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(56) Fremdragne publikationer:
WO-A2-2005/001043
WO-A2-2009/044392
UI-TEI KUMIKO ET AL: "Functional dissection of siRNA sequence by systematic DNA substitution: modified siRNA with a DNA seed arm is a powerful tool for mammalian gene silencing with significantly reduced off-target effect" NUCLEIC ACIDS RESEARCH, vol. 36, no. 7, 1 April 2008 (2008-04-01), pages 2136-2151, XP002598044 ISSN: 0305-1048
HUANG HUANG ET AL: "Profiling of mismatch discrimination in RNAi enabled rational design of allele-specific siRNAs" NUCLEIC ACIDS RESEARCH, vol. 37, no. 22, 8 October 2009 (2009-10-08), pages 7560-7569, XP002624726 ISSN: 0305-1048
DU QUAN ET AL: "A systematic analysis of the silencing effects of an active siRNA at all single-nucleotide mismatched target sites" NUCLEIC ACIDS RESEARCH, vol. 33, no. 5, 2005, pages 1671-1677, XP002624727 ISSN: 0305-1048
UI-TEI K ET AL: "Guidelines for the selection of highly effective siRNA sequences for mammalian and chick RNA interference" NUCLEIC ACIDS RESEARCH, vol. 32, no. 3, 9 February 2004 (2004-02-09), pages 936-948, XP002329955 ISSN: 0305-1048
GRIMSON ANDREW ET AL: "MicroRNA targeting specificity in mammals: Determinants beyond seed pairing" MOLECULAR CELL, vol. 27, no. 1, July 2007 (2007-07), pages 91-105, XP002624728 ISSN: 1097-2765 -&
GRIMSON ANDREW ET AL: "Supplemental Data: MicroRNA Targeting Specificity in Mammals: Determinants

Beyond Seed Pairing" MOLECULAR CELL, [Online] vol. 27, July 2007 (2007-07), pages 1-24, XP002628017
Sciencedirect Retrieved from the Internet: URL:<http://www.sciencedirect.com/science/MiamiMultiMediaURL/B6WSR-4P48CFV-9/B6WSR-4P 48CFV-9-1/7053/html/S1097276507004078/475133524aac81636d402c7400703e69/mmc1.pdf>

DESCRIPTION

FIELD OF THE INVENTION

[0001] The present invention relates to modified siRNA compounds, pharmaceutical compositions comprising same and these compounds for use in methods for the inhibition of gene expression. The compounds and compositions exhibit beneficial properties including knock down activity of target genes and are useful in the treatment of subjects suffering from diseases or conditions and or symptoms associated with such diseases or conditions or at risk of contracting diseases or conditions in which gene expression has adverse consequences.

BACKGROUND OF THE INVENTION

[0002] PCT Patent Publication Nos. WO 2008/104978 and WO 2009/044392 to the assignee of the present invention disclose novel siRNA structures.

[0003] Ui-Tei et al (Nucleic Acids Research, vol. 36 (2008), pages 2136-2151) discloses substitution of ribonucleotides in siRNA with their cognate deoxyribonucleotides.

[0004] Huang et al (Nucleic Acids Research, vol. 37, (2009), pages 7560-7569) discloses single point mutations at internal positions of the antisense strand of siRNAs.

[0005] Du et al (Nucleic Acids Research, vol. 33, (2005), pages 1671-1677) discloses mismatches between the antisense strand of an siRNA and its target site, and recommends the formation of G:U wobble base pair.

[0006] WO 2005/001043 discloses methods and compositions for enhancing the efficacy and specificity of RNAi and recommends lessening the base pair strength between the 5' end of the antisense strand and the 3' end of the sense strand, by introducing for example mismatched base pairs in the duplex.

[0007] Ui-Tei et al (Nucleic Acids Research, vol. 32, (2004), pages 936-948) discloses guidelines for the selection of highly effective siRNA sequences for RNA interference, and more specifically how to select a target sequence in a gene, such that the siRNA targeting this selected sequence can be predicted to be highly effective.

[0008] There remains a need for active and effective siRNA therapeutic agents which exhibit enhanced knock down activity, increased stability and or reduced off target effects.

SUMMARY OF THE INVENTION

[0009] The double stranded RNA (dsRNA) compounds disclosed herein possess structures and modifications which may, for example increase activity, increase stability, reduce immunogenicity,

enhance loading into the RISC complex, and or minimize toxicity; the novel modifications of the siRNAs are beneficially applied to double stranded RNA useful in preventing or attenuating target gene expression.

[0010] Provided herein are double stranded (duplex) oligonucleotide compounds useful for the down regulation of gene expression. The present disclosure is based in part on the unexpected observation that a double stranded nucleic acid molecule comprising a sense strand and a complementary antisense strand and having a mismatch between the 5' terminal nucleotide of the antisense strand and the nucleotide of the target RNA show potent activity in down regulating target genes. In preferred embodiments the dsRNA comprises an adenosine, deoxyadenosine or deoxyuridine, substituted at position 1 (5' terminus) of the antisense strand rather in place of a cytidine or guanine.

[0011] According to one aspect disclosed are modified double stranded nucleic acid molecules having structure (A) set forth below:

(A)	5'	$N^1-(N)x - Z$	3' (antisense strand)
	3'	$Z'-N^2-(N')y - z''$	5' (sense strand)

wherein each of N^2 , N and N' is an unmodified or modified ribonucleotide, or an unconventional moiety;

wherein each of $(N)x$ and $(N')y$ is an oligonucleotide in which each consecutive N or N' is joined to the adjacent N or N' by a covalent bond;

wherein each of x and y is independently an integer between 17 and 39;

wherein the sequence of $(N')y$ has complementarity to the sequence of $(N)x$ and $(N)x$ has complementarity to a consecutive sequence in a target RNA;

wherein N^1 is covalently bound to $(N)x$ and is mismatched to the target RNA or is a complementary DNA moiety to the target RNA;

wherein N^1 is a moiety selected from the group consisting of natural or modified uridine, deoxyribouridine, ribothymidine, deoxyribothymidine, adenosine or deoxyadenosine;

wherein z'' may be present or absent, but if present is a capping moiety covalently attached at the 5' terminus of $N^2-(N')y$; and

wherein each of Z and Z' is independently present or absent, but if present is independently 1-5 consecutive nucleotides, consecutive non-nucleotide moieties or a combination thereof covalently attached at the 3' terminus of the strand in which it is present.

[0012] More specifically, the invention is directed to a double stranded nucleic acid molecule having the structure set forth below:

(A)	5'	$N'-(N)x - Z$	3' (antisense strand)
	3'	$Z'-N^2-(N')y - z''$	5' (sense strand)

wherein each of N², N and N' is an unmodified or modified ribonucleotide, or an unconventional moiety;

wherein each of (N)x and (N')y is an oligonucleotide in which each consecutive N or N' is joined to the adjacent N or N' by a covalent bond;

wherein each of x=y=18;

wherein the sequence of (N)x has complementarity to a consecutive sequence in a target RNA; and the sequence of N²-(N')y is fully complementary to the sequence of N'-(N)x;

wherein N¹ is covalently bound to (N)x and mismatched to the target RNA;

wherein:

when the pairing nucleotide in the target RNA is adenosine, uridine in position 1 of the antisense strand is substituted with an N¹ selected from adenosine or deoxyadenosine; or

when the pairing nucleotide in the target RNA is cytidine, guanosine in position 1 of the antisense strand is substituted with an N¹ selected from adenosine, deoxyadenosine, or deoxyuridine; or

when the pairing nucleotide in the target RNA is guanosine, cytidine in position 1 of the antisense strand is substituted with an N¹ selected from adenosine or deoxyadenosine; or when the pairing nucleotide in the target RNA is uridine, adenosine in position 1 of the antisense strand is substituted with an N¹ which is deoxyuridine;

wherein z" may be present or absent, but if present is a capping moiety covalently attached at the 5' terminus of N²-(N')y; and

wherein each of Z and Z' is independently present or absent, but if present is independently 1-5 consecutive nucleotides, consecutive non-nucleotide moieties or a combination thereof covalently attached at the 3' terminus of the strand in which it is present.

[0013] The sequence of (N')y is fully complementary to the sequence of (N)x. The sequence of N²-(N')y is complementary to the sequence of N¹-(N)x. In some embodiments (N)x comprises an antisense that is fully complementary to about 17 to about 18 consecutive nucleotides in a target RNA. In other embodiments (N)x comprises an antisense that is substantially complementary to about 17 to about 18 consecutive nucleotides in a target RNA.

[0014] In some embodiments N¹ and N² form a Watson-Crick base pair. In some embodiments N¹ and N² form a non-Watson-Crick base pair. In some embodiments a base pair is formed between a ribonucleotide and a deoxyribonucleotide.

[0015] x=y=18.

[0016] N¹ is covalently bound to (N)x and is mismatched to the target RNA.

[0017] In some embodiments a uridine in position 1 of the antisense strand is substituted with an N¹ selected from adenosine or deoxyadenosine.

[0018] In some embodiments guanosine in position 1 of the antisense strand is substituted with an N¹ selected from adenosine, deoxyadenosine or deoxyuridine.

[0019] In some embodiments cytidine in position 1 of the antisense strand is substituted with an N¹ selected from adenosine or deoxyadenosine.

[0020] In some embodiments adenosine in position 1 of the antisense strand is substituted with an N¹ which is deoxyuridine.

[0021] In some embodiments N¹ and N² form a base pair between uridine or deoxyuridine, and adenosine or deoxyadenosine. In other embodiments N¹ and N² form a base pair between deoxyuridine and adenosine.

[0022] In some embodiments the double stranded nucleic acid molecule is a siRNA, siNA or a miRNA.

[0023] The following table provides examples of N1 and corresponding N2.

Target nucleotide	5' terminal nucleotide of AS with full match to target	N1 (5' terminal position of AS)	N2 (3' terminal position of SEN)
A	U	rA, dA	rU, dU, rT, dT
C	G	rA, dA	rU, dU, rT, dT
C	G	dU	rA, dA
G	C	rA, dA	rU, dU, rT, dT
U	A	dU	rA, dA

[0024] In some embodiments of Structure (A), N¹ comprises adenosine. In certain embodiments of structure (A), N² comprises a 2'OMe sugar modified ribonucleotide. In some embodiments of Structure (A), N¹ comprises adenosine and N² comprises a ribouridine or modified ribouridine.

[0025] In some embodiments Z and Z' are absent. In other embodiments one of Z or Z' is present.

[0026] In some embodiments each of N and N' is an unmodified ribonucleotide. In some embodiments at least one of N or N' comprises a chemically modified ribonucleotide or an unconventional moiety. In some embodiments the unconventional moiety is selected from a mirror nucleotide, an abasic ribose moiety and an abasic deoxyribose moiety. In some embodiments the unconventional moiety is a mirror nucleotide, preferably an L-DNA moiety. In some embodiments at least one of N or N' comprises a 2'OMe sugar-modified ribonucleotide.

[0027] The sequence of (N')y is fully complementary to the sequence of (N)x.

[0028] In some embodiments (N)x comprises an antisense sequence that is fully complementary to about 17 to about 18 consecutive nucleotides in a target mRNA. In other embodiments (N)x comprises an antisense that is substantially complementary to about 17 to about 18 consecutive nucleotides in a target mRNA. x=y=18.

[0029] According to certain preferred embodiments siRNA compounds comprising one or more modified nucleotide, wherein the modified nucleotide possesses a modification in the sugar moiety, in the base moiety or in the internucleotide linkage moiety are provided.

[0030] Structure (A) motif is useful with any oligonucleotide pair (sense and antisense strands) to a mammalian or non-mammalian gene. In some embodiments the mammalian gene is a human gene.

[0031] In some embodiments a modified siRNA compound having structure (A) exhibits beneficial properties including enhanced activity (e.g. reduced IC50, increased knock down, reduced residual mRNA) when compared to a control compound, i.e. an siRNA compound wherein the antisense oligonucleotide is fully complementary (including 5' terminal nucleotide base paired e.g. A-U, U-A, C-G, G-C) to a nucleotide sequence in a target mRNA. In some embodiments the activity is enhanced by at least 5%, by at least 10%, by at least 20%, by at least 25% or more when compared to a control compound.

[0032] In another aspect the present invention provides a method of generating a double stranded RNA molecule consisting of a sense strand and an antisense strand comprising the steps of

1. a) selecting a consecutive 17 to 25 nucleotide sequence in a target RNA and synthesizing an antisense strand comprising complementarity to the consecutive 17 to 25 nucleotide sequence of the target mRNA wherein the 5' terminal nucleotide of the antisense strand, if uridine, is substituted with adenosine or deoxyadenosine; or if guanosine, is substituted with adenosine, deoxyadenosine, or deoxyuridine; or if cytidine, is substituted with adenosine or, deoxyadenosine,; or if adenosine, is substituted with a deoxyuridine;
2. b) synthesizing a sense strand of 17 to 25 nucleotides having complementarity to the antisense strand, wherein the 3' terminal nucleotide of the sense strand forms a Watson Crick base pair with the 5' terminal nucleotide of the guide strand; and
3. c) annealing the antisense and sense strands; thereby generating a double stranded RNA molecule.

[0033] According to one aspect, the present description discloses a method of generating a double stranded RNA molecule consisting of a sense strand and an antisense strand exhibiting enhanced RNAi activity when compared to an unmodified a double stranded RNA molecule comprising the steps of

1. a) selecting a consecutive 17 to 25 nucleotide sequence in a target mRNA and synthesizing a sense strand comprising the consecutive 17 to 25 nucleotide sequence of the target mRNA wherein the 3' terminal nucleotide is substituted with adenosine, modified adenosine, deoxyadenosine or modified deoxyadenosine;
2. b) synthesizing an antisense strand of 17 to 25 nucleotides having complementarity to the

sense strand wherein the 5' terminal nucleotide comprises ribouridine, modified ribouridine, deoxyribouridine or modified deoxyribouridine and base pairs with the 3' terminal nucleotide of the passenger strand;

3. c) annealing the sense strand to the antisense strand;

thereby generating a double stranded RNA molecule having enhanced RNAi activity.

[0034] In some embodiments the modified double stranded RNA molecule exhibits enhanced RNAi activity when compared to an unmodified siRNA duplex, i.e. a duplex having full match to the target mRNA.

[0035] According to another aspect, the present description discloses a method of generating a modified a double stranded RNA molecule consisting of a sense strand and antisense strand exhibiting enhanced RNAi activity when compared to an unmodified a double stranded RNA molecule comprising the steps of

1. a) selecting a consecutive 17 to 25 nucleotide sequence in a target mRNA and synthesizing a sense strand comprising the consecutive 17 to 25 nucleotide sequence of the target mRNA wherein the 3' terminal nucleotide is substituted with adenosine, modified adenosine, deoxyadenosine or modified deoxyadenosine;
2. b) synthesizing an antisense strand of 17 to 25 nucleotides having complementarity to the sense strand wherein the 5' terminal nucleotide comprises ribouridine, modified ribouridine, deoxyribouridine or modified deoxyribouridine and base pairs with the 3' terminal nucleotide of the sense strand;
3. c) annealing the sense strand to the antisense strand;

thereby generating a double stranded RNA molecule having enhanced RNAi activity.

[0036] In some embodiments step a) includes selecting a consecutive 17 to 25 nucleotide, or 17, 18, 19, 20 , 21, 22, 23, 24 or 25 nucleotide sequence in a target RNA in a target cell wherein the 3' terminal nucleotide is other than adenosine.

[0037] In another aspect provided is a pharmaceutical composition comprising a compound according to Structure (A), in an amount effective to inhibit mammalian or non-mammalian gene expression; and a pharmaceutically acceptable carrier. In some embodiments the mammalian gene is a human gene. In some embodiments the non-mammalian gene is involved in a mammalian disease, preferably human disease.

[0038] Further provided are compounds for use in methods for treating or preventing the incidence or severity of a disease or condition and/or for reducing the risk or severity of a disease or condition in a subject in need thereof wherein the disease or condition and/or a symptom and/or risk associated therewith is associated with expression of a mammalian or a non-mammalian gene. In a preferred embodiment the subject is a human subject.

[0039] In some embodiments the disease or condition is selected from the group hearing loss, acute renal failure (ARF), Delayed Graft Function (DGF) after kidney transplantation, glaucoma, ocular ischemic conditions including anterior ischemic optic neuropathy, age-related macular degeneration (AMD), Ischemic Optic Neuropathy (ION), dry eye syndrome, acute respiratory distress syndrome

(ARDS) and other acute lung and respiratory injuries, chronic obstructive pulmonary disease (COPD), primary graft failure, ischemia-reperfusion injury, reperfusion injury, reperfusion edema, allograft dysfunction, pulmonary reimplantation response and/or primary graft dysfunction (PGD) after organ transplantation, in particular in lung transplantation, organ transplantation including lung, liver, heart, pancreas, and kidney transplantation, nephro- and neurotoxicity, spinal cord injury, brain injury, neurodegenerative disease or condition, pressure sores, oral mucositis fibrotic conditions including liver fibrosis, lung fibrosis; and cancer. Such methods involve administering to a mammal in need of such treatment a prophylactically or therapeutically effective amount of one or more such compounds, which inhibit or reduce expression or activity of at least one such gene.

[0040] The compounds, methods, materials, and examples that will now be described are illustrative only and are not intended to be limiting; materials and methods similar or equivalent to those described herein can be used in practice or testing of the invention. Other features and advantages of the invention will be apparent from the following detailed description, and from the claims.

BRIEF DESCRIPTION OF THE FIGURES

[0041]

Figure 1 is a diagram of the dsRNA molecules as provided herein. In one embodiment, the 5' terminal G or C nucleotide of the antisense strand is substituted with U, dU, rA, dA, rT, dT; or the 5' terminal U nucleotide of the antisense strand is substituted with dU, rA, dA, rT, dT; or the 5' terminal A nucleotide of the antisense strand is substituted with U, dU, dA, rT, dT. In one embodiment the 5' terminal nucleotide of the antisense strand is mismatched to the target mRNA. In another embodiment the 5' terminal nucleotide of the antisense strand is a deoxyribonucleotide complementary to the target mRNA.

Figures 2-10 show activity (% residual target remaining) bar graphs of dsRNA molecules with different nucleotides in position 1 of the antisense strand to targets that have been modified to individually include the four ribonucleotides at position 19 (position 1 of AS).

DETAILED DESCRIPTION OF THE INVENTION

[0042] The present invention relates generally to oligonucleotide compounds which down-regulate expression of various genes, particularly to small interfering RNAs (siRNA), specifically to modified siRNA compounds and to the use of these modified siRNA compounds in preparation of pharmaceutical compositions and in treatment of a subject suffering from various medical conditions.

[0043] The compounds and compositions exhibit beneficial properties including potent knock down activity of target genes and are useful in the treatment of subjects suffering from diseases or conditions and or symptoms associated with such diseases or conditions or at risk of contracting diseases or conditions in which gene expression has adverse consequences.

[0044] Accordingly, in certain aspects modified siRNA compounds and pharmaceutical compositions

comprising same useful in down regulating gene expression are provided.

[0045] In another aspect, the present description discloses a method of treating a subject suffering from or susceptible to a microvascular disorder, eye disease or disorder, hearing impairment (including hearing loss), a respiratory (including pulmonary) disorder, neurodegenerative disease or disorder, spinal cord injury, brain injury angiogenesis- and apoptosis-related conditions comprising administering to the subject a pharmaceutical composition comprising at least one small interfering RNA of the invention that targets a mammalian or non-mammalian gene, in an amount sufficient to down-regulate the expression of the gene.

[0046] In certain embodiments, the subject compounds are useful in inhibiting expression of a target gene for treatment of *inter alia*, respiratory disorders, microvascular disorders or eye disorders. Particular diseases and conditions to be treated are ARDS; COPD; ALI; Emphysema; Diabetic Neuropathy, nephropathy and retinopathy; DME and other diabetic conditions; Glaucoma; AMD; BMT retinopathy; ischemic conditions including stroke; OIS; neurodegenerative disorders such as Parkinson's, Alzheimer's, ALS; kidney disorders: ARF, DGF, transplant rejection; hearing disorders; spinal cord injuries; oral mucositis; cancer, dry eye syndrome and pressure sores.

Definitions

[0047] For convenience certain terms employed in the specification, examples and claims are described herein.

[0048] It is to be noted that, as used herein, the singular forms "a", "an" and "the" include plural forms unless the content clearly dictates otherwise. Where aspects or embodiments of the invention are described in terms of Markush groups or other grouping of alternatives, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the group.

[0049] An "inhibitor" is a compound, which is capable of reducing (partially or fully) the expression of a gene or the activity of the product of such gene to an extent sufficient to achieve a desired biological or physiological effect. The term "inhibitor" as used herein refers to a siRNA inhibitor. A "siRNA inhibitor" is a compound that is capable of reducing the expression of a gene or the activity of the product of such gene to an extent sufficient to achieve a desired biological or physiological effect. The term "siRNA inhibitor" as used herein refers to one or more of a siRNA, shRNA, siNA, synthetic shRNA; miRNA. Inhibition may also be referred to as down-regulation or, for RNAi, silencing.

[0050] The term "inhibit" as used herein refers to reducing the expression of a gene or the activity of the product of such gene to an extent sufficient to achieve a desired biological or physiological effect. Inhibition is either complete or partial

[0051] As used herein, the term "inhibition" of a target gene means inhibition of the gene expression (transcription or translation) or polypeptide activity of the product of a target gene. "Gene product" as used herein, refers to a product of a gene such as an RNA transcript or a polypeptide. The terms "RNA transcript", "mRNA polynucleotide sequence", "mRNA sequence" and "mRNA" are used interchangeably.

[0052] As used herein, the terms "polynucleotide" and "nucleic acid" may be used interchangeably and refer to nucleotide sequences comprising deoxyribonucleic acid (DNA), and ribonucleic acid (RNA). The terms are to be understood to include, as equivalents, analogs of either RNA or DNA made from nucleotide analogs. Throughout this disclosure, mRNA sequences are set forth as representing the corresponding genes.

[0053] "Oligonucleotide" or "oligomer" refers to a deoxyribonucleotide or ribonucleotide sequence from about 2 to about 50 nucleotides. Each DNA or RNA nucleotide may be independently natural or synthetic, and or modified or unmodified. Modifications include changes to the sugar moiety, the base moiety and or the linkages between nucleotides in the oligonucleotide. The compounds of the present invention encompass molecules comprising deoxyribonucleotides, ribonucleotides, modified deoxyribonucleotides, modified ribonucleotides, nucleotide analogues, modified nucleotide analogues, unconventional and abasic moieties and combinations thereof.

[0054] Substantially complementary refers to complementarity of greater than about 84%, to another sequence. For example in a duplex region consisting of 19 base pairs one mismatch results in 94.7% complementarity, two mismatches results in about 89.5% complementarity and 3 mismatches results in about 84.2% complementarity, rendering the duplex region substantially complementary. Accordingly substantially identical refers to identity of greater than about 84%, to another sequence.

[0055] "Nucleotide" is meant to encompass deoxyribonucleotides and ribonucleotides, which may be natural or synthetic and modified or unmodified. Nucleotides include known nucleotide analogues, which are synthetic, naturally occurring, and non-naturally occurring. Examples of such analogs include, without limitation, phosphorothioates, phosphoramidites, methyl phosphonates, chiral-methyl phosphonates, 2'-O-methyl ribonucleotides, and peptide-nucleic acids (PNAs). Modifications include changes to the sugar moiety, the base moiety and or the linkages between ribonucleotides in the oligoribonucleotide. As used herein, the term "ribonucleotide" encompasses natural and synthetic, unmodified and modified ribonucleotides and ribonucleotide analogues which are synthetic, naturally occurring, and non-naturally occurring. Modifications include changes to the sugar moiety, to the base moiety and/or to the linkages between ribonucleotides in the oligonucleotide.

[0056] The nucleotides are selected from naturally occurring or synthetic modified bases. Naturally occurring bases include adenine, guanine, cytosine, thymine and uracil. Modified bases of nucleotides include inosine, xanthine, hypoxanthine, 2- aminoadenine, 6-methyl, 2-propyl and other alkyl adenines, 5-halouracil, 5-halocytosine, 6-azacytosine and 6-az thymine, pseudouracil, deoxypseudouracil, 4-thiouracil, ribo-2-thiouridine, ribo-4-thiouridine, 8-haloadenine, 8-aminoadenine, 8-thioladenine, 8-thiolalkyl adenines, 8-hydroxyl adenine and other 8-substituted adenines, 8-haloguanines, 8-aminoguanine, 8-thioguanine, 8-thioalkylguanines 8-hydroxylguanine and other substituted guanines, other aza and deaza adenines, other aza and deaza guanines, 5-methylribouridine, 5-trifluoromethyl uracil, 5-methylribocytosine, and 5-trifluorocytosine. In some embodiments one or more nucleotides in an oligomer is substituted with inosine.

[0057] In some embodiments the siRNA compound further comprises at least one modified ribonucleotide selected from the group consisting of a ribonucleotide having a sugar modification, a base modification or an internucleotide linkage modification and may contain DNA, and modified nucleotides such as LNA (locked nucleic acid), ENA (ethylene-bridged nucleic acid), PNA (peptide nucleic acid), arabinoside, phosphonocarboxylate or phosphinocarboxylate nucleotide (PACE

nucleotide), or nucleotides with a 6 carbon sugar.

[0058] Modified deoxyribonucleotide includes, for example 5'OMe DNA (5-methyl-deoxyriboguanosine-3'-phosphate) which may be useful as a nucleotide in the 5' terminal position (position number 1); PACE (deoxyriboadenosine 3' phosphonoacetate, deoxyribocytidine 3' phosphonoacetate, deoxyriboguanosine 3' phosphonoacetate, deoxyribothymidine 3' phosphonoacetate).

[0059] Bridged nucleic acids include LNA (2'-O, 4'-C-methylene bridged Nucleic Acid adenosine 3' monophosphate, 2'-O,4'-C-methylene bridged Nucleic Acid 5-methyl-cytidine 3' monophosphate, 2'-O,4'-C-methylene bridged Nucleic Acid guanosine 3' monophosphate, 5-methyl-uridine (or thymidine) 3' monophosphate); and ENA (2'-O,4'-C-ethylene bridged Nucleic Acid adenosine 3' monophosphate, 2'-O,4'-C-ethylene bridged Nucleic Acid 5-methyl-cytidine 3' monophosphate, 2'-O,4'-C-ethylene bridged Nucleic Acid guanosine 3' monophosphate, 5-methyl-uridine (or thymidine) 3' monophosphate).

[0060] All analogs of, or modifications to, a nucleotide / oligonucleotide are employed with the present invention, provided that said analog or modification does not substantially adversely affect the properties, e.g. function, of the nucleotide / oligonucleotide. Acceptable modifications include modifications of the sugar moiety, modifications of the base moiety, modifications in the internucleotide linkages and combinations thereof.

[0061] A sugar modification includes a modification on the 2' moiety of the sugar residue and encompasses amino, fluoro, alkoxy (e.g. methoxy), alkyl, amino, fluoro, chloro, bromo, CN, CF, imidazole, carboxylate, thioate, C1 to C10 lower alkyl, substituted lower alkyl, alkaryl or aralkyl, OCF₃, OCN, O-, S-, or N- alkyl; O-, S-, or N-alkenyl; SOCH₃; SO₂CH₃; ONO₂; NO₂, N₃; heterocycloalkyl; heterocycloalkaryl; aminoalkylamino; polyalkylamino or substituted silyl, as, among others, described in European patents EP 0 586 520 B1 or EP 0 618 925 B1.

[0062] In one embodiment the modified siRNA compound comprises at least one ribonucleotide comprising a 2' modification on the sugar moiety ("2' sugar modification"). In certain embodiments the siRNA compound comprises 2'O-alkyl or 2'-fluoro or 2'O-allyl or any other 2' modification, optionally on alternate positions. Other stabilizing modifications are also possible (e.g. terminal modifications). In some embodiments a preferred 2'O-alkyl is 2'O-methyl (methoxy) sugar modification.

[0063] In some embodiments the backbone of the oligonucleotides is modified and comprises phosphate-D-ribose entities but may also contain thiophosphate-D-ribose entities, triester, thioate, 2'-5' bridged backbone (also may be referred to as 5'-2'), PACE and the like.

[0064] As used herein, the terms "non-pairing nucleotide analog" means a nucleotide analog which comprises a non-base pairing moiety including but not limited to: 6 des amino adenosine (Nebularine), 4-Me-indole, 3-nitropyrrole, 5-nitroindole, Ds, Pa, N3-Me riboU, N3-Me riboT, N3-Me dC, N3-Me-dT, N1-Me-dG, N1-Me-dA, N3-ethyl-dC, N3-Me dC. In some embodiments the non-base pairing nucleotide analog is a ribonucleotide. In other embodiments the non-base pairing nucleotide analog is a deoxyribonucleotide. In addition, analogues of polynucleotides may be prepared wherein the structure of one or more nucleotide is fundamentally altered and better suited as therapeutic or experimental reagents. An example of a nucleotide analogue is a peptide nucleic acid (PNA) wherein

the deoxyribose (or ribose) phosphate backbone in DNA (or RNA) is replaced with a polyamide backbone which is similar to that found in peptides. PNA analogues have been shown to be resistant to enzymatic degradation and to have enhanced stability in vivo and in vitro. Other modifications include polymer backbones, cyclic backbones, acyclic backbones, thiophosphate-D-ribose backbones, triester backbones, thioate backbones, 2'-5' bridged backbone, artificial nucleic acids, morpholino nucleic acids, glycol nucleic acid (GNA), threose nucleic acid (TNA), arabinoside, and mirror nucleoside (for example, beta-L-deoxyribonucleoside instead of beta-D-deoxyribonucleoside). Examples of siRNA compounds comprising LNA nucleotides are disclosed in Elmen et al., (NAR 2005, 33(1):439-447).

[0065] In some embodiments the compounds of the present invention are synthesized with one or more inverted nucleotides, for example inverted thymidine or inverted adenosine (see, for example, Takei, et al., 2002, JBC 277(26):23800-06).

[0066] Other modifications include 3' terminal modifications also known as capping moieties. Such terminal modifications are selected from a nucleotide, a modified nucleotide, a lipid, a peptide, a sugar and inverted abasic moiety. Such modifications are incorporated, for example at the 3' terminus of the sense and/or antisense strands.

[0067] What is sometimes referred to in the present invention as an "abasic nucleotide" or "abasic nucleotide analog" is more properly referred to as a pseudo-nucleotide or an unconventional moiety. A nucleotide is a monomeric unit of nucleic acid, consisting of a ribose or deoxyribose sugar, a phosphate, and a base (adenine, guanine, thymine, or cytosine in DNA; adenine, guanine, uracil, or cytosine in RNA). A modified nucleotide comprises a modification in one or more of the sugar, phosphate and or base. The abasic pseudo-nucleotide lacks a base, and thus is not strictly a nucleotide.

[0068] The term "capping moiety" as used herein includes abasic ribose moiety, abasic deoxyribose moiety, modifications abasic ribose and abasic deoxyribose moieties including 2' O alkyl modifications; inverted abasic ribose and abasic deoxyribose moieties and modifications thereof; C6-imino-Pi; a mirror nucleotide including L-DNA and L-RNA; 5'O-Me nucleotide; and nucleotide analogs including 4',5'-methylene nucleotide; 1-(β -D-erythofuranosyl)nucleotide; 4'-thionucleotide, carbocyclic nucleotide; 5'-amino-alkyl phosphate; 1,3-diamino-2-propyl phosphate, 3-aminopropyl phosphate; 6-aminohexyl phosphate; 12-aminododecyl phosphate; hydroxypropyl phosphate; 1,5-anhydrohexitol nucleotide; alpha-nucleotide; threo-pentofuranosyl nucleotide; acyclic 3',4'-seco nucleotide; 3,4-dihydroxybutyl nucleotide; 3,5-dihydroxypentyl nucleotide, 5'-5'-inverted abasic moiety; 1,4-butanediol phosphate; 5'-amino; and bridging or non bridging methylphosphonate and 5'-mercapto moieties.

[0069] Certain preferred capping moieties are abasic ribose or abasic deoxyribose moieties; inverted abasic ribose or abasic deoxyribose moieties; C6-amino-Pi; a mirror nucleotide including L-DNA and L-RNA.

[0070] A "hydrocarbon moiety or derivative thereof" refers to straight chain or branched alkyl moieties and moieties *per se* or further comprising a functional group including alcohols, phosphodiester, phosphorothioate, phosphonoacetate and also includes amines, carboxylic acids, esters, amides aldehydes. "Hydrocarbon moiety" and "alkyl moiety" are used interchangeably.

[0071] "Terminal functional group" includes halogen, alcohol, amine, carboxylic, ester, amide, aldehyde, ketone, ether groups.

[0072] The term "unconventional moiety" as used herein refers to abasic ribose moiety, an abasic deoxyribose moiety, a deoxyribonucleotide, a modified deoxyribonucleotide, a mirror nucleotide, a non-base pairing nucleotide analog and a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide phosphate bond; bridged nucleic acids including locked nucleic acids (LNA) and ethylene bridged nucleic acids (ENA).

[0073] Abasic deoxyribose moiety includes for example abasic deoxyribose-3'-phosphate; 1,2-dideoxy-D-ribofuranose-3-phosphate; 1,4-anhydro-2-deoxy-D-ribitol 1-3-phosphate.

[0074] Inverted abasic deoxyribose moiety includes inverted deoxyriboabasic; 3',5' inverted deoxyabasic 5'-phosphate.

[0075] A "mirror" nucleotide is a nucleotide with reversed chirality to the naturally occurring or commonly employed nucleotide, i.e., a mirror image (L-nucleotide) of the naturally occurring (D-nucleotide), also referred to as L-RNA in the case of a mirror ribonucleotide, and "spiegelmer". The nucleotide can be a ribonucleotide or a deoxyribonucleotide and may further comprise at least one sugar, base and or backbone modification. See US Patent No. 6,586,238. Also, US Patent No. 6,602,858 discloses nucleic acid catalysts comprising at least one L-nucleotide substitution. Mirror nucleotide includes for example L-DNA (L-deoxyriboadenosine-3'-phosphate (mirror dA); L-deoxyribocytidine-3'-phosphate (mirror dC); L-deoxyriboguanosine-3'-phosphate (mirror dG); L-deoxyribothymidine-3'-phosphate (mirror dT) and L-RNA (L-riboadenosine-3'-phosphate (mirror rA); L-ribocytidine-3'-phosphate (mirror rC); L-riboguanosine-3'-phosphate (mirror rG); L-ribouridine-3'-phosphate (mirror dU).

[0076] According to one aspect the present invention provides inhibitory modified siRNA compounds comprising unmodified ribonucleotides, modified ribonucleotides and/or unconventional moieties. In some embodiments the modified siRNA compound comprises at least one modified nucleotide selected from the group consisting of a sugar modification, a base modification and an internucleotide linkage modification and may contain modified nucleotides such as LNA (locked nucleic acid) including ENA (ethylene-bridged nucleic acid; PNA (peptide nucleic acid); arabinoside; PACE (phosphonoacetate and derivatives thereof), or nucleotides with a six-carbon sugar or an unconventional moiety selected from an abasic ribose moiety, an abasic deoxyribose moiety, a modified or unmodified deoxyribonucleotide, a mirror nucleotide, and a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide phosphate bond. In some embodiments a modified ribonucleotide is a 2'OMe sugar modified ribonucleotide.

[0077] In some embodiments the siRNA is blunt ended at the 3' terminus of the compound, i.e. the siRNA is blunt ended on the end defined by the 3'-terminus of the sense or passenger strand and the 5'-terminus of antisense or guide strand.

[0078] In other embodiments at least one of the two strands has a 3' overhang of at least one nucleotide at the 3'-terminus; the overhang comprises at least one deoxyribonucleotide. At least one of the strands optionally comprises an overhang of at least one nucleotide at the 3'-terminus. The overhang consists of from about 1 to about 5 nucleotides.

[0079] In various embodiments the overhangs are independently selected from a nucleotide, a non-nucleotide and a combination thereof. In certain embodiments, each overhang, if present, is independently selected from a ribonucleotide, deoxyribonucleotide, abasic deoxyribose moiety, abasic deoxyribose moiety, C3-amino-Pi, C4-amino-Pi, C5-amino-Pi, C6-amino-Pi, a mirror nucleotide.

[0080] In some embodiments each of Z and/or Z' independently includes a C2, C3, C4, C5 or C6 alkyl moiety, optionally a C3 [propane, -(CH₂)₃-] moiety or a derivative thereof including propanol (C3-OH), propanediol, and phosphodiester derivative of propanediol ("C3Pi"). In preferred embodiments each of Z and/or Z' includes two hydrocarbon moieties and in some examples is C3Pi-C3OH or C3Pi-C3Pi. Each C3 is covalently conjugated to an adjacent C3 via a covalent bond, preferably a phospho-based bond. In some embodiments the phospho-based bond is a phosphorothioate, a phosphonoacetate or a phosphodiester bond.

[0081] In some embodiments the C3-C3 overhang is covalently attached to the 3' terminus of (N)x or (N')y via a covalent linkage, for example a phosphodiester linkage. In some embodiments the linkage between a first C3 and a second C3 is a phosphodiester linkage. In some embodiments the 3' non-nucleotide overhang is C3Pi-C3Pi. In some embodiments the 3' non-nucleotide overhang is C3Pi-C3Ps. In some embodiments the 3' non-nucleotide overhang is C3Pi-C3OH (OH is hydroxy). In some embodiments the 3' non-nucleotide overhang is C3Pi-C3OH.

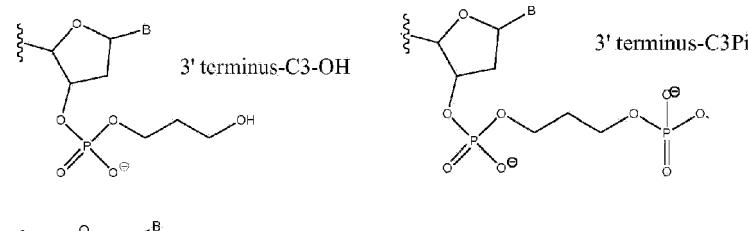
[0082] In various embodiments the alkyl moiety comprises an alkyl derivative including a C3 alkyl, C4 alkyl, C5 alkyl or C6 alkyl moiety comprising a terminal hydroxyl, a terminal amino, or terminal phosphate group. In some embodiments the alkyl moiety is a C3 alkyl or C3 alkyl derivative moiety. In some embodiments the C3 alkyl moiety comprises propanol, propylphosphate, propylphosphorothioate or a combination thereof.

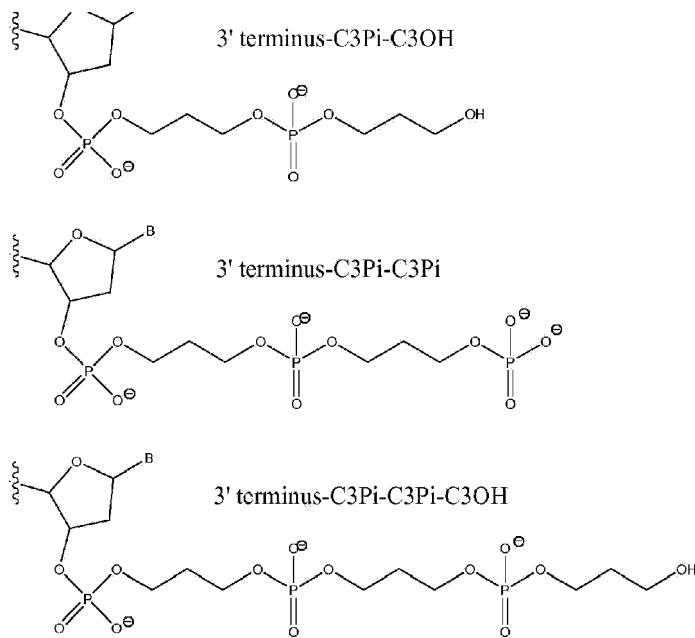
[0083] The C3 alkyl moiety is covalently linked to the 3' terminus of (N')y and/or the 3' terminus of (N)x via a phosphodiester bond. In some embodiments the alkyl moiety comprises propanol, propyl phosphate or propyl phosphorothioate.

[0084] In some embodiments each of Z and Z' is independently selected from propanol, propyl phosphate propyl phosphorothioate, combinations thereof or multiples thereof in particular 2 or 3 covalently linked propanol, propyl phosphate, propyl phosphorothioate or combinations thereof.

[0085] In some embodiments each of Z and Z' is independently selected from propyl phosphate, propyl phosphorothioate, propyl phospho-propanol; propyl phospho-propyl phosphorothioate; propylphospho-propyl phosphate; (propyl phosphate)₃, (propyl phosphate)₂-propanol, (propyl phosphate)₂- propyl phosphorothioate. Any propane or propanol conjugated moiety can be included in Z or Z'.

[0086] The structures of exemplary 3' terminal non-nucleotide moieties are as follows:





[0087] In additional embodiments each of Z and/or Z' comprises a combination of an abasic moiety and an unmodified deoxyribonucleotide or ribonucleotide or a combination of a hydrocarbon moiety and an unmodified deoxyribonucleotide or ribonucleotide or a combination of an abasic moiety (deoxyribo or ribo) and a hydrocarbon moiety. In such embodiments, each of Z and/or Z' comprises C3Pi-rAb or C3Pi-dAb.

[0088] The length of RNA duplex is from about 18 to about 40 ribonucleotides, preferably 19 to 23 ribonucleotides. In some aspects the length of each strand (oligomer) is independently selected from the group consisting of about 18 to about 40 bases, preferably 18 to 23 bases and more preferably 19, 20 or 21 ribonucleotides.

[0089] The antisense strand of the modified siRNA compound and the target nucleic acid are substantially complementary, i.e. having one, two or up to three mismatches between said antisense strand and the target nucleic acid.

[0090] The complementarity between the antisense strand and the sense strand of the modified siRNA compound of present invention is perfect.

[0091] The siRNAs compounds of the present invention possess a terminal moiety covalently bound at the 5'-terminus of the antisense strand which is mismatched to a nucleotide in a target mRNA. In various aspects the moiety at the 5'-terminus of the antisense strand is selected from the group consisting of ribouridine, deoxyribouridine, modified ribouridine, modified deoxyribouridine, pseudouracil, deoxypseudouracil, deoxyribothymidine, modified deoxyribothymidine, ribocytosine, modified ribocytosine, deoxyribocytosine, modified deoxyribocytosine, 5-methylribocytosine, modified 5-methylribocytosine, 5-methylribouridine, ribo-2-thiouridine, ribo-4-thiouridine, abasic ribose moiety and abasic deoxyribose moiety.

[0092] In some embodiments the modified siRNA compounds of the invention exhibit enhanced activity, when compared to an siRNA compound wherein the antisense strand including the 5'-terminal

nucleotide is fully complementary to a consecutive sequence in a target mRNA.

[0093] The siRNA structures of the present invention are beneficially applied to double stranded RNA useful in inhibiting or attenuating mammalian and non-mammalian gene expression.

siRNA Oligonucleotides

[0094] In one aspect the present invention provides a compound having structure (A) set forth below:

(A)	5'	N'-(N)x - Z	3' (antisense strand)
	3'	Z'-N ² -(N')y -z"	5' (sense strand)

wherein each of N², N and N' is an unmodified or modified ribonucleotide, or an unconventional moiety;

wherein each of (N)x and (N')y is an oligonucleotide in which each consecutive N or N' is joined to the adjacent N or N' by a covalent bond;

wherein each of x and y is 18;

wherein the sequence of N²-(N')y is fully complementary to the sequence of N'-(N)x and (N)x has complementarity to a consecutive sequence in a target RNA;

wherein N¹ is covalently bound to (N)x and is mismatched to the target RNA;

wherein N¹ is a moiety selected from the group consisting of natural or modified deoxyribouridine, adenosine or deoxyadenosine;

wherein z" may be present or absent, but if present is a capping moiety covalently attached at the 5' terminus of N²-(N')y; and

wherein each of Z and Z' is independently present or absent, but if present is independently 1-5 consecutive nucleotides, consecutive non-nucleotide moieties or a combination thereof covalently attached at the 3' terminus of the strand in which it is present.

[0095] In some embodiments N¹ is covalently bound to (N)x and is mismatched to the target RNA.

[0096] In some embodiments N¹ and N² together form a Watson-Crick base pair. In other embodiments N¹ and N² together form a non-Watson-Crick base pair.

[0097] In certain embodiments N¹ is selected from the group consisting of deoxyribouridine and modified deoxyribouridine.

[0098] In some aspects N² is a modified ribonucleotide or an unmodified ribonucleotide. In certain aspects N² is selected from the group comprising riboguanine, modified riboguanine, deoxyguanine,

modified deoxyguanine, ribouridine, modified ribouridine, deoxyribouridine, modified deoxyribouridine, adenosine, modified adenosine, deoxyadenosine, modified deoxyadenosine, ribocytosine, modified ribocytosine, deoxyribocytosine and modified deoxyribocytosine.

[0099] In some aspects N² is selected from the group consisting of riboguanine, modified riboguanine, deoxyguanine, ribouridine, deoxyribouridine, adenosine, deoxyadenosine, ribocytosine and deoxyribocytosine. In other aspects N² is selected from the group consisting of ribouridine, modified ribouridine, deoxyribouridine, modified deoxyribouridine, adenosine, modified adenosine, deoxyadenosine, modified deoxyadenosine, ribocytosine, modified ribocytosine, deoxyribocytosine and modified deoxyribocytosine. In yet other aspects N² is selected from the group consisting of ribouridine, modified ribouridine, deoxyribouridine, modified deoxyribouridine, adenosine, modified adenosine, deoxyadenosine, modified deoxyadenosine, ribocytosine, modified ribocytosine, deoxyribocytosine and modified deoxyribocytosine.

[0100] In certain aspects N¹ is selected from the group consisting of ribocytosine, modified ribocytosine, deoxyribocytosine, modified deoxyribocytosine, 5-methylribocytosine and modified 5-methylribocytosine and N² is selected from the group consisting of riboguanine, modified riboguanine, deoxyguanine, ribouridine, deoxyribouridine, adenosine, deoxyadenosine, ribocytosine and deoxyribocytosine.

[0101] In some aspects N¹ is selected from the group consisting of ribouridine, deoxyribouridine, modified ribouridine, modified deoxyribouridine, pseudouracil, deoxypseudouracil, 5-methylribouridine, ribo-2-thiouridine and ribo 4-thiouridine and N² is selected from the group consisting of ribouridine, modified ribouridine, deoxyribouridine, modified deoxyribouridine, adenosine, modified adenosine, deoxyadenosine, modified deoxyadenosine, ribocytosine, modified ribocytosine, deoxyribocytosine and modified deoxyribocytosine.

[0102] In certain aspects N¹ is deoxyribothymidine or modified deoxyribothymidine and N² is selected from the group consisting of ribouridine, modified ribouridine, deoxyribouridine, modified deoxyribouridine, adenosine, modified adenosine, deoxyadenosine, modified deoxyadenosine, ribocytosine, modified ribocytosine, deoxyribocytosine and modified deoxyribocytosine.

[0103] In certain embodiments of structure (A), N² comprises a 2'OMe sugar modified ribonucleotide.

[0104] In some embodiments Z and Z' are absent. In other embodiments one of Z or Z' is present.

[0105] In various embodiments Z and Z' are independently selected from a nucleotide, a non-nucleotide and a combination thereof. In certain embodiments, each of Z and Z', if present, is independently selected from a ribonucleotide, deoxyribonucleotide, abasic deoxyribose moiety, abasic deoxyribose moiety, C3-amino-Pi, C4-amino-Pi, C5-amino-Pi, C6-amino-Pi, a mirror nucleotide. In some embodiments Z is present. In other embodiments Z' is present. In additional embodiments both Z and Z' are present. In some embodiments Z and Z' are present and are identical. In further embodiments Z and Z' are present and are different. In some embodiments Z and Z' are independently 1, 2, 3, 4 or 5 non-nucleotide moieties or a combination of 2, 3, 4, or 5 non-nucleotide moieties and nucleotides. In some embodiments each of Z and or Z' comprises 2 non-nucleotide moieties covalently linked to the 3' terminus of the siRNA strand via a phosphodiester bond. In some

embodiments Z and Z' are present and each one independently comprises one or more alkyl moieties and or derivative thereof. In some embodiments, N² comprises riboadenosine and N¹ comprises uridine (ribouridine).

[0106] A non-nucleotide moiety is selected from the group consisting of an abasic moiety, an inverted abasic moiety, an alkyl moiety or derivative thereof, and an inorganic phosphate. In some embodiments a non-nucleotide moiety is an alkyl moiety or derivative thereof. In some embodiments the alkyl moiety comprises a terminal functional group including alcohol, a terminal amine, a terminal phosphate or a terminal phosphorothioate moiety.

[0107] In some embodiments Z is present and comprises one or more non-nucleotide moieties selected from the group consisting of an abasic moiety, an inverted abasic moiety, hydrocarbon moiety or derivative thereof, and an inorganic phosphate. In some embodiments Z is present and comprises one or more alkyl moieties and or derivative thereof.

[0108] In additional embodiments Z' is present and comprises one or more non-nucleotide moieties selected from the group consisting of an abasic moiety, an inverted abasic moiety, a hydrocarbon moiety, and an inorganic phosphate. In some embodiments Z' is present and comprises one or more alkyl moieties and or derivative thereof.

[0109] In additional embodiments x=y=18 and either Z or Z' is present and independently comprises two non-nucleotide moieties.

[0110] In additional embodiments x=y=18 and Z and Z' are present and each independently comprises two non-nucleotide moieties.

[0111] In some embodiments each of Z and Z' includes an abasic moiety, for example a deoxyriboabasic moiety (referred to herein as "dAb") or riboabasic moiety (referred to herein as "rAb"). In some embodiments each of Z and/or Z' comprises two covalently linked abasic moieties and is for example dAb-dAb or rAb-rAb or dAb-rAb or rAb-dAb. Each moiety is covalently conjugated an adjacent moiety via a covalent bond, preferably a phospho-based bond. In some embodiments the phospho-based bond is a phosphorothioate, a phosphonoacetate or a phosphodiester bond.

[0112] In some embodiments each of Z and/or Z' independently includes an alkyl moiety, optionally propane [(CH₂)₃] moiety or a derivative thereof including propanol (C₃-OH) and phospho derivative of propanediol ("C₃-3'Pi"). In some embodiments each of Z and/or Z' includes two hydrocarbon moieties and in some examples is C₃-C₃. Each C₃ is covalently conjugated an adjacent C₃ via a covalent bond, preferably a phospho-based bond. In some embodiments the phospho-based bond is a phosphorothioate, a phosphonoacetate or a phosphodiester bond.

[0113] In a specific embodiment x=y=18 and Z comprises C₃Pi-C₃OH or C₃Pi-C₃Pi. In a specific embodiment x=y=18 and Z comprises C₃Pi-C₃OH or C₃Pi-C₃Pi. In some embodiments the C₃-C₃ overhang is covalently attached to the 3' terminus of (N)_x or (N')_y via covalent linkage, for example a phosphodiester linkage. In some embodiments the linkage between a first C₃ and a second C₃ is a phosphodiester linkage.

[0114] In various embodiments the alkyl moiety is a C₃ alkyl to C₆ alkyl moiety comprising a terminal

hydroxyl, a terminal amino, terminal phosphate group. In some embodiments the alkyl moiety is a C3 alkyl moiety. In some embodiments the C3 alkyl moiety comprises propanol, propylphosphate, propylphosphorothioate or a combination thereof.

[0115] The C3 alkyl moiety is covalently linked to the 3' terminus of (N') y and/or the 3' terminus of (N) x via a phosphodiester bond. In some embodiments the alkyl moiety comprises propanol, propyl phosphate or propyl phosphorothioate.

[0116] In some embodiments each of Z and Z' is independently selected from propanol, propyl phosphate, propyl phosphorothioate, combinations thereof or multiples thereof.

[0117] In some embodiments each of Z and Z' is independently selected from propyl phosphate, propyl phosphorothioate, propyl phospho-propanol; propyl phospho-propyl phosphorothioate; propylphospho-propyl phosphate; (propyl phosphate) 3 , (propyl phosphate) 2 -propanol, (propyl phosphate) 2 - propyl phosphorothioate. Any propane or propanol conjugated moiety can be included in Z or Z'.

[0118] In additional embodiments each of Z and/or Z' comprises a combination of an abasic moiety and an unmodified deoxyribonucleotide or ribonucleotide or a combination of a hydrocarbon moiety and an unmodified deoxyribonucleotide or ribonucleotide or a combination of an abasic moiety (deoxyribo or ribo) and a hydrocarbon moiety. In such embodiments, each of Z and/or Z' comprises C3-rAb or C3-dAb.

[0119] In preferred embodiments x=y=18, Z' is absent, Z is present and comprises two alkyl moieties covalently linked to each other via a phosphodiester bond, N² comprises riboadenosine and N¹ comprises uridine.

[0120] In some embodiments N and N' comprise an unmodified nucleotide. In some embodiments at least one of N or N' comprises a chemically modified ribonucleotide or an unconventional moiety. In some embodiments the unconventional moiety is selected from the group consisting of a mirror nucleotide, an abasic ribose moiety, an abasic deoxyribose moiety, a deoxyribonucleotide, a modified deoxyribonucleotide, a mirror nucleotide, a non-base pairing nucleotide analog, a bridged nucleic acid and a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide phosphate bond. In some embodiments the unconventional moiety is a mirror nucleotide, preferably an L-DNA moiety. In some embodiments at least one of N or N' is modified at one or more of the sugar, the base or linker. In certain embodiments at least one of N or N' comprises a 2'OMe sugar modified ribonucleotide.

[0121] (N) x and (N') y are fully complementary. In certain embodiments (N) x is fully complementary to a target sequence. In other embodiments (N) x is substantially complementary to a target sequence. According to certain preferred embodiments the present invention provides a modified siRNA compound comprising one or more modified nucleotide, wherein the modified nucleotide possesses a modification in the sugar moiety, in the base moiety or in the internucleotide linkage moiety.

[0122] In certain embodiments of the compound according to Structure (A) alternating ribonucleotides in each of (N) x and (N') y are 2'-OMe sugar modified ribonucleotides. In some embodiments of Structure (A) in (N) x the nucleotides are unmodified or (N) x comprises alternating 2'OMe sugar modified ribonucleotides and unmodified ribonucleotides; and the ribonucleotide located at the middle

position of $N^1-(N)x$ being modified or unmodified preferably unmodified; wherein $(N')y$ comprises unmodified ribonucleotides further comprising one modified nucleotide at a terminal or penultimate position.

[0123] In particular embodiments, $x=y=18$, N^1 comprises 2'OMe sugar modified ribonucleotide, $(N)x$ comprises 2'OMe sugar modified ribonucleotides and the ribonucleotide located at the middle of $N^1-(N)x$ is unmodified. In some embodiments at least one nucleotide at either or both the 5' and 3' termini of $(N')y$ are joined by a 2'-5' phosphodiester bond. In some embodiments N^1 is joined to the 5' terminus of $(N)x$ by a 2'-5' phosphodiester bond. In some embodiments at least one nucleotides at either or both the 5' and 3' termini of $(N')x$ are joined by a 2'-5' phosphodiester bond. In some embodiments N^2 is joined to the 3' terminus of $(N)y$ by a 2'-5' phosphodiester bond. In certain embodiments $x=y=18$; in $(N)x$ the nucleotides alternate between modified ribonucleotides and unmodified ribonucleotides, each modified ribonucleotide is a 2'OMe sugar modified ribonucleotide and the ribonucleotide located at the middle of $N^1-(N)x$ being unmodified; N^2 is joined to the 3' terminus of $(N)y$ by a 2'-5' phosphodiester bond and at least two nucleotides at the 3' terminus of $(N')y$ are joined by two 2'-5' phosphodiester bonds. In other preferred embodiments, $x=y=18$; in $(N)x$ the nucleotides alternate between modified ribonucleotides and unmodified ribonucleotides, each modified ribonucleotide is a 2'-OMe sugar modified ribonucleotide and the ribonucleotide located at the middle of $N^1-(N)x$ being unmodified; and at least three consecutive nucleotides at the 5' terminus of $(N')y$ are joined by three 2'-5' phosphodiester bonds. In a further embodiment, the nucleotide located in the middle position of $N^2-(N)y$, i.e. nucleotide at position 10, is a 2'OMe sugar modified ribonucleotide. In another preferred embodiment, in $(N)x$ the nucleotides alternate between 2'-OMe sugar modified ribonucleotides and unmodified ribonucleotides, and in $(N')y$ four consecutive nucleotides at the 5' terminus are joined by three 2'-5' phosphodiester bonds and the 5' terminal nucleotide or two or three consecutive nucleotides at the 5' terminus comprise 3'-OMe sugar modification.

[0124] In certain preferred embodiments, $x=y=18$ and in $(N')y$ the nucleotide in at least one position comprises a mirror nucleotide, a deoxyribonucleotide and a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide bond.

[0125] In certain embodiments, $x=y=18$ and $(N')y$ comprises a mirror nucleotide. In various embodiments the mirror nucleotide is an L-DNA nucleotide. In certain embodiments the L-DNA is L-deoxyribocytidine. In some embodiments $(N')y$ comprises L-DNA at position 17. In other embodiments $(N')y$ comprises L-DNA at positions 16 and 17. In certain embodiments $(N')y$ comprises L-DNA nucleotides at positions 1 and at one or both of positions 16 and 17.

[0126] In yet other embodiments $(N')y$ comprises a DNA at position 14 and L-DNA at one or both of positions 16 and 17. In that structure, position 1 may further comprise an L-DNA or an abasic unconventional moiety.

[0127] According to various embodiments in $N^2-(N')y$, N^2 is joined to the 3' terminus of $(N)y$ by a 2'-5' phosphodiester bond and in $(N')y$ 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12 or 13 consecutive ribonucleotides at the 3' terminus are linked by 2'-5' internucleotide linkages. In one embodiment, N^2 is joined to the 3' terminus of $(N)y$ by a 2'-5' phosphodiester bond and three consecutive nucleotides at the 3' terminus

of (N')y are joined by three 2'-5' phosphodiester bonds, wherein one or more of the 2'-5' nucleotides which form the 2'-5' phosphodiester bonds further comprises a 3'-OMe sugar modification. In some embodiments N² comprises a 2'-OMe sugar modification. In further embodiments the 3' terminal nucleotide of (N')y comprises a 2'-OMe sugar modification. In certain embodiments x=y=18 and in (N')y two or more consecutive nucleotides at positions 14, 15, 16, 17 and 18 comprise a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide bond. In various embodiments the nucleotide forming the 2'-5' internucleotide bond comprises a 3' deoxyribose nucleotide or a 3' methoxy nucleotide. In some embodiments the nucleotides at positions 16 and 17 in (N')y are joined by a 2'-5' internucleotide bond. In other embodiments the nucleotides at positions 15-16, 16-17, or 15-17 in (N')y are joined by a 2'-5' internucleotide bond.

[0128] In certain embodiments (N')y comprises an L-DNA at position 2 and 2'-5' internucleotide bonds at positions 15-16, 16-17, or 15-17.

[0129] In one embodiment the 3' terminal nucleotide or two or three consecutive nucleotides at the 3' terminus of (N')y are L-deoxyribonucleotides.

[0130] In other embodiments in N²-(N')y, N² is a 2' sugar modified nucleotide and in (N')y (N')y 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12 or 13 consecutive ribonucleotides at either terminus or 1 to 7 modified nucleotides at each of the 5' and 3' termini are independently 2' sugar modified nucleotides. In some embodiments the 2' sugar modification comprises the presence of an amino, a fluoro, an alkoxy or an alkyl moiety. In certain embodiments the 2' sugar modification comprises a methoxy moiety (2'-OMe). In one series of embodiments, three, four or five consecutive nucleotides at the 5' terminus of (N')y comprise the 2'-OMe modification. In another embodiment, N² and two consecutive nucleotides at the 3' terminus of (N')y comprise the 2'-OMe modification.

[0131] In some embodiments in (N')y 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12 or 13 consecutive ribonucleotides at either or 1 to 7 modified nucleotides at each of the 5' and 3' termini are independently bicyclic nucleotides. In various embodiments the bicyclic nucleotide is a locked nucleic acid (LNA) or a species of LNA, e.g. 2'-O, 4'-C-ethylene-bridged nucleic acid (ENA) is a species of LNA.

[0132] In various embodiments (N')y comprises modified nucleotides at the 5' terminus or at both the 3' and 5' termini.

[0133] In some embodiments at least two nucleotides at the 5' of (N')y are joined by P-ethoxy backbone modifications. In some embodiments N² at the 3' of (N')y is joined by P-ethoxy backbone modifications. In certain embodiments x=y=18 and wherein in (N)x the nucleotides alternate between modified ribonucleotides and unmodified ribonucleotides, each modified ribonucleotide being a 2'OMe sugar modified ribonucleotide and the ribonucleotide located at the middle position of N¹-(N)x being unmodified; N1 being a 2'OMe sugar modified ribonucleotide and three consecutive nucleotides at the 3' terminus are joined by two P-ethoxy backbone modifications or four consecutive nucleotides at the 5' terminus of (N')y are joined by three P-ethoxy backbone modifications. In a further embodiment, N² at the 3' of (N')y is joined by P-ethoxy backbone modification.

[0134] In some embodiments in (N')y 1, 2, 3, 4, 5, 6 or 7, consecutive ribonucleotides at each of the

5' and 3' termini are independently mirror nucleotides, nucleotides joined by 2'-5' phosphodiester bond, 2' sugar modified nucleotides or bicyclic nucleotide. In one embodiment, the modification at the 5' and 3' termini of (N')y is identical. In one embodiment, four consecutive nucleotides at the 5' terminus of (N')y are joined by three 2'-5' phosphodiester bonds and two or three consecutive nucleotides at the 3' terminus of (N')y are joined by two 2'-5' phosphodiester bonds. In a further embodiment N² at the 3' of (N')y is joined at the 3' terminus of (N')y by a 2'-5' phosphodiester bond. In another embodiment, the modification at the 5' terminus of (N')y is different from the modification at the 3' terminus of (N')y. In one specific embodiment, the modified nucleotides at the 5' terminus of (N')y are mirror nucleotides and the modified nucleotides at the 3' terminus of (N')y are joined by 2'-5' phosphodiester bond. In another specific embodiment, three consecutive nucleotides at the 5' terminus of (N')y are LNA nucleotides and two consecutive nucleotides at the 3' terminus of (N')y are joined by two 2'-5' phosphodiester bonds. In (N)x the nucleotides alternate between 2'OMe sugar modified ribonucleotides and unmodified ribonucleotides, and the ribonucleotide located at the middle of N¹-(N)x is unmodified.

[0135] In various embodiments, N² is joined at the 3' terminus of (N')y by a 2'-5' phosphodiester bond and two or three consecutive nucleotides at the 3' terminus of (N')x are joined by one or two 2'-5' phosphodiester bonds. In some embodiments, N² is joined at the 3' terminus of (N')y by a 2'-5' phosphodiester bond and two or three consecutive nucleotides at the 3' terminus of (N')x are joined by one or two 2'-5' phosphodiester bonds and one or two or three consecutive nucleotides at the 3' terminus comprise 3'-OMe sugar modification. In a further embodiment N² comprise 3'-OMe sugar modification.

[0136] In another embodiment the present invention provides a compound wherein x=y=18 and wherein in (N)x the nucleotides alternate between modified ribonucleotides and unmodified ribonucleotides, each modified ribonucleotide being a 2'OMe sugar modified ribonucleotide and the ribonucleotide located at the middle of N¹-(N)x being unmodified; two nucleotides at the 3' terminus of (N')y are joined by two 2'-5' phosphodiester bonds and three nucleotides at the 5' terminus of (N')y are LNA such as ENA. In a further embodiment N² is joined at the 3' terminus of (N')y by a 2'-5' phosphodiester bond.

[0137] In another embodiment five consecutive nucleotides at the 5' terminus of (N')y comprise the 2'OMe sugar modification and two consecutive nucleotides at the 3' terminus of (N')y are L-DNA.

[0138] In yet another embodiment, the present invention provides a compound wherein x=y=18 wherein (N)x consists of unmodified ribonucleotides; two consecutive nucleotides at the 3' terminus of (N')y are joined by a 2'-5' phosphodiester bonds and three consecutive nucleotides at the 5' terminus of (N')y are LNA such as ENA. In a further embodiment N² is joined at the 3' terminus of (N')y by a 2'-5' phosphodiester bond.

[0139] According to other embodiments in (N')y the 5' or 3' terminal nucleotide, or 2, 3, 4, 5 or 6 consecutive nucleotides at either termini or 1 to 4 nucleotides at each of the 5' and 3' termini are independently phosphonocarboxylate or phosphinocarboxylate nucleotides (PACE nucleotides). In some embodiments the PACE nucleotides are deoxyribonucleotides. In some preferred embodiments in (N')y, 1 or 2 consecutive nucleotides at each of the 5' and 3' termini are PACE nucleotides.

[0140] In some embodiments, neither strand of the modified siRNA compounds of the invention is phosphorylated at the 3' and 5' termini. In other embodiments the sense and antisense strands are phosphorylated at the 3' termini. In yet another embodiment, the antisense strand is phosphorylated at the terminal 5' termini position using cleavable or non-cleavable phosphate groups. In yet another embodiment, either or both antisense and sense strands are phosphorylated at the 3' termini position using cleavable or non-cleavable phosphate groups.

[0141] Structure (A) is useful with any oligonucleotide pair (sense and antisense strands) to a mammalian or non-mammalian gene. In some embodiments the mammalian gene is a human gene. Examples of oligonucleotide sequence pairs are provided in PCT Patent Publication Nos. WO 2006/023544, WO 2007/084684, WO 2008/050329, WO 2007/141796, WO 2009/044392, WO 2008/106102, WO 2008/152636, WO 2009/001359, WO/2009/090639 assigned to the assignee of the present invention.

[0142] Unless otherwise indicated, in preferred embodiments of the structures discussed herein the covalent bond between each consecutive N and N' is a phosphodiester bond. Unless otherwise indicated, in preferred embodiments of the structures discussed herein the covalent bond between N¹ and (N)x and between N² and (N)y is a phosphodiester bond. In some embodiments at least one of the covalent bond is a phosphorothioate bond.

[0143] For all of the structures above, the oligonucleotide sequence of (N)x is fully complementary to the oligonucleotide sequence of (N')y. In certain embodiments (N)x is fully complementary to a mammalian mRNA or microbial RNA or viral RNA. In other embodiments (N)x is substantially complementary to a mammalian mRNA or microbial RNA or viral RNA.

[0144] In some embodiments a modified siRNA compound having structure (A) exhibits beneficial properties including at least enhanced activity when compared to an siRNA compound wherein N¹ is complementary to a nucleotide in a target mRNA.

[0145] The present invention further provides a pharmaceutical composition comprising a compound of the present invention according to Structure (A), in an amount effective to inhibit mammalian or non-mammalian gene expression, and a pharmaceutically acceptable carrier, and use thereof for treatment of any one of the diseases and disorders disclosed herein. In some embodiments the mammalian gene is a human gene. In some embodiments the non-mammalian gene is involved in a mammalian disease, preferably human disease.

[0146] The present invention further relates to the siRNA compounds of the invention for use in methods for treating or preventing the incidence or severity of any one of the diseases or conditions disclosed herein or for reducing the risk or severity of a disease or a condition disclosed herein in a subject in need thereof, wherein the disease or condition and/or a symptom or risk associated therewith is associated with expression of a mammalian or a non-mammalian gene. In a preferred embodiment the subject is a human subject.

siRNA synthesis

[0147] Using public and proprietary algorithms the sense and antisense sequences of potential siRNAs are generated and N¹ and/or N² of the siRNAs are substituted to generate the modified siRNA compounds according to Structure (A).

[0148] siRNA molecules according to the above specifications are prepared essentially as described herein. The modified siRNA compounds of the present invention are synthesized by any of the methods that are well known in the art for synthesis of ribonucleic (or deoxyribonucleic) oligonucleotides. Synthesis is commonly performed in a commercially available synthesizer (available, *inter alia*, from Applied Biosystems). Oligonucleotide synthesis is described for example in Beauchage and Iyer, *Tetrahedron* 1992; 48:2223-2311; Beauchage and Iyer, *Tetrahedron* 1993; 49: 6123-6194 and Caruthers, et. al., *Methods Enzymol.* 1987; 154: 287-313; the synthesis of thioates is, among others, described in Eckstein, *Ann. Rev. Biochem.* 1985; 54: 367-402, the synthesis of RNA molecules is described in Sproat, in *Humana Press* 2005 edited by Herdewijn P.; Kap. 2: 17-31 and respective downstream processes are, among others, described in Pingoud et al., in *IRL Press* 1989 edited by Oliver R.W.A.; Kap. 7: 183-208.

[0149] Other synthetic procedures are known in the art, e.g. the procedures described in Usman et al., 1987, *J. Am. Chem. Soc.*, 109, 7845; Scaringe et al., 1990, *NAR.*, 18, 5433; Wincott et al., 1995, *NAR.* 23, 2677-2684; and Wincott et al., 1997, *Methods Mol. Bio.*, 74, 59, may make use of common nucleic acid protecting and coupling groups, such as dimethoxytrityl at the 5'-end, and phosphoramidites at the 3'-end. The modified (e.g. 2'-O-methylated) nucleotides and unmodified nucleotides are incorporated as desired.

[0150] In some embodiments the oligonucleotides of the present invention are synthesized separately and joined together post-synthetically, for example, by ligation (Moore et al., 1992, *Science* 256, 9923; Draper et al., International Patent Publication No. WO 93/23569; Shabarova et al., 1991, *NAR* 19, 4247; Bellon et al., 1997, *Nucleosides & Nucleotides*, 16, 951; Bellon et al., 1997, *Bioconjugate Chem.* 8, 204), or by hybridization following synthesis and/or deprotection.

[0151] Overlapping pairs of chemically synthesized fragments can be ligated using methods well known in the art (e.g., see US Patent No. 6,121,426). The strands are synthesized separately and then are annealed to each other in the tube. Then, the double-stranded siRNAs are separated from the single-stranded oligonucleotides that were not annealed (e.g. because of the excess of one of them) by HPLC. In relation to the modified siRNA compounds of the present invention, two or more such sequences can be synthesized and linked together for use in the present invention.

[0152] The siRNAs compounds of the present invention possess a terminal moiety covalently bound at the 5'-terminus of the antisense strand which is mismatched to a nucleotide in a target mRNA. In various aspects the moiety at the 5'-terminus of the antisense strand is selected from the group consisting of ribouridine, deoxyribouridine, modified ribouridine, modified deoxyribouridine, pseudouracil, deoxypseudouracil, deoxyribothymidine, modified deoxyribothymidine, ribocytosine, modified ribocytosine, deoxyribocytosine, modified deoxyribocytosine, 5-methylribocytosine, modified 5-methylribocytosine, 5-methylribouridine, ribo-2-thiouridine, ribo-4-thiouridine, abasic ribose moiety and abasic deoxyribose moiety and the moiety at the 3'-terminus of the sense strand is selected from a ribonucleotide or a modified ribonucleotide or an unconventional moiety. The siRNA structures of the present invention are beneficially applied to double stranded RNA useful in inhibiting or attenuating mammalian and non-mammalian gene expression.

Pharmaceutical Compositions

[0153] While it is possible for the compounds of the present invention to be administered as the raw chemical, it is preferable to present them as a pharmaceutical composition. Accordingly the present invention provides a pharmaceutical composition comprising one or more of the modified siRNA compounds of the invention; and a pharmaceutically acceptable carrier. In some embodiments the pharmaceutical composition comprises two or more modified siRNA compounds of the invention.

[0154] The invention further provides a pharmaceutical composition comprising at least one compound of the invention covalently or non-covalently bound to one or more compounds of the invention in an amount effective to inhibit a target gene; and a pharmaceutically acceptable carrier. In some embodiments the modified siRNA compounds are processed intracellularly by endogenous cellular complexes to produce one or more oligoribonucleotides of the invention.

[0155] The invention further provides a pharmaceutical composition comprising a pharmaceutically acceptable carrier and one or more of the modified siRNA compounds of the invention in an amount effective to down-regulate expression in a cell in a mammal of a target gene, the compound comprising a sequence which is substantially complementary to the sequence of a target mRNA.

[0156] In some embodiments, the modified siRNA compounds according to the present invention are the main active component in a pharmaceutical composition. In other embodiments the modified siRNA compounds according to the present invention are one of the active components of a pharmaceutical composition containing two or more siRNAs, said pharmaceutical composition further being comprised of one or more siRNA compounds which target one or more target genes.

[0157] The present description also discloses a process of preparing a pharmaceutical composition, which comprises:
providing one or more double stranded modified siRNA compound of the invention; and admixing said compound with a pharmaceutically acceptable carrier.

[0158] In a preferred embodiment, the modified siRNA compound used in the preparation of a pharmaceutical composition is admixed with a carrier in a pharmaceutically effective dose. In some embodiments the modified siRNA compound of the present invention is conjugated to a steroid or to a lipid or to another suitable molecule e.g. to cholesterol.

RNA interference

[0159] A number of PCT applications have recently been published that relate to the RNAi phenomenon. These include: PCT publication WO 00/44895; PCT publication WO 00/49035; PCT publication WO 00/63364; PCT publication WO 01/36641; PCT publication WO 01/36646; PCT publication WO 99/32619; PCT publication WO 00/44914; PCT publication WO 01/29058; and PCT publication WO 01/75164.

[0160] RNA interference (RNAi) is based on the ability of dsRNA species to enter a cytoplasmic

protein complex, where it is then targeted to the complementary cellular RNA and specifically degrade it. The RNA interference response features an endonuclease complex containing a siRNA, commonly referred to as an RNA-induced silencing complex (RISC), which mediates cleavage of single-stranded RNA having a sequence complementary to the antisense strand of the siRNA duplex. Cleavage of the target RNA may take place in the middle of the region complementary to the antisense strand of the siRNA duplex (Elbashir et al., *Genes Dev.*, 2001, 15(2):188-200). In more detail, longer dsRNAs are digested into short (17-29 bp) dsRNA fragments (also referred to as short inhibitory RNAs, "siRNAs") by type III RNases (DICER, DROSHA, etc.; Bernstein et al., *Nature*, 2001, 409(6818):363-6; Lee et al., *Nature*, 2003, 425(6956):415-9). The RISC protein complex recognizes these fragments and complementary mRNA. The whole process is culminated by endonuclease cleavage of target mRNA (McManus & Sharp, *Nature Rev Genet*, 2002, 3(10):737-47; Paddison & Hannon, *Curr Opin Mol Ther.* 2003, 5(3):217-24). (For additional information on these terms and proposed mechanisms, see for example Bernstein et al., *RNA* 2001, 7(11):1509-21; Nishikura, *Cell* 2001, 107(4):415-8 and PCT publication WO 01/36646).

[0161] The selection and synthesis of siRNA corresponding to known genes has been widely reported; see for example Ui-Tei et al., *J Biomed Biotechnol.* 2006; 2006: 65052; Chalk et al., *BBRC.* 2004, 319(1): 264-74; Sioud & Leirdal, *Met. Mol Biol.*; 2004, 252:457-69; Levenkova et al., *Bioinform.* 2004, 20(3):430-2; Ui-Tei et al., *Nuc. Acid Res.* 2004, 32(3):936-48. For examples of the use of, and production of, modified siRNA see Braasch et al., *Biochem.*, 2003, 42(26):7967-75; Chiu et al., *RNA*, 2003, 9(9):1034-48; PCT publications WO 2004/015107 (Atugen); WO 02/44321 (Tuschl et al), and US Patent Nos. 5,898,031 and 6,107,094.

siRNA and RNA interference

[0162] RNA interference (RNAi) is a phenomenon involving double-stranded (ds) RNA-dependent gene specific posttranscriptional silencing. Initial attempts to study this phenomenon and to manipulate mammalian cells experimentally were frustrated by an active, non-specific antiviral defense mechanism which was activated in response to long dsRNA molecules (Gil et al. *Apoptosis*, 2000, 5:107-114). Later, it was discovered that synthetic duplexes of 21 nucleotide RNAs could mediate gene specific RNAi in mammalian cells, without stimulating the generic antiviral defense mechanisms (see Elbashir et al. *Nature* 2001, 411:494-498 and Caplen et al. *PNAS USA* 2001, 98:9742-9747). As a result, small interfering RNAs (siRNAs) have become powerful tools in attempting to understand gene function.

[0163] RNA interference (RNAi) in mammals is mediated by small interfering RNAs (siRNAs) (Fire et al, *Nature* 1998, 391:806) or microRNAs (miRNAs) (Ambros, *Nature* 2004, 431(7006):350-355; Bartel, *Cell* 2004, 116(2): 281-97). The corresponding process in plants is commonly referred to as specific post-transcriptional gene silencing (PTGS) or RNA silencing and is also referred to as quelling in fungi.

[0164] A siRNA is a double-stranded RNA or modified RNA molecule which down-regulates or silences (prevents) the expression of a gene/ mRNA of its endogenous (cellular) counterpart. The mechanism of RNA interference is detailed *infra*.

[0165] Several studies have revealed that siRNA therapeutic agents are effective *in vivo* in both mammals and in humans. Bitko et al., have shown that specific siRNA molecules directed against the

respiratory syncytial virus (RSV) nucleocapsid N gene are effective in treating mice when administered intranasally (Nat. Med. 2005, 11(1):50-55). Recent reviews discussing siRNA therapeutics are available (Barik, et al., J. Mol. Med 2005, 83:764-773; Dallas and Vlassov, Med. Sci. Monitor 2006, 12(4):RA67-74; Chakraborty, Current Drug Targets 2007, 8(3):469-82; Dykxhoorn et al., Gene Therapy 2006. 13:541-552).

[0166] Mucke (IDrugs 2007 10(1):37-41) presents a review of current therapeutics, including siRNA to various targets, for the treatment of ocular diseases, for example age related macular degeneration (AMD) and glaucoma.

siRNA Structures

[0167] The selection and synthesis of siRNA corresponding to known genes has been widely reported; (see for example Ui-Tei et al., J Biomed Biotech. 2006; 2006: 65052; Chalk et al., BBRC. 2004, 319(1): 264-74; Sioud & Leirdal, Met. Mol Biol.; 2004, 252:457-69; Levenkova et al., Bioinform. 2004, 20(3):430-2; Ui-Tei et al., NAR. 2004, 32(3):936-48; De Paula et al., RNA 2007, 13:431-56).

[0168] For examples of the use of, and production of, modified siRNA see, for example, Braasch et al., Biochem. 2003, 42(26):7967-75; Chiu et al., RNA, 2003, 9(9):1034-48; PCT publications WO 2004/015107 (atugen AG) and WO 02/44321 (Tuschl et al.). US Patent Nos. 5,898,031 and 6,107,094, describe chemically modified oligomers. US Patent Publication Nos. 2005/0080246 and 2005/0042647 relate to oligomeric compounds having an alternating motif and dsRNA compounds having chemically modified internucleoside linkages, respectively.

[0169] Other modifications have been disclosed. The inclusion of a 5'-phosphate moiety was shown to enhance activity of siRNAs in Drosophila embryos (Boutla, et al., Curr. Biol. 2001, 11:1776-1780) and is required for siRNA function in human HeLa cells (Schwarz et al., Mol. Cell, 2002, 10:537-48). Amarzguioui et al., (NAR, 2003, 31(2):589-95) showed that siRNA activity depended on the positioning of the 2'-O-methyl modifications. Holen et al (NAR. 2003, 31(9):2401-07) report that an siRNA having small numbers of 2'-O-methyl modified nucleosides gave good activity compared to wild type but that the activity decreased as the numbers of 2'-O-methyl modified nucleosides was increased. Chiu and Rana (RNA. 2003, 9:1034-48) describe that incorporation of 2'-O-methyl modified nucleosides in the sense or antisense strand (fully modified strands) severely reduced siRNA activity relative to unmodified siRNA. The placement of a 2'-O-methyl group at the 5'-terminus on the antisense strand was reported to severely limit activity whereas placement at the 3'-terminus of the antisense and at both termini of the sense strand was tolerated (Czauderna et al., NAR. 2003, 31(11):2705-16; WO 2004/015107). The molecules of the present invention offer an advantage in that they are stable and active and are useful in the preparation of pharmaceutical compositions for treatment of various diseases.

[0170] PCT Patent Publication Nos. WO 2008/104978 and WO 2009/044392 to the assignee of the present invention, disclose novel siRNA structures.

[0171] PCT Publication No. WO 2008/050329 and US Serial No. 11/978,089 to the assignee of the present invention relate to inhibitors of pro-apoptotic genes.

[0172] PCT Patent Publication Nos. WO 2004/111191 and WO 2005/001043 relate to methods for enhancing RNAi.

[0173] The description discloses a method of down-regulating the expression of target gene by at least 20%, 30%, 40% or 50% as compared to a control, comprising contacting an mRNA transcript of the target gene with one or more of the compounds of the invention.

[0174] Additionally the description discloses a method of down-regulating the expression of target gene in a mammal by at least 20%, 30%, 40% or 50% as compared to a control, comprising administering one or more of the compounds of the invention to the mammal. In a preferred embodiment of the invention the mammal is a human.

[0175] In various embodiments a double stranded nucleic acid molecule of Structure (A) is down-regulating the expression of a target gene, whereby the down-regulation of the expression of a target gene is selected from the group comprising down-regulation of gene function (which is examined, e.g. by an enzymatic assay or a binding assay with a known interactor of the native gene / polypeptide, *inter alia*), down-regulation of polypeptide product of the gene (which is examined, e.g. by Western blotting, ELISA or immuno-precipitation, *inter alia*) and down-regulation of mRNA expression of the gene (which is examined, e.g. by Northern blotting, quantitative RT-PCR, in-situ hybridisation or microarray hybridisation, *inter alia*).

Delivery

[0176] The modified siRNA compound of the invention is administered as the compound *per se* (i.e. as naked siRNA) or as pharmaceutically acceptable salt and is administered alone or as an active ingredient in combination with one or more pharmaceutically acceptable carrier, solvent, diluent, excipient, adjuvant and vehicle. In some embodiments, the siRNA molecules of the present invention are delivered to the target tissue by direct application of the naked molecules prepared with a carrier or a diluent.

[0177] The term "naked siRNA" refers to siRNA molecules that are free from any delivery vehicle that acts to assist, promote or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. For example, siRNA in PBS is "naked siRNA".

[0178] Pharmaceutically acceptable carriers, solvents, diluents, excipients, adjuvants and vehicles as well as implant carriers generally refer to inert, non-toxic solid or liquid fillers, diluents or encapsulating material not reacting with the active modified siRNA compounds of the invention and they include liposomes and microspheres. For example, the siRNA compounds of the invention may be formulated with polyethylenimine (PEI), with PEI derivatives, e.g. oleic and stearic acid modified derivatives of branched PEI, with chitosan or with poly(lactic-co-glycolic acid) (PLGA). Formulating the compositions in e.g. liposomes, micro- or nano-spheres and nanoparticles, may enhance stability and benefit absorption.

[0179] Additionally, the compositions may include an artificial oxygen carrier, such as perfluorocarbons (PFCs) e.g. perfluoroctyl bromide (perflubron).

[0180] Examples of delivery systems useful in the present invention include U.S. Patent Nos. 5,225,182; 5,169,383; 5,167,616; 4,959,217; 4,925,678; 4,487,603; 4,486,194; 4,447,233; 4,447,224; 4,439,196; and 4,475,196. Many such implants, delivery systems, and modules are well known to those skilled in the art. In one specific embodiment of this invention topical and transdermal formulations are selected.

[0181] Accordingly, in some embodiments the siRNA molecules of the invention are delivered in liposome formulations and lipofectin formulations and the like and can be prepared by methods well known to those skilled in the art. Such methods are described, for example, in U.S. Pat. Nos. 5,593,972, 5,589,466, and 5,580,859.

[0182] Delivery systems aimed specifically at the enhanced and improved delivery of siRNA into mammalian cells have been developed (see, for example, Shen et al FEBS Let. 539: 111-114 (2003), Xia et al., Nat. Biotech. 20: 1006-1010 (2002), Reich et al., Mol. Vision 9: 210-216 (2003), Sorensen et al., J. Mol. Biol. 327: 761-766 (2003), Lewis et al., Nat. Gen. 32: 107-108 (2002) and Simeoni et al., NAR 31, 11: 2717-2724 (2003)). siRNA has recently been successfully used for inhibition of gene expression in primates; (for details see for example, Tolentino et al., Retina 2004. 24(1): 132-138).

[0183] Additional formulations for improved delivery of the compounds of the present invention can include non-formulated compounds, compounds covalently bound to cholesterol, and compounds bound to targeting antibodies (Song et al., Antibody mediated in vivo delivery of small interfering RNAs via cell-surface receptors, Nat Biotechnol. 2005. 23(6):709-17). Cholesterol-conjugated siRNAs (and other steroid and lipid conjugated siRNAs) can be used for delivery (see for example Soutschek et al Nature. 2004. 432:173-177; and Lorenz et al. Bioorg. Med. Chem. Lett. 2004. 14:4975-4977).

[0184] The naked siRNA or the pharmaceutical compositions comprising the chemically modified siRNA of the present invention are administered and dosed in accordance with good medical practice, taking into account the clinical condition of the individual patient, the disease to be treated, the site and method of administration, scheduling of administration, patient age, sex, body weight and other factors known to medical practitioners.

[0185] A "therapeutically effective dose" for purposes herein is thus determined by such considerations as are known in the art. The dose must be effective to achieve improvement including but not limited to improved survival rate or more rapid recovery, or improvement or elimination of symptoms and other indicators as are selected as appropriate measures by those skilled in the art. The siRNA of the invention can be administered in a single dose or in multiple doses.

[0186] In general, the active dose of compound for humans is in the range of from 1ng/kg to about 20-100 mg/kg body weight per day, preferably about 0.01 mg to about 2-10 mg/kg body weight per day, in a regimen of a single dose or a one dose per day or twice or three or more times per day for a period of 1-4 weeks or longer.

[0187] The modified siRNA compounds of the present invention can be administered by any of the conventional routes of administration. The modified siRNA compounds are administered orally, subcutaneously or parenterally including intravenous, intraarterial, intramuscular, intraperitoneally, intraocular, ocular, otic, transtympanic and intranasal administration, intratracheal instillation and intratracheal inhalation, as well as infusion techniques. Implants of the compounds are also useful.

[0188] Liquid forms are prepared for invasive administration, e.g. injection or for topical or local or non-invasive administration. The term injection includes subcutaneous, transdermal, intravenous, intramuscular, intrathecal, intraocular, transtympanic and other parental routes of administration. The liquid compositions include aqueous solutions, with and without organic co-solvents, aqueous or oil suspensions, emulsions with edible oils, as well as similar pharmaceutical vehicles. In a particular embodiment, the administration comprises intravenous administration.

[0189] In some embodiments the compounds of the present invention are formulated for non-invasive administration. In some embodiments the compounds of the present invention are formulated as eardrops for topical administration to the ear. In some embodiments the compounds of the present invention are formulated as eye drops for topical administration to the surface of the eye. Further information on administration of the compounds of the present invention can be found in Tolentino et al., Retina 2004. 24:132-138; and Reich et al., Molecular Vision, 2003. 9:210-216. In addition, in certain embodiments the compositions for use in the treatments of the present invention are formed as aerosols, for example for intranasal administration. In certain embodiments the compositions for use in the treatments of the present invention are formed as nasal drops, for example for intranasal instillation.

[0190] The therapeutic compositions of the present invention are preferably administered into the lung by inhalation of an aerosol containing these compositions / compounds, or by intranasal or intratracheal instillation of said compositions. For further information on pulmonary delivery of pharmaceutical compositions see Weiss et al., Human Gene Therapy 1999. 10:2287-2293; Densmore et al., Molecular therapy 1999. 1:180-188; Gautam et al., Molecular Therapy 2001. 3:551-556; and Shahiwala & Misra, AAPS PharmSciTech 2004. 24;6(3):E482-6. Additionally, respiratory formulations for siRNA are described in U.S. Patent Application Publication No. 2004/0063654. Respiratory formulations for siRNA are described in US Patent Application Publication No. 2004/0063654.

[0191] In certain embodiments, oral compositions (such as tablets, suspensions, solutions) may be effective for local delivery to the oral cavity such as oral composition suitable for mouthwash for the treatment of oral mucositis.

[0192] In a particular embodiment, the modified siRNA compounds of the invention are formulated for intravenous administration for delivery to the kidney for the treatment of kidney disorders, e.g. acute renal failure (ARF), delayed graft function (DGF) and diabetic retinopathy. It is noted that the delivery of the modified siRNA compounds according to the present invention to the target cells in the kidney proximal tubules is particularly effective in the treatment of ARF and DGF.

[0193] Delivery of compounds into the brain is accomplished by several methods such as, *inter alia*, neurosurgical implants, blood-brain barrier disruption, lipid mediated transport, carrier mediated influx or efflux, plasma protein-mediated transport, receptor-mediated transcytosis, absorptive-mediated transcytosis, neuropeptide transport at the blood-brain barrier, and genetically engineering "Trojan horses" for drug targeting. The above methods are performed, for example, as described in "Brain Drug Targeting: the future of brain drug development", W.M. Pardridge, Cambridge University Press, Cambridge, UK (2001).

[0194] In addition, in certain embodiments the compositions for use in the treatments of the present

invention are formed as aerosols, for example for intranasal administration.

[0195] Intranasal delivery for the treatment of CNS diseases has been attained with acetylcholinesterase inhibitors such as galantamine and various salts and derivatives of galantamine, for example as described in US Patent Application Publication No. 2006003989 and PCT Applications Publication Nos. WO 2004/002402 and WO 2005/102275. Intranasal delivery of nucleic acids for the treatment of CNS diseases, for example by intranasal instillation of nasal drops, has been described, for example, in PCT Application Publication No. WO 2007/107789.

Methods of Treatment

[0196] In one aspect the present invention relates to a modified siRNA compound of the present invention for use in a method of treating a subject suffering from a disorder associated with target gene expression comprising administering to the subject a therapeutically effective amount of said modified siRNA compound of the present invention. In preferred embodiments the subject being treated is a warm-blooded animal and, in particular, mammal including human.

[0197] "Treating a subject" refers to administering to the subject a therapeutic substance effective to ameliorate symptoms associated with a disease, to lessen the severity or cure the disease, to slow down the progress of the disease, to prevent the disease from occurring or to postpone the onset of the disease. "Treatment" refers to both therapeutic treatment and prophylactic or preventative measures, wherein the object is to prevent a disorder, to delay the onset of the disorder or reduce the symptoms of a disorder. Those in need of treatment include those already experiencing the disease or condition, those prone to having the disease or condition, and those in which the disease or condition is to be prevented. The compounds of the invention are administered before, during or subsequent to the onset of the disease or condition.

[0198] A "therapeutically effective dose" refers to an amount of a pharmaceutical compound or composition which is effective to achieve an improvement in a subject or his physiological systems including, but not limited to, improved survival rate, more rapid recovery, improvement or elimination of symptoms, delayed onset of a disorder, slower progress of disease and other indicators as are selected as appropriate determining measures by those skilled in the art.

[0199] "Respiratory disorder" refers to conditions, diseases or syndromes of the respiratory system including but not limited to pulmonary disorders of all types including chronic obstructive pulmonary disease (COPD), emphysema, chronic bronchitis, asthma and lung cancer, *inter alia*. Emphysema and chronic bronchitis may occur as part of COPD or independently. In various embodiments the present invention provides methods and compositions useful in preventing or treating primary graft failure, ischemia-reperfusion injury, reperfusion injury, reperfusion edema, allograft dysfunction, pulmonary reimplantation response and/or primary graft dysfunction (PGD) after organ transplantation, in particular in lung transplantation, in a subject in need thereof.

[0200] "Microvascular disorder" refers to any condition that affects microscopic capillaries and lymphatics, in particular vasospastic diseases, vasculitic diseases and lymphatic occlusive diseases. Examples of microvascular disorders include, *inter alia*: eye disorders such as Amaurosis Fugax (embolic or secondary to SLE), apla syndrome, Prot CS and ATIII deficiency, microvascular

pathologies caused by IV drug use, dysproteinemia, temporal arteritis, ischemic optic neuropathy (ION), anterior ischemic optic neuropathy (AION), optic neuritis (primary or secondary to autoimmune diseases), glaucoma, von Hippel Lindau syndrome, corneal disease, corneal transplant rejection cataracts, Eales' disease, frosted branch angiitis, encircling buckling operation, uveitis including pars planitis, choroidal melanoma, choroidal hemangioma, optic nerve aplasia; retinal conditions such as retinal artery occlusion, retinal vein occlusion, retinopathy of prematurity, HIV retinopathy, Purtscher retinopathy, retinopathy of systemic vasculitis and autoimmune diseases, diabetic retinopathy, hypertensive retinopathy, radiation retinopathy, branch retinal artery or vein occlusion, idiopathic retinal vasculitis, aneurysms, neuroretinitis, retinal embolization, acute retinal necrosis, Birdshot retinopathy, long-standing retinal detachment; systemic conditions such as Diabetes mellitus, diabetic retinopathy (DR), diabetes-related microvascular pathologies (as detailed herein), hyperviscosity syndromes, aortic arch syndromes and ocular ischemic syndromes, carotid-cavernous fistula, multiple sclerosis, systemic lupus erythematosus, arteriolitis with SS-A autoantibody, acute multifocal hemorrhagic vasculitis, vasculitis resulting from infection, vasculitis resulting from Behcet's disease, sarcoidosis, coagulopathies, neuropathies, nephropathies, microvascular diseases of the kidney, and ischemic microvascular conditions, *inter alia*.

[0201] Microvascular disorders may comprise a neovascular element. The term "neovascular disorder" refers to those conditions where the formation of blood vessels (neovascularization) is harmful to the patient. Examples of ocular neovascularization include: retinal diseases (diabetic retinopathy, diabetic Macular Edema, chronic glaucoma, retinal detachment, and sickle cell retinopathy); rubeosis iritis; proliferative vitreoretinopathy; inflammatory diseases; chronic uveitis; neoplasms (retinoblastoma, pseudoglioma and melanoma); Fuchs' heterochromic iridocyclitis; neovascular glaucoma; corneal neovascularization (inflammatory, transplantation and developmental hypoplasia of the iris); neovascularization following a combined vitrectomy and lensectomy; vascular diseases (retinal ischemia, choroidal vascular insufficiency, choroidal thrombosis and carotid artery ischemia); neovascularization of the optic nerve; and neovascularization due to penetration of the eye or contusive ocular injury. In various embodiments all these neovascular conditions are treated using the compounds and pharmaceutical compositions of the present invention.

[0202] "Eye disease" refers to conditions, diseases or syndromes of the eye including but not limited to any conditions involving choroidal neovascularization (CNV), wet and dry AMD, ocular histoplasmosis syndrome, angiod streaks, ruptures in Bruch's membrane, myopic degeneration, ocular tumors, retinal degenerative diseases and retinal vein occlusion (RVO). In various embodiments, conditions disclosed herein, such as DR, which are regarded as either a microvascular disorder and an eye disease, or both, under the definitions presented herein, are treated according to the methods of the present invention.

[0203] More specifically, the present invention provides compositions useful in treating a subject suffering from or susceptible to adult respiratory distress syndrome (ARDS); Chronic obstructive pulmonary disease (COPD); acute lung injury (ALI); Emphysema; Diabetic Neuropathy, nephropathy and retinopathy; diabetic macular edema (DME) and other diabetic conditions; Glaucoma; age related macular degeneration (AMD); bone marrow transplantation (BMT) retinopathy; ischemic conditions; ocular ischemic syndrome (OIS); kidney disorders: acute renal failure (ARF), delayed graft function (DGF), transplant rejection; hearing disorders (including hearing loss); spinal cord injuries; oral mucositis; dry eye syndrome and pressure sores; neurological disorders arising from ischemic or hypoxic conditions, such as hypertension, hypertensive cerebral vascular disease, a constriction or

obstruction of a blood vessel- as occurs in the case of a thrombus or embolus, angioma, blood dyscrasias, any form of compromised cardiac function including cardiac arrest or failure, systemic hypotension; stroke, disease, disorders and injury of the CNS, including, without being limited to, epilepsy, spinal cord injury, brain injury and neurodegenerative disorders, including, without being limited to Parkinson's disease, Amyotrophic Lateral Sclerosis (ALS, Lou Gehrig's Disease), Alzheimer's disease, Huntington's disease and any other disease-induced dementia (such as HIV-associated dementia for example).

[0204] The present invention relates to compounds and compositions useful in the treatment of cancer. The terms "cancer" and "cancerous" refer to or describe the physiological condition in mammals that is typically characterized by unregulated cell growth. Examples of cancer include but are not limited to, carcinoma, lymphoma, blastoma, sarcoma, and leukemia or lymphoid malignancies. Other examples of such cancers include kidney or renal cancer, breast cancer, colon cancer, rectal cancer, colorectal cancer, lung cancer including small-cell lung cancer, non-small cell lung cancer, adenocarcinoma of the lung and squamous carcinoma of the lung, squamous cell cancer (e.g. epithelial squamous cell cancer), cervical cancer, ovarian cancer, prostate cancer, liver cancer, bladder cancer, cancer of the peritoneum, hepatocellular cancer, gastric or stomach cancer including gastrointestinal cancer, gastrointestinal stromal tumors (GIST), pancreatic cancer, head and neck cancer, glioblastoma, retinoblastoma, astrocytoma, thecomas, arrhenoblastomas, hepatoma, hematologic malignancies including non-Hodgkins lymphoma (NHL), multiple myeloma and acute hematologic malignancies, endometrial or uterine carcinoma, endometriosis, fibrosarcomas, choriocarcinoma, salivary gland carcinoma, vulval cancer, thyroid cancer, esophageal carcinomas, hepatic carcinoma, anal carcinoma, penile carcinoma, nasopharyngeal carcinoma, laryngeal carcinomas, Kaposi's sarcoma, melanoma, skin carcinomas, Schwannoma, oligodendrolioma, neuroblastomas, rhabdomyosarcoma, osteogenic sarcoma, leiomyosarcomas, urinary tract carcinomas, thyroid carcinomas, Wilm's tumor, as well as B-cell lymphoma (including low grade/follicular non-Hodgkin's lymphoma (NHL); small lymphocytic (SL) NHL; intermediate grade/follicular NHL; intermediate grade diffuse NHL; high grade immunoblastic NHL; high grade lymphoblastic NHL; high grade small non-cleaved cell NHL; bulky disease NHL; mantle cell lymphoma; AIDS-related lymphoma; and Waldenstrom's Macroglobulinemia); chronic lymphocytic leukemia (CLL); acute lymphoblastic leukemia (ALL); Hairy cell leukemia; chronic myeloblastic leukemia; and post-transplant lymphoproliferative disorder (PTLD), as well as abnormal vascular proliferation associated with phakomatoses, edema (such as that associated with brain tumors), and Meigs' syndrome. "Tumor", as used herein, refers to all neoplastic cell growth and proliferation, whether malignant or benign, and all pre-cancerous and cancerous cells and tissues.

[0205] Fibrotic disorders include liver fibrosis, cirrhosis, pulmonary fibrosis including lung fibrosis (including ILF), kidney fibrosis resulting from any condition (e.g., CKD including ESRD), peritoneal fibrosis, chronic hepatic damage, fibrillogenesis, fibrotic diseases in other organs, abnormal scarring (keloids) associated with all possible types of skin injury accidental and iatrogenic (operations); scleroderma; cardiotibrosis, failure of glaucoma filtering operation; and intestinal adhesions.

[0206] Additionally, the description discloses a method of down-regulating the expression of a target gene by at least 20%, 30%, 40% or 50% as compared to a control comprising contacting target mRNA with one or more of the modified siRNA compounds of the present invention. In various embodiments the modified siRNA compound of the present invention down-regulates target gene whereby the down-regulation is selected from the group comprising down-regulation of gene function, down-

regulation of polypeptide and down-regulation of mRNA expression.

[0207] The description discloses a method of inhibiting the expression of the target gene by at least 20%, 30%, or 40%, preferably by 50%, 60% or 70%, more preferably by 75%, 80% or 90% as compared to a control comprising contacting an mRNA transcript of the target gene with one or more of the modified siRNA compounds of the invention.

[0208] In one embodiment the modified siRNA compound of the invention inhibits the target gene polypeptide, whereby the inhibition is selected from the group comprising inhibition of function (which is examined by, for example, an enzymatic assay or a binding assay with a known interactor of the native gene / polypeptide, *inter alia*), inhibition of target protein (which is examined by, for example, Western blotting, ELISA or immuno-precipitation, *inter alia*) and inhibition of target mRNA expression (which is examined by, for example, Northern blotting, quantitative RT-PCR, in-situ hybridization or microarray hybridization, *inter alia*).

[0209] In additional embodiments the invention provides a modified siRNA compound or composition of the invention for use in a method of treating a subject suffering from or susceptible to any disease or disorder accompanied by an elevated level of a mammalian or non-mammalian target gene, the method comprising administering to the subject said modified siRNA compound or composition of the invention in a therapeutically effective dose thereby treating the subject.

[0210] The present invention relates to compounds which down-regulate the expression of a mammalian target gene particularly to modified RNA compounds according to structure (A) for use in the treatment of the following diseases or conditions in which inhibition of the expression of the mammalian target gene is beneficial: ARDS; COPD; ALI; Emphysema; Diabetic Neuropathy, nephropathy and retinopathy; DME and other diabetic conditions; Glaucoma; AMD; BMT retinopathy; ischemic conditions including stroke; OIS; Neurodegenerative disorders such as Parkinson's disease, Alzheimer's disease, ALS; kidney disorders: ARF, DGF, transplant rejection; hearing disorders; spinal cord injuries; oral mucositis; cancer including hematopoietic and solid tumor cancer, dry eye syndrome and pressure sores. In another embodiment the compounds of the present invention are useful in organ storage and /or preservation before transplant.

[0211] By "exposure to a toxic agent" is meant that the toxic agent is made available to, or comes into contact with, a mammal. A toxic agent can be toxic to the nervous system. Exposure to a toxic agent can occur by direct administration, e.g., by ingestion or administration of a food, medicinal, or therapeutic agent, e.g., a chemotherapeutic agent, by accidental contamination, or by environmental exposure, e.g., aerial or aqueous exposure.

[0212] In other embodiments the chemically modified siRNA compounds of the invention are useful for treating or preventing the incidence or severity of other diseases and conditions in a subject. These diseases and conditions include, but are not limited to stroke and stroke-like situations (e.g. cerebral, renal, cardiac failure), neuronal cell death, brain injuries with or without reperfusion, spinal cord injury, chronic degenerative diseases e.g. neurodegenerative disease including, Alzheimer's disease, Parkinson's disease, Huntington's disease, multiple sclerosis, spinobulbar atrophy, prion disease and apoptosis resulting from traumatic brain injury (TBI). In an additional embodiment, the compounds of the invention are directed to providing neuroprotection, and or cerebroprotection.

[0213] Without limitation a target gene is selected from the group consisting of p53 (TP53), TP53BP2, LRDD, CYBA, ATF3, CASP2 (Caspase 2), NOX3, HRK; C1QBP, BNIP3, MAPK8; Rac1, GSK3B, CD38, STEAP4, BMP2a; GJA1, TYROBP, CTGF, SPP1, RTN4R, ANXA2, RHOA, DUOX1, SLC5A1, SLC2A2, AKR1B1, SORD, SLC2A1, MME, NRF2, SRM, REDD2 (RTP801L), REDD1 (RTP801), NOX4, MYC, PLK1, ESPL1, HTRA2, KEAP1, p66, ZNHIT1, LGALS3, CYBB (NOX2), NOX1, NOXO1, ADRB1, HI 95, ARF1, ASPP1, SOX9, FAS, FASLG, Human MLL, AF9, CTSD, CAPNS1, CD80, CD86, HES1, HES5, CDKN1B, ID1, ID2, ID3, CDKN2A, Caspase 1, Caspase 3, Caspase 4, Caspase 5, Caspase 6, Caspase 7, Caspase 8, Caspase 9, Caspase 10, Caspase 12, Caspase 14, Apaf-1, Nod1, Nod2, Ipaf, DEFCAP, RAIDD, RICK, Bcl10, ASC, TUCAN, ARC, CLARP, FADD, DEDD, DEDD2, Cryopirin, PYC1, Pyrin, TRADD, UNC5a, UNC5b, UNC5c, ZUD, p84N5, LRDD, CDK1, CDK2, CDK4, CDK5, CDK9, PITSLRE A, CHK2, LATS1, Prk, MAP4K1, MAP4K2, STK4, SLK, GSK3alpha, GSK3beta, MEKK1, MAP3K5 (Ask1), MAP3K7, MAP3K8, MAP3K9, MAP3K10, MAP3K11, MAP3K12, DRP-1, MKK6, p38, JNK3, DAPK1, DRAK1, DRAK2, IRAK, RIP, RIP3, RIP5, PKR, IRE1, MSK1, PKCalpha, PKCbeta, PKCdelta, PKCepsilon, PKCeta, PKCmu, PKCtheta, PKCzeta, CAMK2A, HIPK2, LKB1, BTK, c-Src, FYN, Lck, ABL2, ZAP70, TrkA, TrkC, MYLK, FGFR2, EphA2, AATYK, c-Met, RET, PRKAA2, PLA2G2A, SMPD1, SMPD2, SPP1, FAN, PLCG2, IP6K2, PTEN, SHIP, AIF, AMID, Cytochrome c, Smac, HtrA2, TSAP6, DAP-1, FEM-, DAP-3, Granzyme B, DIO-1, DAXX, CAD, CIDE-A, CIDE-B, Fsp27, Ape1, ERCC2, ERCC3, BAP31, Bit1, AES, Huntington, HIP1, hSir2, PHAP1, GADD45b, GADD34, RAD21, MSH6, ADAR, MBD4, WW45, ATM, mTOR, TIP49, diubiquitin/FAT10, FAF1, p193, Scythe/BAT3, Amida, IGFBP-3, TDAG51, MCG10, PACT, p52/RAP, ALG2, ALG3, presenelin-1, PSAP, AIP1/Alix, ES18, mda-7, p14ARF, ANT1, p33ING1, p33ING2, p53AIP1, p53DINP1, MGC35083, NRAGE, GRIM19, lipocalin 2, glycodelin A, NADE, Porimin, STAG1, DAB2, Galectin-7, Galectin-9, SPRC, FLJ21908, WWOX, XK, DKK-1, Fzd1, Fzd2, SARP2, axin 1, RGS3, DVL1, NFkB2, IkBalpha, NF-ATC1, NF-ATC2, NF-ATC4, zf3/ZNF319, Egr1, Egr2, Egr3, Sp1, TIEG, WT1, Zac1, Icaros, ZNF148, ZK1/ZNF443, ZNF274, WIG1, HIVEP1, HIVEP3, Fliz1, ZPR9, GATA3, TR3, PPARG, CSMF, RXRa, RARa, RARb, RARg, T3Ra, Erbeta, VDR, GR/GCCR, p53, p73alpha, p63(human [ta alpha, ta beta, ta gamma , da alpha, a beta, da gamma], 53BP2, ASPP1, E2F1, E2F2, E2F3, HIF1 alpha, TCF4, c-Myc, Max, Mad, MITF, Id2, Id3, Id4, c-Jun, c-Fos, ATF3, NF-IL6, CHOP, NRF1, c-Maf, Bach2, Msx2, Csx, Hoxa5, Ets-1, PU1/Spi1, Ets-2, ELK1, TEL1, c-Myb, TBX5, IRF1, IRF3, IRF4, IRF9, AP-2 Ipha, FKHR, FOXO1A, FKHRL1, FOXO3a, AFX1, MLLT7, Tip60, BTG1, AUF1, HNRPD, TIA1, NDG1, PCBP4, MCG10, FXR2, TNFR2, LTbR, CD40, CD27, CD30, 4-1BB, TNFRSF19, XEDAR, Fn14, OPG, DcR3, FAS, TNFR1, WSL-1, p75NTR, DR4, DR5, DR6, EDAR, TNF Ipha, FAS ligand, TRAIL, Lymphotoxin alpha, Lymphotoxin beta, 4-1BBL, RANKL, TL1, TWEAK, LIGHT, APRIL, IL-1-alpha, IL-1-beta, IL-18, FGF8, IL-2, IL-21, IL-5, IL-4, IL-6, LIF, IL-12, IL-7, IL-10, IL-19, IL-24, IFN alpha, IFN beta, IFN gamma, M-CSF, Prolactinm, TLR2, TLR3, TLR4, MyD88, TRIF, RIG-1, CD14, TCR alpha, CD3 gamma, CD8, CD4, CD7, CD19, CD28, CTLA4, SEMA3A, SEMA3B, HLA-A, HLA-B, HLA-L, HLA-Dalpha, CD22, CD33, CALL, DCC, ICAM1, ICAM3, CD66a, PVR, CD47, CD2, Thy-1, SIRPa1, CD5, E-cadherin, ITGAM, ITGAV, CD18, ITGB3, CD9, IgE Fc R beta, CD82, CD81, PERP, CD24, CD69, KLRD1, galectin 1, B4GALT1, C1q alpha, C5R1, MIP1 alpha, MIP1 beta, RANTES, SDF1, XCL1, CCCR5, OIAS/OAS1, INDO, MxA, IFI16, AIM2, iNOS, HB-EGF, HGF, MIF, TRAF3, TRAF4, TRAF6, PAR-4, IKKGamma, FIP2, TXBP151, FLASH, TRF1, IEX-1S, Dok1, BLNK, CIN85, Bif-1, HEF1, Vav1, RasGRP1, POSH, Rac1, RhoA, RhoB, RhoC, ALG4, SPP1, TRIP, SIVA, TRABID, TSC-22, BRCA1, BARD1, 53BP1, MDC1, Mdm4, Siah-1, Siah-2, RoRet, TRIM35, PML, RFWD1, DIP1, Socs1, PARC, USP7, CYLD, SERPINH1 (HSP47). Other useful target genes are genes of microbial origin.

Combination Therapy

[0214] The methods of treating the diseases disclosed herein include administering a modified double stranded nucleic acid molecule disclosed herein in conjunction or in combination with an additional inhibitor, a substance which improves the pharmacological properties of the modified siRNA compound, or an additional compound known to be effective in the treatment of a subject suffering from or susceptible to any of the hereinabove mentioned diseases and disorders, including microvascular disorder, eye disease and condition (e.g. macular degeneration), hearing impairment (including hearing loss), respiratory disorder, kidney disorder, organ transplantation, neurodegenerative disorder, spinal cord injury, brain injury, angiogenesis- and apoptosis-related condition.

[0215] The present invention thus provides in another aspect, a pharmaceutical composition comprising a combination of a therapeutic modified siRNA compound of the invention together with at least one additional therapeutically active agent. By "in conjunction with" or "in combination with" is meant prior to, simultaneously or subsequent to. Accordingly, the individual components of such a combination are administered either sequentially or simultaneously from the same or separate pharmaceutical formulations.

[0216] Combination therapies comprising known treatments for treating microvascular disorders, eye disease and conditions (e.g. macular degeneration), hearing impairments (including hearing loss), respiratory disorders, kidney disorders, organ transplantation, neurodegenerative disorders (e.g. spinal cord injury), angiogenesis- and apoptosis-related conditions, in conjunction with the modified siRNA compounds and therapies described herein are considered part of the current invention.

[0217] Accordingly, in another aspect of present description, an additional pharmaceutically effective compound is administered in conjunction with the pharmaceutical composition of the invention. In addition, the modified siRNA compounds of the invention are used in the preparation of a medicament for use as adjunctive therapy with a second therapeutically active compound to treat such conditions. Appropriate doses of known second therapeutic agents for use in combination with a chemically modified siRNA compound of the invention are readily appreciated by those skilled in the art.

[0218] In some embodiments the combinations referred to above are presented for use in the form of a single pharmaceutical formulation.

[0219] The administration of a pharmaceutical composition comprising any one of the pharmaceutically active siRNA conjugates according to the invention is carried out by any of the many known routes of administration, including intravenously, intra-arterially, subcutaneously, intra-peritoneally or intra-cerebrally, as determined by a skilled practitioner. Using specialized formulations, it is possible to administer the compositions orally or via inhalation or via intranasal instillation. In some embodiments a compound of the present invention is formulated for topical administration, including as eardrops, eye drops, dermal formulation, transdermal formulation and the like.

[0220] By "in conjunction with" is meant that the additional pharmaceutically effective compound is administered prior to, at the same time as, or subsequent to administration of the compounds or the pharmaceutical compositions of the present invention. The individual components of such a combination referred to above, therefore, can be administered either sequentially or simultaneously from the same or separate pharmaceutical formulations. As is the case for the present modified siRNA compounds, a second therapeutic agent can be administered by any suitable route, for example, by

oral, buccal, inhalation, sublingual, rectal, vaginal, transurethral, nasal, otic, ocular, topical, percutaneous (i.e., transdermal), or parenteral (including intravenous, intramuscular, subcutaneous, and intracoronary) administration.

[0221] In some embodiments, a modified siRNA compound of the invention and the second therapeutic agent are administered by the same route, either provided in a single composition as two or more different pharmaceutical compositions. However, in other embodiments, a different route of administration for the modified siRNA compound of the invention and the second therapeutic agent is either possible or preferred. Persons skilled in the art are aware of the best modes of administration for each therapeutic agent, either alone or in combination.

[0222] In various embodiments, the modified siRNA compounds of the present invention are the main active component in a pharmaceutical composition.

[0223] In another aspects, the present invention provides a pharmaceutical composition comprising two or more siRNA molecules for the treatment of a disease and for any of the diseases and conditions mentioned herein. In some embodiments the two or more siRNA molecules or formulations comprising said molecules are admixed in the pharmaceutical composition in amounts that generate equal or otherwise beneficial activity. In certain embodiments the two or more siRNA molecules are covalently or non-covalently bound, or joined together by a nucleic acid linker of a length ranging from 2-100, preferably 2-50 or 2-30 nucleotides.

[0224] In some embodiments the pharmaceutical compositions of the invention further comprise one or more additional siRNA molecule, which targets one or more additional gene. In some embodiments, simultaneous inhibition of said additional gene(s) provides an additive or synergistic effect for treatment of the diseases disclosed herein.

[0225] The treatment regimen according to the invention is carried out, in terms of administration mode, timing of the administration, and dosage, so that the functional recovery of the patient from the adverse consequences of the conditions disclosed herein is improved or so as to postpone the onset of a disorder.

[0226] The invention has been described in an illustrative manner, and it is to be understood that the terminology used is intended to be in the nature of words of description rather than of limitation.

[0227] Modifications and variations of the present invention are possible in light of the above teachings. It is, therefore, to be understood that within the scope of the appended claims, the invention can be practiced otherwise than as specifically described.

[0228] The present invention is illustrated in detail below with reference to examples, but is not to be construed as being limited thereto.

[0229] Citation of any document herein is not intended as an admission that such document is pertinent prior art, or considered material to the patentability of any claim of the present application. Any statement as to content or a date of any document is based on the information available to applicant at the time of filing and does not constitute an admission as to the correctness of such a statement.

EXAMPLES

[0230] Without further elaboration, it is believed that one skilled in the art can, using the preceding description, utilize the present invention to its fullest extent. The following preferred specific embodiments are, therefore, to be construed as merely illustrative, and not limitative of the claimed invention in any way.

[0231] Standard molecular biology protocols known in the art not specifically described herein are generally followed essentially as in Sambrook et al., Molecular cloning: A laboratory manual, Cold Springs Harbor Laboratory, New-York (1989, 1992), and in Ausubel et al., Current Protocols in Molecular Biology, John Wiley and Sons, Baltimore, Maryland (1988), and as in Ausubel et al., Current Protocols in Molecular Biology, John Wiley and Sons, Baltimore, Maryland (1989) and as in Perbal, A Practical Guide to Molecular Cloning, John Wiley & Sons, New York (1988), and as in Watson et al., Recombinant DNA, Scientific American Books, New York and in Birren et al (eds) Genome Analysis: A Laboratory Manual Series, Vols. 1-4 Cold Spring Harbor Laboratory Press, New York (1998) and methodology as set forth in US Patent Nos. 4,666,828; 4,683,202; 4,801,531; 5,192,659 and 5,272,057. Polymerase chain reaction (PCR) was carried out generally as in PCR Protocols: A Guide To Methods And Applications, Academic Press, San Diego, CA (1990). In situ (In cell) PCR in combination with Flow Cytometry is useful for detection of cells containing specific DNA and mRNA sequences (Testoni et al., Blood 1996, 87:3822.) Methods of performing RT-PCR are also well known in the art.

Example 1. Generation of sequences for siRNAs to target genes and production of the modified siRNA compounds

[0232] Using proprietary algorithms and the known sequence of any gene, optionally the pro-apoptotic genes disclosed herein, t19-mer sequences of many potential siRNAs were generated. In addition to the algorithm, some 23-mer oligomer sequences are generated by 5' and/or 3' extension of the 19-mer sequences. The antisense strand sequences that were generated using this method are fully or substantially complementary to a section of target mRNA sequence. In some embodiments the antisense sequence is fully complementary to a section of the corresponding mRNA sequence. For generating the modified siRNA compounds of the invention, the nucleotide at the 5' terminal position of the antisense strand (N)x (position 1; N¹) was substituted to generate a double stranded nucleic acid molecule of embodiments of structure (A). In other examples, the nucleotide at the 5' terminal position of the antisense strand (N)x and the nucleotide at the 3' terminal position of the sense strand (N')y were substituted to generate the double stranded nucleic acid molecule of embodiments of structure (A).

[0233] Table A provides nucleotide sequences of sense and antisense strands of compounds used in some of examples. Table A1 provides exemplary double stranded nucleic acid molecules that were generated as described to provide modified siRNA compounds according to embodiments of structure (A). The compounds shown herein are not meant to be limiting and any target RNA may be used to identify double stranded nucleic acid molecules as described herein. Table A2 provides modified siRNA compounds that were generated for comparative testing. The modified siRNA compounds in

Tables A1 and A2 target TLR2 (Homo sapiens toll-like receptor 2 (TLR2), mRNA, gi|68160956|ref|NM_003264.3|, SEQ ID NO:4) gene and CAPNS1 (calpain, small subunit 1 variant gi|51599152|ref|NM_001749.2|; SEQ ID NO:1 or calpain, small subunit 1 variant 2 gi|51599150|ref|NM_001003962.11, SEQ ID NO:2) gene. Additional compounds target human RhoA mRNA (ras homolog gene family, member A (RhoA), mRNA gi|50593005 |ref|NM_001664.2|, SEQ ID NO:3). mRNA sequences of mammalian target genes are available, for example, on the NCBI web site [<http://www.ncbi.nlm.nih.gov/>].

Table A

dsRNA molecule	SEQ ID	SENSE	SEQ ID	ANTISENSE
CAPNS1_23	5	GAGAUGACAUGGAGGUCAG	10	CUGACCUCCAUGUCAUCUC
TLR2_16	6	GAGUGGUGCAAGUAUGAAC	11	GUUCAUACUUGCACCACUC
TLR2_37	7	GGUGGGAGAACCUUAUGGUC	12	GACCAUAAGGUUCUCCACC
TLR2_46	8	AGAUAAAUGAACACCAAGAC	13	GUCUUGGUGUUCAUUAUCU
RHOA_61	9	GAUCUUCGGAAUGAUGAGA	14	UCUCAUCAUUCCGAAGAUC

Table A1: Exemplary TLR2 double stranded molecules

siRNA Compound	(N')y		
	(N)x		
TLR2_16_S1016	SEN	5' <u>GAGUGGUGCAAGUAUGAAG</u>	3'
	AS	5' UUUCAUACUUGCACCACUC	3'
TLR2_16_S1018	SEN	5' <u>GAGUGGUGCAAGUAUGAAG</u>	3'
	AS	5' CUUCAUACUUGCACCACUC	3'
TLR2_16_S1019	SEN	5' <u>GAGUGGUGCAAGUAUGAAG</u>	3'
	AS	5' dUUUCAUACUUGCACCACUC	3'
TLR2_16_S1020	SEN	5' <u>GAGUGGUGCAAGUAUGAAG</u>	3'
	AS	5' dBUU <u>CAUACUUGCACCACUC</u>	3'
TLR2_16_S1022	SEN	5' <u>GAGUGGUGCAAGUAUGAAU</u>	3'
	AS	5' UUUCAUACUUGCACCACUC	3'
TLR2_16_S1024	SEN	5' <u>GAGUGGUGCAAGUAUGAAU</u>	3'
	AS	5' CUUCAUACUUGCACCACUC	3'
TLR2_16_S1025	SEN	5' <u>GAGUGGUGCAAGUAUGAAU</u>	3'
	AS	5' dUUUCAUACUUGCACCACUC	3'
TLR2_16_S1026	SEN	5' <u>GAGUGGUGCAAGUAUGAAU</u>	3'
	AS	5' dBUU <u>CAUACUUGCACCACUC</u>	3'
TLR2_16_S1028	SEN	5' <u>GAGUGGUGCAAGUAUGAAC</u>	3'
	AS	5' CUUCAUACUUGCACCACUC	3'
TLR2_16_S1029	SEN	5' <u>GAGUGGUGCAAGUAUGAAC</u>	3'
	AS	5' dUUU <u>CAUACUUGCACCACUC</u>	3'
TLR2_16_S1030	SEN	5' <u>GAGUGGUGCAAGUAUGAAC</u>	3'

siRNA Compound	(N')y		
	(N)x		
	AS	5' dB <u>U</u> CAU <u>AC</u> U <u>U</u> GCACCACUC	3'
TLR2_16_S1032	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAA	3'
	AS	5' CUUCAUACUUGCACCACUC	3'
TLR2_16_S1033	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAA	3'
	AS	5' dUU <u>U</u> CAU <u>AC</u> U <u>U</u> GCACCACUC	3'
TLR2_16_S1034	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAA	3'
	AS	5' dB <u>U</u> CAU <u>AC</u> U <u>U</u> GCACCACUC	3'
TLR2_16_S941	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAC	3'
	AS	5' UUUCAUACtTUGCACCACUC	3'
TLR2_16_S938	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAA	3'
	AS	5' UUUCAUACTTUGCACCACUC	3'
CAPNS1_23_S938	SEN	5' G <u>A</u> G <u>A</u> UG <u>A</u> CA <u>U</u> GG <u>G</u> AG <u>U</u> CAA	3'
	AS	5' UUGACCUCCAUGUCAUCUC	3'
CAPNS1_23_S898	SEN	5' G <u>A</u> G <u>A</u> UG <u>A</u> CA <u>U</u> GG <u>G</u> AG <u>U</u> CAA	3'
	AS	5' UUGACCUCCAUGUCAUCUC-dTdT	3'
CAPNS1_23_S939	SEN	5' G <u>A</u> G <u>A</u> UG <u>A</u> CA <u>U</u> GG <u>G</u> AG <u>U</u> CAA	3'
	AS	5' CUGACCUCCAUGUCAUCUC	3'

Table A2: Compounds that were synthesized for comparative testing

siRNA Compound	(N')y		
	(N)x		
TLR2_16_S1015	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UG <u>A</u> AG	3'
	AS	5' GUUCAUACUUGCACCACUC	3'
TLR2_16_S1017	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UG <u>A</u> AG	3'
	AS	5' AUUCAUACUUGCACCACUC	3'
TLR2_16_S1021	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UG <u>A</u> AU	3'
	AS	5' GUUCAUACUUGCACCACUC	3'
TLR2_16_S1023	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UG <u>A</u> AU	3'
	AS	5' AUUCAUACUUGCACCACUC	3'
TLR2_16_S1027	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAC	3'
	AS	5' AUUCAUACUUGCACCACUC	3'
TLR2_16_S1031	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAA	3'
	AS	5' AUUCAUACUUGCACCACUC	3'
TLR2_16_S73	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAC	3'
	AS	5' GUUCAUACUUGCACCACUC	3'
TLR2_16_S939	SEN	5' G <u>A</u> G <u>U</u> GG <u>G</u> UG <u>C</u> A <u>A</u> GU <u>A</u> UGAAA	3'
	AS	5' GUUCAUACUUGCACCACUC	3'

siRNA Compound	(N')y		
	(N)x		
CAPNS1_23_S73	SEN	5' <u>GAGAUGACAU</u> <u>GGAGGU</u> CAG	3'
	AS	5' CUGACCUCCAUGUCAUCUC	3'
CAPNS1_23_S867	SEN	5' <u>GAGAUGACAU</u> <u>GGAGGU</u> CAG	3'
	AS	5' <u>CUGACCUCCAUGUCAUCUC</u> -dTdT	3'

[0234] Table A3 hereinbelow provides a code of the modified nucleotides/unconventional moieties utilized in preparing the siRNA oligonucleotides of Tables A1 and A2.

Table A3: Code of the modified nucleotides/unconventional moieties as used in the Tables herein.

Code	Modification
dT	thymidine-3'-phosphate
A	2'-O-methyladenosine-3'-phosphate
C	2'-O-methylcytidine-3'-phosphate
G	2'-O-methylguanosine-3'-phosphate
U	2'-O-methyluridine-3'-phosphate
dB	Inverted abasic deoxyribose-3'-phosphate
dU	deoxyribouridine-3'-phosphate
LdC	L-deoxyribocytidine-3'-phosphate (enantiomeric dC)
C3	C3-OH
C3C3	C3Pi-C3OH

Example 2: In vitro testing of modified siRNA compounds

Protocol for testing siRNA activity

[0235] The modified siRNA compounds according to the present invention are tested for activity as follows: About 1×10^5 rat REF52 (for TLR2 siRNA activity) or 0.6×10^5 human PC3 cells (for CAPNS1 siRNA activity) were seeded per well in 6 wells plate (30-70% confluent). After about 24h incubation, cells were transfected with dsRNA oligos (see Tables A4 and A5 below), using Lipofectamine™2000 (Invitrogen) at final concentrations of 1-40nM for TLR2 dsRNA and at final concentrations of 5-40nM for CAPNS 1 dsRNA

Table A4: TLR2_16 siRNA compounds (SEQ ID NOS:6 and 11)

TLR2_16_S73 (control compound)	SEN	5' GAGUGGUGCAAGUAUGAAC	3'	G-C base pair.
	AS	5' GUUCAUACUUGCACCACUC	3'	AS full match to target
TLR2_16_S941	SEN	5' GAGUGGUGCAAGUAUGAAC	3'	U-C. AS mismatch to target
	AS	5' GUUCAUACUUGCACCACUC	3'	

				target	
TLR2_16_S939	SEN	5'	<u>GAGUGGUGCAAGUAUGAA</u>	3'	G-A. AS match to target.
	AS	5'	GUUCAUACUUGCACCACUC	3'	Mismatch in duplex
TLR2_16_S938	SEN	5'	<u>GAGUGGUGCAAGUAUGAA</u>	3'	U-A base pair.
	AS	5'	<u>GUUCAUACUUGCACCACUC</u>	3'	AS mismatch to target

Table A5: CAPNS1_23 siRNA compounds (SEQ ID NOS:5 and 10)

CAPNS1_23_S73 (control compound)	SEN	5'	<u>GAGAUGACAUGGAGGUCA</u> G	3'	C-G base pair.
	AS	5'	CUGACCUCCAUGUCAUCUC	3'	AS full match to target
CAPNS1_23_S867	SEN	5'	<u>GAGAUGACAUGGAGGUCA</u> G	3'	C-G base pair.
	AS	5'	<u>CUGACCUCCAUGUCAUCUC</u> -dTdT	3'	AS full match to target
CAPNS1_23_S938	SEN	5'	<u>GAGAUGACAUGGAGGUCA</u> A	3'	U-A base pair.
	AS	5'	<u>UGACCUCCAUGUCAUCUC</u>	3'	AS mismatch to target
CAPNS1_23_S939	SEN	5'	<u>GAGAUGACAUGGAGGUCA</u> A	3'	C-A. AS full
	AS	5'	CUGACCUCCAUGUCAUCUC	3'	match to target

[0236] As positive control for cell transfection efficiency, REDD14-Cy3.5 labeled oligos were used. As negative control for siRNA activity scrambled (CNL) oligos were used (same concentration as mentioned in section 1.2). Transfection efficiency was tested by fluorescent microscopy, 24hrs following cell transfection.

[0237] siRNA Sample preparation: For each transfected well 3 μ l Lipofectamine2000 reagent was diluted in 250 μ l serum free medium, and incubate for 5 min at RT.

[0238] siRNA molecules were prepared as follows:

siRNA working solution:

[0239]

REDD14-Cy3.5 stock 1.5 \times 10⁶ nM (dilute 1:150 to have final concentration of 10 μ M with PBS)

CNL stock 1.6 \times 10⁶nM (dilute 1:160 to have final concentration of 10 μ M with PBS)

Target genes stock 100 μ M (dilute 1:10 to have final concentration of 10 μ M with PBS)

[0240] Samples were diluted to final concentration for TLR2, CAPNS1 and CNL siRNA oligos as

mentioned above (section 1.2 and 1.4) and 20nM of REDD14-Cy3.5 siRNA in 250 DMEM medium, calculated to final volume of 2ml per well. Lipofectamine™ 2000 was combined with siRNA (1:1 volume), sample was mixed gently and incubated at room temperature for 20 minutes.

[0241] Transfection: The Lipofectamine/siRNA complex was added onto the cells (500µl per well), the plate was rocked back and forth (2ml final volume in each well). Cells were incubated at 37°C in a CO₂ incubator.

[0242] Quantification of mRNA levels by qPCR: 48 hrs following cells transfection, cells were harvested and RNA was isolated using EZ-RNA™ kit [Biological Industries (#20-410-100). cDNA Synthesis was performed and mRNA level of was measured by qPCR. Measured mRNA quantities were normalized to the mRNA quantity of the reference gene peptidylprolyl isomerase A (cyclophilin A; CycloA)

siRNA activity Results:

[0243] TLR2_16 siRNA activity: qPCR analysis of rat TLR2 expression in REF52 cells expressing endogenous gene, following TLR2_16 siRNA transfection (data in Table A6 demonstrate residual (%) of Ctrl-un-transfected cells) rat TLR2 expression in REF52 cells.

Table A6

		Exp1	Exp2	Exp3	Exp4	Exp5	Average	Sd
Control		100	100	100	100	100	100	0
CNL	40nM	131			131		131	0
	20nM	115	84		97	89	96	12
	5nM		101			102	102	0
TLR2_16_S73 G-C base pair. AS full match to target	40nM	22			26	13	20	6
	20nM	35	17		17		23	9
TLR2_16_S938 U- A base pair. AS mismatch to target	40nM	17			16	3	12	6
	20nM	13	18		24		19	5
	5nM		24		52		38	14
	1nM		52		15		33	19
TLR2_16_S939 G-A. AS match to target. Mismatch in duplex	40nM	15			32	10	19	10
	20nM	26	39		57		41	13
	5nM		57		50		53	4
	1nM		50		7		29	21
TLR2_16_S941 U-C. AS mismatch to target. Duplex mismatch	40nM	7			18		13	5
	20nM		18	26	132		59	52
	5nM		132	28			80	52

[0244] The most active compound was TLR2_16_S938 which has AS mismatch to target mRNA and fulllly complementary duplex. In TLR2_16_S938 N¹ = U and N² =A and N¹ is mismatched to C in target mRNA.

CAPNS1_23 siRNA activity

[0245] qPCR analysis of CAPNS1 expression in human PC-3 cells expressing endogenous gene, following CAPNS1_23 siRNA transfection (data in Table A7 demonstrate residual (% of Ctrl-untransfected cells) human CAPNS1 expression in PC-3 cells.

Table A7

Exp 1		
Control		100
CNL	40nM	147
	20nM	132
	5nM	133
CAPNS1_23_S73	40nM	32
	20nM	29

to target	5nM	43
CAPNS1_23_S867	40nM	43
C-G base pair. AS full match	20nM	24
to target	5nM	38
CAPNS1_23_S938	40nM	27
U-A base pair. AS mismatch	20nM	14
to target.	5nM	22
CAPNS1_23_S939	40nM	28
C-A. AS full match to target	20nM	29
Duplex mismatch	5nM	13

[0246] The most active compound was CAPNS1_23_S938 which has AS mismatch to target mRNA and fully complementary duplex. In CAPNS1_23_S938 N¹ = U and N² = A and N¹ is mismatched to C in target mRNA.

[0247] The test compounds according to Structure (A) structures, are at least 10% more active than the control compounds, wherein the antisense strand is fully complementary to the target mRNA.

On-target and off-target testing of double stranded RNA molecules

[0248] The purpose of this study was to assess the on-target activity and potential off-target activity of control (unmodified) and test (chemically modified) double stranded nucleic acid molecules.

[0249] The two strands of a siRNA molecule have sequences with configurations that are sense and antisense with respect to the target gene mRNA. Within a cell, the antisense strand of siRNA, known as the guide strand (**GS**) is loaded into the RNA-induced silencing complex (RISC) and serves to guide the RNAi machinery to complementary sequences in target mRNA. The sense strand, known as the passenger strand (**PS**), is destroyed. When exact complementarity exists between the GS and the target mRNA the latter is cleaved by the RNaseH-like slicer activity of RISC.

[0250] In some cases a siRNA molecule may down-regulate unintended genes whose transcripts possess complementarity to the GS seed region (nucleotides at positions 2-8 [5'>3']) in the 3'-UTR. Without wishing to be bound to theory, this off-target effect may be mediated by a mechanism similar to that of target silencing by microRNAs (miRNAs). Another type of off-target activity of siRNA may occur due to loading of the sense strand (PS) into RISC. The unintended off-target effects of synthetic siRNAs can be reduced or abrogated by chemical modification of the initial siRNA sequence in the

siRNA duplex. The test molecules were designed accordingly.

[0251] Test molecules were assessed for both on-target activity (activity to target mRNA) and off-target activity (activity to mRNA other than target mRNA) in the "*guide-seed-sequence-and-passenger-strand-based activity assay*" using the psiCHECK™ (Promega) plasmid constructs. The activity of test and control molecules was tested against either a full target sequence (nucleotide sequence fully complementary to the whole 19-base sequence of either the GS or PS of test molecule) or the seed-target sequence (sequence complementary to nucleotides 1-8 [5'>3'] of either the GS or PS of test molecule).

[0252] The test molecules were at least as active against the GS full target site than was the non-modified control siRNA. Test molecules were inactive towards the PS full target site, whereas the control non-modified siRNA demonstrated activity towards the same site. Both siRNAs were inactive against the GS seed-target sequence and the PS seed-target sequence sites.

[0253] Guide strand (GS) refers to the antisense strand of a double stranded RNA that is able to enter the RISC complex and guide silencing of the target RNA.

[0254] Passenger strand (PS) refers to the sense strand of a double stranded RNA.

[0255] Seed sequence refers to nucleotides 2-8 (5'>3') of the GS and relevant for the off-target recognition.

[0256] CM (complete match) refers to a synthetic DNA fragment with nucleotide sequence fully complementary to the guide strand of the double stranded RNA molecule. This DNA fragment is cloned in 3'UTR of a reporter gene and serves as a target for RNA silencing. (Castanotto & Rossi (2009). *Nature*, 22:426-33)

[0257] SM (seed match) refers to a synthetic DNA fragment with nucleotide sequence with full complementarity to the nucleotides 1-8 (5'>3') of the guide strand of the test molecule siRNA (1st nucleotide + seed). This DNA fragment is cloned in 3'UTR of a reporter gene and serves as a target for the seed-based "off-target" silencing.

[0258] The psiCHECK™ system enables evaluation of the GS (antisense) and the PS (sense strand) to elicit targeted and off-targeted effects, by monitoring the changes in expression levels of their target sequences. Four psiCHECK™-2-based (Promega) constructs were prepared for the evaluation of target activity and potential off-target activity of each test molecule GS and PS strands. In each of the constructs one copy or three copies of either the full target or the seed-target sequence, of test molecule PS or GS, was cloned into the multiple cloning site located downstream of the Renilla luciferase translational stop codon in the 3'-UTR region. The resulting vectors were termed:

1. 1- GS-CM (guide strand, complete-match) vector containing one copy or three copies of the full target sequence (nucleotide sequence fully complementary to the whole 19-base sequence of the GS of the test molecule);
2. 2- PS-CM (passenger strand, complete-match) vector containing one copy or three copies of the full target sequence (nucleotide sequence fully complementary to the whole 19-base sequence of the PS of the test molecule);
3. 3- GS-SM (guide strand, seed-match) vector containing one copy or three copies of the seed

region target sequence (sequence complementary to nucleotides 1-8 of the GS of the test molecule);

4. 4- PS-SM (passenger strand, seed-match) vector containing one copy or three copies of the seed region target sequence (sequence complementary to nucleotides 1-8 of the PS of the test molecule).

[0259] The target sequences, with or without nucleotide substitutions at position 19 (position 1 of AS) were cloned downstream to the coding region of the Renilla luciferase reporter gene. The RNAi or seed-mediated activity of a test molecule toward any of these sequences results either in cleavage and subsequent degradation of the fusion mRNA (GS) or in translational inhibition (PS). In both cases protein expression is attenuated.

Cloning of test molecule GS and PS Seed and Full Target Sites

[0260] A single copy of the relevant target cloned in the 3'UTR of the reporter mRNA, Renilla Luciferase in the psICHECK™-2 (Promega) vector. There are multiple cloning sites in the vector. Typical cloning sites that were used are Xhol and NotI. Vector was prepared for cloning using standard molecular biology techniques. Each strand of CM and SM was chemically synthesized and annealed by heating to 100°C and cooled to room temperature. Ligation was carried out for 3 hours using standard molecular biology techniques. Ligated plasmids were transformed into E. coli DH5a cells

[0261] Resulting colonies were screened for presence of plasmid constructs by colony-PCR using relevant primers. Each of the plasmids (vectors) was purified from one positive colony and its sequence was verified.

Transfection of vectors into Human HeLa Cells.

[0262] About 1.3-2x10⁶ human HeLa cells (ATCC, Cat#CCL-2) were inoculated per 10 cm dish. Cells were then incubated in 37±1°C, 5% CO₂ for 24 hours. Growth medium was replaced one day post inoculation by 8 mL fresh growth medium prepared. Each cell-containing plate was transfected with one of the vectors, using Lipofectamine™2000 reagent (Invitrogen) as follows:

In an Eppendorf tube, 15 µL Lipofectamine™2000 reagent was diluted in 1000 µL DMEM medium and incubated for 5 minutes at room temperature (RT). In a second Eppendorf tube, a vector was diluted to reach a final concentration of 15µg in 1000 µL DMEM medium. The diluted Lipofectamine™2000 reagent was mixed gently with the diluted DNA vector sample and incubated for 20-40 minutes at RT. Following incubation, DNA/ Lipofectamine™2000was added (to reach a 2000 µL final volume) to the cells. The plates were gently rocked. Plates were incubated for 5 hours at 37±1°C and 5% CO₂.

Following a 5-hour incubation, cells were re-plated in a 96-well plate at final concentration of 5x10³ cells per well in 80 µL growth medium. 16 hours later, cells were transfected with test or control molecules using Lipofectamine™2000. Duplicate transfections of each siRNA concentration were performed, as described below: Lipofectamine™2000 was prepared in excess to suffice for 170 wells:

85 μ L of Lipofectamine TM2000 were mixed with 3400 μ L (3.4mL) of DMEM medium and incubated for 5 minutes at RT.

[0263] Preparation of test and control molecule working solutions: working solutions at various concentrations was prepared by diluting a 10 μ M stock solution. This dilution series was prepared for the generation of final transfection concentrations ranging between 0.0095 nM and 100 nM in 100 μ L DMEM transfection medium (0.0095, 0.019, 0.039, 0.07, 0.15, 0.31, 0.625, 1.25, 2.5, 5.0, 20.0, 100.0).

[0264] 100 μ L aliquots of the diluted Lipofectamine 2000 were mixed gently with 100 μ L of each of the diluted test molecule or control molecule working solutions (above) and mixtures were incubated for 20-40 minutes at RT. Following incubation, 20 μ L of the siRNA/ Lipofectamine TM2000 mixture were added on top of the cells (pre-incubated with 80 μ L of cell-culture medium above). The plates were gently rocked. Cells were incubated for 48 hours at 37 \pm 1°C and 5% CO₂.

Determination of Renilla Luciferase Activity in Transfected Cells

[0265] The psiCHECKTM-2 vector enables monitoring of changes in expression of a target sequence fused to the Renilla luciferase reporter gene. The test/control molecule target sequence is cloned into the 3'-untranslated region (3'-UTR) of Renilla luciferase. Measuring the decrease in Renilla luciferase activity thus provides a convenient way of monitoring activity. In addition, the psiCHECKTM-2 vector contains a second reporter gene, Firefly luciferase, transcribed under a different promoter, which allows for normalization of Renilla luciferase expression.

[0266] 48 Hours following test or control molecule, transfection Renilla and FireFly Luciferase activities were measured in each of the siRNA transfected samples, using Dual-Luciferase[®] Assay kit (Promega) according to manufacturer procedure:

Medium was completely removed from cells and cells were then lysed by the addition of 40 μ L/well IX Luciferase lysis solution. Plates were then frozen (-80°C) and thawed at RT. Cell lysates were suspended by pipetting several times and aliquots of 12.5 μ L of each sample were transferred to a separate 96-well plate. 50 μ L Luciferase substrate (LARII) was added to each extract and Firefly Luciferase activity was measured by Absorbance, Fluorescence and Luminescence Reader (Perkin Elmer, VictorTM 1240). 50 μ L of Stop&Glo Reagent was added to each of the samples and Renilla Luciferase activity was measured immediately. Renilla Luciferase activity value was divided by Firefly Luciferase activity value for each sample (normalization) Renilla luciferase activity is finally expressed as the percentage of the normalized activity value in tested sample relative to the normalized value obtained in cells transfected with the corresponding psiCHECKTM-2 plasmid in the absence of test or control molecules.

IC50 Calculation

[0267] The IC50 values of test and control molecule activity against the GS_CM site were determined by constructing a dose-response curve using the activity results obtained with the various final siRNA concentrations. The dose response curve was constructed by plotting the relative normalized values of Renilla luciferase activity versus the logarithm of transfected siRNA concentration. The curve was

calculated by fitting the best sigmoid curve to the measured data. The methods for the sigmoid fit is called 3-point curve fit.

$$Y = Bot + \frac{100 - Bot}{1 + 10^{\frac{LogIC50 - X}{HillSlope}}}$$

Where:

Y is the residual caspase 2 mRNA response,

X is the logarithm of transfected siRNA concentration,

Bot is the Y value at the bottom plateau,

$LogIC50$ is the X value when Y is halfway between bottom and top plateaus and $HillSlope$ is the steepness of the curve.

Results

[0268] For the evaluation of the potential off-target activity of each strand of a double stranded RNA molecule strands, the "guide-seed-sequence-and-passenger-strand-based activity assay" was employed using the psiCHECK (Promega) plasmid constructs. The measurement of Renilla activity provides a convenient way of monitoring double stranded RNA activity.

Measurement of Target Renilla Luciferase Protein Activity

[0269] The activity of both test and control molecules against the four target sequences (GS-CM, guide strand complete match; GS-SM, guide strand seed match; PS-CM, passenger strand complete match; and PS-SM, passenger strand seed match) was assessed at the protein level by measuring the relative activity of the Renilla luciferase reporter protein in cells transfected with various concentrations of molecule. Each assay was repeated three times. The IC50 value of test molecule activity against the GS-CM target site was determined by construction of concentration-response plots.

[0270] Both test and control molecules were active against the target GS-CM site in a dose response.

[0271] Both siRNAs were inactive against the GS-SM and the PS-SM sites.

[0272] Figures 2-10 and Tables 2-10 below show activity as measured by residual target for double stranded molecules targeting the complete match of a target sequence which has a substitution at position 19 (equivalent of position 1 of the antisense) to incorporate one of the ribonucleotides, adenosine (A or rA), cytidine (C or rC), guanosine (G or rG), ribothymidine (rT, also 5'methyluridine) and uridine (U or rU); or deoxyadenosine (dA), deoxycytidine (dC), deoxyguanosine (dG), thymidine (dT) and deoxyuridine (dU); or 2'O methylated adenosine (mA), 2'O methylated cytidine (mC), 2'O methylated guanosine (mG), 2'O methylated uridine (mU). "1st pos AS" refers to position 1 of the antisense strand (5' terminal nucleotide).

Table 2: TLR2_16 siRNA (blunt ended with alternating 2'-OMe on both strands)

		1st pos AS -mC	1st pos AS -mG	1st pos AS - mA	1st pos AS - dU
siRNA concentration	0.4 nM	Target G 81	75	68	58
		Target U 91	93	81	59
	4 nM	Target G 85	61	50	38
		Target U 84	78	56	40
20 nM	Target G 57	55	38	32	
	Target U 62	62	35	32	

[0273] The data shown in Table 2 is presented in Figure 2. Methylated nucleotides at position 1 of the antisense strand show reduced target knockdown, irrespective of base pairing.

Table 3 TLR2_37 siRNA unmodified with C3-C3 overhangs or dTdT overhangs

		dTdT overhang	C3-C3 overhang				
	Position 19 target	1st pos AS -U	1st pos AS -U	1st pos AS - dU	1st pos AS - dT	1st pos AS - mU	1st pos AS - A
siRNA concentration	0.04 nM	Target A 21	18	17	16	48	18
		Target C 20	15	22	21	39	6
	0.4 nM	Target G 35	30	27	27	60	24
		Target U 33	33	28	32	64	27
4 nM	Target A 13	8	8	9	23	11	
	Target C 12	9	9	11	18	10	
	Target G 17	13	12	13	34	16	
	Target U 19	17	12	16	43	14	

[0274] The data in Table 3 is presented in Figure 3. Table 3 and Figure 3 show that a 2'OMe ribonucleotide in the 1st position of AS reduces siRNA activity regardless of the nucleotide in the matching position of the target.

Table 4: TLR2_37 siRNA unmodified with C3-C3 overhangs or dTdT overhangs

		dTdT overhang	C3-C3 overhang			
		1st pos AS -U	1st pos AS -U	1st pos AS - dU	1st pos AS - dT	
siRNA concentration	0.04 nM	Target A 21	18	17	16	
		Target C 20	15	22	21	
	0.4 nM	Target G 35	30	27	27	
		Target U 33	33	28	32	
0.4 nM	Target A 13	8	8	9	9	
	Target C 12	9	9	11	11	
	Target G 17	13	12	12	13	
	Target U 19	17	12	12	16	

[0275] The data in Table 4 is presented in Figure 4. Table 4 and Figure 4 show that

1. siRNA with highest activity is observed when U (dU or dT) in the 1st position of AS is mismatched to C in the target or is matched to A in the target.

2. 2. If U in the 1st position of AS of siRNA (or dU or dT) is mismatched to U in the target, siRNA activity is reduced

3. 3. If U in the 1st position of AS is wobble-paired with G in target, siRNA activity is reduced.

Table 5 TLR2_37 siRNA (non-modified with dTdT overhangs) - 0.04nM

			1st pos AS - G	1st pos AS - U
SiRNA concentration	0.04 nM	Target A	32	21
		Target C	29	20
		Target G	33	35
		Target U	37	33
	0.4 nM	Target A	14	13
		Target C	13	12
		Target G	17	17
		Target U	21	19
	4 nM	Target A	10	10
		Target C	13	8
		Target G	10	13
		Target U	11	12
	20 nM	Target A	9	11
		Target C	8	8
		Target G	8	11
		Target U	10	11
	100 nM	Target A	8	7
		Target C	5	7
		Target G	7	8
		Target U	5	7

[0276] The data in Table 5 is presented in Figure 5. Table 5 and Figure 5 show that

1. U in the 1st position of the AS produced better results with its matched A or mismatched C in the target than with mismatched U or wobble-paired G.
2. G in the 1st position of the AS showed similar results with all possible nucleotides in the matching target position - mismatches or wobble-pairing did not improve activity.

Table 6: TLR2_46 siRNA (unmodified with C3-C3 or dTdT overhangs)

		dTdT overhang	C3-C3 overhang					
			1st pos AS - U	1st pos AS - U	1st pos AS - dU	1st pos AS - dT	1st pos AS - mU	1st pos AS - A
SiRNA concentration	0.04 nM	Target A	70	83	61	69	79	52
		Target C	56	78	49	66	85	58
		Target U	68	85	57	69	95	67
0.4 nM	0.4 nM	Target A	41	30	33	36	83	37
		Target C	35	39	33	41	86	34
		Target U	56	46	43	36	94	48
	4 nM	Target A	13	16	17	20	39	14
		Target C	15	17	11	24	48	13
20 nM	20 nM	Target U	19	21	17	14	59	19
		Target A	8	7	8	8	28	10
	20 nM	Target C	7	10	7	8	27	9
		Target U	10	12	10	6	40	13

[0277] The data in Table 6 is presented in Figure 6. Table 6 and Figure 6 show that substitution of U

with mU reduces activity

Table 7 TLR2_46 siRNA (non-modified with dTdT overhangs)

		1st pos AS - G		1st pos AS - U	
siRNA concentration	0.4 nM	Target A	53	41	
		Target C	61	35	
		Target U	68	56	
	4 nM	Target A	28	13	
		Target C	28	15	
		Target U	32	19	
siRNA concentration	20 nM	Target A	16	8	
		Target C	15	7	
		Target U	17	10	
	100 nM	Target A	16	8	
		Target C	15	7	
		Target U	17	10	

[0278] The data in Table 7 is presented in Figure 7. Table 7 and Figure 7 show that regardless of the nucleotide type of the matching target position, U in the 1st position of AS produced better KD (knock-down). All cases mean - regardless of the complementarity between siRNA and target in the 1st position of the guide strand, or mismatch or wobble pair.

Table 8 RHOA_61 siRNA (non-modified/C3-C3 or dTdT overhangs)

		dTdT overhang	C3-C3 overhang					
			1st pos AS - U	1st pos AS - U	1st pos AS - dU	1st pos AS - dT	1st pos AS - mU	1st pos AS - A
siRNA concentration	0.04 nM	Target A	35	50	34	33	76	33
		Target C	32	38	33	32	86	34
		Target G		41		51		49
	0.4 nM	Target U		53		34		36
		Target A	26	24	26	25	74	24
		Target C	26	22		22	66	23
siRNA concentration	4 nM	Target G		26	26	38		35
		Target U	36			27		27
		Target A	22	22	24	17	37	20
	0.4 nM	Target C	18	19	21	17	34	18
		Target G		21		31		25
		Target U		30		22		22

[0279] The data in Table 8 is presented in Figure 8.

Table 9 RHOA_61 siRNA (non-modified/C3-C3 or dTdT overhangs)

		dTdT overhang	C3-C3 overhang					
			1st pos AS - U	1st pos AS - U	1st pos AS - dU	1st pos AS - dT	1st pos AS - mU	1st pos AS - A
concentration	0.04 nM	Target A	35	50	34	33	76	33
		Target C	32	38	33	32	86	34
		Target G		41		51		49
	0.4 nM	Target U		53		34		36
		Target A	26	24	26	25	74	24
		Target C	26	22		22	66	23
concentration	0.4 nM	Target G		26	26	38		35
		Target U	36			27		27
		Target A	22	22	24	17	37	20

siRNA c		Target C	18	19	21	17	34	18
	4 nM	Target G		21		31		25
		Target U		30		22		22

[0280] The data in Table 9 is presented in Figure 9. Table 9 and Figure 9 show that siRNAs with U in the 1st position of the AS are more active than those with G in the 1st position regardless of the nucleotide in the matching position of the target (match, mismatch, wobble). The results cannot be explained by Wobble base-pairing since having G in siRNA and U in the target is not equivalent to having U in siRNA and G in the target

Table 10 RhoA_61 siRNA (unmodified with dTdT overhangs)

			1st pos AS - G	1st pos AS - U
siRNA concentration	0.04 nM	Target A	52	35
		Target C	52	32
		Target G	90	
		Target U	66	
	0.4 nM	Target A	49	26
		Target C	51	26
		Target G	84	
		Target U	49	
	4 nM	Target A	23	22
		Target C	22	18
		Target G	38	
		Target U	28	
	20 nM	Target A	21	20
		Target C	18	17
		Target G	29	
		Target U	21	
	100 nM	Target A	18	19
		Target C	19	16
		Target G	25	
		Target U	20	

[0281] The data in Table 10 is presented in Figure 10.

Example 3: Model systems of acute renal failure (ARF)

[0282] ARF is a clinical syndrome characterized by rapid deterioration of renal function that occurs within days. Without being bound by theory the acute kidney injury may be the result of renal ischemia-reperfusion injury such as renal ischemia-reperfusion injury in patients undergoing major surgery such as major cardiac surgery. The principal feature of ARF is an abrupt decline in glomerular filtration rate (GFR), resulting in the retention of nitrogenous wastes (urea, creatinine). Recent studies, support that apoptosis in renal tissues is prominent in most human cases of ARF. The principal site of apoptotic cell death is the distal nephron. During the initial phase of ischemic injury, loss of integrity of the actin cytoskeleton leads to flattening of the epithelium, with loss of the brush border, loss of focal cell contacts, and subsequent disengagement of the cell from the underlying substratum.

[0283] The compounds of the invention are tested for efficacy in treating ischemia reperfusion injury in an animal model of ischemia-reperfusion-induced ARF.

Example 4: Model systems of pressure sores or pressure ulcers

[0284] Pressure sores or pressure ulcers including diabetic ulcers, are areas of damaged skin and tissue that develop when sustained pressure (usually from a bed or wheelchair) cuts off circulation to vulnerable parts of the body, especially the skin on the buttocks, hips and heels. The lack of adequate blood flow leads to ischemic necrosis and ulceration of the affected tissue. Pressure sores occur most often in patients with diminished or absent sensation or who are debilitated, emaciated, paralyzed, or long bedridden. Tissues over the sacrum, ischia, greater trochanters, external malleol, and heels are especially susceptible; other sites may be involved depending on the patient's situation.

[0285] The compounds of the invention are tested for efficacy in treating pressure sores, ulcers and similar wounds in, *inter alia*, the mouse model as described in Reid et al., J Surg. Res. 116:172-180, 2004 or in the rabbit model as described by Mustoe et al, JCI, 1991. 87(2):694-703; Ahn and Mustoe, Ann Pl Surg, 1991. 24(1):17-23.

Example 5: Model systems of chronic obstructive pulmonary disease (COPD)

[0286] Chronic obstructive pulmonary disease (COPD) is characterized mainly by emphysema, which is permanent destruction of peripheral air spaces, distal to terminal bronchioles. Emphysema is also characterized by accumulation of inflammatory cells such as macrophages and neutrophils in bronchioles and alveolar structures. Emphysema and chronic bronchitis may occur as part of COPD or independently.

[0287] The compounds of the invention are tested for efficacy in treating COPD/ emphysema/chronic bronchitis in, *inter alia*, animal models such as those disclosed as follows:

Starcher and Williams, 1989. Lab. Animals, 23:234-240; Peng, et al., 2004.; Am J Respir Crit Care Med, 169:1245-1251; Jeyaseelan et al., 2004. Infect. Immunol, 72: 7247-56. Additional models are described in PCT patent publication WO 2006/023544 assigned to the assignee of the present application.

Example 6: Model systems of spinal cord injury

[0288] Spinal cord injury, or myelopathy, is a disturbance of the spinal cord that results in loss of sensation and/or mobility. The two common types of spinal cord injury are due to trauma and disease. Traumatic injury can be due to automobile accidents, falls, gunshot, diving accidents *inter alia*, and diseases which can affect the spinal cord include polio, spina bifida, tumors and Friedreich's ataxia.

[0289] siRNA is injected into the spinal cord following spinal cord contusion and in uninjured rats. Sagittal cryosections are produced and immunostaining using four different groups of antibodies is performed in order to determine whether uptake has occurred in neurons, astroglia, oligodendroglia and/or macrophages/microglia. Markers for neurons include NeuN, or GAP43; markers for astroglia and potential neural stem cells include GFAP, nestin or vimentin; markers for oligodendroglia include NG2 or APC; markers for macrophages/microglia include ED1 or Iba-1 (Hasegawa et al., 2005. Exp

Neurol 193:394-410).

[0290] Rats are injected with two different doses of siRNA (1 μ g/ μ l, 10 μ g/ μ l) and are left for 1 and 3 days before sacrifice. The results indicate that siRNA to spinal cord injury target genes increases motoneuron recovery.

Example 7: Model systems of Glaucoma

[0291] The compounds of the invention are tested for efficacy in treating or preventing glaucoma in the animal model, for example, as described by Pease et al., J. Glaucoma, 2006, 15(6):512-9 (Manometric calibration and comparison of TonoLab and TonoPen tonometers in rats with experimental glaucoma and in normal mice).

Example 7A: Model systems of Ischemic optic neuropathy (ION)

[0292] An animal model for Ischemic optic neuropathy was established in adults Wistar rats using a protocol of optic nerve crush injury. Seven days prior to the optic nerve crush, the retinal ganglion cells (RGC) are selectively labelled by application of the retrograde tracer FluoroGold (2%, Fluorochrome, Englewood, CO) to the superior colliculus. The tracer is transported by retrograde transport along RGC axons resulting in complete and specific labelling of all RGCs within 1 week post injection of the fluorescent tracer. The animals are subjected to the optic nerve crush injury 7 days post retrograde tracing. The orbital optic nerve is exposed through a supraorbital approach and all axons in the optic nerve were transected by crushing with forceps for 10 seconds, 2 mm from the lamina cribrosa. A single dose of 20 μ g/ 5 μ l of PBS of the test modified siRNA compound is microinjected into the vitreous body 2 mm anterior to the nerve head, using a glass micropipette at the time of the optic nerve crush.

[0293] The survival of RGCs is determined 7 days following the optic nerve crush by counting FluoroGold-labelled RGCs on flat-mounted retinas. The experimental animals are perfused transcardially with 4% paraformaldehyde at 1 week after the optic nerve crush. Both retinas are dissected out, fixed for an additional 30 min and flat-mounted on a glass slide for ganglion cell layer quantification. The number of fluorescent RGCs is counted in 16 distinct areas in each retina and the percent of survival of the RGCs is determined compared to samples obtained from rats which did not undergo optic nerve crush injury at all or samples obtained from rats which were injected with PBS, control siRNA or GFP siRNA along with the optic nerve crush injury. Microglia cells that may have incorporated FluoroGold after phagocytosis of dying RGCs were distinguished by their characteristic morphology and excluded from quantitative analyses.

[0294] Another model of optic nerve axotomy where the entire population of RGCs are axotomized by transecting the optic nerve close to the eye is utilized. (Cheng L, et al. J. Neurosci. 2002;22:3977-3986).

Example 8: Model systems of ischemia/reperfusion injury following lung transplantation in rats

[0295] The compounds of the invention are tested for efficacy in treating ischemia/reperfusion injury or hypoxic injury following lung transplantation in one or more of the experimental animal models, for example as described by Mizobuchi et al., 2004. *J. Heart Lung Transplant*, 23:889-93; Huang, et al., 1995. *J. Heart Lung Transplant*. 14: S49; Matsumura, et al., 1995. *Transplantation* 59: 1509-1517; Wilkes, et al., 1999. *Transplantation* 67:890-896; Naka, et al., 1996. *Circulation Research*, 79: 773-783.

Example 9: Model systems of Acute Respiratory Distress Syndrome

[0296] The compounds of the invention are tested for efficacy in treating acute respiratory distress syndrome in *inter alia* in the animal model described by Chen et al (*J Biomed Sci*. 2003;10(6 Pt 1):588-92). Modified siRNA compounds that target genes including CYBA, HRK, BNIP3, MAPK8, MAPK14, RAC1, GSK3B, P2RX7, TRPM2, PARG, SPP1, and DUOX1 are tested in this animal model.

Example 10: Model systems of hearing loss conditions

(i) Chinchilla model of carboplatin-induced or cisplatin-induced cochlea hair cell death

[0297] Chinchillas are pre-treated by direct administration of specific siRNA in saline to the left ear of each animal. Saline is given to the right ear of each animal as placebo. Two days following the administration of the specific modified siRNA compounds of the invention, the animals are treated with carboplatin (75 mg/kg ip) or cisplatin (intraperitoneal infusion of 13mg/kg over 30 minutes). After sacrifice of the chinchillas (two weeks post carboplatin treatment) the % of dead cells of inner hair cells (IHC) and outer hair cells (OHC) is calculated in the left ear (siRNA treated) and in the right ear (saline treated). It is calculated that the % of dead cells of inner hair cells (IHC) and outer hair cells (OHC) is lower in the left ear (siRNA treated) than in the right ear (saline treated).

(ii) Chinchilla model of acoustic-induced cochlea hair cell death

[0298] The activity of specific siRNA in an acoustic trauma model is studied in chinchilla. The animals are exposed to an octave band of noise centered at 4 kHz for 2.5h at 105 dB. The left ear of the noise-exposed chinchillas is pre-treated (48 h before the acoustic trauma) with 30 µg of siRNA in ~10 µL of saline; the right ear is pre-treated with vehicle (saline). The compound action potential (CAP) is a convenient and reliable electrophysiological method for measuring the neural activity transmitted from the cochlea. The CAP is recorded by placing an electrode near the base of the cochlea in order to detect the local field potential that is generated when a sound stimulus, such as click or tone burst, is abruptly turned on. The functional status of each ear is assessed 2.5 weeks after the acoustic trauma. Specifically, the mean threshold of the compound action potential recorded from the round window is determined 2.5 weeks after the acoustic trauma in order to determine if the thresholds in the siRNA-treated ear are lower (better) than the untreated (saline) ear. In addition, the level of inner and outer hair cell loss is determined in the siRNA-treated and the control ear.

[0299] Similar models are used in mice and rats. The modified siRNA compounds that target genes including BNIP3, CAPNS, HES1, HES5, NOX3, ID1-3, HRK, ASPP2 (TP53BP), CASP2, RAC1, HTRA2, CDKN1B are tested in these and other animal model of hearing loss and hearing regeneration.

Example 11. Animal models of Osteoarthritis (OA)

[0300] Collagen induced arthritis (CIA): CIA in mice is described in Trentham et al. (1977. J. Exp. Med. 146: 857-868). Adjuvant-induced arthritis (AA):AA is described in Kong et al., (1999. Nature, 402:304-308). A meniscectomy model is described in Han et al., (1999. Nagoya J Med Sci 62(3-4):115-26).

[0301] The effect of different siRNA inhibitors, such as siRNA to SSP1, on different parameters related to OA such as chondrocyte proliferation, terminal differentiation and development of arthritis, is evaluated using one or more of the above models, in addition to in vitro models known in the art. Modified siRNA compounds directed to proapoptotic genes, in particular to SSP1, are tested in these animal models which show that the modified siRNA compounds treat and/or prevent OA and thus may be used to treat this condition.

Example 12: Rat model systems for transplantation-associated acute kidney injury

[0302] Warm ischemia - In test rats a left nephrectomy is performed, followed by auto transplantation that results in a warm kidney graft preservation period of 45 minutes. Following auto transplantation, a right nephrectomy is performed on the same animal. Chemically modified siRNA to a target is administered intravenously via the femoral vein either before harvesting of the kidney graft (mimicking donor treatment) ("pre"), or after the kidney autotransplantation (mimicking recipient treatment), or both before harvest and after transplantation (combined donor and recipient treatment) ("pre-post").

[0303] Cold ischemia - A left nephrectomy is performed on a donor animal, followed by a cold preservation (on ice) of the harvested kidney for a period of 5 hours. At the end of this period, the recipient rat will undergo a bilateral nephrectomy, followed by transplantation of the cold-preserved kidney graft. The total warm ischemia time (including surgical procedure) is about 30 minutes. Chemically modified siRNA is administered intravenously via the femoral vein, either to the donor animal prior to the kidney harvest ("pre"), or to the recipient animal 15 minutes ("post 15 min") or 4 hours (post 4 hrs) post-transplantation.

[0304] To assess the efficacy of the modified siRNA compounds of the present invention in improving post-transplantation renal function, serum creatinine levels are measured on days 1, 2, and 7 post-transplantation in both warm and cold ischemia models.

SEQUENCE LISTING

[0305]

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REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- [WO2008104978PCT \[0002\]](#)
- [WO2009044392PCT \[0002\] \[0141\]](#)
- [WO2005001043A \[0006\] \[0172\]](#)
- [EP0586520B1 \[0061\]](#)
- [EP0618925B1 \[0061\]](#)
- [US6586238B \[0075\]](#)
- [US6602858B \[0075\]](#)
- [WO2006023544A \[0141\] \[0287\]](#)
- [WO2007084684PCT \[0141\]](#)

- WO2008050329PCT [0141]
- WO2007141796PCT [0141]
- WO2008106102PCT [0141]
- WO2008152636PCT [0141]
- WO2009001359PCT [0141]
- WO2009090639PCT [0141]
- WO9323569A [0150]
- US6121426A [0151]
- WO0044895A [0159]
- WO0049035A [0159]
- WO0063364A [0159]
- WO0136641A [0159]
- WO0136646A [0159] [0160]
- WO9932619A [0159]
- WO0044914A [0159]
- WO0129058A [0159]
- WO0175184A [0159]
- WO2004015107A [0161] [0168] [0169]
- WO244321A [0161] [0168]
- US5898031A [0161] [0168]
- US6107094A [0161] [0168]
- US20050080246A [0168]
- US20050042647A [0168]
- WO2008104978A [0170]
- WO2009044392A [0170]
- WO2008050329A [0171]
- US978089 [0171]
- WO2004111191A [0172]
- US5225182A [0180]
- US5169383A [0180]
- US5167616A [0180]
- US4959217A [0180]
- US4225678A [0180]
- US4487603A [0180]
- US4486194A [0180]
- US4447233A [0180]
- US4447224A [0180]
- US4439196A [0180]
- US4475196A [0180]
- US5593972A [0181]
- US5589466A [0181]
- US5580859A [0181]
- US20040063654 [0190] [0190]
- US2006003989 [0195]
- WO2004002402A [0195]
- WO2005102275PCT [0195]
- WO2007107769A [0195]

- [US4666828A \[0231\]](#)
- [US4683202A \[0231\]](#)
- [US4801531A \[0231\]](#)
- [US5192659A \[0231\]](#)
- [US5272057A \[0231\]](#)
- [WO61264668A \[0305\]](#)
- [WO61295721A \[0305\]](#)
- [WO61372072A \[0305\]](#)

Non-patent literature cited in the description

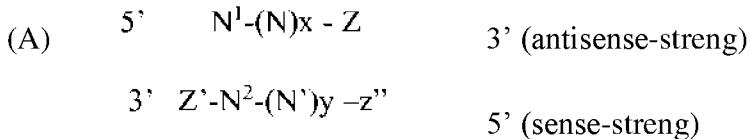
- **UI-TEI et al.** Nucleic Acids Research, 2008, vol. 36, 2136-2151 [\[0003\]](#)
- **HUANG et al.** Nucleic Acids Research, 2009, vol. 37, 7560-7569 [\[0004\]](#)
- **DU et al.** Nucleic Acids Research, 2005, vol. 33, 1671-1677 [\[0005\]](#)
- **UI-TEI et al.** Nucleic Acids Research, 2004, vol. 32, 936-948 [\[0007\]](#)
- **ELMEN et al.** NAR, 2005, vol. 33, 1439-447 [\[0064\]](#)
- **TAKEI et al.** JBC, 2002, vol. 277, 2623800-06 [\[0065\]](#)
- **BEAUCAGEIYERT** Tetrahedron, 1992, vol. 48, 2223-2311 [\[0148\]](#)
- **BEAUCAGEIYERT** Tetrahedron, 1993, vol. 49, 6123-6194 [\[0148\]](#)
- **CARUTHERS** Methods Enzymol., 1987, vol. 154, 287-313 [\[0148\]](#)
- **ECKSTEIN** Ann. Rev. Biochem., 1985, vol. 54, 367-402 [\[0148\]](#)
- **USMAN et al.** J. Am. Chem. Soc., 1987, vol. 109, 7845- [\[0149\]](#)
- **SCARINGE et al.** NAR., 1990, vol. 18, 5433- [\[0149\]](#)
- **WINCOTT et al.** NAR., 1995, vol. 23, 2677-2684 [\[0149\]](#)
- **WINCOTT et al.** Methods Mol. Bio., 1997, vol. 74, 59- [\[0149\]](#)
- **MOORE et al.** Science, 1992, vol. 256, 9923- [\[0150\]](#)
- **SHABAROVA et al.** NAR, 1991, vol. 19, 4247- [\[0150\]](#)
- **BELLON et al.** Nucleosides & Nucleotides, 1997, vol. 16, 951- [\[0150\]](#)
- **BELLON et al.** Bioconjugate Chem., 1997, vol. 8, 204- [\[0150\]](#)
- **ELBASHIR et al.** Genes Dev., 2001, vol. 15, 2188-200 [\[0160\]](#)
- **BERNSTEIN et al.** Nature, 2001, vol. 409, 6818363-6 [\[0160\]](#)
- **LEE et al.** Nature, 2003, vol. 425, 6956415-9 [\[0160\]](#)
- **MCMANUSSHARP** Nature Rev Genet, 2002, vol. 3, 10737-47 [\[0160\]](#)
- **PADDISONHANNON** Curr Opin Mol Ther., 2003, vol. 5, 3217-24 [\[0160\]](#)
- **BERNSTEIN et al.** RNA, 2001, vol. 7, 111509-21 [\[0160\]](#)
- **NISHIKURA** Cell, 2001, vol. 107, 4415-8 [\[0160\]](#)
- **UI-TEI et al.** J Biomed Biotechnol., 2006, vol. 2006, 65052- [\[0161\]](#)
- **CHALK et al.** BBRC, 2004, vol. 319, 1264-74 [\[0161\]](#) [\[0167\]](#)
- **SIoudleIRDAL** Met. Mol Biol., 2004, vol. 252, 457-69 [\[0161\]](#) [\[0167\]](#)
- **LEVENKOVA et al.** Bioinform., 2004, vol. 20, 3430-2 [\[0161\]](#) [\[0167\]](#)
- **UI-TEI et al.** Nuc. Acid Res., 2004, vol. 32, 3936-48 [\[0161\]](#)
- **BRAASCH et al.** Biochem., 2003, vol. 42, 267967-75 [\[0161\]](#) [\[0168\]](#)
- **CHIU et al.** RNA, 2003, vol. 9, 91034-48 [\[0161\]](#) [\[0168\]](#)
- **GIL et al.** Apoptosis, 2000, vol. 5, 107-114 [\[0162\]](#)

- **ELBASHIR** et al. *Nature*, 2001, vol. 411, 494-498 [0162]
- **CAPLEN** et al. *PNAS USA*, 2001, vol. 98, 9742-9747 [0162]
- **FIRE** et al. *Nature*, 1998, vol. 391, 806- [0163]
- **AMBROS** *Nature*, 2004, vol. 431, 7006350-355 [0163]
- **BARTEL** *Cell*, 2004, vol. 116, 2281-97 [0163]
- *Nat. Med.*, 2005, vol. 11, 150-55 [0165]
- **BARIK** et al. *J. Mol. Med.*, 2005, vol. 83, 764-773 [0165]
- **DALLASVLA** *SM* *Med. Sci. Monitor*, 2006, vol. 12, 4RA67-74 [0165]
- **CHAKRABORTY** *Current Drug Targets*, 2007, vol. 8, 3469-82 [0165]
- **DYKXHOORN** et al. *Gene Therapy*, 2006, vol. 13, 541-552 [0165]
- **MUCKE** *Drugs*, 2007, vol. 10, 137-41 [0166]
- **UI-TEI** et al. *J Biomed Biotech.*, 2006, vol. 2006, 65052- [0167]
- **UI-TEI** et al. *NAR*, 2004, vol. 32, 3936-48 [0167]
- **DE PAULA** et al. *RNA*, 2007, vol. 13, 431-56 [0167]
- **BOUTLA** et al. *Curr. Biol.*, 2001, vol. 11, 1776-1780 [0169]
- **SCHWARZ** et al. *Mol. Cell*, 2002, vol. 10, 537-48 [0169]
- **AMARZGUIOUI** et al. *NAR*, 2003, vol. 31, 2589-95 [0169]
- **HOLEN** et al. *NAR*, 2003, vol. 31, 92401-07 [0169]
- **CHIURANARNA**, 2003, vol. 9, 1034-48 [0169]
- **CZAUDERNA** et al. *NAR*, 2003, vol. 31, 112705-16 [0169]
- **SHEN** et al. *FEBS Lett.*, 2003, vol. 539, 111-114 [0182]
- **XIA** et al. *Nat. Biotech.*, 2002, vol. 20, 1006-1010 [0182]
- **REICH** et al. *Mol. Vision*, 2003, vol. 9, 210-216 [0182]
- **SORENSEN** et al. *J. Mol. Biol.*, 2003, vol. 327, 761-766 [0182]
- **LEWIS** et al. *Nat. Gen.*, 2002, vol. 32, 107-108 [0182]
- **SIMEONI** et al. *NAR*, 2003, vol. 31, 112717-2724 [0182]
- **TOLENTINO** et al. *Retina*, 2004, vol. 24, 1132-138 [0182]
- **SONG** et al. *Antibody mediated in vivo delivery of small interfering RNAs via cell-surface receptors* *Nat Biotechnol.*, 2005, vol. 23, 6709-17 [0183]
- **SOUTSCHEK** et al. *Nature*, 2004, vol. 432, 173-177 [0183]
- **LORENZ** et al. *Bioorg. Med. Chem. Lett.*, 2004, vol. 14, 4975-4977 [0183]
- **TOLENTINO** et al. *Retina*, 2004, vol. 24, 132-138 [0189]
- **REICH** et al. *Molecular Vision*, 2003, vol. 9, 210-216 [0189]
- **WEISS** et al. *Human Gene Therapy*, 1999, vol. 10, 2287-2293 [0190]
- **DENSMORE** et al. *Molecular therapy*, 1999, vol. 1, 180-188 [0190]
- **GAUTAM** et al. *Molecular Therapy*, 2001, vol. 3, 551-556 [0190]
- **SHAHIWALAMISRAAAPS** *PharmSciTech*, 2004, vol. 24;6, 3E482-6 [0190]
- **W.M. PARDRIDGE** *Brain Drug Targeting: the future of brain drug development* Cambridge University Press 20010000 [0193]
- **SAMBROOK** et al. *Molecular cloning: A laboratory manual* Cold Springs Harbor Laboratory 19890000 [0221]
- **AUSUBEL** et al. *Current Protocols in Molecular Biology* John Wiley and Sons 19880000 [0231]
- **AUSUBEL** et al. *Current Protocols in Molecular Biology* John Wiley and Sons 19890000 [0231]
- **PERBAL**, *A Practical Guide to Molecular Cloning* John Wiley & Sons 19880000 [0231]
- **WATSON** et al. *Recombinant DNA* Scientific American Books [0231]
- *Genome Analysis: A Laboratory Manual Series* Cold Spring Harbor Laboratory Press 19980000 vol. 1-4, [0231]

- PCR Protocols: A Guide To Methods And Applications Academic Press 19900000 [0231]
- TESTONI et al. *Blood*, 1996, vol. 87, 3822- [0231]
- CASTANOTTO ROSSI *Nature*, 2009, vol. 22, 426-33 [0256]
- REID et al. *J. Surg. Res.*, 2004, vol. 116, 172-180 [0285]
- MUSTOE et al. *JCI*, 1991, vol. 87, 2694-703 [0285]
- AHN MUSTOE *Ann Pl. Surg.*, 1991, vol. 24, 117-23 [0285]
- STARCHER WILLIAMS *Lab. Animals*, 1989, vol. 23, 234-240 [0287]
- PENG et al. *Am J Respir Crit Care Med*, 2004, vol. 169, 1245-1251 [0287]
- JEYASEELAN et al. *Infect. Immunol*, 2004, vol. 72, 7247-56 [0287]
- HASEGAWA et al. *Exp Neurol*, 2005, vol. 193, 394-410 [0289]
- PEASE et al. *J. Glaucoma*, 2006, vol. 15, 6512-9 [0291]
- CHENG L et al. *J. Neurosci.*, 2002, vol. 22, 3977-3986 [0294]
- MIZOBUCHI et al. *J. Heart Lung Transplant*, 2004, vol. 23, 889-93 [0295]
- HUANG et al. *J. Heart Lung Transplant.*, 1995, vol. 14, S49- [0295]
- MATSUMURA et al. *Transplantation*, 1995, vol. 59, 1509-1517 [0295]
- WILKES et al. *Transplantation*, 1999, vol. 67, 890-896 [0295]
- NAKA et al. *Circulation Research*, 1996, vol. 79, 773-783 [0295]
- CHEN et al. *J Biomed Sci.*, 2003, vol. 10, 6588-92 [0296]
- TRENTHAM et al. *J. Exp. Med.*, 1977, vol. 146, 857-868 [0300]
- KONG et al. *Nature*, 1999, vol. 402, 304-308 [0300]
- HAN et al. *Nagoya J Med Sci*, 1999, vol. 62, 3-4115-26 [0300]

PATENTKRAV

1. Dobbeltstrenget nukleinsyremolekyle med nedenstående struktur (A):



5 hvori hver af N^2 , N og N' er et ikke-modificeret eller modificeret ribonukleotid, eller en ikke-konventionel del;

hvor i hver af $(N)x$ og $(N')y$ er et oligonukleotid, hvor i hver konsekutiv N eller N' er bundet til en tilstødende N eller N' med en kovalent binding;

hvor i hver af $x=y=18$;

10 hvori sekvensen af $(N)x$ har komplementaritet med en konsekutiv sekvens i et mål-RNA; og sekvensen af $N^2-(N')y$ er fuldt komplementær til sekvensen af $N^1-(N)x$;

hvor i N^1 er kovalent bundet til $(N)x$ og mismatchet med mål-RNA'et;

hvor i:

15 når parringsnukleotidet i mål-RNA'et er adenosin, uridin i position 1 af antisense-strengen er substitueret med en N^1 udvalgt fra adenosin eller deoxyadenosin; eller

når parringsnukleotidet i mål-RNA'et er cytidin, guanosin i position 1 af antisense-strengen er substitueret med en N^1 udvalgt fra adenosin, deoxyadenosin, eller deoxyuridin; eller

20 når parringsnukleotidet i mål-RNA'et er guanosin, cytidin i position 1 af antisense-strengen er substitueret med en N^1 udvalgt fra adenosin eller deoxyadenosin; eller

når parringsnukleotidet i mål-RNA'et er uridin, adenosin i position 1 af antisense-strengen er substitueret med et N^1 , der er deoxyuridin;

hvor i z'' kan være til stede eller fraværende, men hvis den er til stede, er den en endedel, der er kovalent bundet ved 5'-terminus af $N^2-(N')y$; og

25 hvori hver af Z og Z' er uafhængigt til stede eller fraværende, men hvis de er til stede, er de uafhængigt 1-5 konsekutive nukleotider, konsekutive ikke-nukleotiddele eller en kombination deraf kovalent bundet ved 3'-terminus af den streng, hvor i den er til stede.

30 2. Dobbeltstrenget nukleinsyremolekyle ifølge krav 1, hvor i N^1 er udvalgt fra adenosin eller deoxyadenosin, og hvor i parringsnukleotidet i mål-RNA'et er adenosin.

3. Dobbeltstrenget nukleinsyremolekyle ifølge krav 1, hvor i N^1 er udvalgt fra adenosin, deoxyadenosin, eller deoxyuridin, og hvor i parringsnukleotidet i mål-RNA'et er cytidin.

4. Dobbeltstrengen syremolekyle ifølge et hvilket som helst af krav 1 til 3, hvori N¹ og N² danner et basepar mellem adenosin og uridin eller mellem deoxyuridin og adenosin.

5 5. Dobbeltstrengen syremolekyle ifølge et hvilket som helst af krav 1 til 4, hvori z" er til stede.

6. Dobbeltstrengen nukleinsyremolekyle ifølge et hvilket som helst af krav 1 til 5, hvori Z og Z' er til stede.

10 7. Dobbeltstrengen nukleinsyremolekyle ifølge et hvilket som helst af krav 1 til 5, hvori én af Z eller Z' er til stede.

15 8. Dobbeltstrengen nukleinsyremolekyle ifølge et hvilket som helst af krav 1 til 7, hvori Z eller Z' omfatter en ikke-nukleotiddel.

9. Dobbeltstrengen nukleinsyremolekyle ifølge et hvilket som helst af krav 1 til 8, hvori mindst én af N eller N' omfatter et 2'O-methyl sukermodificeret ribonukleotid.

20 10. Dobbeltstrengen nukleinsyremolekyle ifølge krav 1, hvori x=y=18, Z' er fraværende, hvori Z er til stede og omfatter to alkyldele kovalent bundet til hinanden via en phosphodiesterbinding; hvori N² omfatter adenosin; og hvori N¹ omfatter uridin.

25 11. Dobbeltstrengen nukleinsyremolekyle ifølge krav 1, hvori Z' er til stede og omfatter to alkyldele kovalent bundet til hinanden via en phosphodiesterbinding.

12. Dobbeltstrengen nukleinsyremolekyle ifølge et hvilket som helst af krav 1 til 11 til anvendelse i terapi.

30 13. Farmaceutisk sammensætning, der omfatter mindst ét dobbeltstrengen nukleinsyremolekyle ifølge et hvilket som helst af krav 1 til 11; og en farmaceutisk acceptabel bærer.

14. Fremgangsmåde til generering af et dobbeltstrengen RNA-molekyle bestående af en

sense-streng og en antisense-streng, hvilken fremgangsmåde omfatter følgende trin

a) udvælgelse af en konsekutiv sekvens af 17 til 25 nukleotider i et mål-RNA og syntetisering af en antisense-streng, der omfatter komplementaritet med den konsekutive sekvens af 17 til 25 nukleotider af mål-mRNA'et, hvori antisense-strengens 5'-terminale

5 nukleotid,

hvis uridin, er substitueret med adenosin eller deoxyadenosin; eller

hvis guanosin, er substitueret med adenosin, deoxyadenosin eller deoxyuridin; eller

hvis cytidin, er substitueret med adenosin eller, deoxyadenosin; eller

hvis adenosin, er substitueret med et deoxyuridin;

10 b) syntetisering af en sense-streng med 17 til 25 nukleotider med fuld komplementaritet med antisense-strengen, hvori sense-strengens 3'-terminale nukleotid danner et Watson Crick-basepar med antisense-strengens 5'-terminale nukleotid; og

c) annealing af sense- og antisense-strengene; hvorved der genereres et dobbeltstrenget RNA-molekyle.

15

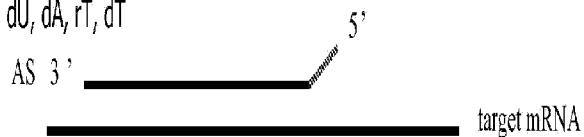
DRAWINGS

1. Guide strand (AS): 5' terminal nucleotide mismatch to target RNA

Target C. AS G > U, dU, rA, dA, rT, dT

Target G. AS C > U, dU, rA, dA, rT, dT

Target U. AS A > U, dU, dA, rT, dT



Passenger strand (SEN): full match to AS

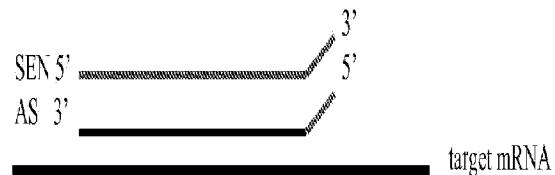
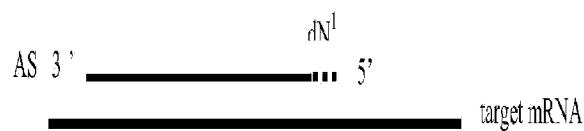


FIGURE 1

2. Guide strand (AS): 5' terminal deoxyribonucleotide (dN^1) complementary to target RNA



TLR2_16 siRNA (blunt ended with alternating 2'-OMe)

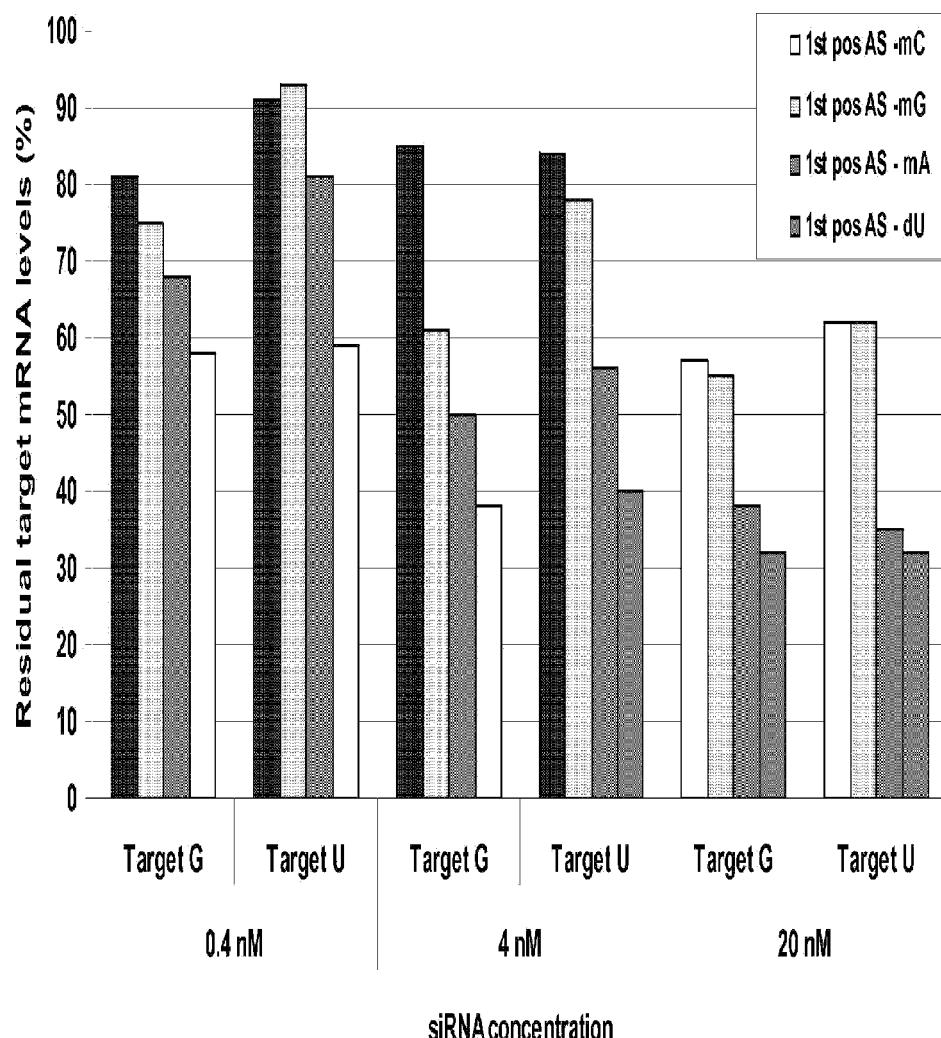
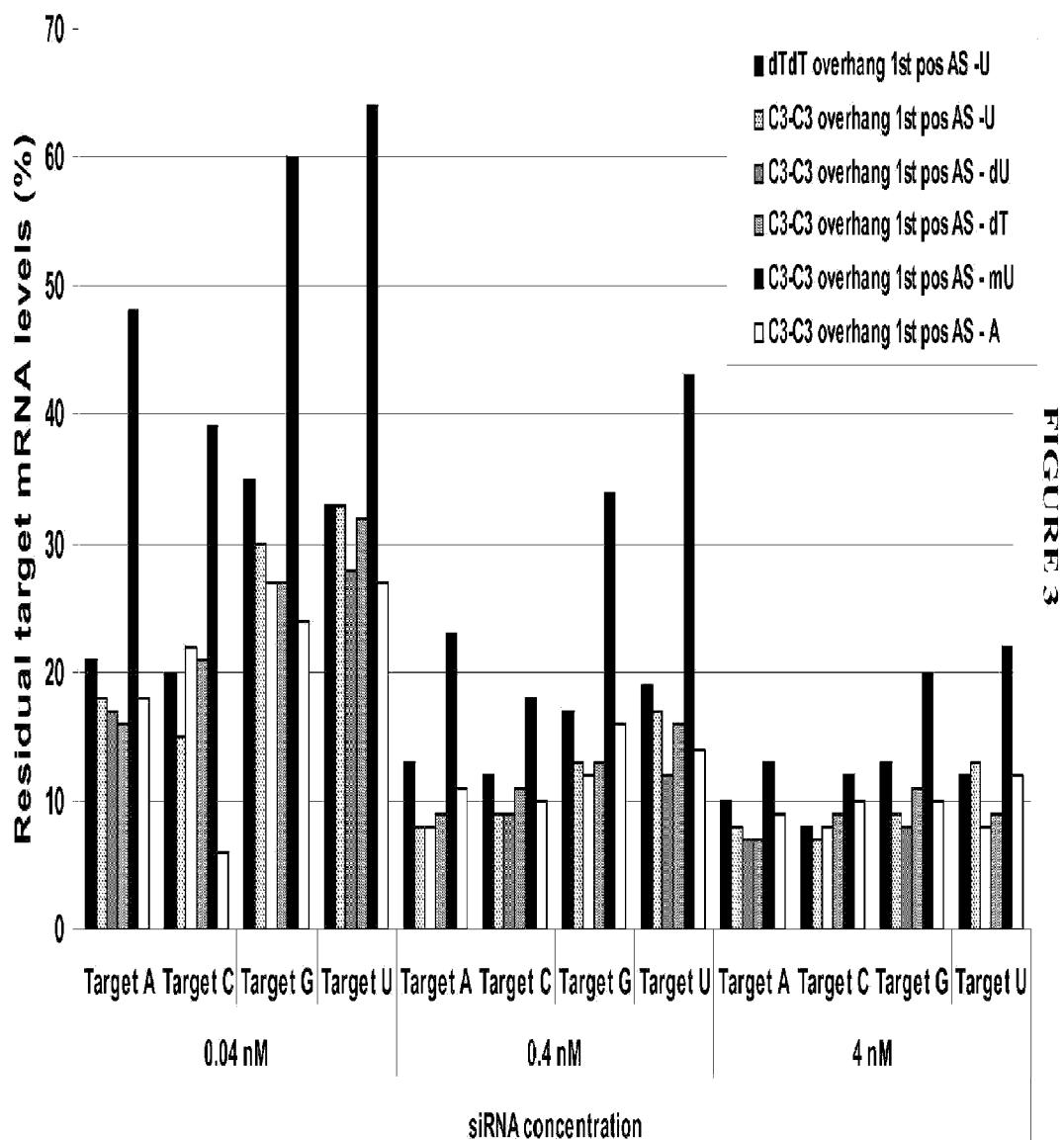
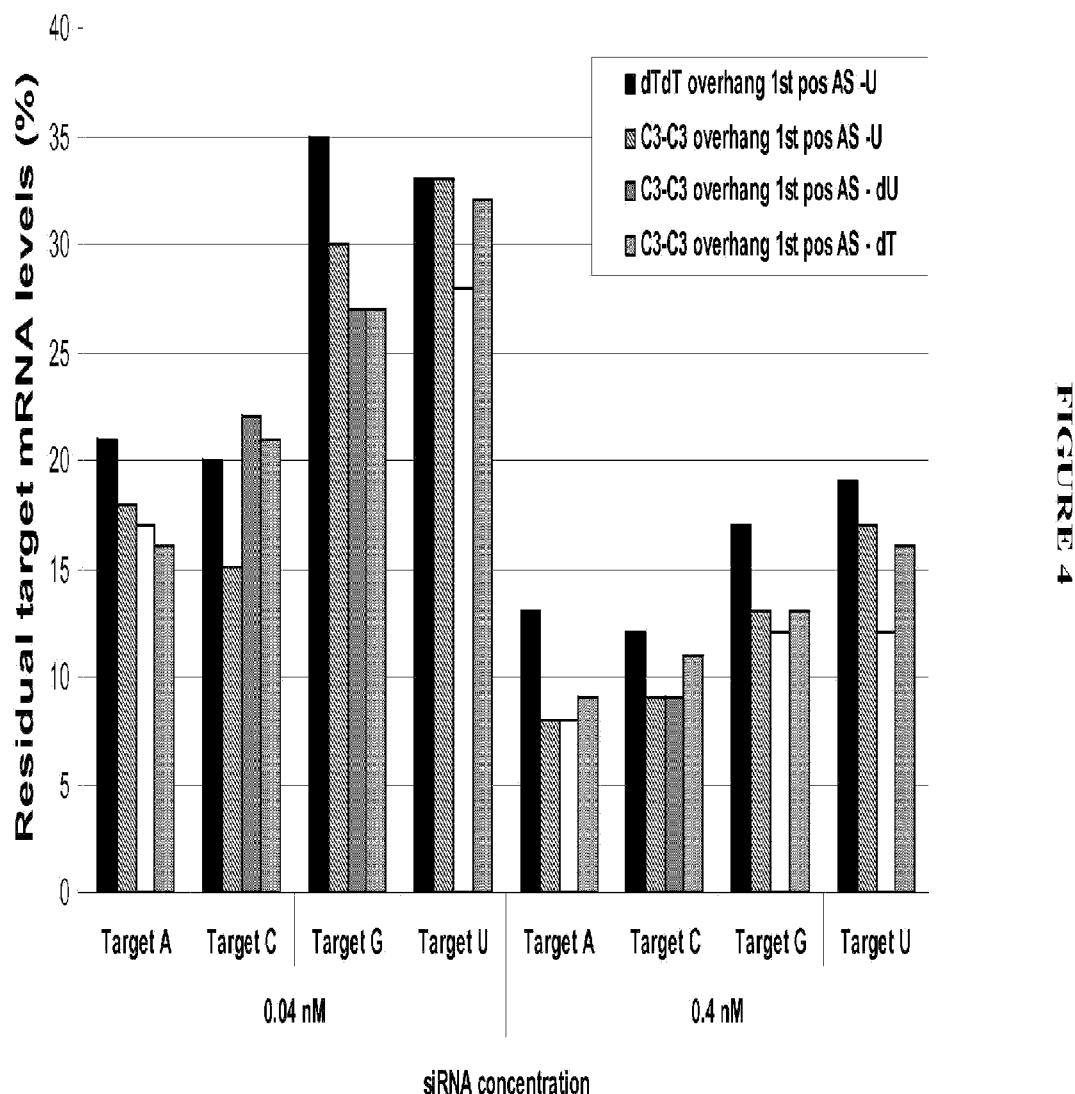


FIGURE 2

TLR2_37 siRNA (non-modified/C3-C3 or dTdT overhangs)



TLR2_37 siRNA (non-modified with C3-C3 overhangs or dTdT overhangs)



TLR2_37 siRNA (non-modified with dTdT overhangs) - 0.04nM

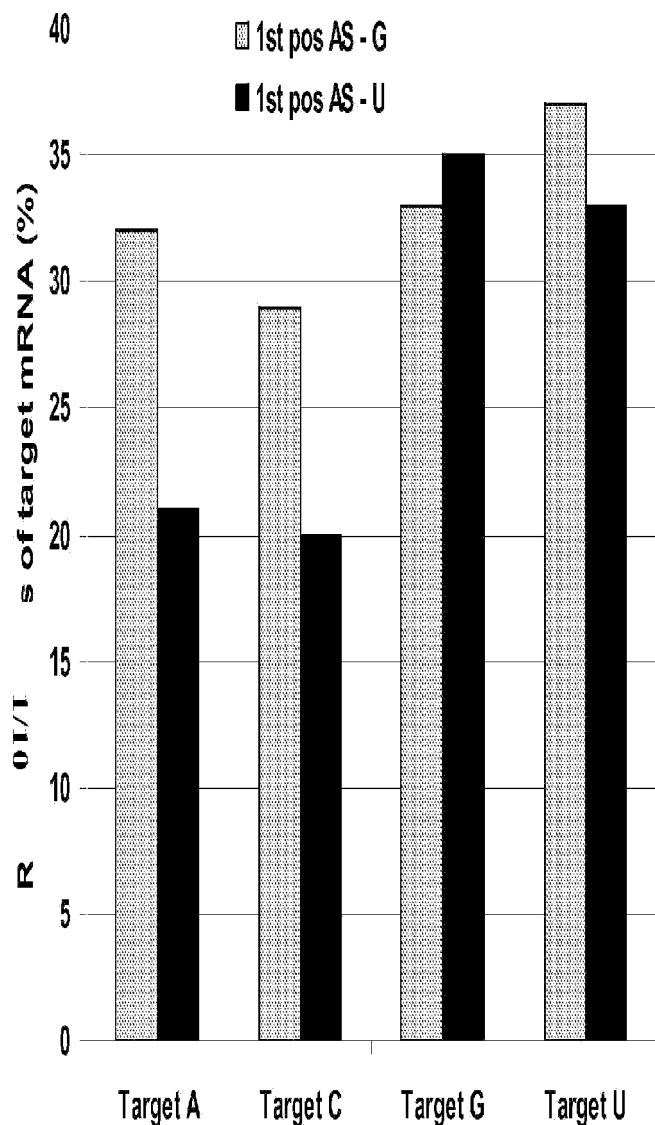
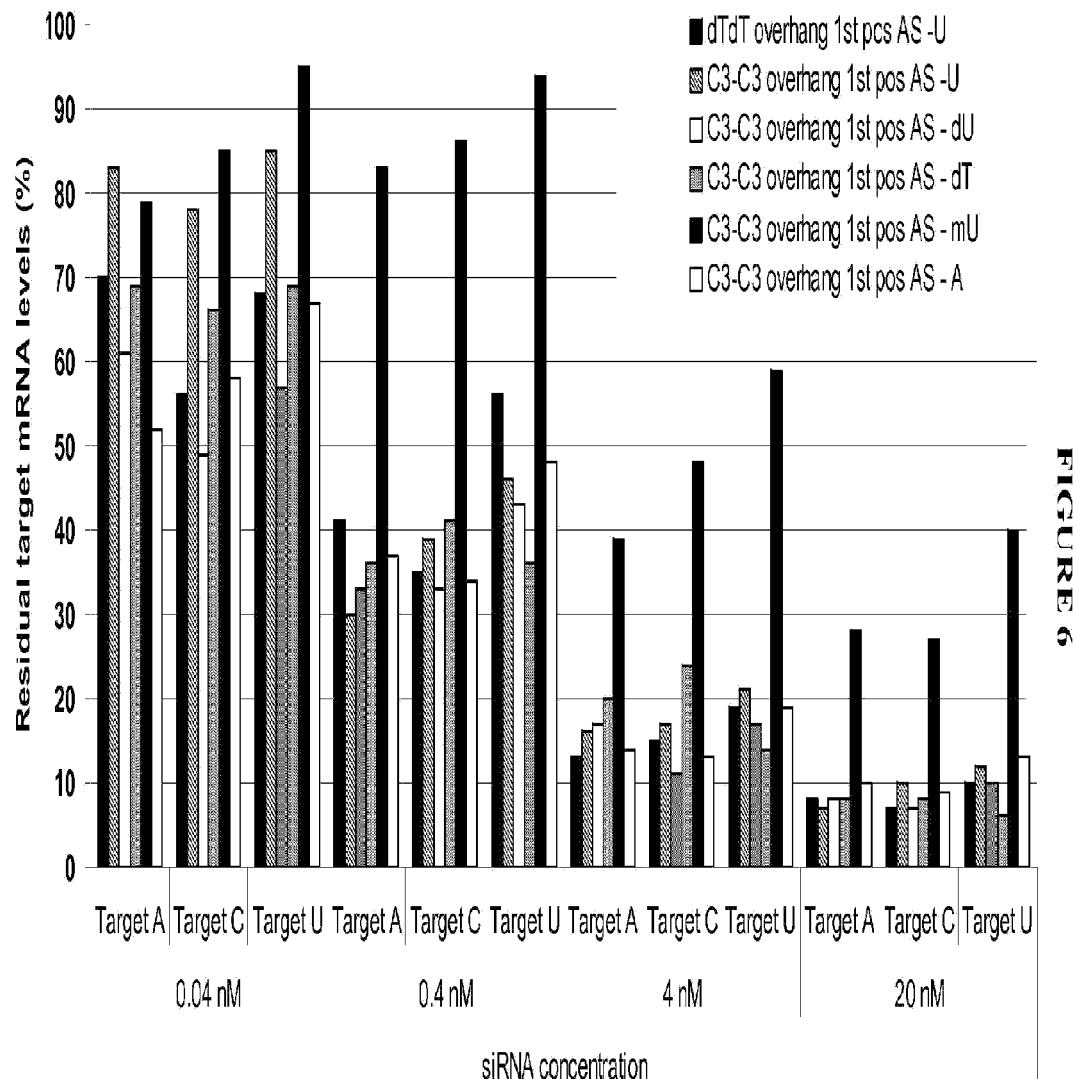


FIGURE 5

TLR2_46 siRNA (unmodified with 3' C3-C3 or dTdT overhangs)



TLR2_46 siRNA (unmodified with dTdT overhangs)

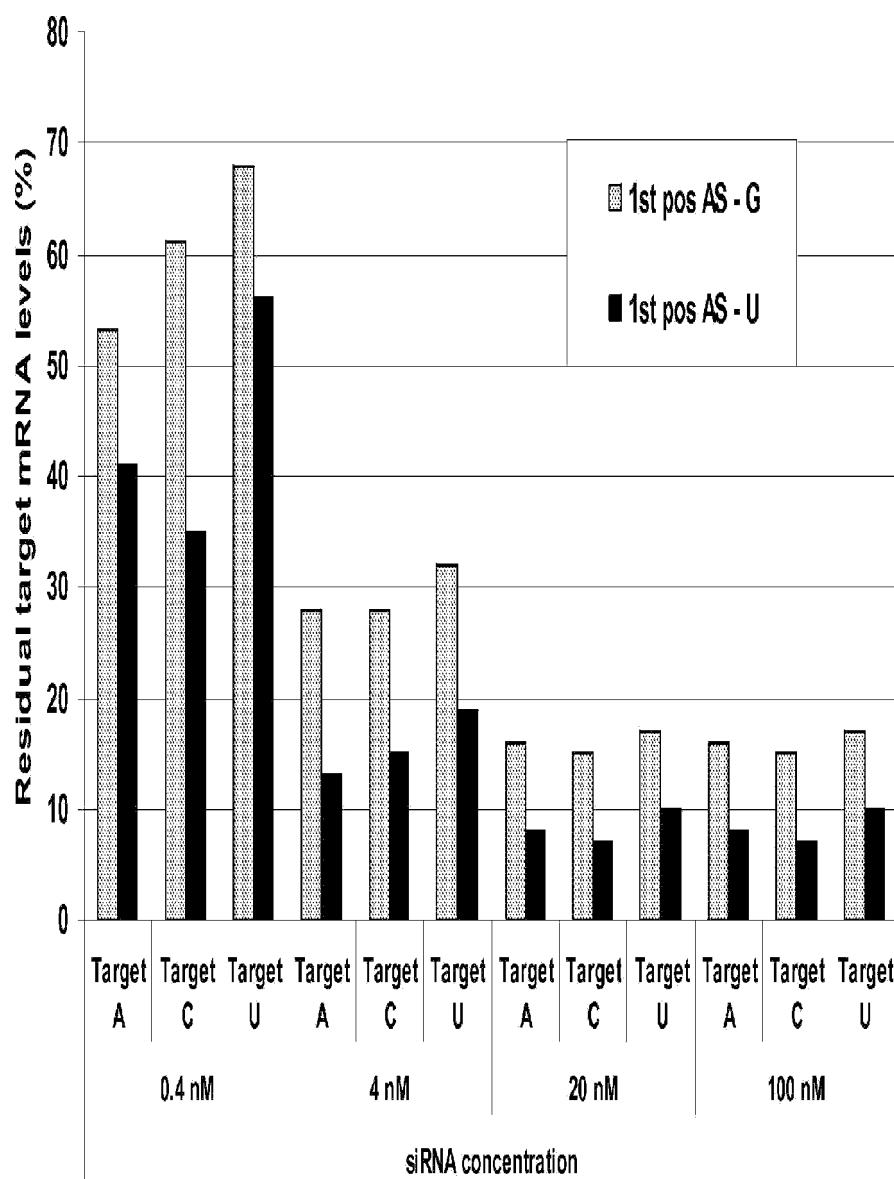
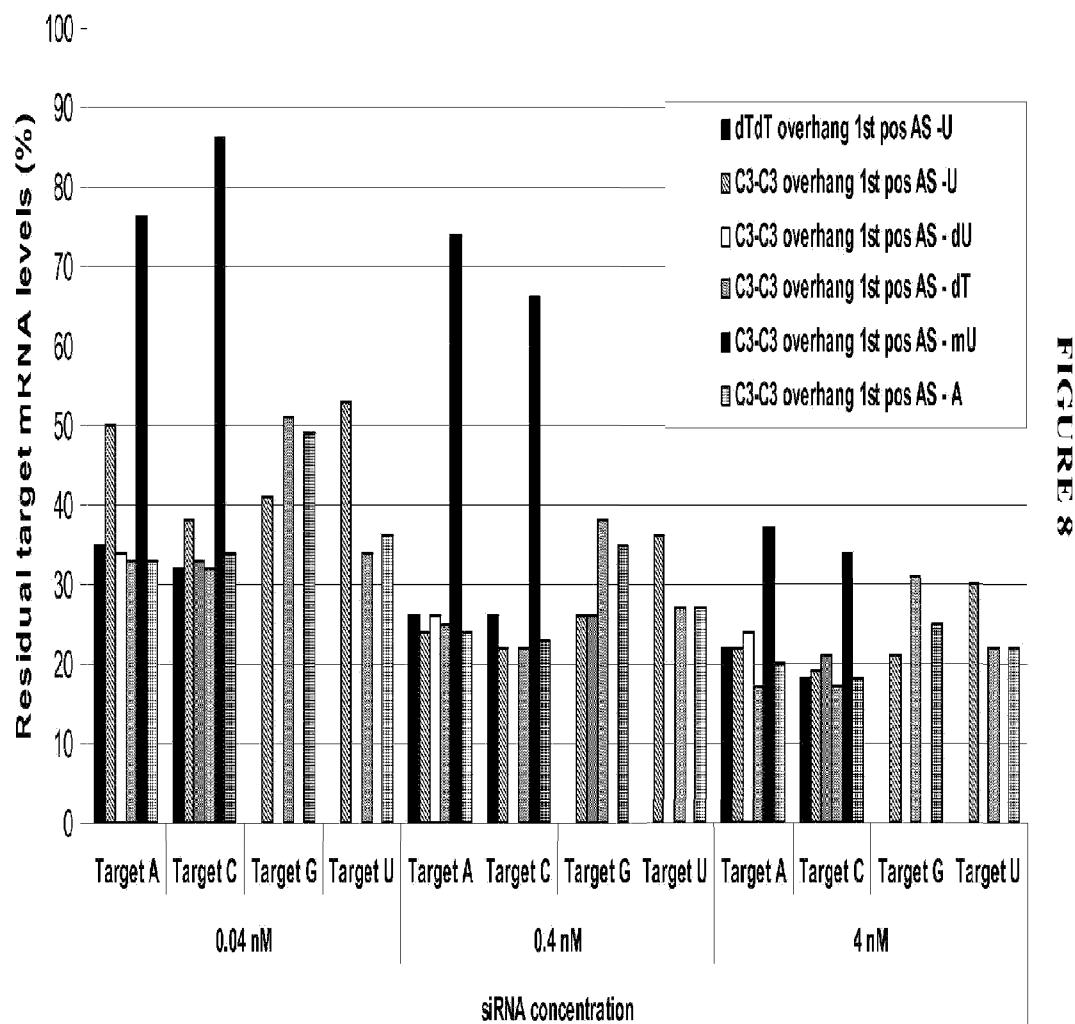


FIGURE 7

RHOA_61 siRNA (unmodified with C3-C3 or dTdT overhangs)



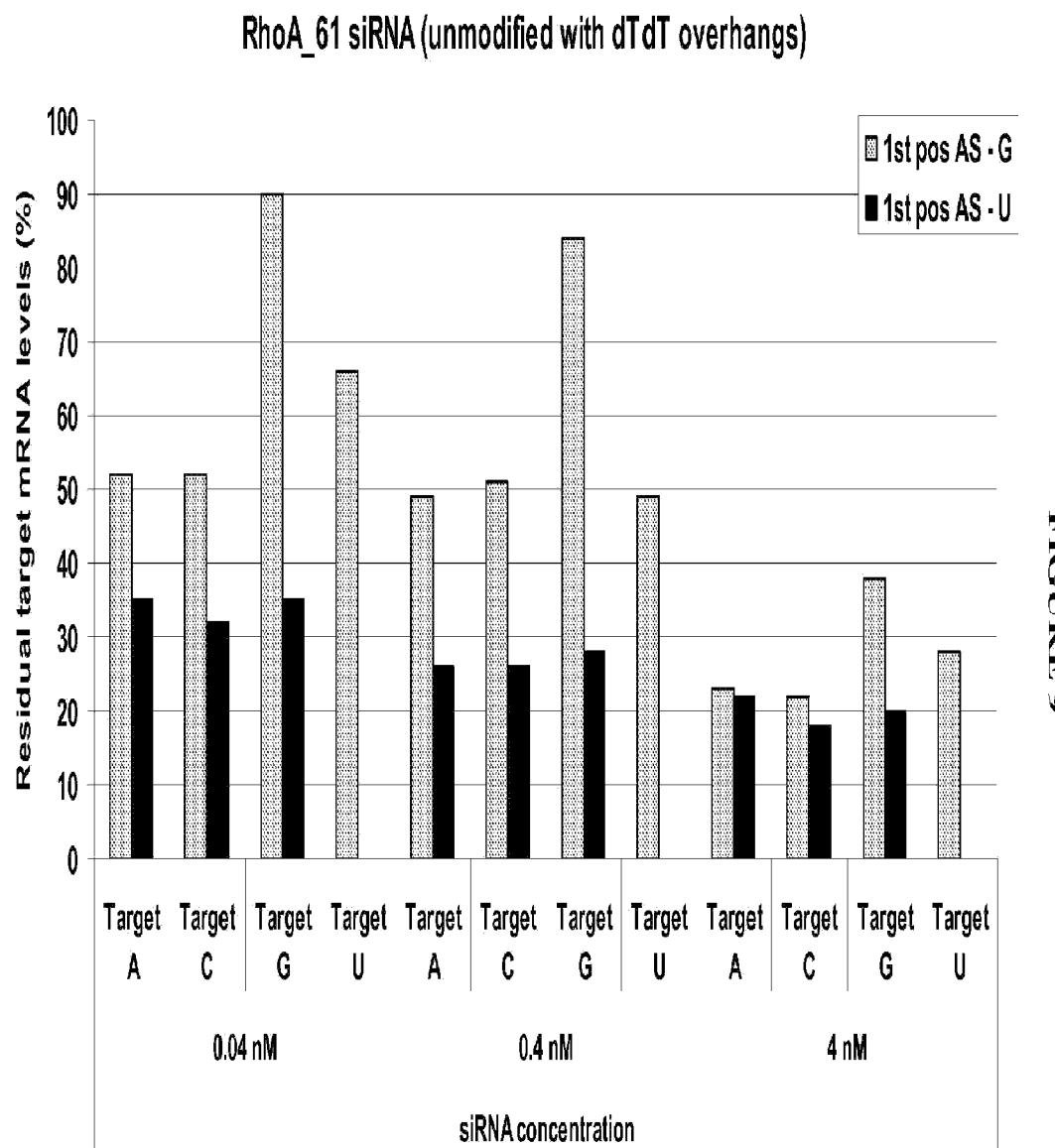


FIGURE 9

RhoA_61 siRNA (unmodified with dTdT overhangs)

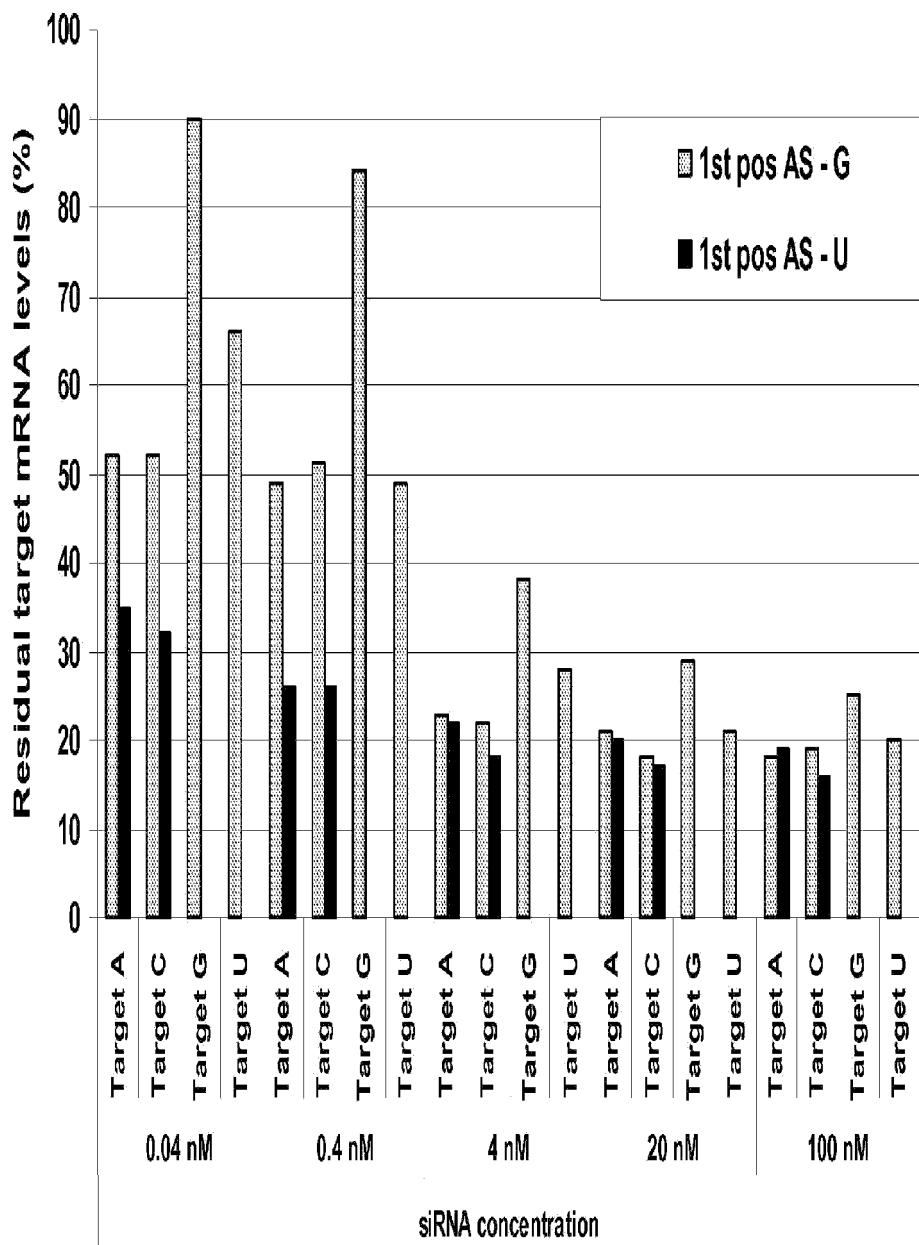


FIGURE 10