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(54) Title: ANTISENSE ANTIVIRAL COMPOUND AND METHOD FOR TREATING PICORNAVIRUS INFECTION

(57) Abstract: The invention provides antisense antiviral compounds and methods of their use and production in inhibition of growth of viruses of the Picornaviridae family and in the treatment of a viral infection. The compounds are particularly useful in the treatment of Enterovirus and/or Rhinovirus infection in a mammal. The antisense antiviral compounds are substantially uncharged, morpholino oligonucleotides have a sequence of 12-40 subunits, including at least 12 subunits having a targeting sequence that is complementary to a region associated with viral RNA sequences within a 32 nucleotide region of the viral 5' untranslated region identified by SEQ ID NO:7.



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ANTISENSE ANTIVIRAL COMPOUND AND METHOD
FOR TREATING PICORNAVIRUS INFECTION

FIELD OF THE INVENTION

This invention relates to antisense oligonucleotide compounds for use in treating a picornavirus infection and antiviral treatment methods employing the compounds.

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BACKGROUND OF THE INVENTION

The Picomaviridae represents a very large family of small RNA viruses responsible for many serious human and animal diseases (Straus and Straus, 2002). The Picomaviridae includes four major genera: *Enterovirus*, *Rhinovirus*, *Aphovirus* and *Hepatovirus*. The *Enterovirus* genus includes polioviruses, coxsackieviruses, echoviruses, and enteroviruses.

Poliovirus is the etiologic agent of the disease poliomyelitis in humans, and there are three known serotypes of the virus. The oral poliovaccine, typically given to children, is a mixture of the Sabin strain of the virus. The oral poliovirus vaccine is safe and effective, yet has two limitations. First, the vaccine is unstable since current vaccines are inactivated by relatively brief (less than 24 hours) exposure to temperatures of 37°C. This necessitates transport in a frozen state to the locale where they are administered. Second, the vaccine occasionally reverts to virulence in vaccine recipients and the reverted virulent virus may then be passed to other individuals who come into contact with the recipient in whom the vaccine has reverted.

The human rhinoviruses consist of at least 100 serotypes and are the primary causative agents of the common cold. Because of the large number of serotypes, development of a vaccine is problematic and antiviral agents may therefore be the best approach to treatment. The Coxsackie viruses and other human enteroviruses (multiple serotypes), are associated with a wide range of human diseases including summer flus, diarrhea, meningitis, hepatitis, pneumonia, myocarditis, pericarditis, and diabetes. These infections occur sporadically in the general population, but are becoming more common among children in day care and their parents and siblings. Other important members of the Picomaviridae family include human hepatitis A virus, Theiler's murine encephalomyelitis virus, foot-and-mouth disease virus, and mengovirus.

The existing drugs which are used against the viruses described above are only moderately effective, and are typically effective against only a limited subset of the rhinovirus serotypes. In general, the available drugs have either failed to demonstrate sufficient prophylactic effects or are converted in the body into inactive metabolites.

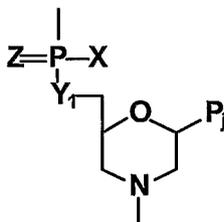
Thus, there remains a need for a more effective antiviral therapy in several members of the Picornoviridae family.

SUMMARY OF THE INVENTION

The invention includes, in one aspect, a method of inhibiting viral infection in mammalian cells by an *Enterovirus* or *Rhinovirus* in the Picomaviridae family. The method includes the steps of exposing the cells to an antisense oligonucleotide compound, thereby to form a heteroduplex structure (i) composed of the virus' positive sense strand and the oligonucleotide compound, and (ii) characterized by a T_m of dissociation of at least 45°C. The oligonucleotide compound is characterized by:

- (i) a substantially uncharged, nuclease-resistant backbone,
- (ii) capable of uptake by mammalian host cells,
- (iii) containing between 12-40 nucleotide bases, and
- (iv) having a targeting sequence of at least 12 subunits complementary to SEQ ID NO:7 in the positive-sense strand of the virus.

The compound to which the host cells are exposed may be composed of morpholino subunits and phosphorus-containing intersubunit linkages joining a morpholino nitrogen of one subunit to a 5' exocyclic carbon of an adjacent subunit. The morpholino subunits may be joined by phosphorodiamidate linkages having the structure:



where $Y_i=O$, $Z=O$, P_j is a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, and X is alkyl, alkoxy, thioalkoxy, amino or alkyl amino, including dialkylamino, e.g., wherein $X=NR_2$, where each R is independently hydrogen or methyl.

The compound may be composed of morpholino subunits linked with the uncharged linkages described above interspersed with linkages that are positively charged at physiological pH. The total number of positively charged linkages is between 2 and no more than half of the total number of linkages. The positively charged linkages have the structure above, where X is 1-piperazine.

The oligonucleotide compound to which the cells are exposed may have a sequence contained in SEQ ID NO:10, such as one of the sequences identified by SEQ ID NO:11, SEQ ID NO:12 or SEQ ID NO:13. The compound may be conjugated to an arginine-rich polypeptide effective to promote uptake of the compound into infected host cells. Exemplary polypeptides have one of the sequences identified as SEQ ID NOS:14-19.

For use in treating a mammalian subject infected by an *Enterovirus* or *Rhinovirus* in the Picornaviridae family, the compound is administered to the subject in a pharmaceutically effective amount. Compound administration may be continued until a significant reduction in viral infection or the symptoms thereof is observed. The subject may be treated with a second anti-viral compound before, after, or during treatment with the oligonucleotide compound.

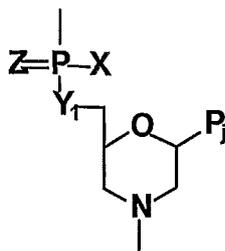
For use in treating a mammalian subject at risk of infection by an *Enterovirus* or *Rhinovirus* in the Picornaviridae family, the compound is administered to the subject in an amount effective to inhibit infection of subject host cells by the virus.

In another aspect, the invention includes an oligonucleotide compound for use in inhibiting viral infection in mammalian cells by an *Enterovirus* or *Rhinovirus* in the Picornaviridae family. The compound is characterized by:

- (i) a substantially uncharged, nuclease-resistant backbone,
- (ii) capable of uptake by mammalian host cells,
- (iii) containing between 12-40 nucleotide bases,
- (iv) having a targeting sequence of at least 12 subunits contained in SEQ ID NO:10; and

- (v) capable of binding to the virus¹ positive sense strand to form a heteroduplex structure having by a T_m of dissociation of at least 45°C.

The compound may be composed of morpholino subunits and phosphorus-containing intersubunit linkages joining a morpholino nitrogen of one subunit to a 5' exocyclic carbon of an adjacent subunit. The morpholino subunits may be joined by phosphorodiamidate linkages having the structure:



where $Y_i=O$, $Z=O$, P_j is a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, and X is alkyl, alkoxy, thioalkoxy, amino or alkyl amino, including dialkylamino, e.g., wherein $X=NR_2$, where each R is independently hydrogen or methyl.

The compound may be composed of morpholino subunits linked with the uncharged linkages described above interspersed with linkages that are positively charged at physiological pH. The total number of positively charged linkages is between 2 and no more than half of the total number of linkages. The positively charged linkages have the structure above, where X is 1-piperazine.

The oligonucleotide compound may have one of the sequences identified by SEQ ID NOS:1 1, 12, and 13. The compound may be conjugated to an arginine-rich polypeptide effective to promote uptake of the compound into infected host cells. Exemplary polypeptides have one of the sequences identified as SEQ ID NOS.-14-19.

The compound may be formulated in combination with another anti-viral compound.

These and other objects and features of the invention will become more fully apparent when the following detailed description of the invention is read in conjunction with the accompanying drawings.

BRIEF DESCRIPTION OF THE FIGURES

Figures 1A-1 D show the repeating subunit segment of exemplary morpholino oligonucleotides, designated A through D.

Figures 2A-2G show examples of uncharged linkage types in oligonucleotide analogs. Figure 2H is an example of a preferred charged, cationic linkage.

Figure 3 shows the sequence conservation across a broad spectrum of picomaviruses for the 5'-32 nt region represented by SEQ ID NOS:1-6, and the combined sequence identified by SEQ ID NO:7.

Figure 4 shows a genetic map of the dual luciferase reporter construct pR&Fluc-PoliolRES.

Figure 5 shows the synthetic steps to produce subunits used to produce +PMO containing the (1-piperazino) phosphinylideneoxy cationic linkage as shown in Fig. 2H.

Figures 6A and 6B show results of cell-free translation assays using exemplary antisense compounds of the invention and their associated EC50 concentrations.

DETAILED DESCRIPTION OF THE INVENTION

I. Definitions

The terms below, as used herein, have the following meanings, unless indicated otherwise:

The term "oligonucleotide analog" refers to an oligonucleotide having (i) a modified backbone structure, e.g., a backbone other than the standard phosphodiester linkage found in natural oligo- and polynucleotides, and (ii) optionally, modified sugar moieties, e.g., morpholino moieties rather than ribose or deoxyribose moieties. The analog supports bases capable of hydrogen bonding by Watson-Crick base pairing to standard polynucleotide bases, where the analog backbone presents the bases in a manner to permit such hydrogen bonding in a sequence-specific fashion between the oligonucleotide analog molecule and bases in a standard polynucleotide {e.g., single-stranded RNA or single-stranded DNA}. Preferred analogs are those having a substantially uncharged, phosphorus containing backbone. An oligonucleotide analog is also referred to herein as an oligonucleotide or oligonucleotide compound or oligonucleotide analog compound.

A substantially uncharged, phosphorus containing backbone in an oligonucleotide analog is one in which a majority of the subunit linkages, e.g., between 50-100%, typically at least 80% of its linkages, are uncharged at physiological pH, and contain a single phosphorous atom. The analog contains between 8 and 40 subunits, typically about 8-25 subunits, and preferably about 12

to 25 subunits. The analog may have exact sequence complementarity to the target sequence or near complementarity, as defined below.

A "subunit" of an oligonucleotide analog refers to one nucleotide (or nucleotide analog) unit of the analog. The term may refer to the nucleotide unit with or without the attached intersubunit linkage, although, when referring to a "charged subunit", the charge typically resides within the intersubunit linkage (e.g. a phosphate or phosphorothioate linkage).

A "morpholino oligonucleotide analog" is an oligonucleotide analog composed of morpholino subunit structures of the form shown in Figs. 1A-1 D where (i) the structures are linked together by phosphorus-containing linkages, one to three atoms long, joining the morpholino nitrogen of one subunit to the 5' exocyclic carbon of an adjacent subunit, and (ii) P_i and P_j are purine or pyrimidine base-pairing moieties effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide. The purine or pyrimidine base-pairing moiety is typically adenine, cytosine, guanine, uracil or thymine. The synthesis, structures, and binding characteristics of morpholino oligomers are detailed in U.S. Patent Nos. 5,698,685, 5,217,866, 5,142,047, 5,034,506, 5,166,315, 5,521,063, and 5,506,337, all of which are incorporated herein by reference in their entirety.

The subunit and linkage shown in Figure 1B are used for six-atom repeating-unit backbones, as shown in Figure 1B (where the six atoms include: a morpholino nitrogen, the connected phosphorus atom, the atom (usually oxygen) linking the phosphorus atom to the 5' exocyclic carbon, the 5' exocyclic carbon, and two carbon atoms of the next morpholino ring). In these structures, the atom Y_i linking the 5' exocyclic morpholino carbon to the phosphorus group may be sulfur, nitrogen, carbon or, preferably, oxygen. The X moiety pendant from the phosphorus is any stable group which does not interfere with base-specific hydrogen bonding. Preferred X groups include fluoro, alky), alkoxy, thioalkoxy, and alkyl amino, including cyclic amines, all of which can be variously substituted, as long as base-specific bonding is not disrupted. Alkyl, alkoxy and thioalkoxy preferably include 1-6 carbon atoms. Alkyl amino preferably refers to lower alkyl (C_1 to C_6) substitution, and cyclic amines are preferably 5- to 7-membered nitrogen heterocycles optionally containing 1-2 additional heteroatoms selected from oxygen, nitrogen, and sulfur. Z is sulfur or oxygen, and is preferably oxygen.

A preferred morpholino oligomer is a phosphorodiamidate-linked morpholino oligomer, referred to herein as a PMO. Such oligomers are composed of morpholino subunit structures such as shown in Fig. 2B, where $X = \text{NH}_2$, NHR , or NR_2 (where R is lower alkyl, preferably methyl), $Y = \text{O}$, and $Z = \text{O}$, and P_i and P_j are purine or pyrimidine base-pairing moieties effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, as seen in Fig. 2G. Also preferred are morpholino oligomers where the phosphordiamidate linkages are uncharged linkages as shown in Fig 2G interspersed with cationic linkages as shown in Fig. 2H where, in Fig. 2B, $X = 1\text{-piperazino}$. In another Fig. 2B embodiment, $X = \text{lower alkoxy}$, such as methoxy or ethoxy, $Y = \text{NH}$ or NR , where R is lower alkyl, and $Z = \text{O}$.

The term "substituted", particularly with respect to an alkyl, alkoxy, thioalkoxy, or alkylamino group, refers to replacement of a hydrogen atom on carbon with a heteroatom-containing substituent, such as, for example, halogen, hydroxy, alkoxy, thiol, alkylthio, amino, alkylamino, imino, oxo (keto), nitro, cyano, or various acids or esters such as carboxylic, sulfonic, or phosphonic. It may also refer to replacement of a hydrogen atom on a heteroatom (such as an amine hydrogen) with an alkyl, carbonyl or other carbon containing group.

As used herein, the term "picornavirus" refers to one or more viral species belonging to the Picomaviridae family and specifically the *Enterovirus* and *Rhinovirus* genera of the Picomaviridae.

As used herein, the term "target" refers to a viral genomic RNA, and specifically, to a region identified by SEQ ID NO:7 within the δ' -untranslated region (5'-UTR) of the positive-sense RNA strand of a member of the Picomaviridae described herein.

The term "target sequence" refers to a portion of the target RNA against which the oligonucleotide analog is directed, that is, the sequence to which the oligonucleotide analog will hybridize by Watson-Crick base pairing of a complementary sequence.

The term "targeting sequence" is the sequence in the oligonucleotide analog that is complementary (meaning, in addition, substantially complementary) to the target sequence in the RNA genome. The entire sequence, or only a portion, of the analog compound may be complementary to the target sequence. For example, in an analog having 20 bases, only 12-14 may be targeting

sequences. Typically, the targeting sequence is formed of contiguous bases from the analog, but may alternatively be formed of non-contiguous sequences that when placed together, e.g., from opposite ends of the analog, constitute a sequence that spans the target sequence.

Target and targeting sequences are described as "complementary" to one another when hybridization occurs in an antiparallel configuration. A targeting sequence may have "near" or "substantial" complementarity to the target sequence and still function for the purpose of the present invention, that is, still be "complementary." Preferably, the oligonucleotide analog compounds employed in the present invention have at most one mismatch with the target sequence out of 10 nucleotides, and preferably at most one mismatch out of 20. Alternatively, the antisense oligomers employed have at least 90% sequence homology, and preferably at least 95% sequence homology, with the exemplary targeting sequences as designated herein. For purposes of complementary binding to an RNA target, and as discussed below, a guanine base may be complementary to either an adenine or uracil RNA base.

An oligonucleotide analog "specifically hybridizes" to a target polynucleotide if the oligomer hybridizes to the target under physiological conditions, with a T_m substantially greater than 45°C, preferably at least 50°C, and typically 60°C-80°C or higher. Such hybridization preferably corresponds to stringent hybridization conditions. At a given ionic strength and pH, the T_m is the temperature at which 50% of a target sequence hybridizes to a complementary polynucleotide. Again, such hybridization may occur with "near" or "substantial" complementarity of the antisense oligomer to the target sequence, as well as with exact complementarity.

A "nuclease-resistant" oligomeric molecule (oligomer) refers to one whose backbone is substantially resistant to nuclease cleavage, in non-hybridized or hybridized form; by common extracellular and intracellular nucleases in the body; that is, the oligomer shows little or no nuclease cleavage under normal nuclease conditions in the body to which the oligomer is exposed.

A "heteroduplex" refers to a duplex between an oligonucleotide analog and the complementary portion of a target RNA. A "nuclease-resistant heteroduplex" refers to a heteroduplex formed by the binding of an antisense oligomer to its complementary target, such that the heteroduplex is substantially resistant to *in*

vivo degradation by intracellular and extracellular nucleases, such as RNAse H, which are capable of cutting double-stranded RNA/RNA or RNA/DNA complexes.

A "base-specific intracellular binding event involving a target RNA" refers to the specific binding of an oligonucleotide analog to a target RNA sequence inside a cell. The base specificity of such binding is sequence specific. For example, a single-stranded polynucleotide can specifically bind to a single-stranded polynucleotide that is complementary in sequence.

An "effective amount" of an antisense oligomer, targeted against an infecting picomavirus, is an amount effective to reduce the rate of replication of the infecting virus, and/or viral load, and/or symptoms associated with the viral infection.

As used herein, the term "body fluid" encompasses a variety of sample types obtained from a subject including, urine, saliva, plasma, blood, spinal fluid, or other sample of biological origin, such as skin cells or dermal debris, and may refer to cells or cell fragments suspended therein, or the liquid medium and its solutes.

The term "relative amount" is used where a comparison is made between a test measurement and a control measurement. The relative amount of a reagent forming a complex in a reaction is the amount reacting with a test specimen, compared with the amount reacting with a control specimen. The control specimen may be run separately in the same assay, or it may be part of the same sample (for example, normal tissue surrounding a malignant area in a tissue section).

"Treatment" of an individual or a cell is any type of intervention provided as a means to alter the natural course of the individual or cell. Treatment includes, but is not limited to, administration of e.g., a pharmaceutical composition, and may be performed either prophylactically, or subsequent to the initiation of a pathologic event or contact with an etiologic agent. The related term "improved therapeutic outcome" relative to a patient diagnosed as infected with a particular virus, refers to a slowing or diminution in the growth of virus, or viral load, or detectable symptoms associated with infection by that particular virus.

An agent is "actively taken up by mammalian cells" when the agent can enter the cell by a mechanism other than passive diffusion across the cell membrane. The agent may be transported, for example, by "active transport", referring to transport of agents across a mammalian cell membrane by e.g. an

ATP-dependent transport mechanism, or by "facilitated transport", referring to transport of antisense agents across the cell membrane by a transport mechanism that requires binding of the agent to a transport protein, which then facilitates passage of the bound agent across the membrane. For both active and facilitated transport, the oligonucleotide analog preferably has a substantially uncharged backbone, as defined below. Alternatively, the antisense compound may be formulated in a complexed form, such as an agent having an anionic backbone complexed with cationic lipids or liposomes, which can be taken into cells by an endocytotic mechanism. The analog also may be conjugated, e.g., at its 5' or 3' end, to an arginine-rich peptide, e.g., a portion of the HIV TAT protein, polyarginine, or combinations of arginine and other amino acids including the non-natural amino acids 6-aminohexanoic acid and beta-alanine. Exemplary arginine-rich delivery peptides are listed as SEQ ID NOS: 14-19. These exemplary arginine-rich delivery peptides facilitate transport into the target host cell as described (Moulton, Nelson *et al.* 2004; Nelson, Stein *et al.* 2005).

Rules for the selection of targeting sequences capable of inhibiting replication of picomaviruses are discussed below.

II. Targeted Viruses

The present invention is based on the discovery that effective inhibition of members of the Picomaviridae family can be achieved with antisense oligonucleotide analog compounds that (i) target the region identified by SEQ ID NO:7 of the 5' untranslated region (UTR) of the virus¹ positive strand, and (ii) have physical and pharmacokinetic features which allow effective interaction between the antisense compound and the virus within host cells. In one aspect, the oligomers can be used in treating a mammalian subject infected with the virus.

The invention targets single stranded, positive sense RNA viruses that are members of the Picomaviridae family. In particular, targeted viruses include members of the *Enterovirus* and *Rhinovirus* genera of the Picomaviridae family. Table 1 lists the viruses targeted by the invention as organized by genus. Various physical, morphological, and biological characteristics of members of the Picomaviridae family can be found, for example, in Textbook of Human Virology, R. Belshe, ed., 2nd Edition, Mosby, 1991, at the Universal Virus Database of the International Committee on Taxonomy of Viruses

(www.ncbi.nlm.nih.gov/ICTVdb/index.htm) and in human virology textbooks (e.g., see Strauss and Strauss, 2002). Some of the key biological characteristics of the Picornaviridae family of viruses are described below.

Table 1. Targeted Viruses of the Invention

Family	Genus	Virus
Picornaviridae	<i>Enterovirus</i>	Poliovirus (PV)
		Human enterovirus A (HEV-A)
		Human enterovirus B (HEV-B)
		Human enterovirus C (HEV-C)
		Human enterovirus D (HEV-D)
	<i>Rhinovirus</i>	Human Rhinovirus A (HRV-89)
		Human Rhinovirus B (HRV-14)

Picomaviruses

GenBank reference entries for exemplary viral nucleic acid sequences representing picomavirus genomic RNA are listed in Table 2 below. The nucleotide sequence numbers in Table 2 are derived from the Genbank reference for the positive-strand RNA. It will be appreciated that these sequence references are only illustrative of other sequences in the Picornaviridae family, as may be available from available gene-sequence databases or literature or patent resources.

Table 2 lists the targets for a 32-base sequence corresponding to nucleotides 443-474 of the poliovirus reference sequence (NC 002058) and contained in the 5' UTR region of several picoriviruses. All the viruses listed in Table 2 are human isolates and are organized into the *Enterovirus* and *Rhinovirus* genera as Human Enteroviruses A-D, Poliovirus, Rhinovirus A and Rhinovirus B according to convention as provided by the International Committee on Taxonomy of Viruses (ICTV). (e.g., see www.ncbi.nlm.nih.gov/ICTVdb/ICTVdB/index.htm). The target sequences (SEQ ID NOS:1-6) are in the Sequence Listing table at the end of the specification.

An important feature of the present invention is the high degree of sequence conservation between viruses in the two genera, *Enterovirus* and *Rhinovirus*, as shown in Figure 3. The prototypic member of the Picornaviridae family is poliovirus and the targeting sequences (described below) are made in reference to the poliovirus sequence. Table 2 lists the corresponding target

regions in a number of clinically relevant Enteroviruses (*Enterovirus* Surveillance - United States, 200-2001 . MMWR 2002;51 :1047-1 049.) and *Rhinoviruses*. The target homologies for the target region is shown in Figure 3. The target sequence identified as SEQ ID NO:7 represents a combined target sequence, where the letter "Y" in the sequence represents a pyrimidine base, *i.e.*, may be either C or T in SEQ ID NOS:1-6.

Targeting sequences are designed to hybridize to a region of the target sequence as listed in Table 3. Selected targeting sequences can be made shorter, *e.g.*, 12 bases, or longer, *e.g.*, 40 bases, and include a small number of mismatches, as long as the sequence is sufficiently complementary to hybridize with the target, and forms with the virus positive-strand, a heteroduplex having a T_m of 45°C or greater.

More generally, the degree of complementarity between the target and targeting sequence is sufficient to form a stable duplex. The region of complementarity of the antisense oligomers with the target RNA sequence may be as short as 8-11 bases, but is preferably 12-15 bases or more, *e.g.* 12-20 bases, or 12-25 bases. An antisense oligomer of about 14-15 bases is generally long enough to have a unique complementary sequence in the viral genome. In addition, a minimum length of complementary bases may be required to achieve the requisite binding T_m , as discussed below.

Oligomers as long as 40 bases may be suitable, where at least a minimum number of bases, *e.g.*, 12 bases, are complementary to the target sequence. In general, however, facilitated or active uptake in cells is optimized at oligomer lengths less than about 30, preferably less than 25. For PMO oligomers, described further below, an optimum balance of binding stability and uptake generally occurs at lengths of 15-22 bases.

Table 2. Exemplary Human Picornavirus Nucleic Acid Target Sequences

Virus	Ref. No.	GB No.	Region	SEQ ID NO
5'-32 Nucleotide Target Region				
Poliovirus-Mahoney strain	NC 002058	V01149	443-474	1
Enterovirus A (CV-A16)	NC 001612	U05876	452-483	1
Enterovirus 71 (HEV-71)		U22521	448-479	2
Enterovirus B (CV-B1)	NC 001472	M16560	446-477	2
Coxsackievirus B3 (CV-B3)		M88483	447-478	2
Coxsackievirus B2 (CV-B2)		AF081485	448-479	2
Coxsackievirus B4 (CV-B4)		X05690	448-479	2
Coxsackievirus B5 (CV-B5)		X67706	448-479	2
Coxsackievirus A9 (CV-A9)		D00627	448-479	1
Echovirus 4 (EV-4)		X89534	331-362	2
Echovirus 6 (EV-6)		U16283	446-477	3
Echovirus 11 (EV-11)		X80059	449-480	4
Echovirus 13 (EV-13)		AF412361	259-290	2
Echovirus 18 (EV-18)		AF412366	259-290	2
Echovirus 25 (EV-25)		AY302549	466-477	4
Enterovirus C (CV-A21)	NC 001428	D00538	441-472	2
Enterovirus D (HEV-70)	NC 001430	D00820	446-477	1
Rhinovirus A (HRV-89)	NC 001617	M16248	442-473	5
Rhinovirus B (HRV-14)	NC 001490	K02121	453-484	6

The oligomer may be 100% complementary to the viral nucleic acid target sequence, or it may include mismatches, e.g., to accommodate variants, as long as a heteroduplex formed between the oligomer and viral nucleic acid target sequence is sufficiently stable to withstand the action of cellular nucleases and other modes of degradation which may occur *in vivo*. Oligomer backbones which are less susceptible to cleavage by nucleases are discussed below. Mismatches, if present, are less destabilizing toward the end regions of the hybrid duplex than in the middle. The number of mismatches allowed will depend on the length of the oligomer, the percentage of G:C base pairs in the duplex, and the position of the mismatch(es) in the duplex, according to well understood principles of duplex stability. Although such an antisense oligomer is not necessarily 100% complementary to the viral nucleic acid target sequence, it is effective to stably and specifically bind to the target sequence, such that a biological activity of the nucleic acid target, e.g., expression of viral protein(s), is modulated.

The oligomer may also incorporate guanine bases in place of adenine when the target nucleotide is a uracil residue. This is useful when the target sequence varies across different viral species and the variation at any given nucleotide residue is either cytosine or uracil. By utilizing guanine in the targeting oligomer at the position of variability, the well-known ability of guanine to base pair with uracil (termed C/U:G base pairing) can be exploited. By incorporating guanine at these locations, a single oligomer can effectively target a wider range of RNA target variability. An example of this is shown in Table 3 below wherein the 5'-32b antisense targeting oligomer (SEQ ID NO:9) has guanine incorporated at positions 7 and 10 compared to the 5'-32a antisense targeting oligomer that has adenine residues at these positions. Comparison of the 5'-32b sequence with the 5'-32 target sequences (SEQ ID NOS: 1-6), demonstrates the broad range of complementarity that C/U:G base pairing provides. Specifically, guanine residues at positions 4, 5, 7, 11 and 14 in the 5'-32b targeting oligomer provide complete complementarity across all the picornavirus 5'-32 target sequences (SEQ ID NOS: 1-6) listed in Table 2. Although the target sequences shown in Figure 3 and listed in Table 2 contain T for thymidine, which is the convention for sequence listings, it will be appreciated that because picornaviruses are RNA viruses, the T residues refer to uracil.

The stability of the duplex formed between the oligomer and the target sequence is a function of the binding T_m and the susceptibility of the duplex to cellular enzymatic cleavage. The T_m of an antisense compound with respect to complementary-sequence RNA may be measured by conventional methods, such as those described by Hames *et al.*, *Nucleic Acid Hybridization*, IRL Press, 1985, pp.107-108 or as described in Miyada CG. and Wallace R.B., 1987, *Oligonucleotide hybridization techniques, Methods Enzymol.* Vol. 154 pp. 94-107. Each antisense oligomer should have a binding T_m , with respect to a complementary-sequence RNA, of greater than body temperature and preferably greater than 50°C. T_m 's in the range 60-80°C or greater are preferred. According to well known principles, the T_m of an oligomer compound, with respect to a complementary-based RNA hybrid, can be increased by increasing the ratio of C:G paired bases in the duplex, and/or by increasing the length (in base pairs) of the heteroduplex. At the same time, for purposes of optimizing cellular uptake, it may be advantageous to limit the size of the oligomer. For this reason,

compounds that show high T_m (50°C or greater) at a length of 20 bases or less are generally preferred over those requiring greater than 20 bases for high T_m values.

The antisense activity of the oligomer may be enhanced by using a mixture of uncharged and cationic phosphorodiamidate linkages as shown in Figures 2G and 2H. The total number of cationic linkages in the oligomer can vary from 1 to 10, and be interspersed throughout the oligomer. Preferably, the number of charged linkages is at least 2 and no more than half the total backbone linkages, e.g., between 2-8 positively charged linkages, and preferably each charged linkages is separated along the backbone by at least one, preferably at least two uncharged linkages. The antisense activity of various oligomers can be measured *in vitro* by fusing the oligomer target region to the 5' end a reporter gene (e.g. firefly luciferase) and then measuring the inhibition of translation of the fusion gene mRNA transcripts in cell free translation assays. The inhibitory properties of oligomers containing a mixture of uncharged and cationic linkages can be enhanced between, approximately, five to 100 fold in cell free translation assays.

Table 3 below shows exemplary targeting sequences, in a 5'-to-3' orientation, that are complementary to a broad spectrum of picornaviruses, specifically members of the *Enterovirus* and *Rhinovirus* genera. The targeting sequences listed below in Table 3 provide a collection of targeting sequences from which targeting sequences may be selected, according to the general class rules discussed above. As seen, the targeting sequences represented by SEQ ID NOS: 8 and 9 can be represented by SEQ ID NO:10, where "R" represents a purine, either A or G.

Table 3. Exemplary Antisense Oligomer Targeting Sequences

PMO	Target Nucleotides	GenBank Acc. No.	Targeting Antisense Oligomer (5' to 3')	SEQ. ID NO.
5'-32a	443-474	V00149	GTTGGGATTAGCCGCATTCAGGGGCCG GAGGA	8
5'-32b	443-474	V00149	GTTGGGGTTGGCCGCATTCAGGGGCCGAGGA	9
5'-32	443-474		GTTGGGRTRGCCGCATTCAGGGGCCG GAGGA	10
PV444	444-463	V00149	CCGCATTCAGGGGCCGAGG	11
PV449	449-470	V00149	GGATTAGCCGCATTCAGGGGCC	12
PV454	454-474	V00149	GTTGGGATTAGCCGCATTCAG	13

III. Antisense Oligonucleotide analog compounds

A. Properties

As detailed above, the antisense oligonucleotide analog compound (the term "antisense" indicates that the compound is targeted against the vims' positive-sense strand RNA) has a base sequence targeting a region that includes one or more of the following; 1) the 5' untranslated region of the positive sense viral RNA and; 2) the internal ribosome entry site within the 5' untranslated region. In addition, the oligomer is able to effectively target infecting viruses, when administered to a host cell, e.g. in an infected mammalian subject. This requirement is met when the oligomer compound (a) has the ability to be actively taken up by mammalian cells, and (b) once taken up, form a duplex with the target RNA with a T_m greater than about 45°C.

As will be described below, the ability to be taken up by cells requires that the oligomer backbone be substantially uncharged, and, preferably, that the oligomer structure is recognized as a substrate for active or facilitated transport across the cell membrane. The ability of the oligomer to form a stable duplex with the target RNA will also depend on the oligomer backbone, as well as factors noted above, the length and degree of complementarity of the antisense oligomer with respect to the target, the ratio of G:C to A:T base matches, and the positions of any mismatched bases. The ability of the antisense oligomer to resist cellular nucleases promotes survival and ultimate delivery of the agent to the cell cytoplasm.

Below are disclosed methods for testing any given, substantially uncharged backbone for its ability to meet these requirements.

B. Active or facilitated uptake by cells

The antisense compound may be taken up by host cells by facilitated or active transport across the host cell membrane if administered in free (non-complexed) form, or by an endocytotic mechanism if administered in complexed form.

In the case where the agent is administered in free form, the antisense compound should be substantially uncharged, meaning that a majority of its intersubunit linkages are uncharged at physiological pH. Experiments carried out in support of the invention indicate that a small number of net charges, e.g., 1-2 for

a 15- to 20-mer oligomer, can in fact enhance cellular uptake of certain oligomers with substantially uncharged backbones. The charges may be carried on the oligomer itself, *e.g.*, in the backbone linkages, or may be terminal charged-group appendages. Preferably, the number of charged linkages is no more than one charged linkage per four uncharged linkages. More preferably, the number is no more than one charged linkage per ten, or no more than one per twenty, uncharged linkages. In one embodiment, the oligomer is fully uncharged.

An oligomer may also contain both negatively and positively charged backbone linkages, as long as opposing charges are present in approximately equal number. Preferably, the oligomer does not include runs of more than 3-5 consecutive subunits of either charge. For example, the oligomer may have a given number of anionic linkages, *e.g.* phosphorothioate or N3'→P5' phosphoramidate linkages, and a comparable number of cationic linkages, such as N,N-diethylenediamine phosphoramidates (Dagle, Littig *et al.* 2000). The net charge is preferably neutral or at most 1-2 net charges per oligomer.

The antisense compound may also be administered in complexed form, where the complexing agent is typically a polymer, *e.g.*, a cationic lipid, polypeptide, or non-biological cationic polymer, having an opposite charge to any net charge on the antisense compound. Methods of forming complexes, including bilayer complexes, between anionic oligonucleotides and cationic lipid or other polymer components, are well known. For example, the liposomal composition Lipofectin (Feigner, Gadek *et al.* 1987), containing the cationic lipid DOTMA (N-[1-(2,3-dioleoyloxy)propyl]-N,N,N-trimethylammonium chloride) and the neutral phospholipid DOPE (dioleoyl phosphatidyl ethanolamine), is widely used. After administration, the complex is taken up by cells through an endocytotic mechanism, typically involving particle encapsulation in endosomal bodies.

The antisense compound may also be administered in conjugated form with an arginine-rich peptide linked covalently to the 5' or 3' end of the antisense oligomer. The peptide is typically 8-16 amino acids and consists of a mixture of arginine, and other amino acids including phenylalanine, cysteine, beta-alanine and 6-aminohexanoic acid. Exemplary arginine-rich delivery peptides are described in the Sequence Listing table as SEQ ID NOS:14-19. The use of arginine-rich peptide-PMO conjugates can be used to enhance cellular uptake of the antisense oligomer (See, *e.g.* Moulton, Nelson *et al.* 2004; Nelson, Stein *et al.* 2005).

In some instances, liposomes may be employed to facilitate uptake of the antisense oligonucleotide into cells. (See, e.g., Williams, S.A., Leukemia 10(1 2):1 980-1 989, 1996; Lappalainen *et al.*, Antiviral Res. 23:1 19, 1994; Uhlmann *et al.*, antisense oligonucleotides: a new therapeutic principle, Chemical Reviews, Volume 90, No. 4, pages 544-584, 1990; Gregoriadis, G., Chapter 14, Liposomes, Drug Carriers in Biology and Medicine, pp. 287-341 , Academic Press, 1979). Hydrogels may also be used as vehicles for antisense oligomer administration, for example, as described in WO 93/01286. Alternatively, the oligonucleotides may be administered in microspheres or microparticles. (See, e.g., Wu, G.Y. and Wu, C.H., J. Biol. Chem. 262:4429-4432, 1987). Alternatively, the use of gas-filled microbubbles complexed with the antisense oligomers can enhance delivery to target tissues, as described in US Patent No. 6,245,747.

Alternatively, and according to another aspect of the invention, the requisite properties of oligomers with any given backbone can be confirmed by a simple *in vivo* test, in which a labeled compound is administered to an animal, and a body fluid sample, taken from the animal several hours after the oligomer is administered, assayed for the presence of heteroduplex with target RNA. This method is detailed in subsection D below.

C. Substantial resistance to RNaseH

Two general mechanisms have been proposed to account for inhibition of expression by antisense oligonucleotides. (See e.g., (Agrawal, Mayrand *et al.* 1990; Bonham, Brown *et al.* 1995; Boudvillain, Guerin *et al.* 1997). In the first, a heteroduplex formed between the oligonucleotide and the viral RNA acts as a substrate for RNaseH, leading to cleavage of the viral RNA. Oligonucleotides belonging, or proposed to belong, to this class include phosphorothioates, phosphotriesters, and phosphodiester (unmodified "natural" oligonucleotides). Such compounds expose the viral RNA in an oligomer:RNA duplex structure to hydrolysis by RNaseH, and therefore loss of function.

A second class of oligonucleotide analogs, termed "steric blockers" or, alternatively, "RNaseH inactive" or "RNaseH resistant", have not been observed to act as a substrate for RNaseH, and are believed to act by sterically blocking target RNA nucleocytoplasmic transport, splicing or translation. This class includes methylphosphonates (Toulme *et al.*, 1996), morpholino oligonucleotides, peptide

nucleic acids (PNA's), certain 2'-O-allyl or 2'-O-alkyl modified oligonucleotides (Bonham, 1995), and N3'-P5' phosphoramidates (Gee, 1998; Ding, 1996).

A test oligomer can be assayed for its RNaseH resistance by forming an RNA:oligomer duplex with the test compound, then incubating the duplex with RNaseH under a standard assay conditions, as described in Stein *et al.* After exposure to RNaseH, the presence or absence of intact duplex can be monitored by gel electrophoresis or mass spectrometry.

D. In vivo uptake

In accordance with another aspect of the invention, there is provided a simple, rapid test for confirming that a given antisense oligomer type provides the required characteristics noted above, namely, high T_m , ability to be actively taken up by the host cells, and substantial resistance to RNaseH. This method is based on the discovery that a properly designed antisense compound will form a stable heteroduplex with the complementary portion of the viral RNA target when administered to a mammalian subject, and the heteroduplex subsequently appears in the urine (or other body fluid). Details of this method are also given in co-owned U.S. Patent applications, Serial No. 09/736,920, entitled "Non-Invasive Method for Detecting Target RNA" (Non-Invasive Method), the disclosure of which is incorporated herein by reference.

Briefly, a test oligomer containing a backbone to be evaluated, having a base sequence targeted against a known RNA, is injected into a mammalian subject. The antisense oligomer may be directed against any intracellular RNA, including a host RNA or the RNA of an infecting virus. Several hours (typically 8-72) after administration, the urine is assayed for the presence of the antisense-RNA heteroduplex. If heteroduplex is detected, the backbone is suitable for use in the antisense oligomers of the present invention.

The test oligomer may be labeled, *e.g.* by a fluorescent or a radioactive tag, to facilitate subsequent analyses, if it is appropriate for the mammalian subject. The assay can be in any suitable solid-phase or fluid format. Generally, a solid-phase assay involves first binding the heteroduplex analyte to a solid-phase support, *e.g.*, particles or a polymer or test-strip substrate, and detecting the presence/amount of heteroduplex bound. In a fluid-phase assay, the analyte sample is typically pretreated to remove interfering sample components. If the

oligomer is labeled, the presence of the heteroduplex is confirmed by detecting the label tags. For non-labeled compounds, the heteroduplex may be detected by immunoassay if in solid phase format or by mass spectroscopy or other known methods if in solution or suspension format.

When the antisense oligomer is complementary to a virus-specific region of the viral genome (such as those regions of picomavirus RNA, as described above) the method can be used to detect the presence of a given picomavirus virus, or reduction in the amount of virus during a treatment method.

E. Exemplary oligomer backbones

Examples of nonionic linkages that may be used in oligonucleotide analogs are shown in Figs. 2A-2G. In these figures, B represents a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, preferably selected from adenine, cytosine, guanine, thymidine and uracil. Suitable backbone structures include carbonate (2A, R=O) and carbamate (2A, R=NH₂) linkages (Mertes and Coats 1969; Gait, Jones *et al.* 1974); alkyl phosphonate and phosphotriester linkages (2B, R=alkyl or -O-alkyl) (Lesnikowski, Jaworska *et al.* 1990); amide linkages (2C) (Blommers, Pieles *et al.* 1994); sulfone and sulfonamide linkages (2D, R₁, R₂ = CH₂); and a thioformacetyl linkage (2E) (Cross, Rice *et al.* 1997). The latter is reported to have enhanced duplex and triplex stability with respect to phosphorothioate antisense compounds (Cross, Rice *et al.* 1997). Also reported are the 3'-methylene-N-methylhydroxyamino compounds of structure 2F. Also shown is a cationic linkage in Fig. 2H wherein the nitrogen pendant to the phosphate atom in the linkage of Fig 2G is replaced with a 1-piperazino structure. The method for synthesizing the 1-piperazino group linkages is described below with respect to Fig. 5. Figure 2H is an example of a cationic linkage group.

As noted above, the substantially uncharged oligomer may advantageously include a limited number of charged backbone linkages. One example of a cationic charged phosphordiamidate linkage is shown in Fig. 2H. This linkage, in which the dimethylamino group shown in Fig 2G is replaced by a 1-piperazino group as shown in Fig. 2G, can be substituted for any linkage(s) in the oligomer. By including between two to eight such cationic linkages, and more generally, at least two and no more than about half the total number of linkages, interspersed

along the backbone of the otherwise uncharged oligomer, antisense activity can be enhanced without a significant loss of specificity. The charged linkages are preferably separated in the backbone by at least 1 and preferably 2 or more uncharged linkages.

Peptide nucleic acids (PNAs) are analogs of DNA in which the backbone is structurally homomorphous with a deoxyribose backbone, consisting of N-(2-aminoethyl) glycine units to which pyrimidine or purine bases are attached. PNAs containing natural pyrimidine and purine bases hybridize to complementary oligonucleotides obeying Watson-Crick base-pairing rules, and mimic DNA in terms of base pair recognition (Egholm, Buchardt *et al.* 1993). The backbone of PNAs are formed by peptide bonds rather than phosphodiester bonds, making them well-suited for antisense applications. The backbone is uncharged, resulting in PNA/DNA or PNA/RNA duplexes which exhibit greater than normal thermal stability. PNAs are not recognized by nucleases or proteases.

A preferred oligomer structure employs morpholino-based subunits bearing base-pairing moieties, joined by uncharged linkages, as described above. Especially preferred is a substantially uncharged phosphorodiamidate-linked morpholino oligomer, such as illustrated in Figs. 1A-1D. Morpholino oligonucleotides, including antisense oligomers, are detailed, for example, in co-owned U.S. Patent Nos. 5,698,685, 5,217,866, 5,142,047, 5,034,506, 5,166,315, 5,185, 444, 5,521,063, and 5,506,337, all of which are expressly incorporated by reference herein.

Important properties of the morpholino-based subunits include: the ability to be linked in a oligomeric form by stable, uncharged backbone linkages; the ability to support a nucleotide base (e.g. adenine, cytosine, guanine, thymidine, uracil or inosine) such that the polymer formed can hybridize with a complementary-base target nucleic acid, including target RNA, with high T_m , even with oligomers as short as 10-14 bases; the ability of the oligomer to be actively transported into mammalian cells; and the ability of the oligomer.RNA heteroduplex to resist RNase degradation.

Exemplary backbone structures for antisense oligonucleotides of the invention include the α -morpholino subunit types are also shown in Figs. 1A-1 D, each linked by an uncharged, phosphorus-containing subunit linkage. Fig. 1A shows a phosphorus-containing linkage which forms the five atom repeating-unit

backbone, where the morpholino rings are linked by a 1-atom phosphoamide linkage. Fig. 1B shows a linkage which produces a 6-atom repeating-unit backbone. In this structure, the atom Y linking the 5' morpholino carbon to the phosphorus group may be sulfur, nitrogen, carbon or, preferably, oxygen. The X moiety pendant from the phosphorus may be fluorine, an alkyl or substituted alkyl, an alkoxy or substituted alkoxy, a thioalkoxy or substituted thioalkoxy, or unsubstituted, monosubstituted, or disubstituted nitrogen, including cyclic structures, such as morpholines or piperidines. Alkyl, alkoxy and thioalkoxy preferably include 1-6 carbon atoms. The Z moieties are sulfur or oxygen, and are preferably oxygen.

The linkages shown in Fig. 1C and 1D are designed for 7-atom unit-length backbones. In Structure 1C, the X moiety is as in Structure 1B, and the moiety Y may be methylene, sulfur, or, preferably, oxygen. In Structure 1D, the X and Y moieties are as in Structure 1B. Particularly preferred morpholino oligonucleotides include those composed of morpholino subunit structures of the form shown in Figure 1B, where $X = \text{NH}_2$ or $\text{N}(\text{CH}_3)_2$, $Y = \text{O}$, and $Z = \text{O}$ and in Figure 2G.

As noted above, the substantially uncharged oligomer may advantageously include a limited number of charged backbone linkages. One example of a cationic charged phosphordiamidate linkage is shown in Fig. 2H. This linkage, in which the dimethylamino group shown in Fig 2G is replaced by a 1-piperazino group as shown in Fig. 2H, can be substituted for any linkage(s) in the oligomer. By including between two to eight such cationic linkages, and more generally, at least two and no more than about half the total number of linkages interspersed along the backbone of the otherwise uncharged oligomer, antisense activity can be enhanced without a significant loss of specificity. The charged linkages are preferably separated in the backbone by at least 1 and preferably 2 or more uncharged linkages.

The antisense compounds can be prepared by stepwise solid-phase synthesis, employing methods detailed in the references cited above. In some cases, it may be desirable to add additional chemical moieties to the antisense compound, e.g. to enhance pharmacokinetics or to facilitate capture or detection of the compound. Such a moiety may be covalently attached, typically to a terminus of the oligomer, according to standard synthetic methods. For example, addition of a polyethyleneglycol moiety or other hydrophilic polymer, e.g., one having 10-

100 monomelic subunits, may be useful in enhancing solubility. One or more charged groups, e.g., anionic charged groups such as an organic acid, may enhance cell uptake. A reporter moiety, such as fluorescein or a radiolabeled group, may be attached for purposes of detection. Alternatively, the reporter label attached to the oligomer may be a ligand, such as an antigen or biotin, capable of binding a labeled antibody or streptavidin. In selecting a moiety for attachment or modification of an antisense oligomer, it is generally of course desirable to select chemical compounds of groups that are biocompatible and likely to be tolerated by a subject without undesirable side effects.

IV. Inhibition of Picomavirus Viral Replication

The antisense compounds detailed above are useful in inhibiting replication of single-stranded, positive-sense RNA viruses of the Picornaviridae family. In one embodiment, such inhibition is effective in treating infection of a host animal by these viruses. Accordingly, the method comprises, in one embodiment, exposing a mammalian cell infected with the virus with an oligonucleotide antisense compound effective to inhibit the replication of the specific virus. In this embodiment, the cells are exposed to the compound either *in vitro* or *in vivo*, where the method is used in the latter case to treat a mammalian subject, e.g., human or domestic animal, infected with a given virus. It is contemplated that the antisense oligonucleotide arrests the growth of the RNA virus in the host. The RNA virus may be decreased in number or eliminated with little or no detrimental effect on the normal growth or development of the host.

In the present invention as described in the Examples, Phosphorodiamidate Morpholino Oligomers (PMOs), designed to hybridize to specific regions of the poliovirus 5' UTR, are evaluated for their ability to inhibit IRES-mediated translation in a cell-free translation system. The region of the virus 5' UTR noted above having the target region is highly conserved within the Picornaviridae family and even more highly conserved within the *Enterovirus* and *Rhinovirus* genera that are specifically targeted. The PMO described herein will target most, if not all, picomavirus virus species because of the high degree of homology between viral species at the respective targets (SEQ ID NOS:1-6) as shown in Figure 3.

A. Identification of the Infective Agent

The specific virus causing the infection can be determined by methods known in the art, e.g. serological, genotyping, or cultural methods, or by methods employing the antisense oligomers of the present invention.

Serological identification employs a viral sample or culture isolated from a biological specimen, e.g., stool, urine, cerebrospinal fluid, blood, nasopharyngeal secretions, *etc.*, of the subject. Immunoassay for the detection of virus is generally carried out by methods routinely employed by those of skill in the art, e.g., ELISA or Western blot. In addition, monoclonal antibodies specific to particular viral strains or species are often commercially available.

Culture methods may be used to isolate and identify particular types of virus, by employing techniques including, but not limited to, comparing characteristics such as rates of growth and morphology under various culture conditions.

Genotyping methods include polymerase chain reaction (PCR) methods using genotype-specific primers or genomic sequencing of viral nucleic acid obtained from the infected individual.

Another method for identifying the viral infective agent in an infected subject employs one or more antisense oligomers targeting broad families and/or genera of viruses. Sequences targeting any characteristic viral RNA can be used. The desired target sequences are preferably (i) common to broad virus families/genera, and (ii) not found in humans. Characteristic nucleic acid sequences for a large number of infectious viruses are available in public databases, and may serve as the basis for the design of specific oligomers.

For each plurality of oligomers, the following steps are carried out: (a) the oligomers) are administered to the subject; (b) at a selected time after said administering, a body fluid sample is obtained from the subject; and (c) the sample is assayed for the presence of a nuclease-resistant heteroduplex comprising the antisense oligomer and a complementary portion of the viral genome. Steps (a) - (c) are carried for at least one such oligomer, or as many as is necessary to identify the virus or family of viruses. Oligomers can be administered and assayed sequentially or, more conveniently, concurrently. The virus is identified based on the presence (or absence) of a heteroduplex comprising the antisense oligomer

and a complementary portion of the viral genome of the given known virus or family of viruses.

Preferably, a first group of oligomers, targeting broad families, is utilized first, followed by selected oligomers complementary to specific genera and/or species and/or strains within the broad family/genus thereby identified. This second group of oligomers includes targeting sequences directed to specific genera and/or species and/or strains within a broad family/genus. Several different second oligomer collections, *i.e.* one for each broad virus family/genus tested in the first stage, are generally provided. Sequences are selected which are (i) specific for the individual genus/species/strains being tested and (ii) not found in humans.

B. Administration of the Antisense Oligomer

Effective delivery of the antisense oligomer to the target nucleic acid is an important aspect of treatment. In accordance with the invention, routes of antisense oligomer delivery include, but are not limited to, various systemic routes, including oral and parenteral routes, *e.g.*, intravenous, subcutaneous, intraperitoneal, and intramuscular, as well as inhalation, transdermal and topical delivery. The appropriate route may be determined by one of skill in the art, as appropriate to the condition of the subject under treatment. For example, an appropriate route for delivery of an antisense oligomer in the treatment of a viral infection of the skin is topical delivery, while delivery of a antisense oligomer for the treatment of a viral respiratory infection is by inhalation. The oligomer may also be delivered directly to the site of viral infection, or to the bloodstream.

The antisense oligomer may be administered in any convenient vehicle which is physiologically acceptable. Such a composition may include any of a variety of standard pharmaceutically accepted carriers employed by those of ordinary skill in the art. Examples include, but are not limited to, saline, phosphate buffered saline (PBS), water, aqueous ethanol, emulsions, such as oil/water emulsions or triglyceride emulsions, tablets and capsules. The choice of suitable physiologically acceptable carrier will vary dependent upon the chosen mode of administration.

In some instances, liposomes may be employed to facilitate uptake of the antisense oligomer into cells. (See, *e.g.*, Williams, S.A., *Leukemia* 10(12):1980-

1989, 1996; Lappalainen *et al.*, *Antiviral Res.* 23:1 19, 1994; Uhlmann *et al.*, antisense oligonucleotides: a new therapeutic principle, *Chemical Reviews*, Volume 90, No. 4, pages 544-584, 1990; Gregoriadis, G., Chapter 14, *Liposomes, Drug Carriers in Biology and Medicine*, pp. 287-341, Academic Press, 1979). As described above, the use of arginine-rich cellular delivery peptides conjugated to the antisense oligomer may also be used. Hydrogels may also be used as vehicles for antisense oligomer administration, for example, as described in WO 93/01286. Alternatively, the oligomers may be administered in microspheres or microparticles. (See, *e.g.*, Wu, G.Y. and Wu, C.H., *J. Biol. Chem.* 262:4429-4432, 1987). Alternatively, the use of gas-filled microbubbles complexed with the antisense oligomers can enhance delivery to target tissues, as described in US Patent No. 6,245,747.

Sustained release compositions may also be used. These may include semipermeable polymeric matrices in the form of shaped articles such as films or microcapsules.

In one aspect of the method, the subject is a human subject, *e.g.*, a patient diagnosed as having a localized or systemic viral infection. The condition of a patient may also dictate prophylactic administration of an antisense oligomer of the invention, *e.g.* in the case of a patient who (1) is immunocompromised; (2) is a burn victim; (3) has an indwelling catheter; or (4) is about to undergo or has recently undergone surgery. In one preferred embodiment, the oligomer is a phosphorodiamidate morpholino oligomer, contained in a pharmaceutically acceptable carrier, and is delivered orally. In another preferred embodiment, the oligomer is a phosphorodiamidate morpholino oligomer, contained in a pharmaceutically acceptable carrier, and is delivered intravenously (*i.v.*).

In another application of the method, the subject is a livestock animal, *e.g.*, a chicken, turkey, pig, cow or goat, *etc.*, and the treatment is either prophylactic or therapeutic. The invention also includes a livestock and poultry food composition containing a food grain supplemented with a subtherapeutic amount of an antiviral antisense compound of the type described above. Also contemplated is, in a method of feeding livestock and poultry with a food grain supplemented with subtherapeutic levels of an antiviral, an improvement in which the food grain is supplemented with a subtherapeutic amount of an antiviral oligonucleotide composition as described above.

The antisense compound is generally administered in an amount and manner effective to result in a peak blood concentration of at least 200-400 nM antisense oligomer. Typically, one or more doses of antisense oligomer are administered, generally at regular intervals, for a period of about one to two weeks. Preferred doses for oral administration are from about 1-100 mg oligomer per 70 kg. In some cases, doses of greater than 100 mg oligomer/patient may be necessary. For i.v. administration, preferred doses are from about 0.5 mg to 100 mg oligomer per 70 kg. The antisense oligomer may be administered at regular intervals for a short time period, e.g., daily for two weeks or less. However, in some cases the oligomer is administered intermittently over a longer period of time. Administration may be followed by, or concurrent with, administration of an antibiotic or other therapeutic treatment. The treatment regimen may be adjusted (dose, frequency, route, *etc.*) as indicated, based on the results of immunoassays, other biochemical tests and physiological examination of the subject under treatment.

C. Monitoring of Treatment

An effective *in vivo* treatment regimen using the antisense oligonucleotides of the invention may vary according to the duration, dose, frequency and route of administration, as well as the condition of the subject under treatment (*i.e.*, prophylactic administration versus administration in response to localized or systemic infection). Accordingly, such *in vivo* therapy will often require monitoring by tests appropriate to the particular type of viral infection under treatment, and corresponding adjustments in the dose or treatment regimen, in order to achieve an optimal therapeutic outcome. Treatment may be monitored, e.g., by general indicators of infection, such as complete blood count (CBC), nucleic acid detection methods, immunodiagnostic tests, viral culture, or detection of heteroduplex.

The efficacy of an *in vivo* administered antisense oligomer of the invention in inhibiting or eliminating the growth of one or more types of RNA virus may be determined from biological samples (tissue, blood, urine *etc.*) taken from a subject prior to, during and subsequent to administration of the antisense oligomer. Assays of such samples include (1) monitoring the presence or absence of heteroduplex formation with target and non-target sequences, using procedures known to those skilled in the art, e.g., an electrophoretic gel mobility assay;

(2) monitoring the amount of viral protein production, as determined by standard techniques such as ELISA or Western blotting, or (3) measuring the effect on viral titer, e.g. by the method of Spearman-Kärber. (See, for example, Pari, G.S. *et al.*, *Antimicrob. Agents and Chemotherapy* 39(5):1 157-1 161 , 1995; Anderson, KP. *et al.*, *Antimicrob. Agents and Chemotherapy* 40(9):2004-201 1, 1996, Cottral, G.E. (ed) in: *Manual of Standard Methods for Veterinary Microbiology*, pp. 60-93, 1978).

A preferred method of monitoring the efficacy of the antisense oligomer treatment is by detection of the antisense-RNA heteroduplex. At selected time(s) after antisense oligomer administration, a body fluid is collected for detecting the presence and/or measuring the level of heteroduplex species in the sample. Typically, the body fluid sample is collected 3-24 hours after administration, preferably about 6-24 hours after administering. As indicated above, the body fluid sample may be urine, saliva, plasma, blood, spinal fluid, or other liquid sample of biological origin, and may include cells or cell fragments suspended therein, or the liquid medium and its solutes. The amount of sample collected is typically in the 0.1 to 10 ml range, preferably about 1 ml or less.

The sample may be treated to remove unwanted components and/or to treat the heteroduplex species in the sample to remove unwanted ssRNA overhang regions, e.g. by treatment with RNase. It is, of course, particularly important to remove overhang where heteroduplex detection relies on size separation, e.g., electrophoresis or mass spectroscopy.

A variety of methods are available for removing unwanted components from the sample. For example, since the heteroduplex has a net negative charge, electrophoretic or ion exchange techniques can be used to separate the heteroduplex from neutral or positively charged material. The sample may also be contacted with a solid support having a surface-bound antibody or other agent specifically able to bind the heteroduplex. After washing the support to remove unbound material, the heteroduplex can be released in substantially purified form for further analysis, e.g., by electrophoresis, mass spectroscopy or immunoassay.

V. Examples

The following examples illustrate but are not intended in any way to limit the invention.

A. Materials and Methods

Standard recombinant DNA techniques are employed in all constructions, as described in Ausubel, FM *et al.*, in CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley and Sons, Inc., Media, PA, 1992 and Sambrook, J. *et al.*, in MOLECULAR CLONING: A LABORATORY MANUAL, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, Vol. 2, 1989).

All peptides are custom synthesized by Global Peptide Services (Ft. Collins, CO) or at AVI BioPharma (Corvallis, OR) and purified to >90% purity. Phosphorodiamidate morpholino oligomers (PMOs) are synthesized at AVI BioPharma in accordance with known methods, as described, for example, in (Summerton and Weller 1997) and U.S. Patent No. 5,185,444.

For Examples 2 and 3 described below, PMO oligomers are conjugated at the 5' end with an arginine-rich peptide (RXR)₄XB-PMO (where R is arginine, X is 6-aminohexanoic acid and B is beta-alanine) to enhance cellular uptake as described (US Patent Application 60/466,703 and Moulton, Nelson *et al.* 2004; Nelson, Stein *et al.* 2005). This peptide is also called P007 and listed as SEQ ID NO:13 in the Sequence Listing table.

Preparation of Morpholino Oligomers having Cationic Linkages

A schematic of a synthetic pathway that can be used to make morpholino subunits containing a (1 piperazino) phosphinylideneoxy linkage is shown in Figure 5; further experimental detail for a representative synthesis is provided in Materials and Methods, below. As shown in the Figure, reaction of piperazine and trityl chloride gave trityl piperazine (1a), which was isolated as the succinate salt. Reaction with ethyl trifluoroacetate (1b) in the presence of a weak base (such as diisopropylethylamine or DIEA) provided 1-trifluoroacetyl-4-trityl piperazine (2), which was immediately reacted with HCl to provide the salt (3) in good yield. Introduction of the dichlorophosphoryl moiety was performed with phosphorus oxychloride in toluene.

The acid chloride (4) is reacted with morpholino subunits (moN), which may be prepared as described in U.S. Patent No. 5,185,444 or in Summerton and Weller, 1997 (cited above), to provide the activated subunits (5,6,7). Suitable protecting groups are used for the nucleoside bases, where necessary; for example, benzoyl for adenine and cytosine, isobutyryl for guanine, and

pivaloylmethyl for inosine. The subunits containing the (1 piperazino) phosphinylideneoxy linkage can be incorporated into the existing PMO synthesis protocol, as described, for example in Summerton and Weller (1997), without modification.

Example 1. Inhibition of Picomavirus virus RNA Translation with
Phosphorodiamidate Morpholino Oligomers

Phosphorodiamidate Morpholino Oligomers (PMOs), designed to hybridize to a specific region of the poliovirus internal ribosome entry site (IRES) region of poliovirus, as described above, are evaluated for their ability to inhibit translation in a rabbit reticulocyte lysate (RRL) assay. A dual reporter expression plasmid construct is derived as follows. A 507 nucleotide fragment of the 5' untranslated region of the poliovirus genome (nets 124 to 630, GenBank No. V01 149) is synthesized (BlueHeron, Inc) and inserted into the intercistronic region of a dual luciferase reporter plasmid described previously (Johannes, Carter *et al.* 1999; Wilson, Powell *et al.* 2000; Brasey, Lopez-Lastra *et al.* 2003) and named herein as pR&Fluc. The resulting plasmid, pR&Fluc-PoliolIRES is shown schematically in Figure 4. For transcription *in vitro*, the dicistronic luciferase pR&Fluc-PoliolIRES construct is digested with *Bam*HI and linear DNA is transcribed with T7 RNA polymerase using the RiboMAX protocol (Promega, Inc) to produce rLuc/fLuc RNA. Uncapped dicistronic RNAs are translated in the RRL, as recommended, and products of translation reactions are measured enzymatically using the dual luciferase reporter assay system (Promega).

Prior to cell-free translation, one nanomolar of the dicistronic RNA is incubated with PMOs (SEQ ID NOS: 11-13) at varying concentrations ranging from 0.01 to 10 micromolar. Relative inhibition of the fLuc downstream reporter gene is measured using a fluorometer. Figure 6A shows the fLuc light output from rLuc/fLuc RNA in the presence of four different PMOs including two that are examples of the present invention, 444 and 449; SEQ ID NOS: 11 and 12, respectively. These two PMOs are compared to two other PMOs, 162 and 544, which are targeted to regions of the poliovirus IRES approximately 280 bases in the 5¹ direction and 100 bases in the 3' direction, respectively. The sequence of PMO 162 is 5'-CCG GGG AAA CAG AAG TGC TTG-3¹ and for PMO 544 it is 5'-GGA AAC ACG GAC ACC CAA AG-3'. The effective concentrations at which 50%

inhibition (EC₅₀) of fLuc output are also listed in Fig. 6. Both the 444 and 449 PMOs have significantly lower EC₅₀s than either of the comparator PMOs. Figure 6B shows similar results to Figure 6A but the data shown is the fLuc to rLuc signal ratio which normalizes the results to the rLuc internal control. The region targeted by the 544 PMO is equally conserved across the Picornaviridae family (See US Provisional Patent Application 60/715,888) as compared to the region targeted by the 444 and 449 PMOs, the focus of the present invention. However, the latter PMOs demonstrate EC₅₀s in this assay that are as much as 45% lower than those observed with the 544 PMO.

Example 2. Inhibition of Coxsackievirus B3 (CVB3) in Tissue Culture with PMOs that Target the 5' UTR of CVB3

The antiviral activity of CVB3-specific PMOs is determined by measuring viral protein expression in PMO-treated, CVB3-infected cells. The test is performed on either cardiomyocytes (HL-1 cells) or HeLa cells. Cell monolayers (6-well plates) are seeded 16 to 20 hours prior to treatment with PMO or infection with virus. Serum-containing medium is replaced with serum-free medium during PMO treatment and infection. The infection is allowed to proceed for either 7h or 24h at 37°C prior to collection of cells and the preparation of cell lysates for immunoblot detection of viral capsid protein by Western blot (*i.e.* immunoblot) assay using a polyclonal antibody to the CVB3 VP1 gene. Antisera against beta-actin is included as a loading control.

In one set of experiments, 6 x 10⁵ HL-1 cells (cardiomyocytes) are seeded in 6-well plates for 16-20h. Culture medium is then replaced with serum-free medium and incubated with individual PMOs at final concentration of 10 μM. Four hours post-infection, cells are infected with CVB3 at a multiplicity of infection (MOI) of 10. At 24 hours post-infection, cell lysates are collected for Western immunoblot analysis to detect viral capsid protein VP1. Immunological detection of beta-actin is used as a loading control. Eight different peptide(P007)-conjugated PMOs are tested.

One of the effective antisense PMOs (PV444, PV449, or PV454; SEQ ID NOS:1 1, 12, and 13) is selected for a dose response assay using HeLa cells under the same conditions as described above. The concentration range analyzed is from 0.01 to 20 micromolar PMO. An estimated effective concentration sufficient

for a reduction in viral replication of 50% (EC50) for this PMO is between 5 and 10 micromolar.

Effective inhibition of CVB3 replication with two PMOs is observed as described above. This inhibition is under conditions where the PMO is introduced four hours prior to infection.

Example 3. Inhibition of CVB3-induced Cytopathic Effects in Tissue Culture with PMOs that Target the 5' UTR of CVB3

Another measure of antiviral activity is to observe a reduction in cytopathic effects (CPE) in tissue culture experiments. Three P007-conjugated PMO are selected for these analyses, PV444, PV449, and PV454 (SEQ ID NOS:1 1, 12, and 13, respectively) are used to treat both HL-1 cells (cardiomyocytes) and HeLa cells under the same conditions described in Example 2 above. Four hours post-treatment with PMO, cells are infected with CVB3 at an MOI of ten. Photomicrographs are taken 24 hours post-infection for HeLa cells.

SEQUENCE LISTING

Name	Target Sequences (5' to 3')	SEQ ID NO
5'-32	TCCTCCGGCCCCTGAATGCGGCTAATCCCAAC	1
5'-32	TCCTCCGGCCCCTGAATGCGGCTAATCCTAAC	2
5'-32	TCCTCCGGCCCCTGAATGCGGCTAATCCTAAC	3
5'-32	TCCTCCGGCCCCTGAATGCGGTTAATCCTAAC	4
5'-32	TCCTCCGGCCCCTGAATGTGGCTAACCTTAAC	5
5'-32	TCCTCCGGCCCCTGAATGCGGCTAACCTTAAC	6
5'-32	TCCTCCGGCCCCTGAATGYGGCTAAYCYAAC	7
	Oligomer Targeting Sequences (5' to 3')	
5'-32a	GTTGGGATTAGCCGCATTCAGGGGCCGGAGGA	8
5'-32b	GTTGGGGTTGGCCGCATTCAGGGGCCGGAGGA	9
5'-32	GTTGGGRTRGCCGCATTCAGGGGCCGGAGGA	10
PV444	CCGCATTCAGGGGCCGGAGG	11
PV449	GGATTAGCCGCATTCAGGGGCC	12
PV454	GTTGGGATTAGCCGCATTCAG	13
	Peptide Sequences (NH ₂ to COOH)	
P003	RRRRRRRRRFFAhxβAla	14
P007	(RAhxR) ₄ AhxβAla	15
P008	(RAhx) ₈ βAla	16
RX4	(RAhx) ₄ βAla	17
RXR2	(RAhxR) ₂ AhxβAla	18
RXR3	(RAhxR) ₃ AhxβAla	19

IT IS CLAIMED:

1. A method of inhibiting viral infection in mammalian cells by an *Enterovirus* or *Rhinovirus* in the Picomaviridae family, comprising

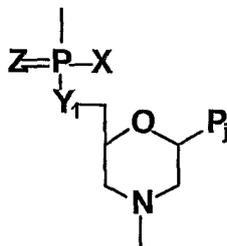
(a) exposing the cells to an antisense oligonucleotide compound characterized by:

- (i) a substantially uncharged, nuclease-resistant backbone,
- (ii) capable of uptake by mammalian host cells,
- (iii) containing between 12-40 nucleotide bases, and
- (iv) having a targeting sequence of at least 12 subunits complementary to SEQ ID NO:7 in the positive-sense strand of the virus, and

(b) by said exposing, forming a heteroduplex structure (i) composed of the virus¹ positive sense strand and the oligonucleotide compound, and (ii) characterized by a T_m of dissociation of at least 45⁰C.

2. The method of claim 1, wherein said oligonucleotide compound to which the host cells are exposed is composed of morpholino subunits and phosphorus-containing intersubunit linkages joining a morpholino nitrogen of one subunit to a 5' exocyclic carbon of an adjacent subunit.

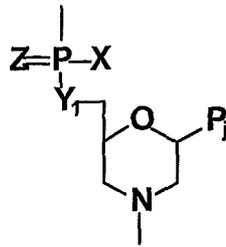
3. The method of claim 2, wherein the morpholino subunits in the oligonucleotide compound to which the host cells are exposed is administered to the subject are joined by phosphorodiamidate linkages having the structure:



where Y_i=O, Z=O, P_j is a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, and X is alkyl, alkoxy, thioalkoxy, amino or alkyl amino, including diaikylamino.

4. The method of claim 2, in which at least 2 and no more than half of the total number of intersubunit linkages are positively charged at physiological pH.

5. The method of claim 2, wherein said morpholino subunits are joined by phosphorodiamidate linkages, in accordance with the structure:



where $Y_i=O$, $Z=O$, P_j is a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, and X for the uncharged linkages is alkyl, alkoxy, thioalkoxy, or an alkyl amino of the form wherein NR_2 , where each R is independently hydrogen or methyl, and for the positively charged linkages, X is 1-piperazine.

6. The method of claim 1, wherein the oligonucleotide compound to which the cells are exposed has a sequence contained in SEQ ID NO:10.

7. The method of claim 6, wherein said sequence of the oligonucleotide compound is selected from the group consisting of SEQ ID NOS:11, 12, and 13.

8. The method of claim 1, wherein the compound to which the cells are exposed is conjugated to an arginine-rich polypeptide effective to promote uptake of the compound into infected host cells.

9. The method of claim 8, wherein the arginine rich peptide has one of the sequences identified as SEQ ID NOS:14-19.

10. The method of claim 1, for use in treating a mammalian subject infected by an *Enterovirus* or *Rhinovirus* in the Picornaviridae family, wherein said exposing includes administering to the subject, a pharmaceutically effective amount of the oligonucleotide compound, and which further includes continuing said administering until a significant reduction in viral infection or the symptoms thereof is observed.

11. The method of claim 10, which further includes administering a second anti-viral compound to the subject.

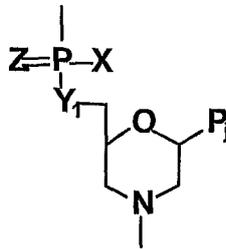
12. The method of claim 1, for use in treating a mammalian subject at risk of infection by an *Enterovirus* or *Rhinovirus* in the Picornaviridae family, wherein said exposing includes administering to the subject, an amount of the oligonucleotide compound effective to inhibit infection of subject host cells by the virus.

13. An antisense oligonucleotide compound for use in inhibiting viral infection in mammalian cells by an *Enterovirus* or *Rhinovirus* in the Picornaviridae family, characterized by:

- (i) a substantially uncharged, nuclease-resistant backbone,
- (ii) capable of uptake by mammalian host cells,
- (iii) containing between 12-40 nucleotide bases,
- (iv) having a targeting sequence of at least 12 subunits contained in SEQ ID NO: 10; and
- (v) capable of binding to the virus' positive sense strand to form a heteroduplex structure having by a T_m of dissociation of at least 45°C.

14. The compound of claim 13, which is substantially uncharged and composed of morpholino subunits and phosphorus-containing intersubunit linkages joining a morpholino nitrogen of one subunit to a 5' exocyclic carbon of an adjacent subunit.

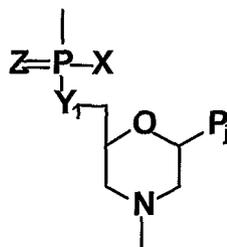
15. The compound of claim 14, wherein said morpholino subunits are joined by phosphorodiamidate linkages having the structure:



where $Y_i=O$, $Z=O$, P_j is a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, and X is alkyl, alkoxy, thioalkoxy, amino or alkyl amino, including dialkylamino.

16. The compound of claim 14, in which at least 2 and no more than half of the total number of intersubunit linkages are positively charged at physiological pH.

17. The compound of claim 16, wherein said morpholino subunits are joined by phosphorodiamidate linkages, in accordance with the structure:



where $Y_i=O$, $Z=O$, P_j is a purine or pyrimidine base-pairing moiety effective to bind, by base-specific hydrogen bonding, to a base in a polynucleotide, and X for the uncharged linkages is alkyl, alkoxy, thioalkoxy, or an alkyl amino of the form wherein NR_2 , where each R is independently hydrogen or methyl, and for the positively charged linkages, X is 1-piperazine.

18. The compound of claim 13, wherein said sequence of the oligonucleotide compound is selected from the group consisting of SEQ ID NO:11, SEQ ID NO:12, and SEQ ID NO:13.

19. The compound of claim 13, which is conjugated to an arginine-rich polypeptide effective to promote uptake of the compound into infected host cells.

20. The compound of claim 19, wherein the arginine rich peptide has one of the sequences identified as SEQ ID NOS: 14-19.

21. The compound of claim 1, in combination with another anti-viral compound.

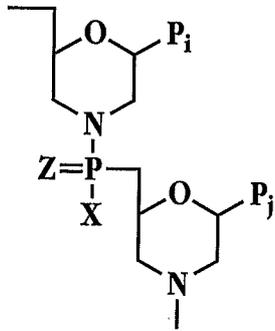


Fig. 1A

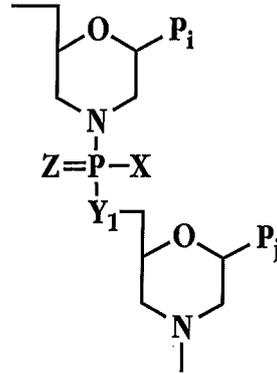


Fig. 1B

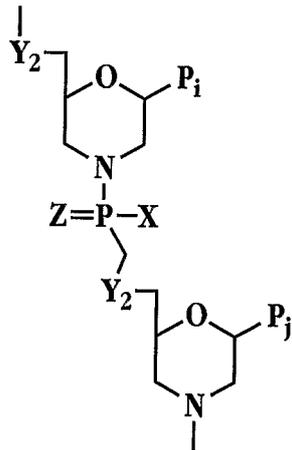


Fig. 1C

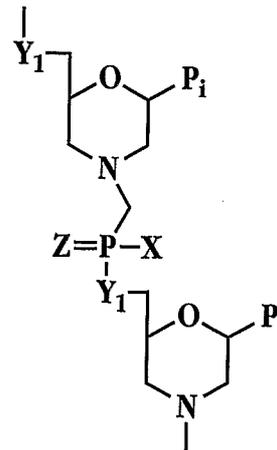


Fig. 1D

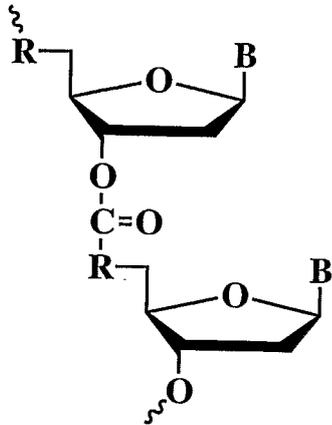


Fig. 2A

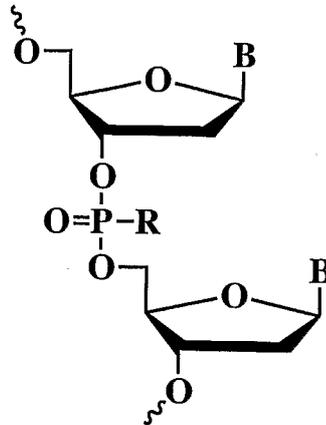


Fig. 2B

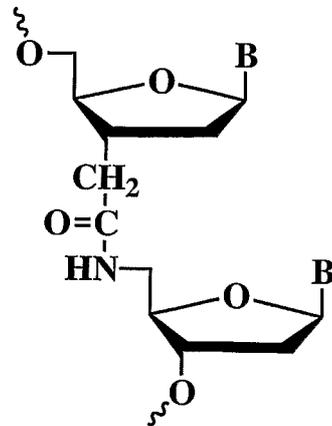


Fig. 2C

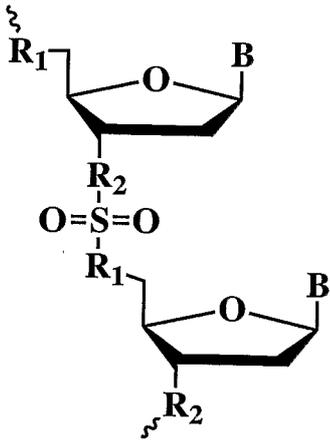


Fig. 2D

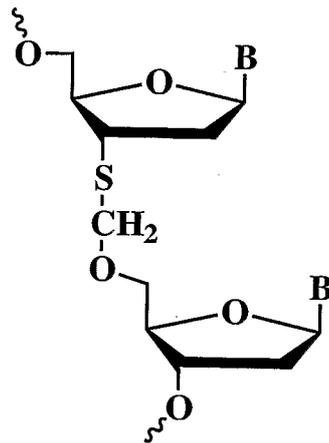


Fig. 2E

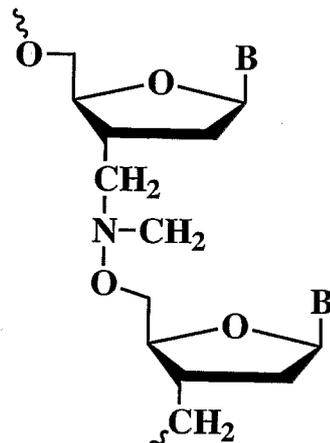


Fig. 2F

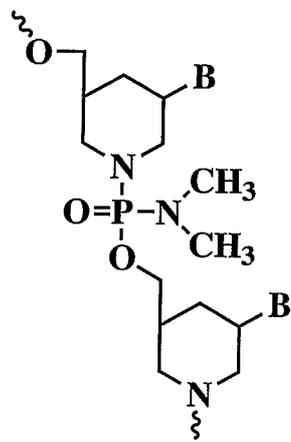


Fig. 2G

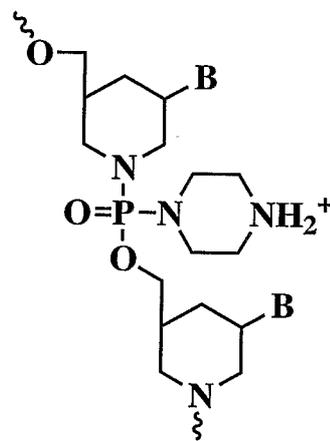


Fig. 2H

***** ** * ** *
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SEQ ID NOS: 4 TCCTCCGGCCCCTGAATGCGGTTAATCCTAAC
2 TCCTCCGGCCCCTGAATGCGGTTAATCCTAAC
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← 5'-32 Target Region →

Fig. 3

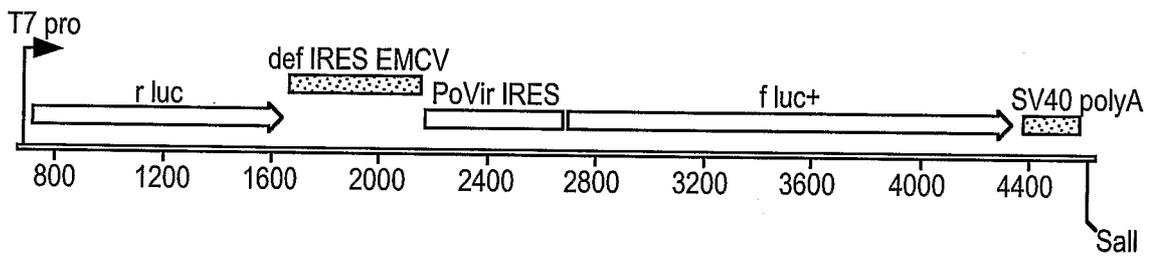


Fig. 4

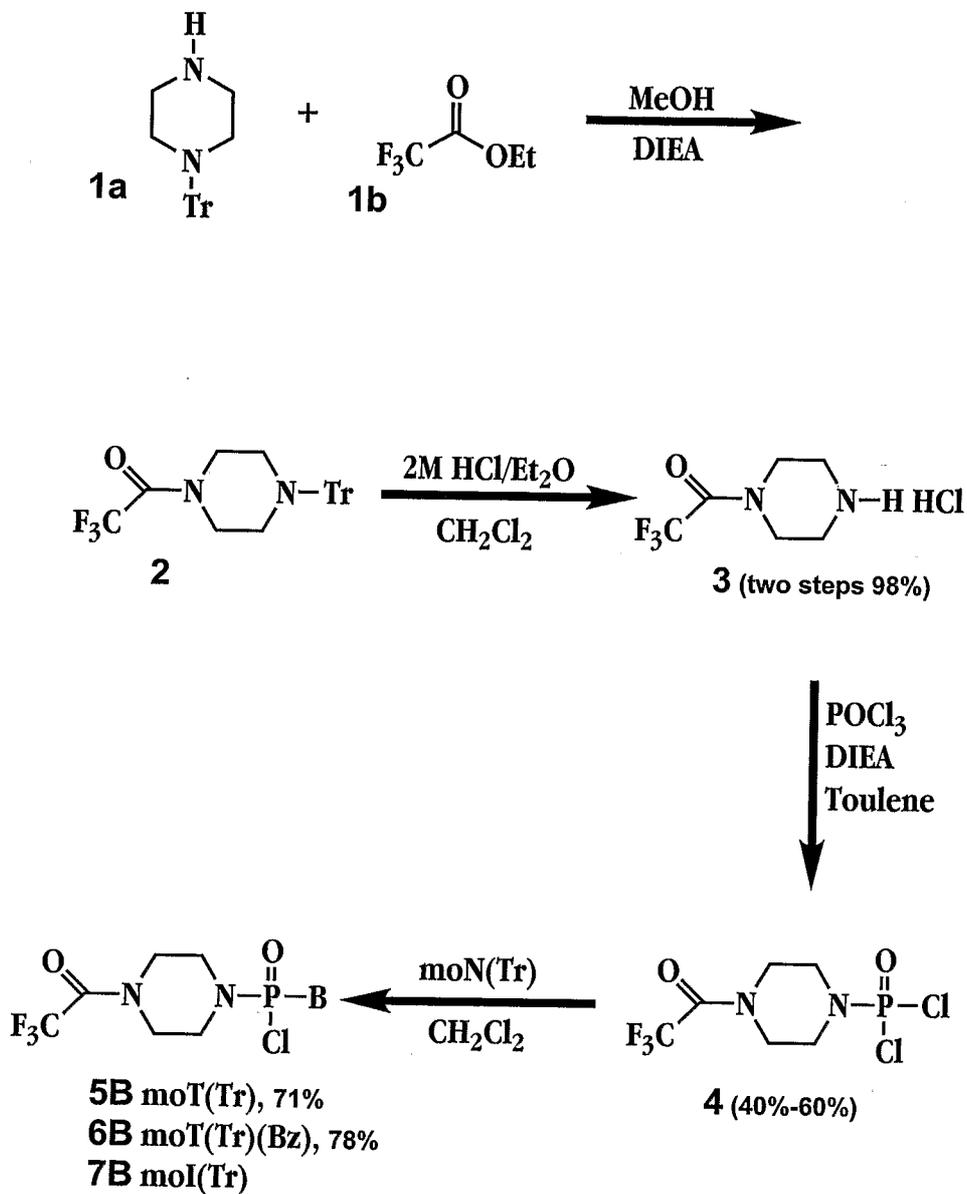


Fig. 5

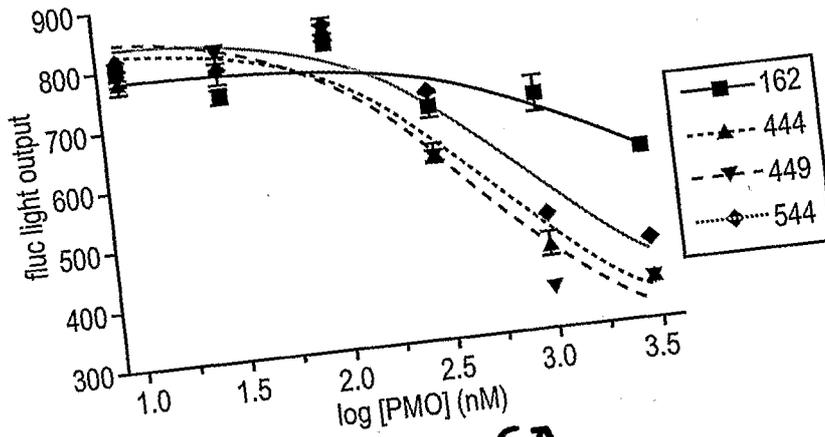


Fig. 6A

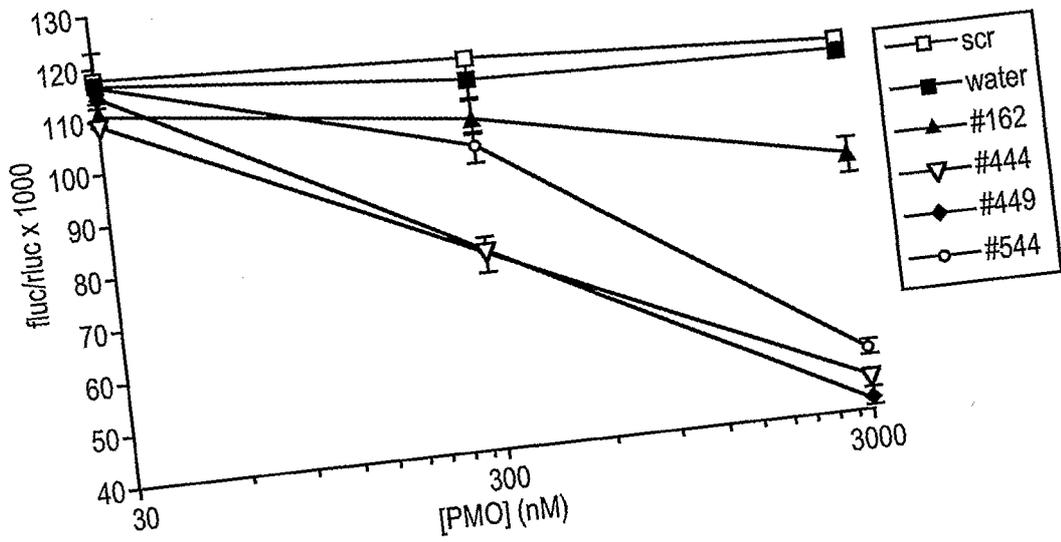


Fig. 6B

SEQUENCE LISTING

<110> Stein, David A .
Bestwick, Richard K .
Iversen, Patrick L .
Weller, Dwight D .

<120> ANTISENSE ANTIVIRAL COMPOUND AND METHOD
FOR TREATING PICORNAVIRUS INFECTION

<130> 504508077W00

<140> Not Yet Assigned
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<150> US 60/800,145
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