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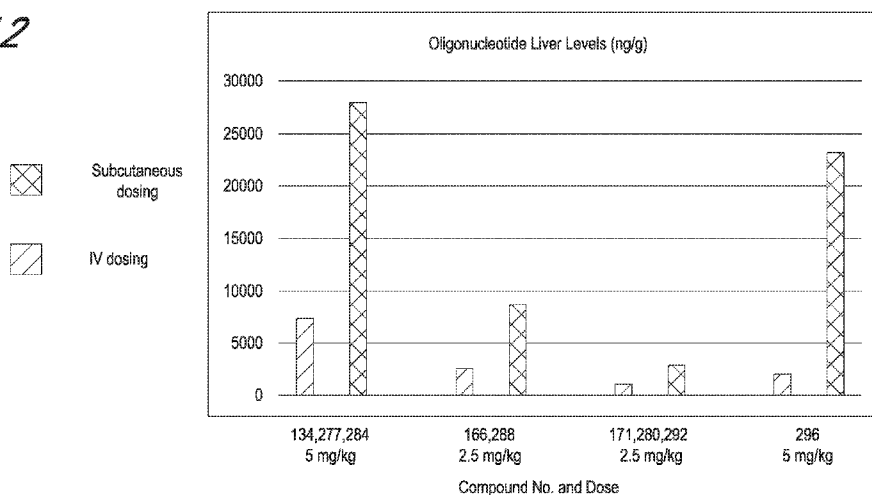
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(54) Title: S-ANTIGEN TRANSPORT INHIBITING OLIGONUCLEOTIDE POLYMERS AND METHODS

FIG. 12



Compound No. (dose)	134,277,284 5 mg/kg		166,288 2.5 mg/kg		171,280,292 2.5 mg/kg		296 5 mg/kg	
Route	IV	SC	IV	SC	IV	SC	IV	SC
Liver Levels (ng/g)	7350.5	27,942.5	2603	8711	1024.5	2938.5	1995.5	23,191.5

(57) Abstract: Various embodiments provide STOPSTTM polymers that are S-antigen transport inhibiting oligonucleotide polymers, processes for making them and methods of using them to treat diseases and conditions. In some embodiments the STOPSTTM modified oligonucleotides include an at least partially phosphorothioated sequence of alternating A and C units having modifications as described herein. The sequence independent antiviral activity against hepatitis B of embodiments of STOPSTTM modified oligonucleotides, as determined by HBsAg Secretion Assay, is greater than that of a reference compound.

HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

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S-ANTIGEN TRANSPORT INHIBITING OLIGONUCLEOTIDE POLYMERS AND METHODS

RELATED APPLICATION INFORMATION

[0001] This application claims priority to U.S. Serial No. 62/757,632, filed November 8, 2018; U.S. Serial No. 62/855,323, filed May 31, 2019; and to U.S. Serial No. 62/907,845, filed September 30, 2019. Each of the foregoing is incorporated herein by reference in its entirety.

BACKGROUND

Field

[0002] This application relates to STOPSTTM antiviral compounds that are S-antigen transport inhibiting oligonucleotide polymers, processes for making them and methods of using them to treat diseases and conditions.

Description

[0003] The STOPSTTM compounds described herein are antiviral oligonucleotides that can be at least partially phosphorothioated and exert their antiviral activity by a non-sequence dependent mode of action. See A. Vaillant, "Nucleic acid polymers: Broad spectrum antiviral activity, antiviral mechanisms and optimization for the treatment of hepatitis B and hepatitis D infection", Antiviral Research 133, 32-40 (2016). The term "Nucleic Acid Polymer" (NAP) has been used in the literature to refer to such oligonucleotides, although that term does not necessarily connote antiviral activity. A number of patent applications filed in the early 2000s disclosed the structures of certain specific compounds and identified various structural options as potential areas for future experimentation. See, e.g., U.S. Patent Nos. 7,358,068; 8,008,269; 8,008,270 and 8,067,385. These efforts resulted in the identification of the compound known to those skilled in the art as REP 2139, a phosphorothioated 40-mer having repeating adenosine-cytidine (AC) units with 5-methylation of all cytosines and 2'-O methyl modification of all riboses, along with the compound known as its clinical progenitor, REP 2055. See I. Roehl et al., "Nucleic Acid Polymers with Accelerated Plasma and Tissue Clearance for Chronic Hepatitis B Therapy", Molecular Therapy: Nucleic Acids Vol. 8, 1-12 (2017). The authors of that publication indicated that the structural features of these

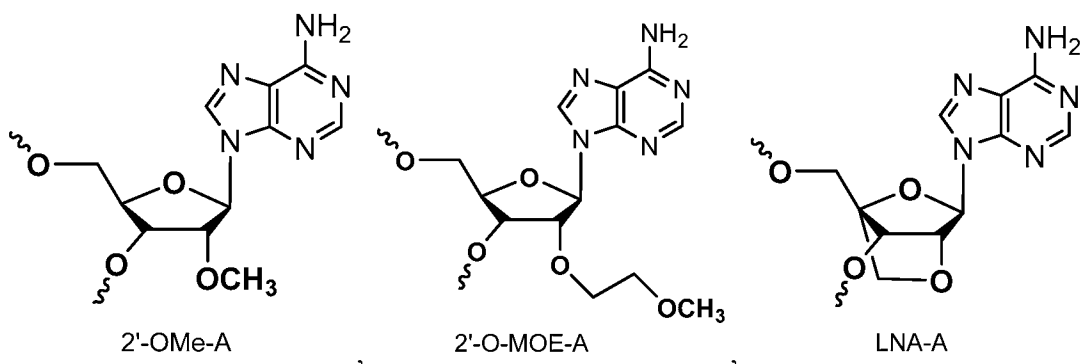
compounds had been optimized for the treatment of hepatitis B (HBV) and hepatitis D (HBD). See also A. Vaillant, "Nucleic acid polymers: Broad spectrum antiviral activity, antiviral mechanisms and optimization for the treatment of hepatitis B and hepatitis D infection", *Antiviral Research* 133 (2016) 32-40. According to these authors and related literature, such compounds preserve antiviral activity against HBV while preventing recognition by the innate immune response to allow their safe use with immunotherapies such as pegylated interferon. However, there remains a long-felt need for more effective compounds in this class.

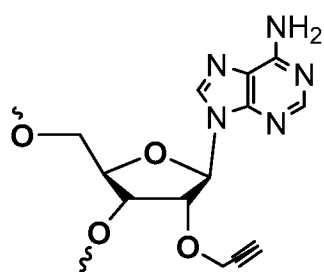
SUMMARY

[0004] It has now been discovered that, contrary to the teachings in the art regarding the optimum combination of desirable structural features for antiviral compounds, significantly improved properties can be obtained by modifying them to provide STOPSTTM compounds as described herein. For example, in some embodiments the sequence independent antiviral activity of the new STOPSTTM compounds against HBV, as determined by HBsAg Secretion Assay, is greater than that of a reference compound. In view of the many years of research culminating in the art-recognized optimized structure of REP 2139, there had been little expectation by those skilled in the art that embodiments of the modified STOPSTTM compounds described herein would be reasonably likely to display such improvements in potency. Thus, the structures of the new STOPSTTM compounds and methods of using them to treat HBV and HBD are surprising and unexpected.

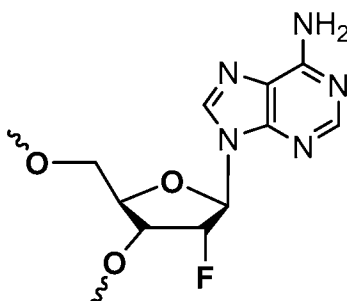
[0005] Some embodiments described herein relate to a modified oligonucleotide or complex thereof having sequence independent antiviral activity against hepatitis B, that can include an at least partially phosphorothioated sequence of alternating A and C units, wherein:

the A units comprise one or more selected from:

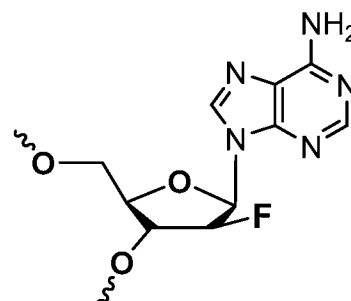




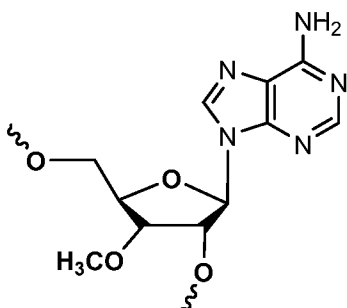
2'-O-Propargyl-A



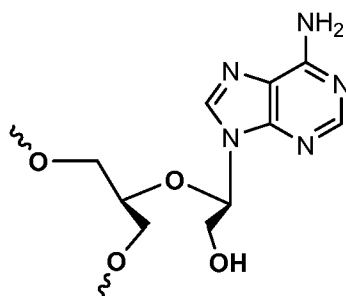
2'-F A



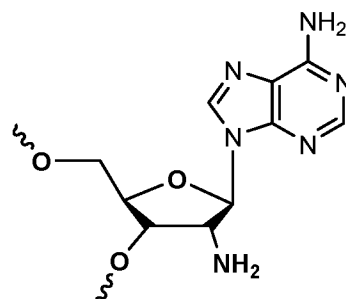
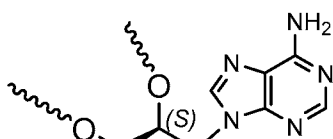
2'-araF A



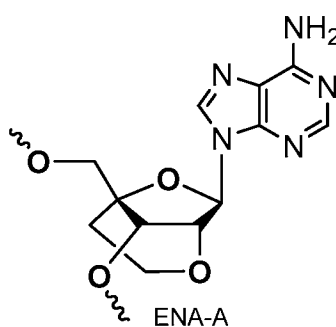
3'-OMe-A



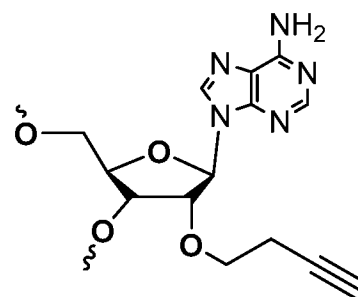
UNA-A

2'-NH₂-A

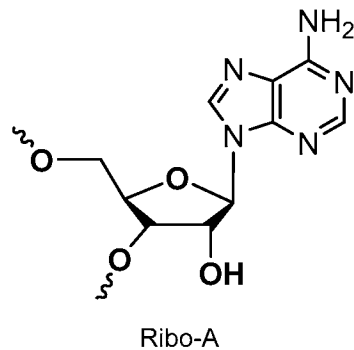
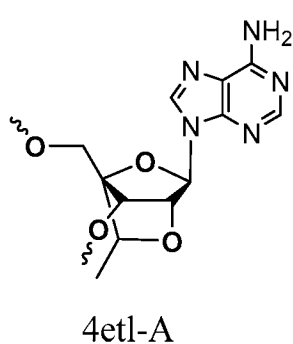
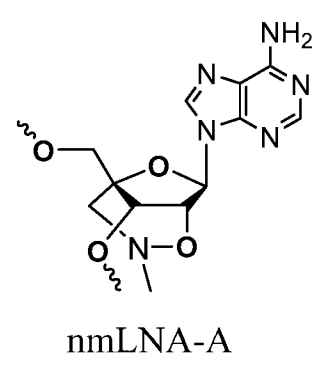
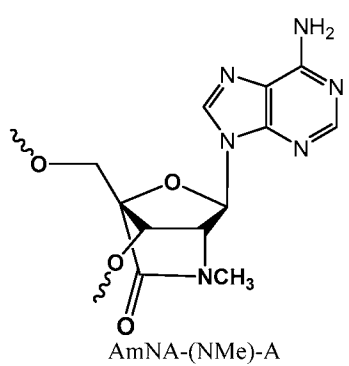
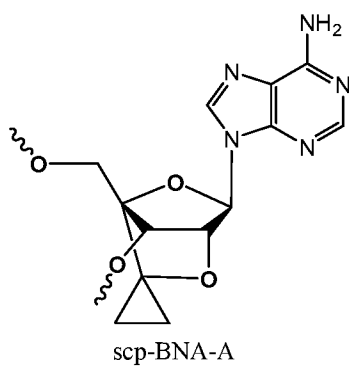
GNA-A



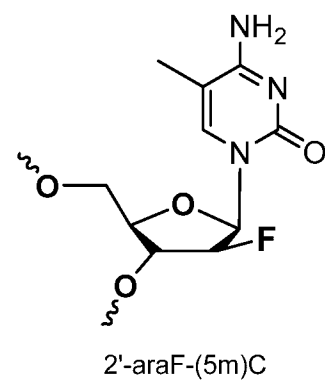
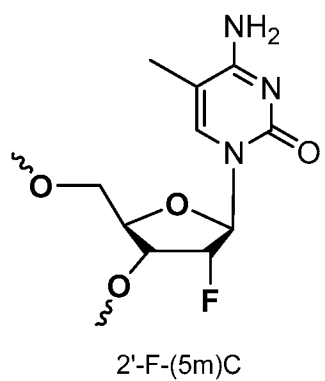
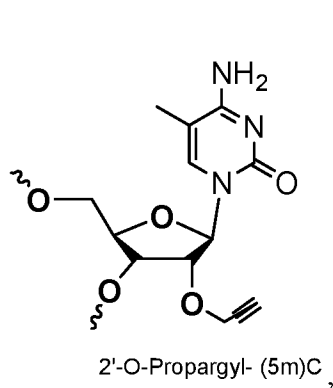
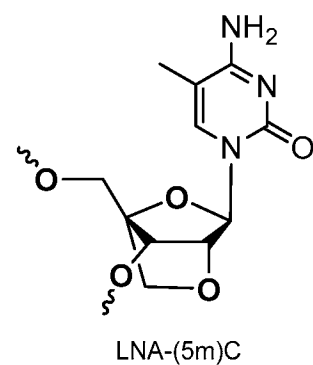
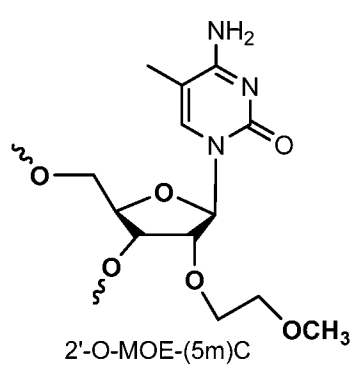
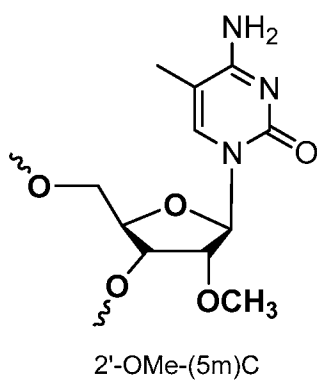
ENA-A

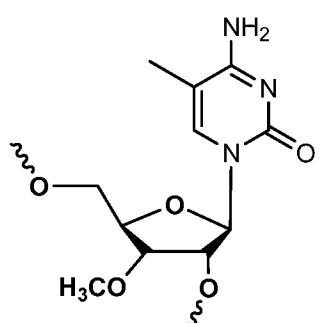


2'-O-Butynyl-A

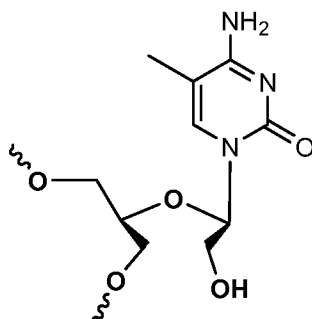


the C units comprise one or more selected from

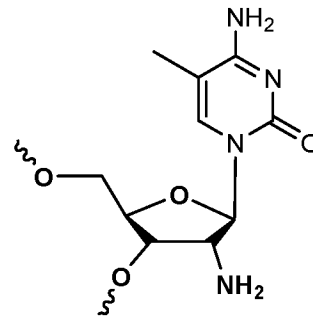




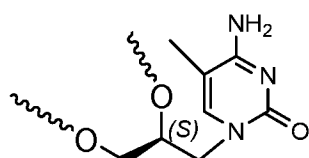
3'-OMe-(5m)C ,



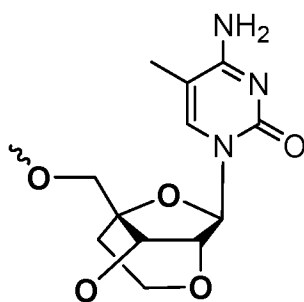
UNA-(5m)C ,



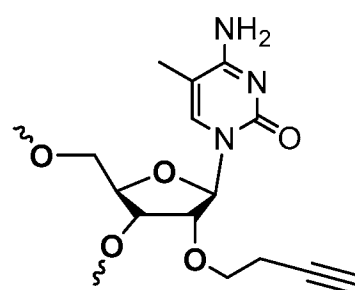
2'-NH2-(5m)C ,



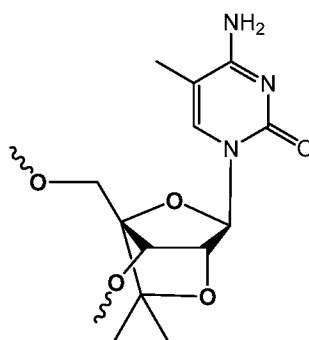
GNA-(5m)C ,



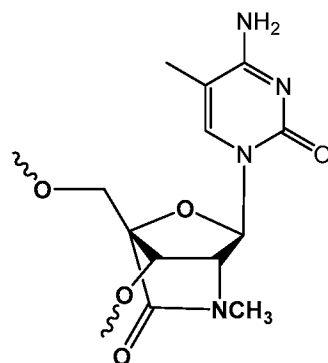
ENA-(5m)C ,



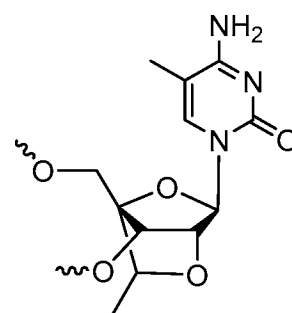
2'-O-Butynyl-(5m)C ,



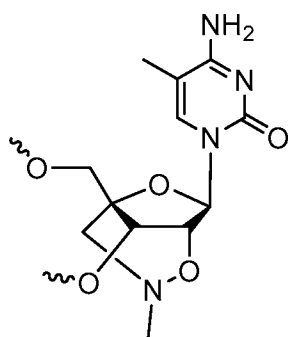
scp-BNA-(5m)C ,



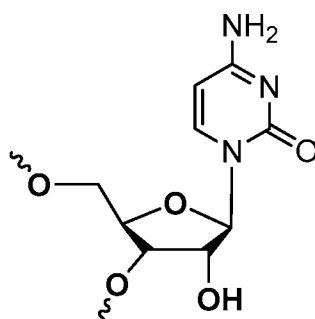
AmNA-(NMe)-(5m)C ,



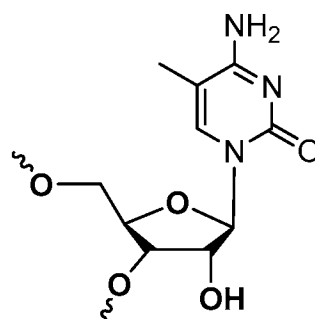
4etl-(5m)C ,



nmLNA-(5m)C ,



Ribo-C ,



Ribo-(5m)C ;

, and

[0006] each terminal $\sim\text{O}$ is independently hydroxyl, an O,O-dihydrogen phosphorothioate, a dihydrogen phosphate, an endcap or a linking group;

[0007] each internal $\sim\text{O}$ is a phosphorus-containing linkage to a neighboring A or C unit, the phosphorus-containing linkage being a phosphorothioate linkage or a modified linkage selected from phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate, 5'-phosphoramidate, 3',5'-phosphordiamidate, 5'-thiophosphoramidate, 3',5'-thiophosphordiamidate or diphosphodiester; and

[0008] the sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, is greater than that of a reference compound;

[0009] with the proviso that, when the sequence of alternating A and C units comprises a Ribo-A unit, the sequence further comprises at least one A unit that is not a Ribo-A unit; and

[0010] with the proviso that, when the sequence of alternating A and C units comprises a Ribo-C unit, the sequence further comprises at least one C unit that is not a Ribo-C unit.

[0011] Some embodiments described herein relate to a method of treating a HBV and/or HDV infection that can include administering to a subject identified as suffering from the HBV and/or HDV infection an effective amount of a modified oligonucleotide modified oligonucleotide as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide as described herein.

[0012] Some embodiments disclosed herein relate to a method of inhibiting replication of HBV and/or HDV that can include contacting a cell infected with the HBV and/or HDV with an effective amount of a modified oligonucleotide modified oligonucleotide as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide as described herein.

[0013] These other embodiments are described in greater detail below

BRIEF DESCRIPTION OF THE DRAWINGS

[0014] FIG. 1 illustrates an embodiment of a modified oligonucleotide that comprises a C₂₋₆alkylene linkage.

[0015] FIG. 2 illustrates an embodiment of a modified oligonucleotide that comprises a propylene oxide linkage.

[0016] FIG. 3A illustrates an embodiment of a modified oligonucleotide having cholesterol attached via a 5' tetraethylene glycol (TEG) linkage.

[0017] FIG. 3B illustrates an embodiment of a modified oligonucleotide having cholesterol attached via a 3' TEG linkage.

[0018] FIG. 3C illustrates an embodiment of a modified oligonucleotide having a tocopherol (Vitamin E) attached via a 5' TEG linkage.

[0019] FIG. 3D illustrates an embodiment of a modified oligonucleotide having a tocopherol (Vitamin E) attached via a 3' TEG linkage.

[0020] FIGS. 4A and 4B illustrate embodiments of modified oligonucleotides having GalNac attached via a linking group.

[0021] FIG. 5 illustrates an embodiment of a reaction scheme for preparing a 5'-EP building block.

[0022] FIG. 6A illustrates embodiments of modified oligonucleotides and corresponding values of sequence independent antiviral activity against hepatitis B (as determined by HBsAg Secretion Assay) and cytotoxicity.

[0023] FIG. 6B illustrates embodiments of modified oligonucleotides and corresponding values of sequence independent antiviral activity against hepatitis B (as determined by HBsAg Secretion Assay) and cytotoxicity.

[0024] FIG. 7 illustrates an embodiment of a reaction scheme for preparing compound 5'-VP.

[0025] FIG. 8 illustrates an embodiment of a reaction scheme for preparing compounds 8-5 and 8-6.

[0026] FIG. 9A illustrates an embodiment of a reaction scheme for preparing compound 9R.

[0027] FIG. 9B illustrates an embodiment of a reaction scheme for preparing compound 9S.

[0028] FIG. 10 illustrates an embodiment of a reaction scheme for preparing compounds 10-5 and 10-6.

[0029] FIG. 11A illustrates an embodiment of a reaction scheme for preparing compound **11R**.

[0030] FIG. 11B illustrates an embodiment of a reaction scheme for preparing compound **11S**.

[0031] FIG. 12 illustrates liver exposure results following subcutaneous administration to non-human primates of embodiments of modified oligonucleotide compounds.

[0032] FIG. 13 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0033] FIG. 14 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0034] FIG. 15 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0035] FIG. 16 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0036] FIG. 17 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0037] FIG. 18 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0038] FIG. 19 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0039] FIG. 20 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0040] FIG. 21 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0041] FIG. 22 illustrates PBMC assay results illustrating the immune reaction of embodiments of modified oligonucleotide compounds.

[0042] FIG. 23 illustrates a graph that is utilized in connection with the HBsAg Secretion Assays described in Examples B3 and B4.

DETAILED DESCRIPTION

Definitions

[0043] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as is commonly understood by one of ordinary skill in the art. All patents, applications, published applications and other publications referenced herein are incorporated by reference in their entirety unless stated otherwise. In the event that there are a plurality of definitions for a term herein, those in this section prevail unless stated otherwise.

[0044] The hepatitis B virus (HBV) is a DNA virus and a member of the *Hepadnaviridae* family. HBV infects more than 300 million worldwide and is a causative agent of liver cancer and liver disease such as chronic hepatitis, cirrhosis, and hepatocellular carcinoma. HBV can be acute and/or chronic. Acute HBV infection can be either asymptomatic or present with symptomatic acute hepatitis. HBV is classified into eight genotypes, A to H.

[0045] HBV is a partially double-stranded circular DNA of about 3.2 kilobase (kb) pairs. The HBV replication pathway has been studied in great detail. *T.J. Liang, Hepatology (2009) 49(5 Suppl):S13-S21*. One part of replication includes the formation of the covalently closed circular (cccDNA) form. The presence of the cccDNA gives rise to the risk of viral reemergence throughout the life of the host organism. HBV carriers can transmit the disease for many years. An estimated 257 million people are living with hepatitis B virus infection, and it is estimated that over 750,000 people worldwide die of hepatitis B each year. In addition, immunosuppressed individuals or individuals undergoing chemotherapy are especially at risk for reactivation of an HBV infection.

[0046] HBV can be transmitted by blood, semen, and/or another body fluid. This can occur through direct blood-to-blood contact, unprotected sex, sharing of needles, and from an infected mother to her baby during the delivery process. The HBV surface antigen (HBsAg) is most frequently used to screen for the presence of this infection. Currently available medications do not cure an HBV and/or HDV infection. Rather, the medications suppress replication of the virus.

[0047] The hepatitis D virus (HDV) is a DNA virus, also in the *Hepadnaviridae* family of viruses. HDV can propagate only in the presence of HBV. The routes of transmission of HDV are similar to those for HBV. Transmission of HDV can occur either via

simultaneous infection with HBV (coinfection) or in addition to chronic hepatitis B or hepatitis B carrier state (superinfection). Both superinfection and coinfection with HDV results in more severe complications compared to infection with HBV alone. These complications include a greater likelihood of experiencing liver failure in acute infections and a rapid progression to liver cirrhosis, with an increased risk of developing liver cancer in chronic infections. In combination with hepatitis B, hepatitis D has the highest fatality rate of all the hepatitis infections, at 20%. There is currently no cure or vaccine for hepatitis D.

[0048] As used herein in the context of oligonucleotides or other materials, the term “antiviral” has its usual meaning as understood by those skilled in the art and thus includes an effect of the presence of the oligonucleotides or other material that inhibits production of viral particles, typically by reducing the number of infectious viral particles formed in a system otherwise suitable for formation of infectious viral particles for at least one virus. In certain embodiments, the antiviral oligonucleotide has antiviral activity against multiple different virus, e.g., both HBV and HDV.

[0049] As used herein the term “oligonucleotide” (or “oligo”) has its usual meaning as understood by those skilled in the art and thus refers to a class of compounds that includes oligodeoxynucleotides, oligodeoxyribonucleotides and oligoribonucleotides. Thus, “oligonucleotide” refers to an oligomer or polymer of ribonucleic acid (RNA) or deoxyribonucleic acid (DNA) or mimetics thereof, including reference to oligonucleotides composed of naturally-occurring nucleobases, sugars and phosphodiester (PO) internucleoside (backbone) linkages as well as “modified” or substituted oligonucleotides having non-naturally-occurring portions which function similarly. Thus, the term “modified” (or “substituted”) oligonucleotide has its usual meaning as understood by those skilled in the art and includes oligonucleotides having one or more of various modifications, e.g., stabilizing modifications, and thus can include at least one modification in the internucleoside linkage and/or on the ribose, and/or on the base. For example, a modified oligonucleotide can include modifications at the 2'-position of the ribose, acyclic nucleotide analogs, methylation of the base, phosphorothioated (PS) linkages, phosphorodithioate linkages, methylphosphonate linkages, linkages that connect to the sugar ring via sulfur or nitrogen, and/or other modifications as described elsewhere herein. Thus, a modified oligonucleotide can include one or more phosphorothioated (PS) linkages, instead of or in addition to PO linkages. Like

unmodified oligonucleotides, modified oligonucleotides that include such PS linkages are considered to be in the same class of compounds because even though the PS linkage contains a phosphorous-sulfur double bond instead of the phosphorous-oxygen double bond of a PO linkage, both PS and PO linkages connect to the sugar rings through oxygen atoms.

[0050] As used herein in the context of modified oligonucleotides, the term “phosphorothioated” oligonucleotide has its usual meaning as understood by those skilled in the art and thus refers to a modified oligonucleotide in which all of the phosphodiester internucleoside linkages have been replaced by phosphorothioate linkages. Those skilled in the art thus understand that the term “phosphorothioated” oligonucleotide is synonymous with “fully phosphorothioated” oligonucleotide. A phosphorothioated oligonucleotide (or a sequence of phosphorothioated oligonucleotides within a partially phosphorothioated oligonucleotide) can be modified analogously, including (for example) by replacing one or more phosphorothioated internucleoside linkages by phosphodiester linkages. Thus, the term “modified phosphorothioated” oligonucleotide refers to a phosphorothioated oligonucleotide that has been modified in the manner analogous to that described herein with respect to oligonucleotides, e.g., by replacing a phosphorothioated linkage with a modified linkage such as phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate, 5'-phosphoramidate, 3',5'-phosphordiamidate, 5'-thiophosphoramidate, 3',5'-thiophosphordiamidate or diphosphodiester. An at least partially phosphorothioated sequence of a modified oligonucleotide can be modified similarly, and thus, for example, can be modified to contain a non-phosphorothioated linkage such as phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate 5'-phosphoramidate, 3',5'-phosphordiamidate, 5'-thiophosphoramidate, 3',5'-thiophosphordiamidate or diphosphodiester. In the context of describing modifications to a phosphorothioated oligonucleotide, or to an at least partially phosphorothioated sequence of a modified oligonucleotide, modification by inclusion of a phosphodiester linkage may be considered to result in a modified phosphorothioated oligonucleotide, or to a modified phosphorothioated sequence, respectively. Analogously, in the context of describing modifications to an oligonucleotide, or to an at least partially phosphodiesterified sequence of a modified oligonucleotide, the inclusion of a phosphorothioated linkage may be considered to result in a modified oligonucleotide or a modified phosphodiesterified sequence, respectively.

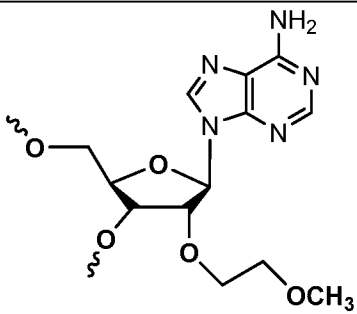
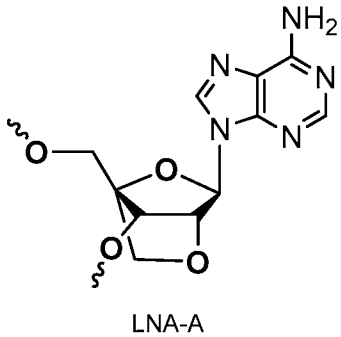
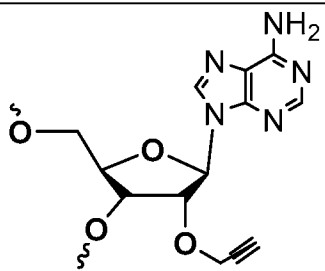
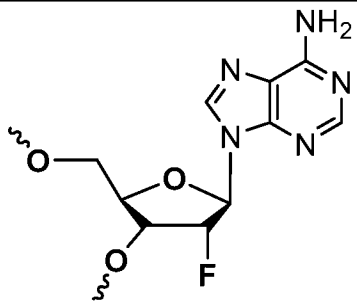
[0051] As used herein in the context of dinucleotides or oligonucleotides, the term “stereochemically defined phosphorothioate linkage” has its usual meaning as understood by those skilled in the art and thus refers to a phosphorothioate linkage having a phosphorus stereocenter with a selected chirality (R or S configuration). A composition containing such a dinucleotide or oligonucleotide can be enriched in molecules having the selected chirality. The stereopurity of such a composition can vary over a broad range, e.g. from about 51% to about 100% stereopure. In various embodiments, the stereopurity is greater than 55%, 65%, 75%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99%; or in a range defined as having any two of the foregoing stereopurity values as endpoints.

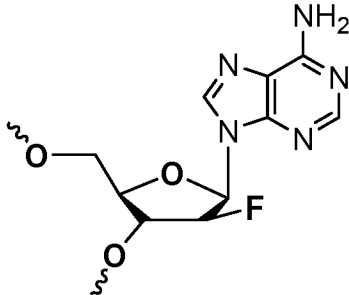
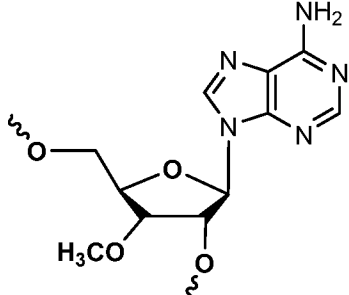
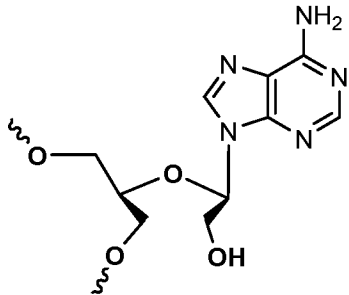
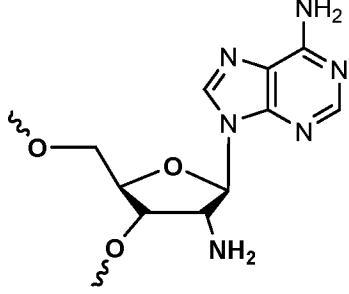
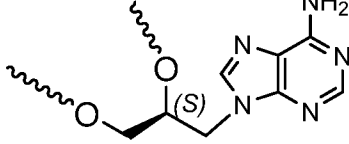
[0052] The term “sequence independent” antiviral activity has its usual meaning as understood by those skilled in the art and thus refers to an antiviral activity of an oligonucleotide (e.g., a modified oligonucleotide) that is independent of the sequence of the oligonucleotide. Methods for determining whether the antiviral activity of an oligonucleotide is sequence independent are known to those skilled in the art and include the tests for determining if an oligonucleotide acts predominantly by a non-sequence complementary mode of action as disclosed in Example 10 of U.S. Patent Nos. 7,358,068; 8,008,269; 8,008,270 and 8,067,385, which is hereby incorporated herein by reference and particularly for the purpose of describing such tests.

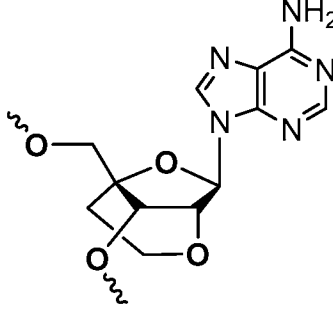
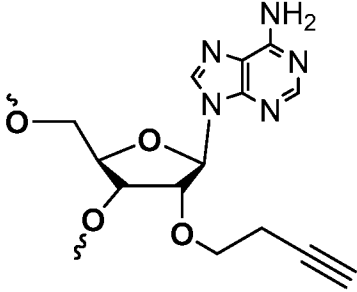
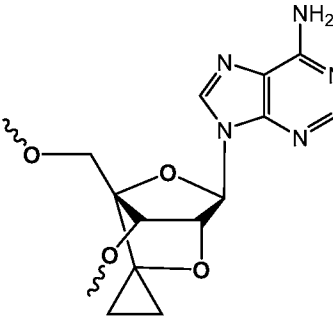
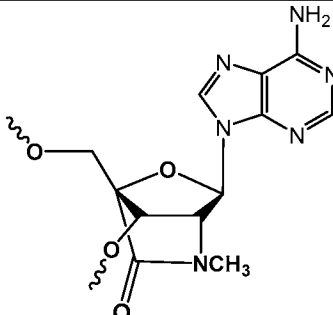
[0053] In the context of describing modified oligonucleotides having sequence independent antiviral activity and comprising a sequence (e.g., an at least partially phosphorothioated sequence) of A and C units (e.g., alternating A and C units, or AC units), the terms “A” and “C” refer to the modified adenosine-containing (A) units and modified cytosine-containing (C) units set forth in Tables 1 and 2 below, respectively.

TABLE 1 – “A” UNITS

Abbreviation (A Unit)	Structure (A Unit)
2'-OMe-A	

Abbreviation (A Unit)	Structure (A Unit)
2'-O-MOE-A	
LNA-A	 LNA-A
2'-O-Propargyl-A	
2'-F-A	

Abbreviation (A Unit)	Structure (A Unit)
2'-araF-A	
3'-OMe-A	
UNA-A	
2'-NH ₂ -A	
GNA-A	

Abbreviation (A Unit)	Structure (A Unit)
ENA-A	
2'-O-Butynyl-A	
scp-BNA-A	
AmNA(NMe)-A	

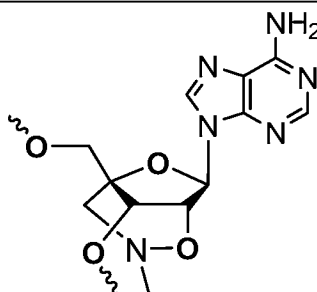
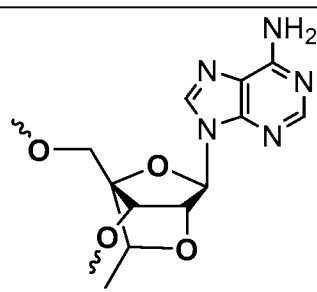
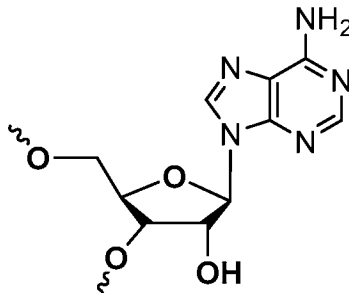
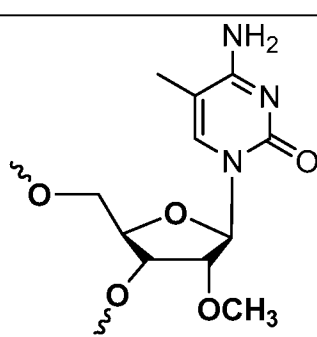
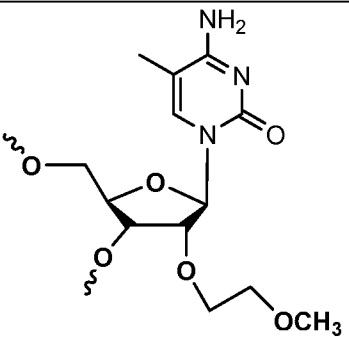
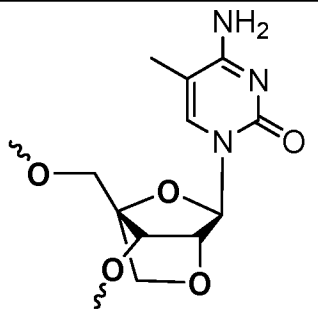
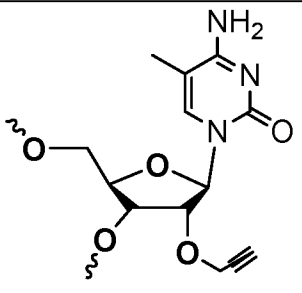
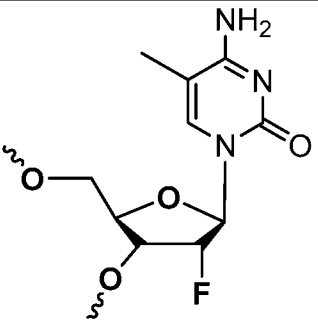
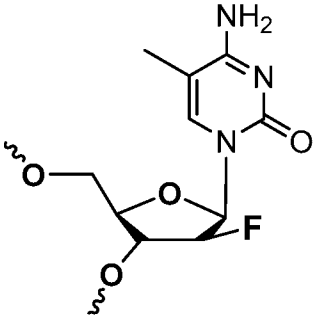
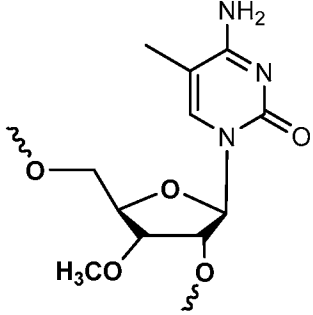
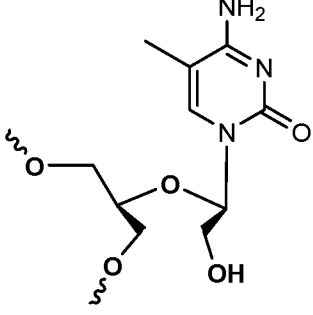
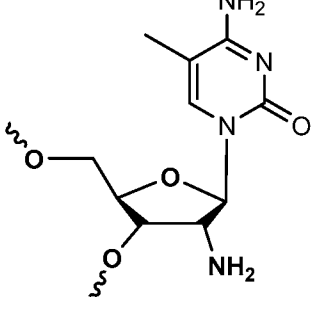
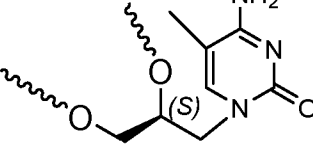
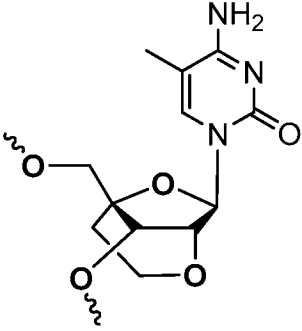
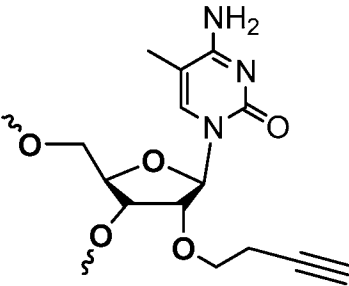
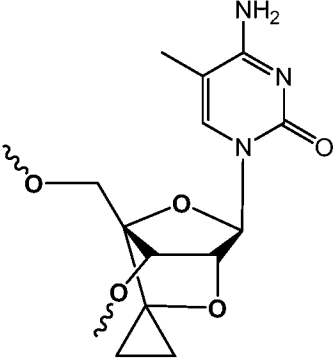
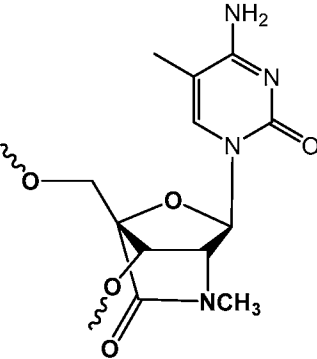
Abbreviation (A Unit)	Structure (A Unit)
nmLNA-A	
4etl-A	
Ribo-A	

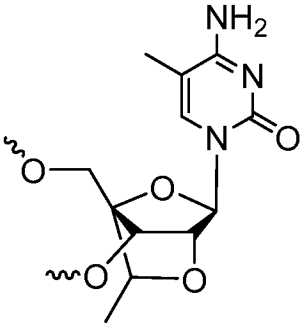
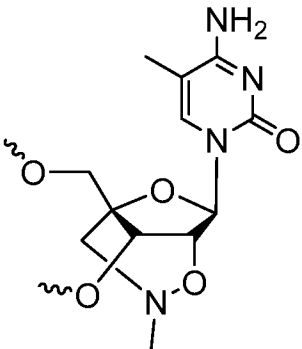
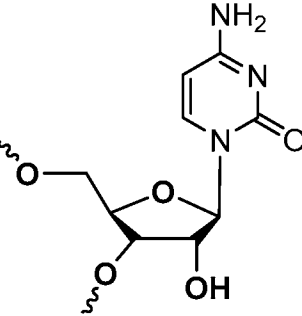
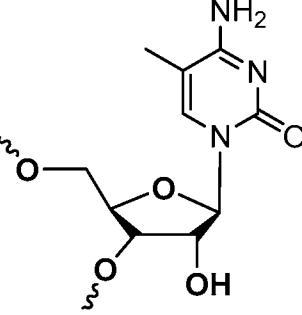
TABLE 2 – “C” UNITS

Abbreviation (C Unit)	Structure (C Unit)
2'-OMe-(5m)C	

Abbreviation (C Unit)	Structure (C Unit)
2'-O-MOE-(5m)C	
LNA-(5m)C	 LNA-5mC
2'-O-Propargyl-(5m)C	
2'-F-(5m)C	

Abbreviation (C Unit)	Structure (C Unit)
2'-araF-(5m)C	
3'-OMe-(5m)C	
UNA-(5m)C	
2'-NH ₂ -(5m)C	
GNA-(5m)C	

Abbreviation (C Unit)	Structure (C Unit)
ENA-(5m)C	
2'-O-Butynyl-(5m)C	
scp-BNA-(5m)C	
AmNA-(NMe)-(5m)C	

Abbreviation (C Unit)	Structure (C Unit)
4etl-(5m)C	
nmLNA-(5m)C	
Ribo-C	
Ribo-(5m)C	

Modified Oligonucleotide Compounds

[0054] An embodiment provides a STOPS™ modified oligonucleotide compound having sequence independent antiviral activity against hepatitis B, comprising an at least

partially phosphorothioated sequence of alternating A and C units, wherein the A units are any one or more selected from those set forth in Table 1 and the C units are any one or more selected from those set forth in Table 2. Various combinations of A and C units can be included in the at least partially phosphorothioated AC sequence, including the combinations described in Table 3 below.

TABLE 3 – EXAMPLES OF AC UNITS

No.	A Unit	C Unit
1	2'-OMe-A	2'-OMe-(5m)C
2	2'-OMe-A	2'-O-MOE-(5m)C
3	2'-OMe-A	LNA-(5m)C
4	2'-OMe-A	ENA-(5m)C
5	2'-OMe-A	scp-BNA-(5m)C
6	2'-OMe-A	AmNA-(NMe)-(5m)C
7	2'-OMe-A	2'-O-Propargyl-(5m)C
8	2'-OMe-A	2'-O-Butynyl-(5m)C
9	2'-OMe-A	2'-F-(5m)C
10	2'-OMe-A	2'-araF-(5m)C
11	2'-OMe-A	3'-OMe-(5m)C
12	2'-OMe-A	UNA-(5m)C
13	2'-OMe-A	2'-NH ₂ -(5m)C
14	2'-OMe-A	GNA-(5m)C
15	2'-OMe-A	4etl-(5m)C
16	2'-OMe-A	nmLNA-(5m)C
17	2'-O-MOE-A	2'-OMe-(5m)C
18	2'-O-MOE-A	2'-O-MOE-(5m)C
19	2'-O-MOE-A	LNA-(5m)C
20	2'-O-MOE-A	ENA-(5m)C

No.	A Unit	C Unit
21	2'-O-MOE-A	scp-BNA-(5m)C
22	2'-O-MOE-A	AmNA-(NMe)-(5m)C
23	2'-O-MOE-A	2'-O-Propargyl-(5m)C
24	2'-O-MOE-A	2'-O-Butynyl-(5m)C
25	2'-O-MOE-A	2'-F-(5m)C
26	2'-O-MOE-A	2'-araF-(5m)C
27	2'-O-MOE-A	3'-OMe-(5m)C
28	2'-O-MOE-A	UNA-(5m)C
29	2'-O-MOE-A	2'-NH ₂ -(5m)C
30	2'- O-MOE-A	GNA-(5m)C
31	2'-O-MOE-A	4etl-(5m)C
32	2'-O-MOE-A	nmLNA-(5m)C
33	LNA-A	2'-OMe-(5m)C
34	LNA-A	2'-O-MOE-(5m)C
35	LNA-A	LNA-(5m)C
36	LNA-A	ENA-(5m)C
37	LNA-A	scp-BNA-(5m)C
38	LNA-A	AmNA-(NMe)-(5m)C
39	LNA-A	2'-O-Propargyl-(5m)C
40	LNA-A	2'-O-Butynyl-(5m)C
41	LNA-A	2'-F-(5m)C
42	LNA-A	2'-araF-(5m)C
43	LNA-A	3'-OMe-(5m)C
44	LNA-A	UNA-(5m)C
45	LNA-A	2'-NH ₂ -(5m)C
46	LNA-A	GNA-(5m)C

No.	A Unit	C Unit
47	LNA-A	4etl-(5m)C
48	LNA-A	nmLNA-(5m)C
49	ENA-A	2'-OMe-(5m)C
50	ENA-A	2'- O-MOE-(5m)C
51	ENA-A	LNA-(5m)C
52	ENA-A	ENA-(5m)C
53	ENA-A	scp-BNA-(5m)C
54	ENA-A	AmNA-(NMe)-(5m)C
55	ENA-A	2'-O-Propargyl-(5m)C
56	ENA-A	2'-O-Butynyl-(5m)C
57	ENA-A	2'-F-(5m)C
58	ENA-A	2'-araF-(5m)C
59	ENA-A	3'-OMe-(5m)C
60	ENA-A	UNA-(5m)C
61	ENA-A	2'-NH ₂ -(5m)C
62	ENA-A	GNA-(5m)C
63	ENA-A	4etl-(5m)C
64	ENA-A	nmLNA-(5m)C
65	scp-BNA-A	2'-OMe-(5m)C
66	scp-BNA-A	2'- O-MOE-(5m)C
67	scp-BNA-A	LNA-(5m)C
68	scp-BNA-A	ENA-(5m)C
69	scp-BNA-A	scp-BNA-(5m)C
70	scp-BNA-A	AmNA-(NMe)-(5m)C
71	scp-BNA-A	2'-O-Propargyl-(5m)C
72	scp-BNA-A	2'-O-Butynyl-(5m)C

No.	A Unit	C Unit
73	scp-BNA-A	2'-F-(5m)C
74	scp-BNA-A	2'-araF-(5m)C
75	scp-BNA-A	3'-OMe-(5m)C
76	scp-BNA-A	UNA-(5m)C
77	scp-BNA-A	2'-NH ₂ -(5m)C
78	scp-BNA-A	GNA-(5m)C
79	scp-BNA-A	4etl-(5m)C
80	scp-BNA-A	nmLNA-(5m)C
81	AmNA(N-Me)-A	2'-OMe-(5m)C
82	AmNA(N-Me)-A	2'- O-MOE-(5m)C
83	AmNA(N-Me)-A	LNA-(5m)C
84	AmNA(N-Me)-A	ENA-(5m)C
85	AmNA(N-Me)-A	scp-BNA-(5m)C
86	AmNA(N-Me)-A	AmNA-(NMe)-(5m)C
87	AmNA(N-Me)-A	2'-O-Propargyl-(5m)C
88	AmNA(N-Me)-A	2'-O-Butynyl-(5m)C
89	AmNA(N-Me)-A	2'-F-(5m)C
90	AmNA(N-Me)-A	2'-ara-F-(5m)C
91	AmNA(N-Me)-A	3'-OMe-(5m)C
92	AmNA(N-Me)-A	UNA-(5m)C
93	AmNA(N-Me)-A	2'-NH ₂ -(5m)C
94	AmNA(N-Me)-A	GNA-(5m)C
95	AmNA(N-Me)-A	4etl-(5m)C
96	AmNA(N-Me)-A	nmLNA-(5m)C
97	2'-O-Propargyl-A	2'-OMe-(5m)C
98	2'-O-Propargyl-A	2'- O-MOE-(5m)C

No.	A Unit	C Unit
99	2'-O-Propargyl-A	LNA-(5m)C
100	2'-O-Propargyl-A	ENA-(5m)C
101	2'-O-Propargyl-A	scp-BNA-(5m)C
102	2'-O-Propargyl-A	AmNA-(NMe)-(5m)C
103	2'-O-Propargyl-A	2'-O-Propargyl-(5m)C
104	2'-O-Propargyl-A	2'-O-Butyne-(5m)C
105	2'-O-Propargyl-A	2'-F-(5m)C
106	2'-O-Propargyl-A	2'-araF-(5m)C
107	2'-O-Propargyl-A	3'-OMe-(5m)C
108	2'-O-Propargyl-A	UNA-(5m)C
109	2'-O-Propargyl-A	2'-NH ₂ -(5m)C
110	2'-O-Propargyl-A	GNA-(5m)C
111	2'-O-Propargyl-A	4etl-(5m)C
112	2'-O-Propargyl-A	nmLNA-(5m)C
113	2'-O-Butynyl-A	2'-OMe-(5m)C
114	2'-O-Butynyl-A	2'- O-MOE-(5m)C
115	2'-O-Butynyl-A	LNA-(5m)C
116	2'-O-Butynyl-A	ENA-(5m)C
117	2'-O-Butynyl-A	scp-BNA-(5m)C
118	2'-O-Butynyl-A	AmNA-(NMe)-(5m)C
119	2'-O-Butynyl-A	2'-O-Propargyl-(5m)C
120	2'-O-Butynyl-A	2'-O-Butynyl-(5m)C
121	2'-O-Butynyl-A	2'-F-(5m)C
122	2'-O-Butynyl-A	2'-araF-(5m)C
123	2'-O-Butynyl-A	3'-OMe-(5m)C
124	2'-O-Butynyl-A	UNA-(5m)C

No.	A Unit	C Unit
125	2'-O-Butynyl-A	2'-NH ₂ -(5m)C
126	2'-O-Butynyl-A	GNA-(5m)C
127	2'-O-Butynyl-A	4etl-(5m)C
128	2'-O-Butynyl-A	nmLNA-(5m)C
129	2'-F A	2'-OMe-(5m)C
130	2'-F A	2'- O-MOE-(5m)C
131	2'-F A	LNA-(5m)C
132	2'-F A	ENA-(5m)C
133	2'-F A	scp-BNA-(5m)C
134	2'-F A	AmNA-(NMe)-(5m)C
135	2'-F A	2'-O-Propargyl-(5m)C
136	2'-F A	2'-O-Butynyl-(5m)C
137	2'-F A	2'-F-(5m)C
138	2'-F A	2'-ara-F-(5m)C
139	2'-F A	3'-OMe-(5m)C
140	2'-F A	UNA-(5m)C
141	2'-F A	2'-NH ₂ -(5m)C
142	2'-F A	GNA-(5m)C
143	2'-F A	4etl-(5m)C
144	2'-F A	nmLNA-(5m)C
145	2'-araF A	2'-OMe-(5m)C
146	2'-araF A	2'- O-MOE-(5m)C
147	2'-araF A	LNA-(5m)C
148	2'-araF A	ENA-(5m)C
149	2'-araF A	scp-BNA-(5m)C
150	2'-araF A	AmNA-(NMe)-(5m)C

No.	A Unit	C Unit
151	2'-araF A	2'-O-Propargyl-(5m)C
152	2'-araF A	2'-O-Butynyl-(5m)C
153	2'-araF A	2'-F-(5m)C
154	2'-araF A	2'-araF-(5m)C
155	2'-araF A	3'-OMe-(5m)C
159	2'-araF A	UNA-(5m)C
157	2'-araF A	2'-NH ₂ -(5m)C
158	2'-araF A	GNA-(5m)C
159	2'-araF A	4etl-(5m)C
160	2'-araF A	nmLNA-(5m)C
161	3'-OMe-A	2'-OMe-(5m)C
162	3'-OMe-A	2'- O-MOE-(5m)C
163	3'-OMe-A	LNA-(5m)C
164	3'-OMe-A	ENA-(5m)C
165	3'-OMe-A	scp-BNA-(5m)C
166	3'-OMe-A	AmNA-(NMe)-(5m)C
167	3'-OMe-A	2'-O-Propargyl-(5m)C
168	3'-OMe-A	2'-O-Butynyl-(5m)C
169	3'-OMe-A	2'-F-(5m)C
170	3'-OMe-A	2'-ara-F-(5m)C
171	3'-OMe-A	3'-OMe-(5m)C
172	3'-OMe-A	UNA-(5m)C
173	3'-OMe-A	2'-NH ₂ -(5m)C
174	3'-OMe-A	GNA-(5m)C
175	3'-OMe-A	4etl-(5m)C
176	3'-OMe-A	nmLNA-(5m)C

No.	A Unit	C Unit
177	UNA-A	2'-OMe-(5m)C
178	UNA-A	2'- O-MOE-(5m)C
179	UNA-A	LNA-(5m)C
180	UNA-A	ENA-(5m)C
181	UNA-A	scp-BNA-(5m)C
182	UNA-A	AmNA-(NMe)-(5m)C
183	UNA-A	2'-O-Propargyl-(5m)C
184	UNA-A	2'-O-Butynyl-(5m)C
185	UNA-A	2'-F-(5m)C
186	UNA-A	2'-araF-(5m)C
187	UNA-A	3'-OMe-(5m)C
188	UNA-A	UNA-(5m)C
189	UNA-A	2'-NH ₂ -(5m)C
190	UNA-A	GNA-(5m)C
191	UNA-A	4etl-(5m)C
192	UNA-A	nmLNA-(5m)C
193	2'-NH ₂ -A	2'-OMe-(5m)C
194	2'-NH ₂ -A	2'-O-MOE-(5m)C
195	2'-NH ₂ -A	LNA-(5m)C
196	2'-NH ₂ -A	ENA-(5m)C
197	2'-NH ₂ -A	scp-BNA-(5m)C
198	2'-NH ₂ -A	AmNA-(NMe)-(5m)C
199	2'-NH ₂ -A	2'-O-Propargyl-(5m)C
200	2'-NH ₂ -A	2'-O-Butynyl-(5m)C
201	2'-NH ₂ -A	2'-F-(5m)C
202	2'-NH ₂ -A	2'-ara-F-(5m)C

No.	A Unit	C Unit
203	2'-NH ₂ -A	3'-OMe-(5m)C
204	2'-NH ₂ -A	UNA-(5m)C
205	2'-NH ₂ -A	2'-NH ₂ -(5m)C
206	2'-NH ₂ -A	GNA-(5m)C
207	2'-NH ₂ -A	4etl-(5m)C
208	2'-NH ₂ -A	nmLNA-(5m)C
209	GNA-A	2'-OMe-(5m)C
210	GNA-A	2'-O-MOE-(5m)C
211	GNA-A	LNA-(5m)C
212	GNA-A	ENA-(5m)C
213	GNA-A	scp-BNA-(5m)C
214	GNA-A	AmNA-(NMe)-(5m)C
215	GNA-A	2'-O-Propargyl-(5m)C
216	GNA-A	2'-O-Butynyl-(5m)C
217	GNA-A	2'-F-(5m)C
218	GNA-A	2'-ara-F-(5m)C
219	GNA-A	3'-OMe-(5m)C
220	GNA-A	UNA-(5m)C
221	GNA-A	2'-NH ₂ -(5m)C
222	GNA-A	GNA-(5m)C
223	GNA-A	4etl-(5m)C
224	GNA-A	nmLNA-(5m)C
225	nmLNA-A	2'-OMe-(5m)C
226	nmLNA-A	2'-O-MOE-(5m)C
227	nmLNA-A	LNA-(5m)C
228	nmLNA-A	ENA-(5m)C

No.	A Unit	C Unit
229	nmLNA-A	scp-BNA-(5m)C
230	nmLNA-A	AmNA-(NMe)-(5m)C
231	nmLNA-A	2'-O-Propargyl-(5m)C
232	nmLNA-A	2'-O-Butynyl-(5m)C
233	nmLNA-A	2'-F-(5m)C
234	nmLNA-A	2'-ara-F-(5m)C
235	nmLNA-A	3'-OMe-(5m)C
236	nmLNA-A	UNA-(5m)C
237	nmLNA -A	2'-NH ₂ -(5m)C
238	nmLNA-A	GNA-(5m)C
239	nmLNA-A	4etl-(5m)C
240	nmLNA-A	nmLNA-(5m)C
241	4etl-A	2'-OMe-(5m)C
242	4etl-A	2'-O-MOE-(5m)C
243	4etl-A	LNA-(5m)C
244	4etl-A	ENA-(5m)C
245	4etl-A	scp-BNA-(5m)C
246	4etl-A	AmNA-(NMe)-(5m)C
247	4etl-A	2'-O-Propargyl-(5m)C
248	4etl-A	2'-O-Butynyl-(5m)C
249	4etl-A	2'-F-(5m)C
250	4etl-A	2'-ara-F-(5m)C
251	4etl-A	3'-OMe-(5m)C
252	4etl-A	UNA-(5m)C
253	4etl-A	2'-NH ₂ -(5m)C
254	4etl-A	GNA-(5m)C

No.	A Unit	C Unit
255	4etl-A	4etl-(5m)C
256	4etl-A	nmLNA-(5m)C

[0055] The length of a modified oligonucleotide as described herein can vary over a broad range. In various embodiments, a modified oligonucleotide as described herein comprises an at least partially phosphorothioated sequence of alternating A and C units that has a sequence length of about 8 units, about 10 units, about 12 units, about 14 units, about 16 units, about 18 units, about 20 units, about 24 units, about 30 units, about 34 units, about 36 units, about 38 units, about 40 units, about 44 units, about 50 units, about 60 units, about 76 units, about 100 units, about 122 units, about 124 units, about 150 units, about 172 units, about 200 units, or a sequence length in a range between any two of the aforementioned values. For example, in an embodiment, the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 8 units to 200 units. In another embodiment, the at least partially phosphorothioated sequence of alternating A and C units has a sequence length that is in any one or more (as applicable) of the following ranges: about 8 units to about 36 units; about 16 units to about 36 units; 20 units to 36 units; 16 units to 30 units; 18 units to 60 units; 20 units to 30 units; 30 units to 50 units; 34 units to 46 units, 36 units to 44 units; 44 units to 200 units; 44 units to 150 units; 44 units to 120 units; 50 units to 200 units; 50 units to 150 units; 50 units to 120 units; 60 units to 200 units; 60 units to 150 units; and/or 60 units to 120 units.

[0056] As described elsewhere herein, a modified oligonucleotide can comprise a single at least partially phosphorothioated sequence of alternating A and C units in some embodiments, or in other embodiments the modified oligonucleotide can comprise a plurality of at least partially phosphorothioated sequences of alternating A and C units that are linked together. Thus, a modified oligonucleotide that contains a single at least partially phosphorothioated sequence of alternating A and C units can have the same sequence length as that sequence. Examples of such sequence lengths are described elsewhere herein. Similarly, a modified oligonucleotide that contains a plurality of at least partially phosphorothioated sequences of alternating A and C units can have sequence length that is the result of linking those sequences as described elsewhere herein. Examples of sequence lengths

for a modified oligonucleotide that contains a plurality of at least partially phosphorothioated sequences of alternating A and C units are expressed elsewhere herein in terms of the lengths of the individual sequences, and also taking into account the length of the linking group.

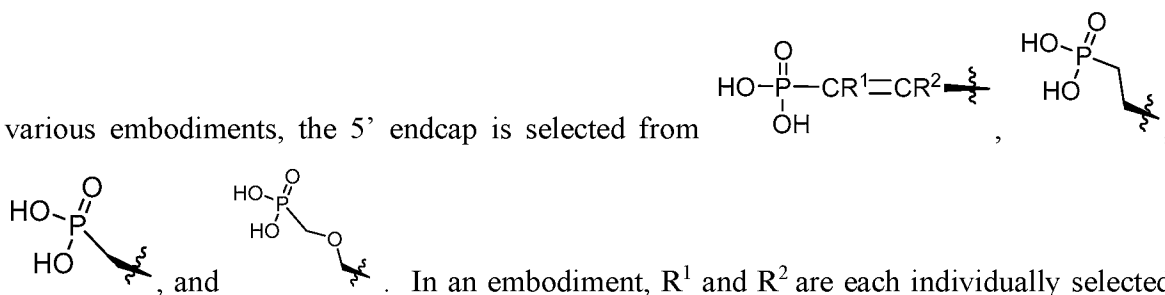
[0057] A modified oligonucleotide as described herein can comprises a plurality of at least partially phosphorothioated sequences of alternating A and C units. In an embodiment, when the sequence of alternating A and C units comprises a Ribo-A unit, the sequence further comprises at least one A unit that is not a Ribo-A unit. Similarly, in an embodiment, when the sequence of alternating A and C units comprises a Ribo-C unit, the sequence further comprises at least one C unit that is not a Ribo-C unit. In an embodiment, the modified oligonucleotide can contain one or more of various nucleotide units (known to those skilled in the art, e.g., thymine, cytosine, adenine, guanine and modified versions thereof) that are not A or C units, e.g., as an end group(s) and/or as a linking group(s) between two or more at least partially phosphorothioated sequences of alternating A and C units. For example, in an embodiment, the modified oligonucleotide comprises one or more cytosine units that link together at least two or more of the at least partially phosphorothioated sequences of alternating A and C units. In an embodiment, the two or more at least partially phosphorothioated sequences of alternating A and C units, which are linked together by a non-A/non-C linking group, are identical to one another. An example of such a modified oligonucleotide is (AC)₈-cytosine-(AC)₈. Such a modified oligonucleotide that comprises a plurality of identical sequences that are joined together may be referred to herein as a concatemer. The two or more at least partially phosphorothioated sequences of alternating A and C units that are linked together can also be different from one another. An example of such a modified oligonucleotide is (AC)₈-cytosine-(AC)₁₆.

[0058] The modified oligonucleotide can contain two or more different A groups and/or two or more different C groups. When an A or C group is replaced by a different A or C group, such a replacement is not ordinarily considered to interrupt the alternating sequence of A and C units. For example, in an embodiment, at least some of the A units are not 2'-O-methylated on the ribose ring and/or at least some of the C units are not 2'-O-methylated on the ribose ring. However, in some embodiments the group linking the two at least partially phosphorothioated sequences of alternating A and C units is itself an A or C unit that interrupts the alternating sequence of A and C units. For example, an at least partially phosphorothioated

16-mer of alternating A and C units may be linked by an A unit to another such 16-mer to form (AC)₈-A-(AC)₈. Similarly, such a 16-mer may be linked by a C unit to another such 16-mer to form (AC)₈-C-(AC)₈. As noted above, when a plurality of at least partially phosphorothioated sequences of alternating A and C units that are identical to one another are joined together by a linking group, the modified oligonucleotide may be referred to herein as a concatemer. As noted above, the two or more at least partially phosphorothioated sequences of alternating A and C units that are linked together can also be different from one another. Examples of such modified oligonucleotides include (AC)₈-A-(AC)₁₆ and (AC)₈-C-(AC)₁₆.

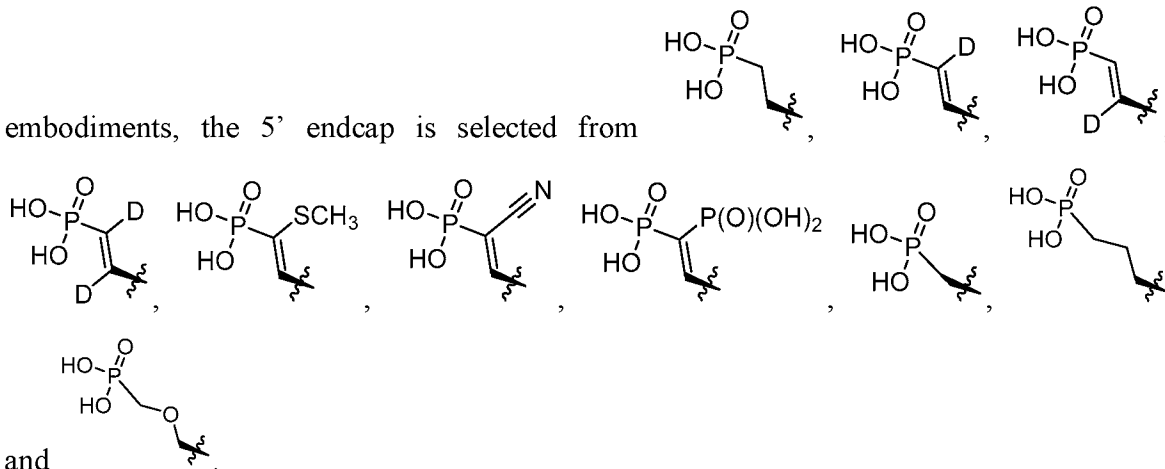
[0059] In an embodiment, the modified oligonucleotide comprises a 5' endcap. In

various embodiments, the 5' endcap is selected from



In an embodiment, R¹ and R² are each individually selected from hydrogen, deuterium, phosphate, thioC₁-alkyl, and cyano. For example, in an embodiment, R¹ and R² are both hydrogen and the modified oligonucleotide comprises a vinyl phosphonate endcap. In other embodiments, R¹ and R² are not both hydrogen. In some

embodiments, the 5' endcap is selected from



and

[0060] In other embodiments, the modified oligonucleotide comprises a 3' and/or 5' linking group. For example, with respect to modified oligonucleotide compounds comprising A and C units as described herein, such as the A and C units of Tables 1 and 2, respectively, at least one terminal $\sim\text{O}$ can be a linking group. Various linking groups known

to those skilled in the art can be used to link the modified oligonucleotide to another moiety (such as one or more second oligonucleotides and/or targeting ligands). In an embodiment, the linking group comprises a non-A/non-C linking group or an A or C unit that interrupts the alternating sequence of A and C units as discussed above, or the linking group comprises a C₂-alkylene linkage (FIG. 1), a C₂-alkylene oxide linkage, such as a propylene oxide linkage (FIG. 2), or a tetraethylene glycol (TEG) linkage (FIGS. 3A-D).

[0061] In various embodiments, two, three, four or more of the modified oligonucleotides can be connected to each other in various ways. For example, the modified oligonucleotides can be connected end-to-end via 3' and/or 5' linking groups, and/or a linking group can be connected to a one 3' or 5' end of multiple modified oligonucleotides, e.g., as illustrated in FIGS. 1 and 2.

[0062] In various embodiments, the modified oligonucleotide further comprises a targeting ligand that is attached to the modified oligonucleotide via the linking group. For example, in various embodiments the targeting ligand is, or comprises, a N-acetylgalactosamine (GalNac) (e.g., triantennary-GalNac), a tocopherol or cholesterol. FIGS. 3A and 3B illustrate embodiments of modified oligonucleotides having cholesterol attached via a 5' TEG linking group and a 3' TEG linking group, respectively. FIGS. 3C and 3D illustrate embodiments of modified oligonucleotides having a tocopherol (Vitamin E) attached via a 5' TEG linking group and a 3' TEG linking group, respectively. FIGS. 4A and 4B illustrate embodiments of modified oligonucleotides having GalNac attached via a linking group. In an embodiment, the GalNac is a triantennary GalNac.

[0063] In various embodiments, the at least partially phosphorothioated sequence of alternating A and C units can include modification(s) to one or more phosphorothioated linkages. The inclusion of such a modified linkage is not ordinarily considered to interrupt the alternating sequence of A and C units because those skilled in the art understand that such a sequence may be only partially phosphorothioated and thus may comprise one or more modifications to a phosphorothioate linkage. In various embodiments, the modification to the phosphorothioate linkage is a modified linkage selected from phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate and diphosphodiester. For example, in an embodiment, the modified linkage is a phosphodiester linkage.

[0064] In various embodiments, the at least partially phosphorothioated sequence of alternating A and C units can have various degrees of phosphorothioation. For example, in an embodiment, the at least partially phosphorothioated sequence of alternating A and C units is at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, or 95% phosphorothioated. In an embodiment, the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated. In an embodiment, the at least partially phosphorothioated sequence of alternating A and C units is fully phosphorothioated.

[0065] In various embodiments, the at least partially phosphorothioated sequence of alternating A and C units can include stereochemical modification(s) to one or more phosphorothioated linkages. In an embodiment, the modified oligonucleotides described herein can comprise at least one stereochemically defined phosphorothioate linkage. In various embodiments, the stereochemically defined phosphorothioate linkage has an R configuration. In various embodiments, the stereochemically defined phosphorothioate linkage has an S configuration.

[0066] Those skilled in the art will recognize that modified oligonucleotide compounds comprising A and C units as described herein, such as the A and C units of Tables 1 and 2, respectively, contain internal linkages between the A and C units as well as terminal groups at the 3' and 5' ends. Thus, with respect to the A and C units described herein, such as the A and C units of Tables 1 and 2, respectively, each  represents an internal  or a terminal . In various embodiments, each terminal  is independently hydroxyl, an O,O-dihydrogen phosphorothioate, a dihydrogen phosphate, an endcap or a linking group. In various embodiments, each internal  is a phosphorus-containing linkage to a neighboring A or C unit, the phosphorus-containing linkage being a phosphorothioate linkage or a modified linkage selected from phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate 5'-phosphoramidate, 3',5'-phosphordiamidate, 5'-thiophosphoramidate, 3',5'-thiophosphordiamidate or diphosphodiester.

[0067] As noted above, the STOPSTTM compounds described herein are antiviral oligonucleotides. In various embodiments, a modified oligonucleotide as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, has sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion

[illegible]

CpsmApsmCpsmApsmCpsmApsmCpsmApsmCpsmApsmCpsmApsmCpsmApsmCpsmAps
mCpsmApsmCpsmApsmCpsmApsmCpsmApsmC 3' (2'-OMe-A, 2'-OMe-C).

[0068] In various embodiments, a modified oligonucleotide as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, has sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nanomolar (nM); in a “B” activity range of 30 nM to less than 100 nM; in a “C” activity range of 100 nM to less than 300 nM; or in a “D” activity range of greater than 300 nM. In some embodiments, a modified oligonucleotide as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, has sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is less than 50 nM.

[0069] The modified oligonucleotides described herein can be prepared in the form of various complexes. Thus, an embodiment provides a chelate complex of a modified oligonucleotide as described herein. For example, in an embodiment such a chelate complex comprises a calcium, magnesium or zinc chelate complex of the modified oligonucleotide. The modified oligonucleotides described herein can also be prepared in the form of various monovalent counterion complexes. For example, in an embodiment such a counterion complex comprises a lithium, sodium or potassium complex of the modified oligonucleotide.

[0070] An embodiment provides a modified oligonucleotide or complex thereof having sequence independent antiviral activity against hepatitis B, comprising an at least partially phosphorothioated sequence of alternating A and C units as described herein, wherein:

[0071] the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;

[0072] the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;

[0073] the A units comprise at least 12 2'-OMe-A units (e.g., at least 15 2'-OMe-A units) and at least 1 Ribo-A unit (e.g., at least 2 Ribo-A units);

[0074] the C units comprise at least 15 LNA-5mC units; and

[0075] the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 100 nM (e.g., less than 50 nM or less than 30 nM).

[0076] An embodiment provides a modified oligonucleotide or complex thereof having sequence independent antiviral activity against hepatitis B, comprising an at least partially phosphorothioated sequence of alternating A and C units as described herein, wherein;

[0077] the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;

[0078] the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;

[0079] the A units comprise at least 15 2'-OMe-A units;

[0080] the C units comprise at least 7 LNA-5mC units; and

[0081] the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 100 nM (e.g., less than 50 nM or less than 30 nM).

[0082] An embodiment provides a modified oligonucleotide or complex thereof having sequence independent antiviral activity against hepatitis B, comprising an at least partially phosphorothioated sequence of alternating A and C units as described herein, wherein;

[0083] the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;

[0084] the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;

[0085] the A units comprise at least 15 2'-OMe-A units;

[0086] the C units comprise at least 3 LNA-5mC units; and

[0087] the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 100 nM (e.g., less than 50 nM or less than 30 nM).

[0088] An embodiment provides a modified oligonucleotide or complex thereof having sequence independent antiviral activity against hepatitis B, comprising an at least partially phosphorothioated sequence of alternating A and C units as described herein, wherein;

[0089] the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;

[0090] the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;

[0091] the A units comprise at least 18 2'-OMe-A units;

[0092] the C units comprise at least 15 LNA-5mC units; and

[0093] the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 100 nM (e.g., less than 50 nM or less than 30 nM).

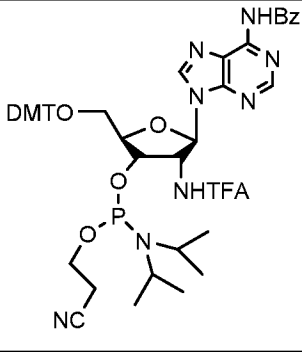
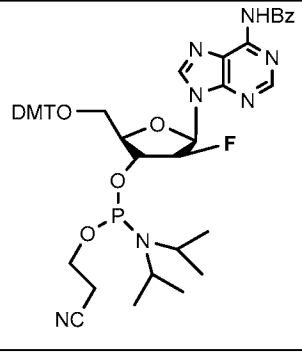
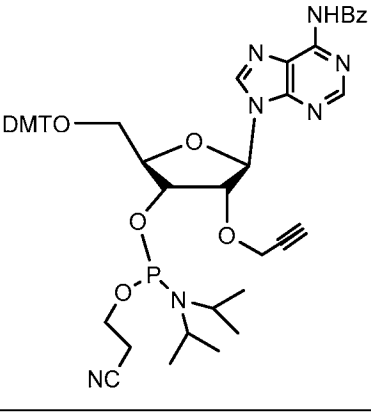
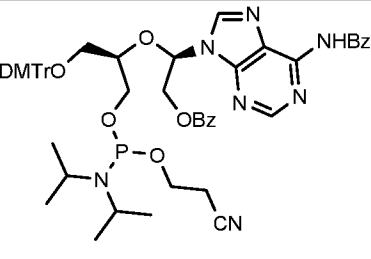
Synthesis

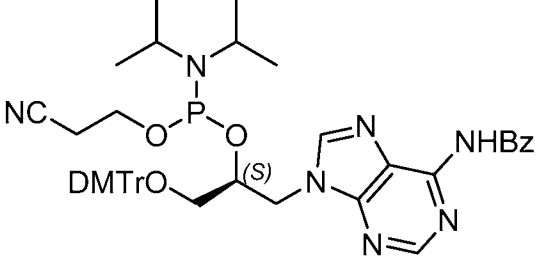
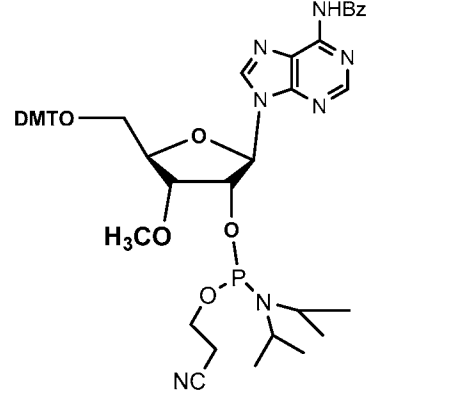
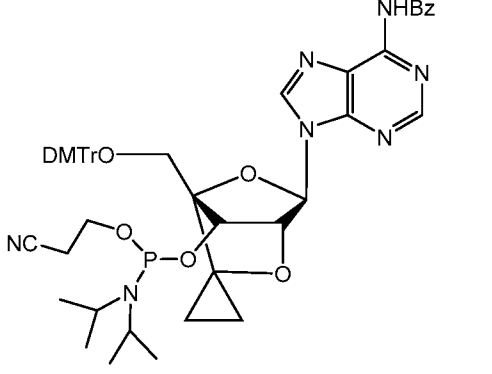
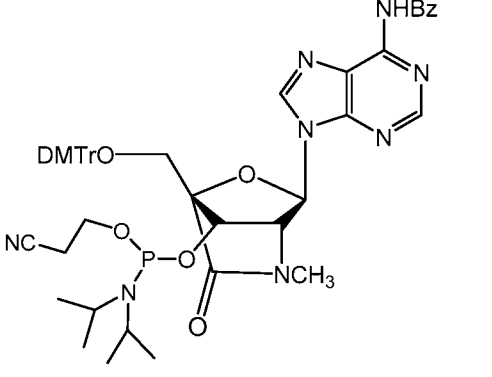
[0094] The modified oligonucleotides described herein can be prepared in various ways. In an embodiment, the building block monomers described in Tables 4 and 5 are employed to make the modified oligonucleotides described herein by applying standard phosphoramidite chemistry. The building blocks described in Tables 4 and 5 and other building block phosphoramidite monomers can be prepared by known methods or obtained from commercial sources (Thermo Fischer Scientific US, Hongene Biotechnology USA Inc., Chemgenes Corporation). Exemplary procedures for making modified oligonucleotides are set forth in the Examples below.

TABLE 4 – BUILDING BLOCKS FOR “A” UNITS

Abbreviation	Structure
2'-OMe-A PHOSPHORAMIDITE	
2'-F-A PHOSPHORAMIDITE	

Abbreviation	Structure
2'-O-MOE-A PHOSPHORAMIDITE	
LNA-A PHOSPHORAMIDITE	
ENA-A PHOSPHORAMIDITE	
2'-O-Butyne-A PHOSPHORAMIDITE	

Abbreviation	Structure
2'-NH₂-A PHOSPHORAMIDITE	
2'-F-Ara-A PHOSPHORAMIDITE	
2'-O-Propargyl-A PHOSPHORAMIDITE	
UNA-A PHOSPHORAMIDITE	

Abbreviation	Structure
GNA-A PHOSPHORAMIDITE	
3'-O-Methyl-A PHOSPHORAMIDITE	
scp-BNA-A PHOSPHORAMIDITE	
AmNA-(N-Me)-A PHOSPHORAMIDITE	

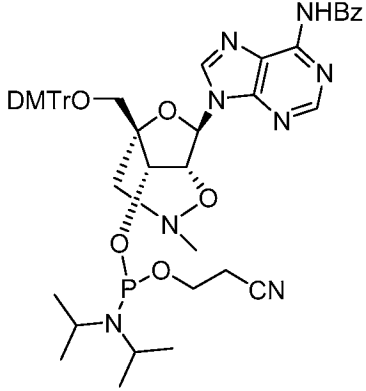
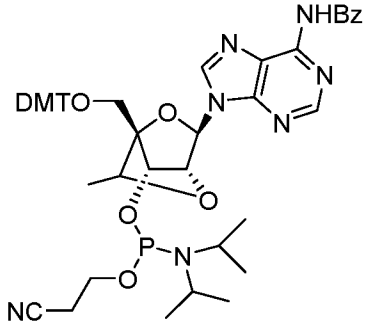
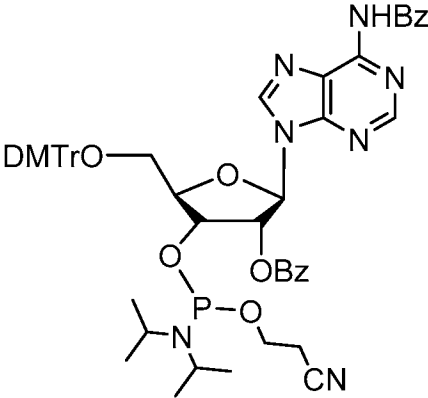
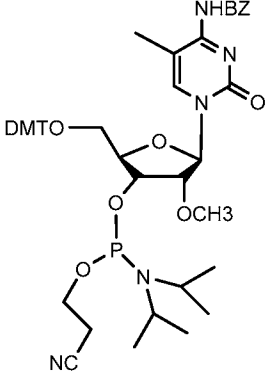
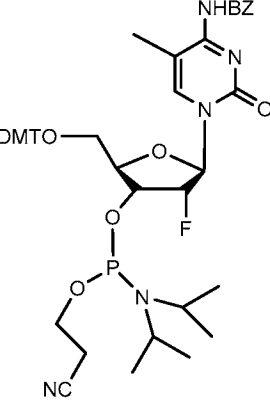
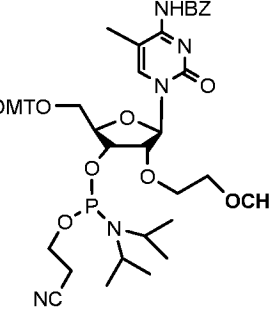
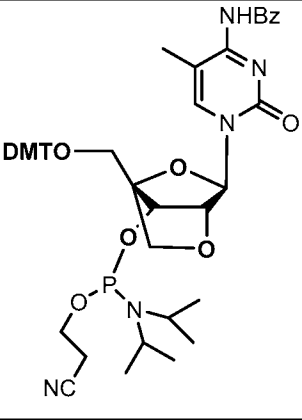
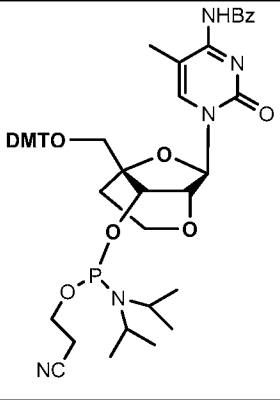
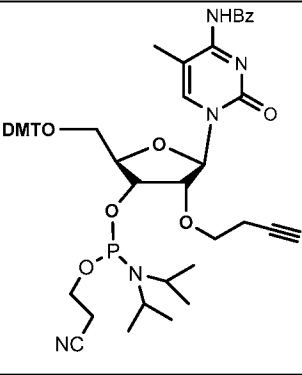
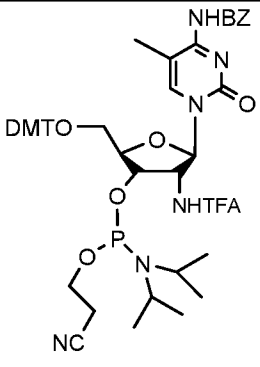
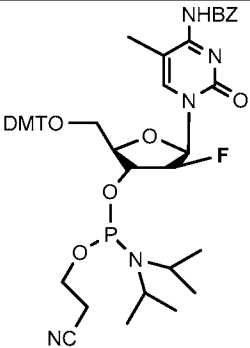
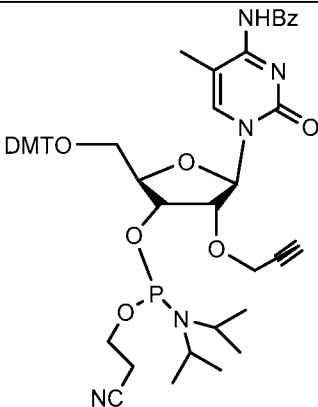
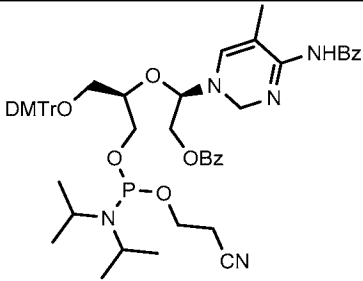
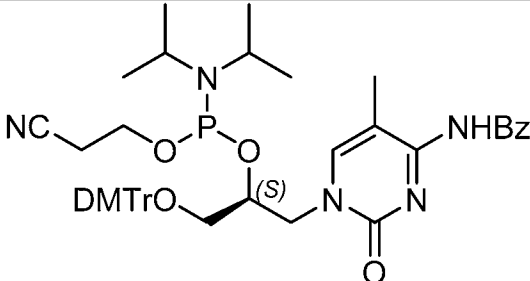
Abbreviation	Structure
nmLNA-A PHOSPHORAMIDITE	
4etl-A PHOSPHORAMIDITE	
Ribo-A PHOSPHORAMIDITE	

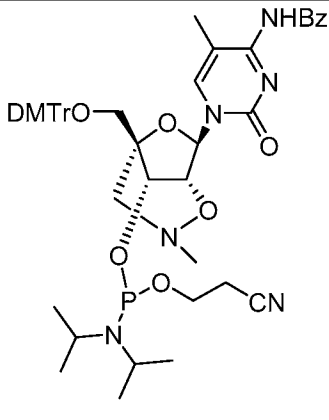
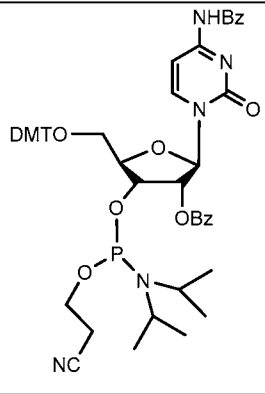
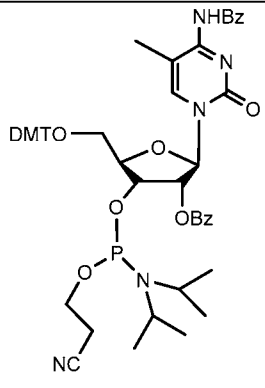
TABLE 5 – BUILDING BLOCKS FOR “C” UNITS

Abbreviation	Structure
2'-OMe-(5m)C PHOSPHORAMIDITE	
2'-F-(5m)C PHOSPHORAMIDITE	
2'-O-MOE-(5m)C PHOSPHORAMIDITE	

Abbreviation	Structure
LNA-(5m)C PHOSPHORAMIDITE	
ENA-(5m)C PHOSPHORAMIDITE	
2'-O-Butyne-(5m)C PHOSPHORAMIDITE	
2'-NH₂-(5m)C PHOSPHORAMIDITE	

Abbreviation	Structure
2'-F-Ara-(5m)C PHOSPHORAMIDITE	
2'-O-Propargyl-(5m)C PHOSPHORAMIDITE	
UNA-(5m)C PHOSPHORAMIDITE	
GNA-(5m)C PHOSPHORAMIDITE	

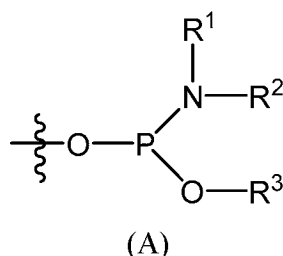
Abbreviation	Structure
3'-O-Methyl-(5m)C PHOSPHORAMIDITE	
scp-BNA-(5m)C PHOSPHORAMIDITE	
AmNA-(NMe)-(5m)C PHOSPHORAMIDITE	
4etl-(5m)C PHOSPHORAMIDITE	

Abbreviation	Structure
nmLNA-(5m)C PHOSPHORAMIDITE	
Ribo-C PHOSPHORAMIDITE	
Ribo-(5m)C PHOSPHORAMIDITE	

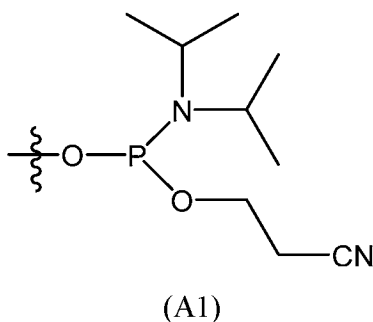
[0095] In various embodiments, the STOPSTM modified oligonucleotides described herein can also be prepared using dinucleotides that comprise or consist of any two of the building block monomers described in Tables 4 and 5. Exemplary procedures for making dinucleotides and the corresponding modified oligonucleotides are set forth in the Examples below.

[0096] An embodiment provides a dinucleotide comprising, or consisting of, an A unit and a C unit connected by a stereochemically defined phosphorothioate linkage, wherein

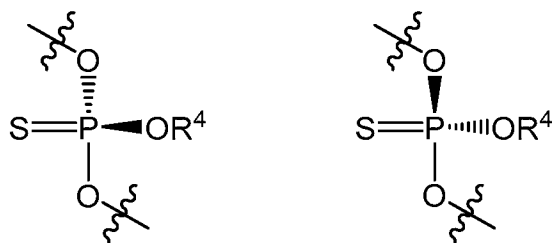
the A unit is selected from any of the building block monomers described in Table 4 and the C unit is selected from any of the building block monomers described in Table 5, and wherein each $\sim\text{O}$ is independently hydroxyl, an O,O-dihydrogen phosphorothioate, an O,O-dihydrogen phosphate, a phosphoramidite, a dimethoxytrityl ether, or the stereochemically defined phosphorothioate linkage. In an embodiment, the $\sim\text{O}$ is a phosphoramidite of the following formula (A):



[0097] In various embodiments R^1 and R^2 of formula (A) are each individually a C_{1-6} alkyl, and R^3 is a C_{1-6} alkyl or a cyano C_{1-6} alkyl. For example, in an embodiment the phosphoramidite of the formula (A) is a phosphoramidite of the following formula (A1):



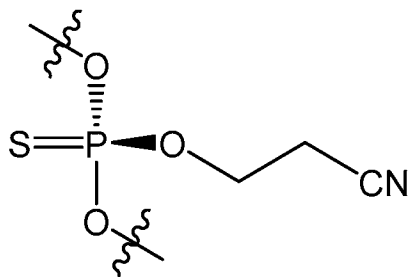
[0098] In another embodiment, the $\sim\text{O}$ is a stereochemically defined phosphorothioate linkage that is a phosphorothioate. For example, in an embodiment, the stereochemically defined phosphorothioate linkage is a phosphorothioate of the following Formula (B1) or (B2):



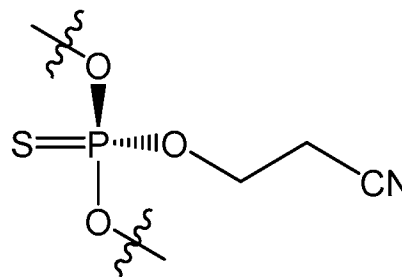
(B1)

(B2)

[0099] In various embodiments R^4 of formulae (B1) and (B2) is a C_{1-6} alkyl or a cyano C_{1-6} alkyl. For example, in an embodiment, the phosphorothioates of the formulae (B1) and (B2) are phosphorothioates of the following Formulae (B3) or (B4), respectively:



(B3)



(B4)

[0100] Various embodiments provide methods of making a modified oligonucleotide as described herein, comprising coupling one or more dinucleotides as described herein. Exemplary methods of carrying out such coupling are illustrated in the Examples below.

Pharmaceutical Compositions

[0101] Some embodiments described herein relate to a pharmaceutical composition, that can include an effective amount of a compound described herein (e.g., a STOPST[™] modified oligonucleotide compound or complex thereof as described herein) and a pharmaceutically acceptable carrier, excipient or combination thereof. A pharmaceutical composition described herein is suitable for human and/or veterinary applications.

[0102] As used herein, a “carrier” refers to a compound that facilitates the incorporation of a compound into cells or tissues. For example, without limitation, dimethyl sulfoxide (DMSO) is a commonly utilized carrier that facilitates the uptake of many organic compounds into cells or tissues of a subject.

[0103] As used herein, a “diluent” refers to an ingredient in a pharmaceutical composition that lacks pharmacological activity but may be pharmaceutically necessary or desirable. For example, a diluent may be used to increase the bulk of a potent drug whose mass is too small for manufacture and/or administration. It may also be a liquid for the dissolution of a drug to be administered by injection, ingestion or inhalation. A common form

of diluent in the art is a buffered aqueous solution such as, without limitation, phosphate buffered saline that mimics the composition of human blood.

[0104] As used herein, an “excipient” refers to an inert substance that is added to a pharmaceutical composition to provide, without limitation, bulk, consistency, stability, binding ability, lubrication, disintegrating ability etc., to the composition. A “diluent” is a type of excipient.

[0105] Proper formulation is dependent upon the route of administration chosen. Techniques for formulation and administration of the compounds described herein are known to those skilled in the art. Multiple techniques of administering a compound exist in the art including, but not limited to, oral, rectal, topical, aerosol, injection and parenteral delivery, including intramuscular, subcutaneous, intravenous, intramedullary injections, intrathecal, direct intraventricular, intraperitoneal, intranasal and intraocular injections. Pharmaceutical compositions will generally be tailored to the specific intended route of administration.

[0106] One may also administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into the infected area, optionally in a depot or sustained release formulation. Furthermore, one may administer the compound in a targeted drug delivery system, for example, in a liposome coated with a tissue-specific antibody. The liposomes may be targeted to and taken up selectively by the organ.

[0107] The pharmaceutical compositions disclosed herein may be manufactured in a manner that is itself known, *e.g.*, by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or tableting processes. As described herein, compounds used in a pharmaceutical composition may be provided as salts with pharmaceutically compatible counterions.

Methods of Use

[0108] Some embodiments described herein relate to a method of treating a HBV and/or HDV infection that can include administering to a subject identified as suffering from the HBV and/or HDV infection an effective amount of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein. Other embodiments described herein relate to using a modified oligonucleotide or complex thereof as described herein in the manufacture of a medicament for treating a HBV and/or HDV infection. Still

other embodiments described herein relate to the use of a modified oligonucleotide or complex thereof as described herein or a pharmaceutical composition that includes a modified oligonucleotide as described herein for treating a HBV and/or HDV infection.

[0109] Various routes may be used to administer a modified oligonucleotide or complex thereof to a subject in need thereof as indicated elsewhere herein. In an embodiment, the modified oligonucleotide or complex thereof is administered to the subject by a parenteral route. For example, in an embodiment, the modified oligonucleotide or complex thereof is administered to the subject intravenously. In another embodiment, the modified oligonucleotide or complex thereof is administered to the subject subcutaneously. Surprisingly, it has now been found that embodiments of a modified oligonucleotide or complex thereof as described herein can be subcutaneously administered to a primate in an amount that is both safe and effective for treatment. Previously, subcutaneous administration of a modified oligonucleotide or complex thereof (such as REP 2139, REP 2055 or those described in U.S. Patent Nos. 7,358,068; 8,008,269; 8,008,270 and 8,067,385) to a primate was considered unlikely to be safe and effective because of the relatively high dosages believed required to achieve efficacy and the concomitant increase in the potential risk of safety concerns such as undesirable injection site reactions. Thus, for example, prior clinical studies involving the administration of REP 2139 to humans are believed to have utilized only intravenous routes. At the dosage levels that were believed to be necessary for efficacy, it is believed that safety concerns such as undesirable injection site reactions would have precluded subcutaneous administration.

[0110] Unexpectedly, as illustrated in FIG. 12 and Example B5 below, it has now been found that liver exposure following subcutaneous administration to non-human primates is much higher than expected based on liver exposure levels resulting from otherwise comparable intravenous dosing. This finding means that embodiments of modified oligonucleotides or complexes thereof as described herein, and particularly embodiments of highly potent STOPS™ compounds or complexes as described herein, can be safely and effectively administered to primates via subcutaneous administration at dosages lower than previously considered likely to be effective. These lower dosages reduce the risk profile (e.g., reduce risk of injection site reactions) and thus provide a clinically acceptable safety profile for human use.

[0111] Some embodiments disclosed herein relate to a method of treating a HBV and/or HDV infection that can include contacting a cell infected with the HBV and/or HDV with an effective amount of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein. In an embodiment, such a method of treating a HBV and/or HDV infection comprises safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0112] Other embodiments described herein relate to using a modified oligonucleotide or complex thereof as described herein in the manufacture of a medicament for treating a HBV and/or HDV infection. Still other embodiments described herein relate to the use of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein for treating a HBV and/or HDV infection. In an embodiment, such uses comprise safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units,

having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0113] Some embodiments disclosed herein relate to a method of inhibiting replication of HBV and/or HDV that can include contacting a cell infected with the HBV and/or HDV with an effective amount of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein. In an embodiment, such a method of inhibiting replication of HBV and/or HDV comprises safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPST[™] compound or complex thereof as described herein. For example, in an embodiment, the STOPST[™] compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0114] Other embodiments described herein relate to using a modified oligonucleotide or complex thereof as described herein in the manufacture of a medicament for inhibiting replication of HBV and/or HDV. Still other embodiments described herein relate to the use of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein, for inhibiting replication of HBV and/or HDV. In an embodiment, such uses for inhibiting replication of HBV and/or HDV comprise safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPST[™] compound or complex thereof as described herein. For example, in an embodiment,

the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0115] In some embodiments, the HBV infection can be an acute HBV infection. In some embodiments, the HBV infection can be a chronic HBV infection.

[0116] Some embodiments disclosed herein relate to a method of treating liver cirrhosis that is developed because of a HBV and/or HDV infection that can include administering to a subject suffering from liver cirrhosis and/or contacting a cell infected with the HBV and/or HDV in a subject suffering from liver cirrhosis with an effective amount of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein. In an embodiment, such a method of treating liver cirrhosis that is developed because of a HBV and/or HDV infection comprises safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0117] Other embodiments described herein relate to using a modified oligonucleotide or complex thereof as described herein in the manufacture of a medicament for treating liver cirrhosis that is developed because of a HBV and/or HDV infection, with an effective amount of the modified oligonucleotide(s). Still other embodiments described herein relate to the use of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein for treating liver cirrhosis that is developed because of

a HBV and/or HDV infection. In an embodiment, such uses for treating liver cirrhosis comprise safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0118] Some embodiments disclosed herein relate to a method of treating liver cancer (such as hepatocellular carcinoma) that is developed because of a HBV and/or HDV infection that can include administering to a subject suffering from the liver cancer and/or contacting a cell infected with the HBV and/or HDV in a subject suffering from the liver cancer with an effective amount of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein. In an embodiment, such a method of treating liver cancer (such as hepatocellular carcinoma) that is developed because of a HBV and/or HDV infection comprises safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0119] Other embodiments described herein relate to using a modified oligonucleotide or complex thereof as described herein in the manufacture of a medicament for treating liver cancer (such as hepatocellular carcinoma) that is developed because of a HBV and/or HDV infection. Still other embodiments described herein relate to the use of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein for treating liver cancer (such as hepatocellular carcinoma) that is developed because of a HBV and/or HDV infection. In an embodiment, such uses for treating liver cancer (such as hepatocellular carcinoma) comprise safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPST[™] compound or complex thereof as described herein. For example, in an embodiment, the STOPST[™] compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0120] Some embodiments disclosed herein relate to a method of treating liver failure that is developed because of a HBV and/or HDV infection that can include administering to a subject suffering from liver failure and/or contacting a cell infected with the HBV and/or HDV in a subject suffering from liver failure with an effective amount of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein. In an embodiment, such a method of treating liver failure that is developed because of a HBV and/or HDV infection comprises safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified

oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0121] Other embodiments described herein relate to using a modified oligonucleotide or complex thereof as described herein in the manufacture of a medicament for treating liver failure that is developed because of a HBV and/or HDV infection. Still other embodiments described herein relate to the use of a modified oligonucleotide or complex thereof as described herein, or a pharmaceutical composition that includes an effective amount of a modified oligonucleotide or complex thereof as described herein for treating liver failure that is developed because of a HBV and/or HDV infection. In an embodiment, such uses for treating liver failure comprise safe and effective subcutaneous administration of the modified oligonucleotide or complex thereof to a human at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration. For example, in an embodiment, the modified oligonucleotide or complex thereof is REP-2139 or a complex thereof. In another embodiment, the modified oligonucleotide or complex thereof comprises a highly potent STOPSTTM compound or complex thereof as described herein. For example, in an embodiment, the STOPSTTM compound or complex thereof is a modified oligonucleotide or complex thereof as described herein, comprising an at least partially phosphorothioated sequence of alternating A and C units, having sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, that is in an “A” activity range of less than 30 nM.

[0122] Various indicators for determining the effectiveness of a method for treating an HBV and/or HDV infection are also known to those skilled in the art. Examples of suitable indicators include, but are not limited to, a reduction in viral load indicated by reduction in HBV DNA (or load), HBV surface antigen (HBsAg) and HBV e-antigen (HBeAg), a reduction in plasma viral load, a reduction in viral replication, a reduction in time to seroconversion (virus undetectable in patient serum), an increase in the rate of sustained viral

response to therapy, an improvement in hepatic function, and/or a reduction of morbidity or mortality in clinical outcomes.

[0123] In some embodiments, an effective amount of a modified oligonucleotide or complex thereof as described herein is an amount that is effective to achieve a sustained virologic response, for example, a sustained viral response 12 month after completion of treatment.

[0124] Subjects who are clinically diagnosed with an HBV and/or HDV infection include “naïve” subjects (e.g., subjects not previously treated for HBV and/or HDV) and subjects who have failed prior treatment for HBV and/or HDV (“treatment failure” subjects). Treatment failure subjects include “non-responders” (subjects who did not achieve sufficient reduction in ALT levels, for example, subject who failed to achieve more than 1 log₁₀ decrease from base-line within 6 months of starting an anti-HBV and/or anti-HDV therapy) and “relapsers” (subjects who were previously treated for HBV and/or HDV whose ALT levels have increased, for example, ALT > twice the upper normal limit and detectable serum HBV DNA by hybridization assays). Further examples of subjects include subjects with a HBV and/or HDV infection who are asymptomatic.

[0125] In some embodiments, a modified oligonucleotide or complex thereof as described herein can be provided to a treatment failure subject suffering from HBV and/or HDV. In some embodiments, a modified oligonucleotide or complex thereof as described herein can be provided to a non-responder subject suffering from HBV and/or HDV. In some embodiments, a modified oligonucleotide or complex thereof as described herein can be provided to a relapser subject suffering from HBV and/or HDV. In some embodiments, the subject can have HBeAg positive chronic hepatitis B. In some embodiments, the subject can have HBeAg negative chronic hepatitis B. In some embodiments, the subject can have liver cirrhosis. In some embodiments, the subject can be asymptomatic, for example, the subject can be infected with HBV and/or HDV but does not exhibit any symptoms of the viral infection. In some embodiments, the subject can be immunocompromised. In some embodiments, the subject can be undergoing chemotherapy.

[0126] Examples of agents that have been used to treat HBV and/or HDV include interferons (such as IFN- α and pegylated interferons that include PEG-IFN- α -2a), and nucleosides/nucleotides (such as lamivudine, telbivudine, adefovir dipivoxil, clevudine,

entecavir, tenofovir alafenamide and tenofovir disoproxil). However, some of the drawbacks associated with interferon treatment are the adverse side effects, the need for subcutaneous administration and high cost. A drawback with nucleoside/nucleotide treatment can be the development of resistance.

[0127] Resistance can be a cause for treatment failure. The term “resistance” as used herein refers to a viral strain displaying a delayed, lessened and/or null response to an anti-viral agent. In some embodiments, a modified oligonucleotide or complex thereof as described herein can be provided to a subject infected with an HBV and/or HDV strain that is resistant to one or more anti-HBV and/or anti-HDV agents. Examples of anti-viral agents wherein resistance can develop include lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir alafenamide and tenofovir disoproxil. In some embodiments, development of resistant HBV and/or HDV strains is delayed when a subject is treated with a modified oligonucleotide as described herein compared to the development of HBV and/or HDV strains resistant to other HBV and/or HDV anti-viral agents, such as those described.

Combination Therapies

[0128] In some embodiments, a modified oligonucleotide or complex thereof as described herein can be used in combination with one or more additional agent(s) for treating and/or inhibiting replication HBV and/or HDV. Additional agents include, but are not limited to, an interferon, nucleoside/nucleotide analogs, a capsid assembly modulator, a sequence specific oligonucleotide (such as anti-sense oligonucleotide and/or siRNA), an entry inhibitor and/or a small molecule immunomodulator. For example, in an embodiment, a modified oligonucleotide or complex thereof as described herein can be used as a first treatment in combination with one or more second treatment(s) for HBV, wherein the second treatment comprises a second oligonucleotide having sequence independent antiviral activity against hepatitis B, an siRNA oligonucleotide (or nucleotides), an anti-sense oligonucleotide, a nucleoside, an interferon, an immunomodulator, a capsid assembly modulator, or a combination thereof. Examples of additional agents include recombinant interferon alpha 2b, IFN- α , PEG-IFN- α -2a, lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir alafenamide, tenofovir disoproxil, JNJ-3989 (ARO-HBV), RG6004, GSK3228836, AB-729, VIR-2218, DCR-HBVS, JNJ-6379, GLS4, ABI-HO731, JNJ-440, NZ-4, RG7907, AB-423, AB-506 and ABI-H2158. In an embodiment, the additional agent is a capsid

assembly modulator (CAM). In an embodiment, the additional agent is an anti-sense oligonucleotide (ASO).

[0129] In some embodiments, a modified oligonucleotide or complex thereof as described herein can be administered with one or more additional agent(s) together in a single pharmaceutical composition. In some embodiments, a modified oligonucleotide or complex thereof as described herein can be administered with one or more additional agent(s) as two or more separate pharmaceutical compositions. Further, the order of administration of a modified oligonucleotide or complex thereof as described herein with one or more additional agent(s) can vary.

EXAMPLES

[0130] Additional embodiments are disclosed in further detail in the following examples, which are not in any way intended to limit the scope of the claims.

EXAMPLES 1-116

[0131] A series of modified oligonucleotides containing phosphorothioated sequences of alternating A and C units were synthesized on an ABI 394 synthesizer using standard phosphoramidite chemistry. The solid support was controlled pore glass (CPG, 1000A, Glen Research, Sterling VA) and the building block monomers are described in Tables 4 and 5. The reagent (dimethylamino-methylidene) amino)-3H-1,2,4-dithiazaoline-3-thione (DDTT) was used as the sulfur-transfer agent for the synthesis of oligoribonucleotide phosphorothioates (PS linkages). An extended coupling of 0.1M solution of phosphoramidite in CH₃CN in the presence of 5-(ethylthio)-1H-tetrazole activator to a solid bound oligonucleotide followed by standard capping, oxidation and deprotection afforded modified oligonucleotides. The stepwise coupling efficiency of all modified phosphoramidites was more than 95%. Several modified oligonucleotides containing sequences of alternating A and C units but having phosphodiester (PO) linkages instead of phosphorothioate (PS) linkages were also made.

Deprotection

[0132] After completion of synthesis the controlled pore glass (CPG) was transferred to a screw cap vial or screw caps RNase free microfuge tube. The oligonucleotide was cleaved from the support with simultaneous deprotection of base and phosphate groups with 1.0 mL of a mixture of ethanolic ammonia (ammonia: ethanol (3:1)) for 5-15 hr at 55°C.

The vial was cooled briefly on ice and then the ethanolic ammonia mixture was transferred to a new microfuge tube. The CPG was washed with 2 x 0.1 mL portions of deionized water, put in dry ice for 10 min then dried in speed vac.

Quantitation of Crude Oligomer or Raw Analysis

[0133] Samples were dissolved in deionized water (1.0 mL) and quantitated as follows: Blanking was first performed with water alone (1 mL). 20 ul of sample and 980 uL of water were mixed well in a microfuge tube, transferred to cuvette and absorbance reading obtained at 260 nm. The crude material is dried down and stored at -20°C.

HPLC Purification of Oligomer

[0134] The crude oligomers were analyzed and purified by HPLC (Dionex PA 100). The buffer system is A = Water B = 0.25 M Tris-HCl pH 8, C: 0.375 M Sodium perchlorate, flow 5.0 mL/min, wavelength 260 nm. First inject a small amount of material (~5 OD) and analyze by LC-MS. Once the identity of this material is confirmed the crude oligomer can then be purified using a larger amount of material, e.g., 60 OD's per run, flow rate of 5mL/min. Fractions containing the full-length oligonucleotides are then pooled together, evaporated and finally desalted as described below.

Desalting of Purified Oligomer

[0135] The purified dry oligomer was then desalted using Sephadex G-25M (Amersham Biosciences). The cartridge was conditioned with 10 mL of water. The purified oligomer dissolved thoroughly in 2.5 mL RNase free water was applied to the cartridge with very slow dropwise elution. The salt free oligomer was eluted with 3.5 ml water directly into a screw cap vial.

HPLC Analysis and Electrospray LC/MS

[0136] Approximately 0.2 OD oligomer is first dried down, redissolved in water (50ul) and then pipetted in special vials for HPLC and LC-MS analysis.

[0137] Table 6 summarizes the sequence length, alternating A and C units and whether the backbone is phosphorothioate (PS) or phosphodiester (PO) for the resulting exemplified modified oligonucleotides.

TABLE 6

No.	Length	A modification	C modification	Backbone
1	(AC)20	2'-OMe	2'-OMe	PS
2	(AC)15	2'-OMe	2'-OMe	PS

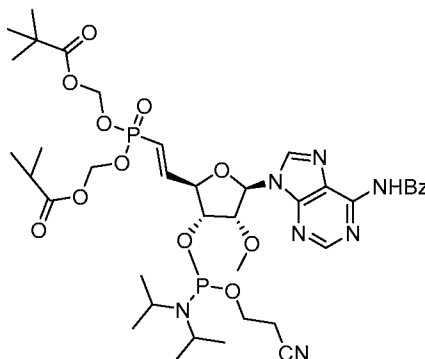
No.	Length	A modification	C modification	Backbone
3	(AC)25	2'-OMe	2'-OMe	PS
4	(AC)30	2'-OMe	2'-OMe	PS
5	(AC)20	2'-O-MOE	2'-O-MOE	PS
6	(AC)20	LNA	LNA	PS
7	(AC)20	2'-F	2'-F	PS
8	(AC)20	2'-O-Propargyl	2'-O-Propargyl	PS
9	(AC)20	2'-O-butyne	2'-O-butyne	PS
10	(AC)20	2'-F-Ara	2'-F-Ara	PS
11	(AC)20	UNA	UNA	PS
12	(AC)20	ENA	ENA	PS
13	(AC)20	2'-OMe	2'-O-MOE	PS
14	(AC)20	2'-OMe	LNA	PS
15	(AC)20	2'-OMe	2'-F	PS
16	(AC)20	2'-OMe	2'-O-Propargyl	PS
17	(AC)20	2'-OMe	2'-O-butyne	PS
18	(AC)20	2'-OMe	2'-F-Ara	PS
19	(AC)20	2'-OMe	UNA	PS
20	(AC)20	2'-OMe	ENA	PS
21	(AC)20	2'-OMe	2'-NH ₂	PS
22	(AC)20	2'-O-MOE	2'-OMe	PS
23	(AC)20	2'-O-MOE	LNA	PS
24	(AC)20	2'-O-MOE	2'-F	PS
25	(AC)20	2'-O-MOE	2'-O-Propargyl	PS
26	(AC)20	2'-O-MOE	2'-O-butyne	PS
27	(AC)20	2'-O-MOE	2'-F-Ara	PS
28	(AC)20	2'-O-MOE	UNA	PS
29	(AC)20	2'-O-MOE	ENA	PS
30	(AC)20	2'-O-MOE	2'-NH ₂	PS
31	(AC)20	LNA	2'-OMe	PS
32	(AC)20	LNA	2'-O-MOE	PS
33	(AC)20	LNA	2'-F	PS
34	(AC)20	LNA	2'-O-Propargyl	PS
35	(AC)20	LNA	2'-O-butyne	PS
36	(AC)20	LNA	2'-F-Ara	PS
37	(AC)20	LNA	UNA	PS
38	(AC)20	LNA	ENA	PS
39	(AC)20	LNA	2'-NH ₂	PS
40	(AC)20	2'-F	LNA	PS

No.	Length	A modification	C modification	Backbone
41	(AC)20	2'-F	2'-OMe	PS
42	(AC)20	2'-F	2'-O-MOE	PS
43	(AC)20	2'-F	2'-O-Propargyl	PS
44	(AC)20	2'-F	2'-O-butyne	PS
45	(AC)20	2'-F	2'-F-Ara	PS
46	(AC)20	2'-F	UNA	PS
47	(AC)20	2'-F	ENA	PS
48	(AC)20	2'-F	2'-NH ₂	PS
49	(AC)20	2'-O-Propargyl	2'-OMe	PS
50	(AC)20	2'-O-Propargyl	2'-O-MOE	PS
51	(AC)20	2'-O-Propargyl	LNA	PS
52	(AC)20	2'-O-Propargyl	2'-F	PS
53	(AC)20	2'-O-Propargyl	2'-O-butyne	PS
54	(AC)20	2'-O-Propargyl	2'-F-Ara	PS
55	(AC)20	2'-O-Propargyl	UNA	PS
56	(AC)20	2'-O-Propargyl	ENA	PS
57	(AC)20	2'-O-Propargyl	2'-NH ₂	PS
58	(AC)20	2'-O-butyne	2'-OMe	PS
59	(AC)20	2'-O-butyne	2'-O-MOE	PS
60	(AC)20	2'-O-butyne	LNA	PS
61	(AC)20	2'-O-butyne	2'-F	PS
62	(AC)20	2'-O-butyne	2'-O-Propargyl	PS
63	(AC)20	2'-O-butyne	2'-F-Ara	PS
64	(AC)20	2'-O-butyne	UNA	PS
65	(AC)20	2'-O-butyne	ENA	PS
66	(AC)20	2'-O-butyne	2'-NH ₂	PS
67	(AC)20	2'-F-Ara	2'-OMe	PS
68	(AC)20	2'-F-Ara	2'-O-MOE	PS
69	(AC)20	2'-F-Ara	LNA	PS
70	(AC)20	2'-F-Ara	2'-F	PS
71	(AC)20	2'-F-Ara	2'-O-Propargyl	PS
72	(AC)20	2'-F-Ara	2'-O-butyne	PS
73	(AC)20	2'-F-Ara	UNA	PS
74	(AC)20	2'-F-Ara	ENA	PS
75	(AC)20	2'-F-Ara	2'-NH ₂	PS
76	(AC)20	UNA	2'-OMe	PS
77	(AC)20	UNA	2'-O-MOE	PS
78	(AC)20	UNA	LNA	PS

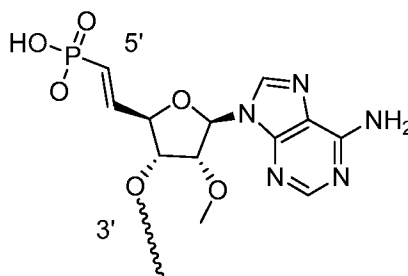
No.	Length	A modification	C modification	Backbone
79	(AC)20	UNA	2'-F	PS
80	(AC)20	UNA	2'-O-Propargyl	PS
81	(AC)20	UNA	2'-O-butyne	PS
82	(AC)20	UNA	2'-F-Ara	PS
83	(AC)20	UNA	ENA	PS
84	(AC)20	UNA	2'-NH ₂	PS
85	(AC)20	ENA	2'-OMe	PS
86	(AC)20	ENA	2'-O-MOE	PS
87	(AC)20	ENA	LNA	PS
88	(AC)20	ENA	2'-F	PS
89	(AC)20	ENA	2'-O-Propargyl	PS
90	(AC)20	ENA	2'-O-butyne	PS
91	(AC)20	ENA	2'-F-Ara	PS
92	(AC)20	ENA	UNA	PS
93	(AC)20	ENA	2'-NH ₂	PS
94	(AC)20	LNA	2'-O-MOE	PS
95	(AC)20	2'-F	LNA	PS
96	(AC)25	2'-OMe	2'-OMe	PO
97	(AC)25	2'-OMe	2'-OMe	PS
98	(AC)20	2'-F	2'-OMe	PS
99	(AC)20	LNA	2'-O-Me	PS
100	(AC)20	2'-OMe	2'-F	PS
101	(AC)20	2'-OMe	2'-OMe	PO
102	(AC)20	2'-F	2'-O-MOE	PS
103	(AC)30	2'-OMe	2'-OMe	PS
104	(AC)15	2'-OMe	2'-OMe	PS
105	(AC)20	2'-OMe	LNA	PS
106	(AC)20	LNA	LNA	PS
107	(AC)20	2'-OMe	2'-O'MOE	PS
108	(AC)20	2'-O-MOE	2'-OMe	PS
109	(AC)20	2'-OMe	2'-OMe	PS
110	(AC)30	2'-OMe	2'-O-butyne	PO
111	(AC)20	2'-F	2'-F	PS
112	(AC)20	2'-OMe	2'-OMe	PS
113	(AC)15	2'-OMe	2'-OMe	PO
114	(AC)20	2'-O-MOE	2'-O-MOE	PS
115	(AC)20	2'-O-MOE	2'-F	PS
116	(AC)20	LNA	2'-F	PS

EXAMPLES 117-130

[0138] The effect of 5' modification was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above in Examples 1-116. End capped oligonucleotides were made by using a 5'-vinyl phosphonate building block to incorporate 5'-vinyl phosphonate endcaps:



5'-vinyl phosphonate building block (5'-VP)



Modified oligo with 5'-vinyl phosphonate endcap

[0139] With reference to FIG. 7, the 5'-vinyl phosphonate building block (5'-VP) was prepared as follows:

[0140] Preparation of compound **7-2**: To a solution of **7-1** (15.0 g, 53.3 mmol) in dry pyridine (150 mL) was added TBSCl (20.0 g, 133.3 mmol) and Imidazole (10.8 g, 159.9 mmol). The mixture was stirred at r.t. for 15h. TLC showed **7-1** was consumed completely. The reaction mixture was concentrated in vacuo to give residue. The residue was quenched with DCM (500 mL). The DCM layer was washed with H₂O (1 L*2) 2 times and brine. The DCM layer concentrated in vacuo to give crude **7-2** (27.2 g, 53.3 mmol) as a yellow oil. The crude **7-2** was used in next step directly. ESI-LCMS m/z 510.5 [M+H]⁺.

[0141] Preparation of compound **7-3**: To a solution of **7-2** (26.2 g, 51.3 mmol) in pyridine (183 mL) was added dropwise the benzoyl chloride (15.8 g, 113.0 mmol) at 5°C. The

reaction mixture was stirred at r.t. for 2 hours. TLC showed the **7-2** was consumed completely. The reaction mixture was quenched with H₂O (4 mL). Then NH₃.H₂O (20 mL) was added to the reaction mixture and stirred at r.t for 30 min. Then the Pyridine was removed from the mixture by concentration under reduced pressure. The residue was added to H₂O (100 mL) and extracted with EA (150 mL*3) and the EA layers combined. The EA layer was washed with brine and dried over Na₂SO₄. Filtered and concentrated to give the crude **7-3** (45.0 g). ESI-LCMS m/z = 614.5 [M+H]⁺.

[0142] Preparation of compound **7-4**: To a mixture solution of **7-3** (44.0 g, crude) in THF (440 mL) was added the H₂O (220 mL) and TFA (220 mL) at 0°C. Then the reaction mixture was stirred at 0°C for 1.5 h. TLC showed the **7-3** was consumed completely. The reaction mixture pH was adjusted to 7-8 with NH₃.H₂O. Then the mixture was extracted with EA (300 mL*7). The combined EA layer was washed with brine and concentrated in vacuo to give crude. The crude was purified by column chromatography (EA: PE = 1:5~1:1) to give compound **7-4** (15.8 g) as a white solid. ¹H-NMR (400 MHz, DMSO-d₆): δ = 11.24 (s, 1H, exchanged with D₂O), 8.77 (s, 2H), 8.04-8.06 (m, 2H), 7.64-7.66 (m, 2H), 7.54-7.58 (m, 2H), 6.14-6.16 (d, *J* = 5.9 Hz, 1H), 5.20-5.23 (m, 1H), 4.58-4.60 (m, 1H), 4.52-4.55 (m, 1H), 3.99-4.01 (m, 1H), 3.69-3.75 (m, 1H), 3.57-3.61 (m, 1H), 3.34 (s, 4H), 0.93 (s, 9H), 0.14-0.15 (d, *J* = 1.44 Hz, 6H). ESI-LCMS m/z = 500.3 [M+H]⁺.

[0143] Preparation of compound **7-5**: To a 500 mL round-bottom flask was added the DMSO (132 mL) and **7-4** (13.2 g, 26.4 mmol), EDCI (15.19 g, 79.2 mmol) in turn at r.t. Then the Pyridine (2.09 g, 26.4 mmol, 2.1 mL) was added to the reaction mixture. After stirring 5 min, the TFA (1.51 g, 13.2 mmol) was added to the reaction mixture. Then reaction mixture was stirred at r.t for 3 hrs. LC-MS showed the **7-4** was consumed completely. The reaction mixture was added to the ice water (500 mL) and extracted with EA (300 mL*3) 3 times. The combined EA layer was washed with H₂O 2 times and brine 1 time. Dried over Na₂SO₄ and filtered. The filtrate was concentrated to get crude **7-5** (14.6 g) as a white solid. ESI-LCMS m/z = 516.3 [M+H]⁺.

[0144] Preparation of compound **7-6**: The **5A** (24.4 g, 38.5 mmol) was added to a mixture solution of NaH (2.5 g, 64.3 mmol, 60% purity) in THF (50 mL) at 0°C. After stirring 15 min, the **7-5** (16.0 g, 32.1 mmol) in THF (60 mL) was added to the reaction mixture. Then the reaction mixture was stirred at r.t for 1 hr. LC-MS showed the **7-5** was consumed

completely. Then the reaction mixture was quenched with sat. NH_4Cl (500 mL) and extracted with EA (400 mL*3) 3 times. The combined EA layer was washed with brine, dried over Na_2SO_4 and filtered. The filtrate was concentrated in vacuo to get crude. The crude was purified by c.c (EA: PE = 1:5 ~ 1:1) to give **7-6** (10.0 g, 12.4 mmol, 38.6% yield) as a white solid. ESI-LCMS $m/z = 804.4$ $[\text{M}+\text{H}]^+$; ^{31}P NMR (162 MHz, $\text{DMSO}-d_6$) δ 17.01.

[0145] Preparation of compound **7-7**: To a 500 mL round-bottom flask was added the **7-6** (9.0 g, 11.2 mmol) and H_2O (225 mL), HCOOH (225 mL) in turn. The reaction mixture was stirred at 26°C for 15 h. LC-MS showed the **7-6** was consumed completely. The reaction mixture was adjusted the pH = 6-7 with $\text{NH}_3\cdot\text{H}_2\text{O}$. Then the mixture was extracted with EA (300 mL*3) 3 times. The combined EA layer was dried over Na_2SO_4 , filtered and filtrate was concentrated to get crude. The crude was purified by column chromatography (DCM/ MeOH = 100:1 ~ 60:1) to give product **7-7** (7.0 g, 10.1 mmol, 90.6% yield). ^1H -NMR (400 MHz, $\text{DMSO}-d_6$): δ = 11.11 (s, 1H, exchanged with D_2O), 8.71-8.75 (d, $J=14.4$, 2H), 8.04-8.06 (m, 2H), 7.64-7.65 (m, 1H), 7.54-7.58 (m, 2H), 6.88-7.00 (m, 1H), 6.20-6.22 (d, $J=5.4$, 2H), 6.06-6.16 (m, 1H), 5.74-5.75 (d, $J=5.72$, 2H), 5.56-5.64 (m, 4H), 4.64-4.67 (m, 1H), 4.58-4.59 (m, 1H), 4.49-4.52 (m, 1H), 3.37 (s, 3H), 1.09-1.10 (d, $J=1.96$, 18H). ^{31}P NMR (162 MHz, $\text{DMSO}-d_6$) δ 17.45. ESI-LCMS $m/z = 690.4$ $[\text{M}+\text{H}]^+$.

[0146] Preparation of compound **5'-VP**: To a solution of **7-7** (5.5 g, 7.9 mmol) in DCM (55 mL) was added the DCI (750 mg, 6.3 mmol), then $\text{CEP}[\text{N}(\text{iPr})_2]_2$ (3.1 g, 10.3 mmol) was added. The mixture was stirred at r.t. for 2 h. TLC showed 3.5% of **7-7** remained. The reaction mixture was washed with H_2O (40 mL*2) and brine (50 mL*2), dried over Na_2SO_4 and concentrated to give crude. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/5 increasing to $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0 within 30 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 3/1; Detector, UV 254 nm. The product was concentrated and extracted with EA (50 mL* 3). The combined EA layer was washed with brine and dried over Na_2SO_4 , filtered and filtrate was concentrated to get resulting **5'-VP** (6.0 g, 98% purity) as a white solid. ^1H -NMR (400 MHz, $\text{DMSO}-d_6$): δ = 11.27 (s, 1H, exchanged with D_2O), 8.72-8.75 (m, 2H), 8.04-8.06 (m, 2H), 7.54-7.68 (m, 3H), 6.85-7.05 (m, 1H), 6.09-6.26 (m, 2H), 5.57-5.64 (m, 4H), 4.70-4.87 (m, 3H), 3.66-3.88 (m, 4H), 3.37-

3.41 (m, 3H), 2.82-2.86 (m, 2H), 1.20-1.21 (m, 12H), 1.08-1.09 (m, 18H). ^{31}P NMR (162 MHz, DMSO- d_6): 149.99, 149.16, 17.05, 16.81. ESI-LCMS m/z = 890.8 $[\text{M}+\text{H}]^+$.

[0147] Table 7 summarizes the sequence length, alternating A and C units, and 5' modification for the resulting exemplified modified phosphorothioated oligonucleotides.

TABLE 7

No.	Length	A	C	5'-Modification
117	(AC)17	LNA-A	LNA-(5m)C	OH
118	(AC)18	LNA-A	LNA-(5m)C	OH
119	(AC)19	LNA-A	LNA-(5m)C	OH
120	(AC)17	LNA-A	LNA-(5m)C	Vinyl-phosphonate-A
121	(AC)18	LNA-A	LNA-(5m)C	Vinyl-phosphonate-A
122	(AC)19	LNA-A	LNA-(5m)C	Vinyl-phosphonate-A
123	(AC)20	LNA-A	LNA-(5m)C	Vinyl-phosphonate-A
124	(AC)17	2'-OMe-A	2'-OMe-(5m)C	OH
125	(AC)18	2'-OMe-A	2'-OMe-(5m)C	OH
126	(AC)19	2'-OMe-A	2'-OMe-(5m)C	OH
127	(AC)17	2'-OMe-A	2'-OMe-(5m)C	Vinyl-phosphonate-A
128	(AC)18	2'-OMe-A	2'-OMe-(5m)C	Vinyl-phosphonate-A
129	(AC)19	2'-OMe-A	2'-OMe-(5m)C	Vinyl-phosphonate-A
130	(AC)20	2'-OMe-A	2'-OMe-(5m)C	Vinyl-phosphonate-A

EXAMPLES 131-174

[0148] The effect of sequence length, LNA incorporation and 5'-modification was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above in Examples 1-116. Table 8 summarizes the sequence length, alternating A and C units, 5' modification, and length and position of LNA units for the resulting exemplified modified phosphorothioated oligonucleotides.

TABLE 8

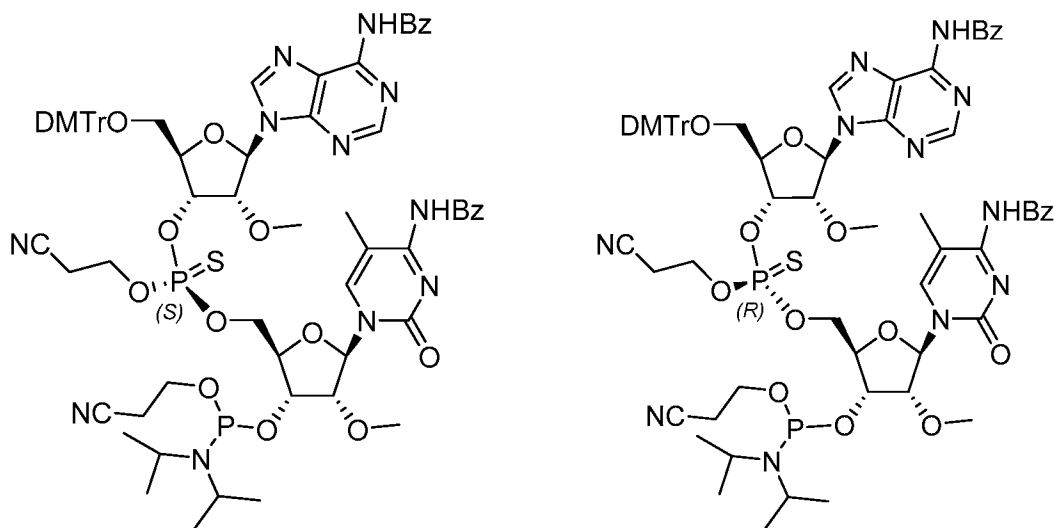
No.	Length	A	C	5'-Modification	LNA Modification
131	(AC)17	2'-OMe-A	LNA-(5m)C	OH	
132	(AC)18	2'-OMe-A	LNA-(5m)C	OH	
133	(AC)19	2'-OMe-A	LNA-(5m)C	OH	
134	(AC)20	2'-OMe-A	LNA-(5m)C	OH	
135	(AC)17	2'-OMe-A	LNA-(5m)C	Vinyl-phosphonate-A	
136	(AC)18	2'-OMe-A	LNA-(5m)C	Vinyl-phosphonate-A	
137	(AC)19	2'-OMe-A	LNA-(5m)C	Vinyl-phosphonate-A	

No.	Length	A	C	5'-Modification	LNA Modification
138	(AC)20	2'-OMe-A	LNA-(5m)C	Vinyl-phosphonate-A	
139	(AC)17	LNA-A	2'-OMe-(5m)C	OH	
140	(AC)18	LNA-A	2'-OMe-(5m)C	OH	
141	(AC)19	LNA-A	2'-OMe-(5m)C	OH	
142	(AC)20	LNA-A	2'-OMe-(5m)C	OH	
143	(AC)17	LNA-A	2'-OMe-(5m)C	Vinyl-phosphonate-A	
144	(AC)18	LNA-A	2'-OMe-(5m)C	Vinyl-phosphonate-A	
145	(AC)19	LNA-A	2'-OMe-(5m)C	Vinyl-phosphonate-A	
146	(AC)20	LNA-A	2'-OMe-(5m)C	Vinyl-phosphonate-A	
147	(AC)17	UNA-A	LNA-(5m)C	OH	
148	(AC)18	UNA-A	LNA-(5m)C	OH	
149	(AC)19	UNA-A	LNA-(5m)C	OH	
150	(AC)20	UNA-A	LNA-(5m)C	OH	
151	(AC)17	UNA-A	LNA-(5m)C	Vinyl-phosphonate-A	
152	(AC)18	UNA-A	LNA-(5m)C	Vinyl-phosphonate-A	
153	(AC)19	UNA-A	LNA-(5m)C	Vinyl-phosphonate-A	
154	(AC)20	UNA-A	LNA-(5m)C	Vinyl-phosphonate-A	
155	(AC)17	LNA-A	UNA-(5m)C	OH	
156	(AC)18	LNA-A	UNA-(5m)C	OH	
157	(AC)19	LNA-A	UNA-(5m)C	OH	
158	(AC)20	LNA-A	UNA-(5m)C	OH	
159	(AC)20	LNA-A	UNA-(5m)C	OH	Block of 4 LNA
160	(AC)17	LNA-A	UNA-(5m)C	Vinyl-phosphonate-A	
161	(AC)18	LNA-A	UNA-(5m)C	Vinyl-phosphonate-A	
162	(AC)19	LNA-A	UNA-(5m)C	Vinyl-phosphonate-A	
163	(AC)20	LNA-A	UNA-(5m)C	Vinyl-phosphonate-A	
164	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	OH	Every 3 rd base is LNA
165	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	Vinyl-phosphonate-A	Every 3 rd base is LNA
166	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	OH	Every 4th base is LNA
167	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	Vinyl-phosphonate-A	Every 4th base is LNA
168	(AC)17	2'-OMe-A	2'-OMe-(5m)C LNA(5m)C	Vinyl-phosphonate-A	5 (5m)InC in the middle
169	(AC)18	2'-OMe-A	2'-OMe-(5m)C	Vinyl-phosphonate-A	6 InAps(5m)C in the middle
170	(AC)19	2'-OMe-A	2'-OMe-(5m)C LNA(5m)C	Vinyl-phosphonate-A	6 InAps(5m)C in the middle

No.	Length	A	C	5'-Modification	LNA Modification
171	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA(5m)C	OH	5 (5m)lnC in the middle
172	(AC)20	LNA-A 2'-OMe-A	2'-OMe-(5m)C LNA(5m)C	OH	10 lnAps(5m)C in the middle
173	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA(5m)C	Vinyl-phosphonate-A	5 (5m)lnC in the middle
174	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	Vinyl-phosphonate-A	10 lnAps(5m)C in the middle

EXAMPLES 175-216

[0149] The effect of sequence length, LNA incorporation, stereochemical modification and 5' modification was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above in Examples 1-116, except that the oligonucleotides were prepared by a modified method using a dinucleotide building block consisting of an A unit and a C unit connected by a stereochemically defined phosphorothioate linkage as follows:



2'-OMeAps $\mathbf{R}$ (5m)mC phosphoramidite (**9R**) 2'-OMeAps $\mathbf{S}$ (5m)mC phosphoramidite (**9S**)

[0150] With reference to FIGS. 8, 9A and 9B, the dinucleotide building blocks **9R** and **9S** were prepared as follows:

[0151] Preparation of compound **8-2**: To a solution of **8-1** (300.0 g, 445.1 mmol) in 3000 mL of dry dioxane with an inert atmosphere of nitrogen was added levulinic acid (309.3 g, 2.67 mol) dropwise at room temperature. Then the dicyclohexylcarbodiimide (274.6 g, 1.33 mol) and 4-dimethylaminopyridine (27.1 g, 222.0 mmol) were added in order at room

temperature. The resulting solution was stirred at room temperature for 5 h and diluted with 5000 mL of dichloromethane and filtered. The organic phase was washed with 2 x 3000 mL of 2% aqueous sodium bicarbonate and 1 x 3000 mL of water respectively. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated under reduced pressure. 345.0 g (crude) of **8-2** was obtained as a white solid and used for next step without further purification. ESI-LCMS: m/z 774 $[M+H]^+$.

[0152] Preparation of compound **8-3**: To a solution of **8-2** (345 g, 445.1 mmol) was dissolved in 3000 mL dichloromethane with an inert atmosphere of nitrogen was added *p*-toluenesulfonic acid (84.6 g, 445.1 mmol) dropwise at 0 °C. The resulting solution was stirred at 0 °C for 0.5 h and diluted with 3000 mL of dichloromethane and washed with 2 x 2000 mL of saturated aqueous sodium bicarbonate and 1 x 2000 mL of saturated aqueous sodium chloride respectively. The organic phase was dried over anhydrous sodium sulfate, and concentrated under reduced pressure and the residue was purified by silica gel column chromatography (SiO₂, dichloromethane: methanol = 30:1) to give **8-3** (210.0 g, 90% over two steps) as a white solid. ¹H-NMR (400 MHz, DMSO-*d*₆) δ = 12.88 (s, 1H), 8.17-8.10 (m, 3H), 7.62-7.60 (m, 1H), 7.58-7.48 (m, , 2H), 5.97-5.91 (m, 1H), 5.42 (d, *J* = 5.9 Hz, 1H), 5.25 (s, 1H), 4.21-4.08 (m, 2H), 3.78-3.59 (m, 2H), 2.75-2.74 (m, 2H), 2.57 (m, 2H), 2.13 (d, *J* = 2.3 Hz, 3H), 2.02 (s, 3H), 1.81 (m, 1H), 1.77-1.56 (m, 1H), 1.33-0.98 (m, 1H). ESI-LCMS: m/z 474 $[M+H]^+$.

[0153] Preparation of compound **8-4**: To a solution of **8-3** (210.0 g, 444.9 mmol) in 2000 mL of acetonitrile with an inert atmosphere of nitrogen was added **8-3a** (360.0 g, 405.4 mmol) and ETT (58.0 g, 445.9 mmol) in order at 0°C. The resulting solution was stirred for 2 h at room temperature. Then the mixture was filtered and used for next step without further purification. ESI-LCMS: m/z 1258 $[M+H]^+$.

[0154] Preparation of compounds **8-5** and **8-6**: To a solution of **8-4** (509.9 g, 405.4 mmol) in 2000 mL of acetonitrile with an inert atmosphere of nitrogen was added pyridine (128.0 g, 1.62 mol) and 5-amino-3H-1,2,4-dithiazole-3-thione (121.8 g, 810.9 mmol) in order at room temperature. The reaction solution was stirred for 30 minutes at room temperature. The resulting solution was filtered and concentrated under reduced pressure. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/1 increasing to CH₃CN/H₂O (0.5%

NH_4HCO_3) = 1/0 within 20 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in a mixture of **8-5** and **8-6** (430.0 g, 90% over two steps) as a white solid. The fractions were diluted with 3000 mL of dichloromethane. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated under reduced pressure. The residue was purified by SFC with the following conditions: CHIRALPAK IB N-5(IB50CD-VD008)/SFC 0.46 cm I.D. $\times$ 25 cm L 10.0 mL Mobile phase: (DCM/EtOAc=80/20(V/V)), Detector, UV 254 nm. The fractions were concentrated until no residual solvent left under reduced pressure. 105.0 g (35.0%) of **8-5** were obtained as a white solid and used to make **9R** as described below. ^1H -NMR (400 MHz, DMSO-d_6) δ = 12.88 (s, 1H), 11.26 (s, 1H), 8.62 (d, J = 8.06 Hz, 2H), 8.18 (m, 2H), 8.05 (d, J = 7.2 Hz, 2H), 7.79 (s, 1H), 7.67-7.48 (m, 6H), 7.40 (d, J = 7.2 Hz, 2H), 7.28-7.18 (m, 7H), 6.86-6.83 (m, 4H), 6.21 (d, J = 6.6 Hz, 1H), 5.91 (d, J = 5.0 Hz, 1H), 5.44-5.41 (m, 1H), 5.28-5.26 (m, 1H), 5.06 (m, 1H), 4.45-4.24 (m, 7H), 3.71 (s, 6H), 3.39 (s, 4H), 3.31 (s, 3H), 2.98 (m, 2H), 2.75 (m, 2H), 2.56 (m, 2H), 2.01 (s, 3H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 67.17. ESI-LCMS: m/z 1292 $[\text{M}+\text{H}]^+$; 170.0 g (56.6%) of **8-6** were obtained as a white solid and used to make **9S** as described below. ^1H -NMR (400 MHz, DMSO-d_6) δ = 12.86 (s, 1H), 11.25 (s, 1H), 8.62 (d, J = 16.6 Hz, 2H), 8.18 (d, J = 7.2 Hz, 2H), 8.05 (m, 2H), 7.78 (s, 1H), 7.67-7.48 (m, 6H), 7.40 (d, J = 7.2 Hz, 2H), 7.28-7.18 (m, 7H), 6.87-6.85 (m, 4H), 6.21 (d, J = 6.8 Hz, 1H), 5.91 (d, J = 5.2 Hz, 1H), 5.43-5.39 (m, 1H), 5.28-5.26 (m, 1H), 5.06 (m, 1H), 4.48-4.21 (m, 7H), 3.72 (s, 6H), 3.36 (s, 4H), 3.26 (s, 3H), 2.95 (m, 2H), 2.73 (m, 2H), 2.55 (m, 2H), 2.04 (s, 3H); ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 66.84; ESI-LCMS: m/z 1292 $[\text{M}+\text{H}]^+$.

[0155] Preparation of compound **9-1**: To a solution of **8-5** (100.0 g, 77.4 mmol) in 700 mL acetonitrile with an inert atmosphere of nitrogen was added 0.5 M hydrazine hydrate (20.0 g, 0.4 mol) in pyridine/acetic acid (3:2) at 0°C. The resulting solution was stirred for 0.5 h at 0°C. Then the reaction was added 2,4-pentanedione at once, the mixture was allowed to warm to room temperature and stirred for additional 15 min. The solution was diluted with DCM (2000 mL) and washed with sat. aq. NH_4Cl twice and washed with brine and dried over Na_2SO_4 . Then the solution was concentrated under reduced pressure and the residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/1 increasing to $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0 within 20 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5%

NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in **9-1** (67.0 g, 80%) as a white solid. ^1H -NMR (400 MHz, DMSO-d_6) δ = 12.97 (s, 1H), 11.26 (s, 1H), 8.62 (d, J = 11.2 Hz, 2H), 8.19 (d, J = 7.2 Hz, 2H), 8.05 (m, 2H), 7.74 (s, 1H), 7.67-7.48. (m, 6H), 7.40 (d, J = 7.2 Hz, 2H), 7.28-7.18 (m, 7H), 6.85 (m, 4H), 6.21 (m, 1H), 5.90 (d, J = 3.2 Hz, 1H), 5.49-5.43 (m, 2H), 5.05 (m, 1H), 4.45 (m, 1H), 4.40-4.30 (m, 4H), 4.18-4.11 (m, 2H), 3.93 (m, 1H), 3.71 (s, 6H), 3.40-3.32 (m, 8H), 2.98 (m, 2H), 2.04 (s, 3H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 67.30. ESI-LCMS: m/z 1194 $[\text{M}+\text{H}]^+$.

[0156] Preparation of compound **9R**: To a solution of **9-1** (58.0 g, 48.6 mmol) in 600 mL of dichloromethane with an inert atmosphere of nitrogen was added $\text{CEP}[\text{N}(\text{iPr})_2]_2$ (18.7 g, 62.1 mmol) and DCI (5.1 g, 43.7 mmol) in order at room temperature. The resulting solution was stirred for 1 hour at room temperature and diluted with 1000 mL dichloromethane and washed with 2 x 1000 mL of saturated aqueous sodium bicarbonate and 1 x 1000 mL of saturated aqueous sodium chloride respectively. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated until no residual solvent left under reduced pressure. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/1 increasing to $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0 within 20 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in **9R** (51.2 g, 70%) as a white solid. ^1H -NMR (400 MHz, DMSO-d_6) δ = 12.94 (m, 1H), 11.26 (s, 1H), 8.62 (m, 2H), 8.19 (d, J = 7.2 Hz, 2H), 8.05 (m, 2H), 7.77 (m, 1H), 7.69-7.46 (m, 6H), 7.39 (d, J = 6.6 Hz, 2H), 7.26-7.20 (m, 7H), 6.84 (m, 4H), 6.20 (m, 1H), 5.90 (m, 1H), 5.43 (m, 1H), 5.06 (s, 1H), 4.46-4.17 (m, 7H), 4.12 (m, 1H), 3.82-3.80 (m, 2H), 3.73-3.66 (s, 6H), 3.64-3.58 (m, 2H), 3.48-3.29 (m, 8H), 2.98 (s, 2H), 2.82-2.77 (m, 2H), 2.03 (s, 3H), 1.24-1.15 (m, 12H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 149.87, 149.80, 67.43, 67.33. ESI-LCMS: m/z 1394 $[\text{M}+\text{H}]^+$.

[0157] Preparation of compound **9-2**: To a solution of **8-6** (110.0 g, 85.1 mmol) in 700 mL acetonitrile with an inert atmosphere of nitrogen was added 0.5 M hydrazine hydrate (21.1 g, 423.6 mmol) in pyridine/acetic acid (3:2) at 0°C. The resulting solution was stirred for 0.5 h at 0°C. Then the reaction was added 2,4-pentanedione at once, the mixture was allowed to warm to room temperature and stirred for additional 15 min, The solution was diluted with DCM (2000 mL) and washed with sat. aq. NH_4Cl twice and washed with brine and dried over Na_2SO_4 . Then the solution was concentrated under reduced pressure and the residue was

purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/1 increasing to CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/0 within 20 min, the eluted product was collected at CH₃CN/ H₂O (0.5% NH₄HCO₃) = 1/0; Detector, UV 254 nm. This resulted in **9-2** (72.0 g, 80%) as a white solid. ¹H-NMR (400 MHz, DMSO-d₆) δ = 12.94 (s, 1H), 11.24 (s, 1H), 8.61-8.57 (m, 2H), 8.18 (d, *J* = 7.6 Hz, 2H), 8.03 (d, *J* = 7.6 Hz, 2H), 7.74 (s, 1H), 7.66-7.47 (m, 6H), 7.40 (d, *J* = 7.1 Hz, 2H), 7.27-7.20 (m, 7H), 6.86 (m, 4H), 6.20 (d, *J* = 6.6 Hz, 1H), 5.87 (d, *J* = 4.0 Hz, 1H), 5.42 (m, 2H), 5.05 (m, 1H), 4.45 (m, 2H), 4.40-4.24 (m, 1H), 4.22-4.06 (m, 4H), 3.92 (m, 1H), 3.71 (s, 6H), 3.40-3.32 (m, 8H), 2.94 (m, 2H), 2.03 (m, 3H). ³¹P-NMR (162 MHz, DMSO-d₆) δ = 66.87. ESI-LCMS: *m/z* 1194 [M+H]⁺.

[0158] Preparation of compound **9S**: To a solution of **9-2** (62.0 g, 51.9 mmol) in 600 mL of dichloromethane with an inert atmosphere of nitrogen was added CEP[N(iPr)₂]₂ (19.0 g, 63.1 mmol) and DCI (5.55 g, 47.0 mmol) in order at room temperature. The resulting solution was stirred for 1 hour at room temperature and diluted with 1000 mL dichloromethane and washed with 2 x 1000 mL of saturated aqueous sodium bicarbonate and 1 x 1000 mL of saturated aqueous sodium chloride respectively. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated until no residual solvent left under reduced pressure. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/1 increasing to CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/0 within 20 min, the eluted product was collected at CH₃CN/ H₂O (0.5% NH₄HCO₃) = 1/0; Detector, UV 254 nm. This resulted in **9S** (51.5 g, 70%) as a white solid. ¹H-NMR (400 MHz, DMSO-d₆) δ = 12.90 (s, 1H), 11.25 (s, 1H), 8.60 (m, 2H), 8.19 (d, *J* = 6.6 Hz, 2H), 8.04 (m, 2H), 7.77 (s, 1H), 7.67-7.48 (m, 6H), 7.41 (d, *J* = 8.0 Hz, 2H), 7.29-7.19 (m, 7H), 6.85 (m, 4H), 6.21 (d, *J* = 6.8 Hz, 1H), 5.91-5.87 (m, 1H), 5.41 (m, 1H), 5.06 (m, 1H), 4.46-4.21 (m, 7H), 4.10 (m, 1H), 3.83-3.75 (m, 2H), 3.73-3.68 (s, 6H), 3.68-3.59 (m, 2H), 3.40-3.32 (m, 8H), 2.93 (m, 2H), 2.80 (m, 2H), 2.02 (s, 3H), 1.18-1.13 (m, 12H). ³¹P-NMR (162 MHz, DMSO-d₆) δ = 149.96, 149.73, 66.99, 66.86. ESI-LCMS: *m/z* 1394 [M+H]⁺.

[0159] The modified method also used a longer coupling time (8 min) and a greater number of equivalents of amidites (8 equivalents). Table 9 summarizes the sequence length, alternating A and C units, the number and type (R or S) of stereochemically defined

phosphorothioate (PS) linkages, and 5'-modification for the resulting exemplified modified phosphorothioated oligonucleotides.

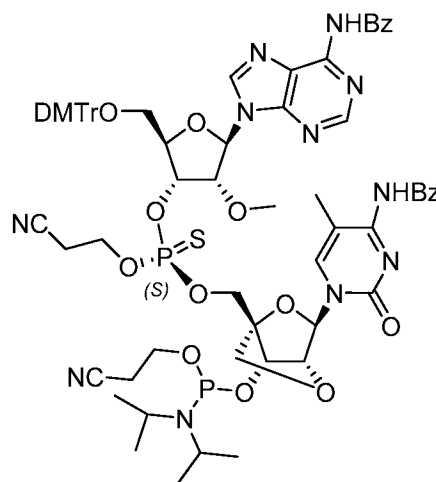
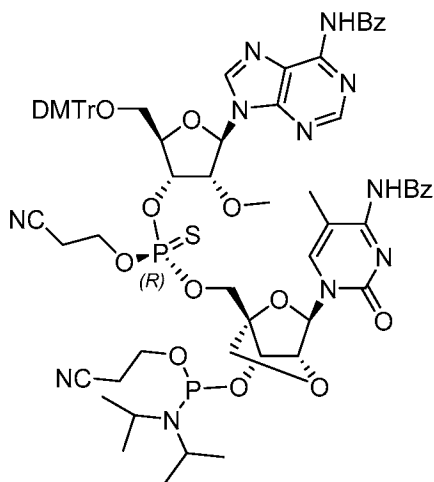
TABLE 9

No.	Length	A	C	PS Modification	5'-Modification
175	(AC)17	2'-OMe-A	2'-OMe-(5m)C	5 R isomer	OH
176	(AC)18	2'-OMe-A	2'-OMe-(5m)C	6 R isomer	OH
177	(AC)19	2'-OMe-A	2'-OMe-(5m)C	4 R isomer	OH
178	(AC)20	2'-OMe-A	2'-OMe-(5m)C	4 R isomer	OH
179	(AC)20	2'-OMe-A	2'-OMe-(5m)C	5 R isomer	OH
180	(AC)20	2'-OMe-A	2'-OMe-(5m)C	6 R isomer	OH
181	(AC)20	2'-OMe-A	2'-OMe-(5m)C	6 R isomer	Vinyl-phosphonate-A
182	(AC)20	2'-OMe-A	2'-OMe-(5m)C	7 R isomer	OH
183	(AC)20	2'-OMe-A	2'-OMe-(5m)C	13 R isomer	OH
184	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20 R isomer	OH
185	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20 R isomer	Vinyl-phosphonate-A
186	(AC)20	2'-OMe-A	2'-OMe-(5m)C	19 R isomer	Vinyl-phosphonate-A
187	(AC)17	LNA-A	LNA-(5m)C	5 R isomer	OH
188	(AC)18	LNA-A	LNA-(5m)C	6 R isomer	OH
189	(AC)19	LNA-A	LNA-(5m)C	6 R isomer	OH
190	(AC)20	LNA-A	LNA-(5m)C	4 R isomer	OH
191	(AC)20	LNA-A	LNA-(5m)C	5 R isomer	OH
192	(AC)20	LNA-A	LNA-(5m)C	6 R isomer	OH
193	(AC)20	LNA-A	LNA-(5m)C	6 R isomer,	Vinyl-phosphonate-A
194	(AC)20	LNA-A	LNA-(5m)C	13 R isomer	OH
195	(AC)20	LNA-A	LNA-(5m)C	20 R isomer	OH
196	(AC)20	LNA-A	LNA-(5m)C	20 R isomer	Vinyl-phosphonate-A
197	(AC)17	2'-OMe-A	2'-OMe-(5m)C	5 S isomer	OH
198	(AC)18	2'-OMe-A	2'-OMe-(5m)C	6 S isomer	OH
199	(AC)19	2'-OMe-A	2'-OMe-(5m)C	6 S isomer	OH
200	(AC)20	2'-OMe-A	2'-OMe-(5m)C	4 S isomer	OH
201	(AC)20	2'-OMe-A	2'-OMe-(5m)C	5 S isomer	OH
202	(AC)20	2'-OMe-A	2'-OMe-(5m)C	6 S isomer	OH
203	(AC)20	2'-OMe-A	2'-OMe-(5m)C	7 S isomer	OH
204	(AC)20	2'-OMe-A	2'-OMe-(5m)C	13 S isomer	OH
205	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20 S isomer	OH
206	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20 S isomer	Vinyl-phosphonate-A
207	(AC)17	LNA-A	LNA-(5m)C	5 S isomer	OH
208	(AC)18	LNA-A	LNA-(5m)C	6 S isomer	OH
209	(AC)19	LNA-A	LNA-(5m)C	6 S isomer	OH

No.	Length	A	C	PS Modification	5'-Modification
210	(AC) ₂₀	LNA-A	LNA-(5m)C	4 S isomer	OH
211	(AC) ₂₀	LNA-A	LNA-(5m)C	5 S isomer	OH
212	(AC) ₂₀	LNA-A	LNA-(5m)C	6 S isomer	OH
213	(AC) ₂₀	LNA-A	LNA-(5m)C	6 S isomer	Vinyl-phosphonate-A
214	(AC) ₂₀	LNA-A	LNA-(5m)C	13 S isomer	OH
215	(AC) ₂₀	LNA-A	LNA-(5m)C	20 S isomer	OH
216	(AC) ₂₀	LNA-A	LNA-(5m)C	20 S isomer	Vinyl-phosphonate-A

EXAMPLES 217-234

[0160] The effect of sequence length, LNA incorporation, stereochemical modification and 5' modification was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above in Examples 175-216, except that the oligonucleotides were prepared by a modified method using a dinucleotide building block consisting of an A unit and a C unit connected by a stereochemically defined phosphorothioate linkage as follows:



2'-OMeApsR(5m)InC phosphoramidite (**11R**) 2'-OMeApsS(5m)InC phosphoramidite (**11S**)

[0161] With reference to FIGS. 10, 11A and 11B, the dinucleotide building blocks **11R** and **11S** were prepared as follows:

[0162] Preparation of compound **10-2**: To a solution of **10-1** (50.0 g, 74.0 mmol) in 500 mL of dry dioxane with an inert atmosphere of nitrogen was added levulinic acid (51.5 g, 44.4 mol) dropwise at room temperature. Then the dicyclohexylcarbodiimide (45.7 g, 0.2 mol) and 4-dimethylaminopyridine (4.6 g, 37.0 mmol) were added in order at room

temperature. The resulting solution was stirred at room temperature for 5 h and diluted with 3000 mL of dichloromethane and filtered. The organic phase was washed with 2 x 1000 mL of 2% aqueous sodium bicarbonate and 1 x 1000 mL of water respectively. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated under reduced pressure. 52.0 g (crude) of **10-2** was obtained as a white solid and used for next step without further purification. ESI-LCMS: m/z 774 $[M+H]^+$.

[0163] Preparation of compound **10-3**: To a solution of **10-2** (52.0 g, 67.0 mmol) was dissolved in 400 mL dichloromethane with an inert atmosphere of nitrogen was added *p*-toluenesulfonic acid (51.5 g, 0.4 mol) dropwise at 0 °C. The resulting solution was stirred at 0 °C for 0.5 h and diluted with 2000 mL of dichloromethane and washed with 2 x 1000 mL of saturated aqueous sodium bicarbonate and 1 x 1000 mL of saturated aqueous sodium chloride respectively. The organic phase was dried over anhydrous sodium sulfate and concentrated under reduced pressure and the residue was purified by silica gel column chromatography (SiO₂, dichloromethane: methanol = 30:1) to give **10-3** (32.0 g, 80% over two steps) as a white solid. ¹H-NMR (400 MHz, DMSO-*d*₆) δ = 13.05 (s, 1H), 8.20-7.91 (m, 4H), 7.60-7.49 (m, 4H), 5.57 (m, 2H), 5.32 (d, J = 10.8 Hz, 1H), 4.88 (s, 1H), 4.49 (s, 1H), 4.18 (s, 1H), 3.91-3.78 (m, 5H), 2.74-2.69 (m, 4H), 2.59-2.49 (m, 7H), 2.10 (s, 5H), 2.06 (s, 4H), 1.74-1.49 (m, 3H), 1.26-1.02 (m, 3H). ESI-LCMS: m/z 472 $[M+H]^+$.

[0164] Preparation of compound **10-4**: To a solution of **10-3** (28.0 g, 59.4 mmol) in 300 mL of acetonitrile with an inert atmosphere of nitrogen was added **8-3a** (50.0 g, 56.3 mmol) and ETT (7.9 g, 59.4 mmol) in order at 0°C. The resulting solution was stirred for 2 h at room temperature. Then the mixture was filtered and used for next step without further purification. ESI-LCMS: m/z 1258 $[M+H]^+$.

[0165] Preparation of compounds **10-5** and **10-6**: To a solution of **10-4** (70.9 g, 56.3 mmol) in 300 mL of acetonitrile with an inert atmosphere of nitrogen was added pyridine (17.8 g, 225.2 mmol) and 5-amino-3H-1,2,4-dithiazole-3-thione (16.9 g, 112.6 mmol) in order at room temperature. The reaction solution was stirred for 30 minutes at room temperature. The resulting solution was filtered and concentrated under reduced pressure. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/1 increasing to CH₃CN/H₂O (0.5% NH₄HCO₃) = 1/0 within 20 min, the eluted product was collected at CH₃CN/ H₂O (0.5%

NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in a mixture of **10-5** and **10-6**. The fractions were diluted with 3000 mL of dichloromethane. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated under reduced pressure. The residue was purified by SFC with the following conditions: CHIRAL CEL OD-H/SFC 20mm*250mmL 5um (Phase A: CO_2 ; Phase B: 50% ethanol-50% acetonitrile), Detector, UV 220 nm. The fractions were concentrated until no residual solvent left under reduced pressure. 9.0 g (25.7%) of **10-5** were obtained as a white solid and used to make **11R** as described below. ^1H -NMR (400 MHz, DMSO-d_6) δ = 13.06 (s, 1H), 11.28 (s, 1H), 8.63 (d, J = 20 Hz, 2H), 8.20 (m, 2H), 8.05 (d, J = 8 Hz, 2H), 7.84 (s, 1H), 7.67-7.39 (m, 8H), 7.28-7.19 (m, 7H), 6.86-6.83 (m, 4H), 6.24 (d, J = 6.6 Hz, 1H), 5.66 (s, 2H), 5.45-5.43 (m, 1H), 5.10-5.03 (m, 2H), 4.82-4.76 (m, 1H), 4.60 (s, 1H), 4.50-4.33 (m, 4H), 4.03-3.96 (m, 2H), 3.72 (s, 6H), 3.41-3.35 (m, 7H), 3.03-3.00 (m, 2H), 2.75-2.72 (m, 2H), 2.56-2.53 (m, 2H), 2.08-2.05 (m, 6H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 67.02. ESI-LCMS: m/z 1290 $[\text{M}+\text{H}]^+$. 15.0 g (42.8%) of **10-6** were obtained as a white solid and used to make **11S** as described below. ^1H -NMR (400 MHz, DMSO-d_6) δ = 13.05 (s, 1H), 11.26 (s, 1H), 8.63 (d, J = 24 Hz, 2H), 8.-7.96 (m, 4H), 7.76 (s, 1H), 7.67-7.39 (m, 8H), 7.28-7.19 (m, 7H), 6.86 (d, J = 7.2 Hz, 4H), 6.24 (d, J = 6.4 Hz, 1H), 5.76 (s, 1H), 5.63 (s, 1H), 5.43-5.41 (m, 1H), 5.12 (m, 1H), 4.97 (s, 1H), 4.82-4.79 (m, 1H), 4.57-4.49 (m, 3H), 4.27-4.25 (m, 2H), 4.07-4.03 (m, 2H), 3.72 (s, 6H), 3.44-3.36 (m, 6H), 2.96 (m, 2H), 2.74-2.71 (m, 2H), 2.55-2.53 (m, 2H), 2.08 (s, 3H), 1.94 (s, 3H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 66.58. ESI-LCMS: m/z 1290 $[\text{M}+\text{H}]^+$.

[0166] Preparation of compound **11-1**: To a solution of **10-5** (10.0 g, 7.7 mmol) in 100 mL acetonitrile with an inert atmosphere of nitrogen was added 0.5 M hydrazine hydrate (1.8 g, 37.5 mmol) in pyridine/acetic acid (3:2) at 0°C. The resulting solution was stirred for 0.5 h at 0°C. Then the reaction was added 2,4-pentanedione at once, the mixture was allowed to warm to room temperature and stirred for additional 15 min. The solution was diluted with DCM (500 mL) and washed with sat. aq. NH_4Cl twice and washed with brine and dried over Na_2SO_4 . Then the solution was concentrated under reduced pressure and the residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/1 increasing to $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0 within 20 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in **11-1** (6.0 g, 65%) as a white solid.

^1H -NMR (400 MHz, DMSO- d_6) δ = 13.13 (s, 1H), 11.28 (s, 1H), 8.63 (d, J = 20 Hz, 2H), 8.21 (d, J = 8 Hz, 2H), 8.06-7.95 (m, 3H), 7.80 (s, 1H), 7.67-7.48 (m, 8H), 7.40 (d, J = 7.6 Hz, 2H), 7.32-7.19 (m, 10H), 6.85 (m, 5H), 6.24 (d, J = 8 Hz, 1H), 6.04 (d, J = 4.0 Hz, 1H), 5.57 (s, 2H), 5.44-5.42 (m, 1H), 5.19-5.17 (m, 2H), 5.10-5.08 (m, 1H), 4.80-4.76 (m, 2H), 4.50 (d, J = 5.6 Hz, 1H), 4.37-4.32 (m, 4H), 4.06-3.99 (m, 2H), 3.81 (m, 1H), 3.72 (s, 7H), 3.40-3.36 (m, 8H), 3.03-3.00 (m, 2H), 2.05 (m, 3H). ^{31}P -NMR (162 MHz, DMSO- d_6) δ = 67.21. ESI-LCMS: m/z 1192 $[\text{M}+\text{H}]^+$.

[0167] Preparation of compound **11R**: To a solution of **11-1** (6.0 g, 5.0 mmol) in 60 mL of dichloromethane with an inert atmosphere of nitrogen was added CEP[N(iPr) $_2$] $_2$ (1.9 g, 6.5 mmol) and DCI (0.6 g, 5.0 mmol) in order at room temperature. The resulting solution was stirred for 1 hours at room temperature and diluted with 1000 mL dichloromethane and washed with 2 x 250 mL of saturated aqueous sodium bicarbonate and 1 x 250 mL of saturated aqueous sodium chloride respectively. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated until no residual solvent left under reduced pressure. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, CH $_3$ CN/H $_2$ O (0.5% NH $_4$ HCO $_3$) = 1/1 increasing to CH $_3$ CN/H $_2$ O (0.5% NH $_4$ HCO $_3$) = 1/0 within 20 min, the eluted product was collected at CH $_3$ CN/ H $_2$ O (0.5% NH $_4$ HCO $_3$) = 1/0; Detector, UV 254 nm. This resulted in **11R** (5.0 g, 70%) as a white solid. ^1H -NMR (400 MHz, DMSO- d_6) δ = 13.10 (s, 1H), 11.28 (s, 1H), 8.20 (d, J = 8.0 Hz, 2H), 8.04 (d, J = 7.2 Hz, 2H), 7.79 (d, J = 14 Hz, 2H), 7.67-7.48 (m, 6H), 7.39 (d, J = 7.2 Hz, 2H), 7.27-7.18 (m, 7H), 6.85-6.82 (m, 4H), 6.23-6.20 (m, 1H), 5.64 (d, J = 6.0 Hz, 1H), 5.44-5.41 (m, 1H), 5.08-5.07 (m, 1H), 4.82-4.77 (m, 1H), 4.56-4.46 (m, 3H), 4.36-4.30 (m, 2H), 4.22 (d, J = 7.2 Hz, 1H), 3.98 (m, 1H), 3.89 (m, 1H), 3.71 (s, 7H), 3.59-3.55 (m, 2H), 3.40-3.34 (m, 10H), 3.02-2.98 (m, 2H), 2.77-2.72 (m, 2H), 2.08-2.05 (m, 3H), 1.13-1.08 (m, 12H). ^{31}P -NMR (162 MHz, DMSO- d_6) δ = 148.71, 148.11, 67.51, 67.44. ESI-LCMS: m/z 1392 $[\text{M}+\text{H}]^+$.

[0168] Preparation of compound **11-2**: To a solution of **10-6** (10.0 g, 7.7 mmol) in 100 mL acetonitrile with an inert atmosphere of nitrogen was added 0.5 M hydrazine hydrate (1.8 g, 37.5 mmol) in pyridine/acetic acid (3:2) at 0°C. The resulting solution was stirred for 0.5 h at 0°C. Then the reaction was added 2,4-pentanedione at once, the mixture was allowed to warm to room temperature and stirred for additional 15 min. The solution was diluted with

DCM (500 mL) and washed with sat. aq. NH_4Cl twice and washed with brine and dried over Na_2SO_4 . Then the solution was concentrated under reduced pressure and the residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/1 increasing to $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0 within 20 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in **11-2** (7.5 g, 80%) as a white solid. ^1H -NMR (400 MHz, DMSO-d_6) δ = 13.11 (s, 1H), 11.26 (s, 1H), 8.63 (d, J = 20 Hz, 2H), 8.20 (d, J = 7.2 Hz, 2H), 8.15 (m, 3H), 7.73 (s, 1H), 7.66-7.47 (m, 8H), 7.41 (d, J = 7.6 Hz, 2H), 7.32-7.19 (m, 10H), 6.85 (m, 5H), 6.24 (m, 1H), 5.99 (s, 1H), 5.54 (s, 1H), 5.41 (m, 1H), 5.10 (m, 1H), 4.79-4.75 (m, 1H), 4.57-4.49 (m, 3H), 4.30-4.24 (m, 4H), 4.02 (m, 2H), 3.85 (m, 1H), 3.72 (s, 7H), 3.38-3.35 (m, 7H), 2.95 (m, 2H), 1.98 (m, 3H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 66.79. ESI-LCMS: m/z 1192 $[\text{M}+\text{H}]^+$.

[0169] Preparation of compound **11S**: To a solution of **11-2** (7.0 g, 5.0 mmol) in 70 mL of dichloromethane with an inert atmosphere of nitrogen was added $\text{CEP}[\text{N}(\text{iPr})_2]_2$ (2.0 g, 6.5 mmol) and DCI (0.6 g, 5.0 mmol) in order at room temperature. The resulting solution was stirred for 1 hours at room temperature and diluted with 1000 mL dichloromethane and washed with 2 x 250 mL of saturated aqueous sodium bicarbonate and 1 x 250 mL of saturated aqueous sodium chloride respectively. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated until no residual solvent left under reduced pressure. The residue was purified by Flash-Prep-HPLC with the following conditions (IntelFlash-1): Column, C18 silica gel; mobile phase, $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/1 increasing to $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0 within 20 min, the eluted product was collected at $\text{CH}_3\text{CN}/\text{H}_2\text{O}$ (0.5% NH_4HCO_3) = 1/0; Detector, UV 254 nm. This resulted in **11S** (6.3 g, 70%) as a white solid. ^1H -NMR (400 MHz, DMSO-d_6) δ = 13.10 (s, 1H), 11.27 (s, 1H), 8.65 (s, 1H), 8.61 (s, 1H), 8.19 (m, 2H), 8.02 (d, J = 7.2 Hz, 2H), 7.76-7.73 (m, 1H), 7.66-7.47 (m, 6H), 7.40 (d, J = 7.2 Hz, 2H), 7.28-7.19 (m, 7H), 6.86-6.85 (m, 4H), 6.24 (d, J = 6.8 Hz, 1H), 5.62 (m, 1H), 5.43-5.41 (m, 1H), 5.10 (s, 1H), 4.84-4.78 (m, 1H), 4.66-4.49 (m, 3H), 4.30-4.18 (m, 3H), 4.04-3.95 (m, 2H), 3.83-3.77 (m, 1H), 3.72 (s, 7H), 3.62-3.54 (m, 2H), 3.44-3.32 (m, 6H), 2.96-2.92 (m, 2H), 2.77-2.72 (m, 2H), 1.98-1.97 (m, 3H), 1.12-1.11 (m, 12H). ^{31}P -NMR (162 MHz, DMSO-d_6) δ = 148.53, 148.09, 67.04. ESI-LCMS: m/z 1392 $[\text{M}+\text{H}]^+$.

[0170] As in Examples 175-216, the modified method also used a longer coupling time (8 min) and a greater number of equivalents of amidites (8 equivalents). Table 10 summarizes the sequence length, alternating A and C units, the number and type (R or S) of stereochemically defined phosphorothioate (PS) linkages, and 5' modification for the resulting exemplified modified phosphorothioated oligonucleotides.

TABLE 10

No.	Length	A	C	PS Modification	5'-Modification
217	(AC)17	2'-OMe-A	2'-OMe-(5m)C	5; 2'-OMeApsR(5m)lnC	OH
218	(AC)18	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsR(5m)lnC	OH
219	(AC)19	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsR(5m)lnC	OH
220	(AC)20	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsR(5m)lnC	OH
221	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20; 2'-OMeApsR(5m)lnC	OH
222	(AC)17	2'-OMe-A	2'-OMe-(5m)C	5; 2'-OMeApsR(5m)lnC	Vinyl-phosphonate-A
223	(AC)18	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsR(5m)lnC	Vinyl-phosphonate-A
224	(AC)19	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsR(5m)lnC	Vinyl-phosphonate-A
225	(AC)20	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsR(5m)lnC	Vinyl-phosphonate-A
226	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20; 2'-OMeApsR(5m)lnC	Vinyl-phosphonate-A
227	(AC)17	2'-OMe-A	2'-OMe-(5m)C	5; 2'-OMeApsS(5m)lnC	OH
228	(AC)18	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsS(5m)lnC	OH
229	(AC)19	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsS(5m)lnC	OH
230	(AC)20	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsS(5m)lnC	OH
231	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20; 2'-OMeApsS(5m)lnC	OH
232	(AC)17	2'-OMe-A	2'-OMe-(5m)C	5; 2'-OMeApsS(5m)lnC	Vinyl-phosphonate-A
233	(AC)18	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsS(5m)lnC	Vinyl-phosphonate-A
234	(AC)19	2'-OMe-A	2'-OMe-(5m)C	6; 2'-OMeApsS(5m)lnC	Vinyl-phosphonate-A

EXAMPLES 235-240

[0171] The effect of branching was evaluated by preparing a series of phosphorothioated oligonucleotides having a branched doubler design in which two of the oligonucleotides are attached to one another via a linking group. An example of a phosphorothioated oligonucleotide having a doubler design is illustrated in FIG. 1. Table 11

summarizes the sequence length, alternating A and C units, and 5' modification for the resulting exemplified phosphorothioated oligonucleotides.

TABLE 11

No.	Length	A	C	5'-Modification
235	(AC)9-(5m)lnC	LNA-A	LNA-(5m)C	5' OH, 19mer
236	(AC)15-(5m)lnC	LNA-A	LNA-(5m)C	5' OH, 31mer
237	(AC)20-(5m)lnC	LNA-A	LNA-(5m)C	5' OH, 41mer
238	(AC)9-(5m)mC	2'-OMe-A	2'-OMe-(5m)C	5' OH, 19mer
239	(AC)15-(5m)mC	2'-OMe-A	2'-OMe-(5m)C	5' OH, 31mer
240	(AC)20-(5m)mC	2'-OMe-A	2'-OMe-(5m)C	5' OH, 41mer

EXAMPLES 241-246

[0172] The effect of branching was evaluated by preparing a series of phosphorothioated oligonucleotides having a branched trebler design in which three phosphorothioated oligonucleotides are attached to one another via a linking group. An example of a phosphorothioated oligonucleotide having a trebler design is illustrated in FIG. 2. Table 12 summarizes the sequence length, alternating A and C units, and 5' modification for the resulting exemplified phosphorothioated oligonucleotides.

TABLE 12

No.	Length	A	C	5'-Modification
241	(AC)10-TREB-(5m)mC	LNA-A	LNA-(5m)C	5' OH, 31mer
242	(AC)13- TREB- (5m)mC	LNA-A	LNA-(5m)C	5' OH, 40mer
243	(AC)15- TREB- (5m)mC	LNA-A	LNA-(5m)C	5' OH, 46mer
244	(AC)10-TREB-(5m)mC	2'-OMe-A	2'-OMe-(5m)C	5' OH, 31mer
245	(AC)13- TREB- (5m)mC	2'-OMe-A	2'-OMe-(5m)C	5' OH, 40mer
246	(AC)15- TREB- (5m)mC	2'-OMe-A	2'-OMe-(5m)C	5' OH, 46mer

EXAMPLES 247-252

[0173] The effect of amido-bridge nucleic acid (AmNA-(N-Me)) modification and spirocyclopropylene-bridged nucleic acid (scp-BNA) modification was evaluated by preparing a series of modified phosphorothioated oligonucleotides. The AmNA-N-Me 6-*N*-benzoyladenine (A^{Bz}), 4-*N*-benzoyl -5-methyl cytidine were obtained from Luxna Biotech Co, Ltd and scp-BNA phosphoramidite monomers with 6-*N*-benzoyladenine (A^{Bz}), 4-*N*-benzoyl -5-methyl cytidine were synthesized by using the procedure described in the

references Takao Yamaguchi, Masahiko Horiba and Satoshi Obika; *Chem. Commun.* 2015, 51, 9737-9740, and Masahiko Horiba, Takao Yamaguchi, and Satoshi Obika; *Journal of Organic Chemistry*, 2016, 81, 11000-11008. The monomers were dried in a vacuum desiccator with desiccant (P_2O_5 , at room temperature for 24 hours). For the AmNA and scp-BNA modifications, the synthesis was carried out on a 1 μ M scale in a 3' to 5' direction with the phosphoramidite monomers diluted to a concentration of 0.12 M in anhydrous CH_3CN in the presence of 0.3 M 5-(benzylthio)-1H-tetrazole activator (coupling time 16-20 min) to a solid bound oligonucleotide followed by modified capping, oxidation and deprotection to afford the modified oligonucleotides. The stepwise coupling efficiency of all modified phosphoramidites was more than 97%. The DDTT (dimethylamino-methylidene) amino)-3H-1, 2, 4-dithiazaoline-3-thione was used as the sulfur-transfer agent for the synthesis of the oligoribonucleotide phosphorothioates. Oligonucleotide-bearing solid supports were washed with 20 % DEA solution in acetonitrile for 15 min then the column was washed thoroughly with AcCN. The support was heated at 65 °C with diisopropylamine:water:methanol (1:1:2) for 5 h in a heat block to cleave from the support and deprotect the base labile protecting groups. Table 13 summarizes the sequence length, alternating A and C units, and 5' modification for the resulting exemplified modified phosphorothioated oligonucleotides.

TABLE 13

No.	Length	A	C	5'-Modification
247	(AmAps(5m)AmC)20	AmNA(NMe)-A	AmNA(NMe)-(5m)C	5' OH, 40mer, All AmNA
248	(ScpAps(5m)scpC)20	Scp-BNA-A	Scp-BNA-(5m)C	5' OH, 40mer, All Scp-BNA
249	AmAps(5m)mC (AC)19	2'-OMe-A	2'-OMe-(5m)C	One AmNA at 5'-end, 40mer
250	(AC)19-mAps(5m)AmC	2'-OMe-A	2'-OMe-(5m)C	One AmNA at 3'-end, 40mer
251	ScpAps(5m)mC (AC)19	2'-OMe-A	2'-OMe-(5m)C	One ScpA at 5'-end, 40mer
252	(AC)19-mAps(5m)ScpC	2'-OMe-A	2'-OMe-(5m)C	One ScpC at 3'-end, 40mer

EXAMPLES 253-256

[0174] The effect of attaching a targeting ligand was evaluated by preparing a series of modified phosphorothioated oligonucleotides. The targeting ligands, cholesterol and a tocopherol (vitamin E), were attached to phosphorothioated oligonucleotides via an alkylene oxide linking group (tetraethylene glycol, TEG) in accordance with the methods described above in Examples 1-116 except that solid phase synthesis was conducted on cholesterol and tocopherol supports with attachment by a TEG linker for 3'-conjugation while final coupling of the phosphoramidite provided the 5'-conjugated oligonucleotides. FIGS. 3A-D and Table 14 illustrate the structures and summarize the sequence length, alternating A and C units, and targeting ligands for the resulting exemplified modified phosphorothioated oligonucleotides.

TABLE 14

No.	Length	A	C	Targeting Ligand
253	Chol-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-Cholesterol, 40mer
254	(AC)20- Chol	2'-OMe-A	2'-OMe-(5m)C	3'-Cholesterol, 40mer
255	Toco-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-Tocopherol, 40mer
256	(AC)20-Toco	2'-OMe-A	2'-OMe-(5m)C	3'-Tocopherol, 40mer

EXAMPLES 257-268

[0175] The effect of attaching a targeting ligand was evaluated by preparing a series of modified phosphorothioated oligonucleotides. N-acetylgalactosamine (GalNAc) was attached to phosphorothioated oligonucleotides via various linking groups by reacting with a GalNAc building block as illustrated in FIG. 4A. GalNAc-3 and GalNAc-5 amidites were purchased from AM Chemicals LLC and Glen Research respectively. GalNAc-4 and GalNAc-6 were obtained from AM Chemicals LLC. Table 15 illustrates the structures and summarizes the sequence length, alternating A and C units, and targeting ligands for the resulting exemplified modified phosphorothioated oligonucleotides.

TABLE 15

No.	Length	A	C	Targeting Ligand
257	GalNAc3ps-GalNAc3ps-GalNAc3po-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-GalNAc-3; 40mer
258	(AC)20-po-GalNAc3ps-GalNAc3ps-GalNAc3	2'-OMe-A	2'-OMe-(5m)C	3'-GalNAc-3; 40mer

No.	Length	A	C	Targeting Ligand
259	GalNAc3ps-GalNAc3ps-GalNAc3po-(AC)20	LNA-A	LNA-(5m)C	5'-GalNAc-3; 40mer
260	(AC)20-po-GalNAc3ps-GalNAc3ps-GalNAc3	LNA-A	LNA-(5m)C	3'-GalNAc-3; 40mer
261	GalNAc4ps-GalNAc4ps-GalNAc4po-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-GalNAc-4; 40mer
262	(AC)20-po-GalNAc4ps-GalNAc4ps-GalNAc4	2'-OMe-A	2'-OMe-(5m)C	3'-GalNAc-4; 40mer
263	GalNAc4ps-GalNAc4ps-GalNAc4po-(AC)20	LNA-A	LNA-(5m)C	5'-GalNAc-4; 40mer
264	(AC)20-po-GalNAc4ps-GalNAc4ps-GalNAc4	LNA-A	LNA-(5m)C	3'-GalNAc-4; 40mer
265	GalNAc5ps-GalNAc5ps-GalNAc5po-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-GalNAc-5; 40mer
266	(AC)20-po-GalNAc5ps-GalNAc5ps-GalNAc5	2'-OMe-A	2'-OMe-(5m)C	3'-GalNAc-5; 40mer
267	GalNAc5ps-GalNAc5ps-GalNAc5po-(AC)20	LNA-A	LNA-(5m)C	5'-GalNAc-5; 40mer
268	(AC)20-po-GalNAc5ps-GalNAc5ps-GalNAc5	LNA-A	LNA-(5m)C	3'-GalNAc-5; 40mer

EXAMPLES 269-272

[0176] The effect of attaching a targeting ligand was evaluated by preparing a series of modified phosphorothioated oligonucleotides. N-acetylgalactosamine (GalNAc) was attached to phosphorothioated oligonucleotides via a linking group by preparing the starting oligonucleotides, forming a precursor by attaching a C₆-NH₂ linking group at the 5'-terminal, and then reacting the precursor with a GalNAc ester. The sequences were synthesized at 10 μmol scale using universal support (Loading 65 μmol/g). The C₆-NH₂ linker was attached to the 5'-terminal to form the precursor by reacting with 6-(4-monomethoxytritylamino)hexyl-(2-cyanoethyl)-(N, N-diisopropyl)-phosphoramidite in 0.1 M acetonitrile was a coupling time of 10 min. The phosphorothioated oligonucleotide-bearing solid supports were heated at room temperature with aqueous ammonia/methylamine (1:1) solution for 3 h in a shaker to cleave from the support and deprotect the base labile protecting groups.

[0177] After IEX purification and desalting, the precursors were dissolved in 0.2 M sodium bicarbonate buffer, pH 8.5 (0.015 mM) and 5-7 mol equivalent of GalNAc ester

dissolved in DMSO was added. The structures of the GalNAc esters are illustrated in FIG. 4B. The reaction mixture was stirred at room temperature for 4 h. The sample was analyzed to confirm the absence of precursor. To this aqueous ammonia (28 wt. %) was added (5× reaction volume) and stirred at room temperature for 2-3 h. The reaction mixture was concentrated under reduced pressure and the resulting residue was dissolved in water and purified by HPLC on a strong anion exchange column.

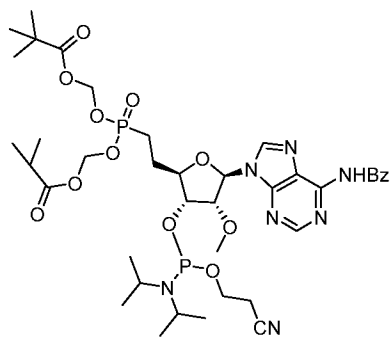
[0178] Table 16 illustrates the structures and summarizes the sequence length, alternating A and C units, and targeting ligands for the resulting exemplified modified phosphorothioated oligonucleotides. GalNAc-1 and GalNAc-2 were prepared in accordance with procedures described in *J. Med. Chem.* 2016 59(6) 2718-2733 and WO 2017/021385A1, respectively

TABLE 16

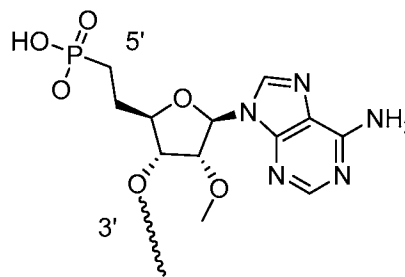
No.	Length	A	C	Targeting Ligand
269	GalNAc1-NH-C6-po-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-GalNAc-1; 40mer
270	GalNAc1-NH-C6-po-(AC)20	LNA-A	LNA-(5m)C	5'-GalNAc-1; 40mer
271	GalNAc2-NH-C6-po-(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-GalNAc-2; 40mer
272	GalNAc2-NH-C6-po-(AC)20	LNA-A	LNA-(5m)C	5'-GalNAc-2; 40mer

EXAMPLES 273-281

[0179] The effect of 5' modification was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above, except that the following 5'-ethyl phosphonate (EP) building block was utilized to incorporate 5'-ethyl phosphonate endcaps:



5'-Ethyl phosphonate (5'-EP) building block



5'-Ethyl phosphonate endcap

[0180] With reference to FIG. 5, the 5'-Ethyl phosphonate building block was prepared as follows:

[0181] To a mixture of **5-1** (3.0 g, 4.35 mmol, 1 eq) in MeOH (5 mL) was added Pd/C (900 mg, 72.50 μ mol, 10% purity) under N₂. The suspension was degassed under vacuum and purged with H₂ for several times. The mixture was stirred under H₂ (15 psi) at 20 °C for 12 hr. ¹H NMR and ³¹P NMR showed **5-1** was consumed completely to form desired product. The reaction mixture was filtered and concentrated to give [2-[(2R,3R,4R,5R)-5-(6-benzamidopurin-9-yl)-3-hydroxy-4-methoxy-tetrahydrofuran-2-yl]ethyl-(2,2-dimethylpropanoyloxymethoxy)phosphoryl]oxymethyl 2,2-dimethylpropanoate, compound **5-2**, (2.8 g, 4.05 mmol, 93.06% yield) as a white solid. ¹H NMR (400 MHz, CD₃OD) δ = 8.75 (s, 1H), 8.53 (s, 1H), 8.08 (d, *J*=7.5 Hz, 2H), 7.68 - 7.61 (m, 1H), 7.59 - 7.50 (m, 2H), 7.23 - 7.17 (m, 1H), 7.15 - 7.10 (m, 1H), 6.15 (d, *J*=4.2 Hz, 1H), 5.71 - 5.61 (m, 4H), 4.57 (t, *J*=4.7 Hz, 1H), 4.41 (t, *J*=5.3 Hz, 1H), 4.09 - 3.99 (m, 1H), 3.49 (s, 3H), 2.16 - 1.97 (m, 4H), 1.17 (d, *J*=3.5 Hz, 18 H); ³¹P NMR (162 MHz, CD₃CN) δ = 32.91 (s, 1P).

[0182] To a solution of **5-2** (2.3 g, 3.33 mmol, 1 eq) in DCM (30 mL) was added 1H-imidazole-4,5-dicarbonitrile (589.06 mg, 4.99 mmol, 1.5 eq) followed by 3-bis(diisopropylamino)phosphanyloxypropanenitrile (2.00 g, 6.65 mmol, 2.11 mL, 2.0 eq), and the mixture was stirred at 25 °C for 2 hr. TLC indicated that majority of **5-2** was consumed and one major new spot was formed. The reaction mixture was washed with H₂O (50 mL*2) and brine (50 mL*2), dried over Na₂SO₄, and concentrated to give a residue. The residue was purified by Flash-C-18 column using the following conditions: Column, C18 silica gel; mobile phase, water and acetonitrile (0%-70% acetonitrile) to give [2-[(2R,3R,4R,5R)-5-(6-benzamidopurin-9-yl)-3-[2-cyanoethoxy-(diisopropylamino)phosphanyl]oxy-4-methoxy-tetrahydrofuran-2-yl]ethyl-(2,2-dimethylpropanoyloxymethoxy)phosphoryl]oxymethyl 2,2-dimethylpropanoate, (**5'-EP** building block), (1.4 g, 1.53 mmol, 45.88% yield, 97.2% purity) as a light yellow solid. LCMS (ESI): RT = 3.785 min, *m/z* calcd. for C₄₀H₆₀N₇O₁₂P₂ 892.37 [M+H]⁺, found 892.0. HPLC: Mobile Phase: 10mMol NH₄Ac in water (solvent C) and acetonitrile (solvent D), using the elution gradient 80%-100% (solvent D) over 10 minutes and holding at 100% for 5 minutes at a flow rate of 1.0 mL/minute; Column30: Waters Xbridge C18 3.5 μ m, 150*4.6mm; ¹H NMR (400MHz, CD₃CN) δ = 9.40 (s, 1H), 8.67 (s, 1H), 8.27 (d, *J*=1.8 Hz, 1H), 8.01 (d, *J*=7.5 Hz, 2H), 7.68 - 7.60 (m, 1H), 7.58 - 7.52 (m, 2H), 6.05 (dd,

$J=5.1, 8.4$ Hz, 1H), 5.62 - 5.54 (m, 4H), 4.68 (t, $J=1.8, 5.0$ Hz, 1H), 4.64 - 4.55 (m, 1H), 4.25 - 4.11 (m, 1H), 3.93 - 3.66 (m, 4H), 3.40 (d, $J=19.2$ Hz, 3H), 2.75 - 2.67 (m, 2H), 2.14 - 1.95 (m, 4H), 1.25 - 1.20 (m, 12H), 1.15 - 1.11 (m, 18H); ^{31}P NMR (162MHz, CD_3CN) $\delta = 149.95, 149.27, 32.29, 32.05$.

[0183] Table 17 summarizes the sequence length, alternating A and C units, the number and type (R or S) of stereochemically defined phosphorothioate (PS) linkages and LNA modification for the resulting exemplified 5'-EP endcapped modified phosphorothioated oligonucleotides.

TABLE 17

No.	Length	A	C	PS Modification	Comments
273	(AC)20	2'-OMe-A	2'-OMe-(5m)C	PS	40mer
274	(AC)20	LNA-A	LNA-(5m)C	PS	40mer
275	(AC)20	2'-OMe-A	2'-OMe-(5m)C	20; 2'-OMeAps R (5m)InC	20 R isomer, 41mer
276	(AC)20	2'-OMe-A	2'-OMe-(5m)C	19; 2'-OMeAps R (5m)InC	19 R isomer, 40mer
277	(AC)20	2'-OMe-A	LNA-(5m)C	PS	40mer Alternate 2'-OMe/LNA
278	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	PS	Every 3rd base is LNA
279	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	PS	Every 4th base is LNA
280	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	PS	5 LNA in the middle
281	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	PS	10 LNA in the middle

EXAMPLES 282-298

[0184] FIG. 6A describes compound nos. 282-295, which were prepared in accordance with the methods described above.

EXAMPLES 296-304

[0185] The effect of sequence length, LNA incorporation, and RNA incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 18.

TABLE 18

No.	Length	A	C	RNA Modification
296	(AC)20	2'-OMe-A	LNA-(5m)C	5 RNA
297	(AC)20	2'-OMe-A	2'-OMe-(5m)C	7 RNA
298	(AC)20	2'-OMe-A	2'-OMe-(5m)C	14 RNA
299	(AC)15	2'-OMe-A	2'-OMe-(5m)C	5 RNA
300	(AC)15	2'-OMe-A	2'-OMe-(5m)C	10 RNA
301	(AC)20	LNA-A	LNA-(5m)C	7 RNA
302	(AC)20	LNA-A	LNA-(5m)C	14 RNA
303	(AC)15	LNA-A	LNA-(5m)C	5 RNA
304	(AC)15	LNA-A	LNA-(5m)C	10 RNA

EXAMPLES 305-313

[0186] The effect of sequence length, LNA incorporation, and backbone was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 19.

TABLE 19

No.	Length	A	C	Backbone
305	(AC)20	LNA-A	LNA-(5m)C	40mer; 20 PO; 19 PS
306	(AC)20	LNA-A	LNA-(5m)C	40mer; 7 PO; 32 PS
307	(AC)20	LNA-A	LNA-(5m)C	40mer; 14 PO; 25 PS
308	(AC)15	LNA-A	LNA-(5m)C	30mer; 5 PO; 24 PS
309	(AC)15	LNA-A	LNA-(5m)C	30mer; 10 PO; 19 PS
310	(AC)20	2'-OMe-A	2'-OMe-(5m)C	40mer; 7 PO; 32 PS
311	(AC)20	2'-OMe-A	2'-OMe-(5m)C	40mer; 14 PO; 25 PS
312	(AC)15	2'-OMe-A	2'-OMe-(5m)C	30mer; 5 PO; 24 PS
313	(AC)15	2'-OMe-A	2'-OMe-(5m)C	30mer; 10 PO; 19 PS

EXAMPLES 314-322

[0187] The effect of sequence length, LNA incorporation, and ethyl phosphonate endcap was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 20.

TABLE 20

No.	Length	A	C	Modification
314	(AC)20	2'-OMe-A	LNA-(5m)C	Ethyl-phosphonate-A

315	(AC)20	2'-OMe-A	2'-OMe-(5m)C	19 R dimer block; Ethyl-phosphonate-A
316	(AC)20	2'-OMe-A	2'-OMe-(5m)C	5 LNA, Ethyl-phosphonate-A
317	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	40mer; Every 4 th base is LNA Ethyl-phosphonate-A
318	(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	40mer; Every 3rd base is LNA Ethyl-phosphonate-A
319	(AC)20	2'-OMe-A	2'-OMe-(5m)C	40mer; Ethyl-phosphonate-A
320	(AC)18	2'-OMe-A	LNA-(5m)C	36mer; Alternate 2'-OMe and LNA
321	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	36mer; Every 3rd base is LNA
322	(AC)20	2'-OMe-A	LNA-(5m)C	36mer; Every 4 th base is LNA

EXAMPLES 323-324

[0188] The effect of LNA incorporation and phosphate endcap was evaluated by preparing phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 21.

TABLE 21

No.	Length	A	C	Endcap
323	(AC)20	LNA-A	LNA-(5m)C	5'-Phosphate
324	(AC)20	2'-OMe-A	2'-OMe-(5m)C	5'-Phosphate

EXAMPLES 325-338

[0189] The effect of level of LNA incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 22.

TABLE 22

No.	Length	A	C	Modification
325	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	40mer; 75% 2'-OMe, 25% LNA
326	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	40mer; 67.5% 2'-OMe 37.5% LNA
327	(AC)20	2'-O-MOE-A LNA-A	2'-O-MOE-(5m)C LNA-(5m)C	40mer; 75% 2'-O-MOE, 25% LNA

No.	Length	A	C	Modification
328	(AC)20	2'-O-MOE-A LNA-A	2'-O-MOE-(5m)C LNA-(5m)C	40mer; 67.5% 2'-O-MOE 37.5% LNA
329	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	40mer; 75% LNA, 25% 2'-OMe (10mer block)
330	(AC)20	2'-OMe-A LNA-A	2'-OMe-(5m)C LNA-(5m)C	40mer; 50% LNA; 50% 2'-OMe(20mer block)
331	(AC)20	2'-O-MOE-A LNA-A	2'-O-MOE-(5m)C LNA-(5m)C	40mer; 75% LNA, 25% 2'-O-MOE (10mer block)
332	(AC)20	2'-O-MOE-A LNA-A	2'-O-MOE-(5m)C LNA-(5m)C	40mer; 50% LNA; 50% 2'-O-MOE (20mer block)
333	(AC)20	LNA-A DNA-A	LNA-(5m)C DNA-(5m)C	40mer; 7 DNA
334	(AC)20	LNA-A DNA-A	LNA-(5m)C DNA-(5m)C	40mer; 14 DNA
335	(AC)20	LNA-A DNA-A	LNA-(5m)C DNA-(5m)C	30mer; 5 DNA
336	(AC)20	LNA-A DNA-A	LNA-(5m)C DNA-(5m)C	30mer; 10 DNA
337	(AC)20	LNA-A DNA-A	LNA-(5m)C DNA-(5m)C	40mer; 50% LNA; 50% DNA (10mer DNA block)
338	(AC)20	LNA-A DNA-A	LNA-(5m)C DNA-(5m)C	40mer; 50% LNA; 50% DNA (20mer DNA block)

EXAMPLES 339-340

[0190] The effect of ScpA and AmNA incorporation was evaluated by preparing phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 23.

TABLE 23

No.	Length	A	C	Modification
339	(AC)20	2'-OMe-A	LNA-(5m)C	One ScpA at 3'-end, 40mer
340	(AC)20	2'-OMe-A	LNA-(5m)C	One AmNA at 3'-end, 40mer

EXAMPLES 341-346

[0191] The effect of GNA and UNA incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 24.

TABLE 24

No.	Length	A	C
341	(AC)20	LNA-A	GNA-(5m)C
342	(AC)20	GNA-A	2'-OMe-(5m)C
343	(AC)20	2'-OMe-A	GNA-(5m)C
345	(AC)20	UNA-A	UNA-(5m)C
346	(AC)20	UNA-A	UNA-(5m)C

EXAMPLES 347-350

[0192] The effect of attaching a targeting ligand was evaluated by preparing a series of modified phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 25.

TABLE 25

No.	Length	A	C	Modification
347	GalNAc5ps-GalNAc5ps-GalNAc5po-(AC)20	2'-OMe-A	LNA-(5m)C	40mer, alternate 2'-OMe-LNA; 5'-GalNac
348	GalNAc5ps-GalNAc5ps-GalNAc5po-(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-(5m)C	40mer, every 4 th base is LNA; 5'-GalNac
349	GalNAc5ps-GalNAc5ps-GalNAc5po-(AC)20	2'-OMe-A	2'-OMe-(5m)C LNA-A	40mer, 5 LNA; 5'-GalNac
350	GalNAc5ps-GalNAc5ps-GalNAc5po-(AC)20	2'-OMe-A	LNA-(5m)C	40mer, alternate 2'-OMe-LNA 5 RNA; 5'-GalNac

EXAMPLES 351-355

[0193] The effect of attaching a cholesterol or tocopherol targeting ligand was evaluated by preparing a series of modified phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 26.

TABLE 26

No.	Length	A	C	Targeting Ligand
351	Chol-(AC)20	2'-OMe-A	(5m)-Propargyl-C	3'-Cholesterol, 40mer
352	(AC)20- Chol	2'-OMe-A	(5m)-Propargyl-C	3'-Palmitoyl, 40mer
353	(AC)20	3'-OMe-A	3'-OMe-(5m)C	3'-OMe, 40mer
354	(AC)20- Chol	3'-OMe-A	3'-OMe-(5m)C	3'-cholesterol, 40mer

No.	Length	A	C	Targeting Ligand
355	(AC)20- Toco	3'-OMe-A	3'-OMe-(5m)C	3'-Tocopherol, 40mer

EXAMPLES 356-358

[0194] The effect of endcap structure (methyl, allyl, cytosine) was evaluated by preparing phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 27.

TABLE 27

No.	Length	A	C	Endcap
356	(AC)20	2'-OMe-A	LNA-(5m)C	40mer, 4'-Me at 5'-end
357	(AC)20	2'-OMe-A 3'-C-allyl-A	LNA-A LNA-(5m)C	40mer, 5 3'-C-allyl-A
358	(AC)20	LNA-A	LNA-(5m)C	40mer, Cy-5 at 3'-end

EXAMPLES 359-362

[0195] The effect of including G and U bases was evaluated by preparing phosphorothioated oligonucleotides in accordance with the methods described above. The compounds are summarized in Table 28.

TABLE 28

No.	Length	Base 1	Base 2	Modification
359	(AG)20	2'-OMe-A	2'-OMe-G	AG repeat
360	(GA)20	2'-OMe-G	2'-OMe-A	GA repeat
361	(CA)20	2'-OMe-(5m)C	2'-OMe-A	CA repeat
362	(AU)20	2'-OMe-A	2'-OMe-U	AU repeat

EXAMPLES 363-376

[0196] The effect of sequence length was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The compounds are summarized in Table 29.

TABLE 29

No.	Length	A	C	Modification
363	(AC)14	2'-OMe-A	2'-OMe-C	28mer
364	(AC)15-A	2'-OMe-A	2'-OMe-(5m)C	31mer

No.	Length	A	C	Modification
365	(AC)17	2'-OMe-A	2'-OMe-(5m)C	34mer
366	(AC)18-A	2'-OMe-A	2'-OMe-(5m)C	37mer
367	(AC)20	2'-OMe-A	2'-OMe-C	20mer
368	(AC)9	2'-OMe-A	2'-OMe-(5m)C	18mer
369	(AC)9-A	2'-OMe-A	2'-OMe-(5m)C	19mer
370	(AC)10	2'-OMe-A	2'-OMe-(5m)C	20mer
371	(AC)9-A	LNA-A	LNA-(5m)C	19mer
372	(AC)9	LNA-A	LNA-(5m)C	18mer
373	(AC)15	LNA-A	LNA-(5m)C	30mer
374	(AC)12-A	2'-OMe-A	2'-OMe-(5m)C	25mer
375	(AC)20	2'-OMe-A	2'-OMe-(5m)C	40mer, 5 S isomers
376	(AC)10	LNA-A	LNA-(5m)C	20 mer

EXAMPLES 377-380 AND 384

[0197] The effect of RNA incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 30.

TABLE 30

No.	Length	A	C	Modification
377	(AC)20	2'-OMe-A Ribo-A	LNA-(5m)C	40mer, 4 RNA
378	(AC)20	2'-OMe-A Ribo-A	LNA-(5m)C	40mer, 3 RNA
379	(AC)20	2'-OMe-A Ribo-A	LNA-(5m)C	40mer, 2 RNA
380	(AC)20	2'-OMe-A UNA-A	2'-OMe-(5m)C UNA-(5m)C	40mer, 4mer blocks of 2'-OMe and UNA
384	(AC)20	2'-OMe-A Ribo-A	LNA-(5m)C	40mer, 1 RNA

EXAMPLES 381-383

[0198] The effect of 4etl (4-ethyl-LNA) incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The 4etl monomers were prepared in accordance with known methods (Seth, P.P., J. Org. Chem. 2010, 75, (5), 1569–1581). The results are summarized in Table 31.

TABLE 31

No.	Length	A	C	Modification
381	(AC)20	4etl-A	4etl-(5m)C	40mer, 100% 4etl
382	(AC)20	2'-OMe-A	4etl-(5m)C	40mer, 50% 4etl
383	(AC)20	2'-OMe-A	2'-OMe-(5m)C 4etl-(5m)C	40mer, 25% 4etl

EXAMPLES 385-389

[0199] The effect of nmLNA (N-methyl LNA) A and C incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The nmLNA monomers were obtained from commercial sources (Bio-Synthesis Inc., Lewisville, TX). The results are summarized in Table 32.

TABLE 32

No.	Length	A	C	Modification
385	(AC)20	2'-OMe-A nmLNA-A	LNA-(5m)C	40mer, 1 nmLNA
386	(AC)20	2'-OMe-A nmLNA-A	LNA-(5m)C	40mer, 3 nmLNA
387	(AC)20	2'-OMe-A nmLNA-A	LNA-(5m)C nmLNA (5m)-C	40mer, 3 nmLNA
388	(AC)20	2'-OMe-A	LNA-(5m)C nmLNA (5m)-C	40mer, 3 nmLNA
389	(AC)20	2'-OMe-A nmLNA-A	LNA-(5m)C nmLNA (5m)-C	40mer, 4 nmLNA

EXAMPLES 390-392

[0200] The effect of AmNA and Scp-BNA A and C incorporation was evaluated by preparing a series of phosphorothioated oligonucleotides in accordance with the methods described above. The results are summarized in Table 33 (also see Table 23).

TABLE 33

No.	Length	A	C	Modification
390	(AC)20	2'-OMe-A	AmNA-(5m)C	40mer, 20 AmNA(50%)

No.	Length	A	C	Modification
391	(AC)20	2'-OMe-A	2'-OMe-(5m)C Scp-(5m)C	40mer, 10 scp-BNA (25%)
392	(AC)20	2'-OMe-A Scp-A	2'-OMe-(5m)C	40mer, 5 scp-BNA (12.5%)

EXAMPLE B1

HBSAG SECRETION ASSAY AND CYTOTOXICITY ASSAY

[0201] The sequence independent antiviral activity against hepatitis B (as determined by HBsAg Secretion Assay) and the cytotoxicity of a number of exemplified modified oligonucleotide compounds was determined as described below and summarized in Tables 34-35 and FIGS. 6A and 6B.

HBsAg Release Assay ProtocolCell Culture

[0202] HepG2.2.15 cells were maintained in DMEM medium with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin, 1% Glutamine, 1% non-essential amino acids, 1% Sodium Pyruvate and 380 ug/ml G418. Cells were maintained at 37°C in a 5% CO₂ atmosphere.

HBsAg Secretion Assay

[0203] HepG2.2.15 cells were grown in DMEM medium as described above. Cells were plated at a concentration of 45,000 cells/well in collagen-I coated 96 well plates. Immediately after addition of the cells, test compounds are added.

[0204] Selected compounds may also be tested following Lipofectamine® RNAiMAX transfection. Lipofectamine® RNAiMAX Transfection Reagent (Thermo Fisher) is used following the manufacturer's instructions.

[0205] The 50% inhibitory concentration (EC₅₀) and 50% cytotoxic concentration (CC₅₀; below) were assessed by solubilizing in 1 X PBS to 100 X the desired final testing concentration. Each compound was then serially diluted (1:3) up to 8 distinct concentrations to 10X the desired final testing concentration in DMEM medium with 10% FBS. A 10 µL sample of the 10X compounds in cell culture media was used to treat the HepG2.2.15 cells in a 96-well format. Cells were initially incubated with compounds for 3 days at 37°C in a 5% CO₂ atmosphere.

[0206] Three days post compound addition/transfection replace media and compound with fresh media/compound with RNAiMax and incubate for a further 3 days for a total incubation time of 6 days. Collect both the cellular supernatant and cell lysate (see below) for quantification of HBsAg.

[0207] Secreted HBsAg was measured quantitatively using HBsAg ELISA kit (Autobio-CL0310).

[0208] The EC₅₀, the concentration of the drug required for reducing HBsAg secretion by 50% in relation to the untreated cell control value was calculated from the plot of the percentage reduction of the HBsAg level against the drug concentrations using Microsoft Excel (forecast function).

[0209] Set up a parallel set of plates that are to be used for testing compound induced cellular cytotoxicity (see below).

Cytotoxicity Assay

[0210] HepG2.2.15 cells were cultured and treated as above. At Day 6, cellular cytotoxicity was assessed using a cell proliferation assay (CellTiter-Glo Luminescent Cell Viability Assay; Promega) according to the manufacturer's instructions or a suitable alternative.

[0211] The CC₅₀, the concentration of the drug required for reducing cell viability by 50% in relation to the untreated cell control value was calculated from the plot of the percentage reduction of viable cells against the drug concentrations using Microsoft Excel (forecast function).

TABLE 34 – POTENCY AND CYTOTOXICITY

Compound No.	EC ₅₀ (μM)	CC ₅₀ (μM)
3	B	A
6	A	B
8	B	A
9	A	A
10	A	A
12	A	A
13	B	A
18	C	A
20	B	B
23	B	B
26	C	A
34	B	A

Compound No.	EC ₅₀ (μM)	CC ₅₀ (μM)
36	B	A
38	A	A
39	B	C
44	A	A
45	A	A
63	B	A
97	B	A
98	B	A
99	B	A
105	B	A
106	B	A
120	C	A
121	B	A
122	B	A
127	B	A
128	D	A
129	D	A
130	B	A
134	A	A
142	C	A
147	D	A
148	D	A
149	B	A
150	A	A
151	D	A
152	D	A
153	B	A
158	B	A
159	C	A
178	A	A
179	A	A
180	A	A
182	A	A
183	A	A
184	A	A
190	B	A
191	B	A
192	A	A
199	B	A
200	C	A
201	B	A
202	B	A
204	B	A
205	B	A

Compound No.	EC ₅₀ (μM)	CC ₅₀ (μM)
220	C	A
221	A	A
223	C	A
235	D	B
236	D	B
237	A	B
238	D	A
239	D	A
240	B	A
241	B	A
242	A	A
243	A	A
244	C	A
245	D	A

[0212] Potency: **A**: $\geq$ 5-fold higher than (2'-OMe-A; 2'-OMe-C); **B**: $\geq$ 2-fold higher than (2'-OMe-A; 2'-OMe-C) and $<$ 5-fold higher than (2'-OMe-A; 2'-OMe-C); **C**: higher than or equal to (2'-OMe-A; 2'-OMe-C) and $<$ 2-fold higher than (2'-OMe-A; 2'-OMe-C); **D**: lower than (2'-OMe-A; 2'-OMe-C).

[0213] Cytotoxicity: **A**: $\geq$ 2 μM; **B**: $<$ 2 μM

TABLE 35 – POTENCY AND CYTOTOXICITY

Compound No. ¹	EC ₅₀	CC ₅₀
6, 274, 283	A	B
376	D	A
371	D	A
372	D	A
273, 282	D	A
367	C	A
368	D	A
369	D	A
370	D	A
345	B	A
346	A	A
351	D	B
352	D	B
373	B	B
308	C	A
239	D	A
235	D	B
236	D	B

Compound No. ¹	EC ₅₀	CC ₅₀
237	A	B
301	A	B
303	B	B
305	C	A
315	C	A
309	D	B
297	C	A
298	D	A
300	D	A
312	D	A
313	D	A
299	D	A
304	D	A
302	D	A
307	D	A
375	B	A
201	C	A
202	C	A
203	B	A
204	D	A
205	D	A
353	B	A
351	D	A
352	D	A
178	A	A
179	A	A
180	C	A
182	A	A
183	D	A
184, 290	A	A
177	B	A
374	D	A
363	D	A
364	D	A
365	D	A
366	D	A
238	D	A
240	B	A
241	B	A
242	A	A

Compound No. ¹	EC ₅₀	CC ₅₀
243	A	A
130	A	A
380	D	A
310	D	A
311	D	A
254	D	A
325	D	A
326	D	A
327	D	A
328	D	A
158	B	A
150	A	A
159	C	A
341	D	A
342	B	A
244	C	A
245	C	A
343	B	A
329	C	A
330	B	B
331	D	A
332	D	A
333	B	A
334	B	A
335	C	A
336	C	A
337	A	B
338	B	B
117	B	A
118	B	A
134, 277, 284	A	A
142	C	A
190	B	A
191	B	A
192	B	A
210	B	A
211	B	A
212	B	A
218	C	A
223	C	A

Compound No. ¹	EC ₅₀	CC ₅₀
221	A	A
127	D	A
128	C	A
129	C	A
120	B	A
121	B	A
122	A	A
181	C	A
147	D	A
148	D	A
149	B	A
151	D	A
152	D	A
153	B	A
294	B	A
276, 291	A	A
275, 295	A	B
173, 293	A	A
165, 287	B	A
167, 289	B	A
164, 286	C	A
166, 288	A	A
171, 280, 292	B	A
314	A	B
281, 316	A	A
296	A	A
285	A	B
251	A	A
356	A	A
320	A	A
321	A	B
322	B	A
317	A	B
318	B	B
319	A	B
357	A	A
339	A	A
252	A	A
340	A	A
250	A	A

Compound No. ¹	EC ₅₀	CC ₅₀
359	D	A
360	D	A
361	D	A
362	D	A
12	A	A
20	B	A
38	A	A
385	A	A
386	A	A
387	A	A
388	A	A
389	A	A
376	A	A
377	A	A
378	A	A
379	A	A
384	A	A
381	A	A
382	A	A
383	B	A
390	A	B
391	A	B
392	B	B

[0214] ¹ A number of compounds described herein are referred to by more than a single compound no. as indicated here and elsewhere throughout the disclosure.

[0215] Potency: A: EC₅₀ < 30 nM; B: EC₅₀ ≥ 30 nM and EC₅₀ < 100 nM; C: EC₅₀ ≥ 100 nM and EC₅₀ < 300 nM; D: EC₅₀ > 300 nM.

[0216] Cytotoxicity: A: CC₅₀ ≥ 1000 nM; B: CC₅₀ < 1000 nM

EXAMPLE B2

LIVE INFECTION ASSAY

[0217] HepG2-NTCP cells were maintained in DMEM/F12 medium with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin, 1% Glutamine, 1% non-essential amino acids, 1% Sodium Pyruvate. Cells were maintained at 37°C in a 5% CO₂ atmosphere.

[0218] HepG2-NTCP cells were resuspended with above mentioned medium and plated at a concentration of 15,000 cells/well in collagen-I coated 96 well plates. On the second day (day 0), the cells were infected with HBV (purified HBV from HepAD38 cells) at 200 moi (ge) in the presence of 4% PEG8000 and 2% DMSO and incubated at 37°C overnight. The inoculum was vacuumed and cells were washed three times with DMEM/F12 with 2% FBS before replacing with the HepG2-NTCP culture medium.

[0219] Treat the cells on day 5. On Day 5, the test compounds were diluted 3-fold with Opti-MEM I media and mixed with Lipofectamine® RNAiMAX transfection reagent following the manufacturer's instructions. After media replacement on Day 8, the test compounds were transfected as described. After incubation for an additional 3 days, the supernatant was harvested and HBsAg was measured by ELISA (Diasino). The cell viability was measured with CellTiter-Glo (Promega).

[0220] The EC₅₀, the concentration of the drug required for reducing HBsAg secretion by 50% in relation to the untreated cell control value, was calculated from the plot of the percent reduction of the HBsAg level against the drug concentrations using the Microsoft Excel forecast function or GraphPad Prism and summarized in Table 36.

TABLE 36 – POTENCY AND CYTOTOXICITY

Compound No.	EC ₅₀	CC ₅₀
6, 274, 283	A	A
273, 282	C	A
315	D	A
290 184	A	A
134, 277, 284	A	A
192	A	A
221	A	A
294	C	A
291 276	A	A
295 275	B	A
173, 293	B	A
165, 287	A	A
167, 289	B	A
164, 286	B	A

Compound No.	EC ₅₀	CC ₅₀
166, 288	B	A
171, 280, 292	B	A
314	A	A
281, 316	C	A
296	A	A
285	A	A
251	B	A
356	A	A
320	A	A

[0221] Potency: A: EC₅₀ < 30 nM; B: EC₅₀ ≥ 30 nM and EC₅₀ < 100 nM; C: EC₅₀ ≥ 100 nM and EC₅₀ < 300 nM; D: EC₅₀ > 300 nM.

[0222] Cytotoxicity: A: CC₅₀ ≥ 1000 nM; B: CC₅₀ < 1000 nM

EXAMPLE B3

HBSAG SECRETION ASSAY FOR COMBINATIONS

[0223] The sequence independent antiviral activity against hepatitis B (as determined by HBsAg Secretion Assay) of exemplified modified oligonucleotide compounds in combination with antisense oligonucleotides (ASOs) was determined as described below and summarized in Table 37.

Cell Culture

[0224] HepG2.2.15 cells were maintained in DMEM/F12 medium with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin, 1% Glutamine, 1% non-essential amino acids, 1% Sodium Pyruvate. Cells were maintained at 37°C in a 5% CO₂ atmosphere.

HBsAg Secretion Assay

[0225] HepG2.2.15 cells were grown in DMEM/F12 medium as described above. Cells were seeded at a concentration of 35,000 cells/well in collagen-I coated 96-well plates. Immediately after addition of the cells, add test compounds. Do double transfections on day 0 and 3.

Transfection method

[0226] Lipofectamine® RNAiMAX transfection. Lipofectamine® RNAiMAX Transfection Reagent (Thermo Fisher, cat#: 13778-150) is used following the manufacturer's instructions.

[0227] A: mix RNAiMAX (0.3ul/well for 96-well plate) with Opti-MEM I (make 20% extra), incubate for 5 min at RT.

[0228] B: dilute combinations of ASOs and modified oligonucleotides in Opti-MEM I to make 40x of final concentration (8-point, 3-fold dilution, include concentration 0nM). The top concentration is about 100 – 200 folds of EC₅₀ value. Then mix equal volume dilutions from both compound1 and compound2 at opposite direction as indicated in the graph shown in FIG. 23.

[0229] Mix A and B at equal volume (make 20% extra volume), incubate another 5-10 min. Then add mixture of A and B at 1/10 of the final culture volume to each well, swirl the plates for 10 seconds by hand. There should be at least triplicate for the plates. Incubate at 37°C for 3 days, refresh medium, repeat the transfection process. On day 6 upon treatment, harvest supernatant for ELISA assay, measure cell viability with CellTiter-Glo (Promega).

Data analysis

[0230] To analyze the synergism, the percentage of HBsAg (or DNA) reduction is calculated for each well. Percentage of reduction = (1-well/average of no drug control)*100. These numbers are input to MacSynergy II software and the synergism/antagonism volume is obtained following the instruction of the software.

[0231] Synergy volume <25 indicates no synergism/antagonism.

[0232] Synergy volume 25-50 indicates minor synergism/antagonism.

[0233] Synergy volume 50-100 indicates moderate synergism/antagonism.

[0234] Synergy volume >100 indicates strong synergism/antagonism.

[0235] Synergy volume >1,000 indicates possible errors, check the data.

[0236] Percentage of cell viability = (well/average of no drug control)*100.

Monitor cytotoxicity as previously described.

HBsAg Quantification

[0237] Secreted HBsAg was measured quantitatively using HBsAg ELISA kit (Autobio-CL0310). Synergy values for combinations of modified oligonucleotides with ASOs are provided in Table 37.

TABLE 37 – SYNERGY OF COMBINATIONS

Compound No.	ASO ¹	HBsAg 95% Synergy Volume
166, 288	ASO-1	335.08

Compound No.	ASO ¹	HBsAg 95% Synergy Volume
134, 277, 284	ASO-2	52.98
296	ASO-2	43.05

¹ASO-1 is an unconjugated HBV ASO SSO-1 as disclosed in in Javanbakht, H. et al.

Molecular Therapy: Nucleic Acids Vol. 11 June 2018, having the following structure: 5'-InApsInGpsIn(5m)CpsGpsApsApsGpsTpsGps(5m)CpsAps(5m)CpsApsIn(5m)CpsInGpsInG-3. ASO-2 is an ASO having a structure as described for the ASO referred to as Sequence #9 in U.S. application serial number 62/855,793, which is hereby incorporated herein by reference and particularly for the purpose of describing the structure of the Sequence #9.

EXAMPLE B4

HBSAG SECRETION ASSAY FOR COMBINATIONS

[0238] The sequence independent antiviral activity against hepatitis B (as determined by HBsAg Secretion Assay) of exemplified modified oligonucleotide compounds in combination with an ASO, capsid assembly modulators (CAM compound 1 or CAM compound 2), or nucleoside analog (Entecavir, ETV) was determined as described below and summarized in Table 38.

Cell Culture

[0239] The following assay procedure describes the HBV antiviral assay. This assay uses HepG2.2.15 cells, which have been transfected with HBV genome, and extracellular HBV DNA quantification as endpoint. Cell viability is assessed in parallel by measuring the intracellular ATP content using the CellTiter-Glo[®] reagent from Promega.

HBsAg Secretion Assay

[0240] HepG2.2.15 cells were grown in DMEM/F12 medium as described above. Cells were seeded at a concentration of 35,000 cells/well in collagen-I coated 96-well plates. Immediately after addition of the cells, add test compounds. Do double transfections on day 0 and 3.

HBV DNA quantification assay

[0241] Extracellular DNA was isolated with QIAamp 96 DNA Blood Kit per the manufacturer's manual. HBV DNA was then quantified by qPCR with HBV specific primers and probes as specified below using the FastStart Universal MasterMix from Roche on an ABI-

7900HT. The PCR cycle program consisted of 95°C for 10 min, followed by 40 cycles at 95°C for 15 sec and 60°C for 1 min.

Items	Name	Sequence (5' → 3')
HBV Primer	HBV-forward	GTGTCTGCGGCGTTTATCA
	HBV-reverse	GACAAACGGGCAACATACCTT
HBV Probe	HBV probe	FAM-CCTCTKCATCCTGCTGCTATGCCTCATC- TAMRA

Transfection method

[0242] Lipofectamine® RNAiMAX transfection. Lipofectamine® RNAiMAX Transfection Reagent (Thermo Fisher, cat#: 13778-150) is used following the manufacturer's instructions.

[0243] A: mix RNAiMAX (0.3ul/well for 96-well plate) with Opti-MEM I (make 20% extra), incubate for 5 min at RT

[0244] B: dilute combinations of a CAM, ASO or ETV with modified oligonucleotides in Opti-MEM I to make 40x of final concentration (8-point, 3-fold dilution, include concentration 0nM). The top concentration is about 100 – 200 folds of EC₅₀ value. Then mix equal volume dilutions from both compound1 and compound2 at opposite direction as indicated in the graph shown in FIG. 23.

[0245] Mix A and B at equal volume (make 20% extra volume), incubate another 5-10 min. Then add mixture of A and B at 1/10 of the final culture volume to each well, swirl the plates for 10 seconds by hands. There should be at least triplicate for the plates. Incubate at 37°C for 4-hrs before adding the ASO, ETV or CAMs to let the cells recover from tranfection. On day 3 upon treatment, harvest supernatant for ELISA assay, measure cell viability with CellTiter-Glo (Promega).

Data analysis

[0246] The synergism data was analyzed as described in Example B3 above.

HBsAg Quantification

[0247] Secreted HBsAg was measured quantitatively using HBsAg ELISA kit (Autobio-CL0310). Synergy values for combinations of modified oligonucleotides with ASOs are provided in Table 38.

TABLE 38 – SYNERGY OF COMBINATIONS

Compound No.	ASO, CAM or ETV ¹	HBV DNA	HBV DNA 95% Synergy Volume
166, 288	ASO-1	Additive	23.99
134, 277, 284	ETV	Synergy	25.91
134, 277, 284	CAM compound 1	Additive	1.35
134, 277, 284	CAM compound 2	Synergy	41.86

¹CAM compound 1 is a CAM having a structure as described for the CAM compound referred to as compound 3 in WO2017/181141, which is hereby incorporated herein by reference and particularly for the purpose of describing the structure of the compound 3. CAM compound 2 is a CAM having a structure as described for the CAM compound referred to as compound 1 in U.S. serial no. 62/805,725, which is hereby incorporated herein by reference and particularly for the purpose of describing the structure of the compound 1. ASO-1 is as described above for Table 37.

EXAMPLE B5

LIVER EXPOSURE IN NON-HUMAN PRIMATES

[0248] Terminal liver exposures in non-human primates were evaluated by dosing exemplified modified oligonucleotide compounds to female cynomolgus monkeys by either the intravenous (IV) or subcutaneous (SC) route. For the IV route, the compound was administered in sterile phosphate-buffered saline (PBS) vehicle and infused over a 2-hr period at 1 mL/kg. For subcutaneous dosing, the vehicle was also sterile PBS and the compound was administered as a single bolus at 1 mL/kg. There were two animals per dose group, and the data shown is the average of the two animals. Liver levels were determined at the 24-hour timepoint. The doses utilized for this study and the data obtained is shown in FIG. 12. Unexpectedly, liver exposure following subcutaneous administration to non-human primates is much higher than expected based on liver exposure levels resulting from otherwise comparable intravenous dosing.

EXAMPLE B6

PBMC ASSAY

[0249] The effect of exemplified modified oligonucleotide compounds on the release of cytokines from peripheral blood mononuclear cells (PBMC) was evaluated as described below and summarized in Table 39 and FIGS. 13-22.

[0250] Buffy coats (N=3) were obtained from Stanford Blood Center and processed to isolate PBMC as per Aragen's standard protocol using Ficoll density gradient centrifugation. PBMC (1 million/mL) were suspended in complete culture (RPMI supplemented with 10% heat inactivated-low IgG FBS and PSG) and plated at 100 μ L/well in a 96-well round bottom plate. PBMC were treated with test articles (list on next slide) (concentration range: 10 μ M to 0 μ M -3 fold dilution) and PHA and Poly IC (concentration range: 10 μ g/mL to 0 μ g/mL -3 fold dilution). All was set up in triplicates. After 24 hours incubation at 37°C/5%CO₂ humidified standard cell culture incubator, supernatants were harvested and stored at -80°C until assayed for cytokines. Cytokines (GM-CSF, IL-1b, IL-2, IL-6, IL-10, IL-8, IL-12p70, IFN γ , TNF α) were tested on Intellicyt iQue Screener and analyzed using ForeCyt analysis software. Cytokine (IFN α) was tested by standard ELISA. Results are expressed as pg/ml calculated based on the standard curve.

TABLE 39

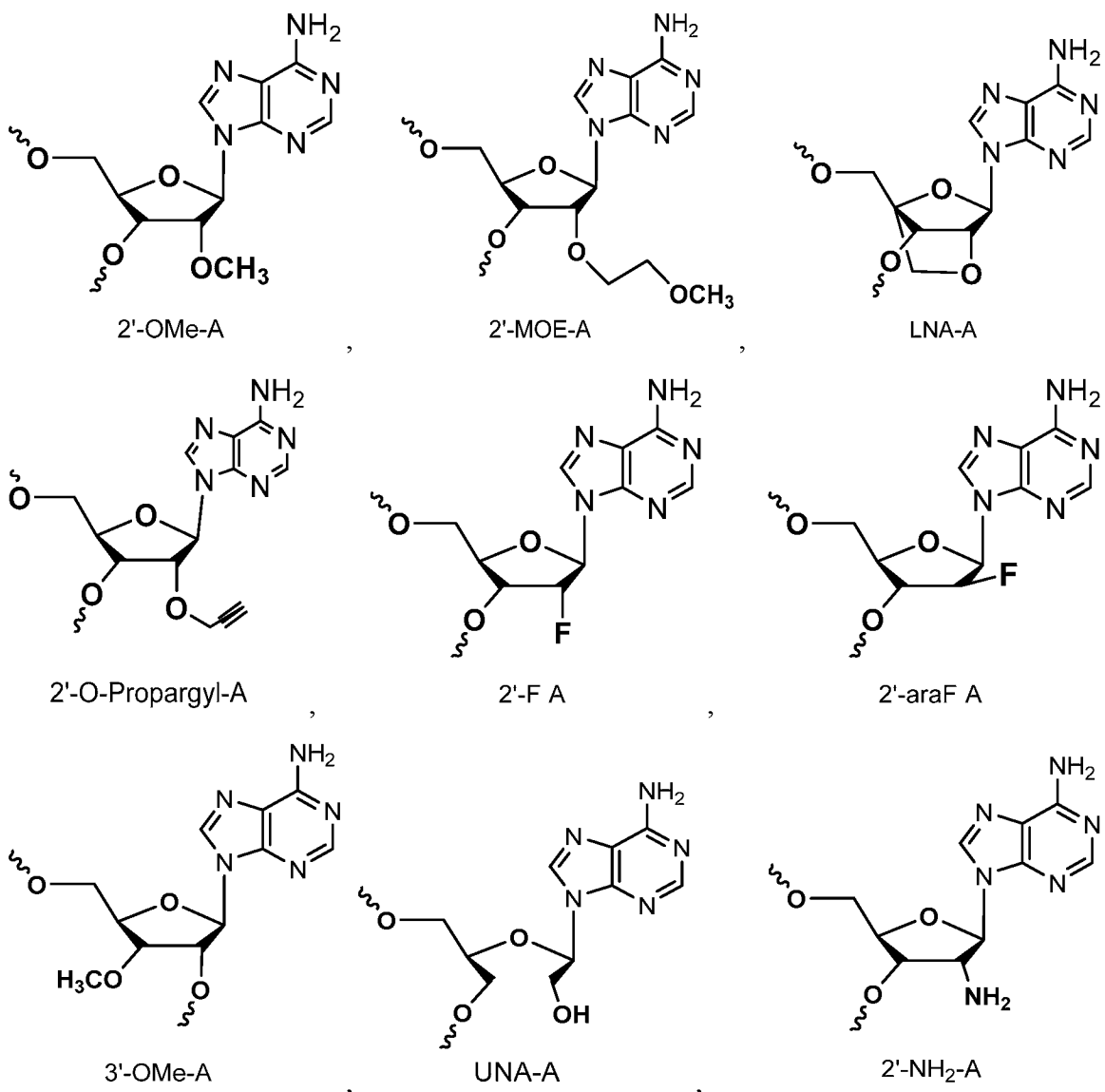
Compound No.	FIG. No.	Immune Reaction ¹
PHA Control	13	Strong
REP-2139	14	Weak
171, 280, 292	15	Weak
296	16	None
134, 277, 284	17	Weak
166, 288	18	None
167, 289	19	None
281, 316	20	None
294	21	Weak
276, 291	22	Weak

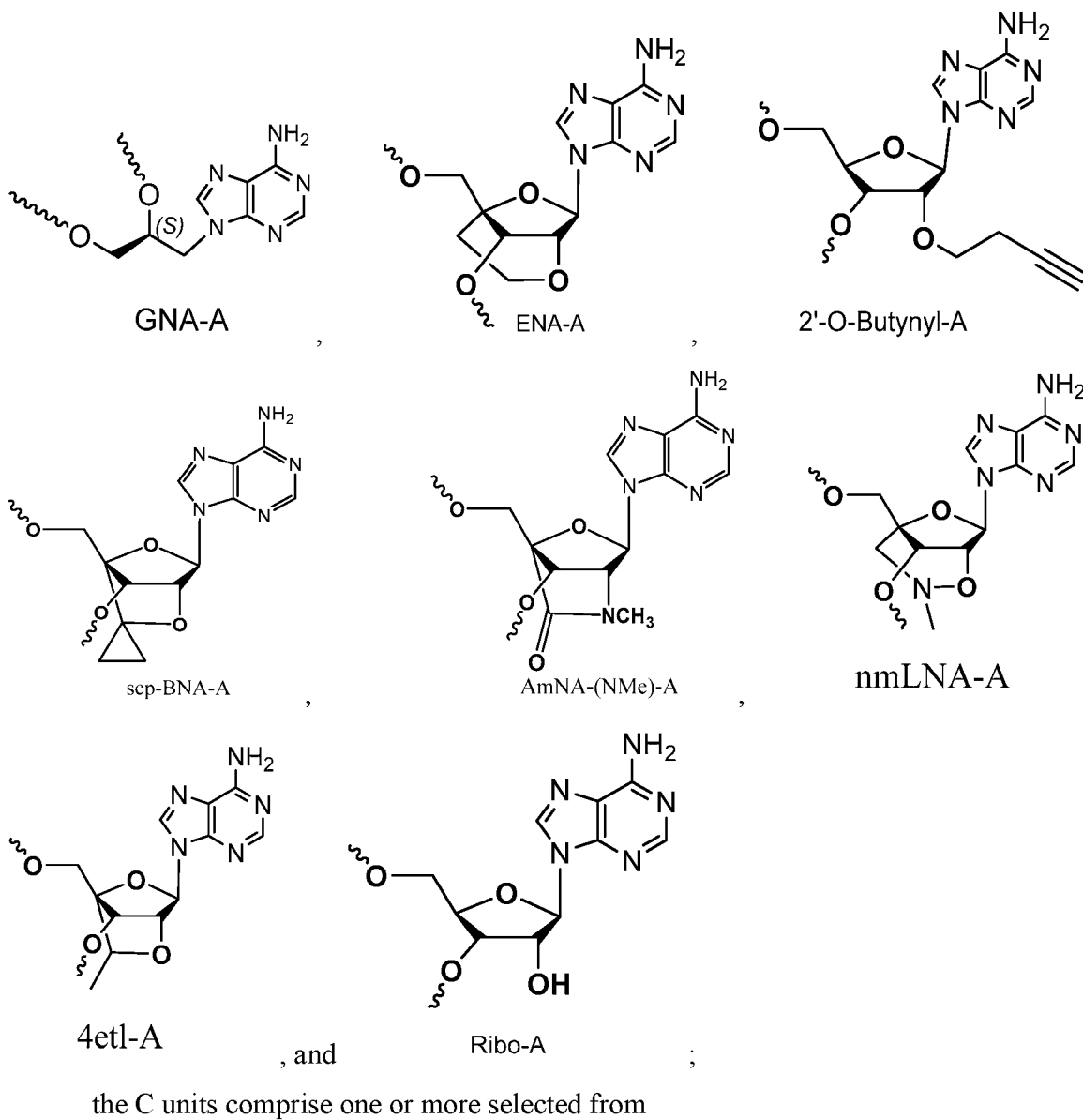
¹Strong: significant induction observed in more than two types of cytokines in the panel tested; Weak: induction observed in one or two types of cytokines in the panel tested; None: no induction observed in any cytokine in the panel tested.

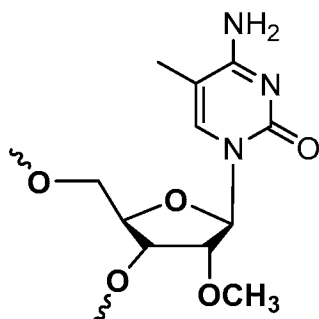
WHAT IS CLAIMED IS:

1. A modified oligonucleotide or complex thereof having sequence independent antiviral activity against hepatitis B, comprising an at least partially phosphorothioated sequence of alternating A and C units, wherein:

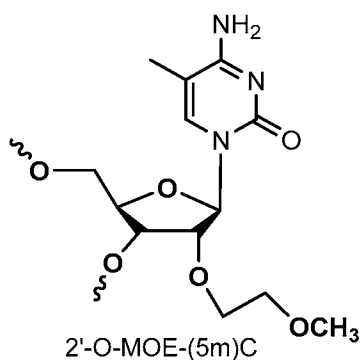
the A units comprise one or more selected from:



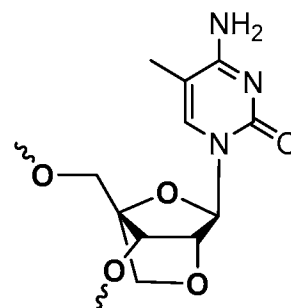




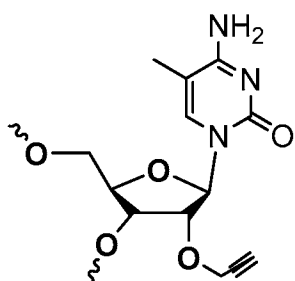
2'-OMe-(5m)C ,



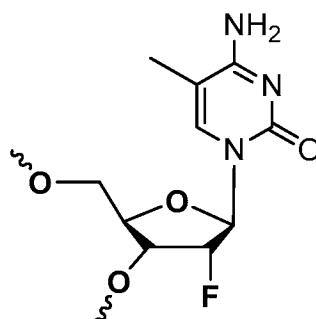
2'-O-MOE-(5m)C ,



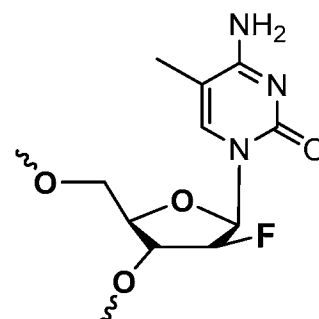
LNA-5mC ,



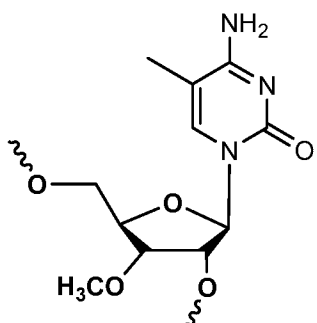
2'-O-Propargyl- (5m)C ,



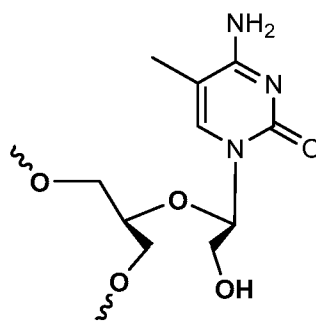
2'-F(5m)C ,



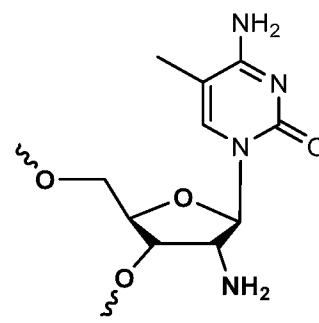
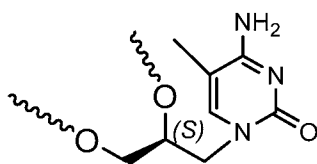
2'-araF(5m)C ,



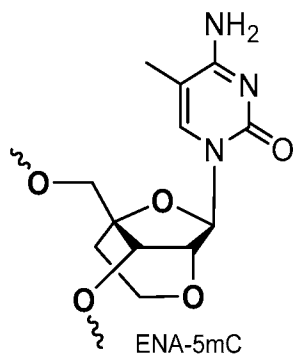
3'-OMe (5m)C ,



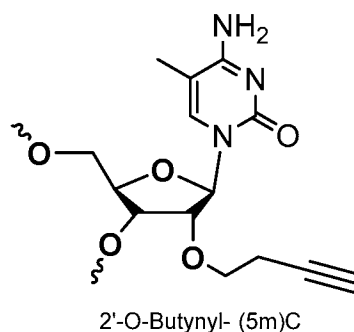
UNA-(5m)C ,

2'-NH₂ (5m)C ,

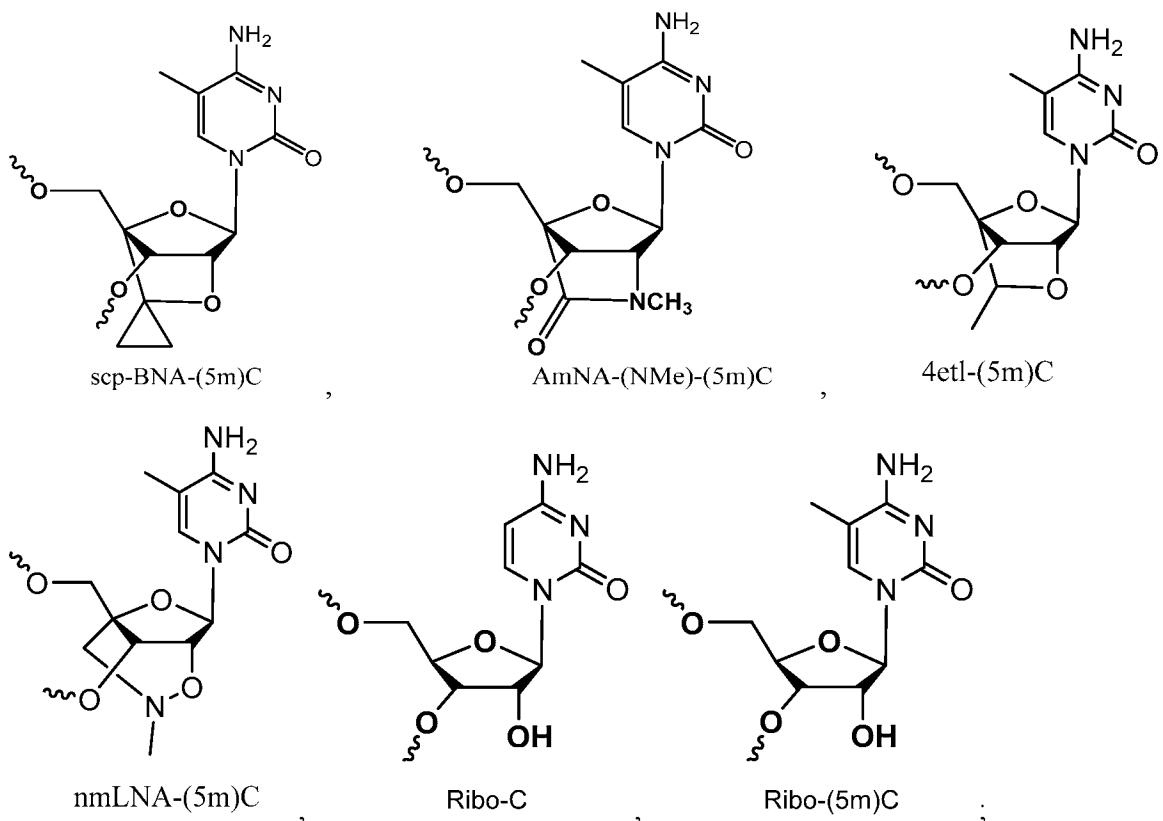
GNA-(5m)C ,



ENA-5mC ,



2'-O-Butynyl- (5m)C ,



each terminal $\sim\text{O}$ is independently hydroxyl, an O,O-dihydrogen phosphorothioate, a dihydrogen phosphate, an endcap or a linking group;

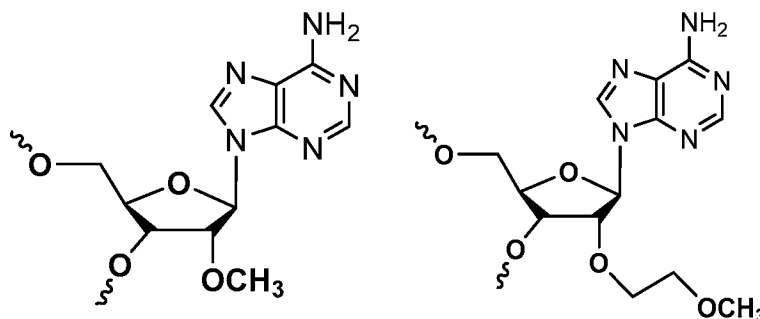
each internal $\sim\text{O}$ is a phosphorus-containing linkage to a neighboring A or C unit, the phosphorus-containing linkage being a phosphorothioate linkage or a modified linkage selected from phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate 5'-phosphoramidate, 3',5'-phosphordiamidate, 5'-thiophosphoramidate, 3',5'-thiophosphordiamidate or diphosphodiester; and

the sequence independent antiviral activity against hepatitis B, as determined by HBsAg Secretion Assay, is greater than that of a reference compound;

with the proviso that, when the sequence of alternating A and C units comprises a Ribo-A unit, the sequence further comprises at least one A unit that is not a Ribo-A unit; and

with the proviso that, when the sequence of alternating A and C units comprises a Ribo-C unit, the sequence further comprises at least one C unit that is not a Ribo-C unit.

2. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



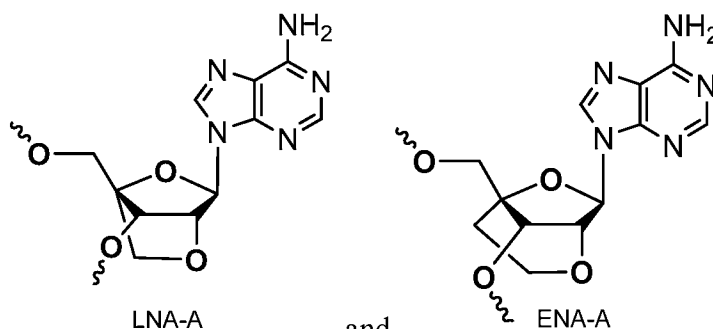
is one or more selected from

2'-OMe-A

and

2'-O-MOE-A

3. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



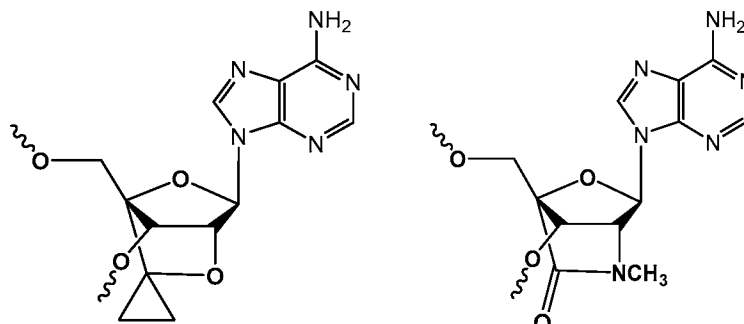
is one or more selected from

LNA-A

and

ENA-A

4. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



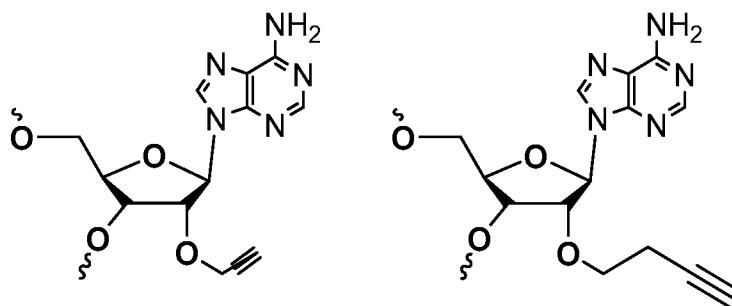
is one or more selected from

scp-BNA-A

and

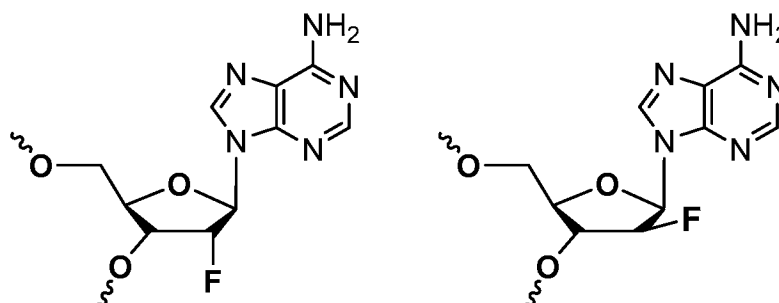
AmNA-(NMe)-A

5. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



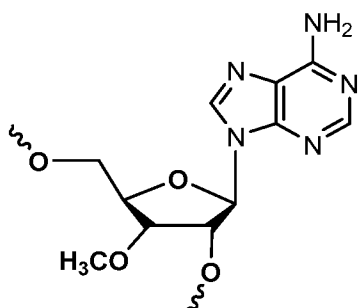
is one or more selected from 2'-O-Propargyl-A and 2'-O-Butynyl-A

6. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



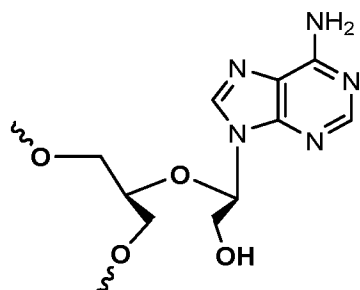
is one or more selected from 2'-F A and 2'-araF A

7. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



is 3'-OMe-A

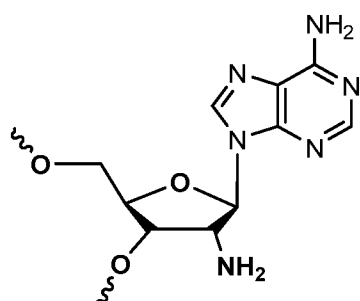
8. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



is

UNA-A

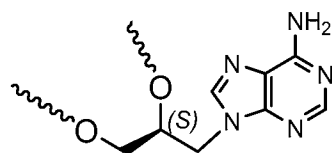
9. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



is

2'-NH₂-A

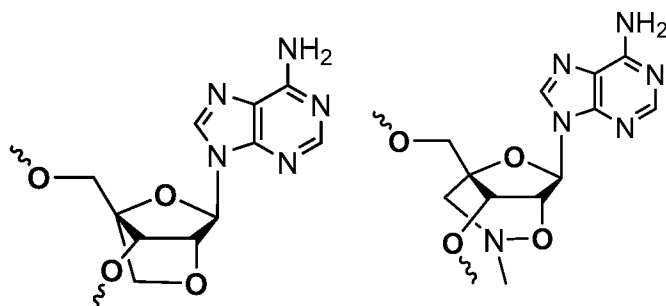
10. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit



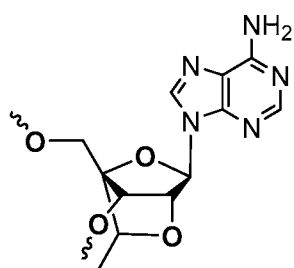
is

GNA-A

11. The modified oligonucleotide or complex thereof of claim 1, wherein the A unit

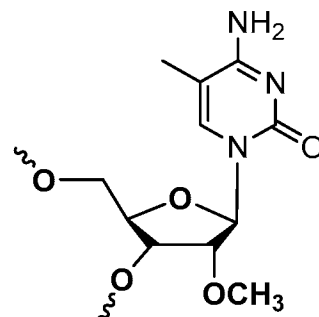


is one or more selected from LNA-A, nmLNA-A, and

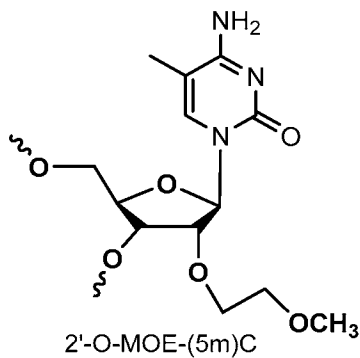


4etl-A

12. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,

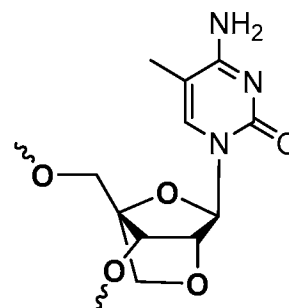


wherein the C unit is one or more selected from 2'-OMe-(5m)C and



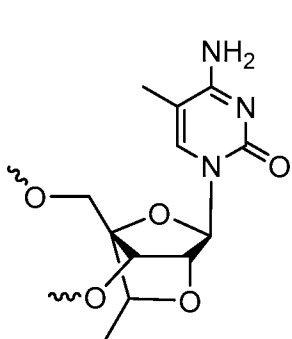
2'-O-MOE-(5m)C

13. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,

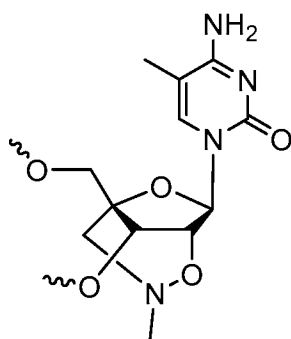


LNA-(5m)C

wherein the C unit is one or more selected from



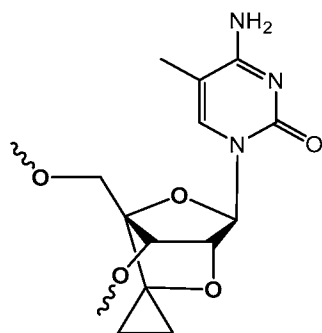
4etl-(5m)C



nmLNA-(5m)C

, and

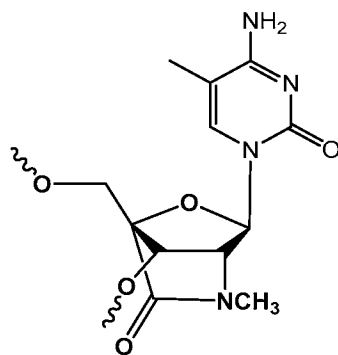
14. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



scp-BNA-(5m)C

wherein the C unit is

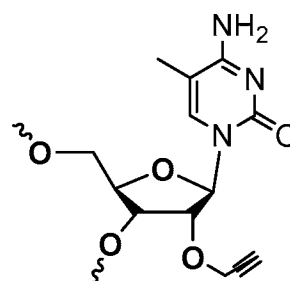
15. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



wherein the C unit is

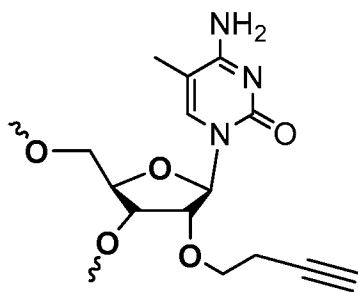
AmNA-(NMe)-(5m)C .

16. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



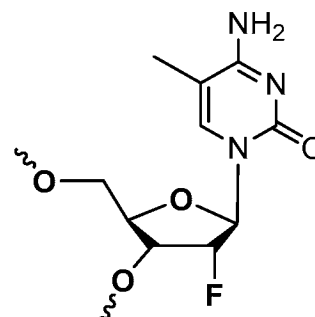
wherein the C unit is one or more selected from

2'-O-Propargyl-(5m)C and



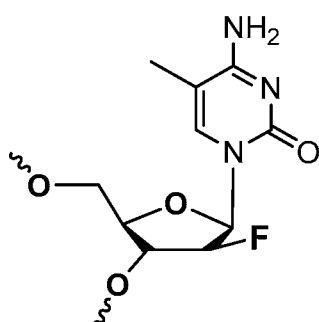
2'-O-Butynyl-(5m)C .

17. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



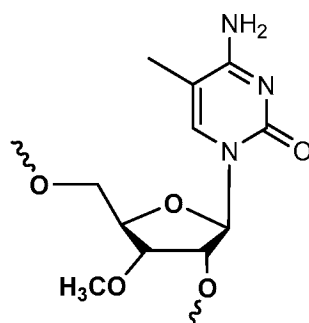
2'-F(5m)C

and



2'-araF(5m)C

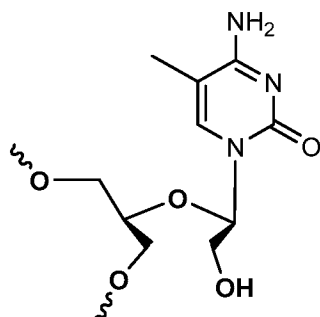
18. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



3'-OMe- (5m)C

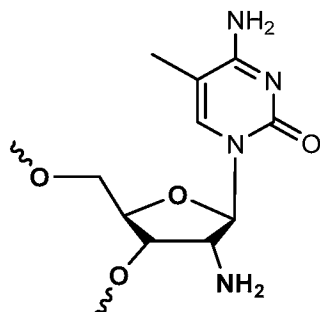
wherein the C unit is

19. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



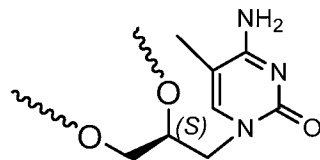
wherein the C unit is UNA-(5m)C

20. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



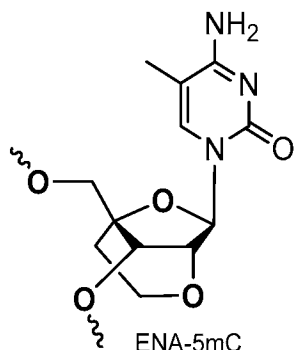
wherein the C unit is 2'-NH₂ (5m)C

21. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



wherein the C unit is GNA-(5m)C

22. The modified oligonucleotide or complex thereof of any one of claims 1 to 11,



wherein the C unit is ENA-5mC

23. The modified oligonucleotide or complex thereof of any one of claims 1 to 22 that is partially phosphorothioated.

24. The modified oligonucleotide or complex thereof of claim 23 that is at least about 85% phosphorothioated.

25. The modified oligonucleotide or complex thereof of any one of claims 1 to 22 that is fully phosphorothioated.

26. The modified oligonucleotide or complex thereof of any one of claims 23 to 25, comprising at least one stereochemically defined phosphorothioate linkage.

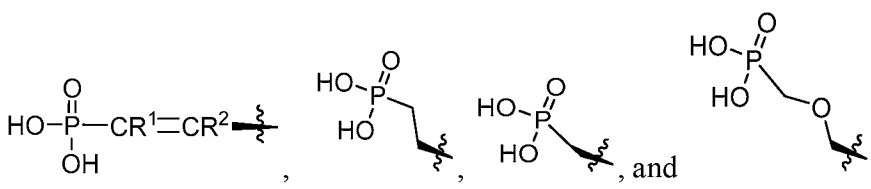
27. The modified oligonucleotide or complex thereof of claim 26, comprising at least 6 stereochemically defined phosphorothioate linkages.

28. The modified oligonucleotide or complex thereof of claim 26 or 27, wherein the at least one stereochemically defined phosphorothioate linkage has an R configuration.

29. The modified oligonucleotide or complex thereof of claim 26 or 27, wherein the at least one stereochemically defined phosphorothioate linkage has an S configuration.

30. The modified oligonucleotide or complex thereof of any one of claims 1 to 29, comprising a 5' endcap.

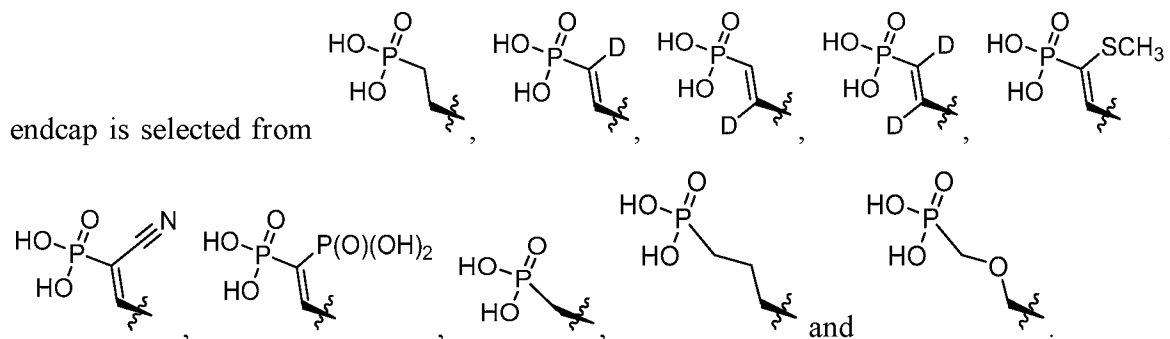
31. The modified oligonucleotide or complex thereof of claim 30, wherein the 5'

endcap is selected from , wherein R¹ and R² are each individually selected from hydrogen, deuterium, phosphate, thioC₁-alkyl, and cyano.

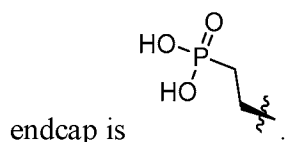
32. The modified oligonucleotide or complex thereof of claim 31, wherein R¹ and R² are both hydrogen.

33. The modified oligonucleotide or complex thereof of claim 31, wherein R¹ and R² are not both hydrogen.

34. The modified oligonucleotide or complex thereof of claim 31, wherein the 5'



35. The modified oligonucleotide or complex thereof of claim 31, wherein the 5'



36. The modified oligonucleotide or complex thereof of any one of claims 1 to 35, wherein the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of about 8 units to about 200 units.

37. The modified oligonucleotide or complex thereof of any one of claims 1 to 35, wherein the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 18 units to 60 units.

38. The modified oligonucleotide or complex thereof of any one of claims 1 to 35, wherein the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 30 units to 50 units.

39. The modified oligonucleotide or complex thereof of any one of claims 1 to 35, wherein the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 34 units to 46 units.

40. The modified oligonucleotide or complex thereof of any one of claims 1 to 35, wherein the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units.

41. The modified oligonucleotide or complex thereof of any one of claims 1 to 40, wherein at least one terminal  is a linking group.

42. The modified oligonucleotide or complex thereof of claim 41, further comprising at least one second oligonucleotide that is attached to the modified oligonucleotide via the linking group.

43. The modified oligonucleotide or complex thereof of claim 41, further comprising a targeting ligand that is attached to the modified oligonucleotide via the linking group.

44. The modified oligonucleotide or complex thereof of claim 43, wherein the targeting ligand comprises N-acetylgalactosamine (GalNAc), triantennary-GalNAc, a tocopherol or cholesterol.

45. The modified oligonucleotide or complex thereof of any one of claims 1 to 44, wherein at least some of the A units are not 2'-O-methylated on the ribose ring.

46. The modified oligonucleotide or complex thereof of any one of claims 1 to 45, wherein at least some of the C units are not 2'-O-methylated on the ribose ring.

47. The modified oligonucleotide or complex thereof of any one of claims 1 to 46, wherein the at least partially phosphorothioated sequence of alternating A and C units further comprises one or more modifications to a phosphorothioate linkage.

48. The modified oligonucleotide or complex thereof of claim 47, wherein the modification to the phosphorothioate linkage is a modified linkage selected from phosphodiester, phosphorodithioate, methylphosphonate, diphosphorothioate 5'-phosphoramidate, 3',5'-phosphordiamidate, 5'-thiophosphoramidate, 3',5'-thiophosphordiamidate or diphosphodiester.

49. The modified oligonucleotide or complex thereof of claim 48, wherein the modified linkage is a phosphodiester linkage.

50. The modified oligonucleotide or complex thereof of any one of claims 1 to 46, further comprising at least two partially phosphorothioated sequences of alternating A and C units linked together to form a concatemer.

51. The modified oligonucleotide or complex thereof of any one of claims 1 to 50, wherein the sequence independent antiviral activity against hepatitis B is at least 2-fold greater than the reference compound.

62. The complex of the modified oligonucleotide of any one of claims 1 to 59, wherein the complex is a monovalent counterion complex.

63. The complex of claim 62, wherein the complex is a lithium, sodium or potassium complex of the modified oligonucleotide.

64. The modified oligonucleotide or complex thereof of claim 1, wherein:
the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;
the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;
the A units comprise at least 12 2'-OMe-A units and at least 1 Ribo-A unit;
the C units comprise at least 15 LNA-5mC units; and
the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 30 nM.

65. The modified oligonucleotide or complex thereof of claim 1, wherein:
the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;
the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;
the A units comprise at least 15 2'-OMe-A units;
the C units comprise at least 7 LNA-5mC units; and
the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 50 nM.

66. The modified oligonucleotide or complex thereof of claim 1, wherein:
the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;
the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;
the A units comprise at least 15-2'-OMe-A units;
the C units comprise at least 3 LNA-5mC units; and
the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 100 nM.

67. The modified oligonucleotide or complex thereof of claim 1, wherein:
the at least partially phosphorothioated sequence of alternating A and C units is at least 85% phosphorothioated;
the at least partially phosphorothioated sequence of alternating A and C units has a sequence length in the range of 36 units to 44 units;
the A units comprise at least 18 2'-OMe-A units;
the C units comprise at least 15 LNA-5mC units; and
the modified oligonucleotide has an EC₅₀ value, as determined by HBsAg Secretion Assay, that is less than 30 nM.
68. The complex of the modified oligonucleotide of any one of claims 64 to 67, wherein the complex is a monovalent counterion complex that comprises a sodium or potassium complex of the modified oligonucleotide.
69. A pharmaceutical composition, comprising an amount of the modified oligonucleotide or complex thereof of any one of claims 1 to 68, that is effective for treating a subject infected with hepatitis B; and a pharmaceutically acceptable carrier.
70. A pharmaceutical composition, comprising an amount of the modified oligonucleotide or complex thereof of any one of claims 1 to 68, that is effective for treating a subject infected with hepatitis D; and a pharmaceutically acceptable carrier.
71. A treatment for hepatitis B, hepatitis D or both, comprising an effective amount of the modified oligonucleotide or complex thereof of any one of claims 1 to 68, or the pharmaceutical composition of claim 69 or 70.
72. A method of treating hepatitis B, comprising administering an effective amount of the modified oligonucleotide or complex thereof of any one of claims 1 to 68, the pharmaceutical composition of claim 69 or 70, or the treatment of claim 71, to a subject in need thereof.
73. The method of claim 72, wherein the modified oligonucleotide or complex thereof is administered to the subject by a parenteral route.
74. The method of claim 72, wherein the modified oligonucleotide or complex thereof is administered to the subject intravenously.
75. The method of claim 72, wherein the modified oligonucleotide or complex thereof is administered to the subject subcutaneously.

76. The method of any one of claims 72 to 75, further comprising administering an effective amount of a second treatment for hepatitis B to the subject.

77. The method of claim 76, wherein the second treatment for hepatitis B comprises a second oligonucleotide having sequence independent antiviral activity against hepatitis B, an siRNA oligonucleotide, an anti-sense oligonucleotide, a nucleoside, an interferon, an immunomodulator, a capsid assembly modulator, or a combination thereof.

78. The method of claim 77, wherein the second treatment for hepatitis B comprises an anti-sense oligonucleotide.

79. The method of claim 77, wherein the second treatment for hepatitis B comprises a capsid assembly modulator.

80. A method of treating hepatitis D, comprising administering an effective amount of the modified oligonucleotide or complex thereof of any one of claims 1 to 68, the pharmaceutical composition of claim 69 or 70, or the treatment of claim 71, to a subject in need thereof.

81. The method of claim 80, wherein the modified oligonucleotide or complex thereof is administered to the subject by a parenteral route.

82. The method of claim 80, wherein the modified oligonucleotide or complex thereof is administered to the subject intravenously.

83. The method of claim 80, wherein the modified oligonucleotide or complex thereof is administered to the subject subcutaneously.

84. The method of any one of claims 80 to 83, further comprising administering an effective amount of a second treatment for hepatitis D to the subject.

85. The method of claim 84, wherein the second treatment for hepatitis D comprises a second oligonucleotide having sequence independent antiviral activity against hepatitis B, an anti-sense oligonucleotide, a nucleoside, an interferon, a capsid assembly modulator, or a combination thereof.

86. The method of claim 85, wherein the second treatment for hepatitis B comprises an anti-sense oligonucleotide.

87. The method of claim 85, wherein the second treatment for hepatitis B comprises a capsid assembly modulator.

88. A method of treating hepatitis B or hepatitis D, comprising subcutaneously administering an effective amount of an antiviral oligonucleotide or complex thereof to a subject in need thereof, wherein the antiviral activity of the oligonucleotide occurs principally by a sequence independent mode of action.

89. The method of claim 88, wherein the antiviral oligonucleotide is REP 2139, REP 2055, REP 2165 or a chelate complex thereof.

90. The method of claim 88, wherein the antiviral oligonucleotide is the modified oligonucleotide or complex thereof of any one of claims 1 to 68, the pharmaceutical composition of claim 69 or 70, or the treatment of claim 71.

91. The method of any one of claims 88 to 90, comprising subcutaneously administering a safe and effective amount of the antiviral oligonucleotide or complex thereof to a human subject in need thereof, at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration.

92. The modified oligonucleotide or complex thereof of any one of claims 1 to 68 for use in the treatment of hepatitis B.

93. The modified oligonucleotide or complex thereof of any one of claims 1 to 68 for use in the treatment of hepatitis D.

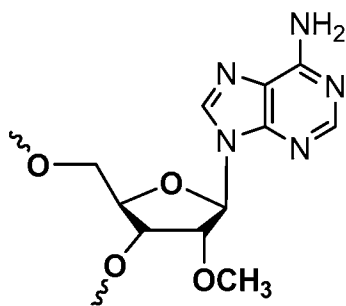
94. Use of the modified oligonucleotide or complex thereof of any one of claims 1 to 68 in the preparation of a medicament for the treatment of hepatitis B.

95. Use of the modified oligonucleotide or complex thereof of any one of claims 1 to 68 in the preparation of a medicament for the treatment of hepatitis D.

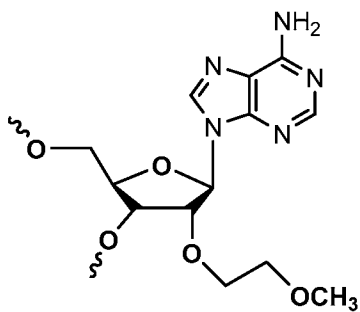
96. The modified oligonucleotide or complex thereof of claims 92 or 93, or the use of claims 94 or 95, wherein the use comprises subcutaneously administering a safe and effective amount of the antiviral oligonucleotide or complex thereof to a human subject in need thereof, at a dosage lower than otherwise expected based on liver levels observed following otherwise comparable intravenous administration.

97. A dinucleotide consisting of an A unit and a C unit connected by a stereochemically defined phosphorothioate linkage, wherein:

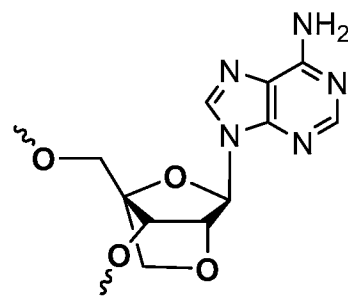
the A units comprise one or more selected from:



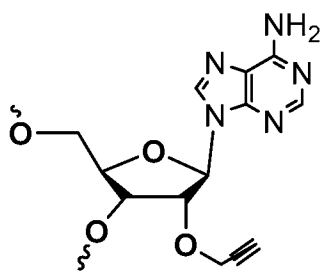
2'-OMe-A



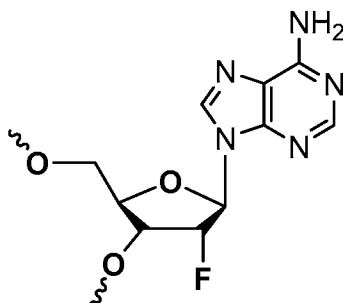
2'-O-MOE-A



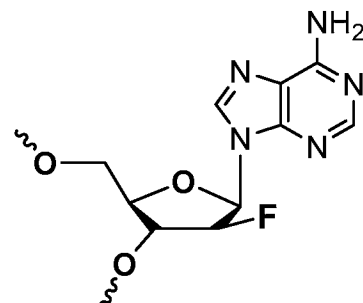
LNA-A



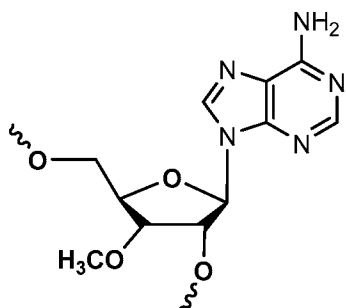
2'-O-Propargyl-A



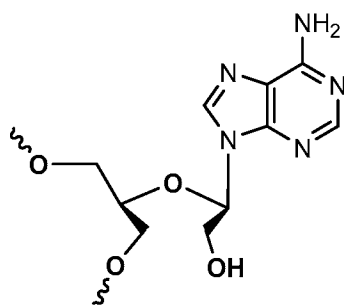
2'-F A



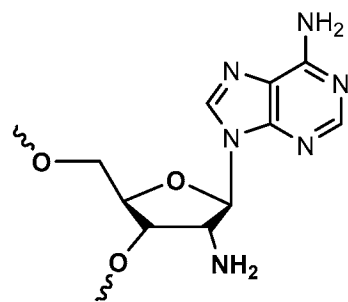
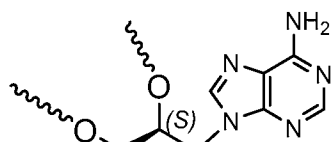
2'-araF A



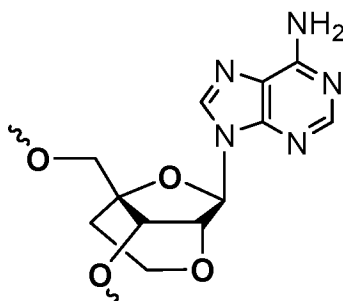
3'-OMe-A



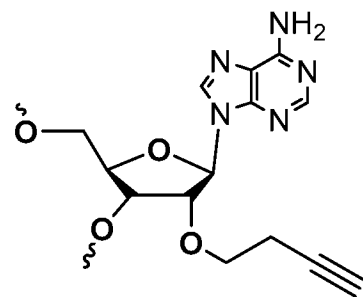
UNA-A

2'-NH₂-A

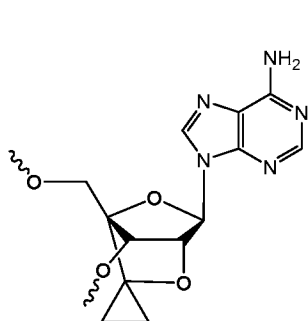
GNA-A



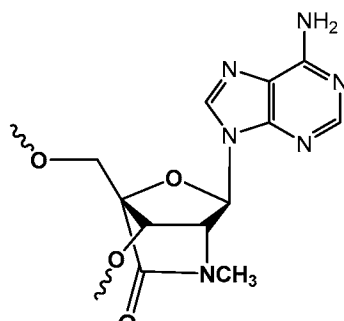
ENA-A



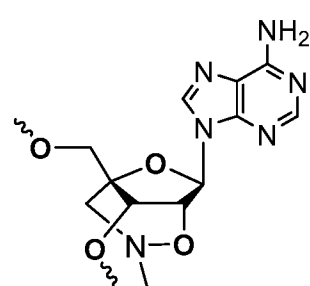
2'-O-Butynyl-A



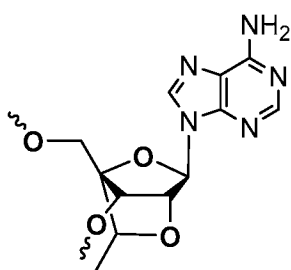
scp-BNA-A



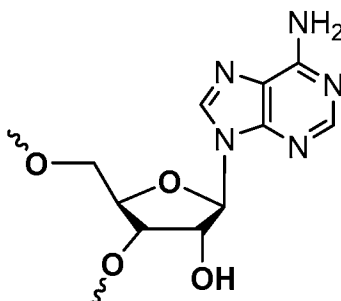
AmNA-(NMe)-A



nmLNA-A



4etl-A

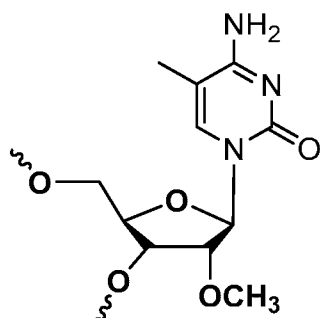


Ribo-A

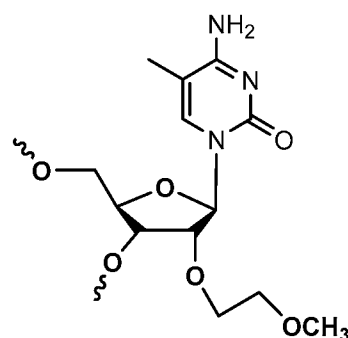
, and

;

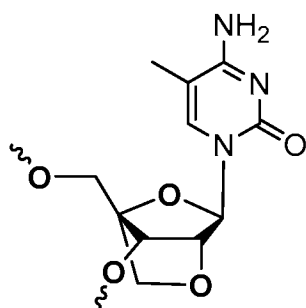
the C units comprise one or more selected from



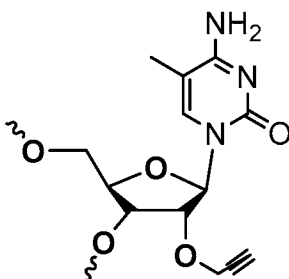
2'-OMe-(5m)C



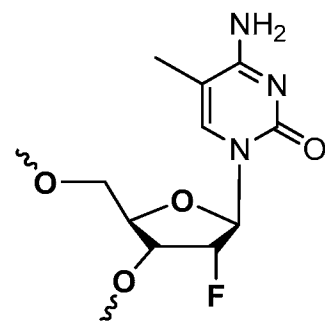
2'-O-MOE-(5m)C



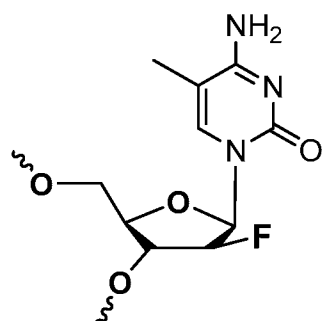
LNA-5mC



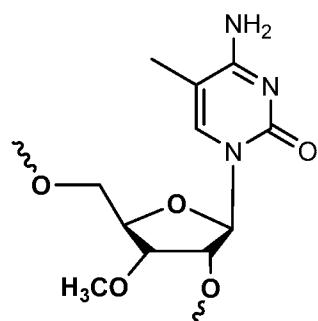
2'-O-Propargyl-(5m)C



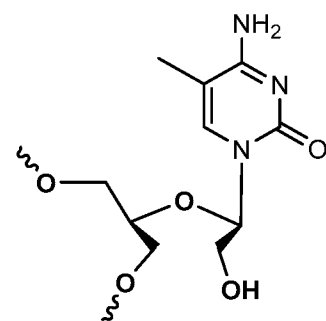
2'-F(5m)C



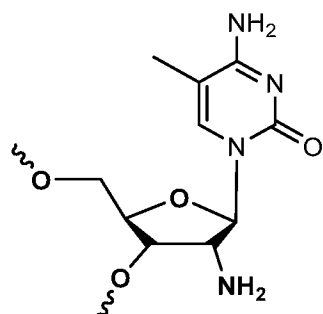
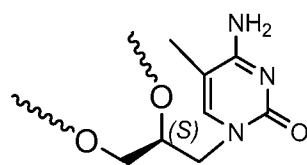
2'-araF(5m)C



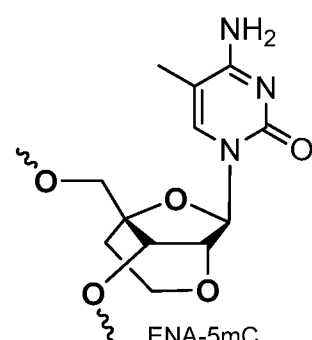
3'-OMe (5m)C



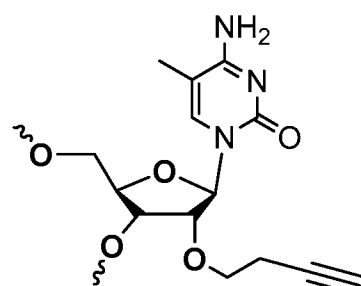
UNA-(5m)C

2'-NH₂ (5m)C

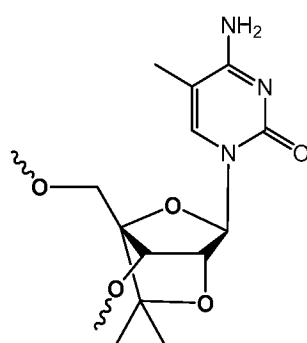
GNA-(5m)C



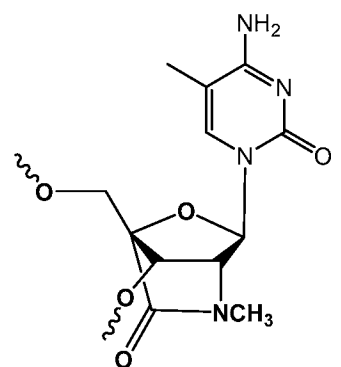
ENA-5mC



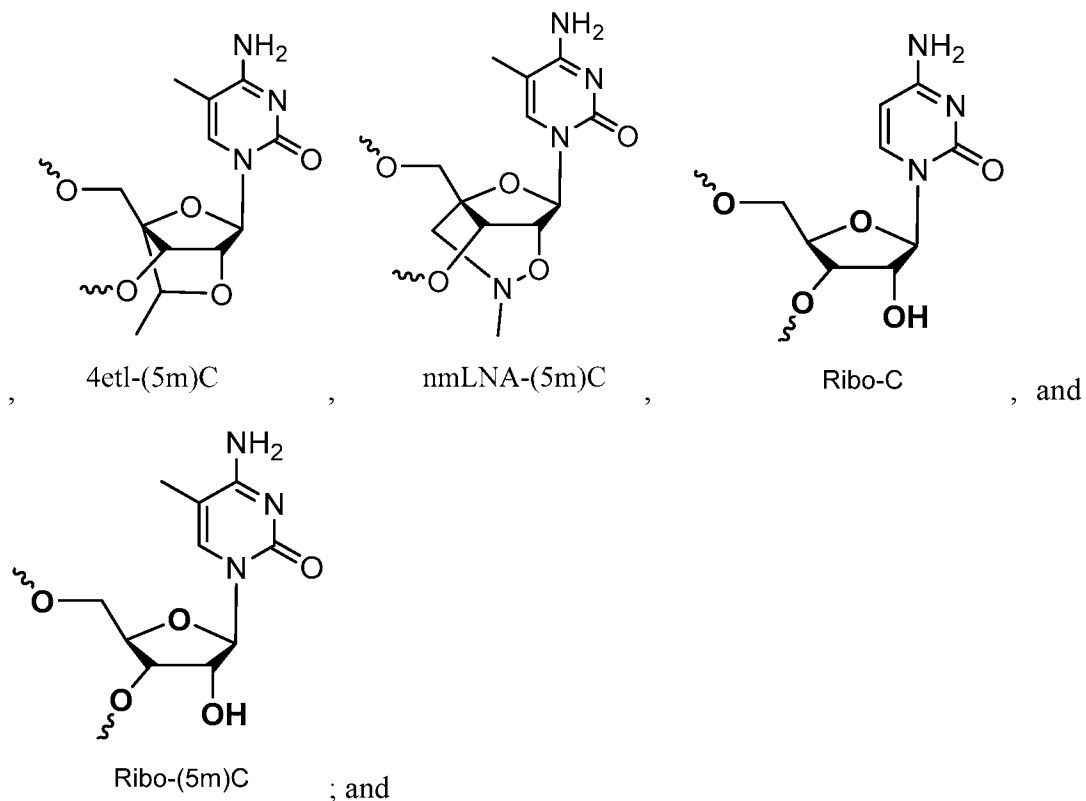
2'-O-Butynyl- (5m)C



scp-BNA-(5m)C



AmNA-(NMe)-(5m)C

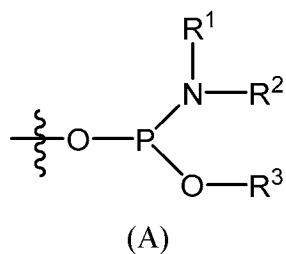


each $\sim\text{O}$ is independently hydroxyl, an O,O-dihydrogen phosphorothioate, a phosphoramidite, a dimethoxytrityl ether, or the stereochemically defined phosphorothioate linkage.

98. The dinucleotide of claim 97, with the proviso that the dinucleotide does not comprise both the Ribo-A unit and the Ribo-C unit.

99. The dinucleotide of claim 97 selected from the dinucleotides that comprise or consist of any two of the building block monomers described in Tables 4 and 5.

100. The dinucleotide of claim 97 or 98, wherein an $\sim\text{O}$ is a phosphoramidite of the following formula (A):

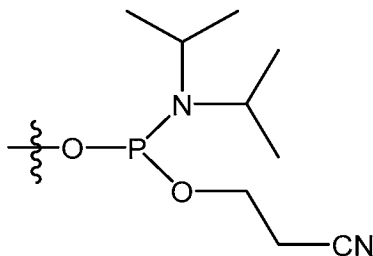


wherein:

R^1 and R^2 are each individually a C_{1-6} alkyl; and

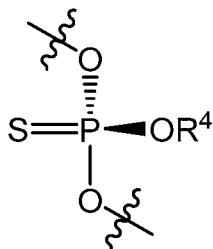
R^3 is a C_{1-6} alkyl or a cyano C_{1-6} alkyl.

101. The dinucleotide of claim 100, wherein the phosphoramidite of the formula (A) is a phosphoramidite of the following formula (A1):

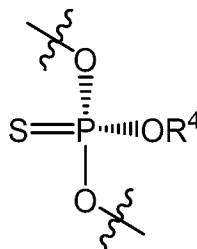


(A1).

102. The dinucleotide of any one of claims 97 to 101, wherein the stereochemically defined phosphorothioate linkage is a phosphorothioate of the following Formulae (B1) or (B2):



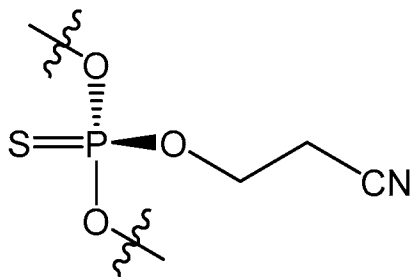
(B1)



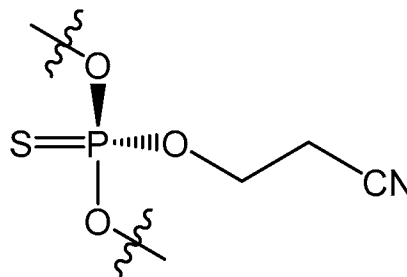
(B2)

wherein R^4 is a C_{1-6} alkyl or a cyano C_{1-6} alkyl.

103. The dinucleotide of claim 102, wherein the phosphorothioate of the formula (B1) is a phosphorothioate of the formula (B3), or the phosphorothioate of the formula (B2) is a phosphorothioate of the formula (B4), as follows:



(B3)



(B4).

104. A method for making the modified oligonucleotide of any one of claims 1 to 68, comprising coupling the dinucleotide of any one of claims 97 to 103.

FIGURE 2

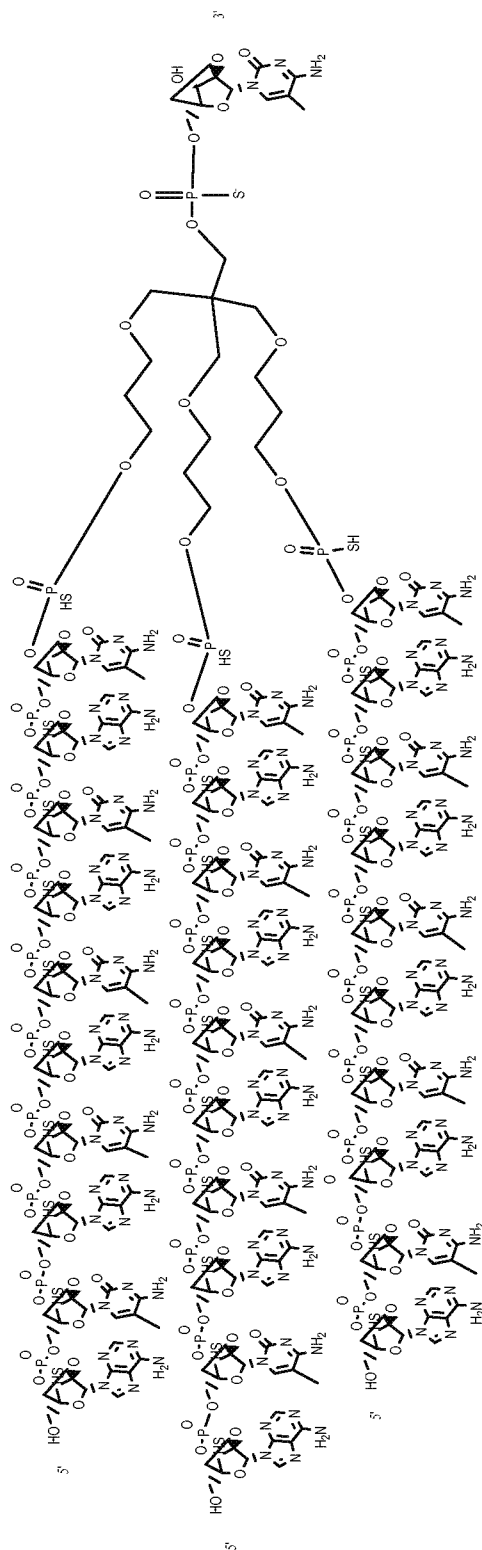


FIGURE 3A

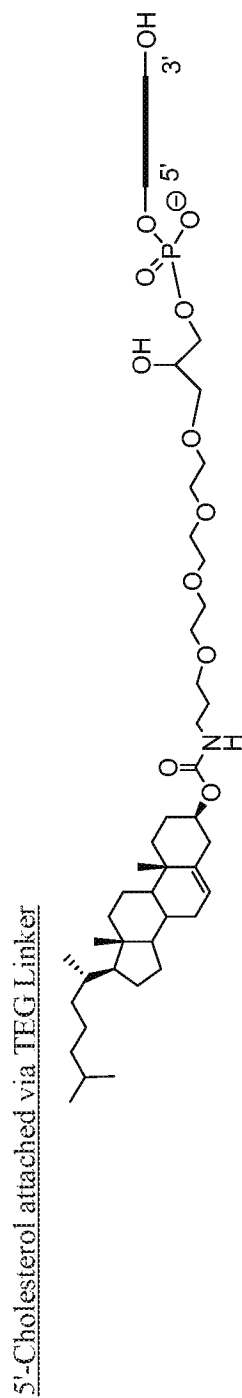


FIGURE 3B

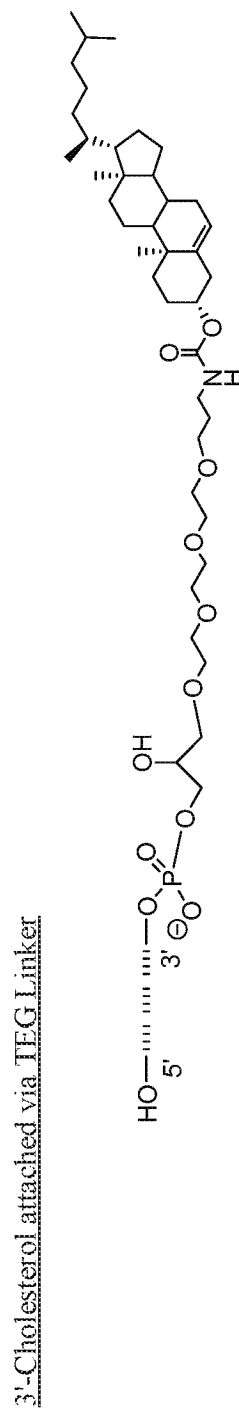


FIGURE 3C

5'-Vitamin E attached via TEG linker

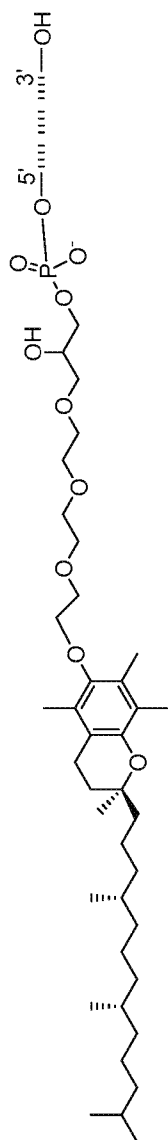


FIGURE 3D

3'-Vitamin E attached via TEG linker

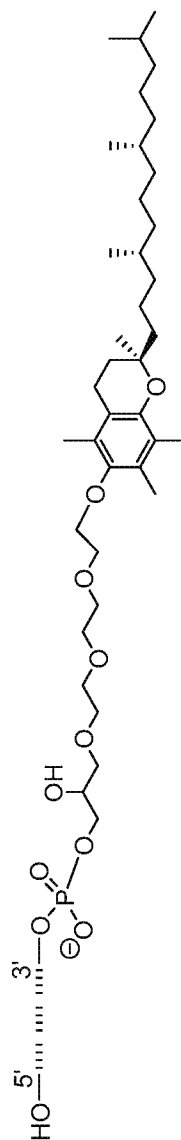


FIGURE 4A

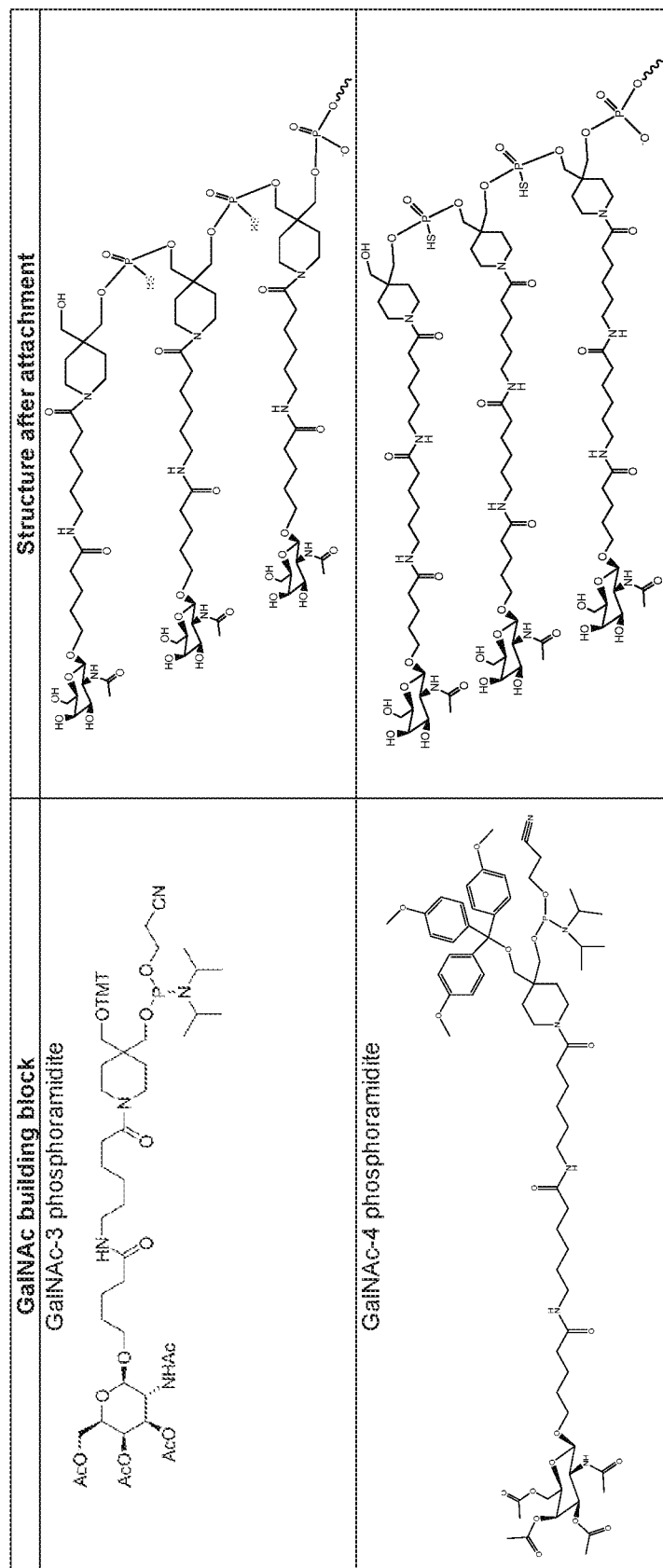


FIGURE 4A (continued)

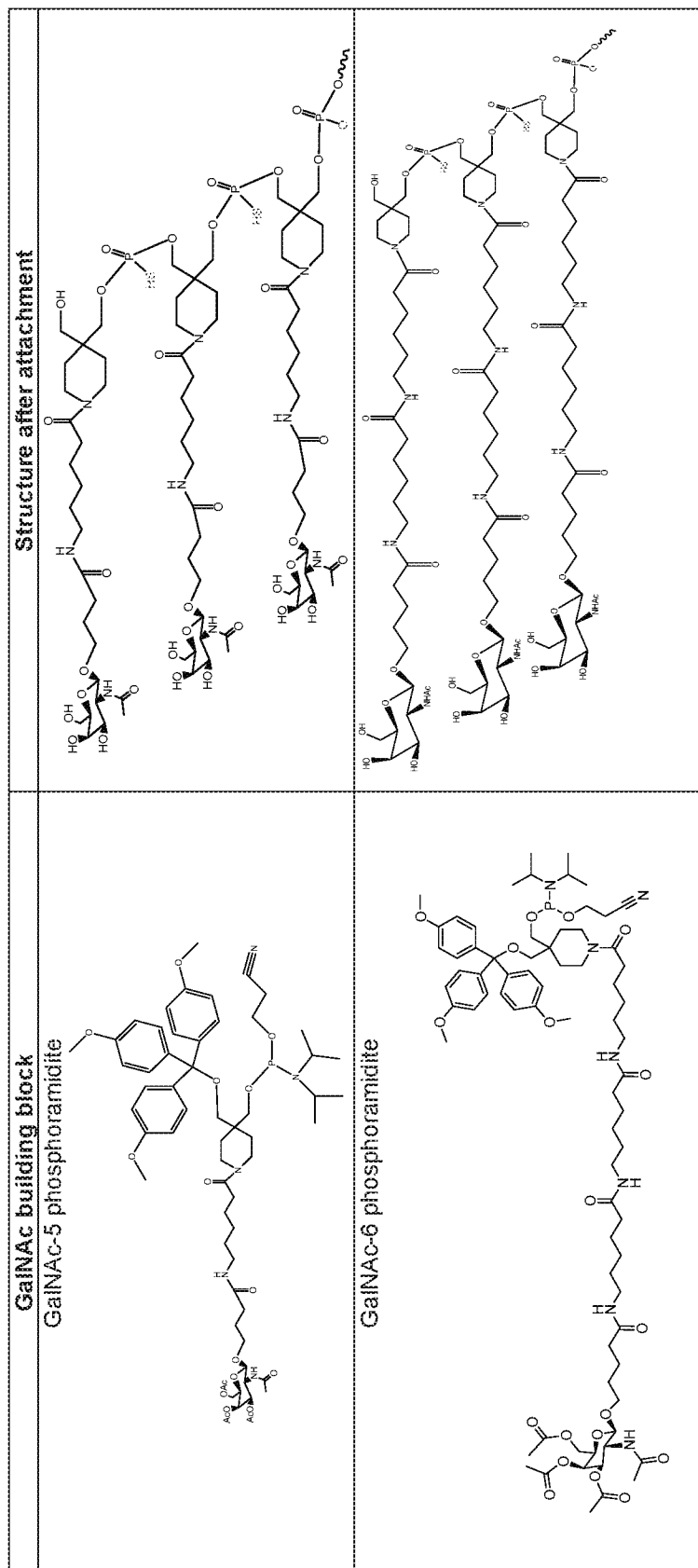


FIGURE 4B

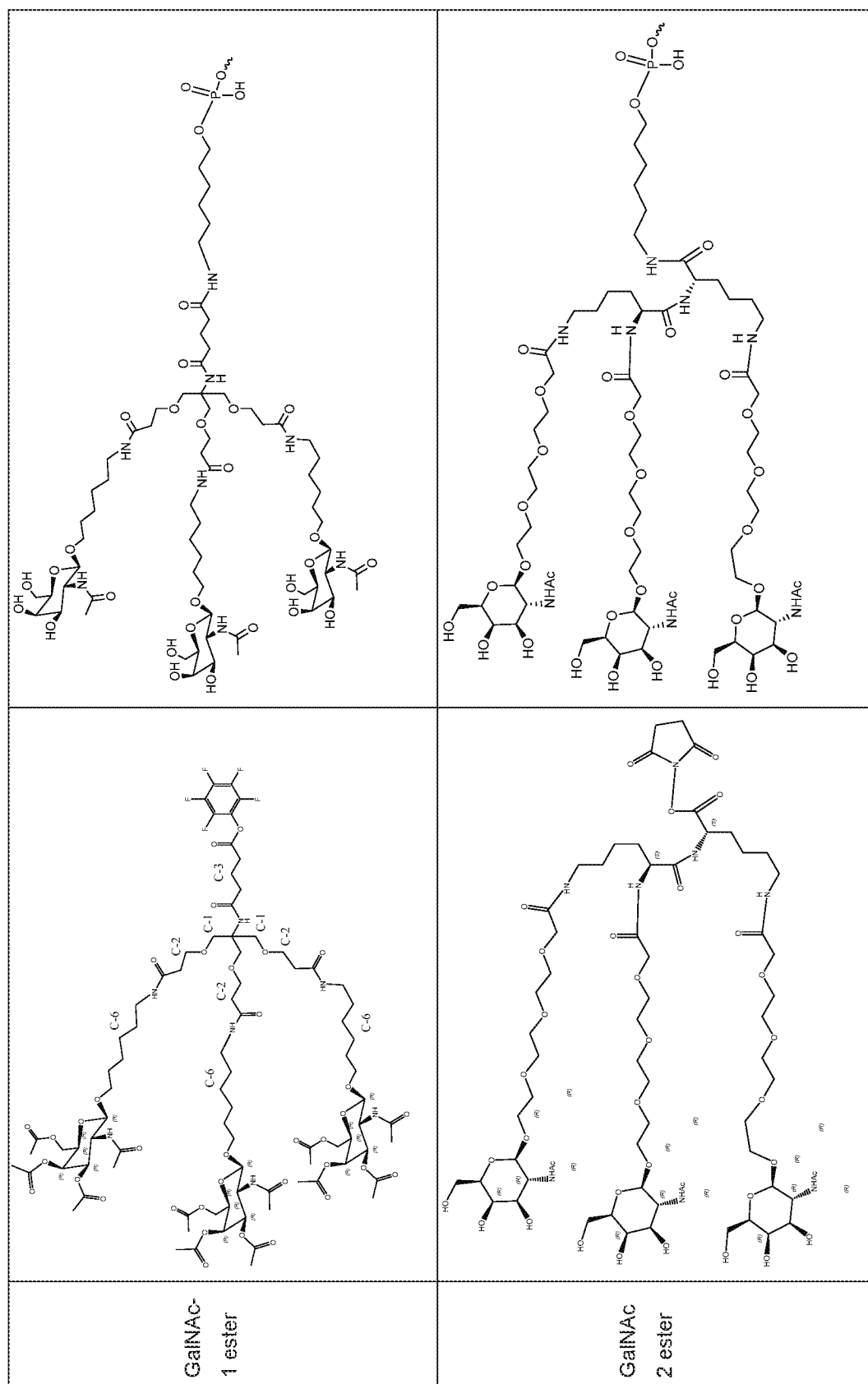


FIGURE 5

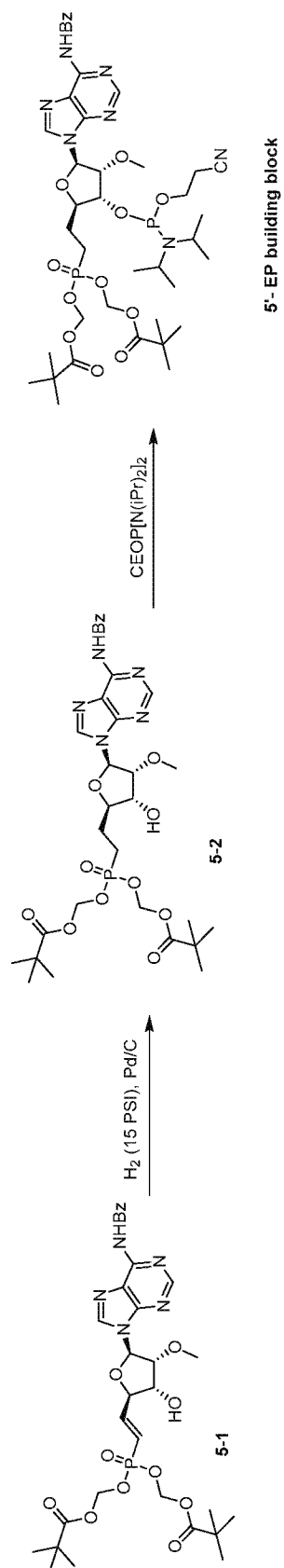


FIGURE 6A (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μ M)	CC ₅₀ (μ M)	Comments
286	5'-mApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)Cps lnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)Cps mApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)Cps lnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)C 3'	AAA		PS 40mer 2'-OMe; 13 LNA (every 3 rd base)
287	5'-VPmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)Cps lnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)Cps mApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)Cps lnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)C 3'	AAAA	A	PS 40mer 2'-OMe; 13 LNA (every 3 rd base) 5'-VP endcap
288	5'-mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)C 3'	AAAA	A	PS 40mer 2'-OMe; 10 LNA (every 4 th base)
289	5'-VP-mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)Cps mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)C 3'	AAAA	A	PS 40mer 2'-OMe; 10 LNA (every 4 th base) 5'-VP endcap

FIGURE 6A (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μ M)	CC ₅₀ (μ M)	Comments
290	5'mApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmA mApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmA psRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsR m(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsR m(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)C 3'	AA	A	PS 40mer 20 R isomer dimer blocks
291	5'VPmApsm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)Cps mApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmA psRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsR m(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsR m(5m)CpsmApsRm(5m)CpsmApsRm(5m)CpsmApsRm(5m)C 3'	AAAA	A	PS 40mer 19 R isomer dimer blocks; 1 m(5m)C 5'-VP endcap
292	5'mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)Cps mApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsln(5m)Cps mApsm(5m)CpsmApsm(5m)C 3'	AAA	A	PS 40mer 2'-OMe; 5' LNA
293	5'VPmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)Cps mApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsln(5m)Cps mApsm(5m)CpsmApsm(5m)C 3'	AAAA	A	PS 40mer 2'-OMe; 5' LNA 5'-VP endcap

FIGURE 6A (continued)

FIGURE 6A (continued)

FIGURE 6B

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
6, 274, 283	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)C pslnApsln(5m)C 3'	A	B
376	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)C 3'	D	A
371	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnA 3'	D	A
372	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)C 3'	D	A
273, 282	5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)C-3'	D	A
367	5' mApsmCpsmApsmCpsmApsmCpsmApsmCpsmApsmCpsmApsmCpsmCpsmCpsmA psmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmAps mCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmCpsmC-3'	C	A
368	5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)C 3'	D	A
369	5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmA 3'	D	A

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
303	5' InApsln(5m)CpslnApsln(5m)CpsrApsln(5m)CpslnApsln(5m)CpslnApsrCps lnApsln(5m)CpslnApsln(5m)CpsrApsln(5m)CpslnApsln(5m)CpslnApsrCps lnApsln(5m)CpslnApsln(5m)CpsrApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)C 3'	B	B
305	5' InApsln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnAps ln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)C polnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnAps ln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)CpolnApsln(5m)C polnApsln(5m)CpolnApsln(5m)C 3'	C	A
309	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpolnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)C pslnApsln(5m)CpolnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)C 3'	D	B
297	5' mApsm(5m)CpsmApsm(5m)CpsrApsm(5m)CpsmApsm(5m)CpsmApsrCps mApsm(5m)CpsmApsm(5m)CpsrApsm(5m)CpsmApsm(5m)CpsmApsrCps mApsm(5m)CpsmApsm(5m)CpsrApsm(5m)CpsmApsm(5m)CpsmApsrCps mApsm(5m)CpsmApsm(5m)CpsrApsm(5m)CpsmApsm(5m)CpsmAps m(5m)C 3'	C	A
298	5' mApsm(5m)CpsmApsm(5m)CpsrApsrCpsmApsm(5m)CpsmApsr(5m)CpsrApsm(5m) CpsmApsm(5m)CpsrApsrCpsmApsm(5m)CpsmApsrCpsrAps m(5m)CpsmApsm(5m)CpsrApsrCpsmApsm(5m)CpsmApsrCpsrApsm(5m)CpsmAps m(5m)CpsrApsrCpsmApsm(5m)CpsmApsm(5m)C 3'	D	A
300	5' mApsm(5m)CpsmApsm(5m)CpsrApsrCpsmApsm(5m)CpsmApsrCpsrAps m(5m)CpsmApsm(5m)CpsrApsrCpsmApsm(5m)CpsmApsrCpsrApsm(5m)CpsmAps m(5m)CpsrApsrCpsmApsm(5m)CpsmApsm(5m)C 3'	D	A

FIGURE 6B (continued)

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
205	5' mApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m) CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsS m(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsS m(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsS m(5m)CpsmApsSm(5m)CpsmApsSm(5m)CpsmApsSm(5m)C 3'	D	A
353	5' 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)C 3'	B	A
351	5' 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)C-Cholesterol 3'	D	A
352	5' 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)Cps 3mAps3m(5m)Cps3mAps3m(5m)Cps3mAps3m(5m)C-Tocopherol 3'	D	A
178	5' mApsRm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsRm(5m)C psmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsRm(5m)CpsmApsm(5m)C psmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsRm(5m)C psmApsm(5m)CpsmApsm(5m)C 3'	A	A

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
238	5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps-DBLR-ps-c4-CONH-c2-psm(5m)C 3'-5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)C 5'	D	A
240	5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)Cps-DBLR-ps-c4-CONH-c2-ps(5m)mC 3'-5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)C 5'	B	A
241	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)Cps-TREB-ps-c3-O-C3-ps(5m)mC 3' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)Cps 5'	B	A
242	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnAps-TREB-ps-c3-O-C3-psm(5m)C 3' lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnAps 5' lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnAps 5'	A	A
243	5' InApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps-TREB-ps-c3-O-C3-psm(5m)C 3'-5' lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnA 5'	A	A

FIGURE 6B (continued)

FIGURE 6B (continued)

[illegible]

FIGURE 6B (continued)

[illegible]

FIGURE 6B (continued)

FIGURE 6B (continued)

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
121	5' VP-mApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m) In(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)C 3'	B	A
122	5' VP-mApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnAps ln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)CpslnApsln(5m)Cps lnApsln(5m)C 3'	A	A
181	5' VP-mApsm(5m)CpsmApsm(5m)CpsmApsRm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsRm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsR m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsRm(5m)CpsmApsm(5m)C psmApsm(5m)CpsmApsRm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsRm(5m)C psmApsm(5m)CpsmApsm(5m)C 3'	C	A
147	5' unApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunAps ln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsl n(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunAps ln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)Cps 3'	D	A
148	5' unApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunAps ln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsl n(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunAps ln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)C 3'	D	A
149	5' unApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunAps ln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsl n(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunAps ln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsln(5m)CpsunApsl n(5m)C 3'	B	A

FIGURE 6B (continued)

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
167, 289	5' VP-mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmAps ln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)C 3'	B	A
164, 286	5' mApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)C 3'	C	A
166, 288	5' mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmAps ln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)C 3'	A	A
171, 280, 292	5' mApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsln(5m)CpsmApsm(5m)CpsmApsm(5m)C 3'	B	A

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
251	5' cpApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm 5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsm 5m)CpsmApsm(5m)C 3'	A	A
356	5' 4'-Me-mApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln 5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln 5m)CpsmApsln(5m)C 3'	A	A
320	5' mApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln 5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)C 3'	A	A
357	5' mApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln 5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpslnApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpslnApsm(5m)C 3'	A	B
322	5' mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln 5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmAps ln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)C 3'	B	A
317	5' EP-mApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln 5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmAps ln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm 5m)CpsmApsln(5m)C 3'	A	B

FIGURE 6B (continued)

FIGURE 6B (continued)

No.	Compound Structure (5' to 3')	EC ₅₀ (μM)	CC ₅₀ (μM)
348	5' GalNAc5mApsm(5m)CpsmApsln(5m)CpsmAps m(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmAps ln(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsm(5m)CpsmApsln(5m)CpsmApsm(5m)CpsmApsln(5m)C 3'		
350	5' GalNAc5mApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsrApsln(5m)CpsmApsln(5m)Cps mApsln(5m)CpsrApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsrAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsrApsln(5m)CpsmApsln(5m)Cps mApsln(5m)CpsrApsln(5m)CpsmApsln(5m)CpsmApsln(5m)C 3'		
349	5' GalNAc5mApsm(5m)CpsmApsm(5m)CpsmAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnAps m(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)CpslnApsm(5m)CpsmApsm(5m)CpsmApsm(5m)C 3'		
339	5' mApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpscpApsln(5m)C 3'	A	A
252	5' mApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmAps ln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApsln(5m)CpsmApscp(5m)C 3'	A	A

FIGURE 6B (continued)

FIGURE 6B (continued)

FIGURE 6B (continued)

In Figure 6B, the following abbreviations are used, and each is described in Table A.

TABLE A

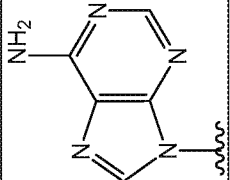
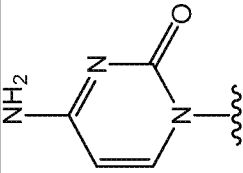
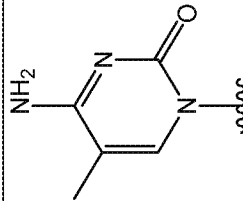
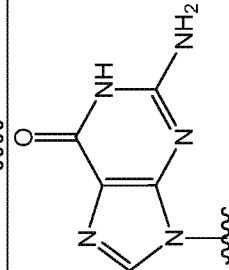
Abbreviation	Name	Structure
A	Adenine	
C	Cytosine	
(5m)C	5-methyl- cytosine	
G	Guanine	

FIGURE 6B (continued)

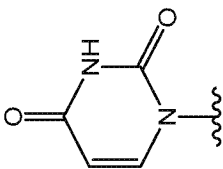
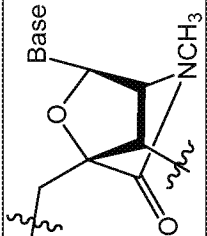
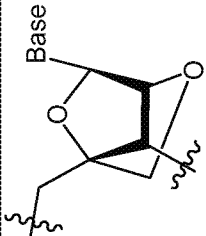
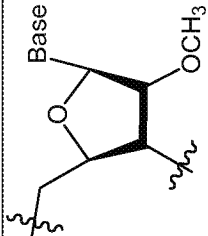
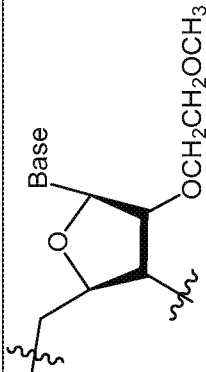
Abbreviation	Name	Structure
U	Uracil	
am	AmNA	
In	LNA	
m	2'-OMe	
moe	2'-MOE	

FIGURE 6B (continued)

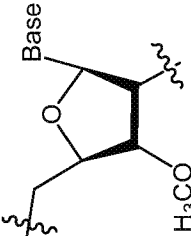
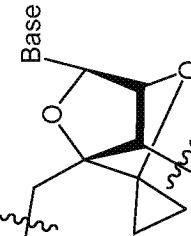
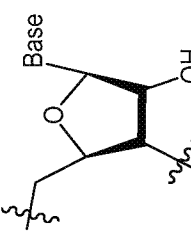
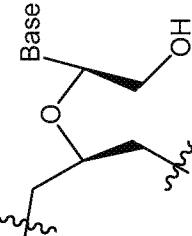
Abbreviation	Name	Structure
3m	3'-OMe	
cp	scp-BNA	
r	RNA	
un	acyclic	

FIGURE 6B (continued)

Abbreviation	Name	Structure
gn	GNA	
po	Phosphate	
ps	Phosphorothioate	
psR	Phosphorothioate with (R)-stereochemistry	
psS	Phosphorothioate with (S)-stereochemistry	
3'-cholesterol (CHOL)		

FIGURE 6B (continued)

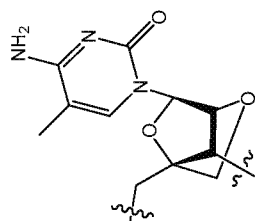
Abbreviation	Name	Structure
Tocopherol		
TREB-ps-c3-O-C3		
GalNAc5		
EP	5'-Ethyl phosphonate endcap	

FIGURE 6B (continued)

Abbreviation	Name	Structure
P		
VP		
4'-Me	4'-Me-2-OMe-A	
3cal	3'-C-allyl-A	
4etl		

FIGURE 6B (continued)

For example, using the abbreviations and structures from Table A, one skilled in the art understands that the abbreviation



“In(5m)C” represents the following:

FIGURE 7

Preparation of (((E)-2-((2R,3R,4R,5R)-5-(6-benzamido-9H-purin-9-yl)-3-(((2-cyanoethoxy)(diisopropylamino)phosphino)oxy)-4-methoxytetrahydrofuran-2-yl)vinyl)phosphoryl)bis(oxy))bis(methylene) bis(2,2-dimethylpropanoate) (5VP)

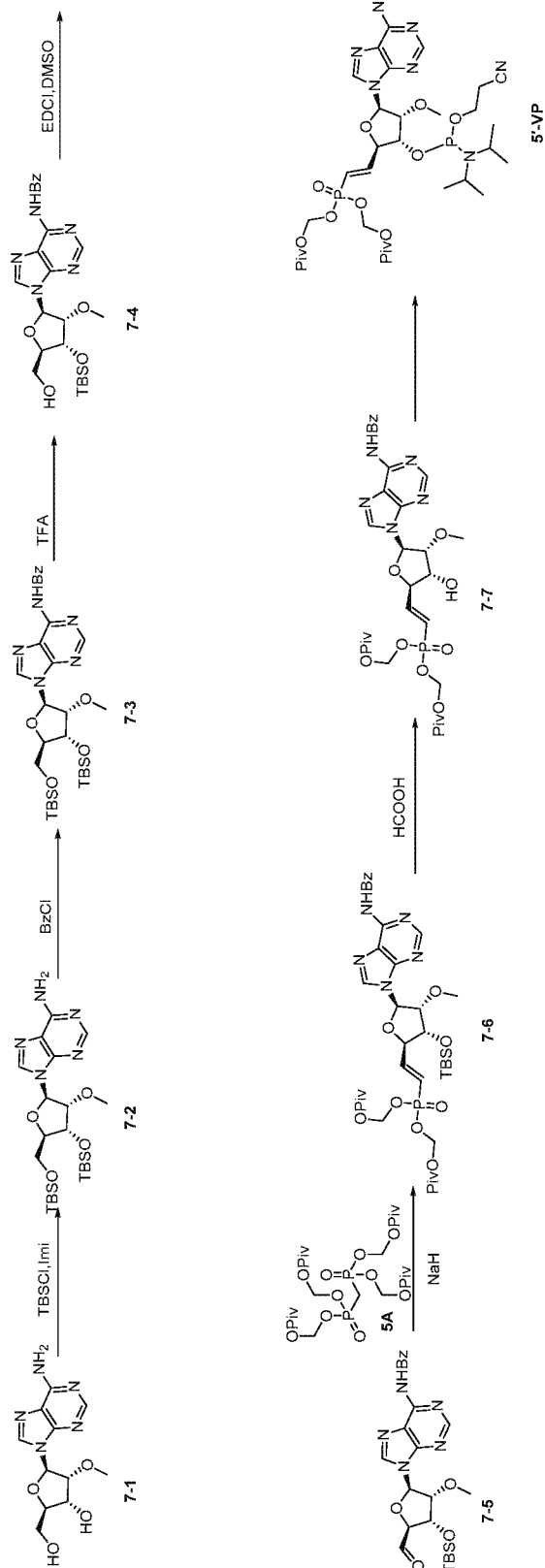


FIGURE 8

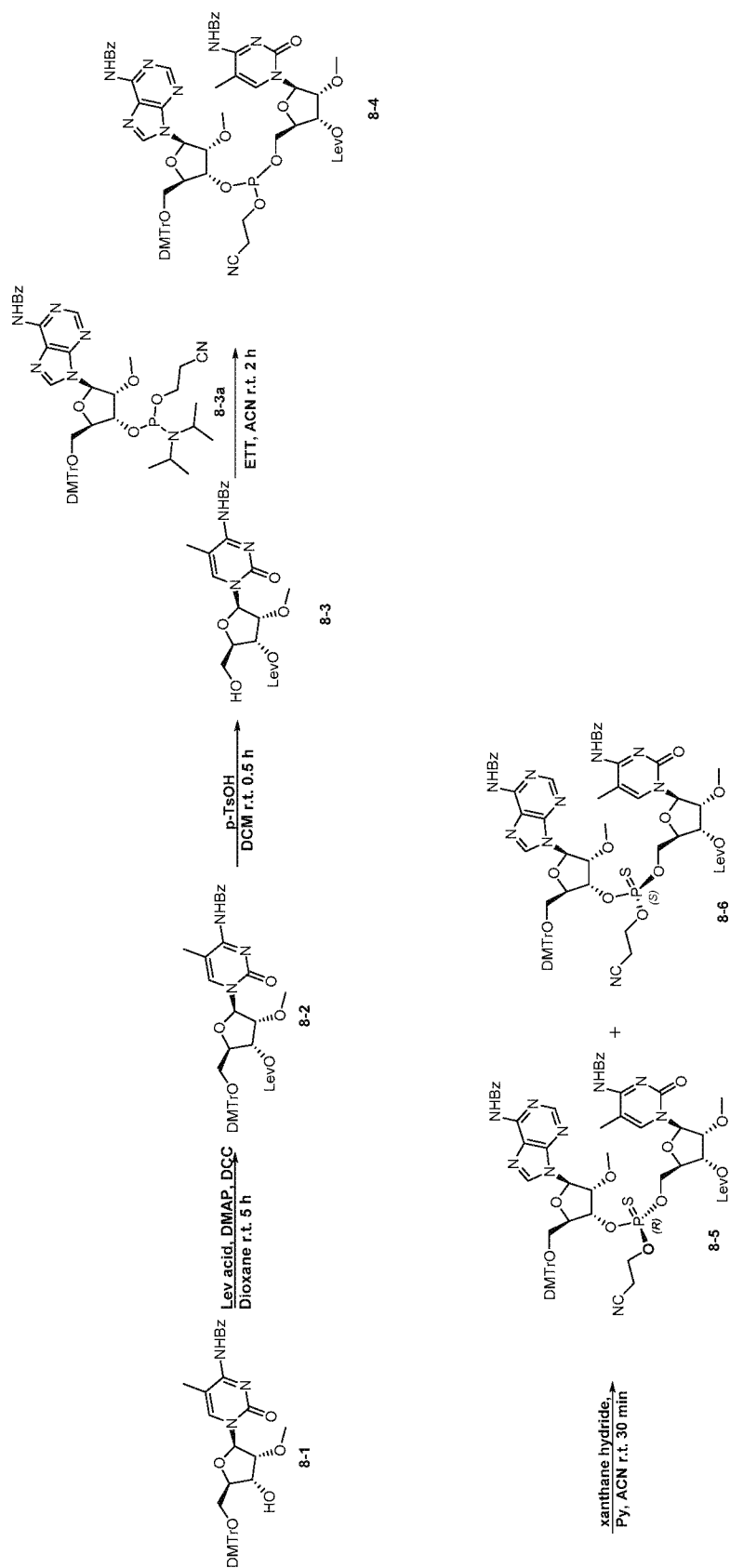


FIGURE 9A

Preparation of (2R,3R,4R,5R)-5-(4-benzamido-5-methyl-2-oxypyrimidin-1(2H)-yl)-2-(((R)-(((2R,3R,4R,5R)-5-(6-benzamido-9H-purin-9-yl)-2-((bis(4-methoxyphenyl)(phenyl)methoxy)methyl)-4-methoxytetrahydrofuran-3-yl)oxy)(2-cyanoethoxy)phosphorothioyl)oxy)methyl)-4-methoxytetrahydrofuran-3-yl(2-cyanoethyl)diisopropylphosphoramidite (9R)

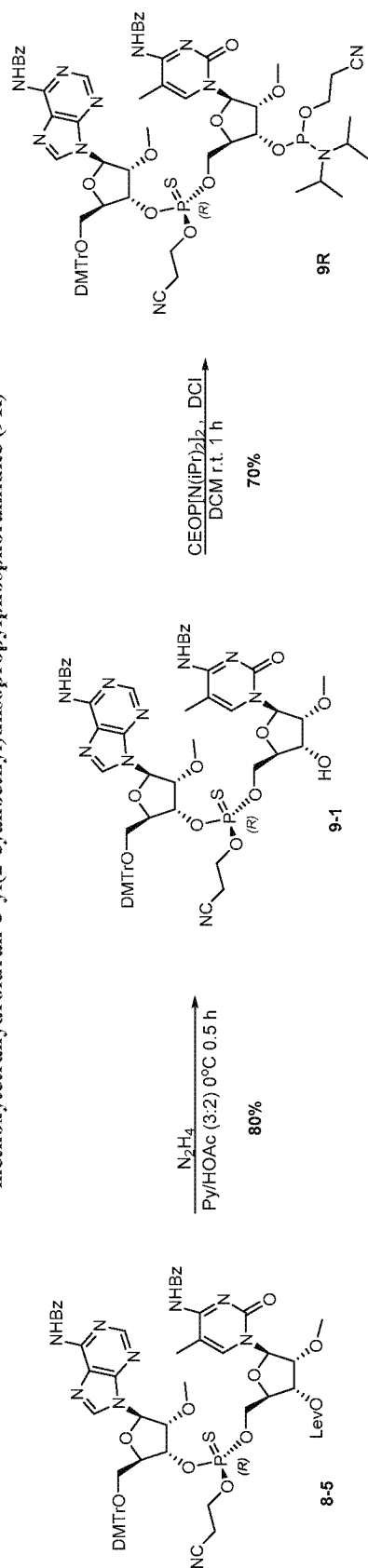


FIGURE 9B

Preparation of (2R,3R,4R,5R)-5-(4-benzamido-5-methyl-2-oxypyrimidin-1(2H)-yl)-2-(((S)-(((2R,3R,4R,5R)-5-(6-benzamido-9H-purin-9-yl)-2-((bis(4-methoxyphenyl)(phenyl)methoxy)methyl)-4-methoxytetrahydrofuran-3-yl)oxy)(2-cyanoethoxy)phosphorothioyl)oxy)methyl)-4-methoxytetrahydrofuran-3-yl(2-cyanoethyl)diisopropylphosphoramidite (9S)

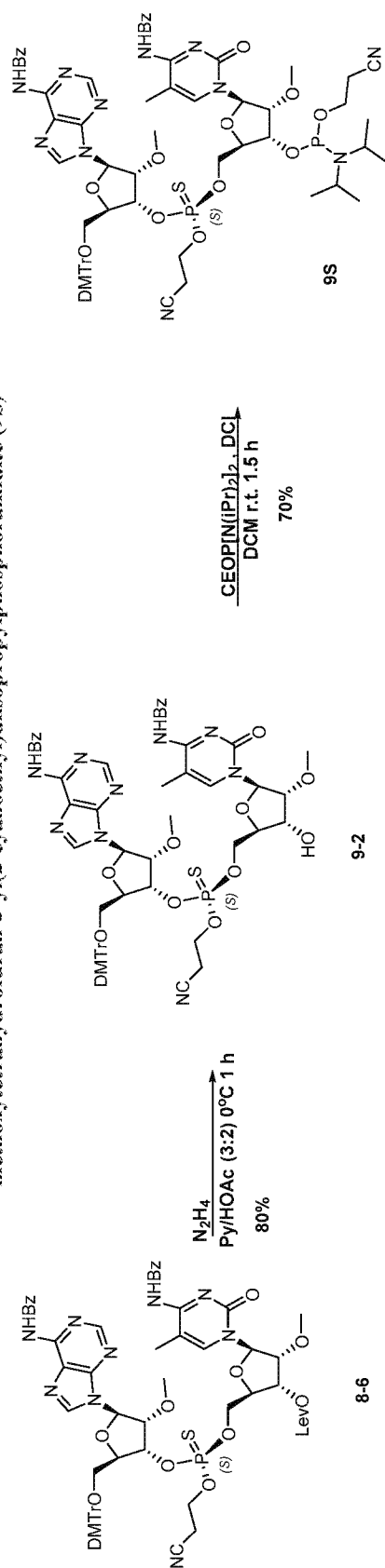


FIGURE 10

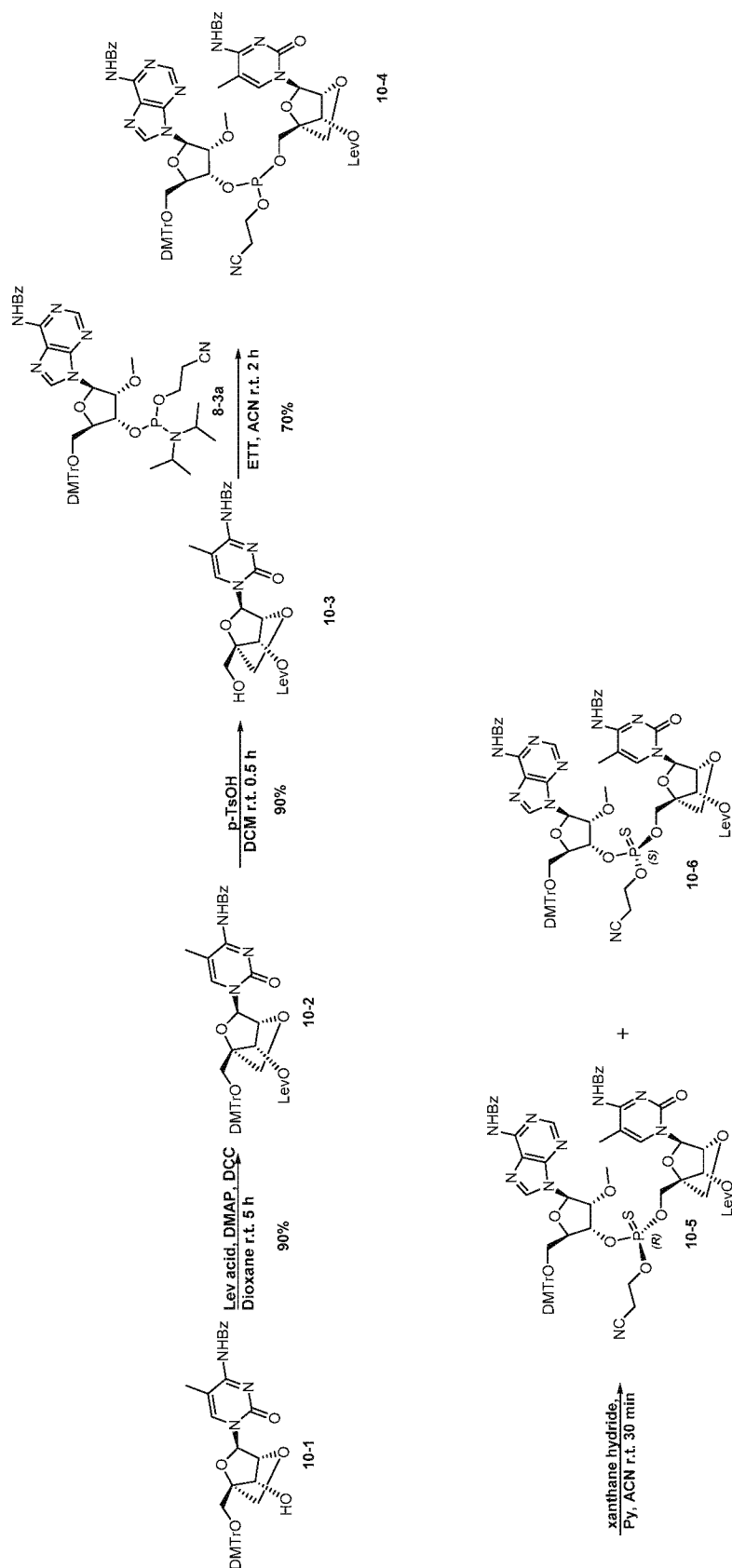


FIGURE 11A

Preparation of (1R,3R,4R,7S)-3-(4-benzamido-5-methyl-2-oxopyrimidin-1(2H)-yl)-1-(((1R)-(((2R,3R,4R,5R)-5-(6-benzamido-9H-purin-9-yl)-2-((bis(4-methoxyphenyl)(phenyl)methoxy)methyl)-4-methoxytetrahydrofuran-3-yl)oxy)(2-cyanoethoxy)phosphorothioyl)oxy)methyl)-2,5-dioxabicyclo[2.2.1]heptan-7-yl (2-cyanoethyl) diisopropylphosphoramidite (11R)

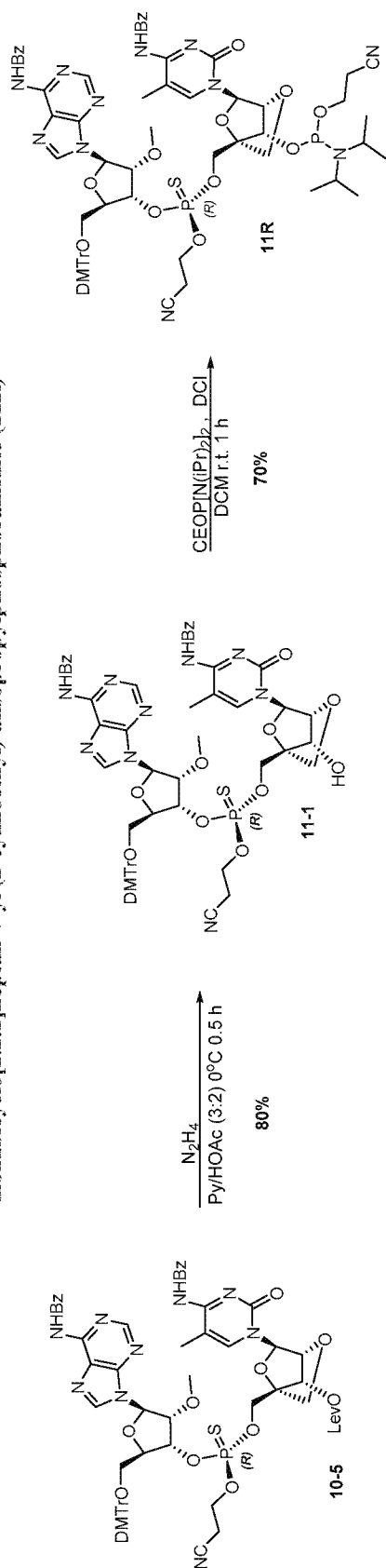
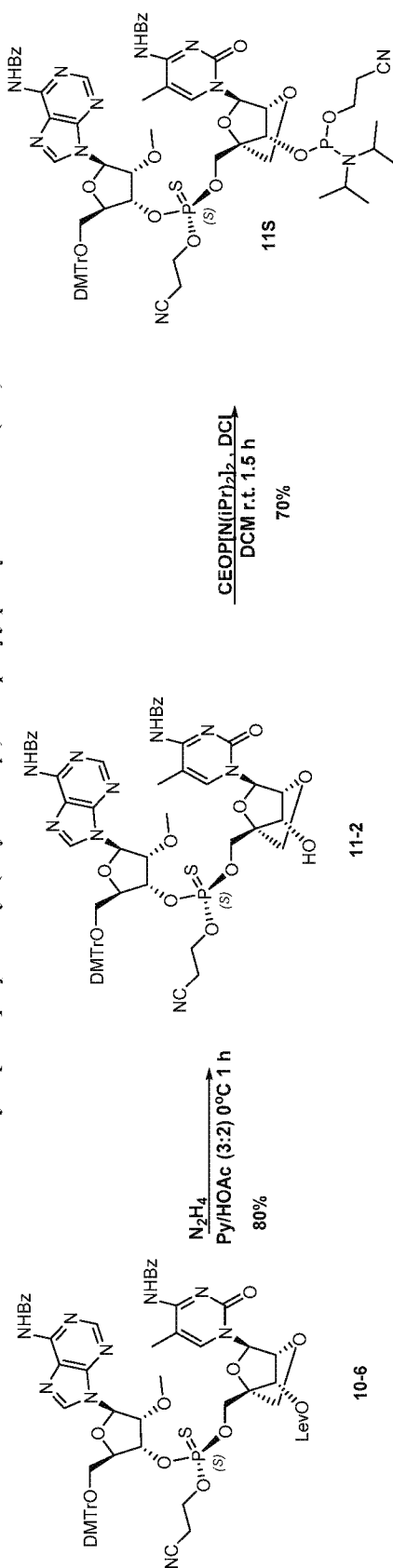
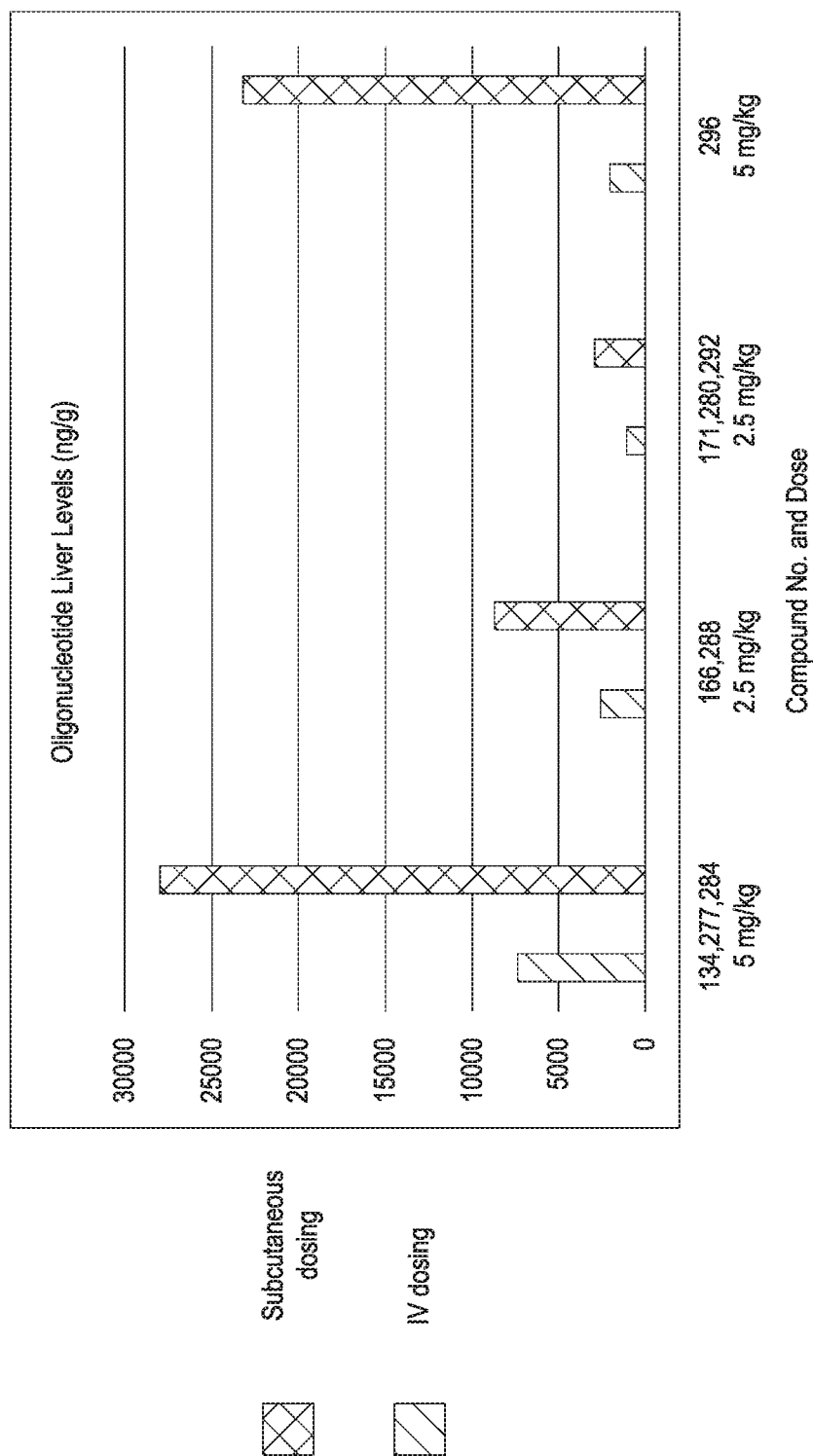


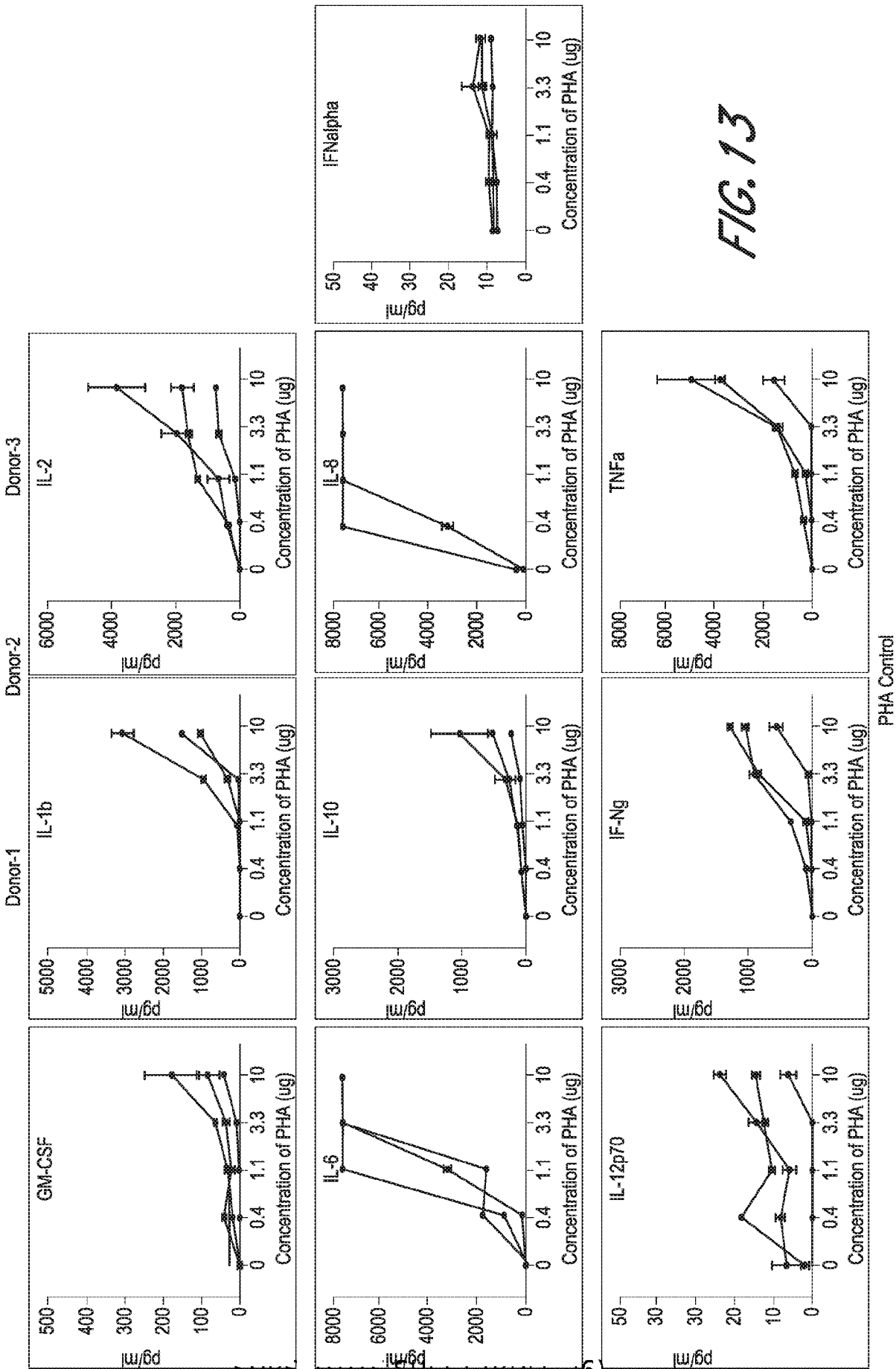
FIGURE 11B

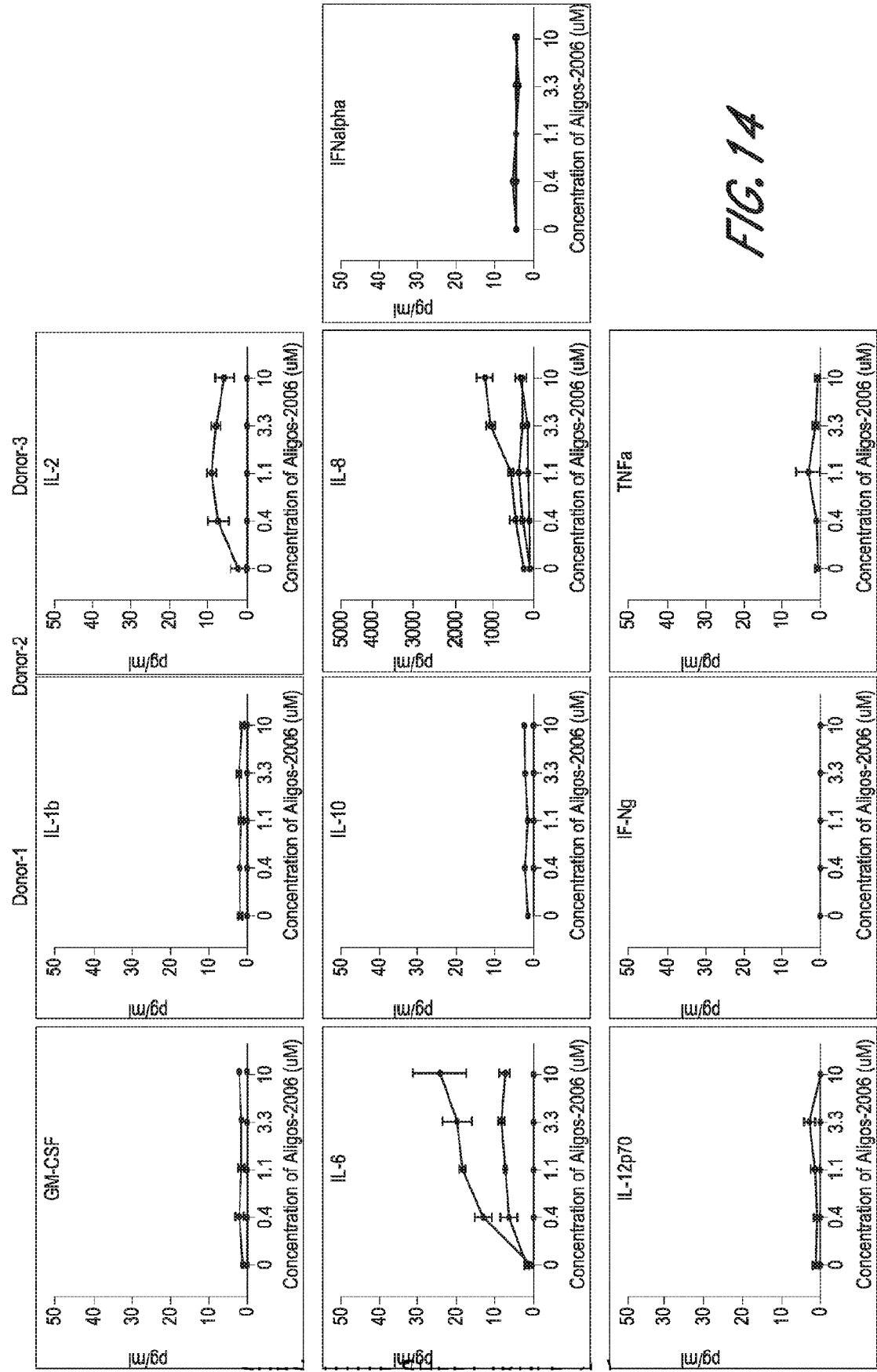
Preparation of (1R,3R,4R,7S)-3-(4-benzamido-5-methyl-2-oxopyrimidin-1(2H)-yl)-1-(((1S)-(((2R,3R,4R,5R)-5-(6-benzamido-9H-purin-9-yl)-2-((bis(4-methoxyphenyl)(phenyl)methoxy)methyl)-4-methoxytetrahydrofuran-3-yl)oxy)(2-cyanoethoxy)phosphorothioyl)oxy)methyl)-2,5-dioxabicyclo[2.2.1]heptan-7-yl (2-cyanoethyl) diisopropylphosphoramidite (11S)

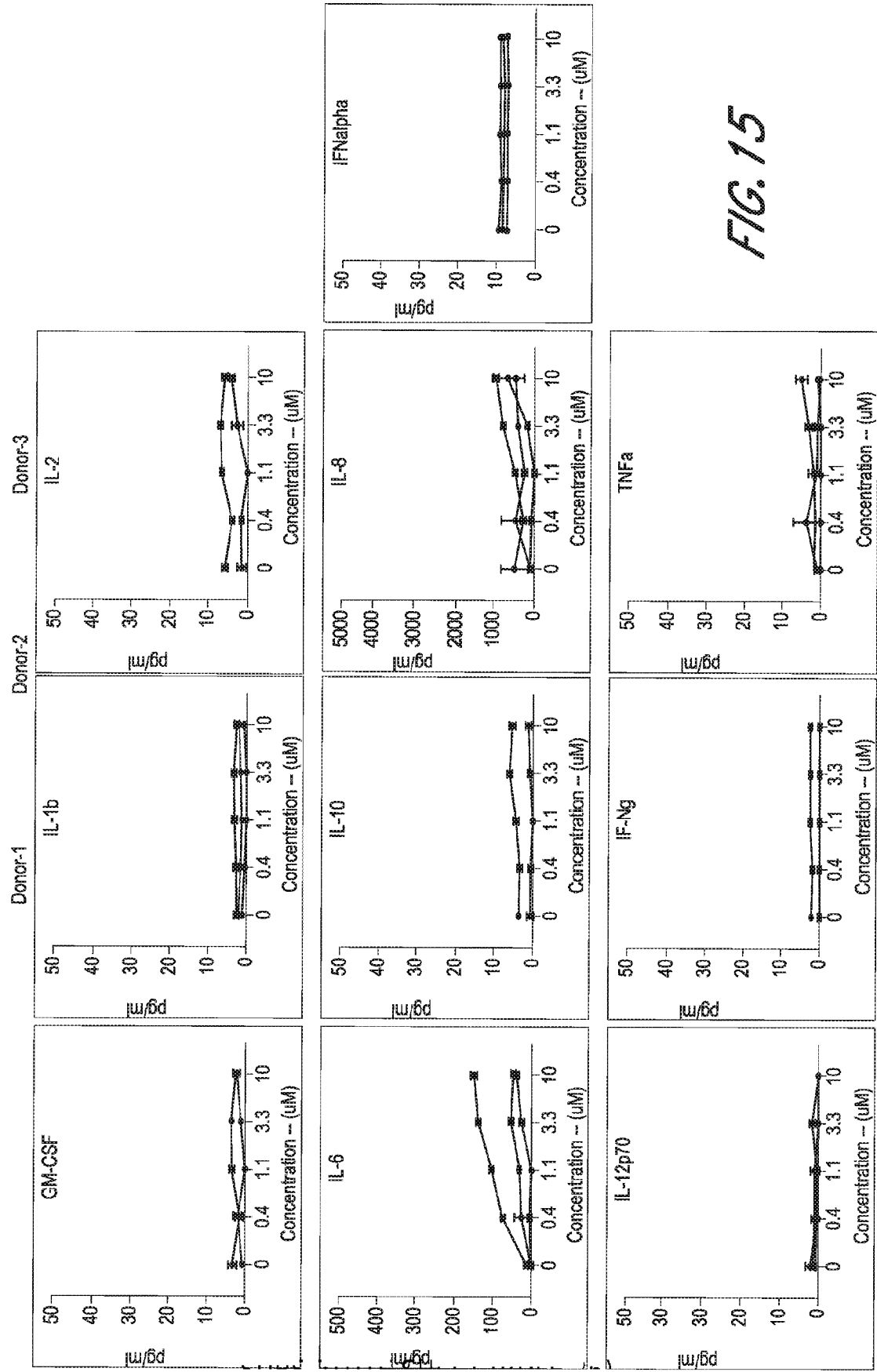




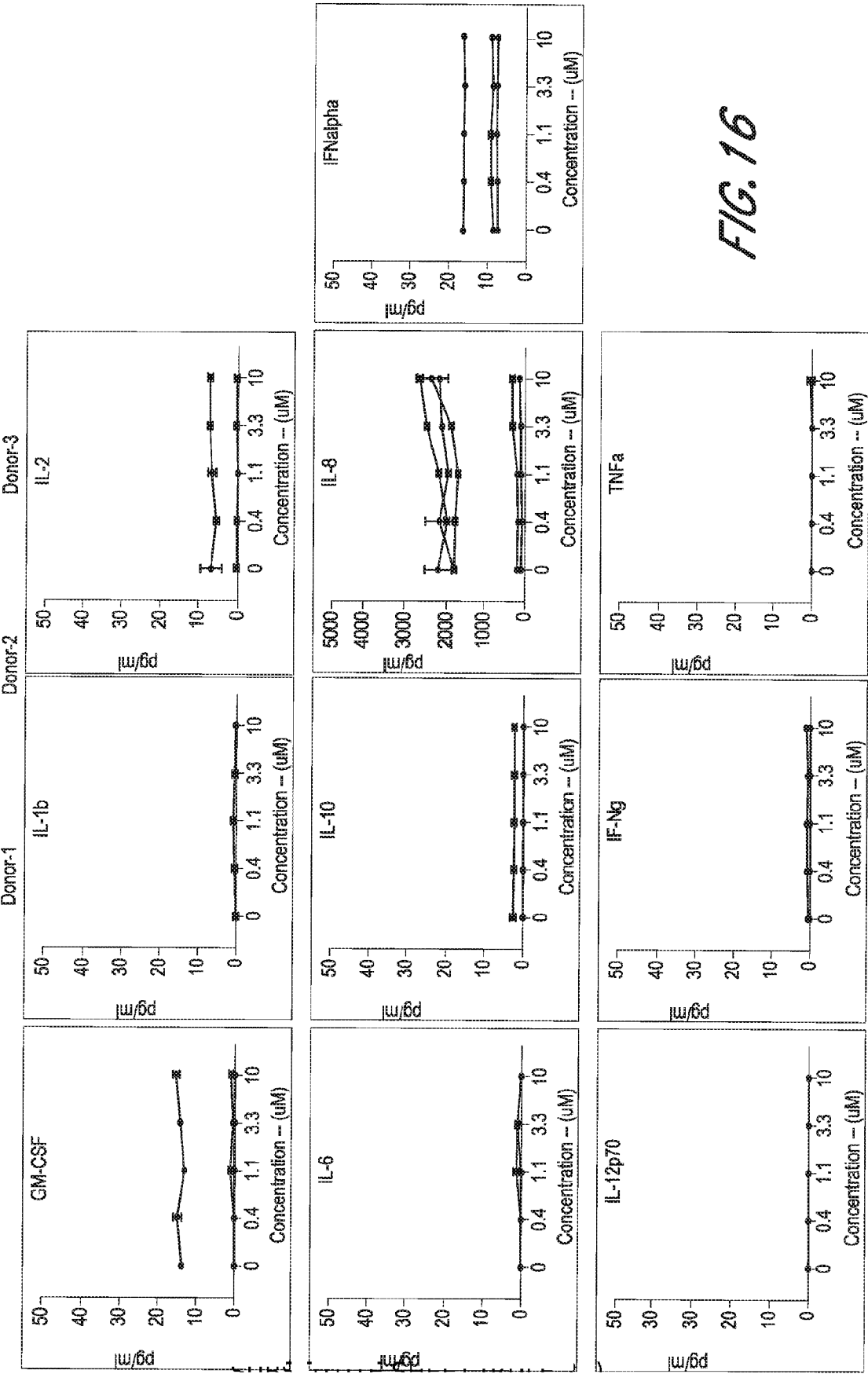
Compound No. (dose)	134,277,284 5 mg/kg		166,288 2.5 mg/kg		171,280,292 2.5 mg/kg		296 5 mg/kg	
Route	IV	SC	IV	SC	IV	SC	IV	SC
Liver Levels (ng/g)	7350.5	27,942.5	2603	8711	1024.5	2938.5	1995.5	23,191.5



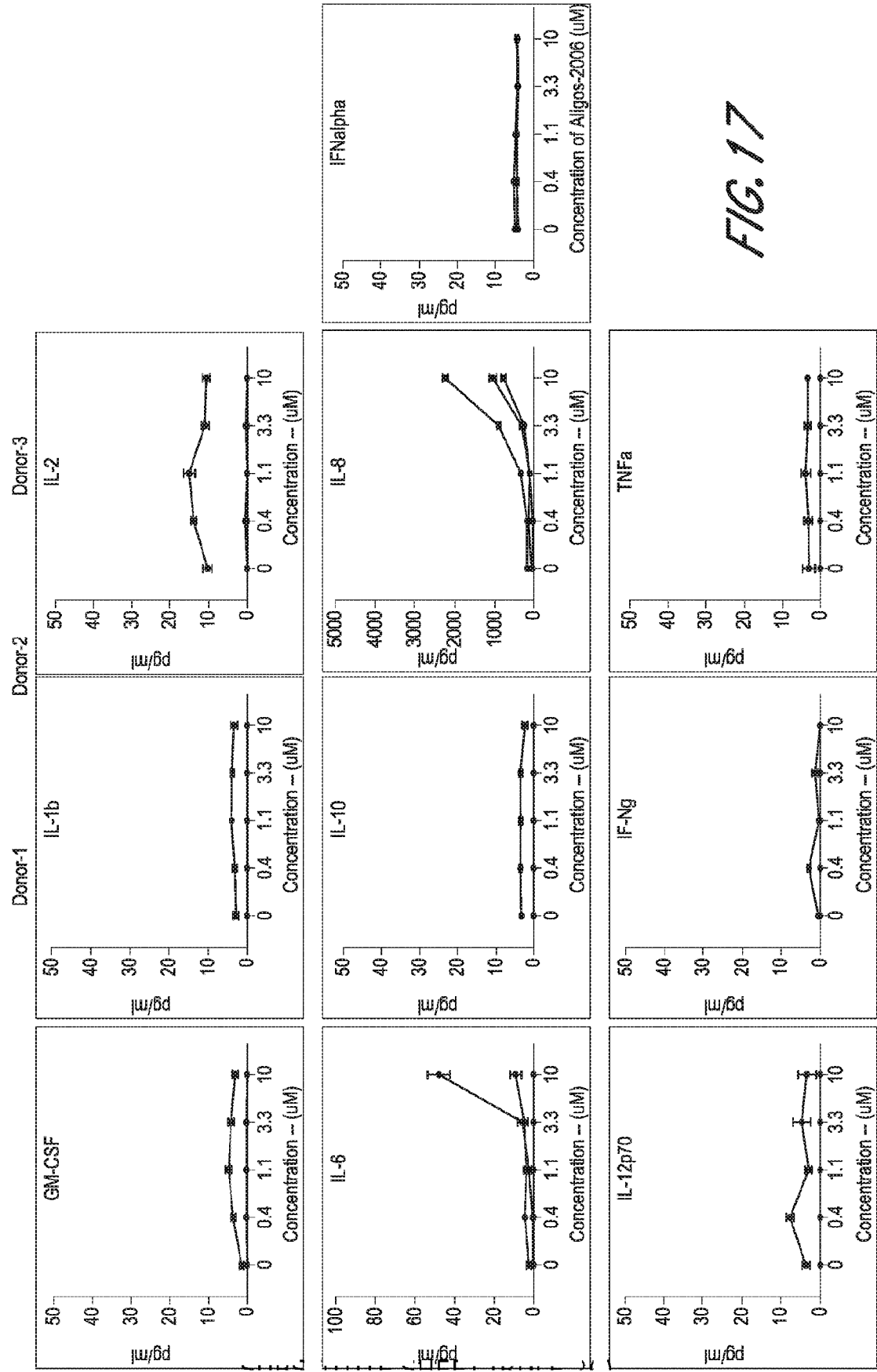




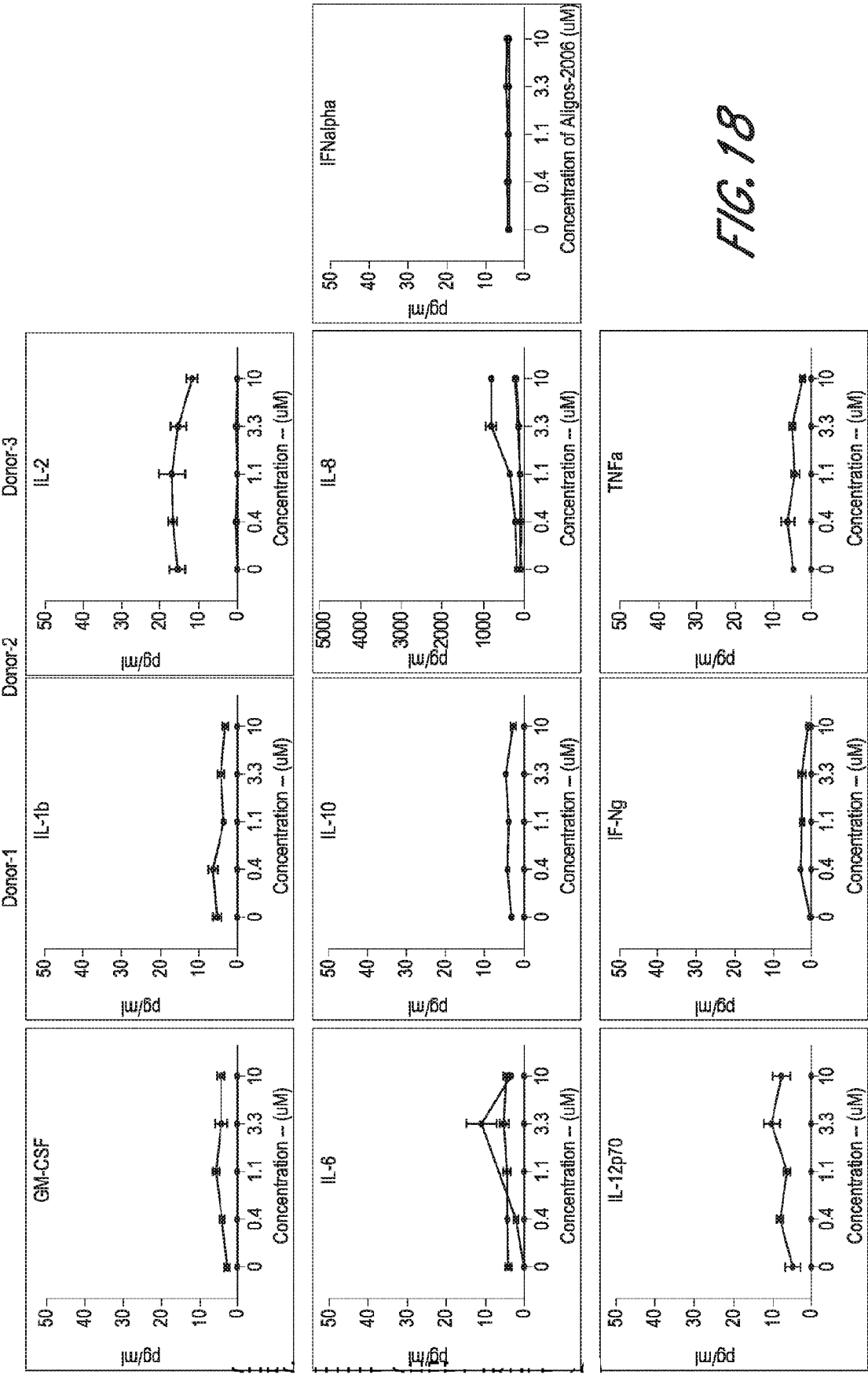
Compound Nos 1 171,280,292



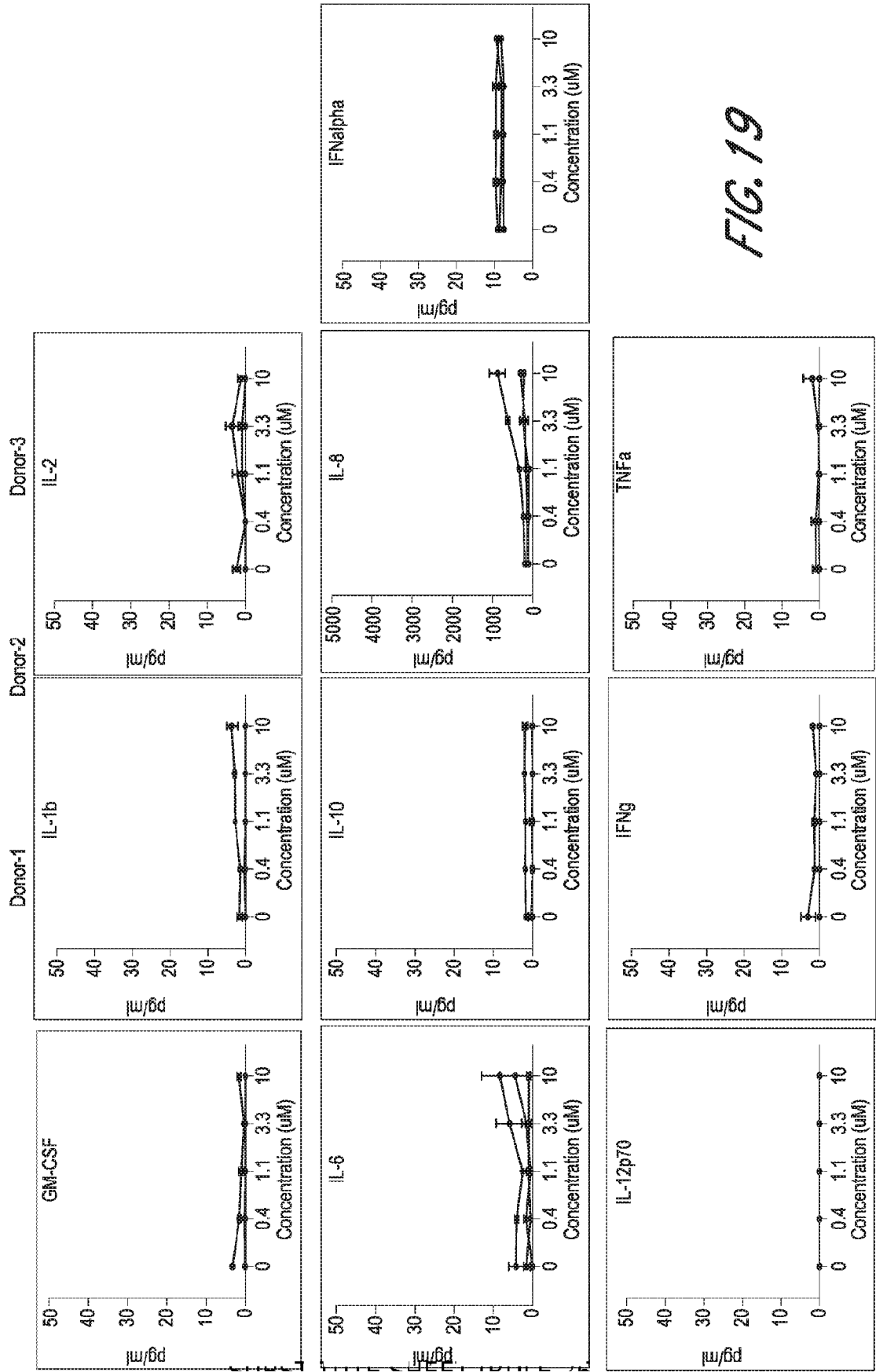
Compound Nos 1 171,280,292



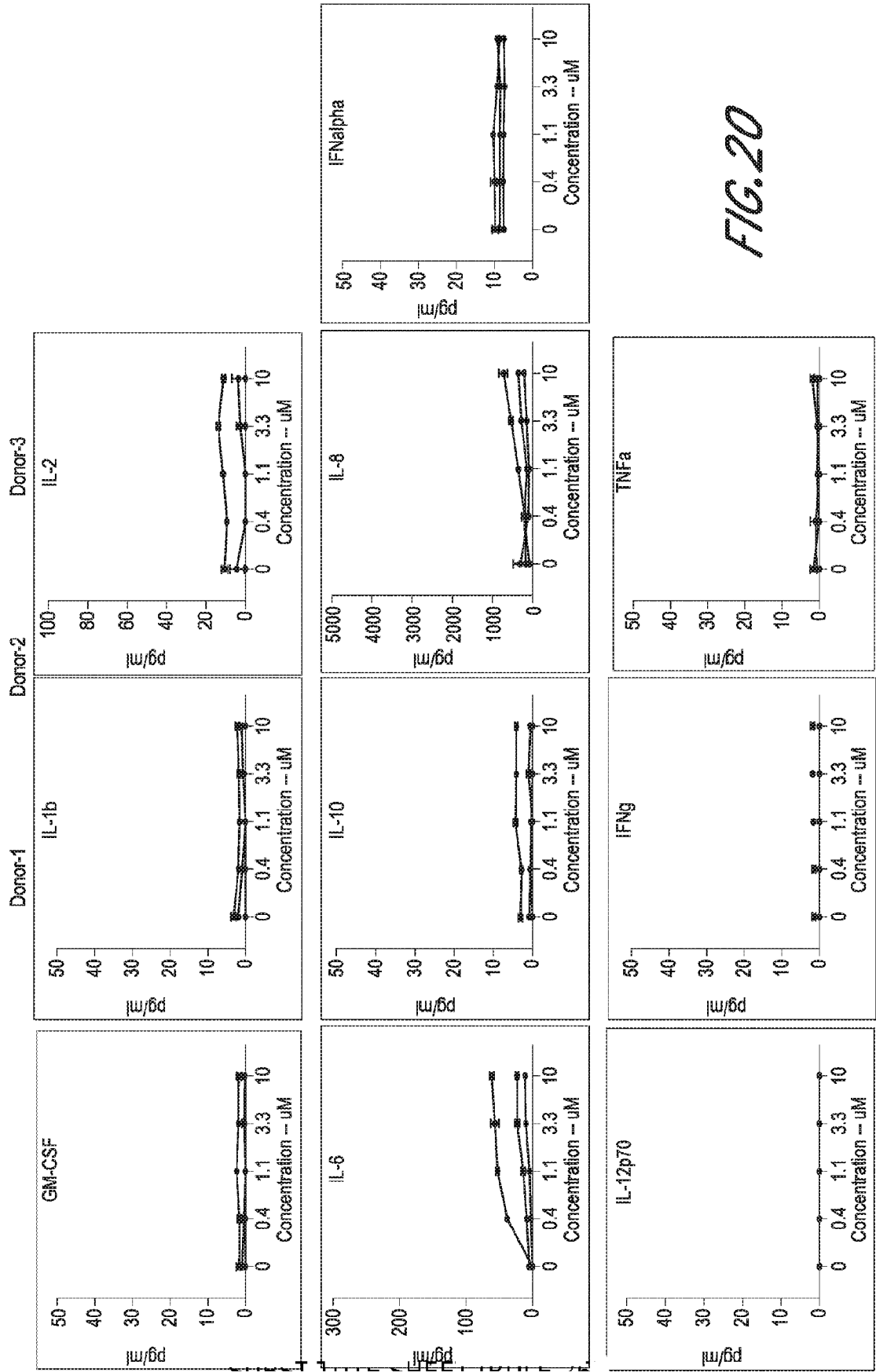
Compound Nos. 134, 277, 284

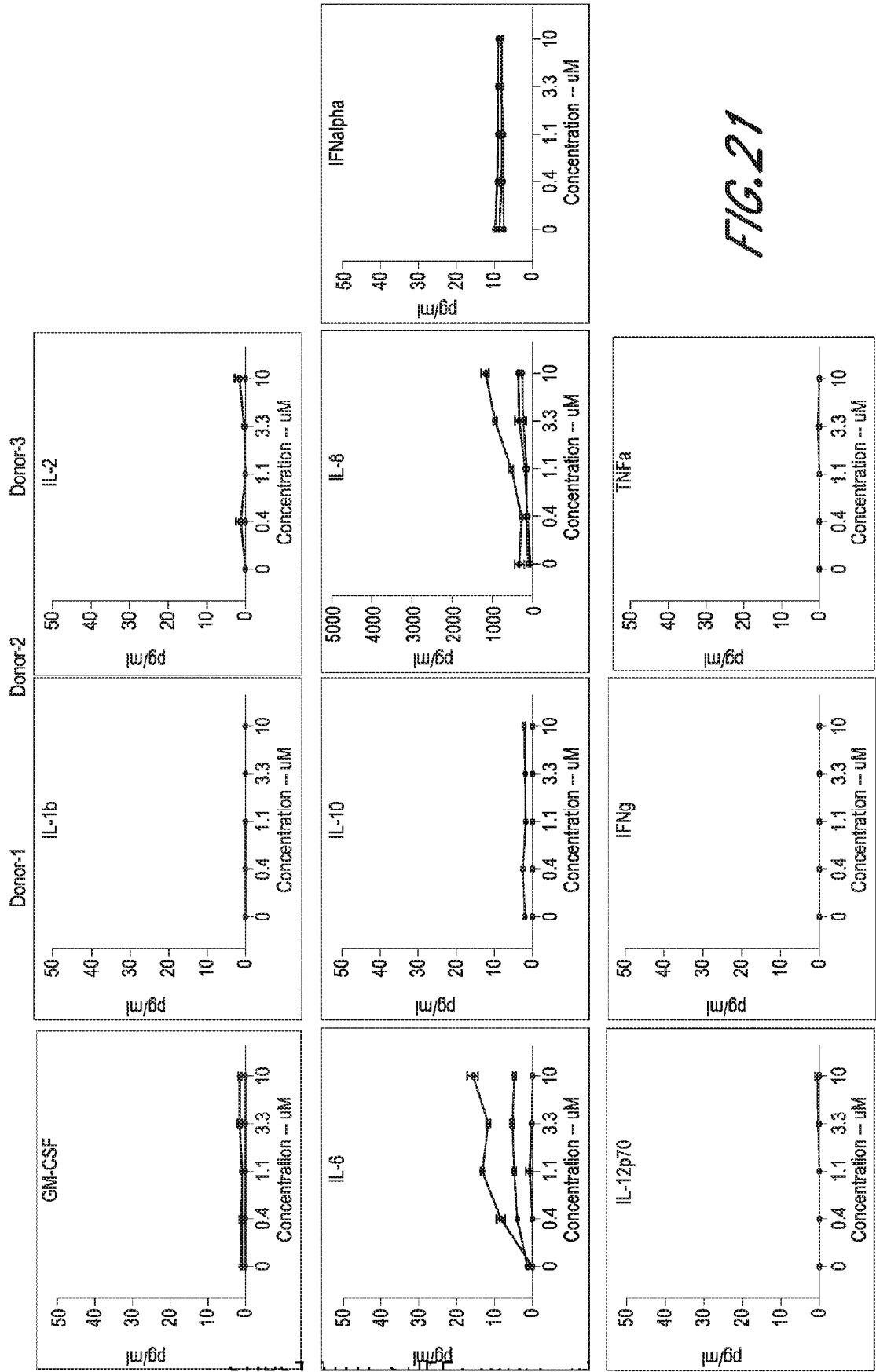


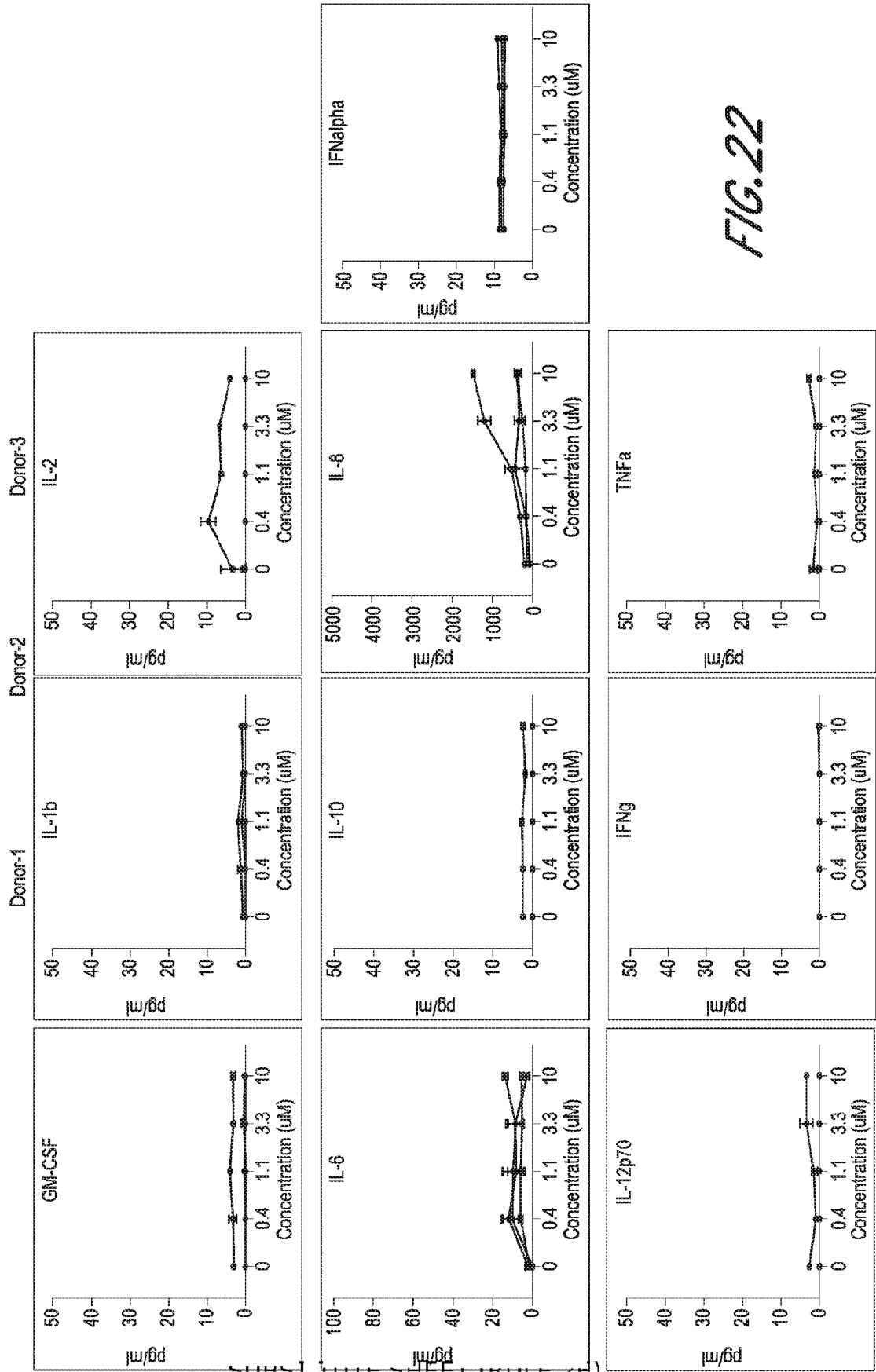
Compound Nos. 166, 288



Compound Nos. 167, 289

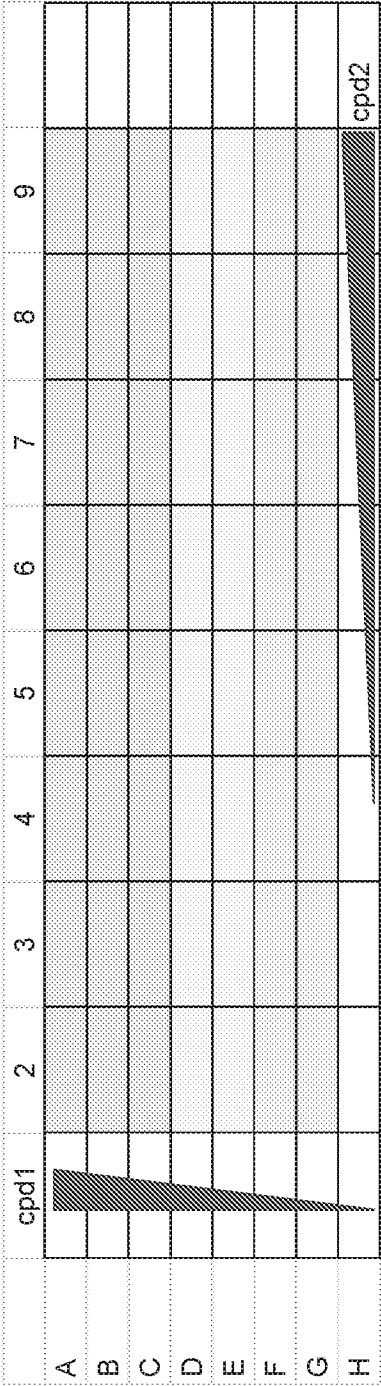






Compound Nos. 276, 291

FIGURE 23



INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2019/060283

A. CLASSIFICATION OF SUBJECT MATTER

C07H 21/02 (2006.01) C07H 21/04 (2006.01) A61K 31/712 (2006.01) A61K 31/7125 (2006.01) A61P 1/16 (2006.01) A61P 31/20 (2006.01)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

PATENTSCOPE, ESPACENET, GOOGLE: Applicant/ Inventor search: ALIGOS THERAPEUTICS, ALIOS BIOPHARMA, BEIGELMAN, Leonid, PANDEY, Rajendra, RAJWANSHI, Vivek, SMITH, David Bernard, BLATT, Lawrence, HONG, Jin; keywords: HBV, antiviral, viral, hepatitis, oligonucleotide, nucleoside, and like terms.; REGISTRY: 1858310-67-4, 1858310-70-9, 740974-33-8, 1858310-68-5, 740974-33-8, 2242457-89-0, 1492109-69-9, 1492109-75-7, 1858310-69-6, 1858310-71-0, 1858310-72-1; CAPLUS, EMBASE, BIOSIS, PATENW: keywords: REP 2139, REP 2055, REP 2154, REP 2033, REP 2153, REP 2164, REP 2166, REP 2167, NUCLEIC ACID POLYMER, MODIFIED, OLIGONUCLEOTIDE, RIBO, DEOXYRIBO, 2-OME, 2-MOE, LNA, 2-O-PROPARGYL, 2-F, 2-ARAF, 2-METHOXY, 2-FLUORO, 3-OME, UNA, 2-NH₂, GNA, ENA, 2-O-BUTYNYL, BNA, AMNA, NMLNA, 4ETL, ANALOG, 2140-79-6, 2140-72-9, A61K31/513, A61K31/52, A61K31/712, A61K31/7125, A61K31/513, A61K31/52, A61P31/12, A61P31/20, C12N15/1131, C12N15/113, HEPATITIS, HBV, HDV, NON SEQUENCE SPECIFIC, SEQUENCE INDEPENDENT, SIRNA, ANTISENSE, INHIBIT, RNA, A61K31/712, A61K31/7125, HOMOLOG, COMPLEMENT, ANTIVIRAL, HEPATITIS, HBV, HDV, HBSAG, HBEAG and like terms

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
	Documents are listed in the continuation of Box C	

☒ Further documents are listed in the continuation of Box C

☒ See patent family annex

* Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"D" document cited by the applicant in the international application	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier application or patent but published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
17 February 2020

Date of mailing of the international search report
17 February 2020

Name and mailing address of the ISA/AU

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INTERNATIONAL SEARCH REPORT		International application No.
C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		PCT/US2019/060283
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2016/004525 A1 (REPLICOR INC.) 14 January 2016 abstract; [0012]-[0018]; [0041]-[0048]; [0069]-[0072]; [0088], table 2; and claims	1-56, 58-98 & 100-104
X	ROEHL, I. ET AL., "Nucleic Acid Polymers with Accelerated Plasma and Tissue Clearance for Chronic Hepatitis B Therapy", Molecular Therapy: Nucleic Acids, 2017, Vol. 8, pages 1-12. whole document, in particular, abstract, introduction, table 1- REP 2139, REP, 2154, REP 2155, REP 2156, REP 2157, REP 2164 and REP 2165, and REP 2055, and results	1-56, 58-98 & 100-104
X	VAILLANT, A., "Nucleic acid polymers: Broad spectrum antiviral activity, antiviral mechanisms and optimization for the treatment of hepatitis B and hepatitis D infection", Antiviral Research, 2016, Vol. 133, pages 32-40 whole document, in particular, figure 1 and table 1	1-56, 58-98 & 100-104
X	BEILSTEIN, F. ET AL., "Nucleic Acid Polymers Are Active against Hepatitis Delta Virus Infection In Vitro," Journal of Virology, 2018, Vol. 92 (4). pii: e01416-17, pages 1-13. whole document, in particular, abstract, results, table 1, and materials and methods, Synthesis of NAPs	1-56, 58-98 & 100-104
X	US 2013/0309201 A1 (REPLICOR INC.) 21 November 2013 [0011]-[0015]; [0264]; [0270]; [0310]; [0320]- [0322]; [0325]; [0330]; [0335]; and claims 1, 2, 12 and 15-21	1-56, 58-98 & 100-104
A	KHVOROVA, A. ET AL., "The Chemical Evolution of Oligonucleotide Therapies of Clinical Utility", Nature Biotechnology, 2017, Vol. 35 (3), pages 238-248 whole document	1-56, 58-98 & 100-104
A	DELEAVEY, G. F. ET AL., "Designing Chemically Modified Oligonucleotides for Targeted Gene Silencing", Chemistry & Biology, 2012, Vol. 19, pages 937-954 whole document, in particular, abstract, page 943, page 945, and page 946, figure 6	1-56, 58-98 & 100-104
Form PCT/ISA/210 (fifth sheet) (July 2019)		

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
the subject matter listed in Rule 39 on which, under Article 17(2)(a)(i), an international search is not required to be carried out, including
2. ☒ Claims Nos.: **57 & 99**
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
See Supplemental Box
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a)

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2019/060283**Supplemental Box****Continuation of Box II**

The claim/s do/does not comply with Rule 6.2(a) because it/they rely on references to the description and/or drawings.

INTERNATIONAL SEARCH REPORT		International application No.	
Information on patent family members		PCT/US2019/060283	
This Annex lists known patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.			
Patent Document/s Cited in Search Report		Patent Family Member/s	
Publication Number	Publication Date	Publication Number	Publication Date
WO 2016/004525 A1	14 January 2016	WO 2016004525 A1	14 Jan 2016
		AR 101126 A1	23 Nov 2016
		AU 2015286199 A1	23 Feb 2017
		BR 112017000320 A2	07 Nov 2017
		CA 2954182 A1	14 Jan 2016
		CL 2017000008 A1	09 Jun 2017
		CN 106659730 A	10 May 2017
		DO P2017000002 A	15 Mar 2017
		EA 201790160 A1	30 Jun 2017
		EP 3166615 A1	17 May 2017
		HK 1231402 A1	22 Dec 2017
		JP 2017521433 A	03 Aug 2017
		KR 20170029577 A	15 Mar 2017
		MD 20170014 A2	31 Jul 2017
		MX 2017000053 A	01 May 2017
		PH 12017500010 A1	15 May 2017
		SG 11201700073P A	27 Feb 2017
		TW 201613617 A	16 Apr 2016
		US 2016008393 A1	14 Jan 2016
		US 9603865 B2	28 Mar 2017
US 2013/0309201 A1	21 November 2013	US 2013309201 A1	21 Nov 2013
		US 9492506 B2	15 Nov 2016
		AU 2013262416 A1	18 Dec 2014
		AU 2013262416 B2	11 May 2017
		BR 112014028654 A2	10 Oct 2017
		CA 2873529 A1	21 Nov 2013
		CL 2014003134 A1	13 Feb 2015
		CN 104349793 A	11 Feb 2015
		CN 104349793 B	10 Nov 2017
		CO 7131387 A2	01 Dec 2014
Due to data integration issues this family listing may not include 10 digit Australian applications filed since May 2001.			
Form PCT/ISA/210 (Family Annex)(July 2019)			

INTERNATIONAL SEARCH REPORT Information on patent family members		International application No. PCT/US2019/060283	
This Annex lists known patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.			
Patent Document/s Cited in Search Report		Patent Family Member/s	
Publication Number	Publication Date	Publication Number	Publication Date
		CR 20140527 A	15 Dec 2014
		DO P2014000264 A	31 Jan 2015
		EA 201401278 A1	30 Apr 2015
		EP 2849798 A1	25 Mar 2015
		HK 1204279 A1	13 Nov 2015
		IL 235548 A	31 Jul 2019
		JP 2015517504 A	22 Jun 2015
		KR 20150013309 A	04 Feb 2015
		KR 102068109 B1	21 Jan 2020
		MX 2014014021 A	10 Feb 2015
		MX 346239 B	13 Mar 2017
		NZ 703095 A	29 Jul 2016
		PH 12014502551 A1	21 Jan 2015
		PH 12014502551 B1	21 Jan 2015
		SG 11201407599S A	30 Dec 2014
		TW 201408308 A	01 Mar 2014
		TW I635864 B	21 Sep 2018
		WO 2013170386 A1	21 Nov 2013
		ZA 201408674 B	27 Jan 2016
End of Annex			
Due to data integration issues this family listing may not include 10 digit Australian applications filed since May 2001. Form PCT/ISA/210 (Family Annex)(July 2019)			