed in a regulated way such as described in United States Patent Application Number 2005/0064489).

[0057] The term “vector” is defined to include any virus, as well as any plasmid, cosmid, phage, binary vector or segment of nucleic acid (DNA or RNA) in double or single stranded linear or circular form that may or may not be self transmissible or mobilizable, and that can transform eukaryotic host cells either by integration into the cellular genome or by existing extrachromosomally (e.g., autonomous replicating plasmid with an origin of replication).

[0058] The gene involved in Huntington’s disease (“HD”) (IT-15) is located on chromosome 4 at the end of the short arm. This gene encodes for the protein huntingtin (also known as “htt”). The mutation in the HD gene responsible for HD is an unstable expanded CAG trinucleotide repeat within the coding region of the gene. This mutation results in a huntingtin protein with an expanded glutamine sequence. While the normal and abnormal functions of huntingtin are not known, the presence of the mutated form of huntingtin has been correlated with the occurrence of HD symptoms. Further, the abnormal huntingtin protein appears to accumulate abnormally in neuronal nuclei. Thus, blocking the expression of huntingtin provides a useful mechanism to study HD pathogenesis and can also provide a potential treatment for HD. A proof-of-principle that suppressing the expression of mutant huntingtin protein can be therapeutic for HD has been shown in mice by Harper et al. who showed that suppression of a mutant HD transgene in mice by use of RNA interference resulted in improvements in the disease phenotype of the transgenic mice. *Proceedings of the National Academy of Sciences*, 2005, 102(16) 5820-5825. It should be noted, however, that the present invention differs from this prior art in that the shNA sequence used and disclosed by Harper et al. is different from any and all of the nucleic acid sequences of the present invention. Furthermore, Harper et al. does not indicate whether the nucleic acid sequence of their study is homologous with both rhesus and human HD genes, whereas the nucleic acid sequences of the present invention are known to be homologous with both rhesus (*Macaca mulatta*) and human (*Homo sapiens*) HD genes. Thus, one important benefit of the present invention is that the same nucleic acid sequences can be studied and characterized for toxicological and pharmacological properties and safety in the non-human primate *Macaca mulatta* and then, without alteration, can also be used for therapeutic benefit in *Homo sapiens*.

[0059] In order to use RNA interference (“RNAi”) to suppress expression of rhesus and human huntingtin, it was first necessary to identify nucleic acid sequences that effectively suppress the expression of human and rhesus HD genes. Towards this goal, a PCR strategy was used to clone and assemble 3437 bp of the 5' portion of the rhesus HD gene. Briefly, a series of PCR primers predicted to anneal to the rhesus HD cDNA were designed based on evolutionarily conserved sequences identified in an alignment of the human (NM_002111) and pig (AB016793) cDNA sequences. These primers were used to amplify partially overlapping portions of the rhesus HD gene from first strand cDNA prepared from rhesus brain and kidney tissues (BioChain Institute, Inc; C1534035 and C1534142 respectively) using high fidelity DNA polymerases. The recovered PCR prod-

ucts were cloned into pCR-Blunt-TOPO (Invitrogen) and sequenced using standard methods. Sequence alignments between multiple, independently derived clones were analyzed to ensure the elimination of potential PCR and sequencing errors. This sequence information was also used to design additional primers to clone gaps in the recovered HD cDNA sequence. A single composite clone containing 3437 bp of the rhesus HD cDNA was assembled using PCR and conventional cloning techniques.

[0060] Next, potential synthetic nucleic acid suppressor sequences for rhesus monkey huntingtin that correspond with portions of the human sequence (i.e., target genes were rhesus and human HD genes) were identified using online software (<http://www.dharmacon.com/>) and also obtained from Dharmacon® (Dharmacon Research, Inc., Boulder, Colo.). These nucleic acid sequences included those in the following Table 1:

TABLE 1

SEQ ID NO:Name	Nucleic Acid Sequence Nucleotide Base Sequence	Position in human cDNA (NM_002111)
1 rh#1-19	TGACAGCAGTGTGATAAA	2071-2089
2 rh#2-19	AAGAACGAGTGCTCAATAA	2862-2880
3 rh#4-19	TTTATGAACTGACGTTACA	1221-1239
4 rh#5-19	GGAGTATTGTGGAAC TTAT	1404-1422
5 rh#6-19	GAGTATTGTGGAAC TTATA	1405-1423
6 rh#11-19	AGACCGTGTGAATCATTGT	442-460
7 rh#12-19	GGTTACAGCTCGAGCTCTA	645-663
8 rh#13-19	GGTTTTGTTAAAGGCCTTC	898-916
9 rh#1-27	TGACAGCAGTGTGATAAAATTTGTGTT	2071-2097
10 rh#2-27	AAGAACGAGTGCTCAATAATGTTGTCA	2862-2888
11 rh#4-27	TTTATGAACTGACGTTACATCATAAC	1221-1247
12 rh#5-27	GGAGTATTGTGGAAC TTATAGCTGGAG	1404-1430
13 rh#6-27	GAGTATTGTGGAAC TTATAGCTGGAGG	1405-1431
14 rh#11-27	AGACCGTGTGAATCATTGTCTGACAAT	442-468
15 rh#13-27	GGTTTTGTTAAAGGCCTTCATAGCGAA	898-924

Note that, as will be understood by one of skill in the art, the nucleic acid molecules of the present invention include the sequences in the preceding table, the reverse complement of these sequences and RNA based sequences including uracils in the place of the listed thymines. Thus, the sequences in the preceding table can be considered target sequences as well as sequences included in the nucleic acid molecules of the present invention. SEQ ID NOS. 1-8 are 19 nucleotide base sequences. SEQ ID NO. 9-15 are the same sequences extended to 27 nucleotide base sequences. The sequence identified as SEQ ID NO. 7 was not extended into a 27 nucleotide base sequence because extending this sequence resulted in a mismatch between the rhesus monkey and human HD sequences. Control sequences also were chosen. The control sequences included a non-sense scrambled

control (SEQ ID NO. 19; TAGCGACTAAACACATCAA) and a TNF α sequence (SEQ ID NO. 16; AATCCTCTCTCGTATTATA) both purchased from Dharmacon.

[0061] To identify which of the above identified 19 nucleotide nucleic acid sequences most effectively suppressed rhesus monkey huntingtin expression, an in vitro co-transfection assay system was developed. Referring to FIG. 1, rhesus monkey HD gene sequences were subcloned into pTracerTM-CMV2 (Invitrogen, Corp., Carlsbad, Calif.) to generate pTracer-rhuntingtin (pTRACER-rhHD). The recombinant plasmid also included a GFP-Zeocin reporter gene for transfection efficiency normalization. The CMV promoter directed constitutive expression of the target gene (rhesus monkey HD) while the EF1 promoter directed constitutive expression of the GFP-Zeocin reporter gene.

[0062] The generated recombinant plasmids were used to facilitate screening of nucleic acid sequences by co-transfection into a eukaryotic cell line. Specifically, HEK293 cell cultures at 60-70% confluency were co-transfected with the appropriate target plasmid (2 μ g/well of a 6-well plate) and test nucleic acid sequences (100 nM) directed against rhesus monkey HD (SEQ ID NO: 1 through SEQ ID NO: 8), TNF α as a control (SEQ ID NO. 16) or a non-sense scrambled control (SEQ ID NO. 19). Forty-eight hours post-transfection, total cellular RNA was harvested from the cells and used to make cDNA by standard methods. The cDNAs were analyzed for target (HD) and reporter (GFP) gene expression levels using realtime PCR methods. The data was normalized for GFP expression levels to control for variation in transfection efficiency.

[0063] FIG. 2 shows the results of the rhesus monkey HD transfection studies related to each of the test nucleic acid sequence's ability to suppress rhesus monkey HD mRNA. As measured by realtime PCR, all non-control sequences effectively suppressed expression of the HD gene when compared to mock treated control cells (set at 100% expression). Because the nucleic acid sequences evaluated in the present experiment have complete sequence identity to portions of the human HD gene sequence, it is reasonable to expect that they also suppress the expression of the human HD gene, thus suppressing production of human huntingtin protein.

[0064] Next, the ability of a subset of the 19 nucleotide sequences was evaluated for its ability to suppress expression of the endogenous rhesus HD gene at two different doses, 100 nM and 10 nM. Specifically, 4MBr5 cell line cultures obtained from American Type Culture Collection ("ATCC") (cells from a rhesus lung cell line) at 70-80% confluency were transfected with test nucleic acid sequences (SEQ ID NO: 1; SEQ ID NO: 2; SEQ ID NO: 4; SEQ ID NO: 6; and SEQ ID NO: 7) directed against rhesus HD mRNA at either 100 nM or 10 nM or control sequences (SEQ ID NO: 19 (scrambled) or SEQ ID NO: 16 (off-target TNF α control)) at 100 nM. Forty-eight hours post transfection total cellular RNA was collected from the cells and used to generate cDNA by standard methods. The cDNAs were used to analyze for rhesus HD and control gene expression using realtime PCR methods. A separate realtime PCR reaction was performed to assess the level of rhesus GAPDH expression. All of the expression data was normalized to GAPDH expression to allow the assessment of the efficacy of siNA-mediated knockdown of rhesus HD in the rhesus

cell lines. As shown in FIG. 3, all sequences directed against rhesus HD mRNA suppressed rhesus HD gene expression at both doses when compared to controls.

[0065] Next, an experiment was performed to evaluate the dependence of rhesus HD mRNA suppression on the length and dose of provided nucleic acid sequences. Specifically, 4MBr5 cells were transfected as described above but with 100 nM, 10 nM, 1.0 nM, 0.2 nM or 0.05 nM of SEQ ID NO: 4 (19 nucleotide sequence); 100 nM, 10 nM, 1.0 nM, 0.2 nM or 0.05 nM of SEQ ID NO: 12 (corresponding 27 nucleotide sequence); 100 nM SEQ ID NO: 19 (scrambled) or 100 nM SEQ ID NO: 16 (off-target TNF α control). As shown in FIG. 4, all doses of both sequence lengths suppressed rhesus HD gene expression when compared to controls. While all doses were effective in 4MBr5 cells, 100 nM, 10 nM and 1 nM were the most effective doses for suppressing rhesus HD gene expression regardless of sequence length.

[0066] A subsequent experiment evaluating the siNA's ability to suppress expression of the endogenous rhesus HD gene also examined the effectiveness of a 27 nucleotide sequence as compared to its previously-evaluated 19 nucleotide sequence counterpart in LLC-MK2 cells. In this experiment, LLC-MK2 cells (a rhesus kidney cell line also obtained from ATCC) at 70-80% confluency were transfected with test nucleic acid sequences (SEQ ID NO: 4 and SEQ ID NO: 12) at one of two doses, 10 nM or 0.2 nM as well as controls (SEQ ID NO: 19 and SEQ ID NO: 16) at 100 nM. Forty-eight hours post transfection total cellular RNA was collected from the cells and used to generate cDNA by standard methods. The cDNAs were used to analyze for rhesus HD and control gene expression using realtime PCR methods. A separate realtime PCR reaction was performed to assess the level of rhesus GAPDH expression. All of the expression data was normalized to GAPDH expression to allow the assessment of the efficacy of siNA-mediated knockdown of rhesus HD in the rhesus cell lines. As shown in FIG. 5, all sequences directed against rhesus HD mRNA suppressed rhesus HD gene expression at both doses when compared to controls.

[0067] Following the experiment described in the preceding paragraph, an identical experiment was conducted with the exception that an additional lower dose of SEQ ID NOS: 4 and 12 (0.05 nM) was compared against the effectiveness of the 0.2 nM dose and control SEQ ID NO: 19 (100 nM). As shown in FIG. 6, all sequences and doses suppressed endogenous rhesus HD gene expression in LLC-MK2 cells.

[0068] The effectiveness of different sequence lengths and doses was also compared using SEQ ID NOS: 1 and 9. Following the same procedures used in the two previously described experiments, LLC-MK2 cells were transfected with 100 nM, 10 nM, 1 nM, 0.1 nM or 0.01 nM SEQ ID NO: 1 (19 nucleotide sequence); 10 nM, 1 nM, 0.1 nM or 0.01 nM SEQ ID NO: 9 (corresponding 27 nucleotide sequence); or control SEQ ID NOS: 19 or 16 (100 nM). As shown in FIG. 7, all doses of both sequences suppressed endogenous rhesus HD gene expression in LLC-MK2 cells with the 1 nM and higher doses of SEQ ID NOS: 1 and 9 providing the most suppression. In addition, from this experiment the IC₅₀ (inhibitor concentration required for 50% suppression) values for these two siNAs are estimated to be 10 pM and 75 pM respectively.

[0069] Next, a subset of the identified siNA sequences was tested for the chosen sequence's ability to suppress expres-

sion of the endogenous human HD gene. In this experiment, HeLa cells (obtained from ATCC) at 70-80% confluency were transfected with test nucleic acid sequences (SEQ ID NO: 1; SEQ ID NO: 2; SEQ ID NO: 4; SEQ ID NO: 6; or SEQ ID NO: 7) or control (SEQ ID NO: 19) at 100 nM. Forty-eight hours post transfection total cellular RNA was collected from the cells and used to generate cDNA by standard methods. The cDNAs were used to analyze for human HD and control gene expression using realtime PCR methods. A separate realtime PCR reaction was performed to assess the level of human GAPDH expression. All of the expression data was normalized to GAPDH expression to allow the assessment of the efficacy of siNA-mediated knockdown of human HD in the HeLa cell line. As shown in **FIG. 8A**, all sequences directed against human HD mRNA suppressed human HD gene expression when compared to controls.

[0070] The experiment just described was next repeated using doses of either 1 nM, 0.1 nM or 0.01 nM SEQ ID NOS. 1, 2, 4, 6 or 7 or 100 nM of control SEQ ID NO: 19. As shown in **FIGS. 8B-8D**, all doses of each of these SEQ ID NOS. 1, 2, 4, 6 and 7 were effective in suppressing endogenous human HD gene expression in HeLa cells. In only one instance was human HD suppression not observed, SEQ ID NO: 2 at the lowest dose (0.01 nM).

[0071] Following the previously described experiments, studies were undertaken to determine whether sequences of the present invention suppress exogenous huntingtin protein expression. In this first protein suppression experiment, suppression of the partial rhesus huntingtin protein expressed from the pTRACER plasmid depicted in **FIG. 1** was measured by western blot analysis. Specifically, LLC-MK2 cells were cotransfected with the pTRACER-rhHD plasmid (2 μ g/well of a 6-well plate) and 100 nM of either SEQ ID NO: 19 (control); SEQ ID NO: 1 or SEQ ID NO: 4. Controls in this experiment also consisted of LLC-MK2 cells transfected solely with the pTRACER-rhHD plasmid. Protein was harvested 48 hours post-transfection and was analyzed using western blot analysis. In this analysis, 10 μ g of total protein was loaded per well. Rhesus huntingtin protein expression was measured using the Chemicon® International (Temecula, Calif.) huntingtin antibody MAB2168 and β -actin was measured with abcam® (Cambridge, Mass.) antibody 20272-100. To quantify huntingtin protein-expression the intensities of the bands for huntingtin and β -actin on the autoradiogram of the western blot were measured by densitometry. Huntingtin protein expression was normalized to β -actin expression. As can be seen in **FIG. 9**, both SEQ ID NO: 1 and SEQ ID NO: 4 were effective to suppress exogenous huntingtin protein expression when compared to controls transfected with the pTRACER-rhHD plasmid and SEQ ID NO: 19 or the pTRACER-rhHD plasmid alone.

[0072] Next the ability of SEQ ID NO: 1 and SEQ ID NO: 4 to suppress endogenous protein expression was examined in LLC-MK2 cells again using western blot analysis. LLC-MK2 cells were transfected with 100 nM of either SEQ ID NO: 1, SEQ ID NO: 4 or SEQ ID NO: 19. An additional control consisted of extracts from untransfected cells. Protein was harvested and analyzed 48 hours post-transfection with 10 μ g of total protein loaded per western blot well. Again, rhesus huntingtin protein expression was measured using the Chemicon® International (Temecula, Calif.) hun-

tingtin antibody MAB2168 and β -actin was measured with abcam® (Cambridge, Mass.) antibody 20272-100. As is evident on the western blot for rhesus huntingtin and β -actin shown in **FIG. 10** both SEQ ID NO: 1 and SEQ ID NO: 4 suppress huntingtin protein expression. To quantify the amount of protein suppression, the level of rhesus huntingtin protein expression was normalized to β -actin expression. To do this the intensities of the bands on the autoradiogram of the western blot (**FIG. 10**) were measured by densitometry. As can be seen in **FIG. 11**, SEQ ID NO: 1 and SEQ ID NO: 4 both suppressed endogenous huntingtin protein expression when compared to cells transfected with control SEQ ID NO: 19 or to extract from untransfected cells.

[0073] After effective nucleic acid sequences are identified as described above, effective sequences can be formatted into short hairpin nucleic acid ("shNA") structures. **FIG. 12** shows SEQ ID NO: 4 reformatted and constructed as an anti-huntingtin shNA sequence (SEQ ID NO: 17; **FIG. 12A**). **FIG. 12** also shows a control shNA sequence (SEQ ID NO: 18; **FIG. 12B**). The shNA sequences contain the siNA nucleic acid sequence (boxed sequence shown in black; in **FIG. 12A**, an originally effective nucleic acid sequence), a nine nucleotide loop (underlined), the reverse complement of the boxed nucleic acid sequence (italicized), and a Pol III terminator sequence (TTTTTT (bolded)). Four partially overlapping synthesized oligonucleotides (-A1, -A3, -A2, and -A4 in **FIG. 12A** and -B1, -B3, B2 and -B4 in **FIG. 12B**) can be used to assemble the shNAs. In two separate reactions, the $\frac{1}{3}$ and $\frac{2}{4}$ oligonucleotides for each shNA can be annealed. Referring to **FIGS. 12A and 12B**, the A1 and B1 oligonucleotides include the boxed sequences. The A2 and B2 oligonucleotides include the underlined nine nucleotide loop, the reverse complement of oligonucleotides A1 and B1 respectively, the bolded Pol III terminator sequence (TTTTTT) and the first G of the EcoRI site depicted in these **FIGS.** The A3 and B3 oligonucleotides of **FIGS. 12A and 12B** respectively include the Apal overhang region, the reverse complement of the boxed A1/B1 sequences, and the reverse complement of the underlined nine nucleotide loop sequence. The A4 and B4 oligonucleotides include the reverse complement of the italicized portion of A2 and B2 respectively and the reverse complement (AAAAAA) of the bolded Pol III terminator sequence. Finally, the A4 and B4 oligonucleotides contain the EcoRI overhang region. Apal and EcoRI restriction enzyme-compatible ends are included in the shNA structures for directional subcloning into a murine U6 promoter-containing shuttle vector (pSilencer1.0-U6; Ambion, Inc., Austin, Tex.; **FIG. 15**). The full-length shNAs (SEQ ID NOS: 17 and 18) then can be cloned into a Apal/EcoRI-digested pSilencer vector using a three-way ligation reaction. The U6 promoter (murine or human) is used for constitutive high level expression of the nucleic acid sequences. In keeping with the present invention, a human H1 promoter can also be used. One non-limiting example of a shuttle vector containing the human U6 promoter is psiSTRIKE™ hmGFP from Promega Corporation (Madison, Wis.). One non-limiting example of a shuttle vector containing the human H1 promoter is pSilencer 3.0-H1 from Ambion, Inc. (Austin, Tex.).

[0074] While this example provides one embodiment of the nucleic acid sequences of the present invention in shNA format and a method for creating them, other configurations and methods also fall within the scope of the present invention. For example, in one embodiment, the loop struc-

ture of the shNA hairpin is 4 to 13 nucleotides in length. In another embodiment, the arms of the hairpin structures are less than approximately 30 nucleotides in length. In another embodiment, the arms of the hairpin structure are between 19 and 27 nucleotides in length. Thus, while a specific example is given, this example should not be interpreted to limit the scope of the present invention.

[0075] In the next described experiment, the ability of plasmids expressing shNA to suppress rhesus HD gene expression was examined. LLC-MK2 cells were transfected with a psiSTRIKE™ hmGFP plasmid (Promega Corporation, Madison, Wis.) with one of two stem lengths (SEQ ID NO: 4 (19 nucleotides) or SEQ ID NO: 12 (27 nucleotides)) and one of two different loop structures (EB4 or mir23). A scrambled control was also generated for each of the loop structures. Specifically, these scrambled control sequences included a 27 nucleotide sequence control (SEQ ID NO: 20) for EB4 (**FIG. 13A**) and a 19 nucleotide sequence control (SEQ ID NO: 21) for mir23 (**FIG. 13B**; SEQ ID NO: 21 is a modified version of control SEQ ID NO: 18). Thus, the whole of the transfected sequences in this study included SEQ ID NO: 20; SEQ ID NO: 21; SEQ ID NO: 22 (**FIG. 13C**; modified SEQ ID NO: 17-EB4 (as will be recognized by one of ordinary skill in the art, SEQ ID NO: 22 includes SEQ ID NO: 17 with modified overhang regions for subcloning into a different vector)); SEQ ID NO: 23 (**FIG. 13D**; modified SEQ ID NO: 17-mir23); SEQ ID NO: 24 (**FIG. 13E**; shNA-modified SEQ ID NO: 12-EB4); and SEQ ID NO: 25 (**FIG. 13F**; shNA-modified SEQ ID NO: 12-mir23). As can be seen in **FIG. 14**, all test plasmids suppressed rhesus HD gene expression when compared to control plasmids.

[0076] Next, a method to prepare viral production plasmids is described. This method is based on the method for producing the nucleic acid sequences in shNA formats described above. As will be understood by one of skill in the art, however, various known modifications of the described technique also fall within the scope of the present invention. First, the BamHI fragment (**FIG. 15**) containing the shNA expression cassette (murine U6 promoter, shNA sequence, and Pol III terminator sequence) from each of the pSilencer shuttle vectors (huntingtin and control) is recovered, blunted with T4 DNA polymerase, and subcloned into an AAV1/2 expression vector (deprAVE™; GeneDetect.com Ltd, Bradenton, Fla.). As shown in **FIG. 16**, in this example, in the final viral expression vector (used for virus production), the U6 promoter (murine or human) can drive the expression of the shNA and the Woodchuck enhancer/chicken β -actin promoter can drive the expression of the enhanced green fluorescent protein (WPRE/CAG-eGFP). These expression cassettes can be flanked by viral inverted terminal repeats (ITR). The Woodchuck post-transcriptional regulatory element (WPRE) and the presence of a bovine growth hormone (BGH) polyadenylation sequence ensure high transcription following cellular transduction. In alternative methods of the present invention, the expression cassette additionally contains a polyadenylation signal, such as a synthetic minimal polyadenylation signal. In another alternative embodiment of the present invention, the H1 promoter can be used.

[0077] **FIG. 17** depicts a different plasmid, the plasmid used to construct pAAV vectors used in the following experiments to generate AAV (pAAV-hrGFP, Stratagene, La Jolla, Calif.). Six different plasmids were constructed; 3 with

an H1 promoter and 3 with a U6 promoter (human H1 promoter from Ambion (Austin, Tex.) and human U6 promoter from Promega (Madison, Wis.)). Specifically, the following were subcloned into the depicted vector (1) U6-SEQ ID NO: 17; (2) U6-SEQ ID NO: 26 (SEQ ID NO: 26 is SEQ ID NO: 1 in an shRNA format; see **FIG. 18A**); (3) H1-SEQ ID NO: 17; (4) H1-SEQ ID NO: 26; (5) U6-SEQ ID NO: 18 (scrambled shNA control); and (6) H1-SEQ ID NO: 27 (SEQ ID NO: 27 is a scrambled SEQ ID NO: 1 shNA control; see **FIG. 18B**). In this instance this was done by removing a cassette containing the promoter and the shNA sequence from the appropriate psiSTRIKE™ or pSilencer 3.0 shNA expressing vector with PvuII. This PvuII fragment was then subcloned into the blunted MluI site of pAAV-hrGFP (Stratagene, La Jolla, Calif.). Directionality was confirmed by restriction digest mapping and sequence analysis.

[0078] These six pAAV plasmids were co-transfected into HEK293T cells along with pTRACER-rhHD (**FIG. 1**). As controls, two siRNAs (SEQ ID NOS: 1 or 4) and the non-sense scrambled control (SEQ ID NO: 19) were also transfected into HEK293T cells. pTRACER-rhHD was also transfected alone as a control. RNA was collected 48 hours post transfection and the suppression of rhesus HD gene expression (on the pTRACER plasmid) was measured by realtime PCR and normalized to the amount of GFP expressed from the pTRACER-rhHD plasmid. As can be seen in **FIG. 19**, pAAV plasmids containing U6-SEQ ID NO: 17; U6-SEQ ID NO: 26; H1-SEQ ID NO: 17; and H1-SEQ ID NO: 26 were as effective at suppressing exogenous rhesus HD gene expression as previously tested SEQ ID NO: 1 and SEQ ID NO: 4. pAAV plasmids expressing scrambled control sequences did not suppress exogenous rhesus HD gene expression.

[0079] Next, the described pAAV plasmids along with the required pHELPER and REPCAP plasmids were used to generate recombinant adeno-associated virus (AAV) with serotype 1. Five separate AAV viruses were generated. These viruses were engineered to express U6-SEQ ID NO: 26; U6-SEQ ID NO: 17; H1-SEQ ID NO: 26; U6-SEQ ID NO: 18 (shNA control) or H1-SEQ ID NO: 27 (shNA control). The ability of these viruses to suppress endogenous human HD gene expression in HeLa cells and HEK293T cells was examined. 5e5 cells in a well of a 6-well plate were transduced with ~5e9 virions by directly adding the virus to the well of cells in 1 ml of Dulbecco's Modified Eagle Medium (DMEM) cell culture media supplemented with 2% fetal bovine serum (FBS). This plate was incubated at 37° C., 5% CO₂ for 2 hours with gentle mixing by rocking every 30 minutes. Following this 2 hour incubation, 1 ml of DMEM cell culture media supplemented with 18% FBS was added to each well yielding a final FBS concentration of 10%. Growth of the transduced cells was continued at 37° C., 5% CO₂. RNA was collected 72 hours post transduction and used to make cDNA by standard methods. The cDNAs were analyzed for HD and GAPDH expression levels by realtime PCR methods. Endogenous HD expression was normalized to endogenous GAPDH expression. As shown in **FIG. 20**, U6-SEQ ID NO: 26 and U6-SEQ ID NO: 17 were effective at suppressing endogenous human HD gene expression. P values were determined by comparison to the mock transduced data set. In this experiment, even though H1-SEQ ID NO: 26 did not significantly suppress endogenous human HD expression in vitro, its potential effectiveness in vivo

should not be discounted especially in light of literature indicating a potential disconnect between *in vitro* and *in vivo* activity. See, for example, Wooddell et al., 334 Biochem. and Biophys. Res. Comm. 117-27 (2005).

[0080] The siNA sequences and molecules of the present invention can be manipulated to enhance their uptake into the RISC complex. Specifically, manipulating the 3 prime terminal nucleotide of the sense strand can be highly advantageous. Preferential entry of the guide, or antisense, strand into RISC can be achieved by introducing 3 prime mismatches in the sense strand while maintaining perfect base pairing (of the antisense strand and the intended mRNA target) at the 5 prime terminus of the antisense strand. This maximizes entry of the antisense strand into the RISC complex, while also reducing potential off-target inhibition by the sense strand.

[0081] Physical methods to introduce nucleic acid molecules and/or their carriers (i.e. vectors) into eukaryotic cells are known in the art. Some of these methods include, without limitation, calcium phosphate or calcium chloride co-precipitation, DEAE-dextran-mediated transfection, cationic lipid-mediated transfection, lipofection, protoplast fusion, particle bombardment, microinjection, liposome fusion, biolistics and other suitable methods found in, for example, Sambrook, et al. (Molecular Cloning: A Laboratory Manual, 2nd, ed, Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1989), and other laboratory manuals. Further, biological methods to introduce nucleic acid molecules into a cell include the use of viral vectors. For mammalian gene therapy, viral vectors, and especially retroviral vectors, have become the most widely used method for inserting genes into mammalian, e.g., rhesus monkey or human cells. Other viral vectors can be derived from poxviruses, herpes simplex virus I, adenoviruses and adeno-associated viruses, and the like (see, for example, Boenisch et al., U.S. Pat. No. 5,350,674 and Wilson et al., U.S. Pat. No. 5,585,362 which are hereby incorporated by reference). Embodiments of the present invention also can be delivered through the use of liposomes, polyethyleneimine, by iontophoresis, or by incorporation into other vehicles, such as hydrogels, cyclodextrins, biodegradable nanocapsules, and bioadhesive microspheres. Nucleic acid molecules also can be directly delivered to cells or tissues with or without the aforementioned vehicles. The nucleic acid molecules/vehicle combinations can be locally delivered by catheter and drug pump systems, delivered by direct local injection or through the use of polymers and/or drug-eluting stents.

[0082] The nucleic acid sequences, molecules, expression cassettes and vectors of the present invention can be used to

suppress HD gene expression and resulting huntingtin formation. The nucleic acid sequences, molecules, expression cassettes and vectors of the present invention also can be used in the study of HD pathogenesis. Further, the nucleic acid sequences, molecules, expression cassettes and vectors of the present invention also may be administered to prevent or treat the symptoms of HD. When used as a potential treatment, the amount of these agents administered as well as the timing of their delivery will vary depending on various factors including, for example and without limitation, the composition chosen, the weight, physical condition, and age of the patient, and whether prevention or treatment is desired as well as other factors known to skilled practitioners. Administration of the therapeutic agents can be continuous or intermittent. The administration of the agents of the invention can be essentially continuous over a pre-selected period of time or can be in a series of spaced doses. Further, both local and systemic administration can be appropriate for use within the present invention. Such factors can be determined by the researcher or treating physician.

[0083] A pre-clinical study of the safety of suppressing huntingtin protein expression in the normal adult rhesus monkey brain was conducted. In this study, a 6 year old female rhesus monkey weighing approximately 3.9 kg was anesthetized and bilaterally infused with an anti-HD shRNA/GFP-containing recombinant adeno-associated virus (AAV) (AAV-U6-SEQ ID NO: 17 (serotype AAV1; 2.07×10^{12} vg/ml; lot #J0531)) by stereotactic injection. Stereotactic targets were determined by presurgical MRI and included putamen (AP=17; L=11; DV1=20 and DV2=17) and caudate nucleus (AP=20.5; L=5.6; DV=17) sites. Injections (Hamilton syringe (100 μ l) with 27 Ga needle (compression fitting)) were performed at 2 putamen sites (50 μ l/site) using a single needle tract and 1 caudate site (50 μ l) at a rate of 1 μ l/min. Following this injection, the needle was left in place for at least 20 minutes and then withdrawn at a rate of 1 mm/min.

[0084] Twenty-eight days later, the animal was deeply anesthetized with pentobarbital (1.5 ml intramuscular; 2 ml, intravenous) and transcardially perfused with heparinized ice-cold saline (4-6 L). Following perfusion and removal, the brain was placed into a container of ice-cold saline.

[0085] The right hemisphere of the brain was sectioned into 14-4 mm thick coronal sections and numbered 1-14 (rostral to caudal). A 14 G biopsy needle was then used to collect tissue punches from various brain regions. A total of 103 tissue punches was collected from the right hemisphere. Each tissue punch contained about 6 mg of tissue and was snap frozen in liquid nitrogen and stored at -80° C. until RNA was isolated.

Caudate; 20 punches (3 slabs)	Putamen; 20 punches (3 slabs)	Nigra, 5 punches (1 slab)
Accumbens; 8 punches (2 slabs)	GP; 5 punches (1 slab)	Front. cx; 9 punches (1 s)
Cerebellum; 6 punches (1 slab)	Occipital cx; 9 punches (1 slab)	Motor cx; 9 punches (1 s)
Thalamus; 6 punches (1 slab)	Hippocampus; 6 punches (1 slab)	Total n = 103 punches

RNA was isolated from tissue punches using the RNeasy Lipid Kit from Qiagen (Valencia, Calif.). Total RNA was used to generate cDNA using the Stratascript® First Strand cDNA synthesis kit from Stratagene (La Jolla, Calif.). The cDNAs were used to analyze expression levels of rhesus HD and GAPDH by realtime PCR. In addition, AAV expression was monitored by measuring the level of GFP expression. Rhesus HD expression was normalized to rhesus GAPDH expression and compared to a region not expressing the virus (GFP-).

[0086] The left hemisphere of the brain was sectioned into 4 blocks (rostral[to caudal). Block 1 was 10 mm; blocks 2 and 3 were 16 mm; and block 4 was 20 mm. Each block was embedded in OCT, snap frozen in isopentane and stored at -80° C. Ten micron sections were then collected and placed onto laser microdissection (LMD) compatible slides. Tissue on the slides was fixed in ethanol (30 seconds in 75% ethanol; 30 seconds in 95% ethanol; and 30 seconds in 100% ethanol) with an air dry between steps. Slides were then screened to identify the ones containing brain regions of interest (i.e. caudate and putamen) and data on viral spread was obtained by following the expression of the marker gene (GFP). GFP positive regions were collected by LMD from the caudate and putamen.

[0087] RNA was isolated from GFP collected material using the PicoPure RNA isolation kit from Arcturus (Mountain View, Calif.). The RNA was quantified prior to the generation of cDNA. Typically about 10-80 ng of total RNA from the LMD collected material was isolated. cDNA was prepared using the Stratascript® QPCR cDNA synthesis kit from Stratagene (La Jolla, Calif.). HD expression was measured by realtime PCR and normalized to GAPDH expression in the same sample. To verify RNA collected from virally transduced cells were being analyzed, GFP realtime PCR to monitor GFP expression was also performed.

[0088] As can be seen in FIG. 21, when measured from samples obtained from tissue punches, in vivo HD gene expression was suppressed in the putamen but not the caudate. The likely reason suppression was not observed in the caudate is that the percent of cells expressing the virus within the collected punch is too low to elicit a detectable level of rhesus HD suppression. This was addressed by collecting GFP+ transduced cells by LMD. When measured from samples obtained through LMD, in vivo HD gene expression was suppressed in the putamen, caudate and corona radiate (FIG. 22; R1=replicate 1; R2=replicate 2). Together, these data demonstrate the feasibility and safety of in vivo HD gene expression in accordance with the methods and sequences of the present invention.

PROPHETIC EXAMPLE 1

Pre-Clinical Test of the Safety of Suppressing Huntingtin Protein Expression in the Rhesus Monkey Brain using shNA of the Invention

[0089] A pre-clinical study of the safety of suppressing huntingtin protein expression in the normal adult rhesus monkey brain is conducted over a 12-month period. During the first three months of the study, six rhesus macaques (*Macaca mulatta*) are trained to perform the tasks of a commercially available computerized behavioral test battery (Monkey CANTAB, Model 80650, Lafayette Instruments,

Lafayette, Ind.) that is based upon human neuropsychological tests (CANTAB, CeNeS, Cambridge, UK). The performance of these six monkeys on the test battery is compared to previously published performance norms for rhesus monkeys to verify that the monkey's cognitive abilities are within normal limits. See, for example, Weed et al., *Brain Research: Cognitive Brain Research*, Oct. 25, 1999; 8(3) 185-201.

[0090] Next, the monkeys are randomly assigned to one of two experimental groups: the first group of three monkeys receives intracranial injections of an adeno-associated virus (AAV) vector comprising an expression cassette containing an RNA polymerase III promoter driving expression of a short hairpin RNA transcript encoding for the shNA of SEQ ID NO: 17. A second expression cassette in the same vector encodes for a green fluorescent protein reporter gene. The second group of three monkeys receives intracranial injections of an adeno-associated virus vector comprising an expression cassette containing the same RNA polymerase III promoter driving expression of a control shNA sequence (SEQ ID NO:18). Also, a second expression cassette in this vector encodes for a green fluorescent protein reporter gene. The AAV vector is administered to the caudate nucleus and putamen of each hemisphere of each monkey's brain. The amount of AAV vector administered in each intracranial injection, and the surgical method and procedures for making these injections, are well-known to those skilled in the art.

[0091] After a one-week post-operative recovery time, the monkeys are repeatedly tested on a periodic basis using the Monkey CANTAB computerized testing battery for the remainder of the twelve-month study period. The cognitive and behavioral performance of each monkey over time is compared to its own pre-operative performance. No statistically significant decline in cognitive or behavioral performance measures is obtained for either the monkeys treated with the control shNA vector, or the monkeys treated with the anti-rhesus-HD shNA vector. This indicates that there are no significant cognitive or behavioral consequences of suppressing the expression of huntingtin protein in the caudate and putamen of the adult primate brain. Periodic neurological examinations of each monkey by skilled and qualified veterinarians unaware of the experimental group membership of each monkey also shows no neurological deficits in the monkeys.

[0092] At the end of the twelve-month study period, the monkeys are humanely terminated, and the brain of each monkey is removed and examined for abnormalities by trained pathologists. In addition, it is studied by fluorescence microscopy, immunohistochemistry, and molecular and biochemical assays. The regions of the brain in which the AAV vector effectively delivered the shNA are identified by fluorescence microscopy. This reveals expression of the green fluorescent protein reporter gene in substantial regions of the caudate nucleus and putamen in each hemisphere of each monkey's brain. Immunohistochemical staining for huntingtin protein, using antibodies and methods well-known to those skilled in the art, shows reduced levels of huntingtin protein in the caudate and putamen of the monkeys who received the anti-rhesus-HD shNA vector, compared to the monkeys who received the control shNA vector, indicating suppression of huntingtin protein expression by the shNA of the invention. Finally, homogenized brain tissue

samples from the caudate and putamen of each monkey's brain are assayed for levels of rhesus HD mRNA measured by quantitative reverse transcriptase polymerase chain reaction (qRT-PCR) assay, and for levels of rhesus huntingtin protein measured by Western blotting, using methods well-known to those skilled in the art. These assays show the expression of HD mRNA and huntingtin protein is significantly reduced in the caudate and putamen of the monkeys who received the anti-rhesus-HD shNA vector, compared to the monkeys who received the control shNA vector. These data further confirm that the shNA vector of the invention suppresses the expression of huntingtin protein in the caudate and putamen of the monkeys receiving the anti-rhesus-HD shNA vector. Because this suppression of huntingtin protein in the caudate and putamen of the rhesus monkeys does not result in corresponding cognitive or behavioral deficits, or other observable neurological abnormalities or brain tissue pathology, the safety of suppressing huntingtin protein in these regions of the adult primate brain is established.

PROPHETIC EXAMPLE 2

Treatment of Huntington's Disease ("HD") in Human Patients using shNA of the Invention

[0093] Once the safety of suppressing huntingtin protein in the caudate and putamen of the adult primate brain is established in rhesus monkeys, the same shNA expression cassette that is used in the pre-clinical safety tests in rhesus monkeys is engineered into an AAV vector using methods well-known to those skilled in the art. This AAV vector comprises an expression cassette containing an RNA polymerase III promoter driving expression of a short hairpin RNA transcript encoding for the shNA of SEQ ID NO: 17. The AAV vector optionally does not contain a second expression cassette encoding for a green fluorescent protein reporter gene, because this expression cassette is not needed for the treatment of HD in patients.

[0094] It is to be especially understood that because the shNA sequence encoded by SEQ ID NO:17 contains the siNA target region of SEQ ID NO: 4, and the sequence of SEQ ID NO: 4 is 100% homologous with corresponding sequences within both the *Macaca mulatta* and the *Homo sapiens* HD gene, an AAV vector comprising an expression cassette containing SEQ ID NO: 17 is effective at suppressing the expression of the *Homo sapiens* HD gene and huntingtin protein, as well as the *Macaca mulatta* HD gene and huntingtin protein.

[0095] Following safety and dose-escalation trials in human patients in which the safety of administering the AAV vector comprising SEQ ID NO: 17 is established, clinical

trials evaluating the clinical efficacy of the invention in treating HD in human patients are conducted. In these trials, the AAV vector can be administered to the human patient's brain according to the systems and methods disclosed in U.S. Patent Application Numbers 20040162255 and 20040220132. These trials establish that suppression of huntingtin protein in the caudate and putamen of human patients afflicted with HD using the nucleic acid molecules, expression cassettes, or vectors of the present invention is effective at arresting and partially reversing the cognitive and motor deficits in patients treated after onset of these symptoms of the disease. These results are consistent with findings obtained in conditional transgenic mouse models of HD, in which expression of a mutant HD transgene results in a disease phenotype, after which suppressing the expression of the mutant HD transgene results in reversal of the disease phenotype and improvement of the mice. Yamamoto et al., *Cell*, Mar. 31, 2000; 101(1):57-66.

[0096] It is to be understood that the present invention is not limited to the particular embodiments, materials, and examples described herein, as these can vary. For example, the nucleic acid molecules of the present invention can be created in a variety of formats and lengths. Indeed, those skilled in the art will recognize that the most important attribute of the nucleic acid molecules of the present invention is their ability to enter the RISC complex and also complementarily bind to the mRNA sequences of interest to induce an RNA interference effect, resulting in reduction of the mRNA stability and/or the translation rate of these sequences. The phrase "complementarily bind" as used herein, refers to the abilities of the nucleic acid molecules to form hydrogen bond(s) with mRNA sequences by either traditional Watson-Crick pairing or other non-traditional types.

[0097] It also is to be understood that the terminology used herein is used for the purpose of describing particular embodiments only, and is not intended to limit the scope of the present invention. It must be noted that as used herein and in the appended claims, the singular forms "a," "an," and "the" include the plural reference unless the context clearly dictates otherwise. Thus, for example, a reference to "a sequence" or "an shNA" is a reference to one or more sequences or shNAs and includes equivalents thereof known to those skilled in the art and so forth.

[0098] Unless defined otherwise, all technical terms used herein have the same meanings as commonly understood by one of ordinary skill in the art to which this invention belongs. Specific methods, devices, and materials are described, although any methods and materials similar or equivalent to those described herein can be used in the practice or testing of the present invention.

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What is claimed is:

1. A nucleic acid molecule comprising a first strand and a second strand wherein said first strand comprises a nucleotide sequence and wherein said second strand comprises the reverse complement of said first strand and wherein said nucleic acid molecule suppresses the expression of both *Macaca mulatta* and *Homo sapiens* mRNA sequences that encode for huntingtin.

2. An isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 19 contiguous nucleotides encoded by the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, and SEQ ID NO: 15 and wherein the second strand is complementary to at least 15 contiguous nucleotides of the first strand.

3. An isolated nucleic acid duplex comprising a first strand of nucleic acid and a second strand of nucleic acid, wherein the first strand comprises at least 27 contiguous nucleotides encoded by the group consisting of SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, and SEQ ID NO: 15 and

wherein the second strand is complementary to at least 23 contiguous nucleotides of the first strand.

4. A nucleic acid duplex according to claim 2, wherein said duplex is between 19 and 30 base pairs in length.

5. A nucleic acid duplex according to claim 2 or 3, wherein said first and/or said second strand further comprises an overhang region.

6. A nucleic acid duplex according to claim 2 or 3, wherein said first and/or second strand further comprises a 3' overhang region, a 5' overhang region, or both 3' and 5' overhang regions.

7. A nucleic acid duplex according to claim 5, wherein said overhang region is from 1 to 10 nucleotides in length.

8. A nucleic acid duplex according to claim 2 or 3, wherein said first strand and said second strand are operably linked by means of a nucleic acid loop strand to form a hairpin structure comprising a duplex structure and a loop structure.

9. A nucleic acid duplex according to claim 8, wherein said loop structure contains from 4 to 13 nucleotides.

10. An expression cassette comprising a nucleic acid encoding at least one strand of the nucleic acid duplex of claim 2 or 3.

11. An expression cassette according to claim 10, further comprising a promoter.

12. An expression cassette according to claim 11, wherein said promoter is a regulatable promoter.

13. An expression cassette according to claim 11, wherein said promoter is a constitutive promoter.

14. An expression cassette according to claim 11, wherein said promoter is a CMV, RSV, pol II or pot III promoter.

15. An expression cassette according to claim 10, wherein said expression cassette further comprises a polyadenylation signal.

16. An expression cassette according to claim 15, wherein said polyadenylation signal is a synthetic minimal polyadenylation signal.

17. An expression cassette according to claim 10, further comprising a marker gene.

18. A vector comprising the expression cassette of claim 10.

19. A vector comprising two expression cassettes, a first expression cassette comprising a nucleotide sequence encoding said first strand of said nucleic acid duplex of claim 2 and a second expression cassette comprising a nucleotide sequence encoding the second strand of the nucleic acid duplex of claim 2.

20. A vector comprising two expression cassettes, a first expression cassette comprising a nucleotide sequence encoding said first strand of said nucleic acid duplex of claim 3 and a second expression cassette comprising a nucleotide sequence encoding the second strand of the nucleic acid duplex of claim 3.

21. A vector according to claim 19 or 20, wherein said vector is a viral vector.

22. A vector according to claim 21, wherein said vector is an adenoviral, lentiviral, adeno-associated viral (AAV), poliovirus, herpes simplex virus, feline immunodeficiency virus or murine Maloney-based viral vector.

23. A vector according to claim 19 or 20, wherein said vector further comprises a promoter.

24. A vector according to claim 23, wherein said promoter is an inducible promoter.

25. A cell comprising the expression cassette of claim 10.

26. A cell of claim 25, wherein said cell is a mammalian cell.

27. A non-human mammal comprising the expression cassette of claim 10.

28. A method of suppressing the accumulation of huntingtin in a cell comprising:

introducing a nucleic acid duplex of claim 2 or 3 into said cell in an amount sufficient to suppress accumulation of huntingtin in said cell.

29. A method according to claim 28, wherein said accumulation of said huntingtin is suppressed by at least 10%.

30. A method of preventing cytotoxic effects of mutant huntingtin in a cell comprising:

introducing a nucleic acid duplex of claim 2 or 3 into said cell in an amount sufficient to suppress accumulation of said mutant huntingtin and wherein said nucleic acid duplex prevents cytotoxic effects of said mutant huntingtin in said cell.

31. A method to inhibit expression of a huntingtin gene in a cell comprising:

introducing a nucleic acid duplex of claim 2 or 3 into said cell in an amount sufficient to inhibit expression of said huntingtin protein, and wherein said nucleic acid duplex inhibits expression of said huntingtin gene.

32. A method to inhibit expression of huntingtin in *Macaca mulatta* and *Homo sapiens* comprising:

providing a *Macaca mulatta* or *Homo sapiens* containing a neuronal cell, wherein said neuronal cell contains the huntingtin gene and said neuronal cell is susceptible to nucleic acid interference and said huntingtin gene is expressed in said neuronal cell; and

contacting said *Macaca mulatta* or *Homo sapiens* with a nucleic acid duplex of claim 2 or 3, thereby inhibiting expression of said huntingtin gene.

33. A method according to claim 32 wherein expression of said huntingtin is inhibited by at least 10%.

34. A method of preventing cytotoxic effects of Huntington's disease ("HD") in a *Macaca mulatta* or *Homo sapiens* comprising:

introducing the vector of claim 18 into a cell in an amount sufficient to suppress accumulation of a protein associated with HD, and wherein the nucleic acid duplex prevents cytotoxic effects of HD.

35. A method to inhibit expression of huntingtin in a *Macaca mulatta* or *Homo sapiens* comprising:

introducing the vector of claim 18 into a cell in an amount sufficient to inhibit expression of huntingtin wherein said nucleic acid duplex inhibits expression of the protein associated with HD.

36. A method to inhibit expression of huntingtin in a *Macaca mulatta* or *Homo sapiens* comprising:

providing a *Macaca mulatta* or *Homo sapiens* containing a neuronal cell, wherein said neuronal cell contains the huntingtin gene and said neuronal cell is susceptible to nucleic acid interference, and the huntingtin gene is expressed in said neuronal cell; and

contacting said *Macaca mulatta* or *Homo sapiens* with the vector of claim 19 or 20, thereby inhibiting expression of the huntingtin gene.

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