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#### ANTIVIRAL AGENTS FOR THE (54)TREATMENT, CONTROL AND PREVENTION OF INFECTIONS BY CORONAVIRUSES

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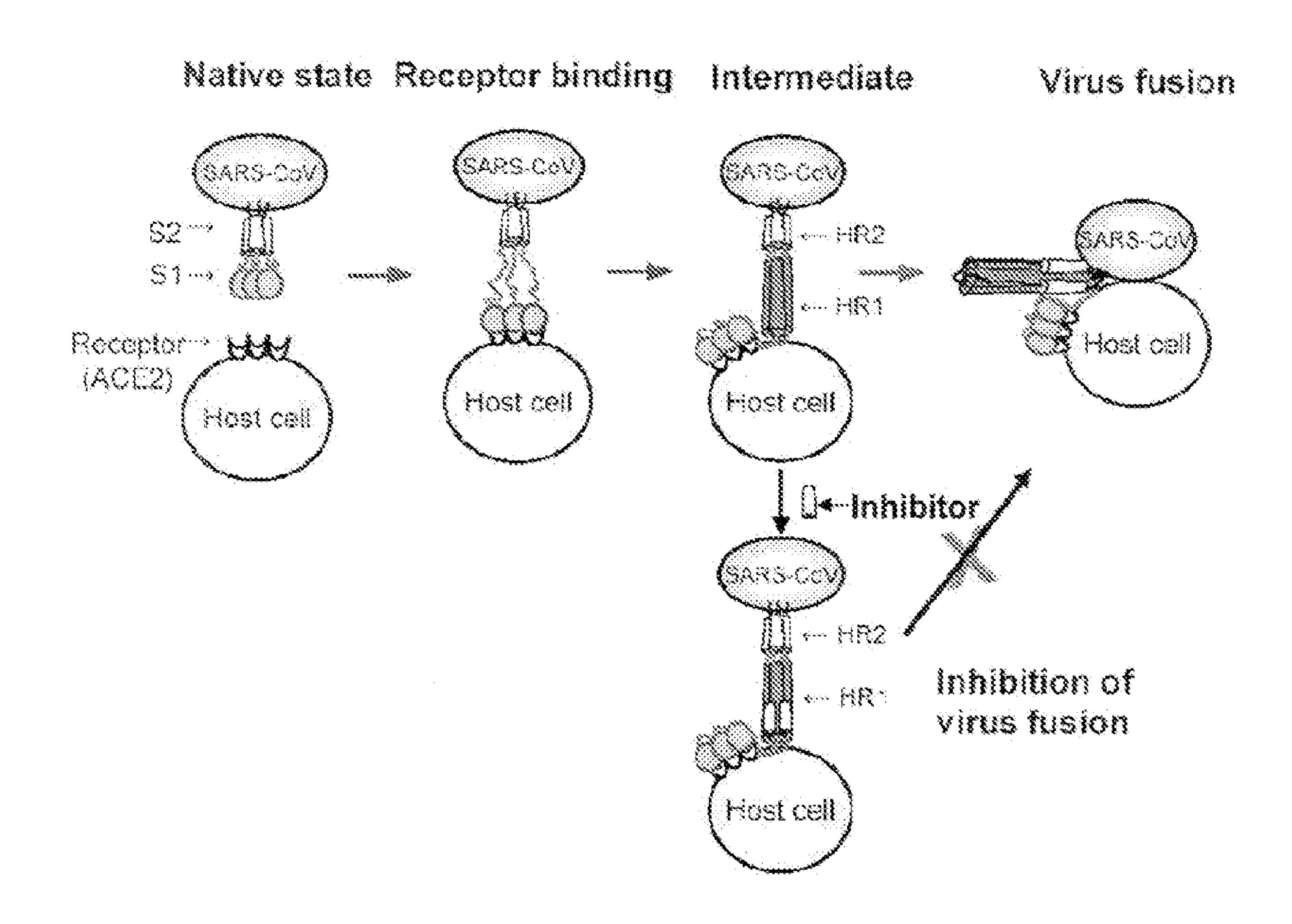
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#### **ABSTRACT** (57)

The invention provides compositions and methods that are useful for preventing and treating a coronavirus infection in a subject. More specifically, the invention provides peptides and conjugates and pharmaceutical compositions containing those peptides and conjugates that block fusion of a coronavirus, such as the SARS virus, to a target cell. This blocking mechanism prevents or treats a coronavirus infection, such as a SARS infaction, in a subject, such as a human subject.

# Figure 1



# Mignre 2

# ANTIVIRAL AGENTS FOR THE TREATMENT, CONTROL AND PREVENTION OF INFECTIONS BY CORONAVIRUSES

[0001] This is a continuation of prior application Ser. No. 10/833,304, filed Apr. 28, 2004, which claims priority to 60/466,432, filed Apr. 30, 2003 and 60/465,782 filed Apr. 28, 2003, the entireties of which are hereby incorporated by reference.

# BACKGROUND

[0002] Severe Acute Respiratory Syndrome (SARS) is an emerging new infectious disease caused by a novel coronavirus that infects humans. See Ksiazek et al., New Engl. J (http://content.nejm.org/cgi/reprint/ Med.NEJMoa030781v2.pdf, published Apr. 10, 2003). SARS is fatal in about 4-10% of cases reported so far. Initially described in mid February, 2003 in China's Guangdong province as atypical pneumonia, by mid-March, 2003 the World Health Organization (WHO) had received reports of more than 150 new suspected cases of unknown origin or cause. By mid April, 2003, over 4400 cases with 263 deaths of patients diagnosed with symptoms of SARS have been documented from 26 different countries, including Canada, China, Hong Kong, Indonesia, Philippines, Singapore, Thailand, Viet Nam and the United States. In light of the rapid spread of SARS to several countries in a short period of time, the World Health Organization issued a global alert and provided emergency guidance for travellers and airlines. In only a few months after the outbreak was first recognized, SARS became a worldwide threat to global health and global economies. There are presently no known therapies that are effective against SARS, and no vaccine is available. Accordingly, there is an urgent need for antiviral agents that can control or prevent SARS in infected individuals, and that can prevent SARS from spreading.

[0003] In general, SARS begins with a fever greater than 100.4° F. [>38.0° C.]. Other symptoms may include headache, an overall feeling of discomfort, and body aches. Some people also experience mild respiratory symptoms. After 2 to 7 days, SARS patients may develop a dry cough and have trouble breathing.

[0004] The primary way that SARS appears to spread is by close person-to-person contact. Most cases of SARS have involved people who cared for or lived with someone with SARS, or had direct contact with infectious material (for example, respiratory secretions) from a person who has SARS. Potential ways in which SARS can be spread include touching the skin of other people or objects that are contaminated with infectious droplets followed by touching of eye(s), nose, or mouth. This can happen when someone who is sick with SARS coughs or sneezes droplets onto themselves, other people, or nearby surfaces. It also is possible that SARS can be spread more broadly through the air or by other ways that are currently not known.

[0005] Scientists at the Centers for Disease Control and Prevention (CDC) and other laboratories around the world have detected a previously unrecognized coronavirus in patients with SARS. The evidence for a coronavirus was based on genetic fingerprint and electron microscopic ultrastructural studies and was widely reported in the popular press. Viologists at the CDC, WHO and numerous academic

laboratories all reported that a coronavirus is the leading hypothesis for the cause of SARS.

[0006] The CDC recently reported sequencing the genome for SARS-CoV (Urbani strain), a strain of a novel human coronavirus believed to be responsible for SARS. The sequence data confirm that the SARS virus is a previously unrecognized coronavirus. The virus was cultured from cells taken from a throat culture taken from a SARS patients and grown in Vero cells (African green monkey kidney cells) in order to reproduce the ribonucleic acid (RNA) of the disease-causing coronavirus. The new sequence has 29,727 nucleotides, which places it well within the typical RNA boundaries for coronaviruses. Members of this viral family tend to have between 29,000 and 31,000 nucleotides. See Lai et al., *Adv. Virus Res.* 48:1, (1997). The genome organization of the SARS virus also is similar to that of other coronaviruses.

[0007] The genome sequence of SARS-CoV (Urbani) is available from GenBank at the Web site for the National Center for Biotechnology Information, National Library of Medicine httfp://www.ncbi.nlm.nih.gov/. The accession number for the sequence of SARS-CoV (Urbani strain) is ay278741. The present inventors have used these sequence data to identify molecular targets that can be exploited to design safe and effective novel antiviral therapies that can be used to treat SARS and to stem the tide of the growing epidemic.

# SUMMARY OF THE INVENTION

[0008] In accordance with a first aspect of the invention there is provided an antiviral peptide having between 7 and 50 amino acids, where the peptide exhibits antiviral activity against a coronavirus, and where the peptide contains a sequence comprising at least 7 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 1)

DV**DL**GDISGINAS**V**VNIQKEIDR**L**NE**VA**KN**L**NES**L**IDLQE**LG**K**Y**EQY**I**K;

(SEQ ID NO: 2)

QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3)

ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4)

GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5)

RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

[0009] In accordance with a second aspect of the invention there is provided an antiviral peptide having between 7 and 50 amino acids, where the peptide exhibits antiviral activity against a coronavirus, and where the peptide contains a sequence comprising at least 7 contiguous amino acids from the sequence:

(SEQ ID NO: 1)

DV<u>DL</u>GD<u>I</u>SG<u>I</u>NAS<u>V</u>VN<u>I</u>QKE<u>I</u>DR<u>L</u>NE<u>VA</u>KN<u>L</u>NES<u>L</u>IDLQE<u>LG</u>K<u>Y</u>EQY<u>I</u>K;;

[0010] where the amino acids at bold letter positions can be substituted with an amino acid selected from the group

consisting of I, L, V, W, Y, F, N, Q, S, T, D, E, G, H, and M, and where amino acids in non-bold positions can be any amino acid except proline.

[0011] The peptides above may contain a sequence selected from the group consisting of

SVVNIQK (SEQ ID		6)	VVNIQKE (SEQ ID		7)	VNIQKEI (SEQ ID	NO:	8)
NIQKEID (SEQ ID	NO:	9)	IQKEIDR (SEQ ID	NO:	10)	QKEIDRL (SEQ ID	NO:	11)
KEIDRLN (SEQ ID	NO:	12)	EIDRLNE (SEQ ID	NO:	13)	IDRLNEV (SEQ ID	NO:	14)
DRLNEVA (SEQ ID	NO:	15)	RLNEVAK (SEQ ID	NO:	16)	LNEVAKN (SEQ ID	NO:	17)
NEVAKNL (SEQ ID	NO:	18)	EVAKNLN (SEQ ID	NO:	19)	VAKNLNE (SEQ ID	NO:	20)
AKNLNES (SEQ ID	NO:	21)	KNLNESL (SEQ ID	NO:	22)	NLNESLI (SEQ ID	NO:	23)
LNESLID (SEQ ID	NO:	24)	NESLIDL (SEQ ID	NO:	25)	ESLIDLQ (SEQ ID	NO:	26)
SLIDLQE (SEQ ID	NO:	27)	LIDLQEL (SEQ ID	NO:	28)	IDLQELG (SEQ ID	NO:	29)
DLQELGK (SEQ ID	NO:	30)	LQELGKY (SEQ ID	NO:	31)	QELGKYE (SEQ ID	NO:	32)
ELGKYEQ (SEQ ID		33)	LGKYEQY (SEQ ID			GKYEQYI (SEQ ID		35)
KYEQYIK (SEQ ID			QIPFAMQ (SEQ ID		37)	IPFAMQM (SEQ ID		38)
PFAMQMA (SEQ ID		39)	FAMQMAY (SEQ ID		40)	AMQMAYR (SEQ ID	NO:	41)
MQMAYRF (SEQ ID			QMAYRFN (SEQ ID			MAYRFNG (SEQ ID	NO:	44)
AYRFNGI (SEQ ID	NO:	45)	YRFNGIG (SEQ ID	NO:		RFNGIGV (SEQ ID	NO:	47)
FNGIGVT (SEQ ID		48)	NGIGVTQ (SEQ ID		49)	IGVTQNV (SEQ ID	NO:	50)
GVTQNVL (SEQ ID	NO:	51)	VTQNVLY (SEQ ID	NO:	52)	TQNVLYE (SEQ ID	NO:	53)
QNVLYEN (SEQ ID	NO:	54)	NVLYENQ (SEQ ID			VLYENQK (SEQ ID		56)
LYENQKQ (SEQ ID	NO:	57)	YENQKQI (SEQ ID		58)	ENQKQIA (SEQ ID		59)
NQKQIAN (SEQ ID	NO:		QKQIANQ (SEQ ID		61)	KQIANQF (SEQ ID		62)
QIANQFN (SEQ ID	NO:	63)	IANQFNK (SEQ ID			ANQFNKA (SEQ ID		65)
NQFNKAI (SEQ ID			QFNKAIS (SEQ ID					68)
NKAISQI (SEQ ID		69)	KAISQIQ (SEQ ID					71)
ISQIQES (SEQ ID	NO:	72)	SQIQESL (SEQ ID	NO:	73)	QIQESLT (SEQ ID	NO:	74)
ESLTTTS			SLTTTST			LTTTSTA		

					_			
(SEQ II	NO:	75)	-cont (SEQ ID			(SEQ ID	NO:	77)
TTTSTAI (SEQ II		78)	TTSTALG (SEQ ID	NO:	79)	TSTALGK (SEQ ID	NO:	80)
STALGKI (SEQ II		81)	TALGKLQ (SEQ ID		82)	ALGKLQD (SEQ ID	NO:	83)
LGKLQDV		84)	GKLQDVV (SEQ ID	NO:	85)	KLQDVVN (SEQ ID	NO:	86)
LQDVVNÇ (SEQ II		87)	QDVVNQN (SEQ ID			DVVNQNA (SEQ ID	NO:	89)
VVNQNAÇ (SEQ II		90)	VNQNAQA (SEQ ID			NQNAQAL (SEQ ID	NO:	92)
QNAQALI (SEQ II		93)	NAQALNT (SEQ ID				NO:	95)
QALNTLV		96)	ALNTLVK (SEQ ID	NO:		LNTLVKQ (SEQ ID		98)
NTLVKQI (SEQ II		99)	TLVKQLS (SEQ ID			LVKQLSS	NO:	101)
VKQLSSI (SEQ II		102)	KQLSSNF (SEQ ID			QLSSNFG (SEQ ID	NO:	104)
LSSNFGA (SEQ II		105)	SSNFGAI (SEQ ID		106)	SNFGAIS	NO:	107)
NFGAISS (SEQ II		108)	LQDVVNQ (SEQ ID				NO:	110)
DVVNQNA (SEQ II		111)	VVNQNAQ (SEQ ID			VNQNAQA (SEQ ID	NO:	113)
NQNAQAI (SEQ II		114)	QNAQALN (SEQ ID	NO:	115)	NAQALNT (SEQ ID	NO:	116)
AQALNTI (SEQ II		117)	QALNTLV (SEQ ID				NO:	119)
LNTLVKÇ (SEQ II		120)	NTLVKQL (SEQ ID				NO:	122)
			VKQLSSN (SEQ ID				NO:	125)
QLSSNFO		126)	LSSNFGA (SEQ ID		127)	SSNFGAI (SEQ ID	NO:	128)
SNFGAIS (SEQ II			NFGAISS (SEQ ID				NO:	131)
GAISSVI (SEQ II		132)	AISSVLN (SEQ ID	NO:	133)	ISSVLND (SEQ ID	NO:	134)
SSVLNDI (SEQ II			SVLNDIL (SEQ ID			VLNDILS (SEQ ID	NO:	137)
LNDILSE (SEQ II		138)	NDILSRL (SEQ ID	NO:	139)	DILSRLD (SEQ ID	NO:	140)
ILSRLDE (SEQ II		141)	LSRLDK (SEQ ID	NO:	142)	SRLDKV (SEQ ID	NO:	143)
RLDKVE (SEQ II	NO:	144)	LDKVEA,	NO:	145)	RLITGRL (SEQ ID	NO:	146)
LITGRLÇ (SEQ II	_	147)	ITGRLQS (SEQ ID		148)	TGRLQSL (SEQ ID	NO:	149)
GRLQSLÇ (SEQ II			RLQSLQT (SEQ ID			LQSLQTY (SEQ ID	NO:	152)

OCT OFFI		-cont			т опши	7.TI.O		
QSLQTYV (SEQ ID NO:	153)	SLQTYVT (SEQ ID					NO:	155)
QTYVTQQ		TYVTQQL			YVTQQ	T.T		
(SEQ ID NO:	156)		NO:			_	NO:	158)
VTQQLIR		TQQLIRA			QQLIF	RAA		
(SEQ ID NO:	159)	(SEQ ID	NO:	160)	(SEQ	ID	NO:	161)
QLIRAAE		LIRAAE			IRAAE			
(SEQ ID NO:	162)	(SEQ ID	NO:	163)	(SEQ	ID	NO:	164)
RAAEIRA	165)	AAEIRAS					NO.	167)
(SEQ ID NO:	165)	(SEQ ID	NO:	100)	(SEQ	ΤD	NO:	167)
EIRASAN		IRASANL			RASAN	ILA		
(SEQ ID NO:	168)	(SEQ ID	NO:	169)	(SEQ	ID	NO:	170)
ASANLAA		SANLAAT			ANLAA	ΥK		
(SEQ ID NO:	171)						NO:	173)
NLAATKM		LAATKMS			AATKM	ISE		
(SEQ ID NO:	174)	(SEQ ID	NO:	175)	(SEQ	ID	NO:	176)
ATKMSEC		TKMSECV			KMSEC	VL	and	
(SEQ ID NO:	177)							179)
MSECVLG.								
(SEQ ID NO:	180).							

[0012] The peptide may contain at least 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from one of the sequences:

(SEQ ID NO: 181)
VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2)
QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3)

ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE;

and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

[0013] Any of the peptides above may be linked to a carrier protein, such as human serum albumin, for example.

[0014] In accordance with a third aspect of the invention there is provided an antiviral composition comprising a peptide X having between 7 and 50 amino acids, where the peptide exhibits antiviral activity against a coronavirus, and where the composition has the structure:

B-X-Z,

[0015] where B is an amino acid sequence containing up to about 43 amino acids, or B is an amino group, an acetyl group, a 9-fluorenylmethoxy-carbonyl group, a hydrophobic group, or a macromolecule carrier group, or B is a carrier protein, in which case B may contain more than 8 amino acids, and may also comprises a linker peptide sequence that connects the antiviral sequence to the carrier protein; Z is an amino acid sequence containing up to about 43 amino acids, or Z comprises a carboxyl group, an amido group, a hydrophobic group, or a macromolecular carrier group, or Z is a carrier protein, such as HSA, in which case Z may contain more than 8 amino acids, and may also comprise a linker

peptide sequence that connects the antiviral sequence to the carrier protein; where when considered together B and Z must contain at least 8 amino acids between the B and Z groups; and where X is a peptide sequence comprising at least 7 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 181)
VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2)
QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLYKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

[0016] X may contain, for example, at least 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 181)
VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2)
QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

[0017] In accordance with a fourth aspect of the invention there is provided an antiviral peptide having between 7 and 50 amino acids, where the peptide exhibits antiviral activity against a coronavirus, and where the peptide comprises a sequence that exhibits identity in any two of the seven positions of a contiguous heptapeptide, where the contiguous heptapeptide comprises 7 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 181)
VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2)
QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

[0018] The sequence identity may be located, for example, in the ith and i+4th positions in the contiguous heptapeptides.

[0019] In accordance with another aspect of the invention there is provided a pharmaceutical composition comprising a peptide or composition as described above and a pharmaceutically acceptable diluent, adjuvant and/or excipient.

[0020] In accordance with yet another aspect of the invention there is provided a method of treating or preventing a coronavirus infection in a subject, comprising administering to a patient suspected of suffering from the infection an effective amount of a peptide or composition as described above. The subject may be a human, a cow, pig, or chicken.

[0021] Other objects, features and advantages of the present invention will become apparent from the following detailed description. It should be understood, however, that the detailed description and the specific examples, while indicating preferred embodiments of the invention, are given by way of illustration only, since various changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art from this detailed description.

# BRIEF DESCRIPTION OF THE DRAWINGS

[0022] FIG. 1 shows a model of SARS-CoV fusing with the host cell and of an inhibitor blocking fusion between SARS-CoV and the host cell membrane. SARS-CoV contains a surface spike protein consisting of S1 and S2 domains. SARS-CoV binds to the host cell through interaction between S1 domain and the host cell receptor, ACE2. The HR2 helices of the S2 protein fold over and interact with the HR1 helices to form "hairpin-like" structures, which draw both the viral and host cell membranes together for fusion. An inhibitor is shown in the lower portion binding to the HR1 trimer and blocking association of HR1 and HR2, thereby inhibiting SARSCoV fusion with the host cell. WWW.nybloodcenter.org/pdf/Anti-Model from SARS%20Peptide%20Model.pdf by Dr. Shibo Jiang.

[0023] FIG. 2 shows some representative examples of results of secondary structure prediction and homology analyses on peptides (SEQ ID NOS 197 and 2, respectively in order of appearance) from SARS coronavirus, isolate Tor2, E2 glycoprotein precursor. Amino acids in the most highly predicted helical regions are listed in bold. E, H and L designations are from the secondary structure prediction algorithm and refer to extended or coil, helix and loop regions, respectively. Numbers refer to the probable accuracy of the prediction, from lowest (0) to highest(9). The last row indicates percent amino acid sequence homology among the various S2 proteins of coronaviruses tested. Residues shown are conserved in over 70% of all S2 coronavirus proteins; residues in SARS S2 protein that are identical to the consensus are highlighted in bold.

# DESCRIPTION OF THE INVENTION

[0024] The invention provides compositions and methods that are useful for preventing and treating a coronavirus infection in a subject. More specifically, the invention provides peptides and conjugates and pharmaceutical compositions containing those peptides and conjugates that block fusion of a coronavirus, such as the SARS virus, to a target cell. This blocking mechanism prevents or treats a coronavirus infection, such as a SARS infection, in a subject, such as a human subject.

SARS-CoV is a Novel Coronavirus

[0025] The sequence of the genome of the SARS-CoV was downloaded from the CDC website and translated into

ten putative open reading frames (ORFs). The amino acid sequences corresponding to putative proteins encoded by the ten ORFs were analyzed for homology to existing proteins in the proteome using BLAST, a protein database searching program. See Altschul et al, *Nucleic Acids Res.* 25: 3389 (1997). A number of open reading frames were found to encode proteins with significant sequence homology to proteins from known coronaviruses. For example, ORF1 corresponds to a coronavirus polymerase protein (polymerase 1a, 1b), and ORF3 corresponds to a coronavirus spike protein (S). The homology and organization of the genome provide additional convincing evidence that the SARS virus is a coronavirus.

[0026] Coronaviruses previously have been grouped into three categories based on cross-reactivity of antibodies backed up by genetic data. The two previously identified human coronaviruses fall into two different groups. One of these groups includes a number of enteric coronaviruses that cause gastroenteritis. The other includes coronaviruses that cause respiratory or neurological diseases in diverse species. The third group includes coronaviruses isolated from avian species.

[0027] The sequence of the S protein from SARS-CoV shares about 30-35% identity throughout its 1260 or so amino acid length with S proteins from all three groups of coronaviruses, including coronaviruses from humans, cows, pigs, mice and chickens. Based on the sequence homology analysis, the SARS coronavirus represents the first, and so far only, member of a new fourth coronavirus group.

Spike Proteins are Required for Viral Entry

[0028] Numerous studies have shown that entry of enveloped viruses into host cells requires membrane fusion between virus and host cell. For most animal viruses, this fusion function is mediated by a single envelope glycoprotein on virions. The S protein has been shown to be the fusion protein that mediates cellular entry for coronavirus. See Spaan et al., *J Gen. Virol.* 69:2939 (1988).

[0029] The S protein forms the peplomer projections that protrude from the virion surface as seen in electron micrographs. Peplomers are thought to be composed of three oligomerized S protein molecules. See Delmas et al., *J Virol*. 64:5367 (1990). The S protein is cleaved by host proteases during virus assembly into two similarly-sized subunits: S1 and S2. The C-terminal S2 subunit, which associates non-covalently with the N-terminal S1, anchors the S protein to the membrane through a transmembrane domain, while the S1 subunit contains the receptor binding activity of the S protein.

Helical Heptad Repeats in the S2 Subunit are Required for Fusion

[0030] Reviews of the roles of the coronavirus spike proteins in viral entry and pathogenesis can be found in Gallagher et al., *Virology* 279:371 (2001) and Luo et al, *J. Virol.* 73:8152 (1999). Several studies suggest that the S2 subunit is required for viral fusion. Functional mutagenesis studies indicate that critical residues for fusion are located within two regions in S2 that have been identified as heptad repeat regions. Helical heptad repeats are found in fusion proteins from other enveloped viruses, including paramyx-oviruses, such as influenza virus, retroviruses, such as HIV, and filoviruses, such as Ebola virus. The existence of two heptad repeats, HR1 and HR2, separated by a non-helical

spacer in the S2 subunit of coronaviruses is suggestive of the formation of a coiled-coil or "trimer of hairpins" fusogenic complex similar to the fusogenic structures thought to be formed by the helical heptad repeats in the fusion glycoproteins of, for example, HIV, influenza virus and Ebola virus.

[0031] In the case of HIV, a peptide sequence that mimics either HR1 or HR2 can prevent HIV fusion and block viral replication. Enfurvitide, a 38-residue peptide based on the sequence of HR2 of the HIV glycoprotein, is used clinically to treat HIV/AIDS. The present inventors reasoned that peptides that can bind to coiled-coil intermediates of coronaviruses should block the formation of a productive fusogenic complex and prevent virus entry (FIG. 1).

SARS-CoV Contains High Homology Regions with HR1 and HR2 of Coronaviruses that are Predicted to be Helical.

[0032] The amino acid sequence homology between the S protein of SARS-CoV and other coronaviruses was evaluated throughout its length. The highest homology resides in regions that overlap with the HR1 and HR2 heptad repeats in known coronoaviruses. Analysis of the S protein using secondary structure prediction methods (see for example program PROF as implemented on the CUBIST protein prediction server @http://cubic.bioc.columbia.edu)<sub>13</sub> revealed that the HR1 and HR2 regions are strongly predicted to be helical (FIG. 2). The HR1 and HR2 regions were divided into five contiguous amino acids segments that are most strongly predicted to be helical. In the same fashion as observed with HIV gp41, heptad-containing sequences derived from the HR1 segments bind to HR2 helices. Similarly, heptad-containing sequences derived from the HR2 segments can bind to the HR1 helices. Peptides that bind to either HR1 or HR2 prevent virus entry, possibly by disrupting formation of the fusogenic complex. Such peptides, and compositions containing these peptides, are useful for treating infections caused by coronaviruses, and as prophylactics against coronavirus infection. These peptides are particularly useful for preventing and treating SARS infection.

[0033] The N peptides are predicted to form a trimeric coiled-coil with 3-fold symmetry similar to HR1 of HIV-gp41. The C peptides are predicted to form helices that bind in the grooves formed by adjacent N-helices in the coiled coil, similar to HR2 in HIV gp41. These peptide sequences are predicted to form continuous alpha helices constructed of series of contiguous, or nearly contiguous, helical heptad repeats in which the ith and i+4th residues are important or critical for oligomer formation.

[0034] The peptides contained in the S protein have the following sequences (most strongly predicted helical sequences are shown in bold)

C-terminal, HR2 peptide 1 (SEQ ID NO: 182)
PDVDLGDISGINASVVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK

N-Terminal, HR1 peptide 2 (SEQ ID NO: 2) QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT

N-Terminal, HR1 peptide 3 (SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS

N-Terminal, HR1 peptide 4 (SEQ ID NO: 4)
GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE

-continued
N-Terminal, HR1 peptide 5 (SEQ ID NO: 183)
QALNTLVKQLSSNFGAISSVLNDILSRLDKVEAEVQIDRLITGRLQS

N-Terminal, HR1 peptide 6 (SEQ ID NO: 5)
RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF

[0035] These sequences were used to design peptide inhibitors having between about 15 and about 50 amino acids, advantageously having 15 to 50 amino acids and containing at least 7 contiguous amino acids from one of the sequences shown above. Thus, for example, the peptides may contain one of the following 7 amino acid sequences:

[0036] for the C-terminal HR2 peptide 1 the peptide may contain one of the following sequences

PDVDLGD (SEQ ID									NO •	186)
									NO.	100,
DLGDISG (SEQ ID									NO:	189)
DISGINA			ISGIN			101)			110	100
(SEQ ID	NO:	190)	(SEQ	ID	NO:	191)	(SEQ	ID	NO:	192)
GINASVV (SEQ ID							NASVV		NO •	195)
									110.	100,
ASVVNIQ (SEQ ID							VVNIÇ (SEQ	-	NO:	7)
VNIQKEI			NIQKE				IQKE			
(SEQ ID	NO:	8)	(SEQ	ID	NO:	9)	(SEQ	ID	NO:	10)
QKEIDRL						4.0.	EIDRI			10)
(SEQ ID	NO:	11)	(SEQ	ID	NO:	12)	(SEQ	ID	NO:	13)
IDRLNEV			DRLNE		N/O -		RLNEV		NO -	16)
(SEQ ID	NO:	14)	(SEQ	תד	NO:	15)	(SEQ	TD	NO:	16)
LNEVAKN			NEVAK	NL			EVAKI	$1\Gamma N$		
(SEQ ID	NO:	17)	(SEQ	ID	NO:	18)	(SEQ	ID	NO:	19)
VAKNLNE			AKNLN	IES			KNLNE	EST.		
(SEQ ID									NO:	22)
NLNESLI			LNESI	JD			NESLI	DL		
(SEQ ID	NO:	23)	(SEQ	ID	NO:	24)	(SEQ	ID	NO:	25)
ESLIDLQ			SLIDI	ΩE			LIDLÇ	ŒL		
(SEQ ID	NO:	26)	(SEQ	ID	NO:	27)	(SEQ	ID	NO:	28)
IDLQELG			DLOET	GK			LQELO	KY:		
(SEQ ID					NO:	30)			NO:	31)
QELGKYE			ELGKY	ÆQ.			LGKYE	EQY		
(SEQ ID	NO:	32)	(SEQ	ID	NO:	33)	(SEQ	ID	NO:	34)
GKYEQYI			KYEOY	ZIK						
(SEQ ID					NO:	36				
for the	N-Te	rmina	al, HI	R1 ]	pept:	ide 2 t	he pe	epti	lde m	ıay
contain	one	of th	ne foi	llo	wing	sequer	ices			
QIPFAMQ										
(SEQ ID	NO:	37)	(SEQ	ID	NO:	38)	(SEQ	ID	NO:	39)
FAMQMAY			AMQMA	ΔYR			MQMAY	/RF		
(SEQ ID									NO:	42)
QMAYRFN			MAYRF	'nG			AYRFI	IGI		
(SEQ ID	NO:	43)	(SEQ	ID	NO:	44)	(SEQ	ID	NO:	45)
YRFNGIG			RFNGI	GV			FNGIO	TV		
(SEQ ID	NO:				NO:	47)			NO:	48)

				<del> </del>	
NGIGVTQ	-continued IGVTQNV	GVTQNVL	VVNQNAQ	-continued VNQNAQA	NQNAQAL
(SEQ ID NO: 49)	(SEQ ID NO: 50)	(SEQ ID NO: 51)		(SEQ ID NO: 113)	(SEQ ID NO: 114)
(012 10 10)	(019 10 100)	(OLG ID NOT)	(DIG ID NOT III)	(019 10 100	(010 10 111)
VTQNVLY	TQNVLYE	QNVLYEN	QNAQALN	NAQALNT	AQALNTL
(SEQ ID NO: 52)	(SEQ ID NO: 53)	(SEQ ID NO: 54)	(SEQ ID NO: 115)	(SEQ ID NO: 116)	(SEQ ID NO: 117)
NVLYENQ	VLYENQK	LYENQKQ	QALNTLV	ALNTLVK	LNTLVKQ
(SEQ ID NO: 55)	(SEQ ID NO: 56)	(SEQ ID NO: 57)	(SEQ ID NO: 118)	(SEQ ID NO: 119)	(SEQ ID NO: 120)
YENQKQI	ENQKQIA	NQKQIAN	NTLVKQL	TLVKQLS	LVKQLSS
(SEQ ID NO: 58)	(SEQ ID NO: 59)	(SEQ ID NO: 60)		(SEQ ID NO: 122)	(SEQ ID NO: 123)
, ~	, ~	, ~	, ~	` ~ ,	, ~ ,
QKQIANQ	KQIANQF	QIANQFN	VKQLSSN	KQLSSNF	QLSSNFG
(SEQ ID NO: 61)	(SEQ ID NO: 62)	(SEQ ID NO: 63)	(SEQ ID NO: 124)	(SEQ ID NO: 125)	(SEQ ID NO: 126)
TANGUNIZ	A NICHTALIZ A	NICENIE N. T.	T C CNIEC N	CCMECAT	CNECATO
IANQFNK	ANQFNKA (SEQ ID NO: 65)	NQFNKAI (SEQ ID NO: 66)	LSSNFGA (SEQ ID NO: 127)	SSNFGAI (SEQ ID NO: 128)	SNFGAIS
(SEQ ID NO: 64)	(SEQ ID NO. 05)	(SEQ ID NO. 00)	(SEQ ID NO. 127)	(SEQ ID NO. 120)	(SEQ ID NO: 129)
QFNKAIS	FNKAISQ	NKAISQI	NFGAISS	FGAISSV	GAISSVL
(SEQ ID NO: 67)	(SEQ ID NO: 68)	(SEQ ID NO: 69)	(SEQ ID NO: 130)	(SEQ ID NO: 131)	(SEQ ID NO: 132)
KAISQIQ	AISQIQE	ISQIQES	AISSVLN	ISSVLND	SSVLNDI
(SEQ ID NO: 70)	(SEQ ID NO: 71)	(SEQ ID NO: 72)	(SEQ ID NO: 133)	(SEQ ID NO: 134)	(SEQ ID NO: 135)
SQIQESL	QIQESLT		SVLNDIL	VLNDILS	LNDILSR
	(SEQ ID NO: 74)			(SEQ ID NO: 137)	
,	,		, ~ ,	,	,
[0027] for the N	I Towasia al IID 1 as as	atida 2 tha mantida	NDILSRL	DILSRLD	ILSRLDK
<b>-</b> -	N-Terminal, HR1 per	• • •	(SEQ ID NO: 139)	(SEQ ID NO: 140)	(SEQ ID NO: 141)
may contain one of	of the following sequ	ences	T GDT DI	ant nuut	D.T. D.III.
			LSRLDK	SRLDKV	RLDKVE
			(SEQ ID NO: 142)	(SEQ ID NO: 143)	(PEÖ ID MO: 144)
ESLTTTS	SLTTTST	LTTTSTA	LDKVEA,		
(SEQ ID NO: 75)	(SEQ ID NO: 76)	(SEQ ID NO: 77)	(SEO ID NO: 145)		

(SEQ ID NO: 145)

[0039] and for the N-Terminal, HR1 peptide 5 the peptide may contain one of the following sequences:

RLITGRL LITGRLQ	ITGRLQS
(SEQ ID NO: 146) (SEQ ID NO: 147)	
, , , , , , , , , , , , , , , , , , , ,	, ~
TGRLQSL GRLQSLQ	RLQSLQT
(SEQ ID NO: 149) (SEQ ID NO: 150)	(SEQ ID NO: 151)
LQSLQTY QSLQTYV	SLQTYVT
(SEQ ID NO: 152) (SEQ ID NO: 153)	(SEQ ID NO: 154)
	шуушоот
LQTYVTQ QTYVTQQ	TYVTQQL
(SEQ ID NO: 155) (SEQ ID NO: 156)	(SEQ ID NO: 137)
YVTQQLI VTQQLIR	TQQLIRA
(SEQ ID NO: 158) (SEQ ID NO: 159)	
(	(
QQLIRAA QLIRAAE	LIRAAEI
(SEQ ID NO: 161) (SEQ ID NO: 162)	(SEQ ID NO: 163)
IRAAEIR RAAEIRA	AAEIRAS
(SEQ ID NO: 164) (SEQ ID NO: 165)	(SEQ ID NO: 166)
AEIRASA EIRASAN	IRASANL
(SEQ ID NO: 167) (SEQ ID NO: 168)	(SEQ ID NO: 169)
RASANLA ASANLAA	SANLAAT
(SEQ ID NO: 170) (SEQ ID NO: 171)	
(BEQ ID NO. 170) (BEQ ID NO. 171)	(BEQ ID NO. 172)
ANLAATK NLAATKM	LAATKMS
(SEQ ID NO: 173) (SEQ ID NO: 174)	(SEQ ID NO: 175)
· - / · · · · /	, - ,
AATKMSE ATKMSEC	TKMSECV
(SEQ ID NO: 176) (SEQ ID NO: 177)	(SEQ ID NO: 178)
KMSECVL MSECVLG	

(SEQ ID NO: 179) (SEQ ID NO: 180)

TSTALGK TTTSTAL TTSTALG (SEQ ID NO: 80) (SEQ ID NO: 79) STALGKL TALGKLQ ALGKLQD (SEQ ID NO: 81) (SEQ ID NO: 82) (SEQ ID NO: 83) LGKLQDV GKLQDVV KLQDVVN (SEQ ID NO: 84) (SEQ ID NO: 85) (SEQ ID NO: 86) LQDVVNQ QDVVNQN DVVNQNA (SEQ ID NO: 87) (SEQ ID NO: 88) (SEQ ID NO: 89) VVNQNAQ VNQNAQA NQNAQAL (SEQ ID NO: 90) (SEQ ID NO: 91) (SEQ ID NO: 92) QNAQALN NAQALNT AQALNTL (SEQ ID NO: 93) (SEQ ID NO: 94) (SEQ ID NO: 95) ALNTLVKQALNTLV LNTLVKQ(SEQ ID NO: 96) (SEQ ID NO: 97) (SEQ ID NO: 98) NTLVKQL TLVKQLS LVKQLSS (SEQ ID NO: 99) (SEQ ID NO: 100) (SEQ ID NO: 101) VKQLSSN KQLSSNF QLSSNFG (SEQ ID NO: 102) (SEQ ID NO: 103) (SEQ ID NO: 104) LSSNFGA SSNFGAI SNFGAIS (SEQ ID NO: 105) (SEQ ID NO: 106) (SEQ ID NO: 107) NFGAISS (SEQ ID NO: 108)

[0038] for the N-Terminal, HR1 peptide 4, the peptide may contain one of the following sequences:

LQDVVNQ QDVVNQN DVVNQNA (SEQ ID NO: 109) (SEQ ID NO: 110) (SEQ ID NO: 111)

[0040] Moreover, because only certain of the amino acids of the peptide make contact in the grooves formed by adjacent N-helices in the coiled coil, amino acids at nongroove binding positions can be replaced with essentially any other amino acid to make "mutated" peptide inhibitors. In addition, amino acids at positions that make groove contact also may be replaced with certain preferred amino acids. Thus, in the peptide

(SEQ ID NO: 1)

DVDLGDISGINASVVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

[0041] the amino acids at bold letter positions can be substituted with an amino acid selected from the group consisting of I, L, V, W, Y, F, N, Q, S, T, D, E, G, H, and M. Of these amino acids, I, L, V, W, and Y are most preferred, and F, N, Q, S, and T are next most preferred, although D, E, G, H, and M also may be used. The amino acids in non-bold positions can be any amino acid except proline, which is predicted to break the helical structure and therefore prevent groove binding. As with the above peptide, a peptide containing any seven contiguous residues of such a mutated peptide can be used.

[0042] These peptide sequences are contained within a longer peptide sequence containing at least 15 and up to 50 amino acids. The additional amino acids may be at the N and/or C termini of the sequences shown.

[0043] The peptide inhibitors may also contain at least 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from one of the sequences shown above, while remaining within the length limitations described above. The peptides may also be linked to carriers, such as carrier proteins, described in more detail below, in which case the entire molecule may contain more than 50 amino acids, but the portion of the molecule responsible for cell binding will still contain up to 50 amino acids.

The peptide of the invention may have the following general structure:

B-X-Z,

[0045] where B is an amino acid sequence containing up to about 43 amino acids, or B is an amino group, an acetyl group, a 9-fluorenylmethoxy-carbonyl group, a hydrophobic group, or a macromolecule carrier group. B also may comprise a carrier protein, such as HSA, in which case B may contain more than 8 amino acids, and may also comprises a linker peptide sequence that connects the antiviral sequence to the carrier protein.

[0046] Z is an amino acid sequence containing up to about 43 amino acids, or Z comprises a carboxyl group, an amido group, a hydrophobic group, or a macromolecular carrier group. Z also may comprise a carrier protein, such as HSA, in which case Z may contain more than 8 amino acids, and may also comprise a linker peptide sequence that connects the antiviral sequence to the carrier protein. The peptide and the carrier may also be linked as a chemical conjugate, via a linker such as a maleimide linker of the type that is commercially available from, for example, Pierce (Rockford, Ill.).

[0047] When considered together B and Z must contain at least 8 amino acids between the B and Z groups. Typically,

only one of B and Z is a macromolecule or carrier protein X is any 7, 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from the C or N peptides identified above.

The peptides of the invention also may comprise peptide sequences that exhibit 70% or more sequence identity with at least 7 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from one of the sequences shown above, while remaining within the length limitations described above.

The peptides of the invention also may comprise [0049] peptide sequences that exhibit identity in any two of the seven positions of the contiguous heptapeptide peptides described above, while remaining within the length limitations described above. This sequence identity advantageously may be located in the ith and i+4th positions in the contiguous heptapeptide peptides described above.

[0050] Ex vivo conjugation of the peptides of the invention moiety to a macromolecule such as HSA produces a highly soluble conjugate that can be purified and administered in tightly controlled dosage. The cloaked conjugate is biologically active as the conjugate, i.e. it does not act as a prodrug that releases the peptide moiety from the conjugate and cleavage of the conjugate is not required for biological activity. Moreover, once administered to a subject the conjugate has a surprisingly long in vivo half-life, has excellent tissue distribution and produces sustained activity corresponding to the activity of the biologically active moiety of the conjugate.

[0051] Advantageously, the peptide and the carrier protein and the macromolecule are linked in an approximately 1:1 ratio, to avoid "haptenization" of the biologically active moiety and generation of an immune response to the conjugate. Moreover, the peptide is advantageously appended to a single site in the macromolecule. For example, selective linkage to the unusually reactive cysteine 34 (C34) of HSA may be used. Methods for selective linkage to C34 using, for example, a maleimide containing linker, are known in the art.

[0052] In the event that more than one molecule of peptide is linked to the macromolecule, this is advantageously achieved via a "multivalent" linker that is attached to a single point of the macromolecule. For example, a linker can be appended to C34 of HSA that permits attachment of a plurality of peptides to the linker. Multivalent linkers are known in the art and can contain, for example, a thiophilic group for reaction with C34 of HSA, and multiple nucleophilic (such as NH or OH) or electrophilic (such as activated ester) groups that permit attachment of a plurality of peptides to the linker.

Preparation of Peptides of the Invention

[0053] The peptides of the invention may be synthesized or prepared by techniques well known in the art. Peptide synthesizers are commercially available from, for example, Applied Biosystems or Milligen/Biosearch. See also, for example, Creighton, 1983, Proteins: Structures and Molecular Principles, W. H. Freeman and Co., N.Y., which is incorporated herein by reference in its entirety. Short peptides, for example, can be synthesized on a solid support or in solution. Longer peptides, or fusions of longer peptides with carrier proteins such as human serum albumin, may be made using recombinant DNA techniques. Nucleotide sequences encoding the desired peptides or fusion proteins

containing the peptides may be synthesized, and/or cloned, and expressed according to techniques well known to those of ordinary skill in the art. See, for example, Sambrook, et al., 1989, Molecular Cloning, A Laboratory Manual, Vols. 1-3, Cold Spring Harbor Press, N.Y.

[0054] The peptides also may be synthesized such that one or more of the bonds linking the amino acid residues of the peptides are non-peptide bonds. Alternative non-peptide bonds may be formed by reactions well known to those in the art, and may include, but are not limited to imino, ester, hydrazide, semicarbazide, and azo bonds. In yet another embodiment of the invention, peptides comprising the sequences described above may be synthesized with additional chemical groups present at their amino and/or carboxy termini, such that, for example, the stability, bioavailability, and/or inhibitory activity of the peptides is enhanced. For example, hydrophobic groups such as carbobenzoxyl, dansyl, or t-butyloxycarbonyl groups, may be added to a peptide's amino terminus. Likewise, an acetyl group or a 9-fluorenylmethoxy-carbonyl group may be placed at a peptide's amino terminus. Additionally, a hydrophobic group, t-butyloxycarbonyl, or an amido group may be added to a peptide's carboxy terminus. Further, non-naturally occurring amino acids can be used to improve a peptide's stability, bioavailability, or binding/inhibitory characteristics. For example, methionine can be replaced with norleucine. Other non-naturally occurring amino acid residues are well known.

[0055] The peptides of the invention also may contain amino acid substitutions, which may be of a conserved or non-conserved nature. Conserved amino acid substitutions consist of replacing one or more amino acids in a peptide sequence with amino acids of similar charge, size, and/or hydrophobicity characteristics, such as, for example, a glutamic acid (E) to aspartic acid (D) amino acid substitution. When only conserved substitutions are made, the resulting peptide retains the functionality of the unsubstituted peptide. Non-conserved substitutions consist of replacing one or more amino acids of a peptide sequence with amino acids possessing dissimilar charge, size, and/or hydrophobicity characteristics, such as, for example, a glutamic acid (E) to valine (V) substitution. The peptides of the present invention may advantageously contain amino acid substitutions of a conserved nature.

[0056] The stability of the peptides of the invention may be increased by either in vivo or ex vivo linkage to a carrier protein, such as a blood component. Suitable blood components for use in the present invention are known in the art. Human serum albumin ("HSA") is a predominant component of human blood and is particularly suited for use in the present invention. In particular, HSA has an exposed surface cysteine residue that provides a reactive thiol moiety for covalent linkage of the peptides compounds to the protein. Activated linkers that are particularly suited for linkage to thiols include unsaturated cyclic imides such as maleimides,  $\alpha$ -halo esters, such as  $\alpha$ -iodo- and  $\alpha$ -bromo acetates, and vinyl pyridine derivative. Such linkers can be added to the peptides during synthesis and can be added at any point in the sequence although the N and/or C terminus advantageously is used. Suitable activated linkers are commercially available from, for example, Pierce Chemical (Rockford, Ill.). Methods for preparing suitable activated compounds

for linking to HSA are known in art. See for example, U.S. Pat. No. 5,612,034, which is incorporated herein in its entirety.

[0057] Moreover, the gene for HSA has been cloned, which permits the ready preparation of fusion proteins of the peptides and HSA. Methods of making fusion proteins are known in the art. See, for example, WO01/79271 and WO01/79258, the contents of which are hereby incorporated by reference in their entirety. The preparation of fusion proteins is useful for preparing persistent derivatives of the present anti-viral peptides.

[0058] Another blood component that is suitable for linkage to the anti-viral compounds is an immunoglobulin ("Ig") molecule. Igs are persistent and are present in relatively high concentration in the blood. For in vitro coupling, Igs have the advantage of being readily stable and readily isolated, and methods of making Ig conjugates are well known in the art. Moreover, Ig genes may readily be cloned and recombinant Ig and Ig fusion proteins prepared. Methods for obtaining fully human Igs are well known in the art. See for example, U.S. Pat. Nos. 5,969,108 and 6,300,064, the contents of which are hereby incorporated by reference in their entirety. In addition, phage display methods for selecting Igs having a particularly desired binding activity, for example, for binding to HSA, are well known in the art. See U.S. Pat. Nos. 5,885,793, 5,969,108 and 6,300,064. In the context of the present invention, an Ig refers to any suitable immunoglobulin or immunogolobulin derivative known in the art, and includes, for example, whole IgG, IgM, Fab fragments, F(ab')2 fragments, and single chain Fv fragments.

[0059] Other blood components suitable for use in the present invention include transferrin, ferritin, steroid binding proteins, thyroxin binding protein, and  $\alpha$ -2-macroglobulin.

[0060] In the peptides, the activated linkers also may be coupled to reactive side chain residues, such as lysine side chains. For example, a linker containing an active ester moiety and a maleimide moiety can be selectively reacted at the active ester (such as an N-hydroxysuccinimidyl ester) via lysine side chains or at the N-terminus of the peptide.

[0061] Both natural and recombinant HSA and human Igs are commercially available and are suitable for use in the present invention.

[0062] The peptides also may have a non-peptide macro-molecular carrier group covalently attached to their amino and/or carboxy termini. Such macromolecular carrier groups may include, for example, lipid-fatty acid conjugates, polyethylene glycol, or carbohydrates.

Use of the Peptides

[0063] The peptides of the invention exhibit potent antiviral activity against coronaviruses, such as for example, the SARS virus. As such, the peptides may be used as inhibitors of human and non-human coronoviruses, especially SARS, transmission to uninfected cells. Various peptides from the C-terminal HR2 domains of S2 proteins of SARS virus and murine hepatitis virus (MHV) have been shown to exhibit antiviral activity against these viruses in cell culture assays(see for example Liu et al., *Lancet* 363: 938 (2004); and Bosch et al., *J. Virol.* 77:8801 (2003). The human SARS viruses whose transmission may be inhibited by the peptides

of the invention include, but are not limited to all strains of the SARS virus. The non-human retroviruses whose transmission may be inhibited by the peptides of the invention include, but are not limited to coronaviruses that infect domestic animals and livestock, for example, coronaviruses from cows, pigs, mice and chickens. However, as will be appreciated by one skilled in the art, the peptides used for preventing coronaviruses will be most effective when derived using the specific sequence of the infecting virus strain.

[0064] With respect to SARS in humans, the peptides of the invention may be used as a therapeutic in the treatment of SARS infections. The peptides of the invention may be administered using techniques well known to those in the art. Preferably, agents are formulated and administered systemically. Techniques for formulation and administration may be found in "Remington's Pharmaceutical Sciences" 18th ed., 1990 Mack Publishing Co., Easton, Pa. Suitable routes may include oral, rectal, transmucosal, or intestinal administration; parenteral delivery, including intramuscular, subcutaneous, intramedullary injections, as well as intrathecal, direct intraventricular, intravenous, intraperitoneal, intranasal, or intraocular injections, just to name a few. Most preferably, administration is intravenous. For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks' solution, Ringer's solution, or physiological saline buffer. For such transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art. Other diluents, adjuvants, and excipients are known in the art.

[0065] In addition, the peptides may be used as a prophylactic measure in previously uninfected individuals after acute exposure to a SARS virus. Examples of such prophylactic use of the peptides may include, but are not limited to, settings where the likelihood of SARS transmission exists, such as, for example, in hospitals and transport termini such as airports and train stations. The peptides of the invention in such cases may serve the role of a prophylactic vaccine, wherein the host raises antibodies against the peptides of the invention, which then serve to neutralize SARS viruses by, for example, inhibiting further SARS infection. Administration of the peptides of the invention as a prophylactic vaccine, therefore, would comprise administering to a host a concentration of peptides effective in raising an immune response which is sufficient to neutralize SARS or a related coronavirus, by, for example, inhibiting SARS ability to infect cells. The exact concentration will depend upon the specific peptide to be administered, but may be determined by using standard techniques for assaying the development of an immune response which are well known to those of ordinary skill in the art. The peptides to be used as vaccines are usually administered intramuscularly.

[0066] The peptides may be formulated with a suitable adjuvant in order to enhance the immunological response. Such adjuvants may include, but are not limited to mineral gels such as aluminum hydroxide; surface active substances such as lysolecithin, pluronic polyols, polyanions; other peptides; oil emulsions; and potentially useful human adjuvants such as BCG and *Corynebacterium parvum*. Many methods may be used to introduce the vaccine formulations described here. These methods include but are not limited to oral, intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, and intranasal routes.

[0067] Alternatively, an effective concentration of polyclonal or monoclonal antibodies raised against the peptides of the invention may be administered to a host so that no uninfected cells become infected by the SARS virus or other coronavirus. The exact concentration of such antibodies will vary according to each specific antibody preparation, but may be determined using standard techniques well known to those of ordinary skill in the art. Administration of the antibodies may be accomplished using a variety of techniques, including, but not limited to those described in this section.

[0068] Effective dosages of the peptides of the invention to be administered may be determined through procedures well known to those in the art which address such parameters as biological half-life, bioavailability, and toxicity.

[0069] The antiviral activity of the peptides of the invention may show a pronounced type and subtype specificity, i.e., specific peptides may be effective in inhibiting the activity of only specific coronaviruses. This feature of the invention presents many advantages. One such advantage, for example, lies in the field of diagnostics, wherein one can use the antiviral specificity of the peptide of the invention to ascertain the identity of a viral isolate. With respect to coronaviruses, one may easily determine whether a viral isolate consists of a coronavirus that causes SARS or a virus that causes milder cold-like symptoms. For example, uninfected cells may be co-infected with a coronavirus isolate which has been identified as containing a SARS virus. A peptide of the invention may be added which is known to be active against the SARS virus, after which the retroviral activity of cell supernatants may be assayed, using known methods. Those isolates whose viral activity is completely or nearly completely inhibited contain the SARS virus. Those isolates whose viral activity is unchanged or only reduced by a small amount, may be considered to not contain the SARS virus. Such an isolate may then be treated with one or more of the other peptides of the invention, and subsequently be tested for its viral activity in order to determine the identify of the viral isolate.

<sup>&</sup>lt;210> SEQ ID NO 1

<sup>&</sup>lt;211> LENGTH: 49

<sup>&</sup>lt;212> TYPE: PRT

```
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Glu Ser Leu Ile Asp Leu Gln Glu Leu Gly Lys Tyr Glu Gln Tyr Ile
Lys
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Gln Ile Pro Phe Ala Met Gln Met Ala Tyr Arg Phe Asn Gly Ile Gly
Val Thr Gln Asn Val Leu Tyr Glu Asn Gln Lys Gln Ile Ala Asn Gln
            20
                                25
                                                    30
Phe Asn Lys Ala Ile Ser Gln Ile Gln Glu Ser Leu Thr
                                                45
        35
                            40
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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      peptide
<400> SEQUENCE: 3
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                                    10
Val Asn Gln Asn Ala Gln Ala Leu Asn Thr Leu Val Lys Gln Leu Ser
            20
                                                    30
Ser Asn Phe Gly Ala Ile Ser Ser
        35
<210> SEQ ID NO 4
<211> LENGTH: 45
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      peptide
<400> SEQUENCE: 4
Gly Lys Leu Gln Asp Val Val Asn Gln Asn Ala Gln Ala Leu Asn Thr
                                    10
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Asn Asp Ile Leu Ser Arg Leu Asp Lys Val Glu Ala Glu
```

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<210> SEQ ID NO 5
<211> LENGTH: 48
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      peptide
<400> SEQUENCE: 5
Arg Leu Ile Thr Gly Arg Leu Gln Ser Leu Gln Thr Tyr Val Thr Gln
Gln Leu Ile Arg Ala Ala Glu Ile Arg Ala Ser Ala Asn Leu Ala Ala
Thr Lys Met Ser Glu Cys Val Leu Gly Gln Ser Lys Arg Val Asp Phe
        35
                            40
<210> SEQ ID NO 6
<211> LENGTH: 7
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      peptide
<400> SEQUENCE: 6
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<210> SEQ ID NO 7
<211> LENGTH: 7
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      peptide
<400> SEQUENCE: 7
Val Val Asn Ile Gln Lys Glu
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Arg Leu Asn Glu Val Ala Lys
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Ala Lys Asn Leu Asn Glu Ser
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Ser Leu Ile Asp Leu Gln Glu
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Leu Ile Asp Leu Gln Glu Leu
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Ile Asp Leu Gln Glu Leu Gly
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Gln Ile Pro Phe Ala Met Gln
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Ile Pro Phe Ala Met Gln Met
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Pro Phe Ala Met Gln Met Ala
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Phe Ala Met Gln Met Ala Tyr
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Ala Met Gln Met Ala Tyr Arg
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Met Ala Tyr Arg Phe Asn Gly
<210> SEQ ID NO 45
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Ala Tyr Arg Phe Asn Gly Ile
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Tyr Arg Phe Asn Gly Ile Gly
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Arg Phe Asn Gly Ile Gly Val
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Phe Asn Gly Ile Gly Val Thr
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Asn Gly Ile Gly Val Thr Gln
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Ile Gly Val Thr Gln Asn Val
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Gly Val Thr Gln Asn Val Leu
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Val Thr Gln Asn Val Leu Tyr
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Thr Gln Asn Val Leu Tyr Glu
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Gln Asn Val Leu Tyr Glu Asn
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Val Leu Tyr Glu Asn Gln Lys
<210> SEQ ID NO 57
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Leu Tyr Glu Asn Gln Lys Gln
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<220> FEATURE:
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Glu Asn Gln Lys Gln Ile Ala
<210> SEQ ID NO 60
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Asn Gln Lys Gln Ile Ala Asn
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Lys Gln Ile Ala Asn Gln Phe
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Gln Ile Ala Asn Gln Phe Asn
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Ala Asn Gln Phe Asn Lys Ala
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Asn Gln Phe Asn Lys Ala Ile
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Gln Phe Asn Lys Ala Ile Ser
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Phe Asn Lys Ala Ile Ser Gln
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Asn Lys Ala Ile Ser Gln Ile
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Lys Ala Ile Ser Gln Ile Gln
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Ala Ile Ser Gln Ile Gln Glu
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Ser Gln Ile Gln Glu Ser Leu
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Gln Ile Gln Glu Ser Leu Thr
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Glu Ser Leu Thr Thr Thr Ser
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Ser Leu Thr Thr Thr Ser Thr
<210> SEQ ID NO 77
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Leu Thr Thr Ser Thr Ala
<210> SEQ ID NO 78
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Thr Thr Ser Thr Ala Leu
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Thr Thr Ser Thr Ala Leu Gly
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Thr Ser Thr Ala Leu Gly Lys
<210> SEQ ID NO 81
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Thr Ala Leu Gly Lys Leu Gln
<210> SEQ ID NO 83
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Ala Leu Gly Lys Leu Gln Asp
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Leu Gly Lys Leu Gln Asp Val
<210> SEQ ID NO 85
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Gly Lys Leu Gln Asp Val Val
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Lys Leu Gln Asp Val Val Asn
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Leu Gln Asp Val Val Asn Gln
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Gln Asp Val Val Asn Gln Asn
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Asp Val Val Asn Gln Asn Ala
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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
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Val Asn Gln Asn Ala Gln Ala
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Asn Gln Asn Ala Gln Ala Leu
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Gln Asn Ala Gln Ala Leu Asn
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<400> SEQUENCE: 94
Asn Ala Gln Ala Leu Asn Thr
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Ala Gln Ala Leu Asn Thr Leu
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Ala Leu Asn Thr Leu Val Lys
<210> SEQ ID NO 98
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Asn Thr Leu Val Lys Gln Leu
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Thr Leu Val Lys Gln Leu Ser
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Leu Val Lys Gln Leu Ser Ser
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<220> FEATURE:
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Lys Gln Leu Ser Ser Asn Phe
<210> SEQ ID NO 104
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Gln Leu Ser Ser Asn Phe Gly
<210> SEQ ID NO 105
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Leu Ser Ser Asn Phe Gly Ala
<210> SEQ ID NO 106
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Ser Ser Asn Phe Gly Ala Ile
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<220> FEATURE:
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Asn Phe Gly Ala Ile Ser Ser
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Val Val Asn Gln Asn Ala Gln
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Asn Gln Asn Ala Gln Ala Leu
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Gln Asn Ala Gln Ala Leu Asn
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Asn Ala Gln Ala Leu Asn Thr
<210> SEQ ID NO 117
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Ala Gln Ala Leu Asn Thr Leu
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Gln Ala Leu Asn Thr Leu Val
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Ala Leu Asn Thr Leu Val Lys
<210> SEQ ID NO 120
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Asn Thr Leu Val Lys Gln Leu
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Thr Leu Val Lys Gln Leu Ser
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<220> FEATURE:
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Leu Val Lys Gln Leu Ser Ser
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Val Lys Gln Leu Ser Ser Asn
<210> SEQ ID NO 125
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Lys Gln Leu Ser Ser Asn Phe
<210> SEQ ID NO 126
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<220> FEATURE:
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Gln Leu Ser Ser Asn Phe Gly
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<220> FEATURE:
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Leu Ser Ser Asn Phe Gly Ala
<210> SEQ ID NO 128
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<220> FEATURE:
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Ser Ser Asn Phe Gly Ala Ile
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<212> TYPE: PRT
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Ser Asn Phe Gly Ala Ile Ser
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Asn Phe Gly Ala Ile Ser Ser
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Phe Gly Ala Ile Ser Ser Val
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Gly Ala Ile Ser Ser Val Leu
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Ala Ile Ser Ser Val Leu Asn
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Ile Ser Ser Val Leu Asn Asp
<210> SEQ ID NO 135
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<212> TYPE: PRT
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<400> SEQUENCE: 135
Ser Ser Val Leu Asn Asp Ile
<210> SEQ ID NO 136
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Ser Val Leu Asn Asp Ile Leu
<210> SEQ ID NO 137
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<400> SEQUENCE: 137
Val Leu Asn Asp Ile Leu Ser
<210> SEQ ID NO 138
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Leu Asn Asp Ile Leu Ser Arg
<210> SEQ ID NO 139
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Asn Asp Ile Leu Ser Arg Leu
<210> SEQ ID NO 140
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Asp Ile Leu Ser Arg Leu Asp
<210> SEQ ID NO 141
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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<400> SEQUENCE: 141
Ile Leu Ser Arg Leu Asp Lys
<210> SEQ ID NO 142
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<400> SEQUENCE: 142
Leu Ser Arg Leu Asp Lys
<210> SEQ ID NO 143
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
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Ser Arg Leu Asp Lys Val
<210> SEQ ID NO 144
<211> LENGTH: 6
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<400> SEQUENCE: 144
Arg Leu Asp Lys Val Glu
<210> SEQ ID NO 145
<211> LENGTH: 6
<212> TYPE: PRT
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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      peptide
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<400> SEQUENCE: 145
Leu Asp Lys Val Glu Ala
<210> SEQ ID NO 146
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<400> SEQUENCE: 146
Arg Leu Ile Thr Gly Arg Leu
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<211> LENGTH: 7
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Leu Ile Thr Gly Arg Leu Gln
<210> SEQ ID NO 148
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Ile Thr Gly Arg Leu Gln Ser
<210> SEQ ID NO 149
<211> LENGTH: 7
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<220> FEATURE:
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Thr Gly Arg Leu Gln Ser Leu
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Gly Arg Leu Gln Ser Leu Gln
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<212> TYPE: PRT
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Arg Leu Gln Ser Leu Gln Thr
<210> SEQ ID NO 152
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<213> ORGANISM: Artificial Sequence
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Leu Gln Ser Leu Gln Thr Tyr
<210> SEQ ID NO 153
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Gln Ser Leu Gln Thr Tyr Val
<210> SEQ ID NO 154
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Ser Leu Gln Thr Tyr Val Thr
<210> SEQ ID NO 155
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Leu Gln Thr Tyr Val Thr Gln
<210> SEQ ID NO 156
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Gln Thr Tyr Val Thr Gln Gln
<210> SEQ ID NO 157
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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Thr Tyr Val Thr Gln Gln Leu
<210> SEQ ID NO 158
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Tyr Val Thr Gln Gln Leu Ile
<210> SEQ ID NO 159
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Val Thr Gln Gln Leu Ile Arg
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Thr Gln Gln Leu Ile Arg Ala
<210> SEQ ID NO 161
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<220> FEATURE:
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Gln Gln Leu Ile Arg Ala Ala
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Gln Leu Ile Arg Ala Ala Glu
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<220> FEATURE:
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Leu Ile Arg Ala Ala Glu Ile
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Ile Arg Ala Ala Glu Ile Arg
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<211> LENGTH: 7
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<400> SEQUENCE: 165
Arg Ala Ala Glu Ile Arg Ala
<210> SEQ ID NO 166
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<212> TYPE: PRT
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<400> SEQUENCE: 166
Ala Ala Glu Ile Arg Ala Ser
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<211> LENGTH: 7
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Ala Glu Ile Arg Ala Ser Ala
<210> SEQ ID NO 168
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Glu Ile Arg Ala Ser Ala Asn
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Ile Arg Ala Ser Ala Asn Leu
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Arg Ala Ser Ala Asn Leu Ala
<210> SEQ ID NO 171
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<213> ORGANISM: Artificial Sequence
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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Ala Ser Ala Asn Leu Ala Ala
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<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<210> SEQ ID NO 173
<211> LENGTH: 7
<212> TYPE: PRT
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<220> FEATURE:
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<400> SEQUENCE: 173
Ala Asn Leu Ala Ala Thr Lys
<210> SEQ ID NO 174
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Asn Leu Ala Ala Thr Lys Met
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Leu Ala Ala Thr Lys Met Ser
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Ala Ala Thr Lys Met Ser Glu
<210> SEQ ID NO 177
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
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<400> SEQUENCE: 177
Ala Thr Lys Met Ser Glu Cys
<210> SEQ ID NO 178
<211> LENGTH: 7
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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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<210> SEQ ID NO 179
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<220> FEATURE:
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Lys Met Ser Glu Cys Val Leu
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Met Ser Glu Cys Val Leu Gly
<210> SEQ ID NO 181
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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Val Val Asn Ile Gln Lys Glu Ile Asp Arg Leu Asn Glu Val Ala Lys
Asn Leu Asn Glu Ser Leu Ile Asp Leu Gln Glu Leu Gly Lys Tyr Glu
            20
                                                     30
Gln Tyr Ile Lys
        35
<210> SEQ ID NO 182
<211> LENGTH: 50
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
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Glu Gln Tyr Ile Lys
```

1. An antiviral peptide having between 7 and 50 amino acids, wherein said peptide exhibits antiviral activity against a coronavirus, and where said peptide contains a sequence comprising at least 7 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 1) DVDLGDISGINASVVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2) QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4)

GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

2. The peptide of claim 1, wherein said peptide contains a sequence selected from the group consisting of

SVVNIQK VVNIQKE VNIQKEI (SEQ ID NO: 7) (SEQ ID NO: 8) (SEQ ID NO: 6) NIQKEID IQKEIDR QKEIDRL (SEQ ID NO: 9) (SEQ ID NO: 10) (SEQ ID NO: 11) KEIDRLN EIDRLNE IDRLNEV (SEQ ID NO: 12) (SEQ ID NO: 13) (SEQ ID NO: 14) DRLNEVA RLNEVAK LNEVAKN (SEQ ID NO: 15) (SEQ ID NO: 16) (SEQ ID NO: 17) NEVAKNL EVAKNLN VAKNLNE (SEQ ID NO: 18) (SEQ ID NO: 19) (SEQ ID NO: 20) KNLNESL AKNLNES NLNESLI (SEQ ID NO: 21) (SEQ ID NO: 22) (SEQ ID NO: 23) LNESLID NESLIDL ESLIDLQ (SEQ ID NO: 24) (SEQ ID NO: 25) (SEQ ID NO: 26) SLIDLQE LIDLQEL IDLQELG (SEQ ID NO: 27) (SEQ ID NO: 28) (SEQ ID NO: 29) QELGKYE DLQELGK LQELGKY (SEQ ID NO: 30) (SEQ ID NO: 31) (SEQ ID NO: 32) ELGKYEQ LGKYEQY GKYEQYI (SEQ ID NO: 33) (SEQ ID NO: 34) (SEQ ID NO: 35) KYEQYIK IPFAMQM QIPFAMQ (SEQ ID NO: 36) (SEQ ID NO: 37) (SEQ ID NO: 38) PFMAQMA FAMQMAY AMQMAYR (SEQ ID NO: 39) (SEQ ID NO: 40) (SEQ ID NO: 41) MQMAYRF QMAYRFN MAYRFNG (SEQ ID NO: 42) (SEQ ID NO: 43) (SEQ ID NO: 44) AYRFNGI YRFNGIG RFNGIGV (SEQ ID NO: 45) (SEQ ID NO: 46) (SEQ ID NO: 47) FNGIGVT NGIGVTQ IGVTQNV (SEQ ID NO: 48) (SEQ ID NO: 49) (SEQ ID NO: 50) GVTQNVL VTQNVLY TQNVLYE (SEQ ID NO: 51) (SEQ ID NO: 52) (SEQ ID NO: 53) QNVLYEN NVLYENQVLYENQK

(SEQ ID NO: 54) (SEQ ID NO: 55) (SEQ ID NO: 56)

LYELNQKQ (SEQ ID NO:					59)
NQKQIAN (SEQ ID NO:		QKQIANQ (SEQ ID N	NO: 61)	KQIANQF (SEQ ID NO:	62)
		IANQFNK (SEQ ID N	NO: 64)	ANQFNKA (SEQ ID NO:	65)
NQFNKAI (SEQ ID NO:			NO: 67)	FNKAISQ (SEQ ID NO:	68)
NKAISQI (SED ID NO:	69)	KAISQIQ (SEQ ID N	NO: 70)	AISQIQE (SEQ ID NO:	71)
ISQIQES (SEQ ID NO:		SQIQESL (SEQ ID N	NO: 73)	QIQESLT (SEQ ID NO:	74)
ESLTTTS (SEQ ID NO:	75)	SLTTTST (SEQ ID N	NO: 76)	LTTTSTA (SEQ ID NO:	77)
TTTSTAL (SEQ ID NO:		TTSTALG (SEQ ID N	NO: 79)	TSALGAK (SEQ ID NO:	80)
STALGKL (SEQ ID NO:	81)	TALGKLQ (SEQ ID N	NO: 82)	ALGKLQD (SEQ ID NO:	83)
LGKLQDV (SEQ ID NO:			NO: 85)	KLQDVVN (SEQ ID NO:	86)
LQDVVNQ (SEQ ID NO:		QDVVNQN (SEQ ID N	NO: 88)	DVVNQNA (SEQ ID NO:	89)
		VNQNAQA (SEQ ID N	NO: 91)	NQNAQAL (SEQ ID NO:	92)
		NAQALNT (SEQ ID N	NO: 94)	AQALNTL (SEQ ID NO:	95)
QALNTLV (SEQ ID NO:	96)	ALNTLVK (SEQ ID N	NO: 97)	LNTLVKQ (SEQ ID NO:	98)
NTLVKQL (SEQ ID NO:		TLVKQLS	NO: 100)	LVKQLSS (SEQ ID NO:	101)
VKQLSSN		KQLSSNF		QLSSNFG (SEQ ID NO:	
LSSNFGA		SSNFGAI		SNFGAIS	
(SEQ ID NO:		(SEQ ID N		(SEQ ID NO: QDVVNQN	107)
		(SEQ ID N	NO: 109)	(SEQ ID NO: VNQNAQA	110)
(SEQ ID NO:	111)	(SEQ ID N	NO: 112)	(SEQ ID NO:	113)
NQNAQAL (SEQ ID NO:		QNAQALN (SEQ ID N	NO: 115)	NAQALNT (SEQ ID NO:	116)
AQALNTL (SEQ ID NO:			NO: 118)	ALNTLVK (SEQ ID NO:	119)
LNTLVKQ (SEQ ID NO:			NO: 121)	TLVKQLS (SEQ ID NO:	122)
LVKQLSS (SEQ ID NO:		VKQLSSN (SEQ 1D N	NO: 124)	KQLSSNF (SEQ ID NO:	125)
QLSSNFG (SEQ ID NO:		LSSNFGA (SEQ ID N	NO: 127)	SSNFGAI (SEQ ID NO:	128)
SNFGAIS (SEQ ID NO:	129)	NEGAISS (SEQ ID N	NO: 130)	FGAISSV (SEQ ID NO:	131)

	AISSVLN 132) (SEQ ID NO: 133)	ISSVLND (SEQ ID NO: 134)
SSVLNDI (SEQ ID NO:	SVLNDIL 135) (SEQ ID NO: 136)	VLNDILS (SEQ ID NO: 137)
	NDILSRL 138) (SEQ ID NO: 139)	DILSRLD (SEQ ID NO: 140)
ILSRLDK (SEQ ID NO:	LSRLDK 141) (SEQ ID NO: 142)	SRLDKV (SEQ ID NO: 143)
RLDKVE (SEQ ID NO:	LDKVEA, 144) (SEQ ID NO: 145)	RLITGRL (SEQ ID NO: 146)
	ITGRLQS 147) (SEQ ID NO: 148)	TGRLQSL (SEQ ID NO: 149)
	RLQSLQT 150) (SEQ ID NO: 151)	LQSLQTY (SEQ ID NO: 152)
	SLQTYVT 153) (SEQ ID NO: 154)	
	TYVTQQL 156) (SEQ ID NO: 157)	YVTQQLI (SEQ ID NO: 158)
	TQQLIRA 159) (SEQ ID NO: 160)	
~	LIRAAEI 162) (SEQ ID NO: 163)	IRAAEIR (SEQ ID NO: 164)
	AAEIRAS 165) (SEQ ID NO: 166)	AEIRASA (SEQ ID NO: 167)
	IRASANL 168) (SEQ ID NO: 169)	RASANLA (SEQ ID NO: 170)
ASANLAA (SEQ ID NO:	SANLAAT 171) (SEQ ID NO: 172)	ANLAATK (SEQ ID NO: 173)
NLAATKM (SEQ ID NO:	LAATKMS 174) (SEQ ID NO: 175)	AATKMSE (SEQ ID NO: 176)
ATKMSEC (SEQ ID NO:	TKMSECV 177) (SEQ ID NO: 178)	KMSECVL (SEQ ID NO: 179)
MSECVLG.	180)	

3. A peptide according to claim 1, wherein said peptide comprises at least 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from one of the sequences:

(SEQ ID NO: 181)
VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2) QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3)

ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

4. The peptide according to claim 1, wherein said peptide is linked to a carrier protein.

- 5. The peptide according claim 4, wherein said peptide is linked to human serum albumin.
- 6. An antiviral peptide having between 7 and 50 amino acids, wherein said peptide exhibits antiviral activity against a coronavirus, and wherein said peptide has the structure:

B-X-Z,

- where B is an amino acid sequence containing up to about 43 amino acids, or B is an amino group, an acetyl group, a 9-fluorenylmethoxy-carbonyl group, a hydrophobic group, or a macromolecule carrier group, or B is a carrier protein, in which case B may contain more than 8 sequence to said carrier protein;
- Z is an amino acid sequence containing up to about 43 amino acids, or Z comprises a carboxyl group, an amindo group, a hydrophobic group, or a macromolecular carrier group, or Z is a carrier protein, such as HSA, in which case Z may contain more than 8 amino acids, and may also comprises a linker peptide sequence that connects the antiviral sequence to said carrier protein;
- wherein when considered together B and Z must contain at least 8 amino acids between the B and Z group;
- and wherein X is a peptide sequence comprising at least 7 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 181)
VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2)
QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

7. The peptide according to claim 6, comprising at least 10, 15, 20, 25, 30, 35, or 40 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 181)

VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2) QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4) GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5) RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

8. An antiviral peptide having between 7 and 50 amino acids, wherein said peptide exhibits antiviral activity against a coronavirus, and wherein said peptide comprises a sequence that exhibits identity in any two of the seven

positions of a contiguous heptapeptide, wherein said contiguous heptapeptide comprises 7 contiguous amino acids from one of the following sequences:

(SEQ ID NO: 181) VVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIK;

(SEQ ID NO: 2)

QIPFAMQMAYRFNGIGVTQNVLYENQKQIANQFNKAISQIQESLT;

(SEQ ID NO: 3) ESLTTTSTALGKLQDVVNQNAQALNTLVKQLSSNFGAISS;

(SEQ ID NO: 4)

GKLQDVVNQNAQALNTLVKQLSSNFGAISSVLNDILSRLDKVEAE; and

(SEQ ID NO: 5)

RLITGRLQSLQTYVTQQLIRAAEIRASANLAATKMSECVLGQSKRVDF.

- 9. The peptide according to claim 8, wherein said sequence identity is located in the ith and i+4th positions in said contiguous heptapeptides.
- 10. A pharmaceutical composition comprising a peptide according to claim 1 and a pharmaceutically acceptable diluent, adjuvant and/or excipient.
- 11. A method of treating or preventing a coronavirus infection in a subject, comprising administering to a patient suspected of suffering from said infection an effective amount of a peptide or composition according to claim 10.
- 12. The method according to claim 11, wherrein said subject is a human.
- 13. The method according to claim 11, wherein said subject is a cow, pig, or chicken.

**14**. An antiviral peptide having between 7 and 50 amino acids, wherein said peptide exhibits antiviral activity against a coronavirus, and wherein said peptide contains a sequence comprising at least 7 contiguous amino acids from the sequence:

(SEQ ID NO: 1) DV**DL**GDISGINAS**V**VNIQKEIDRLNE**VA**KNLNESLIDLQE**LG**K**Y**EQYIK;

wherein the amino acids at bold letter positions can be substituted with an amino acid selected from the group consisting of I, L, V, W, Y, F, N, Q, S, T, D, E, G, H, and M, and wherein amino acids in non-bold positions can be any amino acid except proline.

- 15. A compound comprising a peptide according to claim 14 covalently linked to a carrier protein.
- 16. The peptide according to claim 14 wherein at least 8 of said amino acids at said bold letter positions are selected from the group consisting of I, L, V, W, and Y.
- 17. A compound according to claim 15 wherein said compound is a fusion protein.
- 18. A compound according to claim 15 wherein said peptide is linked to said carrier protein by a non-peptide linker.
- 19. A pharmaceutical composition comprising a peptide according to claim 14 and a pharmaceutically acceptable excipient, diluent or adjuvant.
- 20. A method of preventing or treating a coronavirus infection in a subject, comprising administering to said subject a pharmaceutical composition according to claim 19.