



US007897744B2

(12) **United States Patent**
Plummer et al.

(10) **Patent No.:** **US 7,897,744 B2**
(45) **Date of Patent:** **Mar. 1, 2011**

(54) **SARS VIRUS NUCLEOTIDE AND AMINO ACID SEQUENCES AND USES THEREOF**

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(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 906 days.

(21) Appl. No.: **10/555,073**

(22) PCT Filed: **Apr. 28, 2004**

(86) PCT No.: **PCT/CA2004/000626**

§ 371 (c)(1),
(2), (4) Date: **Dec. 19, 2006**

(87) PCT Pub. No.: **WO2004/096842**

PCT Pub. Date: **Nov. 11, 2004**

(65) **Prior Publication Data**

US 2007/0258999 A1 Nov. 8, 2007

Related U.S. Application Data

(60) Provisional application No. 60/465,783, filed on Apr. 28, 2003, provisional application No. 60/466,733, filed on May 1, 2003.

(51) **Int. Cl.**
C12N 15/50 (2006.01)
C12N 7/00 (2006.01)

(52) **U.S. Cl.** **536/23.72**; 435/235.1; 435/320.1; 514/44

(58) **Field of Classification Search** None
See application file for complete search history.

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(57) **ABSTRACT**

The invention provides, in part, the genomic sequence of a putative coronavirus, the SARS virus, and provides novel nucleic acid and amino acid sequences that may be used, for example, for the diagnosis, prophylaxis, or therapy of a variety of SARS virus related disorders.

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Replicase 1A

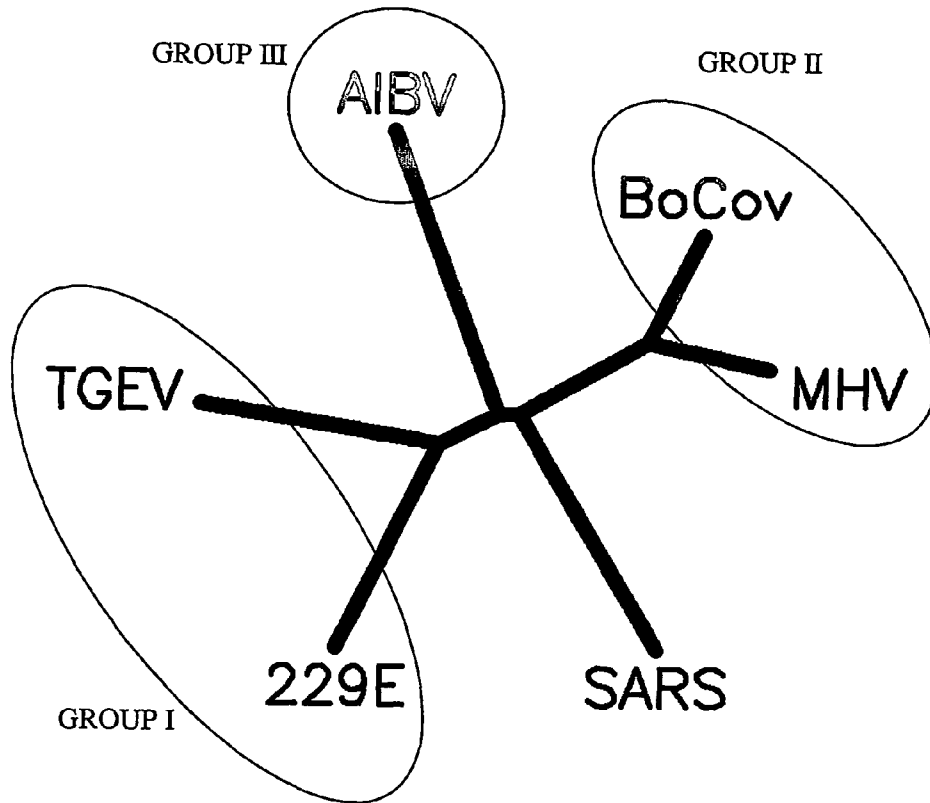


Figure 1A

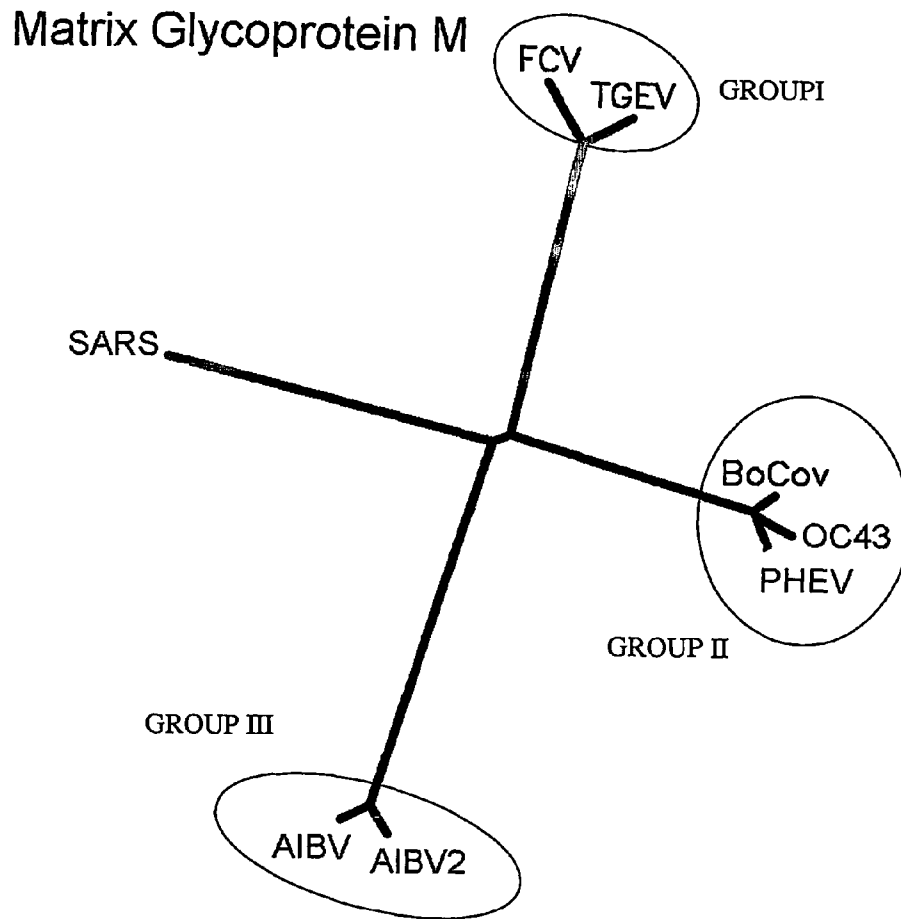


Figure 1B

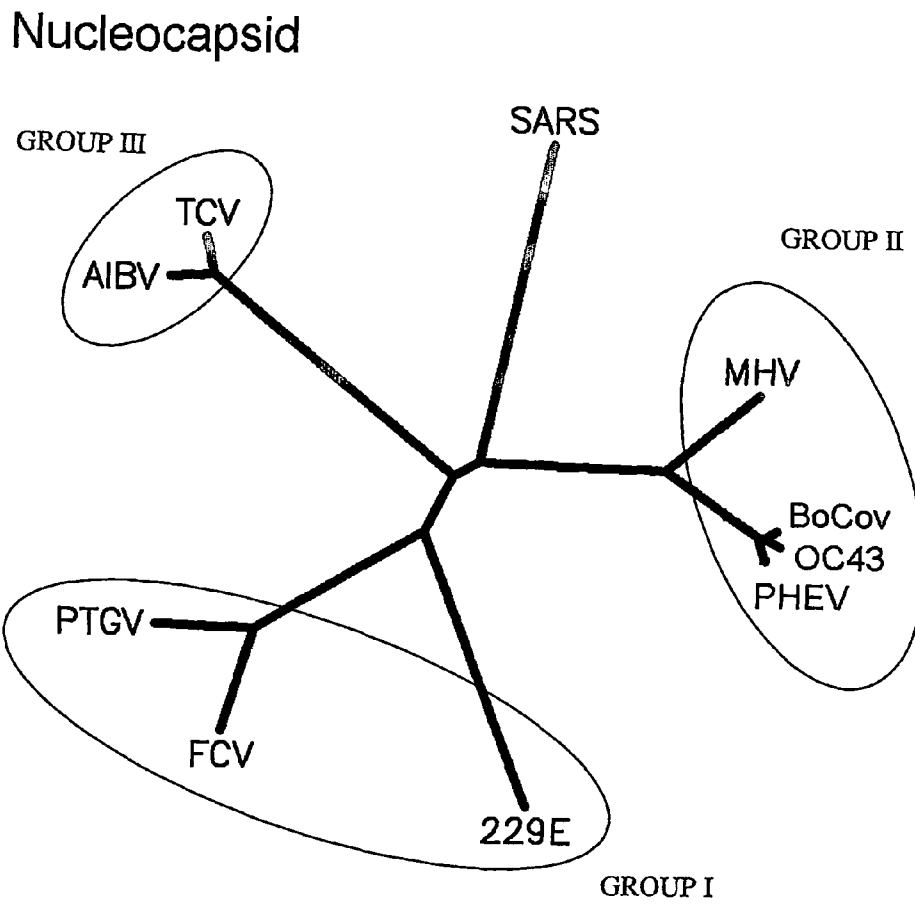


Figure 1C

S (Spike) Glycoprotein

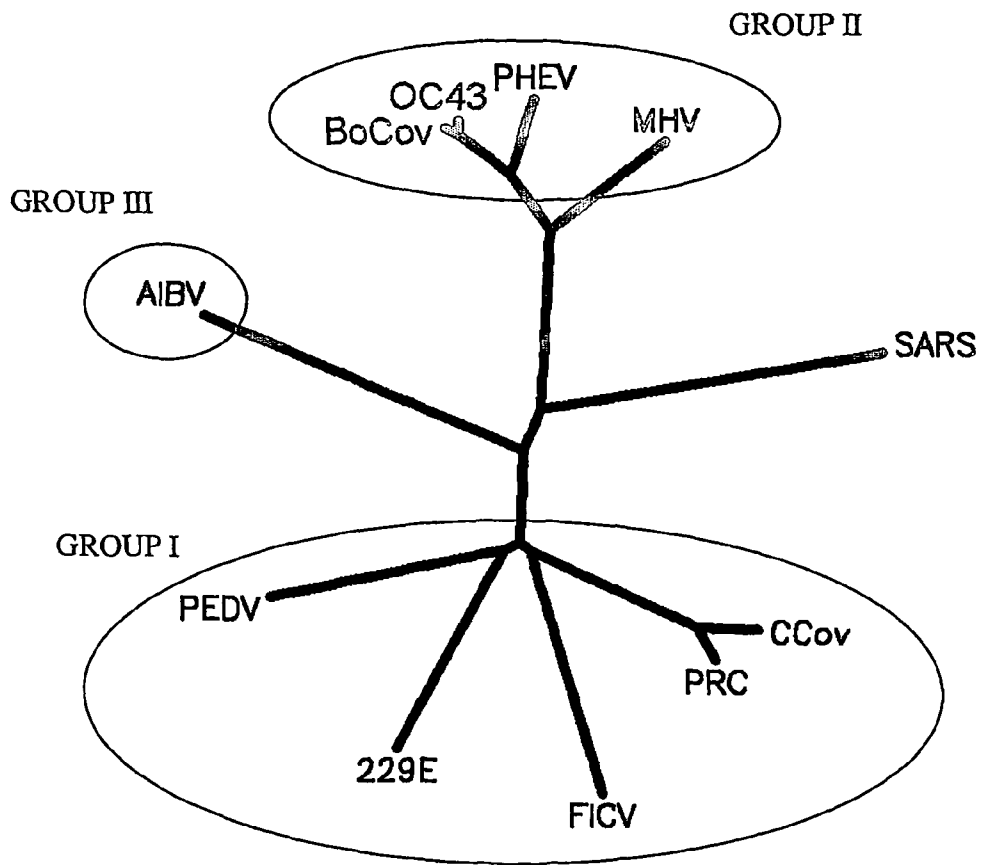


Figure 1D

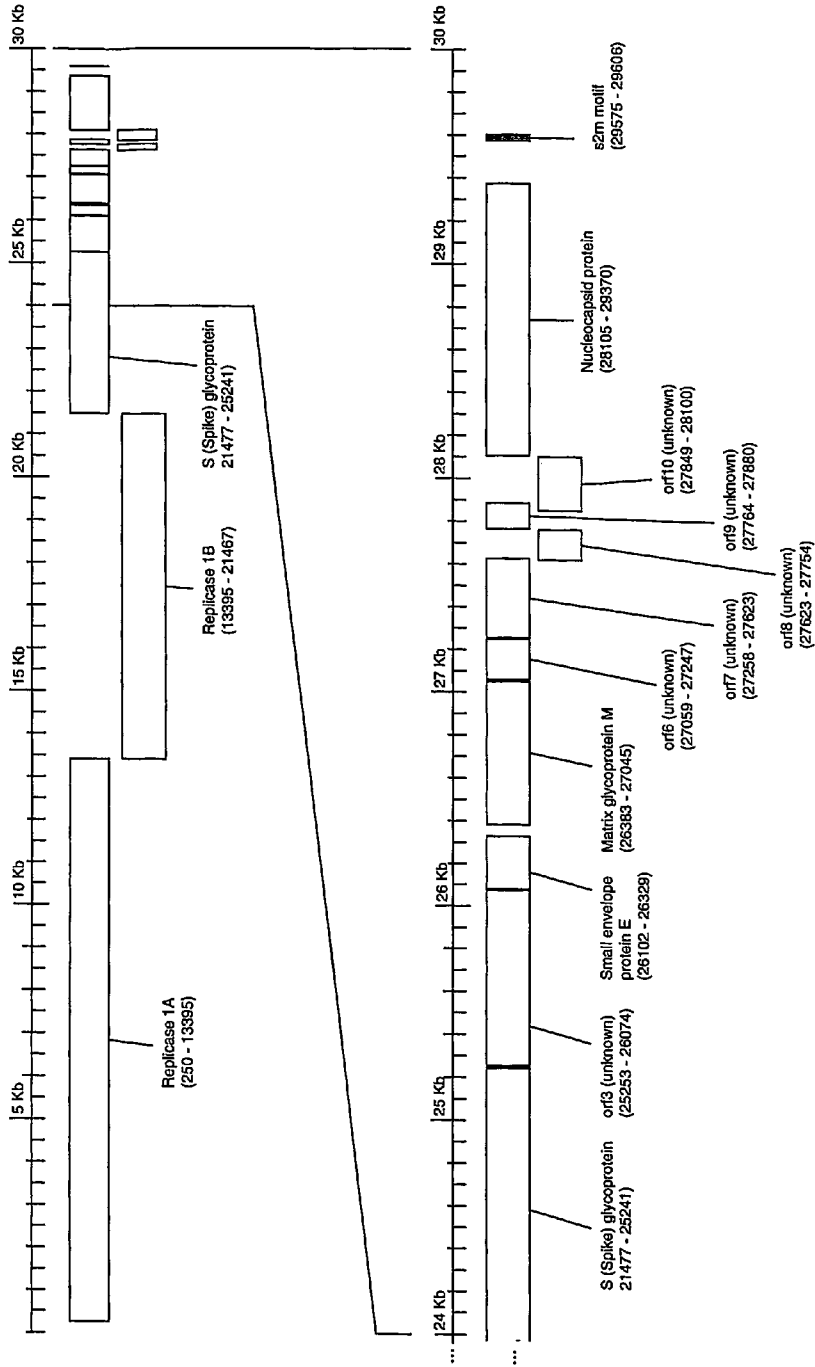


Figure 2

CTACCCAGGAAAAGCCAACCAACCTCGATCTCTTGTAGATCTGTTCTCTAAACGAACTTTAAAATCTGTGT
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AGAAACGAGTAACCTCGTCCCTCTTCTGCAGACTGCTTACGGTTTCGTCCGTGTTGCAGTCGATCATCAGCA
TACCTAGGTTTCGTCCGGGTGTGACCGAAAGGTAAGATGGAGAGCCTTGTCTTGGTGTCAACGAGAAAAC
ACACGTCCAACCTCAGTTTGCCTGTCTTTCAGGTTAGAGACGTGCTAGTGCCTGGCTTCGGGGACTCTGTGG
AAGAGGCCCTATCGGAGGCACGTGAACACCTCAAAAATGGCACTTGTGGTCTAGTAGAGCTGGAAAAAGGC
GTACTGCCCCAGCTTGAACAGCCCTATGTGTTTCAATTAACGTTTCTGATGCCCTAAGCACCAATCACGGCCA
CAAGGTCGTTGAGCTGGTTCAGAAATGGACGGCATTACAGTACGGTTCGTAGCGGTATAACACTGGGAGTAC
TCGTGCCACATGTGGGCGAAACCCCAATTGCATACCGCAATGTTCTTCTTCGTAAGAACGGTAATAAGGGA
GCCGGTGGTCATAGCTATGGCATCGATCTAAAGTCTTATGACTTAGGTGACGAGCTTGGCACTGATCCCAT
TGAAGATTATGAACAAAACCTGGAACACTAAGCATGGCAGTGGTGCACCTCCGTGAACCTCACTCGTGCATCA
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CATTGGGCGTATAGCTTGTGTACCTGTTGCATCCACAGGAGTGAACAATATGCACCTTGTCTACCT
TGATGAAATGTAATCATTGCGATGAAGTTTTCATGGCAGCTGCGACTTTCTGAAAGCCACTTGTGAACAT
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GGGTTGCACCAATTAAGGTTAACTTTGGAGAAGTACTGTTTGGGAAGTTCAGGTTACAAGAATGTG
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GGAAGAACTAAGTTTCTTACCAATAAGTTACTCTTGTGTTGCTGATATCAATGGTAAGCTTTACCATGATT
CTCAGAACATGCTTAGAGGTGAAGATATGCTTTTCTTGGAAAGGATGCACCTTACATGGTAGGTGATGTT

FIGURE 3A

ATCACTAGTGGTGATATCACTTGTGTTGTAATACCCCTCCAAAAGGCTGGTGGCACTACTGAGATGCTCTC
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CACCAGCACTTCAAGAGGCTTATTATAGAGCCCGTGTGGTGTGCTGCTAACTTTTGTGCACTCATACTC
GCTTACAGTAATAAACTGTTGGCGAGCTTGGTGTATGTCAGAGAACTATGACCCATCTTCTACAGCATGC
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FIGURE 3B

CAAGGTGTTGTTGATACCGATGTTGACACAAAGGATGTTATGAATGTCTCAAACCTTTCACATCACTCTGA
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TCCAAGACATGTCATTGACAGCAGAGAAGACATGCTTAATCCTAACAATGAAGATCTGCTCATTCCAAAT
CCAACCATAGCTTCTTGTTCAGGCTGGCAATGTTCAACTTCGTTGTTATTGGCCATTCTATGCAAAAATGT
CTGCTTAGGCTTAAAGTTGATACTTCTAACCCTAAGACACCAAGTATAAATTTGTCGGTATCCAACCTGG
TCAAACATTTTCAAGTTCTAGCATGCTACAATGGTTTCCATCTGGTGTATTACAGTGTGCCATGAGACCTA
ATCATACCATTAAAGTTCTTTCTTAATGGATCATGTGGTAGTGTGGTTTTAACATTGATATGATGCTG
GTGCTTTCTGCTATATGCATCATATGGAGCTTCCAACAGGAGTACACGCTGGTACTGACTTAGAAGGTAA
ATTCTATGGTCCATTGTTGACAGACAAAAGTGCACAGGCTGCAGGTACAGACACAAACATAACATTAAGT
TTTTGGCATGGTGTATGCTGCTGTTATCAATGGTGTATAGGTTTCTTAATAGATTACCACTACTTTG
AATGACTTTAACCTTGTGGCAATGAAGTACAACATGAACTTTGACACAAAGTATGTTGACATATTGGG
ACCTCTTCTGCTCAAACAGGAATGGCGTCTTAGATATGTGTGCTGCTTTGAAAGAGCTGCTGCAGAATG
GTATGAATGGTTCGTACTATCTTGGTAGCACTATTTAGAAGATGAGTTTACACCATTTGATGTTGTTAGA
CAATGCTCTGGTGTACCTTCCAAGGTAAGTTCAAGAAAATTGTTAAGGGCACTCATATTGGATGCTTTT
AACTTTCTTGACATCACTATTGATTTCTGTTCAAAGTACACAGTGGTCACTGTTTCTTTGTTTACGAGA
ATGCTTTCTTGCCATTTACTCTTGGTATTATGGCAATTGCTGCTATGCTATGCTGCTTGTAAAGCATAAG
CAGCATTCTTGTGCTTGTTCGTTACCTTCTCTGCAACAGTTGCTTACTTTAATATGGTCTACATGCC
TGCTAGCTGGGTGATGCGTATCATGACATGGCTTGAATGGCTGACACTAGCTTGTCTGGTTATAGGCTTA
AGGATTGTGTTATGATGCTTCAGCTTTAGTTTTGTCTTATCTCATGACAGCTCGCACTGTTTATGATGAT
GCTGTAGACGTTTGGACACTGATGAATGTCATTACACTGTTTACAAGTCTACTATGGTAATGCTTT
AGATCAAGCTATTTCCATGTGGCCCTTAGTATTTCTGTAACCTCTAATCTTGGTGTGCTTACGACTA
TCATGTTTTTAGCTAGAGCTATAGTGTGTTGTGTGTTGAGTATTACCCATTGTTATTTACTGGCAAC
ACCTTACAGTGTATCATGCTTGTATTATGTTTCTTAGGCTATTGTTGCTGCTACTTTGGCCTTTCTG
TTTACTCAACCGTTACTTCAGGCTTACTCTTGGTGTATTATGACTACTTGGTCTCTACACAAGAATTTAGGT
ATATGAACTCCAGGGGCTTTTGCCCTTAAGAGTAGTATTGCTTTCAAGCTTAAACATTAAGTTGTTG
GGATTGGAGGTAACCATGTATCAAGTTGCTACTGTACAGTCTAAAATGCTGACGTAAGTGCACATC
TGTGGTACTGCTCTCGGTTCTTCAACAACCTAGAGTAGAGTCACTTCTTAAATTTGTTGGGCACAATGTGTAC
AACTCCACAATGATATTCTTCTGCAAAAGACACAACGAAGCTTTCGAGAAGATGGTTTCTTTTGTCT
GTTTTGCTATCCATGCAGGGTCTGTAGACATTAATAGGTTGTGCGAGGAAATGCTCGATAACCGTCTAC
TCTTCAGGCTATTGCTTCAGAATTTAGTCTTTTACCATCATATGCCGCTTATGCCACTGCCAGGAGGCT
ATGAGCAGGCTGTAGCTAATGGTATTTCTGAAGTCTTCTCAAAGGTTAAAGAAATCTTTGAATGTGGCT

FIGURE 3C

AAATCTGAGTTTGACCGTGATGCTGCCATGCAACGCAAGTTGGAAAAGATGGCAGATCAGGCTATGACCCA
AATGTACAAACAGGCAAGATCTGAGGACAAGAGGGCAAAGTAAGTACTAGTGTATGCAAACAATGCTCTTCA
CTATGCTTAGGAAGCTTGATAATGATGCACTTAACAACATTATCAACAATGCGCGTGATGGTTGTGTTCCA
CTCAACATCATACCATTGACTACAGCAGCCAAACTCATGGTTGTTGTCCCTGATTATGGTACCTACAAGAA
CACTTGTGATGGTAACACCTTTACATATGCATCTGCACTCTGGGAAATCCAGCAAGTTGTTGATGCGGATA
GCAAGATTGTTCAACTTAGTGAATTAACATGGACAATTCACCAAATTTGGCTTGGCCTCTTATTGTTACA
GCTCTAAGAGCCAACCTCAGCTGTTAAACTACAGAATAATGAACTGAGTCCAGTAGCACTACGACAGATGTC
CTGTGCGGCTGGTACCACACAAACAGCTTGTACTGATGACAATGCACCTTGCTACTATAACAATTCGAAGG
GAGGTAGGTTTGTGCTGGCATTACTATCAGACCACCAAGATCTCAAATGGGCTAGATTCCCTAAGAGTGAT
GGTACAGGTACAATTTACACAGAACCTGGAACCACCTTGTAGGTTTGTTCAGACACACCAAAAAGGGCTTAA
AGTGAATACTTGTACTTTCATCAAAGGCTTAAACAACCTAAATAGAGGTATGGTGTGCTGGGCAGTTTAGCTG
CTACAGTACGCTTCAGGCTGGAAATGCTACAGAAGTACCTAGCAAGTGGAGGACAACCAATCACCACATGTGT
TTTGCAGTAGACCCCTGCTAAAGCATATAAGGATTACCTAGCAAGTGGAGGACAACCAATCACCACATGTGT
GAAGATGTTGTGTACACACACTGGTACAGGACAGGCAATTACTGTAACACCAGAAGCTAACATGGACCAAG
AGTCCCTTGGTGGTGCCTTCATGTTGCTGTATTGTAGATGCCACATTGACCATCCAAATCCTAAAGGATTC
TGTGACTTGAAGGTAAGTACGTCCAAATACCTACCCTTGTGCTAATGACCCAGTGGGTTTTACACTTAG
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TGATGCACTCTGCGGATGCATCAACGTTTAAACGGGTTTGGCGTGAAGTGCAGCCCGTCTTACACCGT
GCGGCACAGGCACTAGTACTGATGTCGCTTACAGGCTTTTGTATTTACAACGAAAAGTTGCTGGTTTT
GCAAAGTTCCCTAAAACTAATTGCTGTCGCTTCCAGGAGAAGGATGAGGAAGGCAATTTATFAGACTCTTA
CTTTGTAGTTAAGAGGCATACTATGCTAACTACCAACATGAAGAGACTATTTATAACTTGGTTAAAGATT
GTCCAGCGGTTGCTGTCCATGACTTTTTCAAGTTTLAGAGTAGATGGTGACATGGTACCACATATACACGT
CAGCGTCTAACTAAATACACAATGGCTGATTTAGTCTATGCTCTACGTCATTTTGTATGAGGGTAATTGTGA
TACATTAAGAAATACTCGTCAACATAACAATGCTGTGATGATGATTTTCAATAAGAAGGATTGGTATG
ACTTCGTAGAGAATCCGTGACATCTTACCGGTATATGCTAAGTGGTACGCGTGTACGCCAATCATTATTA
AAGACTGTACAATTTCTGCGATGCTATGCGTGTATGCGGATTTGTAGGCGTACTGACATTAGATAATCAGGA
TCTTAATGGGAACCTGGTACGATTTCCGGTATTTCTGTACAAGTAGCACCAGGCTGCGGAGTTCCCTATTGTGG
ATTCATATTACTCATTTGCTGATGCCCATCCCTCATTGACTAGGGCATTGGCTGCTGAGTCCCATATGGAT
GCTGATCTCGCAAACCACTTATTAAGTGGGATTTGCTGAAATATGATTTTACGGAAGAGAGACTTTGTCT
CTTCGACCGTTATTTAAATATTGGGACCAGACATACCATCCCAATTTGATTAACCTGTTTGGATGATAGGT
GTATCCTTCATTTGTCAAACCTTTAATGTGTTATTTCTACTGTGTTCCACCTACAAGTTTGGACCACATA
GTAAGAAAATATTTGTAGATGGTGTTCCTTTTGTGTTTCAACTGGATACCATTTTTCGTGAGTTAGGAGT
CGTACATAATCAGGATGTAACTTACATAGCTCGCGTCTCAGTTTCAAGGAACCTTTTAGTGTATGCTGCTG
ATCCAGCTATGCATGCAGCTTCTGGCAATTTATTGTAGATAAAGCCTACATGCTTTTTCAGTAGCTGCA
CTAACAAACAATGTTGCTTTTCAAACCTGTCAAACCCGTAATTTTAAATAAGACTTTTATGACTTTGCTGT
GTCTAAAGGTTTCTTTAAGGAAGGAAGTCTGTGAACTAAAACACTTCTTCTTTGCTCAGGATGGCAACG
CTGCTATCAGTGATTTATGACTATTATCGTTATAATCTGCCAACATGTGTGATATCAGACAACCTCCTATT
GTAGTTGAAGTTGTTGATAAATACTTTGATTGTTACGATGGTGGCTGTATTAATGCCAACCAAGTAATCGT
TAACAATCTGGATAAATCAGCTGGTTTCCATTTAATAAATGGGGTAAGGCTAGACTTTTATTTAGCTCAA
TGAGTTATGAGGATCAAGATGCATTTTTCGCGTATACTAAGCGTAATGTCATCCCTACTATAACTCAAATG
AATCTTAAGTATGCCATTAGTGCAAGAATAGAGCTCGCACCGTAGCTGGTGTCTCTATCTGTAGTACTAT
GACAAATAGACAGTTTCATCAGAAATATTGAAGTCAATAGCCGCCACTAGAGGAGCTACTGTGGTAATTG
GAACAAGCAAGTTTACGGTGGCTGGCATAATATGTTAAACCTGTTTACAGTGATGTAGAACTCCACAC
CTTATGGGTTGGGATTTATCCAAAATGTGACAGAGCCATGCCAATGCTTAGGATAATGGCCTCTCTTGT
TCTTGCTCGCAAACATAACACTTGTGTAACCTATCACACCGTTTCTACAGGTTAGCTAACGAGTGTGCGC
AAGTATTAAGTGAATGGTTCATGTTGCGGCTCCTATATGTTAAACCAGGTGGAACATCATCCGGTGTAT
GCTACAACCTGCTTATGCTAATAGTGTCTTTAACATTTGTCAAGCTGTTACAGCCAATGTAATGCACCTCT
TTCACACTGATGGTAATAAGATAGCTGACAAGTATGTCGCAATCTACAACACAGGCTCTATGAGTGTCTCT
ATAGAAATAGGGATGTTGATCATGAATTCGTGGATGAGTTTACGCTTACCTGCGTAAACATTTCTCCATG
ATGATTCTTCTGATGATGCCGTTTGTGCTATAACAGTAACTATGCGGCTCAAGGTTTAGTACTAGACTAGCA
TAAGAAGTTTAAAGGAGTTCTTTATATCAAATAATGTGTTTCAATGCTGCTGAGGCAAAATGTTGGACTGAGA
CTGACCTTACTAAAGGACCTCACGAATTTGCTCACAGCATACAATGCTAGTTAAACAAGGAGATGATTAC
GTGTACCTGCCCTTACCAGATCCATCAAGAATATTAGGCGCAGGCTGTTTTGTGCTGATGATTTGTCAAAC
AGATGGTACACTTATGATTGAAAGGTTCTGTGCTACTGGCTATTGATGCTTACCACCTTCAAACATCCCTA
ATCAGGAGTATGCTGATGCTTTTCACTTGTATTTACAATACATTAGAAAAGTTACATGATGAGCTTACTGGC
CACATGTTGGACATGATTTCCGTAATGCTAACAATGATAACACCTCACGGTACTGGGAACCTGAGTTTAA
TGAGGCTATGTACACACCACATACAGTCTTGCAGGCTGTAGGTGCTTGTGATTTGTGCAATTCACAGACTT

FIGURE 3D

CACTTCGTTGCGGTGCCTGTATTAGGAGACCATTCCATGTTGCAAAGTGCTGCTATGACCATGTCATTTCA
ACATCACACAAATAGTGTGTCTGTAAATCCCTATGTTGCAATGCCCCAGGTTGTGATGTCACCTGATGT
GACACAACGTATCTAGGAGGTATGAGCTATTATTGCAAGTCACATAAGCCTCCCATAGTTTCCATAT
GTGCTAATGGTCAGGTTTTGGTTTATACAAAAACACATGTGTAGGCAGTGACAATGTCACCTGACTTCAAT
GCGATAGCAACATGTGATTGGACTAATGCTGGCGATTACATACTTGCCAACACTTGTACTGAGAGACTCAA
GCTTTTCGAGCAGAAACGCTCAAAGCCACTGAGGAAACATTTAAGCTGTCTATGGTATTGCCACTGTAC
GCGAAGTACTCTGACAGAGAATTGCATCTTTCATGGGAGGTTGGAACCCTAGACCACCATTTGAACAGA
AACTATGCTTTACTGGTTACCGTGTAACFAAAAAATAGTAAAGTACAGATTGGAGAGTACACCTTTGAAAA
AGGTGACTATGGTGTGCTGTTGTGTACAGAGGTACTACGACATACAAGTTGAATGTTGGTGTACTTTG
TGTGACATCTCACACTGTAATGCCACTTAGTGCACCTACTCTAGTGCCACAAGAGCACTATGTGAGAATT
ACTGGCTGTACCCAACTCAACATCTCAGATGAGTTTCTAGCAATGTTGCAAATTTCAAAAGGTCGG
CATGCAAAAGTACTCTACACTCCAAGGACCACCTGGTACTGGTAAGAGTCAATTTGCCATCGGACTTGCCT
TCTATTACCCTATGCTCGCATAGTGTATACGCGATGCTCTCATGCAGCTGTTGATGCCCTATGTGAAAG
GCATTAATAATTTGCCCATAGATAAATGTAGTAGAATCATACTGCGCGTGCCGCGTAGAGTGTTTTGA
TAAATTCAAAGTGAATFCAACACTAGAACAGTATGTTTCTGCACTGTAAATGCATTTGCCAGAAACA
ACTGCTGACATTTAGTCTTTGATGAAATCTCTATGGCTACTAATTATGACTTGAGTGTGTCAATGCTAG
ACCTCGTGCAAAACACTACGCTCTATATTGGCGATCTGCTCAATTACCAGCCCCCGCACATTTGCTGACT
AAAGCACACTAGAACAGAAATATTTAATFCAGTGTGCAGACTTATGAAAACAATAGGTCAGACATGTTCT
TGGAACCTTGTCCTGCTGFAAATGTTGACACTGTGAGTGTCTTAGTTTATGACAATAAGCTAAAA
GCACACAAGGATAAGTCACTCAATGCTTCAAATGTTCTACAAAGGTTTATTACACATGATGTTTCATC
TGCAATCAACAGACCTCAAATAGGCGTGTAGAGAAATTTCTTACACGCAATCCGCTTGGAGAAAAGCT
TTTTTATCTCACCTTATAATTCACAGAAGCTGTAGCTTCAAAAATCTTAGGATTTGCCATCGCAGACT
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TTGTAA
TGTCACCCGCTTCAATGTGGCTATCACAAGGGCAAAAATTTGGCATTGTTGTGCATAATGTCTGATAG
AGATC
TTTTATGACAACTGCAATTTACAAGTCTAGAAAATACCACGTCGCAATGTGGCTACATTACAAGCAGAA
AAT
GTAAGTGGACTTTTTAAGGACTGTAGTAAGATCAATCTGCTTTCATCTTACACAGGCACCTACACACCT
CAGCGTTGATATAAAGTTCAAGACTGAAGGATTTATGTTGACATACCAGGCATACCAAAGGACATGACCT
ACCGTAGACTCATCTCTATGATGGGTTTCAAATGAATTAACAAGTCAATGGTTACCTAATATGTTTATC
ACCCGGAAGAAGCTATTCGTCACGTTTCGTGCGTGGATTGGCTTTGATGTAGAGGGCTGTCAATGCA
ACTAG
AGATGCTGTGGTACTAACCTACCTCTCCAGCTAGGATTTTCTACAGGTGTTAACTTAGTAGCTGTACCG
A
CTGGTTATGTTGACACTGAAAATAACACAGAATTCACCAGAGTTAATGCAAAACCTCCACCAGGTGACC
G
TTTAAACATCTTATACCCTCATGTATAAAGGCTTGCCCTGGAATGTAGTCCGTATTAAGATAGTACA
AAT
GCTCAGTGATACACTGAAAGGATTTGTGACAGAGTTCGTGTTTCGTCCTTTGGGCGCATGGCTTTGAG
CTTA
CATCAATGAAGTACTTTGTCAAGATTTGGACCTGAAAGAACGTTGTCTGTGTGACAAACGTCACACT
TGC
TTTTCTACTTCATCAGATACTTATGCCCTGCTGGAATCATTCGTGGGTTTGGACTATGCTATAACCC
ATT
TATGATTTGATGTTTCAGCAGTGGGGCTTTACGGGTAACTTCAGAGTAACCATGACCAACATTGCC
AGTAC
ATGGAAATGCACATGTGGCTAGTTGTGATGCTATCATGACTAGATGTTTAGCAGTCCATGAGTGTCT
TGT
AAGCGCGTTGATTTGGTCTGTTGAATACCCCTATTATAGGAGATGAACGAGGTTAATTTCTGCTG
CAGAAA
AGTACAACACATGGTGTGGAAGTCTGCATTTGCTGATAAGTTCCAGTCTTTCATGACATTGGAAAT
C
CAAAGGCTATCAAGTGTGTGCCCTCAGGCTGAAGTGAATGGAAGTTCTACGATGCTCAGCCATGTAG
TGAC
AAAGCTTACAAAATAGAGGAACTTCTCTATTCTTATGCTACACATCACGATAAATTCACATGAT
GGTGT
TTTGTTTTGGAAATGTAACGTTGATCGTTACCAGCCAATGCAATTTGTGTGTAGGTTTGACACA
AGAGTCT
TGTCAAACCTTGAACCTTACCAGGCTGTGATGGTGGTAGTTTGTATGTGAATAAGCATGCATTTCC
ACACTCA
GCTTTCGATAAAAAGTGCATTTACTAATTTAAAGCAATTTGCCTTTCTTTTACTATTTCTGATAG
TCTTT
GTCTCATGGCAACAAGTAGTGTCCGATATGATTAATGTTCCACTCAAATCTGCTACGTGTATTACAC
GAT
GCAATTTAGGTGGTGTCTGTTTCGACACACCATGCAAAATGAGTACCAGCAGTACTTGGATGCAT
ATAATG
ATGATTTCTGCTGGATTTAGCCTATGGATTTACAACAATTTGATACTTATAACCTGTGGAATACAT
TTAC
CAGGTTACAGAGTTTGAAAAATGTGGCTTATAATGTTGTTAATAAAGGACACTTTGATGGACAGC
CCGCG
AAGCACCTGTTTCCATCATTAAATAATGCTGTTTACACAAAGGTAGATGGTATTGATGTGGAGAT
CTTTG
AAATAAGACAACACTTCTGTTAATGTTGCATTTGAGCTTTGGGCTAAGCGTAACATTAAACCAGT
GCCAG
GATTAAGATACTCAATAATTTGGGTGTTGATATCGCTCTAATACTGTAATCTGGGACTACAAA
GAGAAG
CCCCAGCACATGTATCTACAATAGGTGCTGCACAATGACTGACATTTGCCAAGAAACCTACTGAG
AGTGT
TGTCTTCACTACTGTCTTGTGTTGATGGTAGAGTGGAGGACAGGTAGACCTTTTGA
AAACCC
TGGTGTTTTAATAACAGAAGGTTTCAAGTCAAAGGCTAACACCTTCAAAGGGACCAGCAAGCTAG
CGTCA
ATGGAGTCAATTAATTTGGAGAAATCAGTAAAAACACAGTTTAACTACTTTAAGAAAGTAGAC
GGCATT
CAACAGTTGCCTGAAACCTACTTTACTCAGAGCAGAGACTTAGAGGATTTTAAAGCCAGAT
CAAAAT
AACTGACTTTCTCGAGCTCGCTATGGATGAATTCATACAGCGATATAAGCTCGAGGGCTATGCCT
TCGAAC

FIGURE 3E

ACATCGTTTATGGAGATTTTCAGTCATGGACAACCTGGCGGTCTTCATTTAATGATAGGCTTAGCCAAGCGC
TCACAAGATTACCACCTTAAATTAGAGGATTTTATCCCTATGGACAGCACAGTGAAAAATTACTTCATAAC
AGATGCGCAAACAGGTTTCATCAAAATGTGTGTGTTCTGTGATGATCTTTACTTGTACTTTGTGCGAGA
TAATAAAGTCACAAGATTTGTCAAGTATTCAAAAAGTGGTCAAGGTTACAATGACTATGCTGAAATTTCA
TTCATGCTTTGGTGTAAAGATGGACATGTTGAAACCTTCTACCCAAAACATAAGCAAGTCTGAGCGTGGCA
ATCAGGTGTTGCGATGCTTAACCTGTACAAGATGCAAAGAATGCTTCTTGAAGAAGTGTGACCTTCAGAATT
ATGGGTGAAAATGCTGTTATACCAAAGGAATAATGATGAATGTGCGAAAGTATACTCAACTGTGTCAATAC
TTAAATACACTTACTTTAGCTGTACCCCTACAACATGAGAGTTATTACCTTTGGTGTGCTGGCTCTGATAAAGG
AGTTGCACCAGGTACAGCTGTGCTCAGACAATGGTTGCCAAGTGGCACACTACTTGTGCGATTTCAGATCTTA
ATGACTTCGTCTCCGACGCATATTTACTTTAATTTGGAGACTGTGCAACAGTACATACGGCTAATAAATGG
GACCTTATTTATTAGCGATATGATGACCCTAGGACCAAACATGTGACAAAAGAGAATGACTCTAAAAGAAGG
GTTTTTCACTTATCTGTGTGGATTATAAAGCAAAAACCTAGCCCTGGGTGGTTCTATAGCTGTAAAGATAA
CAGAGCATTTTGGAAATGCTGACCTTTACAAGCTTATGGGCCATTTCTCATGGTGGACAGCTTTTGTTACA
AATGTAATGTCATCATCGGAAGCATTTTTAATTTGGGGCTAACTATCTTGGCAAGCCGAAGGAACAAAT
TGATGGCTATACCATGTCATGCTAACTACATTTTCTGGAGGAACAAATCCATCCAGTTGTCTTCTTAT
CACTCTTTGACATGAGCAAATTTCCCTTAAATTAAGAGGAACCTGCTGTAATGTCTCTTAAGGAGAATCAA
ATCAATGATATGATTTATTTCTTCTTGGAAAAGGTAGGCTTATCATTAGAGAAAACAACAGAGTTGTGGT
TTCAAGTGATATTTCTTGAACAACATAACGAACATGTTTATTTCTTATTTATTTCTTACTCTCACTAGTG
GTAGTGACCTTGGCCGTGACCACTTTTGTGATGATGTTCAAGCTCCTAATTACACTCAACATACCTTCATCT
ATGAGGGGGGTTTACTATCCTGATGAAATTTTTAGATCAGACACTCTTTAATTTAACTCAGGATTTATTTCT
TCCATTTTATTTAATGTTACAGGTTTCTACTATTAATCATACGTTTGGCAACCCTGTCATACCTTTTA
AGGATGTTATTTATTTGCTGCCACAGAGAAATCAAATGTTGTCCGTGGTTGGGTTTTTGGTTCTACCATG
AACAAAGTCACAGTCGGTGATTATTTAACAATTTCTACTAATGTTGTTATACGAGCATGTAACCTTTGA
ATTTGTGTGACAACCCCTTTCTTTGCTGTTCTAAACCCATGGGTACACAGACACATACTATGATATTCGATA
ATGCATTTAATGTCACCTTCGAGTACATATCTGATGCTTTTCCCTTGTGATGTTTCAGAAAAGTCAGGTAAT
TTAAACACTTACGAGAGTTGTGTTTAAAAATAAGATGGGTTTCTCTATGTTTATAAGGGCTATCAACC
TATAGATGTAGTTCGTGATCTACCTTCTGGTTTTAACACTTTGAAACCTATTTTTAAGTTGCCCTTGGTA
TTAACATTACAAATTTTAGAGCCATTCTTACAGCCTTTTCACCTGCTCAAGACATTTGGGGCACGTCAGCT
GCAGCCTATTTTGTGGCTATTTAAAGCCAACCTACATTTATGCTCAAGTATGATGAAAATGGTACAATCAC
AGATGCTGTTGATGTTCTCAAATCCACTTGCTGAACTCAAATGCTCTGTTAAGAGCTTTGAGATGACA
AAGGAATTTACCAGACCTCTAATTTTCAGGGTGTTCCTCAGGAGATGTTGTGAGATTCCCTAATATTACA
AACTTGTGTCTCTTTGGAGAGGTTTTAATGCTACTAAATCCCTTCTGTCTATGCATGGGAGAGAAAAAA
AATTTCTAATTTGTTGCTGATTACTCTGTGCTCAACTCAACATTTTTTTCAACCTTTAAGTGTCTATG
GCTTTCTGCCACTAAGTTGAATGATCTTTGCTTCTCCAATGTCTATGCAGATTCTTTTGTAGTCAAGGGA
GATGATGTAAGACAAATAGCGCCAGGACAACTGGTGTATTGCTGATATAAATTAATAATGGCAGATGA
TTTCATGGGTTGTGCTTGTGGAATACTAGGAACATTTGATGCTACTTCAACTGGTAATATAAATATA
AATATAGGTATCTTAGACATGGCAAGCTTAGGCCCTTTGAGAGAGACATATCTAATGTGCTTTCTCCCT
GATGGCAAACCTTGCACCCCACTGCTCTTAATTTGCTTATTTGGCCATTTAAATGATTATGTTTGTTCACAC
TACTGGCATTGGCTACCAACCTTACAGAGTTGTAGTACTTTCTTTTGAACTTTTAAATGCACCGGCCACGG
TTTGTGGACCAAATTTACTGACCTTATTAAGAACCAGTGTGTCAATTTAATTTAATGGACTCACT
GGTACTGGTGTGTTAACTCCCTCTTCAAAGAGATTTCAACCATTTCAACAATTTGGCCGTGATGTTCTGA
TTTCACTGATTCGGTTCGAGATCCTAAAACATCTGAAATATTAGACATTTCACTTGGCGCTTTTGGGGGTG
TAAGTGAATTACACCTGGAACAAATGCTTCATCTGAAGTTGCTGTTCTATATCAAGATGTTAACTGCAC
TACTGTTCTACAGCAATTCATGCAGATCAACTCACACCAGCTTGGCGCATATATTTACTGGAAACAATGT
ATTCCAGACTCAAGCAGGCTGTCTTATAGGAGCTGAGCATGTGACACCTCTTATGAGTGGACATTCCTA
TTGGAGCTGGCATTGTGCTAGTTACCATACAGTTTCTTTATTACGTAGTACTAGCCAAAATCTATTTGTG
GCTTATACTATGCTTTAGGTGCTGATAGTTCAATTTGCTTACTCTAATAACACCATTGCTATACCTACTAA
CTTTTCAATTAGCATTACTACAGAAGTAATGCCTGTTTCTATGGCTAAAACCTCCGTAGATTGTAATATGT
ACATCTGCGGAGATTTACTGAAATGTGCTAATTTGCTTCTCAAATATGGTAGCTTTTGCACACAACATAAT
CGTGCCTCTCAGGTATTGCTGCTGAACAGGATCGCAACACACGTGAAGTGTTCGCTCAAGTCAAACAAAT
GTACAAAACCCCAACTTTGAAATATTTGGTGGTTTTAATTTTTCACAAATATTACCTGACCCCTCAAAGC
CAACTAAGAGGTCTTTTATTGAGGACTTGCTCTTTAATAAGGTGACACTCGCTGATGCTGGCTTCATGAAG
CAATATGGCAATGCCTAGGTGATATAATGCTAGAGATCTCATTTGTGGCAGAAAGTTCAATGGACTTAC
AGTGTGGCCACTCTGCTCACTGATGATATGATTTGCTGCTTACTGCTTACTGCTGCTAGTTAGTGGTACTGCCA
CTGCTGGATGGACATTTGGTGTGGCGCTGCTCTTCAAATACCTTTTGGCTATGCAAATGGCATATAGGTTT
AATGGCATTTGGAGTTACCCAAAATGTTCTCTATGAGAACCAAAACAAATCGCCAAACCAATTTAAACAGGC
GATTAGTCAAAATCAAGAATCACTTACAACAACATCAACTGCATTTGGCAAGCTGCAAGACGTTGTTAACC

FIGURE 3F

AGAATGCTCAAGCATTAAACACACTTGTTAAACACTTAGCTCTAATTTGGTGCAATTTCAAGTGTGCTA
AATGATATCCTTTTCGCGACTTGATAAAGTCGAGGCGGAGGTACAAATTTGACAGGTTAATTACAGGCAGACT
TCAAAGCCTTCAAACCTATGTAAACAACAACATAATCAGGGCTGCTGAAATCAGGGCTTCTGCTAATCTTG
CTGCTACTAAAATGTCTGAGTGTGTTCTTGGACAATCAAAAAGAGTTGACTTTTGTGGAAAGGGCTACCAC
CTTATGTCTTCCCACAAGCAGCCCCGCATGGTGTGCTTTCCTACATGTCACGTATGTGCCATCCCAGGA
GAGGAACCTTACCACAGCGCCAGCAATTTGTCTATGAAGGCAAAGCATACTTCCCTCGTGAAGGTGTTTTTG
TGTTAATGGCACTTCTTGGTTTATTACACAGAGGAACCTCTTTTCTCCACAAATAATTACTACAGACAA
ACATTTGTCTCAGGAAATGTGATGTGCTTATTGGCATCATTAAACAACACAGTTTATGATCCTCTGCAACC
TGAGCTTGACTCATTCAAAGAAGAGCTGGACAAGTACTTCAAATAATCATACATCACCAGATGTTGATCTTG
GCGACATTTACAGGCATTAACGCTTCTGTGCTCAACATTCAAAAAGAAATGACCCCTCAATGAGGTGCGCT
AAAAATTTAAATGAATCACTCATGACCTTCAAGAATTTGGGAAAATATGAGCAATATATTAATGGCCTTG
GTATGTTTGGCTCGGCTTCATTGCTGGACTAATGGCAATCGTTCATGGTTACAATCTTGCTTTGTTGCATGA
CTAGTTGTTGCAAGTTGCCCTCAAGGGTGCATGCTCTTGTGTTCTTGCTGCAAGTTGATGAGGATGACTCT
GAGCCAGTTCCTCAAGGGTGTCAAATACATTACACATAAACGAACCTTATGGATTTGTTTATGAGATTTTT
ACTCTGGATCAATTAAGCAGCCAGTAAATAATGACAATGCTTCTCCTGCAAGTACTGTTTACTGCTAC
AGCAACCTGACTACCGCTACAAGCCTCACTCCCTTTCGGATGGCTTGTATTGGCGTTGCATTTCTTGCTTT
TTCAGAGCGCTACCAAATAAATGGCGCTCAATAAAGATGGCAGCTAGCCCTTTATAAGGGCTTCCAGTTC
ATTTGCAATTTACTGCTGCTATTTGTTTACCATCTATTACATCTTTTGCTTGTGCTGCAGGTATGGAGGC
GCAATTTTTGTACTCTATGCCCTTGATATATTTTCTACAATGCATCAACGCATGTAGAAATATTATGAGAT
GTTGGCTTTGTTGGAAGTGCAAATCCAAGAACCATTACTTTATGATGCCAATCTTTGTTGCTGGCAC
ACACATAACTATGACTACTGTATACCATAAACAGTGTACAGATACAATGTGCTTACTGAAGGTGACGG
CAATTTCAACACCAAACTCAAAGAAGACTACCAAATTTGGTGGTTATTCTGAGGATAGGCACTCAGGTGTTA
GACTATGTGCTGTGACTGCTATTTACCGAAGTTTACTACCAGCTTGAGTCTACACAAATTAACATA
GACTGCTGATTTGAAAATGCTACATCTTTCATCTTTAAACAAGCTTGTAAAGACCACCGAATGTGCAAAT
ACACACAATCGACGGCTCTTCAGGAGTTGCTAATCCAGCAATGGATCCAATTTATGATGAGCCGACGACGA
CTACTAGCGTGCCTTTGTAAGCACAAGAAAGTGAAGTACGAACCTTATGACTCATTGCTTTGGAAGAAACA
GGTACGTTAATAGTTAATAGCGTACTTCTTTTCTTGCTTTGCTGCTATCTTGCTAGTACACTAGCCAT
CCTTACTGCGCTTCGATTTGTGTGCGTACTGCTGCAATATTGTTAACGTGAGTTTAGTAAAACCAACGGTTT
ACGCTACTCGCGTGTAAAAATCTGAACCTTCTGAAGGAGTTCTGATCTTCTGGTCTAAACGAACCTAA
CTATTATTATTATTCTGTTTGAACCTTAAACATTTGCTTATCATGCGAGACAACGGTACTTATCCGTTGAG
GAGCTTAAACAACCTCCCTGGAACAATGGAACCTAGTAATAGGTTTCCCTATTCCTAGCCTGGATTATGTTACT
ACAATTTGCCTATTCTAATCGGAACAGGTTTTTGTACATAATAAAGCTTGTTTTCTCTGGCTCTTGTGGC
CAGTAACACTTGCTTGTTTGTGCTTGTGCTGCTTACAGAATTAATGGGTGACTGGCGGGATTGCGATT
GCAATGGCTTGTATTGTAGGCTTGATGTGGCTTAGCTACTTCGTTGCTTCCCTTCAGGCTGTTTGTCTGTA
CCGCTCAATGTGGTCAATCAACCCAGAAACAACATTTCTTCTCAATGTGCTCTCCGGGGACAATTTGTA
CCAGACCGCTGATGGAAGTGAACCTTGTCTATGGTGTCTGTGATCATTCGTTGGTCACTTGCCAATGGCCGA
CACTCCCTAGGGCGCTGTGACATTAAGGACCTGCCAAAGAGATCACTGTGGCTACATCAGCAATGCTTTC
TTATTACAAATTAGGAGCGTGCAGCGTGTAGGCTACTGATTCAGGTTTTGCTGCATACAACCGCTACCCTA
TTGGAACTATAAATTAATAACAGACCACCGCGGTAGCAACGACAATATGCTTTGCTAGTACAGTAAGTG
ACAACAGATGTTTCACTTGTGACTTCCAGGTTACAATAGCAGAGATATTGATTATCATTATGAGGACTT
TCAGGATGCTATTTGGAATCTGACGTTATAATAAGTTCAATAGTGAGACAATTAATTAAGCCTCTAAT
AAGAAGATTTATTCGGAGTTAGATGATGAAGAACCCTATGGAGTTAGATTATCCATAAAACGAACATGAAA
TTATCTCTTCTGACATGATTTGATTTACATCTTGGAGCTATATCATTATCAGGAGTGTGTTAGAGGT
ACGACTGTACTACTAAAAGAACCTTGCCCATCAGGAACATACGAGGGCAATTCACCATTTACCCTCTTGC
TGACAATAAATTTGCACATAACTTGCACTAGCACACACTTTGCTTTTGTGCTGACGGTACTCGACATA
CCTATCAGCTGCGTGCAAGATCAGTTTACCAAACCTTTTTCATCAGACAAGAGGAGGTTCAACAAGAGCTC
TACTCGCCACTTTTCTCATTTGTTGCTGCTCTAGTATTTTAATACTTTGCTTACCATTAAGAGAAAGAC
AGAATGAATGAGCTCACTTTAATGACTTCTATTTGTGCTTTTGTAGCCTTTCTGCTATTCCTTGTTTTAA
AATGCTTATTATATTTTGGTTTTCACTCGAAATCCAGGATCTAGAAGAACCTTGTACCAAAGTCTAAACGA
ACATGAAACTTCTCATTGTTTTGACTTGTATTCTCTATGCAGTTGCATATGCAGTGTAGTACAGCGCTGT
GCATCTAATAAACCTCATGTGCTTGAAGATCCTTTGAAGGTACAACACTAGGGGTAATFACTATAGCACATG
CTTGGCTTTGCTGCTTAGGAAAGTTTTACCTTTTCAATAGATGGCACACTATGGTTCAAACATGCACACCT
AATGTTACTATCAACTGTCAAGATCCAGCTGGTGGTGGCTTATAGCTAGGTGTTGGTACCTTCATGAAGG
TCACCAAACGCTGCATTTAGAGACGTAATTTGTTGTTTTAAATAAACGAACAAATTAATAATGCTGATAAT
GGACCCCAATCAAACCAACGTAGTGCCCCCGCATTACATTTGGTGGACCCACAGATTCAACTGACAATAA
CCAGAATGGAGGACGCAATGGGGCAAGGCCAAAACAGCGCCGACCCCAAGGTTTACCCATAAATACTGCGT
CTTGGTTTACAGCTCTCACTCAGCATGGCAAGGAGGAACCTTAGATTCCTCGAGGCCAGGGCGTTCCAATC

FIGURE 3G

AACACCAATAGTGGTCCAGATGACCAAAATGGCTACTACCGAAGAGCTACCCGACGAGTTCGTGGTGGTGA
CGGCAAAATGAAAGAGCTCAGCCCCAGATGGTACTTCTATTACCTAGGAACTGGCCCAGAAGCTTCACCTC
CCTACGGCGCTAACAAAGAAGGCATCGTATGGGTTGCAACTGAGGGAGCCTTGAATACACCCAAAGACCAC
ATTGGCACCCGCAATCCTAATAACAATGCTGCCACCGTGCTACAACCTCCTCAAGGAACAACATTGCCAAA
AGGCCTTCTACGCAGAGGGAAGCAGAGGGCGGCAGTCAAGCCTCTTCTCGCTCCTCATCACGTAGTCGCGGTA
ATTCAAGAAATTCACCTCTGGCAGCAGTAGGGGAAATTCCTGCTCGAATGGCTAGCGGAGGTGGTGAA
ACTGCCCTCGCGCTATTGCTGCTAGACAGATTGAACCAGCTTGAGAGCAAAGTTTCTGGTAAAGGCCAACAA
ACAACAAGGCCAAACTGTCACCTAAGAAATCTGCTGCTGAGGCATCTAAAAAGCCTCGCCAAAAACGTACTG
CCACAAAACAGTACAACGTCACCTCAAGCATTTGGGAGACGTGGTCCAGAACAACCCAAAGGAAATTTCTGGG
GACCAAGACCTAATCAGACAAGGAACTGATTACAAACATTGGCCGCAAATTGCACAATTTGCTCCAAGTGC
CTCTGCATTTCTTGGAAATGTCACGCATTGGCATGGAAGTCACACCTTCGGGAACATGGCTGACTTATCATG
GAGCCATTAATTTGGATGACAAAGATCCACAATTCAAAGACAACGTCATACTGCTGAACAAGCACATTGAC
GCATACAAAACATTTCCACCACAGAGCCTAAAAAGGACAAAAAGAAAAGACTGATGAAGCTCAGCCTTT
GCCGAGAGACAAAAGAAGCAGCCCACTGTGACTCTTCTTCTCGCGCTGACATGGATGATTTCTCCAGAC
AACTTCAAAATTCATGAGTGGAGCTTCTGCTGATTCAACTCAGGCATAAACACTCATGATGACCACACAA
GGCAGATGGGCTATGTAAACGTTTTCGCAATTCGGTTTACGATACATAGTCTACTCTTGTGCAGAATGAAT
TCTCGTAACTAAACAGCACAAAGTAGGTTTAGTTAACTTTAATCTCACATAGCAATCTTTAATCAATGTGTA
ACATTAGGGAGGACTTGAAAGAGCCACCACATTTTCATCGAGGCCACGCGGAGTACGATCGAGGGTACAGT
GAATAATGCTAGGGAGAGCTGCCATATATGGAAGAGCCCTAATGTGTAAAATTAATTTTAGTAGTGCTATCC
CCATGTGATTTTAATAGCTTCTTAGGAGAATGACAAAAAAGAAAAAAGAAAAAAGAAAAAAGAAAAAAGAAAAA

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FIGURE 3H

CTACCCAGGAAAAGCCAACCAACCTCGATCTCTTGTAGATCTGTTCTCTAAACGAACTTTAAAATCTGTGT
AGCTGTCGCTCGGCTGCATGCCTAGTGCACCTACGCAGTATAAAACAATAATAAATTTACTGTCGTTGACA
AGAAACGAGTAACCTCGTCCCTCTTCTGCAGACTGCTTACGGTTTCGTCGCTGTTGCAGTCCAGTATCAGCA
TACCTAGGTTTTCGTCGGGTGTGACCGAAAGTAAAGTGGAGAGCCTTGTCTTGGTGTCAACGAGAAAAC
ACACGTCCAACTCAGTTTGCCTGTCTTACAGGTTAGAGACGTCTAGTGCCTGGCTTCGGGGACTCTGTGG
AAGAGGCCCTATCGGAGGCAGTGAACACCTCAAAAATGGCAGTCTGTGGTCTAGTAGAGCTGGAAAAGGC
GTACTGCCCCAGCTTGAACAGCCCTATGTGTTCAATTAACGTTCTGATGCCTTAAGCACAATCAGGCCA
CAAGGTCGTTGAGCTGGTTGCAGAAATGGACGGCATTAGTACGGTCGTAGCGGTATAACACTGGGAGTAC
TCGTGCCACATGTGGGCGAAACCCCAATTGCATACCGCAATGTTCTTCTTCGTAAGAACGGTAATAAGGGA
GCCGTTGGTTCATATGGCATCGATCTAAAGTCTTATGACTTAGGTGACGAGCTTGGCACTGATCCCAT
TGAAGATTATGAACAAAACGGAAACACTAAGCATGGCAGTGGTGCCTCCGTAACCTCACTCGTGCATCA
ATGGAGGTGCAGTCACTCGCTATGTGACACAACATTTCTGTGGCCAGATGGGTACCCTCTTGTATTGCATC
AAAGATTTTCTCGCACGCGGGCAAGTCAATGTGACTCTTCCGAACAACCTTGATTACATCGAGTCGAA
GAGAGGTGCTACTGCTGCCGTGACCATGAGCATGAAATTGCCCTGGTTCAGTGCAGCTCTGATAAGAGCT
ACGAGCACCAGACACCCTTCGAAAATAAGAGTGCACGAAATTTGACACTTTCAAAGGGGAATGCCCAAAG
TTTGTGTTTCTCTTAACCTAAAAGTCAAAGTCAATTAACCAGTGTGAAAAGAAAAGACTGAGGGTTT
CATGGGGCGTATACGCTCTGTGTACCCTGTGTCATCTCCACAGGAGTGAACAATATGCACCTGTCTACCT
TGATGAAATGTAATCATTGCGATGAAGTTTCAATGGCAGACGTGCGACTTTCTGAAAGCCACTTGTGAACAT
TGTGGCACTGAAAATTTAGTTATTGAAGGACCTACTACATGTGGGTACCTACCTACTAATGCTGTAGTGAA
AATGCCATGCTCTGCCTGTCAAGACCCAGAGATTGGACCTGAGCATAGTGTGTCAGATTATCACAACCCT
CAAACATTTGAACTCGACTCCGCAAGGGAGGTAGGACTAGATGTTTGGAGGCTGTGTGTTTGCCTATGTT
GGCTGTATAATAAGCGTGCTACTGGGTTCTCGTGCTAGTGTGATATTGGCTCAGGCCATACTGGCAT
TACTGGTGACAAATGGAGACCTTGAATGAGGATCTCTTGGAGTACCTGAGTGTGAACTGTTAACTTA
ACATTTGTTGGCGATTTTCAATTTGAATGAAGAGGTTGCCATCATTTTGGCATCTTCTCTGCTTACAAAGT
GCCTTTATTGACACTATAAAGAGTCTTGATTACAAGTCTTCAAACCATTGTTGAGTCTCGCGTAACCTA
TAAAGTTACCAAGGGAAGCCCGTAAAAGGTGCTTGGACATTTGGACAACAGAGATCAGTTTTAACACCAC
TGTGTGGTTTTTCCCTCACAGGCTGCTGGTGTATCAGATCAATTTTGGCGGCACACTTGATCGAGCAAAC
CACTCAATTCCTGATTTGCAAAGAGCAGCTGTCAACATACTTGATGGTATTTCTGAACAGTCAATACGCTCT
TGTCGACGCCATGGTTTATACCTCAGACCTGCTACCAACAGTGTCAATATTATGGCATATGTAACCTGGT
GCTTTGTACAACAGACTTCTCAGTGGTGTGCTAATCTTTTGGGCACTACTGTTGAAAACCTCAGGCCCTATC
TTTGAATGGATTGAGGCGAAACTTAGTGCAGGAGTTGAATTTCTCAAGGATGCTTGGGAGATTCTCAAATT
TCTCATACAGGTGTTTTTGCATCGTCAAGGGTCAAATACAGGTTGCTTCAGATAACATCAAGGATTGTG
TAAAATGCTTCATTGATGTTGTTAAACAAGGCACCTCGAAATGTGCATTGATCAAGTCACTATCGCTGGCGCA
AAGTTGCGATCACTCAACTTAGGTGAAGTCTTCATCGCTCAAAGCAAGGGACTTTACCGTCAGTGTATACG
TGCAAGGAGCAGCTGCAACTACTCATGCCCTTAAAGGCACCAAAGAAGTAACTTCTTGAAGGTGATT
CACATGACACAGTACTTACCTCTGAGGAGGTTGTTCTCAAGAACGGTGAACCTCGAAGCAGCTCGAGACCC
GTTGATAGCTTACAAATGGAGCTATCGTTGGCACACCAGTCTGTGTAATGGCCTCATGCTCTTAGAGAT
TAAGGACAAGAACAATACTGCGCATTTGCTCCTGGTTTACTGGCTACAACAATGTCTTTCCGTTAAAAG
GGGTGCACCAATTAAGGTGTAACCTTTGGAGAAGATACGTTTGGGAAGTTCAAGGTTACAAGAAATGTG
AGAATCACATTTGAGCTGATGAACGTGTTGACAAAGTGTAAATGAAAAGTGTCTGTCTACACTGTTGA
ATCCGGTACCGAAGTTACTGAGTTTGCATGTGTTGTAGCAGAGGCTGTTGTGAAGACTTTACAACCAGTTT
CTGATCTCCTTACCAACATGGGTATTGATCTGTGATGAGTGGAGTGTAGCTACATTTCTACTTATTTGATGAT
GCTGGTGAAGAAAACTTTTCATCACGTATGTATTGTTCTTTTACCCTCCAGATGAGGAAGAAGAGGACGA
TGCAGAGTGTGAGGAAGAAGAAATTTGATGAAACCTGTGAACATGAGTACGGTACAGAGGATGATTATCAAG
GTCTCCCTCTGGAATTTGGTGCCTCAGCTGAAACAGTTCGAGTTGAGGAAGAAGAGGGAAGACTGGCTG
GATGATACTACTGAGCAATCAGAGATTGAGCCAGAACCAGAACCTACACCTGAAGAACCAGTTAATCAGTT
TACTGGTTATTTAAAACCTACTGACAATGTGCCATTAATGTGTTGACATCGTTAAGGAGGCACAAAGTG
CTAATCCTATGGTGATTGTAATGTCTGCTAACAATACACTGAAACATGGTGGTGGTGTAGCAGTGCACCTC
AACAAGGCAACCAATGGTGCATGCAAAAGGAGAGTGAATGATTACATTAAGCTAAAATGGCCCTCTTACAGT
AGGAGGGTCTTGTGTTGCTTCTGACATAATCTTGCTAAGAAGTGTCTGCATGTTGTTGGACCTAACCTAA
ATGCAGGTGAGGACATCCAGCTTCTTAAGGCAGCATATGAAAATTTCAATTCACAGGACATCTTACTTGCA
CCATTGTTGTCAGCAGGCATATTTGGTGTCAAACCACTTCACTTTTACAAGTGTGCGTGCAGACGGTTCG
TACACAGGTTTATATTGCAGTCAATGACAAAGCTCTTTATGAGCAGGTTGTCAATGGATTATCTTGATAACC
TGAAGCTTAGAGTGAAGCACCATAACAAGAGGACCCACCAACACAGAAGATCCAAAACCTGAGGAGAAA
TCTATCCGTACAGAAGCCTGTGATGTGAAGCCAAAATAAGGCCATGATTGATGAGGTTACCACAACACT
GGAAGAACTAAGTTTCTTACCAATAAGTTACTCTTGTGTTGCTGATATCAATGGTAAGCTTTACCATGATT
CTCAGAACATGCTTAGAGGTGAAGATATGTCTTCTTGGAGAAGGATGCACCTTACATGGTAGGTGATGTT

FIGURE 3I

ATCACTAGTGGTGATATCACTTGTGTGTAATAACCCCTCCAAAAAGGCTGGTGGCACTACTGAGATGCTCTC
AAGAGCTTTGAAGAAAGTGCCAGTTGATGAGTATATAACCACGTACCCTGGACAAGGATGTGCTGGTTATA
CACTTGAGGAAGCTAAGACTGCTCTTAAGAAATGCAAATCTGCATTTTATGTACTACCTTCAGAAGCACCT
AATGCTAAGGAAGAGATCTAGGAACGTATCCTGGAATTTGAGAGAAATGCTTGCTCATGCTGAAGAGAC
AAGAAAATTAATGCCTATATGCATGGATGTAGAGCCATAATGGCAACCATCCAACGTAAGTATAAAGGAA
TTAAAATTC AAGAGGGCATCGTTGACTATGGTGTCCGATTCTCTTTTATACTAGTAAAGAGCCTGTAGCT
TCTATTATTACGAAGCTGAACTCTCTAAATGAGCCGCTTGTCCAAATGCCAATTGGTTATGTGACACATGG
TTTTAATCTTGAAGAGGCTGCGCGCTGTATGCGTTCCTTAAAGCTCCTGCCGTAGTGTCACTATCATCAC
CAGATGCTGTTACTACATATAATGGATACCTCACTTCGTATCAAAGACATCTGAGGAGCACTTTGTAGAA
ACAGTTTCTTTGGCTGGCTTTACAGAGATTGGTCTATTTCAGGACAGCGTACAGAGTTAGGTGTTGAATT
TCTTAAGCGTGGTGACAAAATTTGTGTACCACACTCTGGAGAGCCCGTCGAGTTTCATCTTGACGGTGAGG
TTCTTTCACTTGACAACTAAAGAGTCTCTTATCCCTGCGGGAGGTTAAGACTATAAAAGTGTTCACAACT
GTGGACAACACTAATCTCCACACACAGCTTGTGGATATGCTATGACATATGGACAGCAGTTTGGTCCAAC
ATACTTGGATGGTGTGATGTACAAAATTAACCTCATGTAATCATGAGGGTAAGACTTTCTTTGTAC
TACCTAGTGATGACACACTACGTAGTGAAGCTTTCGAGTACTACCATACTCTTGATGAGAGTTTTCTTGGT
AGGTACATGTCTGCTTTAAACCACACAAAGAAATGGAAATTTCCCAAGTTGGTGGTTAACTTCAATTA
ATGGGCTGATAAACAATTTGTTATTTGTCTAGTGGTTTATTAGCACTTCAACAGCTTGAAGTCAAATTAAG
CACCAGCACTTCAAGAGGCTTATTATAGAGCCCGTCTGGTGTGATGCTGCTAACTTTTGTGCACTCATACTC
GCTTACAGTAATAAACTGTTGGCGAGCTTGGTGTGTCAGAGAACTATGACCCATCTTCTACAGCATGC
TAATTTGGAATCTGCAAAGCGAGTCTTAAATGTGGTGTGTAACATTTGGTTCAGAAACTACTACTTAA
CGGGTGTAGAAGCTGTGATGTATATGGGTACTCTATCTTATGATAATCTTAAGACAGGTGTTTCCATTTCA
TGTGTGTGGTGTGATGCTACACAATATCTAGTACAACAAGAGTCTTCTTTTGTATGATGCTGCACC
ACCTGCTGAGTATAAATACAGCAAGGTACATCTTATGTGCGAATGAGTACACTGGTAACATACAGTGTG
GTCATTACACTCATATAACTGCTAAGGAGACCCTCTATCGTATTGACGGAGCTCACCTTACAAAGATGTCA
GAGTACAAAGGACCAGTACTGATGTTTTCTACAAGGAAACATCTTACACTACAACCATCAAGCCTGTGTC
GTATAAATCGATGGAGTTACTTACACAGAGATTGAACCAAAATTTGGATGGGTATTATAAAAAGGATAATG
CTTACTATACAGAGCAGCCTATAGACCTTGTACCAACTCAACCATTACCAAAATGCGAGTTTTGATAATTTCA
AAACTCACATGTTCTAACACAAAATTTGCTGATGATTTAAATCAAATGACAGGCTTCACAAAGCCAGCTTC
ACGAGAGCTATCTGTACATTTCTCCAGACTTGAATGGCGATGTAGTGGCTATTGACTATAGACACTATT
CAGCGAGTTCAAGAAAGGTGCTAAATTAATGCTGATAAGCAATTTGTTGGCACATTAACCAGGCTACAACC
AAGCAACGTTCAAAACAAACACTTGGTGTTCAGTGTCTTTGGAGTACAAAGCCAGTAGACTTCAAAA
TTCATTTGAAGTTCTGGCAGTAGAAGACACACAAGGAATGGACAATCTTGCCTTGTGAAAAGTCAACAACCCA
CCTCTGAAGAAGTAGTGGAAAATCTACCATACAGAAGGAAGTCATAGAGTGTGACGTGAAAACCTACCGAA
GTTGTAGGCAATGTCATACTTAAACCATCAGATGAAGGTGTTAAAGTAACACAAGAGTTAGGTCATGAGGA
TCTTATGGCTGCTTATGTGGAAAACACAAGCATTACCAATTAAGAAACCTAATGAGCTTTCACTAGCCTTAG
GTTTAAAAACAATGGCACTCATGGTATTGCTGCAATTAATAGTGTTCCTTGGAGTAAAATTTTGGCTTAT
GTCAAACCTTCTTAGGACAAGCAGCAATTAACAATCAAATTTGGCTAAGAGATTAGCAACAACGTTGCTG
TAACAATATATAGCCTTATGTGTTTACATTTATTGTTTCAATTTGGTACTTTTACTAAAAGTACCAATCTA
GAATTAGAGCTTCACTACCTACAACATTTGCTAAAAATAGTGTAAAGAGTGTGCTAAAATATGTTTGGAT
GCCGGCATTAAATATGTAAGTCAACCAAAATTTTCTAAATTTGTTACAATCGCTATGTGGCTATTGTTGTT
AAGTATTTGCTTAGGTTCTTAATCTGTGTAAGTGTGCTTTTGGTGTACTCTTATCTAATTTTGGTGTCTC
CTTCTTATTGTAATGGCGTTAGAGAATGTATCTTAATTCGTCTAACGTTACTACTATGGATTTCTGTGAA
GGTCTTTTCTTGCAGCATTTGTTAAGTGGATTAGACTCCCTTGATTCTTATCCAGCTCTTGAACCAT
TCAGGTGACGATTTTCATCGTACAAGCTAGACTTGAACAATTTAGGTCTGGCCGCTGAGTGGGTTTTGGCAT
ATATGTGTTTCAAAAATCTTTTATTTATTAGGCTTTTCAGCTATAATGACAGGTGTTCTTTGGCTATTTT
GCTAGTCAATTTTCATCAGCAATTTCTGGCTCATGTGGTTTTATCATTAGTATTGTACAAATGGCACCCGTTTC
TGCAATGGTTAGGATGTACATCTTCTTTGCTTCTTCTACTACATATGGAAGAGCTATGTTTATATCATGG
ATGGTTGCACCTCTTCGACTTGCATGATGTGCTATAAGCGCAATCGTGCCACACGCGTTGAGTGTACAAC
ATTTGTTAATGGCATGAAGAGATCTTTCTATGCTATGCAAAATGGAGCCGTGGCTTCTGCAAGACTCACAA
TTGGAATTTGCTCAATTTGTGACACATTTGCACTGGTAGTACATTCATTAGTGTGAGTTGCTCGTGTAT
GTTCACTCCAGTTTTAAAGACCAATCAACCTACTGACCAGTCACTGATATATTGTTGATGATGTTGTTG
AAAATGGCGCGCTTCACTTACTTTGACAAGCTGGTCAAAGACCTATGAGAGACATCCGCTCTCCCA
TTTTGTCAATTTAGACAATTTGAGAGCTAACAACACTAAAGGTTCACTGCCTATTAATGTCAATGTTTTG
ATGGCAAGTCCAAATGCGACGAGTCTGCTTCTAAGTCTGCTTCTGTGACTACAGTCAAGTGTGTTGTTG
CCTATTTCTGTTGCTTGGACAGCTCTTGTATCAGAGCTTGGAGATAGTACTGAAGTTTTCCGTTAAGATGTT
TGATGCTTATGTGACACCTTTTCAGCAACTTTTATGTTTCTTATGAAAAACTTAAGGCACCTTGTGTTGTA
CAGCTCACAGCGAGTTAGCAAAGGGTGTAGCTTTAGATGGTGTCTTTCTACATTCGTGTGAGCTGCCCGA

FIGURE 3J

CAAGGTGTTGTTGATACCGATGTTGACACAAAGGATGTTATTGAATGTCTCAAACCTTTCACATCACTCTGA
CTTAGAAGTGACAGGTGACAGTTGTAACAATTTTCATGCTCACCTATAATAAGGTTGAAAACATGACGCCCA
GAGATCTTGGCGCATGTATTGACTGTAATGCAAGGCATATCAATGCCAAGTAGCAAAAAGTCACAATGTT
TCACTCATCTGGGAATGTAAAAGACTACATGCTCTTATCTGAACAGCTGCGTAAACAATTCGTAGTGTGC
CAAGAAGAACAACATAACCTTTTAGACTAACTTGTGCTACAAC TAGACAGGTTGTCAATGTCATAACTACTA
AAATCTCACTCAAGGGTGGTAAGATTGTTAGTACTTGTTTTAAACTTATGCTTAAAGCCACATTATTGTGC
GTTCTTGCTGCATTGGTTTGTATATCGTTATGCCAGTACATACATTGTCATCCATGATGGTTACACAAA
TGAAATCATTGGTTACAAAGCCATTCAGGATGGTGTCACTCGTGACATCATTTCTACTGATGATTGTTTTG
CAAATAAACATGCTGGTTTTGACGCATGGTTTAGCCAGCGTGGTGGTTCATACAAAATGACAAAAGCTGC
CCTGTAGTAGCTGCTATCATTACAAGAGAGATTGGTTTCATAGTGCCCTGGCTTACCGGGTA CTGTGCTGAG
AGCAATCAATGGTGACTTCTTGCATTTTCTACCTCGTGTTTTTAGTGCTGTGGCAACATTTGCTACACAC
CTTCCAACTCATTGAGTATAGTATTTGCTACCTCGCTTGGTCTTGCTGCTGAGTGTACAATTTTT
AAGGATGCTATGGGCAAACCTGTGCCATATGTTATGACACTAATTTGCTAGAGGGTCTATTTCTTATAG
TGAGCTTCGTCCAGACACTCGTTATGTGCTTATGGATGGTTCCATCATAACAGTTTCCCTAACACTTACCTGG
AGGGTCTGTAGAGTAGTAACAACTTTTGATGCTGAGTACTGTAGACATGGTACATGCCGAAAGGTCAGAA
GTAGGTATTTGCCCTATCTACCAGTGGTAGATGGGTTCTTAATAATGAGCATTACAGAGCTCTATCAGGAGT
TTTTCTGTGGTGTGATGCGATGAATCTCATAGCTAACATCTTTACTCCTCTTGTGCAACCTGTGGGTGCTT
TAGATGTCTGCTTCAGTAGTGGCTGGTGGTATTATTGCCATATTGGTGACTTGTGCTGCCCTACTACTTT
ATGAAATTCAGACGTGTTTTGGTGAGTACAACCATGTTGTTGCTGCTAATGCACTTTTGTTTTGATGTC
TTTTACTATACTCTGTCTGGTACCAGCTTACAGCTTTCTGCCGGGAGTCTACTCAGTCTTTTACTTGTACT
TGACATTTCTATTTACCAATGATGTTTTCACTTTGGCTCACCTTCAATGGTTTGGCATGTTTTCTCCTATT
GTGCCTTTTTGGATAACAGCAATCTATGTATTCTGTATTTCTCTGAAGCACTGCCATTTGGTTCTTTAACAA
CTATCTTAGGAAAAGAGTCATGTTAATGGAGTTACATTTAGTACCTTCGAGGAGGCTGCTTTGTGTACCT
TTTTGCTCAACAAGGAAATGTACCTAAAATGCGTAGCGAGACACTGTTGCCACTTACACAGTATAACAGG
TATCTTGCTCTATATAACAAGTACAAGTATTTAGTGGAGCCTTAGATACTACCAGCTATCGTGAAGCAGC
TTGCTGCCACTTAGCAAAGGCTCTAAATGACTTTAGCAACTCAGGTGCTGATGTTCTCTACCAACCACCAC
AGACATCAATCACTTCTGCTGTTCTGCAGAGTGGTTTTAGGAAAATGGCATTCCCGTCAGGCAAAGTTGAA
GGGTAGCATGGTACAAGTAACCTGTGGAACATACTTAAATGGATGTGGTTGGATGACACAGTATACTG
TCCAAGACATGTCATTTGCACAGCAGAAGACATGCTTAACTTAACTATGAAGATCTGCTCATTCGCAAA
CCAACCATAGCTTTCTTGTTCAGGCTGGCAATGTTCAACTTCGTGTTATTGGCCATTCTATGCCAAAATGTT
CTGCTTAGGCTTAAAGTTGATACTTCTAACCTAAGACACCCCAAGTATAAAATTTGTCCGATCCAACCTGG
TCAAACATTTTTCAGTCTTAGCATGCTACAATGGTTCACCATCTGGTGTATTATCAGTGTGCCATGAGACCTA
ATCATAACCATTAAAGGTTCTTTCCCTAATGGATCATGTGGTAGTGTGGTTTTAACATTGATTATGATTGC
GTGTCTTTCTGCTATATGCATCATATGGAGCTTCCAACAGGAGTACACGCTGGTACTGACTTAGAAGGTAA
ATTCTATGGTCCATTTGTTGACAGACAAACTGCACAGGCTGCAGGTACAGACACAACCATAACATTAATG
TTTTGGCATGGCTGTATGCTGCTGTTATCAATGGTGATAGGTGGTTTTCTTAATAGATTACCACACTACTTTG
AATGACTTTAACCTTTGGCAATGAAGTACAACATGAACTTTGACACAAGATCATGTTGACATATGGG
ACCTTTTCTGCTCAAACAGGAATTCGGTCTTAGATATGTGTGCTGCTTTGAAAGAGCTGCTGACAGAATG
GTATGAATGGTGCCTATCCTTGGTAGCACTATTTTAGAAGATGAGTTTACACCATTTGATGTTGTTAGA
CAATGCTCTGGTGTACCTTCCAAGGTAAGTTCAAGAAAATGTTAAGGGCACTCATCATTGGATGCTTTTT
AACTTTCTTGACATCACTATTGATTTCTGTTCAAAGTACACAGTGGTCACTGTTTTTCTTTGTTTACGAGA
ATGCTTTCTTGCCATTTACTCTTGGTATTATGGCAATGCTGCATGTGCTATGCTGCTTGTTAAGCATAAG
CACGCATTTCTGTGCTTGTCTGTTACCTTCTCTTGAACAGTTGCTTACTTTAATATGGTCTACATGCC
TGCTAGCTGGGTGATGCGTATCATGACATGGCTTGAATGGCTGACACTAGCTTGTCTGGTTATAGGCTTA
AGGATGTGTTATGTATGCTTCAGCTTTAGTTTTGCTTATTCATGACAGCTCGCACTGTTTATGATGAT
GCTGCTAGACGTGTTTTGGACACTGATGAATGTCATTACACTTGTTTACAAAGTCTACTATGGTAATGCTTT
AGATCAAGCTATTTCCATGTGGGCTTAGTTATTTCTGTAACCTCTAACTATCTGGTGTGTTACGACTA
TCATGTTTTTAGCTAGAGCTATAGTGTGTTGTGTTGAGTATTACCCATTGTTATTTATTA CTGTTGCAAC
ACCTTACAGTGTATCATGCTTGTATTGTTTCTTAGGCTATTGTTGCTGCTGCTACTTTGGCCTTTTCTG
TTTACTCAACCGTTACTTCAGGCTTACTCTGGTGTATTGACTACTTGGTCTCTACACAAGAATTTAGGT
ATATGAACCTCCAGGGCTTTTGCCTCCTAAGAGTAGTATTGATGCTTTCAAGCTTAAACATTAAGTTGTTG
GGTATTGGAGGTAAACCATGTATCAAGGTTGCTACTGTACAGTCTAAAATGTCTGACGTAAAAGTGCACATC
AATCCACAATGATATTCTTCTTGCAAAAGACACAACAGTGAAGCTTTGAGAAGATGGTTTCTCTTTGCT
GTTTTGCTATCCACTCAGGGTGTGTAGACATTAATAGTTGTGCGAGGAAATGCTCGATAACGGTGCCTAC
TCTTCAGGCTATTGCTTCAAGATTTAGTTCTTTACCATCATATGCCGCTTATGCCACTGCCAGGAGCCCT
ATGAGCAGGCTGTAGCTAATGGTGATTTCTGAAGTCTTCTCAAAAAGTTAAAGAAATCTTTGAATGTGGCT

FIGURE 3K

AAATCTGAGTTTGACCGTGATGCTGCCATGCAACGCAAGTTGGAAAAGATGGCAGATCAGGCTATGACCCA
AATGTACAAACAGGCAAGATCTGAGGACAAGAGGGCAAAAAGTAACTAGTGCATGCAAAACATGCTCTTCA
CTATGCTTAGGAAGCTTGATAATGATGCACCTAACCAACATTATCAACAATGCGCGTGATGGTTGTGTTCCA
CTCAACATCATACCATTGACTACAGCAGCCAAACTATGGTTGTGTCCTGATATATGGTACCTACAAGAA
CACTTGTGATGGTAACACCTTTACATATGCATCTGCACTCTGGGAAATCCAGCAAGTTGTTGATGCGGATA
GCAAGATTGTTCAACTTAGTGAATTAACATGGACAATTCACCAAATTTGGCTTGGCCTCTTATTGTTACA
GCTCTAAGAGCCAACCTCAGCTGTTAAACTACAGAATAATGAACCTGAGTCCAGTAGCACTACGACAGATGTC
CTGTGCGGCTGGTACCACACAAACAGCTTGTACTGATGACAAATGCACTTGCCTACTATAACAATTCGAAGG
GAGGTAGGTTTGTGCTGGCATTACTATCAGACCACCAAGATCTCAAATGGGCTAGATTCCCTAAGAGTGAT
GGTACAGGTACAATTTACACAGAACTGGAACCACCTTGTAGGTTTGTACAGACACACCAAAGGGCCCTAA
AGTGAAATACCTTGTACTTCATCAAAGGCTTAAACAACCTAAATAGAGGTATGGTGTGCTGGGCAGTTTAGCTG
CTACAGTACGCTTTCAGGCTGGAAATGTACAGAAGTACCTGCCAATTCAACTGTGCTTTCCTTCTGTGCT
TTTGCAGTAGACCTTGCTAAAGCATATAAGGATTACCTAGCAAGTGGAGGACAACCAATCACCAACTGTGT
GAAGATGTTGTGACACACACTGGTACAGGACAGGCAATTACTGTAACACCAGAAGCTAACATGGACCAAG
AGTCCTTTGGTGGTGCCTTCATGTTGCTGTATGTTAGATGCCACATFGACCATCCAAATCCTAAAGGATTC
TGTGACTTGAAGGTAAGTACGTCCAAATACCTACCCTTGTGCTAATGACCCAGTGGGTTTTACACTTAG
AAACACAGTCTGTACCGTCTGCGGAATGTGAAAGGTTATGGCTGTAGTTGTGACCAACTCCGCGAACCTT
TGATCGAGTCTGCGGATGCATCAACGTTTTTAAACGGTTTTCGGTGTAAAGTGCAGCCCGCTTTACACCGT
GCGGCACAGGCACTAGTACTGATGTCGCTACAGGCTTTTGATATTTACAACGAAAAGTTGCTGGTTTT
GCAAAGTTCCTAAAACCTAATGCTGTGCTTCCAGGAGAAGGATGAGGAAGGCAATTTATTAGACTCTTA
CTTTGTAGTTAAGAGGCATACTATGCTTAACCTACCAACATGAAGAGACTATTTATAACTTGGTTAAAGATT
GTCCAGCGGTTGCTGTCCATGACTTTTTCAAGTTTAGAGTAGATGGTGACATGGTACCACATATATCACGT
CAGCGTCTAACTAAATACACAATGGCTGATTTAGTCTATGCTCTACGTCATTTTGTAGAGGGTAATTTGTGA
TACATTAAGAAAATACTCGTCACATACAATGCTGTGATGATGATTTTCAATAAGAAGGATTTGGTATG
ACTTCGTAGAGAATCCTGCATCTTACCGTATATGCTAACTTAGGTGAGCGGTGACGCCAATCATTATTA
AAGACTGTACAATTCGCGATGCTATGCGTGTGATGCGAGGATTTGTAGGCGTACTGACATTAGATAATCAGGA
TCTTAATGGGAACGGTACGATTTCCGGTATTTGCTACAAGTAGCACCAGGCTGCGGAGTTCCTATTGTGG
ATTCATATTACTCATGCTGATGCCATCCTCACTTTGACTAGGGCATTTGGCTGCTGAGTCCCATATGGAT
GCTGATCTCGCAAAACCCTTATTAAGTGGGATTTGCTGAAATATGATTTTACGGAAGAGAGACTTTGTCT
CTTCGACCGTTATTTAAATATTGGGACCAGACATACCATCCCAATTTGATTAACCTGTTGGATGATAGGT
GTATCCTTCATTGTGCAAACTTTAATGTGTTATTTCTACTGTGTTTTCCACCTACAAGTTTTGGACCATA
GTAAGAAAATATTTGTAGATGGTGTCTCTTTTGTGTTTCAACTGGATACCATTTTCGTGAGTTAGGAGT
CGTACATAATCAGGATGTAACCTTACATAGCTCGCGTCTCAGTTTCAAGGAACCTTTTAGTGTATGCTGCTG
ATCCAGCTATGCATGCAGCTTCTGCAATTTATGCTAGATAAACGCCTACATGCTTTTTCAGTAGCTGCA
CTAACAAACAAATGTGCTTTTCAACTGTCAAACCGGTAATTTAATAAAGACTTTTATGACTTTGCTGT
GTCATAAGGTTCTTTAAGGAAGGAGTTCTGTTGAACCTAAAACACTTCTTCTTTGCTCAGGATGGCAAG
CTGCTATCAGTGATGACTATTATCGTTATAATCTGCCAACAATGTGTGATATCAGACAACTCCTATTC
GTAGTTGAAGTTGTTGATAAATACTTTGATTGTTACGATGGTGGCTGTATTAATGCCAACCAAGTAATCGT
TAACAATCTGGATAAATCAGCTGGTTTCCCATTTAATAAATGGGGTAAGGCTAGACTTTATATGACTCAA
TGAGTTATGAGGATCAAGATGCACCTTTTCGCTATACTAAGCGTAATGTCACTCCCTACTATAACTCAAATG
AATCTTAAGTATGCCATTAGTGCAAGAATAGAGCTCGCACCGTAGCTGGTGTCTCTATCTGTAGTACTAT
GACAAAATAGACAGTTTTTCATCAGAAATTTATGAAGTCAATAGCCGCACTAGAGGAGCTACTGTGGTAATTG
GAACAAGCAAGTTTTACGGTGGCTGGCATAATATGTTAAAACTGTTTACAGTGATGTAGAACTCCACAC
CTTATGGGTTGGGATTTACCAAATGTGACAGAGCCATGCCFAACATGCTTAGGATAATGGCCTCTCTTGT
TCTTGTCTCGCAACATAACACTTGTGTAACCTTATCACACCGTTTTCTACAGGTTAGCTAACGAGTGTGCGC
AAGTATTAAGTGAGATGGTCAATGTTGCGGCTCACTATATGTTAAACCAGGTGGAACATCATCCGGTGTG
GCTACAACCTGCTTATGCTAATAGTGTCTTTAACATTTGTCAAGCTGTTACAGCCAATGTAATGCACTTCT
TTCAACTGATGGTAATAAGATAGCTGACAAGTATGTCGCAATCTACAACACAGGCTCTATGAGTGTCTCT
ATAGAAATAGGGATGTTGATCATGAATTCGTGGATGAGTTTTACGCTTACCTGCGTAAACATTTCTCCATG
ATGATTCCTTCTGATGATGCCGTTGTGTGCTATAACAGTAACATATGCGGCTCAAGGTTTAGTAGCTAGCAT
TAAGAACCTTAAGGCAGTCTTTTATTAATAAATAATGTGTTCATGCTGAGGCAAAATGTTGGACTGAGA
CTGACCTTACTAAAGGACCTCACGAATTTGCTCACAGCATAACATGCTAGTTAAACAAGGAGATGATTAC
GTGTACCTGCTTACCCAGATCCATCAAGAATATTAAGGCGCAGGCTGTTTTGTGCTGATGATATTGTCAAAC
AGATGGTACACTTATGATGAAAGGTTGCTGTCACTGGCTATTGATGCTTACCCACTTACAAAACATCCTA
ATCAGGAGTATGCTGATGCTTTTCACTGTATTTACAATACATTAGAAAAGTTACATGATGAGCTTACTGGC
CACATGTTGGACATGATTTCCGTAATGCTAACAATGATAACCTCACGGTACTGGGAACCTGAGTTTTA
TGAGGCTATGTACACACCACATACAGTCTTGCAGGCTGTAGGTGCTTGTGTATTGTGCAATTCACAGACTT

FIGURE 3L

CACTTCGTTGCGGTGCCTGTATTAGGAGACCATTCCATGTTGCAAGTGCTGCTATGACCATGTCATTTCA
ACATCACACAAATTAGTGTGCTGTTAATCCCTATGTTTGCATGCCCCAGGTTGTGATGTCACGTGATGT
GACACAACGTATCTAGGAGGTATGAGCTATTATTGCAAGTCACATAAGCCTCCCATTAGTTTTCCATTAT
GTGCTAATGGTCAGGTTTTGGTTTTATACAAAACACATGTGTAGGCAGTGACAATGTCACGTGACTTCAAT
CGGATAGCAACATGTGATGGACTAATGCTGGCGATTACATACTTGCCAACACTTGTACTGAGAGACTCAA
GCTTTTCGAGCAGAAACGCTCAAAGCCACTGAGGAAACATTTAAGCTGTCATATGGTATTGCCACTGTAC
GCGAAGTACTCTCTGACAGAGAATTGCATCTTTTCATGGGAGGTTGGAAAACCTAGACCACCATTGAACAGA
AACTATGCTTTACTGGTTACCGTGTAACTAAAAATAGTAAAGTACAGATTGGAGAGTACACCTTTGAAAA
AGGTGACTATGGTGTATGCTGTTGTGTACAGAGGTACTACGACATACAGTTGAATGTTGGTGTATTACTTTG
TGTGACATCTCACACTGTAATGCCACTTAGTGCACCTACTCTAGTGCCACAAGAGCACTATGTGAGAATT
ACTGGCTTGTACCCAACACTCAACATCTCAGATGAGTTTCTAGCAATGTTGCAAAATATCAAAAAGGTCGG
CATGCAAAAGTACTCTACACTCCAAGGACCCTGGTACTGGTAAGAGTCATTTTGCCATCGGACTTGCTC
TCTATTACCCATCTGCTCGCATAGTGTATACGGCATGCTCTCATGCAGCTGTTGATGCCCTATGTGAAAAG
GCATTAAAATATTTGCCCATAGATAAATGTAGTAGAATCATACCTGCCCGTGCAGCGCTAGAGTGTTTGA
TAAATTCAAAAGTGAATTCACACTAGAACAGTATGTTTCTGCACGTGAAAATGCATTTGCCAGAAAACCTG
CTGACATTGTAGTCTTTGATGAAATCTCTATGGCTACTAATATGACTTGAGTGTGTCATGCTAGACTT
CGTGCAAAACACTACGCTCTATATTGGCGATCCTGCTCAATTACCAGCCCCCGCACATTGCTGACTAAAAGG
CACACTAGAACCAGAATATTTAATTCAGTGTGCAGACTTATGAAAACAATAGGTCCAGACATGTTCCCTG
GAACTTGTGCGGTTGTCTGCTGAAATTTGTGACACTGTGAGTGTCTTAGTTTATGACAATAAGCTAAAA
GCACACAAGGATAAGTCAGCTCAATGCTTCAAATGTTCTACAAAGGTTTATTACACATGATGTTTCATC
TGCAATCAACAGACCTCAAATAGGCGTTGTAAGAGAATTTCTTACACGCAATCCTGCTTGGAGAAAAGCTG
TTTTTATCTCACCTTATAATTCACAGAACGCTGTAGCTTCAAATCTTAGGATTGCCACGACAGCTGTT
GATTCATCACAGGTTCTGAAATAGACTATGTCATATTCACACAAACTACTGAAAACAGCACACTCTTGTA
TGTCACCCGTTCAATGTGGCTATFACAAGGGCAAAAATTTGGCATTTTGTGCATAATGTCGTGATAGAGATC
TTTTATGACAACTGCAATTTACAAGTCTAGAAAATACCAGCTCGCAATGTGGCTACATTACAAGCAGAAAAT
GTAACCTGGACTTTTAAAGACTGTAGTAAGATCATTACTGGTCTTCATCCTACACAGGCACCTACACACCT
CAGCGTTGATATAAAGTTCAAGACTGAAGGATATGTGTTGACATACCAGGCATACCAAAGGACATGACCT
ACCGTAGACTCATCTCTATGATGGGTTTCAAATGAATACCAGTCAATGGTACCCTAATATGTTTATC
ACCCGCAAGAAGCTATTTCGTCACGTTTCGTGCGTGGATTGGCTTTGATGTAGAGGGCTGTCTAGCAACTAG
AGATGCTGTGGGTACTAACCTACCCTCCAGCTAGGATTTTCTACAGGTGTTAACTTAGTAGCTGTACCGA
CTGGTTATGTTGACTGAAAATAACACAGAATTCACCAGAGTTAATGCAAAACCTCCACCAGGTGACCAG
TTTAAACATCTTATACCCTCATGTATAAAGGCTTGCCCTGGAATGTAGTGCCTATTAAGATAGTACAAAT
GCTCAGTGATACACTGAAAGGATGTFCAGACAGAGTCTGTTGCTCCTTTGGGCGCATGGCTTTGAGCTTA
CATCAATGAAGTACTTTGTCAAGATGGACCTGAAGAACGCTGTTGTCTGTGTGACAAACGTCGCAACTGC
TTTTCTACTTCATCAGATACTTATGCCTGCTGGAATCATTCTGTGGGTTTTGACTATGTCTATAACCCATT
TATGATGATGTTTACAGCTGGGGCTTTACGGGTACCTTCAGAGTAACCATGACCAACATTGCCAGGTAC
ATGGAAATGCACATGTGGCTAGTTGTGATGCTATCATGACTAGATGTTTAGCAGTCCATGAGTCTTTGTT
AAGCGGTTGATTGGTCTGTTGAATACCCTATTATAGGAGATGAACTGAGGGTTAATTTCTGCTTGACAGAA
AGTACAACACATGGTTGTGAAGTCTGCATTGCTTGATAAGTTCCAGTTCTTCATGACATTTGAAATC
CAAAGGCTATCAAGTGTGTGCCTCAGGCTGAAGTAGAATGGAAGTTCTACGATGCTCAGCCATGTAGTGAC
AAAGCTTACAAAATAGAGGAACCTTCTTATTCTTATGCTACACATCAGATAAAATCACTGATGGTGTGTTG
TTTTTTTTGGAATTTGTAACGTTGATCGTTACCCAGCCAATGCAATTTGTGTGTAGGTTTGACACAAGAGTCT
TGTCAAACTTGAACTTACCAGGCTGTGATGGTGGTAGTTGTATGTGAATAAGCATGCATTTCCACACTCCA
GCTTTTCGATAAAAGTGCATTTACTAATTTAAAGCAATTCGCTTTCTTTTACTATTCTGATAGTCTTTGTGA
GTCTCATGGCAAAACAAGTAGTGTCCGATATTGATTATGTTCCACTCAAATCTGCTACGTTGATTACACGAT
GCAATTTAGGTGGTGTGTTTGACAGACACCATGCAATGAGTACCGACAGTACTTTGGATGCATATAATATG
ATGATTTCTGCTGGATTTAGCCTATGGATTTACAACAATTTGATACTTATAACCTGTGGAATACATTTAC
CAGGTTACAGAGTTTAGAAAATGTGGCTTATAATGTTGTTAATAAAGGACACTTTGATGGACACGCCGGCG
AAGCACCTGTTTCCATCATTAATAATGCTGTTTACACAAAGGTAGATGGTATTGATGTGGAGATCTTTGAA
AATAAGACAACACTTCTGTTAATGTTGCATTTGAGCTTTGGGCTAAGCGTAACATTAACCAAGTGCAG
GATTAAGATACTCAATAAATTTGGGTTGATATCGCTGCTAATACTGTAATCTGGGACTACAAAAGAGAAG
CCCCAGCACATGTATCTACAATAGGTGCTGCACAATGACTGACATTTGCCAAGAAACCTACTGAGAGTGTCT
TGTTCTTCACTTACTGTCTGTGTTGATGGTAGAGTGGAAAGGACAGGTAGACCTTTTTAGAAAACGCCGTAA
TGGTGTTTTAAATAACAGAAGGTTTCAGTCAAAGGCTTAAACACTTCAAAGGGACCAGCAAGCTAGCGTCA
ATGGAGTCACATTAATGGAGAATCAGTAAAAACAGTTTAACTACTTTAAGAAAGTAGACGGCATTATT
CAACAGTTGCCGTGAAACCTACTTTACTCAGACAGAGACTTAGAGGATTTTAAAGCCAGATCACAATGGA
AACTGACTTTCTCGAGCTCGCTATGGATGAATTCATACAGGATATAAGCTCGAGGGCTATGCCTTCGAAAC

FIGURE 3M

ACATCGTTTATGGAGATTTAGTATGACAACTTGGCGGTCTTCATTTAATGATAGGCTTAGCCAAGCGC
TCACAAGATTCACCACCTTAAATAGAGGATTTATCCCTATGGACAGCACAGTGAAAAATTACTTCATAAC
AGATGCGCAAAACAGGTTTATCAAAATGTGTGTGTTCTGTGATTGATCTTTACTTGATGACTTTGTGCGAGA
TAATAAAGTCACAAGATTTGTGAGTATTTCAAAGTGTGCAAGGTTACAATTGACTATGCTGAAATTTCA
TTCATGCTTTGGTGTAGGATGGACATGTTGAAACCTTCTACCCAAAACACTACAAGCAAGTCAAGCGTGGCA
ACCAGGTGTGCGATGCCTAACTTGTACAAGATGCAAGAATGCTTCTTGAAAAGTGTGACCTTCAGAATT
ATGGTGAAAAATGCTGTATACCAAAAAGGAATAATGATGATGTGCGCAAGTATACTCAACTGTGTCAATAC
TTAAATACACTTACTTTAGCTGTACCCACAAATGAGAGTTATTCACCTTTGGTGTGCTGCTGATAAAGG
AGTTGCACCAGGTACAGTGTGCTCAGACAATGGTTGCCAACTGGCACACTACTTGTGATTGATGATCTTA
ATGACTTCGCTCCGACCGAGATTTACTTTAATGGAGACTGTGCAACAGTACATACGGCTAATAAATGG
GACCTTATTTATAGCGATATGTATGACCCTAGGACCAAAACATGTGACAAAAGAGAATGACTCTAAAGAAGG
GTTTTTCACTTATCTGTGTGGATTTATAAAGCAAAAACACTAGCCCTGGTGGTTCATAGCTGTAAAGATAA
CAGAGCATTTCTGGAATGCTGACCTTTACAAGCTTATGGGCCATTTCTCATGGTGGACAGCTTTTGTTACA
AATGTAAATGCATCATATCGGAAGCATTTTTAATTTGGGGTAACTATCTTGGCAAGCCGAGGAACAAT
TGATGGCTATACCATGCATGCTAATACATTTTCTGAGGAACACAAATCCTATCCAGTTGTCTTCCATTT
CACTCTTTGACATGAGCAAAATTTCTCTTAAATTAAGAGGAAGTGTGTAATGTCTCTTAAGGAGAATCAA
ATCAATGATATGATTTATCTCTTCTGGAAAAAGGTAGGCTTATCATTAGAGAAAACAACAGAGTTGTGGT
TTCAAGTGATATTTCTGTAAACAATAACGAACATGTTTATTTCTTATTTCTTACTCTCACTAGTG
GTAGTGACCTTGACCGGTGACCCACTTTTGTGATGATGTTCAAGCTCCTAATACACTCAACATCTTCACTCT
ATGAGGGGGGTTTACTATCTGATGAAATTTTTAGATCAGACACTCTTTATTTAACTCAGGATTTATTTCT
TCCATTTTATCTAATGTTACAGGGTTTACATATTAATCATACGTTTGGCAACCTGTATACCTTTTA
AGGATGGTATTTATTTTGTGCTGACAGAAAATCAAATGTTGCTGCTGGTGGGTTTTGGTTCACCATG
AACAAACAGTCACAGTCGGTGTATTATTAACAATTTACTAATGTTGTTATACGAGCATGTAACCTTTGA
ATTGTTGACAAACCTTTCTTGTGCTTCTTAAACCCATGGGTACACAGACACATATGATATTCGATA
ATGCATTTAATTCGACTTTCGAGTACATATCTGATGCTTTTTCGCTTGTGTTTTCAGAAAAGTCAGSTAAT
TTTAAACACTTACGAGAGTTGTGTTTAAAAATAAAGATGGGTTTCTCTATGTTTATAAGGGCTATCAACC
TATAGATGTAGTTTTCGTGATCTACCTTCTGGTTTTAACACTTTGAAACCTATTTTTAAGTTGCTCTGTA
TTAACATTACAAATTTAGAGCCATTTTACAGCCTTTTCACCTGCTCAAGACATTTGGGGCACGTCAGCT
GCAGCTATTTTGTGGCTATTTAAAGCCAACACTACATTTATGCTCAAGTATGATGAAAATGGTACAAATCAC
AGATGCTGTTGATTTCTCAAATCCACTTGTGTAACCTAAATGCTCTGTTAAGAGCTTTGAGATTTGACA
AAGGAATTTACCAGACCTTAATTTCCAGGTTGTCCCTCAGGAGATGTTGTGAGATTCCTAATATTTACA
AAGTTGTCTCTTTTGGAGAGGTTTTTAAATGCTACTTAAATCCCTTCTGCTATGATGGAGAGAAAAA
AATTTCTAATTTGTTGCTGATTAATCTGCTCTACAACCTCAACATTTTTTTCAACCTTTAAGTGTATG
GCGTTTCTGCCACTAAGTTGAATGATCTTTGCTTCTCAATGCTATGACAGATTTCTTTTGTAGTCAAGGA
GATGATGTAAGACAAATAGCGCCAGGACAAACTGGTGTATTGCTGATTATAAATATAAATTTGCCAGATGA
TTTCATGGGTTGTCTCTTGGTAACTAGGAACATTTGATGCTACTTCAACTGGTAATTAATAATATA
AATAAGGATCTTTAGACATGGCAAGCTTAGGCCCTTTGAGAGAGACATATCTAATGTGCTTTCTCCCT
GATGGCAAACTTGCACCCACCTGCTCTTAATTTGTTATTTGGCCATTAATGATTATGGTTTTTACACCAC
TACTGGCATTGGCTACCAACCTTACAGAGTTGTAGTACTTTCTTTTGAACTTTTAAATGCACCGGCCACGG
TTTGTGGACCAAAATTTACCACTGACCTTATTAAGAACCAGTGTGTCATTTTAAATTTTAAATGGACTCACT
GGTACTGGTGTGTTAATCTCTTCAAAGAGATTTCAACATTTCAACAATTTGGCCGTGATGTTTCTGTA
TTTTACTGATTTCCGTTTCGAGATCCTAAAACATCTGAATATTTAGACATTTACCTTGCCTTTTGGGGTG
TAAGTGTAAATACACCTGGAACAAATGCTTCATCTGAAGTTGCTGTTCTATATCAAGATGTTAACTGCCT
GATGTTTCTACAGCAATTCATGAGATCAACTCACACCAGCTTGGCGCATATATCTACTGGAACAATGT
ATTCAGACTCAAGCAGGCTGTCTTATAGGAGCTGAGCATGTGACACTTCTTATGAGTGGACATTTCTTA
TTGGAGCTGGCATTGTGCTAGTTACCATAACAGTTTCTTATTACCTAGTACTAGCCAAAAATCTATTGTG
CTTATACTATGCTTTAGGTGCTGATAGTTCAATTTAGCTTACTCTAATAACACCATTGCTATACCTACTAA
CTTTTCAATTAGCATTACTACAGAAGTAATGCCTGTTTCTATGGCTAAAACCTCCGTAGATTGTAATATGT
ACATCTGCGGAGATTTACTGAAATGTGCTAATTTGCTTCTCAATATGGTAGCTTTTGCACACAACATAAAT
CGTGCCTCTCAGGTATTTGCTGTGAACAGGATCGCAACACAGTGAAGTGTTCGCTCAAGTCAAAACAAT
GTACAAAACCCCAACTTTGAAATATTTGGTGGTTTTAATTTTTCACAAATATTACCTGACCTCTAAAGC
CAACTAAGAGGCTTTTATTTAGGACTTGTCTTTAATTAAGGTGACACTCGCTGATGCTGGCTTCATGAAG
CAATATGGCGAATGCCATAGGTGATATTAATGCTAGAGATCTCATTTGTGCGCAGAAGTTCAATGGACTTAC
AGTGTGGCACCTCTGCTCACTGATGATGATGCTGCTACACTGCTGCTCTAGTTAGTGGTACTGCCA
CTGCTGGATGGACATTTGGTGTGCGCTGCTCTTCAAATACCTTTTGTCTATGCAAAATGGCATATAGGTTT
AATGGCATTTGAGTTACCCAAAATGTTCTCTATGAGAACCAAAAACAATCGCCAACCAATTTAAACAAGG
GATTAGTCAAATTAAGAATCACTTACAACAACATCAACTGCATTGGGCAAGCTGCAAGACGTTGTTAACC

FIGURE 3N

AGAATGCTCAAGCATTAACACACTTGTAAACAACCTTAGCTCTAATTTGGTGCAATTC AAGTGTGCTA
AATGATATCCCTTTCGCGACTTGATAAAGTCGAGGCGGAGGTACAAATGACAGGTTAATTACAGGCAGACT
TCAAAGCCTTCAAACCTATGTAACACAACACTAATCAGGGCTGCTGAAATCAGGGCTTCTGCTAATCTTG
CTGCTACTAAAATGTCTGAGTGTGTCTTTGGACAATCAAAAAGAGTTGACTTTTGTGGAAAGGGCTACCAC
CTTATGTCTTCCACAAGCAGCCCGCATGGTGTCTTCTTACATGTCACGTATGTGCCATCCAGGA
GAGGAACCTCACCACAGCGCCAGCAATTTGTCATGAAGGCAAGCATACTTCCCTCGTGAAGGTGTTTTG
TGTTTAAATGGCACTTCTTGGTTTATACACAGAGGAACCTTCTTCTCCACAAAATAATTACTACAGACAAT
ACATTTGTCTCAGGAAATTTGTGATGTCGTTATTTGGCATCATTAACAACACAGTTTATGATCCTCTGCAACC
TGAGCTTGACTCATTCAAAGAAGAGCTGGACAAGTACTTCAAAAATCATACATCACCAGATGTTGATCTTG
GCGACATTTCAGGCATTAACGCTTCTGTCTCAACATTCAAAAGAAATFGACCGCTCAATGAGGTCGCT
AAAATTTAAATGAATCACTCATTGACCTTCAAGAAATGGGAAAATATGAGCAATATATTAATGAGCTTG
GTATGTTTGGCTCGGCTTCAATGCTGGACTAATGCCATCGTCATGGTTACAATCTGCTTTGTTGTCATGA
CTAGTTGTTGCGAGTTGCCCTCAAGGGTGCATGCTCTTGTGGTTCTTGCTGCAAGTTTGTATGAGGATGACTCT
GAGCCAGTTCTCAAGGGTGTCAAATTCATTAACATTAACAACTTATGGATTGTTTATGAGATTTTTT
ACTCTTAGATCAATTAAGTGCACAGCCAGTAAAATTTGACAATGCTTCTTCCAGTCAAGTACTGTTACTAC
AGCAACGATACCGCTACAAGCCTCACTCCCTTTCGGATGGCTTGTATTTGGCGTTGCATTTCTGCTGTTT
TTCAGAGCGTACCAAATAAATFGCGTCAATAAAGATGGCAGCTAGCCCTTATAAGGGCTTCCAGTTC
ATTTGCAATTTACTGCTGCTATTTGTTACCATCTATTACATCTTTTGGCTTGTCTGCTGCAGGTATGGAGGC
GCAATTTTGTACTCTATGCTTGTATATATTTCTACAATGCATCAACGCATGTAGAATTAATATGAGAT
GTTTGGCTTTGTTGGAAGTGAATCCAGAACCCATTACTTTATGATGCCAACTACTTTGTTGCTGGCAC
ACACATAACTATGACTACTGTATACCATATAACAGTGTACAGATACAATTTGCTGTTACTGAAGGTGACGG
CATTTCAACACCAAACTCAAAGAAGACTACCAAATTTGGTGGTTATTTCTGAGGATAGGCACCTCAGGTGTTA
AAGACTATGTCGTTGTACATGGCTATTTCAACGAAGTTTACTACCAGCTGAGTCTACACAAATTAATACACA
GACACTGGTATTTGAAAATGCTACATCTTCAATTTTAAACAAGCTTGTAAAGACCCACCGAATGTCAAAAT
ACACACAATCGACGGCTCTTCAAGGAGTTGCTAATCCAGCAATGGATCCAATTTATGATGAGCCGACGACGA
CTACTAGCGTGCCTTTGTAAGCACAAGAAAGTGAAGTACGAACTTATGTACTCATTCGTTTCGGAAAGAAACA
GGTACGTTAATAGTTAATAGCGTACTTCTTTTCTTGTCTTTCGTTGGTATTTCTGCTAGTCACACTAGCCAT
CCTTACTGCGCTTTCGATTTGTTGCTGCTACTGCTGCAATAATGTTAAACGTTGAGTTTATGTAACCAACCGT
ACGCTACTCGCGTGTAAAAATCTGAACTTCTTGAAGGAGTTCTGATCTTCTGGTCTAAACGAACATAA
CTATTATATATTTCTGTTTGGAACTTTAACATTTGCTTATCATGGCAGACAACGGTACTATTACCGTTGAG
GAGCTTAAACAACCTCTGGAACAATGGAACCTAGTAATAGGTTTCCATTTCTAGCCTGGATTATGTTACT
ACAACAGATGTTTTCATCTTGTGACTTCCAGGTTACAATAGCAGAGATATTGATATCATATTATGAGGACTT
CAGTAACACTTGTCTGTTTGTGCTTGTCTGTCTACAGAATTAATTTGGGTGACTGGCGGGATTGCGATT
GCAATGGCTTGTATTGTAGGCTTGTATGCTTACTTTCGTTGCTTCCCTCAGGCCTGTTTCTGCTGCTAC
CCGCTCAATGTGGTCATTCACCCAGAAACAACATTTCTTCAATGTGCCCTTCCGGGGGACAATTTGTGA
CCAGACGCTCATGGAAAGTGAACCTTGTCAATGGTGTGATCATTCGTTGCTCACTGCGAATGCGAATGCGGA
CACTCCCTAGGGCGCTGTGACATTAAGGACCTGCCAAAAGAGATCACTGTGGCTACATCAGAACCGCTTTC
TTATTACAAATTAGGAGCGTGCACGCTGTAGGCCTGATTACAGGTTTGTGCTGCATACAACCGCTACCGTA
TTGGAACCTATAAATTAATACAGACCACCGGTTAGCAACGACAATATTGCTTTGCTAGTACAGTAAGT
ACAACAGATGTTTTCATCTTGTGACTTCCAGGTTACAATAGCAGAGATATTGATATCATATTATGAGGACTT
TCAGGATTGCTATTTGGAATCTTGGCTTATAAATAGTTCAATAGTGAAGACAATTTTAAAGCCTCTAAT
AAGAAGAATTTTCGGAGTTAGATGATGAAGAACCATGGAGTTAGATTATCCATAAAAACGAACATGAAAA
TTATTTCTCTTCCGACATTTGATTTGATTTTACATCTTGGAGCTATATCACTATCAGGAGTGTGTTAGAGGT
ACGACTGTACTACTAAAAGAACCCTTGCCTCAGGAACATACGAGGGCAATTCACCATTTCAACCTCTTGC
TGACAATAAATTTGCACTAATTTGCACTAGCACACTTTGCTTTTGTGCTGCTGACGGTACTCGACATA
CCTATCAGCTGCGTGAAGATCAGTTTCCACAAAACCTTTTCAATCAGACAAGAGGAGGTTCAACAAGAGCTC
TACTCGCCACTTTTCTCATTTGTTGCTGCTAGTATTTTTAATACTTTGCTTACCATTAAAGAGAAAGAC
AGAATGAATGAGCTCACTTTAATGACTTCTATTTGTGCTTTTGTAGCTTCTGCTATTCTTGTGTTTAAAT
AATGCTTATATATTTTGGTTTTCATCTCGAAATCCAGGATCTAGAAGAACCTTGTACCAAAGTCTAAACGA
ACATGAAACTTCTCATTTGTTGACTTGTATTTCTCTATGCAGTTGCATATGCACTGTAGTACAGCGCTGT
GCATCTAATAAACCTCATGTGCTTGAAGATCCTTGTAAAGGTACAACACTAGGGGTAATACTTATAGCACTG
CTTTGGCTTTGTGCTCTAGGAAAGGTTTACCTTTTCAATAGATGGCACACTATGGTTCAACATGCACACCT
AATGTTACTATCAACTGTCAAGATCCAGTGGTGGTGGCTTATAGTACTAGGTTGGTACCTTATGAAGG
TCACCAACTGCTGCATTTAGAGACGTAATTTGTTTAAATAAACGAACAATTAATATGCTGATAAT
GGACCCCAATCAAACAACGTAAGTGCCTCCCGCATTACATTTGGTGGACCCACAGATTCACCTGACAATAA
CCAGAATGGAGGACGCAATGGGGCAAGGCCAAAACAGCGCCGACCCCAAGGTTTACCAATAATACTGCGT
CTTGGTTACAGCTCTCACTCAGCATGGCAAGGAGAACTTAGATTTCCCTCGAGGCGAGGGCTTCCAATC

FIGURE 30

AACACCAATAGTGGTCCAGATGACCAAAATTGGCTACTACCGAAGAGCTACCCGACGAGTTCGTGGTGGTGA
CGGCAAAATGAAAGAGCTCAGCCCCAGATGGTACTTCTATTACCTAGGAACTGGCCCAGAAGCTTCACTTC
CCTACGGCGCTAACAAAGAAGGCATCGTATGGGTGCAACTGAGGGAGCCTTGAATACACCCAAAGACCAC
ATTGGCACC CGCAATCCTAATAACAATGCTGCCACCGTGCTFACAACCTCCTCAAGGAACAACATTGCCAAA
AGGCTTCTACGCAGAGGGGAAGCAGAGGGCGGCACTCAAGCCTCTTCTCGCTCCTCATCACGTAGTCGCGGTA
ATTCAAGAAATTCAACTCCTGGCAGCAGTAGGGGAAATTCCTCTGCTCGAATGGCTAGCGGAGGTGGTGA
ACTGCCCTCGCGCTATTGCTGCTAGACAGATTGAACCAGCTTGAGAGCAAAGTTTCTGGTAAAGGCCAACA
ACAACAAGGCCAAACTGTACTAAGAAATCTGCTGCTGAGGCATCTAAAAAGCCTCGCCAAAACGTA
CCACAAAACAGTACAACGTCCTCAAGCATTTGGGAGACGTGGTCCAGAACAACCCAAAGGAAATTTCCGGG
GACCAAGACCTAATCAGACAAGGAACGTATPACAAACATTTGGCCGCAAATGACACAATTTGCTCCAAGTGC
CTCTGCATTCTTTGGAATGTCACGCATTGGCATGGAAGTCAACCTTCGGGAACATGGCTGACTTATCATG
GAGCCATTAAATTTGGATGACAAAAGATCCACAATTCAAAGACAACGTCATACGCTGAACAAGCACATTGAC
GCATACAAAACATTCCCACCAACAGAGCCTAAAAAGGACAAAAAGAAAAGACTGATGAAGCTCAGCCTTT
GCCGAGAGACAAAAGAAGCAGCCACTGTGACTCTTCTTCTGCGGCTGACATGGATGATTTCTCCAGAC
AACTTCAAAATTCATGAGTGGAGCTTCTGCTGATTCAACTCAGGCATAAACACTCATGATGACCACACAA
GGCAGATGGGCTATGTAAACGTTTTTCGCAATTCGGTTTACGATACATAGTCTACTCTTGTGCAGAATGAAT
TCTCGTAACTAAACAGCACAAAGTAGGTTTAGTTAACTTTAATCTCACATAGCAATCTTTAATCAATGTGTA
ACATTAGGGAGGACTTGAAAGAGCCACCACATTTTCATCGAGGCCACGCGGAGTACGATCGAGGGTACAGT
GAATAATGCTAGGGAGAGCTGCCTATATGGAAGAGCCCTAATGTGTAATAATTAATTTTAGTAGTGCTATCC
CCATGTGATTTTAATAGCTTCTTAGGAGAATGACAAAAA
AAAAAAAAAAAAAAAAAAAAAAAAA

GenBank Accession No. AY274119.2.; SEQ ID NO: 2

FIGURE 3P

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ERV-2 -----
TOR2  ACACTCATGATGACCACACAAGGCAGATGGGCTATGTAACGTTTTCGCAATCCGTTTA
AIBV  -----

ERV-2 -----
TOR2  CGATACATAGTCTACTCTTGTGCAGAAATGAAATTCGTAACAAACAGCACAGTAGGTT
AIBV  -----

ERV-2 -----ACCCGTTACCCATAAAATCCCTCC
TOR2  TAGTTAACTTTAATCTCACATAGCAATCTTTAATCAATGTTAATACATTAGGGAGGACTTG
AIBV  -----TAGTTAGTTTAAAGTTAGTTAG
                * * * * *

ERV-2 CCTTCTCTTCAC-----TCGCCGAGGCCACGCCGAGTAGGACCGAGGGTACAGC----
TOR2  AAAGAGCCACCACATTT--TCATCGAGGCCACGCCGAGTACGATCGAGGGTACAGT----
AIBV  AGTAGGTATAAAGATGCCAGTGCCGGGGCCACGCCGAGTACGATCGAGGGTACAGCACTA
                *                ** ***** ** * *****

ERV-2 -GAGTCTTT-TAGTTTAAAGTGT-TAGATGTAAGGTACGTGGGCTTTCT--TTTGGTTTA
TOR2  -GAATAATGCTAGGGAGAGCTGCCATATGGAAGAGCCCTAATGTTAAAATTAATTTTA
AIBV  GGACGCCCATTAGGGGAAGA-GCTAAATTTAGTTTAAAGTTAAGTTAA--TTGGCTAA
                **      ***      ** * * * * * * * * * * * * * * *

ERV-2 CTTCTTC----- GenBank: AF361253 (SEQ ID NO: 31)
TOR2  GTAGTGCTATCCCATGTGATTTTAAATAGCTTCTTAGGAGAATGAC (SEQ ID NO: 18)
AIBV  GTATAGTTAAAATTTATAGGCTAGTATAGAGTTAGAGCA----- GenBank: NC_001451 (SEQ ID NO: 32)
                *
```

Figure 4

MFIFLLFLTLTSGSDLDRCCTTFDDVQAPNYTQHTSSMRGVVYPDEIFRSD
TLYLTQDLFLPFYSNVTGFHTINHTFGNPVIFPKDGIYFAATEKSNVVRG
WVFGSTMNKSQSVIIINNSTNVVIRACNFELCDNPFPAVSKPMGTQHT
MIFDNAFNCTFEYISDAFSLDVSEKSGNFKHLREFVFKNKDGFYVYKGY
QPIDVVRDLPSGFNTLKPFIKFLPLGINITNFRAILTAFSPAQDIWGTSA
AYFVGYLKPTTFMLKYDENGTTITDAVDCSQNPLAELKCSVKSFEIDKGIY
QTSNFRVVPSPGDVVRFPNITNLCPFGEVFNATKFPVYAWERKKISNCVA
DYSVLYNSTFFSTFKCYGVSATKLNLDLCSNVYADSFVVKGDDVRQIAPG
QTGVIADYNYKLPDDFMGCVLAWNTRNIDATSTGNYNKYRYLRHGKLRP
FERDISNVPFSPDGKPCPPALNCYWPLNDYGFYTTTGIGYQPYRVVLS
FELLNAPATVCGPKLSTDLIKNQCWNFNFNGLTGTGVLTPSSKRFQPFQO
FGRDVSDF'TDSVRDPKTSEILDISPCAFGGVSVITPGTNASSEVAVLYQD
VNCTDVSTAIHADQLTPAWRIYSTGNNVFQTOAGCLIGAEHVDTSYECDI
PIGAGICASYHTVSLLRSTSQKSIVAYTMSLGADSSIAYSNNTIAIPTNF
SISITTEVMPVSMAKTSVDCNMYICGDSTECANLLLQYGSFCTQLNRALS
GIAAEQDRNTREVFAQVKQMYKPTLKYFGGFNFSQILPDPLKPTKRSFI
EDLLFNKVTLADAGFMKQYGECLGDINARDLICAQKFNGLTVLPPLTDD
MIAAYTAALVSGTATAGWTFGAGAALQIPFAMQMAYRFNGIGVTQNVLYE
NOKQIANQFNKAI SQIQESLTTTSTALGKLQDVVNQNAQALNTLVKQLSS
NFGAISSVLNDILSRDKVEAEVQIDRLITGRLQSLQTYVTQQLIRAAEI
RASANLAATKMSECVLGQSKRVDFCGKGYHLMSPQAAPHGVVFLHVTYV
PSQERNFTTAPAICHEGKAYFPREGVVFVNGT SWFITQRNFFSPQIITTD
NTFVSGNCDVVIGIINNTVYDPLQPELDSFKEELDKYFKNHTSPDVLGD
ISGINASVVNIQKEIDRLNEVAKNLNESLIDLQELGKYEQYIKWPWYVWL
GFIAGLIAIVMTILLCCMTSCCSCLKGACSCGSCCKFDEDDSEPVLKGV
KLHYT (SEQ ID NO: 33)

Figure 5

MADNGTITVEELKQLLEQWNLVIGFLFLAWIMLLQFAYSNRNRFLYIIKL
VFLWLLWPVTLACFVLAAYRINWVTGGIAIAMACIVGLMWLSYFVASFR
LFARTRSMWSFNPETNILLNVPLRGTIIVTRPLMESELVIGAVIIRGHLRM
AGHSLGRCDIKDLPKEITVATSRTLSYYKLGASQRVGTDSGFAAYNRYRI
GNYKLNTDHAGSNDNIALLV (SEQ ID NO: 34)

Figure 6

MYSFVSEETGTLIVNSVLLFLAFVVFLLVTLAILTALRLCAYCCNIVNVS
LVKPTVYVYSRVKNLNSSEGVPELLV (SEQ ID NO: 35)

Figure 7

MSDNGPQSNQRSAPRITFGGPTDSTDNNQNGGRNGARPKQRRPQGLPNNT
ASWFTALTQHGKEELRFPRGQVPINTNSGPDDQIGYYRRATRVRGGDG
KMKELSPRWYFYLLGTGPEASLPYGANKEGIVWVATEGALNTPKDHIGTR
NPNNNAATVLQLPQGTTLPKGFYAEGSRGGSQASSRSSRSRGNSRNSTP
GSSRGNSPARMASGGGETALALLLDRLNQLESKVSQKQQQQGQTVTKK
SAAEASKKPRQKRTATKQYNVTQAFGRRGPEQTQGNFGDQDLIRQGTDYK
HWPQIAQFAPSASAFFGMSRIGMEVTPSGTWLTYHGAIKLDDKDPQFKDN
VILLNKHIDAYKTFPPTEPKKDKKKKTDEAQPLPQRQKKQPTVTLLPAAD
MDDFSRQLQNSMSGASADSTQA (SEQ ID NO: 36)

Figure 8

```

BoCov -----MSSVTPAP--VYTWTADEAIKFLKEWNFSL
OC43 -----MSSKTPAP--VYIWTADEAIKFLKEWNFSL
PHEV -----MSSPTTVP--VTSWTADEAIKFLKEWNFSL
FCV MKILLILACAVACVYGEQIRYCAMQ-BTGLSCRNCTASDCBSCFNGGDLIWHLANWNFSW
TGEV MKILLILACVLIACACGE--RYCAMKSDTDLSCRNSTASDCESCFCNGGDLIWHLANWNFSW
TOR2_M -----MAD--NGTITVEELKQLLEQWNLVI
ORF5 -----MAD--NGTITVEELKQLLEQWNLVI
AIBV2 -----MMEN---CTLNLEQATLFLFKEYNLFI
AIBV -----MSNGTEN---CTLSTQQAELPKKEYNLFI
      . : : : : * :

```

```

BoCov GIILLPITVILQFGYTSRSMFVYVVKMVLWMLWPLTIIILTFNVCV--YALNN-VYLGFS
OC43 GIILLPITVILQFGYTSRSMFVYVVKMVLWMLWPLTIIILTFNVCV--YALNN-VYLGFS
PHEV GIIVLFTITILQFGYTSRSMFVYVVKMVLWMLWPLTIIILTFNVCV--YALNN-VYLGFS
FCV SIILIVFITVILQYGRPQFSWFVYGIKMLIMWLLWPLVLAALTFNAYSEYVSRVYVMFGFS
TGEV SILLIVFITVILQYGRPQFSWFVYGIKMLIMWLLWPLVLAALTFNAYSEYVSRVYVMFGFS
TOR2_M IAGAIIVFVILWIMYFVRSIQLYRRTKSWWSFNPETNAILCVNAL--GRSYVLPDGTPTGV
ORF5 GFLFLAWIMLLQFAYSNNRNFYIILKLVFLWLLWPLVLAALTFNAYSEYVSRVYVMFGFS
AIBV2 TAFLLFLTILLQYGYATRSRFYIILKMLVWLCFPLNIAVGVISCI--YPPNT--GGLVAA
AIBV TAFLLFLTILLQYGYATRSRFYIILKMLVWLCFPLNIAVGVISCI--YPPNT--GGLVAA
      : : : * : : : * : : : * : : : * : : : * : : :

```

```

BoCov IVFTIVAII MWIVYFVNSIRLFI RTGSWWSFNPETNMLCIDMK--GRMYVRPIIEDYHTL
OC43 IVFTIVAII MWIVYFVNSIRLFI RTGSWWSFNPETNMLCIDMK--GRMYVRPIIEDYHTL
PHEV IVFTIVAII MWVIVYFVNSIRLFI RTGSWWSFNPETNMLCIDMK--GRMYVRPIIEDYHTL
FCV VAGAVVTFALWMMYFVRSIQLYRRTKSWWSFNPETNAILCVNAL--GRSYVLPDGTPTGV
TGEV IAGAIIVFVILWIMYFVRSIQLYRRTKSWWSFNPETNAILCVNAL--GRSYVLPDGTPTGV
TOR2_M IAMACIVGLMWLSYFVASFRLPARTRSMWWSFNPETNILLNVPLR--GTIVTRPLMESELVI
ORF5 IAMACIVGLMWLSYFVASFRLPARTRSMWWSFNPETNILLNVPLR--GTIVTRPLMESELVI
AIBV2 IILTVFACLSFVGYWYIQSRLFKRCSRWSWWSFNPESNAVGSILLTNGQQCNFAIESVPMVL
AIBV IILTVFACLSFVGYWYIQSRLFKRCSRWSWWSFNPESNAVGSILLTNGQQCNFAIESVPMVL
      : : . . : * : : * : * * : : : * : : : * : : :

```

```

BoCov TVTIIIRGHLYMQGKIKLGTGYSLSDLFPAYVTVAKVSHLLTYKR---GFLDKIGDTSGFVAV
OC43 TVTIIIRGHLYIQGKIKLGTGYSWADLFPAYVTVAKVTHLCTYKR---GFLDRISDTSGFVAV
PHEV TATIIRGHLYIQGKIKLGTGYSLSDLFPAYVTVAKVTHLCTYKR---GFLDRIGDTSGFVAV
FCV TLTLLSGNLYAEGFKMAGGLTIEHLPKYVMIRTPNRTIVYTLV--GKQLKATTATGWAYY
TGEV TLTLLSGNLYAEGFKIAGGMNIDNLPKYVMVALPSRTIVYTLV--GKLLKASSATGWAYY
TOR2_M GAVIIRGHLMAGHSLGR-CDIKDLPEKIVTAT--SRTLSTYKLL--GASQRVGTDSGFAAY
ORF5 GAVIIRGHLMAGHSLGR-CDIKDLPEKIVTAT--SRTLSTYKLL--GASQRVGTDSGFAAY
AIBV2 APIIKNGVLYCEGQWLAK-CEPDHLPKIDIFVCTPDRRNIYRMVQKYTGDSGNKKRVATF
AIBV SPIIKNGALYCEGQWLAK-CEPDHLPKIDIFVCTPDRRNIYRMVQKYTGDSGNKKRFATF
      : * * * : . . * : : : : * : : : * : : :

```

```

BoCov VKSKVGNRYRLPSTQKSGGLDTALLRNNI
OC43 VKSKVGNRYRLPSTQKSGGMDTALLRNNI
PHEV VKSKVGNRYRLPSTHKSGGMDTALLRNNI
FCV VKSKAGDYSTEARTDNLSEHEKLLHMV-
TGEV VKSKAGDYSTEARTDNLSEHEKLLHMV-
TOR2_M NRYRIGNYKLNTHAGSNDNIALLVQ--
ORF5 NRYRIGNYKLNTHAGSNDNIALLVQ--
AIBV2 VYAKQSVDTGELGSVATGSSLYT---
AIBV VYAKQSVDTGELGSVATGSSLYT---
      : . .

```

Key	Name	Genbank	%ID
PHEV	Porcine hemagglutinating encephalomyelitis virus	AAL80035	40.4% (SEQ ID NO: 37)
BoCov	matrix protein [Bovine coronavirus].	NP_150082	40.0% (SEQ ID NO: 38)
AIBV	membrane protein [Avian infectious bronchitis virus].	AAF35863	31.3% (SEQ ID NO: 39)
TGEV	membrane protein [Transmissible gastroenteritis virus].	NP_058427	28.5% (SEQ ID NO: 40)
FCV	membrane [feline coronavirus].	BAC01160	27.7% (SEQ ID NO: 41)
OC43	membrane glycoprotein [Human coronavirus OC43].	AAA45462	39.1% (SEQ ID NO: 42)
AIBV2	membrane protein [Avian infectious bronchitis virus].	AAK83027	32.0% (SEQ ID NO: 43)
TOR2_M/ORF 5	Sars associated coronavirus M glycoprotein	(SEQ ID NO: 34)	

Figure 9

```

BoCov MSFTPGKQSS-SRASSGNRSGNGILK---WADQSDQSRNVQTRGRRAQP--KQTATSQQP
OC43 MSFTPGKQSS-SRASSGNRSGNGILK---WADQSDQSRNVQTRGRRAQP--KQTATSQQP
PHEV MSFTPGKQSS-SRASSGNRSGNGILK---WADQSDQSRNVQTRGRRAQP--KQTATSQQP
MHV MSFVPGQENAGSRSSSVNRAGNGLKKTWADQTERGPNQNRGRRNQP--KQTATTQ-P
AIBV2 -----MASGKAAGK---TDAPAPVIK----LGGPKFP--KVGSSGN--
TCV -----MASGKATGK---TDAPAPIK----LGGPKFP--KVGSSGN--
AIBV -----MASGKAAGK---TDAPAPVIK----LGGPKFP--KVGSSGN--
FCV -----MATQGGQRVN---WGDEFSKRR-----GRSNSR--GRKNNLIP-
PTGV -----MANQGGQRVN---WGDESTKTR-----GRSNSR--GRKNNNIP-
229E -----MATVK---WADASEPQR-----GRQ-----GRIPYSL--
TOR2_N -----MSDNGPQSNQRSAPRITFGGPTDSTDDNNQNGRNGARPKQRREQGLPN

```

```

BoCov SGGNVVPPYYSWFSGITQFQKQKFEFAEGQGVPIAPGVPAATEAKGYWYRHNRRSFKTADG
OC43 SGGNVVPPYYSWFSGITQFQKQKFEFAEGQGVPIAPGVPAATEAKGYWYRHNRRSFKTADG
PHEV SGGTVVPPYYSWFSGITQFQKQKFEFAEGQGVPIAPGVPAATEAKGYWYRHNRRSFKTADG
MHV NSGSVPPHYSWFSGITQFQKQKFEFAEGQGVPIANGIPASEQKGYWYRHNRRSFKTPDG
AIBV2 AS-----WFQAIKAKKLNTPPKFEGSGVDPDENIKPSQQHGYWRRQAR--FKPGKG
TCV AS-----WFQSIKAKKLNTPPKFEGSGVDPDENIKPSQQHGYWRRQAR--FKPGKG
AIBV AS-----WFQALKAKKLNTPPKFEGSGVDPDENIKPSQQHGYWRRQAR--FKPGKG
FCV LS-----YFNPITLDQGSKFWNLCPRDFVPGIGNK--DQQIGYWNRQAR--YRIVKG
PTGV LS-----FFNPITLDQGSKFWNLCPRDFVPGIGNR--DQQIGYWNRQAR--YRIVKG
229E -Y-----SPLLVDS-EQPWKVIPRNLVPIKKDK--NKLIGYWNVQKR--FRTKRG
TOR2_N NTAS-----WFTALTQHG-KEELRFRGQGVPIINTNSGPDQIGYRRATRR--VRGGDG

```

```

BoCov NQRQLLPRWYFYLLGTGPHAKDQYGTDLIDGVVWASNQADVNTPADILDRDPSSDEAIPT
OC43 NQRQLLPRWYFYLLGTGPHAKDQYGTDLIDGVVWASNQADVNTPADIVDRDPSSDEAIPT
PHEV NQRQLLPRWYFYLLGTGPHAKDQYGTDLIDGVVWASNQADINTPADIVDRDPSSDEAIPT
MHV QQRQLLPRWYFYLLGTGPHAGAEGDDIDGVVWASNQADTKTTADIVERDPSSHEAIPT
AIBV2 GRKPVPAWYFYLLGTGPAADLNWGDTPQDQIVVAAKADTKSRNSQGTTRDPKDFDQYPL
TCV GRKPVPAWYFYLLGTGPAADLNWGDTPQDQIVVAAKADVKSRNSQGTTRDPKDFDQYPL
AIBV GRKPVPAWYFYLLGTGPAADLNWGDSDQDQIVVAAKADVKSRNSQGTTRDPKDFDQYPL
FCV QRVELPERWFFYLLGTGPHADAKFKDKLDGVVWVAKDGAMN-KPTSLGTRG-TNNESKPL
PTGV QRRELPERWFFYLLGTGPHADAKFKDKLDGVVWVAKDGAMN-KPTTLGSRG-ANNESKAL
229E KRVDLSPKLFYLLGTGPHKDAKFRERVEGVVWAVDGAKT-EPTGYGVR--KNSEPEIP
TOR2_N KMKELSPRWYFYLLGTGPEASLPYGANKEGIVWVATEGALNTPKDHIGTRNPNNAATVL

```

```

BoCov RFPPTVLPQGYIEGS-GRSAPNSRSTSRASSRASSA---GSRSRANSQNR---TPTSG
OC43 RFPPTVLPQGYIEGS-GRSAPNSRSTSRASSRASSA---GSRSRANSQNR---TPTSG
PHEV RFPPTVLPQGYIEGS-GRSAPNSRSTSRAPNAPS---GSRSRANSQNR---TSTPG
MHV RFAPPTVLPQGYIEGS-GRSAPASRSGSRSSRSGF-----NNRARSSSNQR---QFAS
AIBV2 RFSDG--GPDGNFRWDF-IPLKNRGRSG-RSTAASSAA---ASRAPSRGSR---GRRSD
TCV RFSDG--GPDGNFRWDF-IPLN-RGRSG-RSTAASSAA---SSRAPSRGSR---GRRSG
AIBV RFSDG--GPDGNFRWDF-IPLN-RGRSG-RSTAASSAA---SSRAPSRGSR---GRLNG
FCV KFDGK-I PPQPQLEVN- SRNNSRSGSQSRVSRNRS---QSRGRQSNQ---NTNVED
PTGV KFDGK-VPGEPQLEVN- SRNNSRSGSQSRVSRNRS---QSRGRQSNQ---DDSVEQ
229E HFNQK--LPNGVTVVEE-PDSRAPSRSGSRSGRSGESKPSRNPSSDRNHSQDDIMK
TOR2_N QLPQGTTLPKGFYAEGSRGSSQASSRSSSRSGNSRNPSTPGSSRGNSPARMAS-GGETA

```

```

BoCov VTPDMADQIASLVLAKLGDAAKP-----QQVTKQTAKEVRQK--IL
OC43 VTPDMADQIASLVLAKLGDATKP-----QQVTKHTAKEVRQK--IL
PHEV VTPDMADQIASLVLAKLGDATKP-----QQVTKQTAKEVRQK--IL
MHV VKPDMAEETALVLAKLGDAGQP-----KQVTKQSAKEVRQK--IL
AIBV2 SGDDL IARA AKI IQDQKKGS-----RITKAKADEMAHR--RY
TCV SEDDL IARA AKI IQDQKKGS-----RITKAKADEMAHR--RY
AIBV AEDDL IARA AKI IQDQKKGS-----RITKAKAEEMIHR--RY
FCV TIVAVLQKLGVTDK---QRSRSKS-----GERSQSKSRDTPK--NA
PTGV AVL AALKLGVYTEKQQRSRSKS-----KERSNSKTRDTPK--NE
229E AVAALKSLGPKQEKDKKSAKTGTPKPSRNQSPASSQTSAKSLARSQSSETKEQKHEM
TOR2_N LALLLLDRLNQLLESKVSQKQQQQG-----QTVTKKSAEASKK--PR

```

FIGURE 10A

```

BoCov      NKPRQKRSPNKQCT--VQQCFGKR---GPNQNFGGGEMLKLGTSDFPFI LAELAPTAGA
OC43       NKPRQKRSPNKQCT--VQQCFGKR---GPNQNFGGGEMLKLGTSDFPFI LAELAPTAGA
PHEV       NKPRQKRSPNKQCT--VQQCFGKR---GPNQNFGGGEMLKLGTSDFPFI LAELAPTAGA
MHV        NKPRQKRSPNKQCF--VQQCFGKR---GPNQNFGGGEMLKLGTSDFPFI LAELAPTPSA
AIBV2      CK---RTIIPNYR--VDQVFGPRT--KGKEGNFGDDKMNEEGIKDGRVTAMLNLVPSHA
TCV        CK---RTVPPGYK--VDQVFGPRT--KGKEGNFGDDKMNEEGIKDGRVTAMLNLVPSHA
AIBV       CK---RTVPPGVS--IDKVFGRPT--KGKEGNFGDDKMNEEGIKDGRVTAMLNLVPSHA
FCV        NKHTWKRKTAGKGD---VTNFGAR---SSSANFGSDLVANGNAAKCYPQI AECVPSVSS
PTGV       NKHTWKRKTAGKGD---VTRFYGTR---SNSANFGSDLVANGSSAKHYQI AECVPSVSS
229E      QKPRMKRQPNDDVTSNVTQCFGPR---DLDNHFGSAGVAVANGVAKGYVQPAELVPSATA
TOR2_N     QK---RTATKQYN--VTQAFGRRGPEQTQGNFGDQLIRQGTDYKHWPIAQFAPSASA
          *   :   :   : *   .   *   :   *   :   :   : *   :

```

```

BoCov      FFFGSRLELAKVQNLGSLNDEPQKDVYELRYNGAIR----FDSTLSGFETIMKVLNENL
OC43       FFFGSRLELAKVQNLGSLNDEPQKDVYELRYNGAIR----FDSTLSGFETIMKVLNENL
PHEV       FFFGSRLELAKVQNLGSLNDEPQKDVYELRYNGAIR----FDSTLSGFETIMKVLNENL
MHV        FFFGSKLELVKKN--SGGADDPKDVYELQYSGAIR----FDSTLPGFETIMKVLNENL
AIBV2      CLFGSRVTPKQL--DGLHLRFETTVVPCDDPQFDNYVKICDQCVDGVTGTRPKDDEPKP
TCV        CLFGSRVTPKQLP--DGLHLRFETTVVPRDDPQFDNYVTICDQCVDGVTGTRPKDNEFRP
AIBV       CLFGSOVTPKLP--DGLHLTFRTTVVSRDDPQFDNYVKICDCECDVGVGTRPKDEVVPR
FCV        ILFGSQWSAEAG--DQVKVTLTHNYLPKDDAKTS-----QFLEQI
PTGV       ILFGSYWTSKEDG--DQIBVTFTHKYHLPKDDPKTG-----QFLEQI
229E      MLFDSHIVSKESG--NTVVLTFTRVTVPKDHPHLG-----KFLLEEL
TOR2_N     FFGMSRIGMEVTP--SGTWLTVHGAIKLDDKDPQFK-----DN-----VILLNKHI
          :   *

```

```

BoCov      NAYQQQ-DGTMNMSPKPQRQRG---QKNGQGENDNISVAAPKSRVQONKIRELTAEDIS
OC43       NAYQQQ-DGMNMSPKPQRQRG---HKNGQGENDNISVAVPKSRVQONKSRELTAEDIS
PHEV       NAYQHQEDGMNMSPKPQRQRG---QKNGQVENDNISVAAPKSRVQONKSRELTAEDIS
MHV        DAYQDQAGGADVSPKPKRKRGT--KQKALKQEVNDVSVAKPKSSVQRNVSRELTPEDRS
AIBV2      KSRSSSRPATRGNSPAPRQQRPK--KEKKLKKQDDEADKALTSDEERNNAQLEFVDEP-K
TCV        KSRSSSRPATRGNSPAPRQQRPK--KEKKPKKQDDEVDKALTSDEERNNAQLEFVDEP-K
AIBV       KSRSSSRPATRGTSAPKQQRPK--KEKKPKKQDDEVDKALTSDEERNNAQLEFVDEP-K
FCV        DAYKRP-----SEVAKDQRQ---RKSRSKSAADKPKPELS--VTLEAYTDFVDDTQVE
PTGV       NAYARF-----SEVAKQQRK---RKSRSKSAERSEQEVVVDALIEYITDFVDDTQVE
229E      NAPTRE-----MQQHP-----LLNPSALEFNPSTPATAPVVRDEVSIEET-D
TOR2_N     DAYKTFPP---TEPKKDKKKTDEAQLPQRQKQKPTVTLPAADMDFSRQLQNSMSG
          :   :

```

```

BoCov      LLKKMDEP----FTEDTSEI
OC43       LLKKMDEP----YTEDTSEI
PHEV       LLKKMDEP----YTEDTSEI
MHV        LLAQILDDGVVVDGLEDDSNV
AIBV2      VINWGDA-----LGENEL--
TCV        VINWGDSA-----LGENHL--
AIBV       VINWGDA-----LGENEL--
FCV        MIDEVTN-----
PTGV       MIDEVTN-----
229E      IIDEVN-----
TOR2_N     ASADSTQA-----

```

Key	Genbank	*%ID	
MHV	NUCLEOCAPSID PROTEIN	P18446	34.3% (SEQ ID NO: 44)
BoCov	nucleocapsid protein [Bovine coronavirus].	NP_150083	34.4% (SEQ ID NO: 45)
AIBV	nucleocapsid protein [Avian infectious bronchitis virus].	AAK27162	28.3% (SEQ ID NO: 46)
FCV	nucleocapsid [Feline coronavirus].	CAA74230	29.4% (SEQ ID NO: 47)
PTGV	nucleoprotein [porcine transmissible gastroenteritis virus].	AAMB7563	28.0% (SEQ ID NO: 48)
229E	nucleocapsid protein [Human coronavirus 229E].	NP_073556	24.6% (SEQ ID NO: 49)
OC43	NUCLEOCAPSID PROTEIN.	F33469	33.9% (SEQ ID NO: 50)
PHEV	nucleocapsid protein [porcine hemagglutinating encephalomyelitis]	AAL80036	33.3% (SEQ ID NO: 51)
TCV	nucleocapsid protein [turkey coronavirus].	AAF23873	28.2% (SEQ ID NO: 52)
TOR_N	SARS associated virus nucleocapsid protein (SEQ ID NO: 36)		

FIGURE 10B

ATATTAGGTTTTTACCTACCCAGGAAAAGCCAACCAACCTCGATCTCTTG
TAGATCTGTTCTCTAAACGAACTTTAAAATCTGTGTAGCTGTCCGCTCGGC
TGCATGCCCTAGTGCACCTACGCAGTATAAACAAATAAAATTTTACTGTC
GTTGACAAGAAACGAGTAACTCGTCCCTCTTCTGCAGACTGCTTACGGTT
TCGTCCGTGTGCAGTCGATCATCAGCATAACCTAGGTTTCGTCCGGGTGT
GACCGAAAGSTAAGATGGAGAGCCTTGTTCTTGGTGTCAACGAGAAAACA
CACGTCCAACCTAGTTTGCCTGTCTTTCAGGTTAGAGACGTGCTAGTGCG
TGGCTTCGGGGACTCTGTGGAAGAGGCCCTATCGGAGGCACGTGAACACC
TCAAAAATGGCACTTGTGGTCTAGTAGAGCTGGAAAAGGCGTACTGCC
CAGCTTGAACAGCCCTATGTGTTTATTAAACGTTCTGATGCCTTAAGCAC
CAATCACGGCCACAAGGTCGTTGAGCTGGTTGCAGAAATGGACGGCATT
AGTACGGTTCGTAGCGGTATAACACTGGGAGTACTCGTGCCACATGTGGGC
GAAACCCCAATTGCATACCGCAATGTTCTTCTTTCGTAAAGACGGTAATA
GGGAGCCGGTGGTCATAGCTATGGCATCGATCTAAAGTCTTATGACTTAG
GTGACGAGCTTGGCACTGATCCCATTGAAGATTATGAACAAAACCTGGAAC
ACTAAGCATGGCAGTGGTGCACCTCCGTGAACCTCACCTCGTGAGCTCAATGG
AGGTGCAGTCACTCGCTATGTGACAAACAATTTCTGTGGCCAGATGGGT
ACCCTCTTGATTGCATCAAAGATTTTCTCGCACGCCGGGCAAGTCAATG
TGCACCTCTTCCGAACAACCTTGATTACATCGAGTCGAAGAGAGGTGTCTA
CTGCTGCCGTGACCATGAGCATGAAATTCGCTGGTTCACTGAGCGCTCTG
ATAAGAGCTACGAGCACCAGACACCCCTTCGAAATTAAGAGTGCCAAGAAA
TTTGACACTTTCAAAGGGGAATGCCCAAAGTTTGTGTTTCTCTTAACTC
AAAAGTCAAAGTCAATCAACCACGTGTTGAAAAGAAAAAGACTGAGGGTT
TCATGGGGCGTATACGCTCTGTGTACCCCTGTTGCATCTCCACAGGAGTGT
AACAAATATGCACCTTGTCTACCTTGATGAAATGTAATCATTGCGATGAAGT
TTCATGCGCAGACGTGCGACTTTCTGAAAGCCACTTGTGAACATTGTGGCA
CTGAAAATTTAGTTATTAAGGACCTACTACATGTGGGTACCTACCTACT
AATGCTGTAGTGAAAATGCCATGTCTGCTGTCAAGACCCAGAGATTGG
ACCTGACATAGTGTGTCAGATTATCACACCCTCAAACATTGAAAACCTC
GACTCCGCAAGGGAGGTAGGACTAGATGTTTTGAGGGCTGTGTCTTGCC
TATGTTGGCTGCTATAATAAGCGTGCCCTACTGGGTTCTCGTGTAGTGC
TGATATTGGCTCAGGCCACTTGGCATTACTGGTGACAATGTGGAGACCT
TGAATGAGGATCTCCTTGAGATACTGAGTCGTGAACGTTTAAACATTAAC
ATTTGTTGGCATTCTTCAATTTGAATGAAGAGGTTGCCATCATTCTGGCATC
TTTCTCTGCTTCTACAAGTGCCCTTATTGACACTATAAAGAGTCTTGATT
ACAAGTCTTTCAAACCATGTTGAGTCTTCCGGTAACTATAAAGTTACC
AAGGGAAAGCCCGTAAAAGGTGCTTGGAACATTGGACAACAGAGATCAGT
TTTAAACACCCTGTGTGGTTTCCCTCACAGGCTGCTGGTGTATCAGAT
CAATTTTTCGCGCACACTTGATGCAGCAAACCACTCAATTCCTGATTTG
CAAAGAGCAGCTGTCACCATACTTGATGGTATTTCTGAACAGTCATTACG
TCTTGTGACGCCATGGTTTACTTTCAGACCTGCTCACCAACAGTGTCA
TTATTATGGCATATGTAACGTGGTCTTGTACAACAGACTTCTCAGTGG
TTGTCTAATCTTTGGGCACACTGTGAAAACCTCAGGCCTATCTTTGA
ATGGATTGAGGCGAAACTTAGTGCAGGAGTTGAATTTCTCAAGGATGCTT
GGGAGATTTCAAATTTCTCATACAGGTGTTTTTGACATCGTCAAGGGT
CAAATACAGGTTGCTTCAGATAACATCAAGGATTTGTGTAATGCTTCAT
TGATGTTGTTAAACAAGGCACCTCGAAATGTGCATTGATCAAGTCATATCG
CTGGCGCAAAGTTGCGATCACTCAACTTAGGTGAAGTCTTCATCGCTCAA
AGCAAGGGACTTTACCGTCAGTGTATACGTGGCAAGGAGCAGCTGCAACT
ACTCATGCCCTTAAAGGCACAAAAGAAGTAACCTTTCTTGAAGGTGATT
CACATGACACAGTACTTACCTCTGAGGAGGTTGTTCTCAAGAACGGTGAA
CTCGAAGCACTCGAGACGCCCGTTGATAGCTTACAAATGGAGCTATCGT
TGGCACACCAGTCTGTGTAATGGCTCATGCTCTTAGAGATTAAGGACA
AAGAACAATACTGGCATTGTCTCTTGGTTTACTGGCTACAACAATGTC
TTTTCGCTTAAAGGGGGTGCACCAATTAAGGTGTAACCTTTGGAGAAGA
TACTGTTTGGGAAGTTCAAGGTTACAAGAATGTGAGAATCAATTTGAGC
TTGATGAACGTGTTGACAAAGTGTAAATGAAAAGTGTCTGTCTACACT

FIGURE 11A

GTTGAATCCGGTACCGAAGTTACTGAGTTTGCATGTGTTGTAGCAGAGGC
TGTTGTGAAGACTTTACAACCACTTCTGATCTCCTTACCAACATGGGTA
TTGATCTTGATGAGTGGAGTGTAGCTACATTTACTTATTTGATGATGCT
GGTGAAGAAAACCTTTTCATCACGTATGTATTGTTCCCTTTACCCCTCCAGA
TGAGGAAGAAGAGGACGATGCAGAGTGTGAGGAAGAAGAAATTTGATGAAA
CCTGTGAACATGAGTACGGTACAGAGGATGATTATCAAGGTCTCCCTCTG
GAATTTGGTGCCTCAGCTGAAACAGTTCGAGTTGAGGAAGAAGAAGAGGA
AGACTGGCTGGATGATACTACTGAGCAATCAGAGATTGAGCCAGAACCAG
AACCTACACCTGAAGAACCAGTTAATCAGTTTACTGGTTATTTAAAACCTT
ACTGACAATGTTGCCATTAATGTGTTGACATCGTTAAGGAGGCACAAAAG
TGCTAATCCTATGGTGATTGTAATGCTGCTAACATACACCTGAAACATG
GTGGTGGTGTAGCAGGTGCACCAACAAGGCAACCAATGGTGCCATGCAA
AAGGAGAGTGATGATTACATTAAGCTAAATGGCCCTTTACAGTAGGAGG
GTCCTTGTGTTGCTTTCTGGACATAATCTTGCTAAGAAGTGTCTGCATGTTG
TTGGACCTAACCTAAATGCAGGTGAGGACATCCAGCTTCTTAAGGCAGCA
TATGAAAATTTCAATTCACAGGACATCTTACTTGCACCATTGTTGTCAGC
AGGCATATTTGGTGCTAAACCACCTCAGTCTTTACAAGTGTGCGTGCAGA
CGGTTGTCATGGATTATCTTGATAACCTGAAGCCTAGAGTGGAAAGCACC
TAAACAAGAGGAGCCACCAACACAGAAGATTCCAAAACCTGAGGAGAAAT
CTGTCTACAGAAGCCTGTGATGTGAAGCCAAAATTAAGGCCCTGCATT
GATGAGGTTACCACAACACTGGAAGAACTAAGTTTCTTACCAATAAGTT
ACTCTTGTGTTGCTGATATCAATGGTAAGCTTACCATGATTCTCAGAACA
TGCTTAGAGGTGAAGATATGCTTTCCCTTGAGAAGGATGCACCTTACATG
GTAGGTGATGTTATCACTAGTGGTGATATCACTTGTGTTGTAATACCCCTC
CAAAAAGGCTGGTGGCACTACTGAGATGCTCTCAAGAGCTTTGAAGAAAAG
TGCCAGTTGATGAGTATATAACCACGTACCCTGGACAAGGATGTGCTGGT
TATACACTGAGGAAGCTAAGACTGCTCTTAAGAAATGCAAATCTGCATT
TTATGTACTACCTTCAGAAGCACCTAATGCTAAGGAAGAGATTCTAGGAA
CTGTATCCTGGAATTTGAGAGAAATGCTTGCTCATGCTGAAGAGACAAGA
AAATTAATGCCTATATGCATGGATGTTAGAGCCATAATGGCAACCATCCA
ACGTAAGTATAAAGGAATTAATAATCAAGAGGGCATCGTTGACTATGGTG
TCCGATTCCTCTTTTATACTAGTAAAGAGCCTGTAGCTTCTATTATACG
AAGCTGAACCTCTTAAATGAGCCGCTTGTCACAATGCCAATTTGGTTATGT
GACACATGGTTTAACTCTGAAGAGGCTGCGCGCTGTATGCGTTCTCTTA
AAGCTCCTGCCGTAGTGTGATGATCATCACCAGATGCTGTTACTACATAT
AATGGATACCTCACTTCGTCATCAAAGACATCTGAGGAGCACCTTGTAGA
AACAGTTCTTTGGCTGGCTCTTACAGAGATTGGTCTATTTCAGGACAGC
GTACAGAGTTAGGTGTTGAATTTCTTAAGCGTGGTGACAAAATTTGTGTAC
CACACTCTGGAGAGCCCCGTGAGTTTCATCTTGACGGTGAGGTTCTTTC
ACTTGACAACTAAAGAGTCTCTTATCCCTGCGGGAGGTTAAGACTATAA
AAGTGTTCACAACCTGTGGACAACACTAATCTCCACACACAGCTTGTGGAT
ATGTCTATGACATATGGACAGCAGTTTGGTCCAACATACTTGGATGGTGC
TGATGTTACAAAATTAACCTCATGTAATCATGAGGGTAAGACTTTCCT
TTGTACTACCTAGTGATGACACACTACGTAGTGAAGCTTTCGAGTACTAC
CATACTCTTGATGAGAGTTTTCTTGGTAGGTACATGTCTGCTTTAAACCA
CACAAAGAAATGGAATTTCCCTCAAGTTGGTGGTTAACTTCAATTAAT
GGGCTGATAACAAATGTTATTTGTCTAGTGTTTTATTAGCACTTCAACAG
CTTGAAGTCAAATTCATGCACCAGCACTTCAAGAGGCTTATTATAGAGC
CCGTGCTGGTGTGCTGCTAACTTTTGTGCACTCATACTCGCTTACAGTA
ATAAAAACCTGTTGGCGAGCTTGGTGTGTCAGAGAACTATGACCCTCTT
CTACAGCATGCTAATTTGGAATCTGCAAGCGAGTTCTTAATGTGGTGTG
TAAACATTTGGTGTGAGAAAACACTACTACCTTAACGGGTGTAGAAGCTGTGA
TGTATATGGGTACTCTATCTTATGATAATCTTAAGACAGGTGTTTCCATT
CCATGTGTGTGTGGTCTGATGCTACACAATATCTAGTACAACAAGAGTC
TTCTTTTGTATGATGCTGCACCACCTGCTGAGTATAAATACAGCAAG
GTACATTCCTTATGTGCGAATGAGTACACTGGTAACTATCAGTGTGGTCAT

FIGURE 11B

TACACTCATATAACTGCTAAGGAGACCCCTCTATCGTATTGACGGAGCTCA
CCTTACAAAGATGTCAGAGTACAAAGGACCAGTGACTGATGTTTTCTACA
AGGAAACATCTTACACTACAACCATCAAGCCTGTGTCGTATAAACTCGAT
GGAGTTACTTACACAGAGATTGAACCAAATTTGGATGGGTATTATAAAAA
GGATAATGCTTACTATACAGAGCAGCCTATAGACCTTGTACCAACTCAAC
CATTACCAAATGCGAGTTTTGATAATTTCAAACTCACATGTTCTAACACA
AAATTTGCTGATGATTTAAATCAAATGACAGGCTTCACAAAGCCAGCTTC
ACGAGAGCTATCTGTCCACATTCCTCCAGACTTGAATGGCGATGTAGTGG
CTATTGACTATAGACACTATTTCAGCGAGTTTCAAGAAAGGTGCTAAATTA
CTGCATAAGCCAATTGTTTGGCACATTAACCAGGCTACAACCAAGACAAC
GTTCAAACCAAACACTTGGTGTTCAGTTGCTTTGGAGTACAAAGCCAG
TAGATACTTCAAATTCATTTGAAGTTCTGGCAGTAGAAGACACACAAGGA
ATGGACAATCTTGC TTGTGAAAGTCAACAACCCACCTCTGAAGAAGTAGT
GGAAAATCCTACCATACAGAAGGAAGTCATAGAGTGTGACGTGAAAAC TA
CCGAAGTTGTAGGCAATGTCATACTTAAACCATCAGATGAAGGTGTTAAA
GTAACACAAGAGTTAGGTTCATGAGGATCTTATGGCTGCTTATGTGGAAA
CACAAGCATTACCATTAAGAAAACCTAATGAGCTTTCCTAGCCTTAGGTT
TAAAAACAATTGCCACTCATGGTATTGCTGCAATTAATAGTGTTCCTTGG
AGTAAAAATTTGGCTTATGTCAAACCATTCCTAGGACAAGCAGCAATTAC
AACATCAAATTGGCGCTAAGAGATTAGCACAACGTTGTGTTAAACAATTATA
TGCCCTATGTGTTTACATTAATGTTCCAATTTGTGTACTTTTACTAAAAGT
ACCAATCTAGAATTAGAGCTTCACTACCTACAAC TATGCTAAAAATAG
TGTTAAGAGTGTGCTAAATTAATGTTGGATGCCGGCATAAATTATGTGA
AGTCACCCAAATTTCTAAATGTTTCACAATCGCTATGTGGCTATGTGTTG
TTAAGTATTTGCTTAGGTTCTCTAATCTGTGTAAC TGCTGCTTTTGGTGT
ACTCTTATCTAATTTTGGTGCTCCTTCTTATTGTAATGGCGTTAGAGAAT
TGTATCTTAATTCGTCTAACGTTACTACTATGGATTTCTGTGAAGGTTCT
TTTCTTGCAGCATTGTTTAAGTGGATTAGACTCCCTTGATTCCTATCC
AGCTCTGAAACCATT CAGGTGACGATTT CATCGTACAAGCTAGACTTGA
CAATTTTAGGTC TGCCGCTGAGTGGGTTTGGCATATATGTTGTT CACA
AAATCTTTTATTTATTAGGTC TTT CAGCTATAATGCAGGTGTTCTTTGG
CTATTTTGCTAGTCATTT CATCAGCAATCTTGGCTCATGTGGTTTATCA
TTAGTATTGTACAAATGGCACCCGTTCTGCAATGGTTAGGATGTACATC
TCTTTTGCTTCTTCTACTACATATGGAAGAGCTATGTT CATATCATGGA
TGGTTGCACCTCTTCGACTTGCATGATGTGCTATAAGCGCAATCGTGCCA
CACGCGTTGAGTGTACAAC TATGTTAATGGCATGAAGAGATCTTCTAT
GTCTATGCAAATGGAGGCCGTTGGCTTCTGCAAGACTCACAATTGGAATTG
TCTCAATTTGTGACACATTTTGCAC TGGTAGTACATTCATTAGTGATGAAG
TTGCTCGTGATTTGTCAC TCCAGTTTAAAAGACCAATCAACCCTACTGAC
CAGTCATCGTATATTGTTGATAGTGTGCTGTGAAAAATGGCGCGCTTCA
CCCTACTTTGACAAGGCTGGTCAAAGACCTATGAGAGACATCCGCTCT
CCCATTTTGTCAATTTAGACAATTTGAGAGCTAACAACACTAAAGGTTCA
CTGCCTATTAATGTCATAGT TTTTGTGATGGCAAGTCCAAATGCGACGAGTC
TGCTTCTAAGCTGCTTCTGTG TACTACAGTCAGCTGATGTGCCAACCTA
TTCTGTTGCTTGACCAAGCTCTTGTATCAGACGTTGGAGATAGTACTGAA
GTTTCCGTTAAGATGTTTGTATGCTTATGTCGACACCTTTT CAGCAACTTT
TAGTGTCCCTATGGA AAAACTTAAGGCAC TTTGCTACAGCTCACAGCG
AGTTAGCAAAGGGTGTAGCTTTAGATGGTGTCTTTCTACATTCGTGTCA
GCTGCCCGACAAGGTGTTGTTGATACCGATGTGACACAAGGATGTTAT
TGAATGTCTCAAAC TTT CACATCACTCTGACTTAGAAGTGACAGGTGACA
GTTGTAACAATTT CATGCTCACCTATAATAAGGTTGAAAACATGACGCC
AGAGATCTTGGCGCATGTATTGACTGTAATGCAAGGCATATCAATGCCCA
AGTAGCAAAAAGTCACAATGTTTCACTCATCTGGAATGTAAAAGACTACA
TGTCTTTATCTGAACAGCTGCGTAAACAAATTCGTAGTGCTGCCAAGAAG
AACACATACCTTTTAGACTA ACTTGTGCTACAAC TAGACAGGTTGTCAA
TGTCTAATACTACTAAAATCTCACTCAAGGTTGGTAAGATTGTTAGTACTT
GTTTTAAACTTATGCTTAAGGCCACATTATTGTGCGTTCTTGCTGCATTG

FIGURE 11C

GTTTGTATATATCGTTATGCCAGTACATACATTGTCAATCCATGATGGTTA
CACAAATGAAATCATTGGTTACAAAGCCATTCAGGATGGTGTCACTCGTG
ACATCATTCTACTGATGATTGTTTTGCAAATAAACATGCTGGTTTTGAC
GCATGGTTTAGCCAGCGTGGTGGTTCATACAAAATGACAAAAGCTGCC
TG TAGTAGCTGCTATCATTACAAGAGAGATPGGTTTCATAGTGCCGCT
TACCGGGTACTGTGCTGAGAGCAATCAATGGTGACTTCTTGCATTTTCTA
CCTCGTGTTTTTAGTGCTGTGGCAACATTTGCTACACACCTTCCAACT
CATTGAGTATAGTGATTTGCTACCTCTGCTTGCCTTCTGCTGCTGAGT
GTACAATTTTTAAGGATGCTATGGGCAAACCTGTGCCATATTGTTATGAC
ACTAATTTGCTAGAGGGTCTATTTCCTTATAGTGAGCTTCGTCCAGACAC
TCGTTATGTGCTTATGGATGGTTCCATCATAACAGTTTCCCTAACACTTACC
TGGAGGGTCTGTAGAGTAGTAACAACCTTTGATGCTGAGTACTGTAGA
CATGGTACATGCGAAAGGTCAGAAGTAGGATTTGCCATCTACCAGTGG
TAGATGGGTTCTTAATAATGAGCATTACAGAGCTCTACAGGAGTTTTCT
GTGGTGTGATGCGATGAATCTCATAGCTAACATCTTACTCCTCTTGTG
CAACCTGTGGGTGCTTAGATGTGTCTGCTTTCAGTAGTGGCTGGTGGTAT
TATGGCCATATTGGTGACTTGTGCTGCCACTACTTTATGAAATTCAGAC
GTGTTTTGGTGAGTACAACCATGTTGTTGCTGCTAATGCATTTTGT
TTGATGCTTTTCACTATACTCTGCTGGTACCAGCTTACAGCTTTCTGCC
GGGAGTCTACTCAGTCTTTTACTTGTACTTGACATTTCTATTTACCAATG
ATGTTTCATTCTTGGCTCACCTTCAATGGTTGCCATGTTTTCTCCTATP
GTGCCTTTTGGATAACAGCAATCTATGATTCTGTATTTCTCTGAAGCA
CTGCCATGGTTCTTTAACAACCTATCTTAGGAAAAGAGTCATGTTAATG
GAGTTACATTTAGTACCTTCGAGGAGGCTGCTTTGTGTACCTTTTTGCTC
AACAGGAAATGTACCTAAAATGCCGTAGCGAGACACTGTTGCCACTTAC
ACAGTATAACAGGTATCTTGCTCTATATAACAAGTACAAGTATTTACAGT
GAGCCTTAGATACTACCAGCTATCGTGAAGCAGCTTCTGCCACTTAGCA
AAGGCTCTAAATGACTTTAGCAACTCAGGTGCTGATGTTCTTACCAACC
ACCACAGACATCAATCACTTCTGCTGTCTGTCAGAGTGGTTTTAGGAAAA
TGGCATTCCCGTCAGGCAAAGTTGAAGGGTGCATGGTACAAGTAACTGT
GGAATACAACCTTAATGGATTGTGGTTGGATGACACAGTATACTGTCC
AAGACATGTCATTTGCACAGCAGAAGACATGCTTAATCCTAACTATGAAG
ATCTGCTCATTGCGAAATCCAACCATAGCTTTCTTGTTCAGGCTGGCAAT
GTTCAACTTCGTGTTATTGGCCATTTCTATGCAAAATTTGCTGCTTAGGCT
TAAAGTTGATACTTCTAACCCCTAAGACACCCAAGTATAAATTTGTCCGTA
TCCAACCTGGTCAAACATTTTCACTTCTAGCATGCTACAATGGTTCAACA
TCTGGTGTATTATCAGTGTGCCATGAGACCTAATCATAACATTAAAGGTT
TTTCTTAATGGATCATGTGGTAGTGTGGTTTTAACATTGATTATGATT
GCGTGTCTTCTGCTATATGCATCATATGGAGCTTCCAACAGGAGTACAC
GCTGGTACTGACTTAGAAGGTAATTTCTATGGTCCATTTGTTGACAGACA
AACTGCACAGGCTGCAGGTACAGACACAACCATAACATTAATGTTTTGG
CATGGCTGTATGCTGCTGTTATCAATGGTGATAGGTTGTTTTCTTAATAGA
TTCACCACTACTTTGAATGACTTTAACCTTGTGGCAATGAAGTACAACCTA
TGAACCTTTGACACAAGATCATGTTGACATATTGGGACCTCTTTCTGCTC
AAACAGGAATGGCGTCTTAGATATGTGTGCTGCTTTGAAAGAGCTGCTG
CAGAAATGGTATGAAATGGTCTGCTATCCTTGGTAGCACTATTTTAGAAGA
TGAGTTTACACCATTTGATGTTGTTAGACAATGCTCTGGTGTACCTTCC
AAGGTAAGTTCAAGAAAATTTGTTAAGGGCACTCATATTGGATGCTTTTA
ACTTTCTTGACATCACTATTGATTCTTGTTCAAAGTACACAGTGGTCACT
GTTTTCTTTGTTTACGAGAATGCTTTCTTGGCAATTTACTCTTGGTATTA
TGGCAATTGCTGCATGTGCTATGCTGCTTGTAAAGCATAAGCAGCATTC
TTGTGCTTGTTCGTTACCTTCTCTTGAACAGTTGCTTACTTTAATAT
GGTCTACATGCCGCTAGCTGGGTGATGCGTATCATGACATGGCTTGAAT
TGGCTGACACTAGCTTGTCTGGTTATAGGCTTAAGGATTGTGTTATGAT
GCTTCAGCTTTAGTTTTGCTTATTTCTCATGACAGCTCGCACTGTTTATGA
TGATGCTGCTAGACGTGTTTGGACACTGATGAATGTCATTACACTTGT
ACAAAGTCTACTATGGTAATGCTTTAGATCAAGCTATTTCCATGTGGCC

FIGURE 11D

TTAGTTATTTCTGTAACCTCTAACTATTCTGGTGTGCTTACGACTATCAT
GTTTTTAGCTAGAGCTATAGTGTGTTGTGTGTGTTGAGTATTACCCATTGT
TATTTATTAAGTGGCAACACCTTACAGTGTATCATGCTTGTATTATTGTTTC
TTAGGCTATTGTTGCTGCTGCTACTTTGGCCTTTTCTGTTTACTCAACCG
TTACTTCAGGCTTACTCTTGGTGTGTTATGACTACTTGGTCTCTACACAAG
AATTTAGGTATATGAACTCCCAGGGGCTTTTGCCTCCTAAGAGTAGTATT
GATGCTTTCAAGCTTAACATTAAGTTGTTGGGTATTGGAGGTAACCATG
TATCAAGGTTGCTACTGTACAGTCTAAAATGTCTGACGTAAAGTGCACAT
CTGTGGTACTGCTCTCGGTCTTCAACAACCTTAGAGTAGAGTCATCTTCT
AAATTGTGGGCACAATGTGTACAACCTCCACAATGATATTCTTCTTGCAA
AGCACAACTGAAGCTTTCGAGAAGATGGTTCCTCTTTTGTCTGTTTTGC
TATCCATGCAGGGTGTGTAGACATTAATAGGTTGTGCGAGGAAATGCTC
GATAACCGTGTACTCTTCAGGCTATTGCTTCAGAATTTAGTTCTTTACC
ATCATATGCCGCTTATGCCACTGCCAGGAGGCTTATGACAGGCTGTAG
CTAATGGTGAATCTGAAGTCGTTCTCAAAAAGTTAAAGAAATCTTTGAAT
GTGGCTAAATCTGAGTTTGACCGTGATGCTGCCATGCAACGCAAGTTGGA
AAAGATGGCAGATCAGGCTATGACCCAAATGTACAAACAGGCAAGATCTG
AGGACAAGAGGGCAAAAGTAAC TAGTGCTATGCAACAATGCTCTTCACT
ATGCTTAGGAAGCTTGATAATGATGCACTTAACAACATTTACAACAATGC
GCGTGATGGTTGTGTTCCACTCAACATCATACCATTGACTACAGCAGCCA
AACTCATGGTTGTTGTCCCTGATTATGGTACCTACAAGAACACTTGTGAT
GGTAACACCTTTACATATGCATCTGCACCTCTGGGAAATCCAGCAAGTTGT
TGATGCGGATAGCAAGATTGTTCAACTTAGTGAAATTAACATGGACAATT
CACCAAATTTGGCTTGGCCTCTTATTGTTACAGCTCTAAGAGCCAACCTCA
GCTGTAAACTACAGAATAATGAAC TGAGTCCAGTAGCACTACGACAGAT
GTCTGTGCGGCTGGTACCACACAACAGCTTGTACTGATGACAATGCAC
TTGCCACTATAACAATTCGAAGGGAGGTAGGTTTGTGCTGGCATTACTA
TCAGACCACCAAGATCTCAAATGGGCTAGATTCCCTAAGAGTGATGGTAC
AGGTACAATTTACACAGAAGTGAACCACTTGTAGGTTTGTACAGACA
CACCAAAGGGCTAAAGTGAATACTTGTACTTCATCAAAGGCTTAAAC
AACCTAAATAGAGGTATGGTGTGCTGGGCAGTTTAGCTGCTACAGTACGCTC
TCAGGCTGGAAATGCTACAGAAGTACCTGCCAATTCAACTGTGCTTTTCT
TCTGTGCTTTTGCAGTAGACCTGCTAAAGCATATAAGGATTACCTAGCA
AGTGGAGGACAACCAATCACCACCTGTGTGAAGATGTTGTGTACACACAC
TGGTACAGGACAGGCAATTAAGTAAACACCAGAGCTAACATGGACCAAG
AGTCTTTGGTGGTGTCTCATGTTGTCTGTATTGTAGATGCCACATTTGAC
CATCCAAATCTAAAGGATTTCTGTGACTTGAAGGTAAGTACGTTCCAAAT
ACCTACCCTTGTGCTAATGACCCAGTGGGTTTTTACACTTAGAAAACAG
TCTGTACCGTCTGCGGAATGTGGAAGGTTATGGCTGTAGTTGTGACCAA
CTCCGGAACCCCTTGATGCAGTCTGCGGATGCATCAACGTTTTTAAACGG
GTTTGGCGTGAAGTGCAGCCGCTTTACACCGTGGGCACAGGCACTAG
TACTGATGTCGTCTACAGGGCTTTTGTATTTTACACGAAAAAGTTGCTG
GTTTTGCAAAGTTCTTAAAACTAATTGCTGTGCTTCCAGGAGAAGGAT
GAGGAAGGCAATTTATTAGACTCTTACTTTGTAGTTAAGAGGCATACTAT
GTCTAACTACCAACATGAAGAGACTATTTATAACTTGGTTAAAGATTGTC
CAGCGGTTGCTGTCCATGACTTTTTCAAGTTTAGAGTAGATGGTGACATG
GTACCACATATATCACGTCAGCGTCTAACTAAATACACAATGGCTGATTT
AGTCTATGCTCTACGTCATTTTGTGATGAGGGTAATTGTGATACATTAAG
AAATACTCGTCACATACAATGCTGTGATGATGATTAATTTCAATAAGAAG
GATTGGTATGACTTCGTAGAGAATCCTGACATCTTACCGGTATATGCTAA
CTTAGGTGAGCGTGTACGCCAATCATTTAATAAGACTGTACAATCTGCG
ATGCTATGCTGATGAGGCAATGTTAGGCGTACTGACATTAAGATAATCAG
GATCTTAATGGGAACCTGGTACGATTTCCGGTGAATTCGTACAAGTAGCACC
AGGCTGCGGAGTTCCTATTGTGGATTCATATTACTCATGCTGATGCCCA
TCCTCACTTTGACTAGGGCATTTGGCTGTGAGTCCCATATGGATGCTGAT
CTCGCAAACCACTTATTAAGTGGGATTTGCTGAAATATGATTTTACGGA
AGAGAGACTTTGTCTCTTCGACCGTTATTTTAAATATTTGGGACCAGACAT

FIGURE 11E

ACCATCCCAATTGTATTAAC TGTGGATGATAGGTGATCCTTCATTGT
GCAAAC TTTAATGTGTTATTTCTACTGTGTTCCACCTACAAGTTTGG
ACCAC TAGTAAGAAAAATATTTGTAGATGGTGTCCCTTTTGTGTTCAA
CTGGATACCATTTCGTGAGTTAGGAGTCGTACATAATCAGGATGTAAAC
TTACATAGCTCGCGTCTCAGTTCAAGGAAC TTTAGTGTATGCTGCTGA
TCCAGCTATGCATGCAGCTTCTGGCAATTTATTGCTAGATAAACGCCTA
CATGCTTTTCAGTAGCTGCAC TAACAAACAATGTTGCTTTTCAAAC TGC
AAACCCGGTAATTTAATAAAGACTTTTATGACTTTCGTGTGCTAAAGG
TTTCTTTAAGGAAGGAAGTTCGTGAAC TAAACACTTCTTCTTTGCTC
AGGATGGCAACGCTGCTATCAGTGATTATGACTATTATCGTTATAATCTG
CCAACAATGTGTGATATCAGACAAC TCTATTTCGTAGTTGAAGTTGTGA
TAAATAC TTTGATTGTTACGATGGTGGCTGTATTAATGCCAACCAAGTAA
TCGTTAACAATCTGGATAAATCAGCTGGTTTCCCATT AATAAATGGGGT
AAGGCTAGACTTTATTATGACTCAATGAGTTATGAGGATCAAGATGCACT
TTTCGCGTATACTAAGCGTAATGTCATCCCTACTATAACTCAAATGAATC
TTAAGTATGCCATTAGTGCAAAGAATAGAGCTCGCACCGTAGCTGGTGT
TCTATCTGTAGTACTATGACAAATAGACAGTTTCATCAGAAATTATTGAA
GTCAATAGCCGCCACTAGAGGAGCTACTGTGGTAATTGGAACAAGCAAGT
TTTACGGTGGCTGGCATAATATGTTAAAAACTGTTTACAGTGTGTAGAA
ACTCCACACTTATGGGTGGGATTATCCAAAATGTGACAGAGCCATGCC
TAACATGCTTAGGATAATGGCCTCTCTTGTTC TCGCTCGCAAACATAACA
CTTGCTGTAAC TTTACACCCGTTTCTACAGGTTAGCTAACGAGTGTCCG
CAAGTATTAAGTGAGATGGTCATGTGTGGCGGCTCACTATATGTTAAACC
AGGTGGAACATCATCCGGTGATGCTACAAC TCGCTTATGCTAATAGTGTCT
TTAACATTTGTCAAGCTGTTACAGCCAATGTAATGCAC TTTCTTCAA
GATGGTAATAAGATAGCTGACAAGTATGTCGCAATCTACAACACAGGCT
CTATGAGTGTCTCTATAGAAATAGGGATGTTGATCATGAATTCGTGGATG
AGTTTACGCTTACCTGCGTAAACATTTCTCCATGATGATTCTTTCTGAT
GATGCCGTTGTGTGCTATAACAGTAACTATGCGGCTCAAGGTTTAGTAGC
TAGCATTAAGAACTTTAAGGCAGTTCTTTATTATCAAATAATGTGTTCA
TGTCTGAGGCAAAATGTTGGACTGAGACTGACCTTACTAAAGGACCTCAC
GAATTTGCTCACAGCATAACAATGCTAGTTAAACAAGGAGATGATTACGT
GTACCTGCCTTACCCAGATCCATCAAGAATATTAGGCGCAGGCTGTTTTG
TCGATGATATTGTCAAACAGATGGTACACTTATGATTGAAAGGTTTCGTG
TCACTGGCTATTGATGCTTACCCACTTACAAAACATCCTAATCAGGAGTA
TGCTGATGCTTTTCACTTGTATTTACAATACATTAGAAAGTTACATGATG
AGCTTACTGGCCACATGTTGGACATGTATTCGCTAATGCTAAC TAATGAT
AACACCTACGGTACTGGGAACCTGAGTTTATGAGGCTATGTACACACC
ACATACAGTCTTGCAGGCTGTAGGTGCTTGTGTATTGTGCAATTCACAGA
CTTCACTTCGTTGCGGTGCCTGTATTAGGAGACCATTCCATGTTGCAAG
TGCTGCTATGACCATGTCA TTTCAACATCACACAAATTAGTGTGCTGT
TAATCCCTATGTTTGAATGCCCCAGGTTGTGATGTCACTGATGTGACAC
AACTGTATCTAGGAGGTATGAGCTATTATTGCAAGTCACATAAGCCTCCC
ATTAGTTTTCATTATGTGCTAATGGTCAAGTTT TTTGGTTTATACAAAA
CACATGTGTAGGCAGTGACAATGTCACTGACTTCAATGCGATAGCAACAT
GTGATGGACTAATGCTGGCGATTACATAC TTGCCAACACTTGTACTGAG
AGACTCAAGCTTTTCGCAGCAGAAACGCTCAAAGCCACTGAGGAAACATT
TAAGCTGTCATATGGTATTGCCACTGTACGCGAAGTACTCTCTGACAGAG
AATTGCATCTTTTCATGGGAGGTTGGAAAACCTAGACCACCATGAAACAGA
AACTATGCTTTTACTGGTTACCGTGTAAC TAAAAATAGTAAAGTACAGAT
TGGAGAGTACACCTTTGAAAAAGGTGACTATGGTGTGCTGTTGTGTACA
GAGGTACTACGACATACAAGTTGAATGTTGGTGTGATTACTTTGTGTTGACA
TCTCACACTGTAATGCCACTTAGTGCACCTACTCTAGTGCCACAAGAGCA
CTATGTGAGAATTACTGGCTTGTACCCAACACTCAACATCTCAGATGAGT
TTTCTAGCAATGTTGCAAATATCAAAGGTCGGCATGCAAAAGTACTCT
ACACTCCAAGGACCAC TGGTACTGGTAAGAGTCATTTTGCCATCGGACT
TGCTCTCTATTACCCATCTGCTCGCATAGTGTATACGGCATGCTCTCATG

FIGURE 11F

CAGCTGTTGATGCCCTATGTGAAAAGGCATTAATAATTTGCCCATAGAT
AAATGTAGTAGAATCATACCTGCGCGTGCGCGCTAGAGTGTTTTGATAA
ATTCAAAAGTGAATTC AACACTAGAACAGTATGTTTCTGCACTGTAAATG
CATTGCCAGAAACAAC TGCTGACATTGTAGTCTTTGATGAAATCCTATG
GCTACTAATTATGACTTGAGTGTGTCAATGCTAGACTTCGTGCAAAACA
CTACGTCTATATTGGCGATCCTGCTCAATTACCAGCCCCCGCACATTGC
TGACTAAAGGCACACTAGAACAGAAATTTTAATTCAGTGTGCAGACTT
ATGAAAACAATAGGTCCAGACATGTTCCCTGGAACTTGTCGCCGTTGTCC
TGCTGAAATGTTGACACTGTGAGTGTCTTAGTTTATGACAATAAGCTAA
AAGCACACAAGGATAAGTCAGCTCAATGCTTCAAATGTTCTACAAAGGT
GTTATTACACATGATGTTTCATCTGCAATCAACAGACCTCAAATAGGCGT
TGTAAGAGAATTTCTTACACGCAATCCTGCTTGGAGAAAAGCTGTTTTTA
TCTCACCTTATAATTCACAGAACGCTGTAGCTTCAAATCTTAGGATTG
CCTACGCAGACTGTTGATTCATCACAGGGTCTGAAATATGACTATGTCAT
ATTCACACAAACTACTGAAACAGCACACTCTGTAATGTCAACCGCTTCA
ATGTGGCTATCACAAGGGCAAAAATGGCATTGTTGTCATAATGTCTGAT
AGAGATCTTTATGACAACTGCAATTTACAAGTCTAGAAATACCACGTCG
CAATGTGGCTACATTACAAGCAGAAAATGTAAGTGGACTTTTTAAGGACT
GTAGTAAGATCATTACTGGTCTTTCATCCTACACAGGCACCTACACACCTC
AGCGTTGATATAAAGTTC AAGACTGAAGGATTAATGTTGACATACCAGG
CATACCAAAGGACATGACCTACCGTAGACTCATCTCTATGATGGGTTTCA
AAATGAATTACCAAGTCAATGGTTACCCTAATATGTTTATCACCCCGGAA
GAAGCTATTCGTACGTTGTTGCGTGGATTGGCTTTGATGTAGAGGGCTG
TCATGCAACTAGAGATGCTGTGGGTACTAACCCTACCTCTCCAGCTAGGAT
TTCTACAGGTGTTAACTTAGTAGCTGTACCAGCTGGTTATGTTGACACT
GAAAATAACACAGAATTCACCAGAGTTAATGCAAAACCTCCACCAGGTGA
CCAGTTTAAACATCTTATACCACCTCATGTATAAAGGCTTGCCCTGGAATG
TAGTGCGTATTAAGATAGTACAAATGCTCAGTGATACACTGAAAGGATTG
TCAGACAGAGTCTGTTGCTCCTTTGGGGCGCATGGCTTTGAGCTTACATC
AATGAAGTACTTTGTC AAGATTGGACCTGAAAGAACGTGTTGCTGTGTG
ACAAACGTGCAACTTGCTTTTCTACTTCATCAGATACTTATGCCTGCTGG
AATCATTCTGTGGGTTT GACTATGTCTATAACCCATTTATGATTGATGT
TCAGCAGTGGGGCTT TACGGGTAACCTTCAGAGTAACCATGACCAACATT
GCCAGGTACATGGAAAATGCACATGTGGCTAGTTGTGATGCTATCATGACT
AGATGTTTAGCAGTCCATGAGTGTCTTGTTAAGCGGTTGATTGGTCTGT
TGAATACCCTATTTATAGGAGATGAACTGAGGGTTAATCTGCTTGCAGAA
AAGTACAACACATGGTTGTGAAGTCTGCATTGCTTGCTGATAAGTTTCCA
GTTCTTCATGACATTGGAATCCAAAGGCTATCAAGTGTGTGCCTCAGGC
TGAAGTAGAATGGAAGTTCACGATGCTCAGCCATGTAGTGACAAAGCTT
ACAAAATAGAGGAAC TCTTCTATTCTTATGCTACACATCAGATAAATTC
ACTGATGGTGTGTTGTTGTTTGGAAATGTAACGTTGATCGTTACCCAGC
CAATGCAATTGTGTGTAGGTTTGACACAAGAGTCTTGTCAAAC TTGAAC
TACCAGGCTGTGATGGTGGTAGTTTGTATGTGAATAAGCATGCATTCCAC
ACTCCAGCTTTCGATAAAAAGTGCATTTACTAATTTAAAGCAATTGCCTTT
CTTTTACTATTCTGATAGTCCCTTGTGAGTCTCATGGCAAACAAGTAGTGT
CGGATATTGATTATGTTCCACTCAAATCTGCTACGTGTATTACACGATGC
AATTTAGGTGGTGTGTTTGCAGACACCATGCAAATGAGTACCAGACAGTA
CTTGATGCATATAATATGATGATTTCTGCTGGATTTAGCCTATGGATTT
ACAAACAATTTGATACTTATAACCTGTGGAATACATTTACCAGGTTACAG
AGTTTAGAAAATGTGGCTTATAATGTGTTAATAAAGGACACTTTGATGG
ACACGCCGGCGAAGCACCTGTTTCCATCATTAATAATGCTGTTTACACAA
AGGTAGATGGTATTGATGTGGAGATCTTTGAAAATAAGACAACACTTCCCT
GTTAATGTTGCATTTGAGCTTTGGGCTAAGCGTAACATTAACCCAGTGCC
AGAGATTAAGATACTCAATAATTTGGGTGTGATATCGCTGCTAATACTG
TAATCTGGGACTACAAAAGAGAAGCCCCAGCACATGTATCTACAATAGGT
GTCTGCACAATGACTGACATTGCCAAGAAACCTACTGAGAGTGTGTTTCT
TTCACCTTACTGCTTGTGTTGATGGTAGAGTGAAGGACAGGTAGACCTTT

FIGURE 11G

TTAGAAACGCCCGTAATGGTGTTTTAATAACAGAAGGTTCAAGGTT
CTAACACCTTCAAAGGGACCAGCACAAGCTAGCGTCAATGGAGTCACATT
AATTGGAGAATCAGTAAAAACACAGTTTAACTACTTTAAGAAAGTAGACG
GCATTATTCAACAGTTGCCTGAAACCTACTTTACTCAGAGCAGAGACTTA
GAGGATTTTAAGCCCAGATCACAATGGAACTGACTTTCFCGAGCTCGC
TATGGATGAATTCATACAGCGATATAAGCTCGAGGGCTATGCCCTCGAAC
ACATCGTTTATGGAGATTTCACTCATGGACAACCTGGCGGTCTTCATTTA
ATGATAGGCTTAGCCAAGCGCTCACAAGATTCACCACTTAAATAGAGGA
TTTTATCCCTATGGACAGCACAGTGAAAAATTACTTCATAACAGATGCGC
AAACAGGTTCAATAATGTGTGTCTGTGATTGATCTTTACTTGAT
GACTTTGTGAGATAATAAAGTCACAAGATTTGTCAGTGATTTCAAAGT
GGTCAAGGTTACAATTGACTATGCTGAAATTCATTCATGCTTTGGTGTA
AGGATGGACATGTTGAAACCTTCTACCCAAAACCTACAAGCAAGTCAAGCG
TGGCAACCAGGTGTTGCGATGCCCTAACCCTGTACAAGATGCAAAGAATGCT
TCTTGAAAAGTGTGACCTTCAGAATTATGGTGAAAATGCTGTTATACCAA
AAGGAATAATGATGAATGTGCGCAAAGTATACTCAACTGTGTCAATACTTA
AATACACTTACTTTAGCTGTACCTTACAACATGAGAGTTATTCACPTTGG
TGCTGGCTCTGATAAAGGAGTTGCACCAGGTACAGCTGTGCTCAGACAAT
GGTTGCCAACTGGCACACTACTGTGCGATTAGATCTTAATGACTTCGCTC
TCGACGCAGATCTACTTTAATGGAGACTGTGCAACAGTACATACGGC
TAATAAATGGGACCTTATTATTAGCGATATGTATGACCCTAGGACCAAAC
ATGTGACAAAAGAGAATGACTCTAAAGAAGGGTTTTTCACTTATCTGTGT
GGATTTATAAAGCAAAAACCTAGCCCTGGGTGGTTCTATAGCTGTAAAGAT
AACAGAGCATCTTGGAAATGCTGACCTTTACAAGCTTATGGGCCATTTCT
CATGGTGGACAGCTTTTGTACAAATGTAATGCATCATCGGAAGCA
TTTTTAATTTGGGGTAACTATCTTGGCAAGCCGAAGGAACAAATGATGG
CTATACCATGCATGCTAACTACATTTTCTGGAGGAACACAAATCCTATCC
AGTGTCTTCTTACTCTTTGACATGAGCAAATTTCTCTTAAATTA
AGAGAACTGCTGTAATGTCTCTTAAGGAGAATCAATCAATGATATGAT
TTATTTCTTCTGGAAAAGGTAGGCTTATCATTAGAGAAAACAACAGAG
TTGTGGTTTCAAGTGATATTCTTGTAAACAACTAAACGAACATGTTTAT
TCTTATTATTCTTACTCTCAGTAGTGGTAGTGACCTTGACCGGTGCAC
CACTTTTGATGATGTTCAAGCTCCTAATTACACTCAACATACTTCATCTA
TGAGGGGGGTTTACTATCCTGATGAAATTTTAGATCAGACACTCTTTAT
TTAATCAGGATTTATTTCTTCCATTTTATTCTAATGTTACAGGGTTTCA
TACTATTAATCATACTTTGGCAACCCTGTACATACCTTTTAAAGGATGGTA
TTTATTTTGGCTGCCACAGAGAAATCAAATGTTGTCCGTGGTTGGGTTTTT
GGTTCTACCATGAACAACAAGTCACAGTCGGTGATTTATTAACAATTC
TACTAATGTTGTTATACGAGCATGTAACCTTGAATGTTGTGACAACCCTT
TCTTTGCTGTTTCTAAACCATGGGTACACAGACACATACTATGATATTC
GATAATGCATTTAATGCACTTTCCAGTACATATCTGATGCCCTTTTCGCT
TGATGTTTCAGAAAAGTCAGGTAATTTTAAACACTTACGAGAGTTTGTGT
TTAAAAATAAAGATGGGTTTCTCTATGTTTATAAGGGCTATCAACCTATA
GATGTAGTTCGTGATCTACCTTCTGGTTTAAACACTTTGAAACCTATTTT
TAAGTTGCCCTCTTGGTATTAACATTACAATTTTAGAGCCATTTCTACAG
CCTTTTCACCTGCTCAAGACATTTGGGGCACGTGAGCTGCAGCCTATTTT
GTTGGCTATTTAAAGCCAACTACATTTATGCTCAAGTATGATGAAAATGG
TACAATCACAGATGCTGTGATGTTCTCAAATCCACTGCTGAACTCA
AATGCTCTGTTAAGAGCTTTGAGATTGACAAAAGGAATTTACCAGACCTCT
AATTCAGGGTTGTTCCCTCAGGAGATGTTGTGAGATTCCTAATATTAC
AAACTTGTCTCTTTTGGAGAGGTTTTTAAATGCTACTAAATTCCTTCTG
TCTATGCATGGGAGAGAAAAAAATTTCTAATGTTGTGCTGATTACTCT
GTGCTCTACAACCTAACATTTTTTCAACCTTAAAGTGTATGGCGTTTC
TGCCACTAAGTTGAATGATCTTTGCTTCTCCAATGTCTATGCAGATTCTT
TTGTAGTCAAGGGAGATGATGTAAGACAAATAGCGCCAGGACAAACTGGT
GTTATTGCTGATTATAATTATAAATGCCAGATGATTTTATGGGTTGTGT
CCTTGCCTTGAATACTAGGAACATTGATGCTACTTCAACTGGTAATTATA

FIGURE 11H

ATTATAAATATAGGTATCTTAGACATGGCAAAGCTTAGGCCCTTTGAGAGA
GACATATCTAATGTGCCTTTCTCCCCTGATGGCAAACCTTGCACCCACC
TGCTCTTAATTTGTTATTTGGCCATTAATGATTATGGTTTTTACACCACTA
CTGGCATTTGGCTACCAACCTTACAGAGTTGTAGTACTTTCTTTGAACTT
TTAAATGCACCGGCCACGGTTTGTGGACCAAAATTATCCACTGACCTTAT
TAAGAACCAGTGTGTCAATTTAATTTAATGGACTCACTGGTACTGGTG
TGTTAACCTCTTTCAAAGAGATTTCAACCATTTCAACAATTTGGCCGT
GATGTTTTCTGATTTCACTGATTCGGTTTCGAGATCCTAAAACATCTGAAAT
ATTAGACATTTCACTTGGCGCTTTTGGGGGTGTAAGTGTAAATTACACCTG
GAACAAATGCTTCATCTGAAGTTGCTGTTCTATATCAAGATGTTAACTGC
ACTGATGTTTCTACAGCAATTCATGCAGATCAACTCACACCAGCTTGGCG
CATATATTTCTACTGAAACAATGTATTCCAGACTCAAGCAGGCTGTCTTA
TAGGAGCTGAGCATGTGCACACTTCTTATGAGTGCACATTCCTATTGGA
GCTGGCATTTGTGCTAGTTACCATACAGTTTCTTTATTACGTAGTACTAG
CCAAAAATCTATTGTGGCTTATACTATGTCTTTAGGTGCTGATAGTTCAA
TTGCTTACTCTAATAACACCATTGCTATACTACTAACTTTTCAATTAGC
ATTACTACAGAAGTAATGCCCTGTTTCTATGGCTAAAACCTCCGTAGATTG
TAATATGTACATCTGCGGAGATTTCTACTGAATGTGCTAATTTGCTTCTCC
AATATGGTAGCTTTTGCACACAACCTAAATCGTGCCTCTCAGGTATTGCT
GCTGAACAGGATCGCAACACACCTGAAGTGTTCGCTCAAGTCAAACAAAT
GTACAAAACCCCAACTTTGAAATATTTTGGTGGTTTTAATTTTTTCACAAA
TATFACCTGACCCTCTAAAGCCAACCTAAGAGGCTTTTATTGAGGACTTG
CTCTTTAATAAGGTGACACTCGCTGATGCTGGCTTCATGAAGCAATATGG
CGAATGCCTAGGTGATATTAATGCTAGAGATCTCATTTGTGCGCAGAAAGT
TCAATGGACTTACAGTGTGGCCACCTCTGCTCACTGATGATATGATTGCT
GCCTACACTGCTGCTCTAGTTAGTGGTACTGCCACTGCTGGATGGACATT
TGGTGTGGCGCTGCTCTTCAAATACCTTTTGTATGCAAATGGCATAATA
GGTTCAATGGCATTTGGAGTTACCCAAAATGTTTCTCTATGAGAACCAAAA
CAAAATCGCCAACCAATTTAACAAGGGGATTAGTCAAATTCAGAATCACT
TACAACAACATCAACTGCATTGGGCAAGCTGCAAGACGTTGTTAACCAGA
ATGCTCAAGCATTAACACACTTGTAAACAACCTTAGCTCTAATTTTGGT
GCAATTTCAAGTGTGCTAAATGATATCTTTTCGCGACTTGATAAAGTCGA
GGCGGAGGTACAAATGACAGGTTAATTTACAGGCAGACTTCAAAGCCTTC
AAACCTATGTAACACAACAACCTAATCAGGGCTGCTGAAATCAGGGCTTCT
GCTAATCTTGCTGTACTAAAATGTCTGAGTGTGTTCTTTGGACAATCAA
AAGAGTTGACTTTTGTGGAAAGGGCTACCACCTTATGTCTTCCCACAAG
CAGCCCCGCATGGTGTGTTCTTCTACATGTCACGTATGTGCCATCCCAG
GAGAGGAACTTACCACAGCGCCAGCAATTTGTTCATGAAGGCAAAGCATA
CTTCCCCTGGAAGGTGTTTTTGTGTTAATGGCACTTCTTGGTTTATTA
CACAGAGAACTTCTTTCTCCACAATAATTTACTACAGACAATAACATTT
GTCTCAGGAAATTTGTGATGTCGTTATTGGCATCATTAAACAACACAGTTTA
TGATCTCTGCAACCTGAGCTTGACTCATTCAAAGAAGAGCTGGACAAGT
ACTTCAAAAATCATAACATCACCAGATGTTGATCTTGGCGACATTTTCAGGC
ATTAACGCTTCTGTGCTCAACATTCAAAAGAAAATTGACCGCTCAATGA
GGTCGCTAAAAATTTAAATGAATCACTCATTTGACCTTCAAGAATTTGGAA
AATATGAGCAATATATTAATGGCCTTGGTATGTTTGGCTCGGCTTCATTT
GCTGGACTAATTTGCCATCGTTCATGGTTACAATCTTGTCTTTGTGTCATGAC
TAGTTGTGTCAGTTGCCTCAAGGGTGCATGCTCTTGTGGTTCTTTGCTGCA
AGTTTGTATGAGGATGACTCTGAGCCAGTTCTCAAGGGTGTCAAATTACAT
TACACATAAACGAACTTATGGATTTGTTTATGAGATTTTTTACTCTTAGA
TCAATTTACTGCACAGCCAGTAAAAATGACAATGCTTCTCCTGCAAGTAC
TGTTTCATGCTACAGCAACGATACCGCTACAAGCCTCACTCCCTTTCGGAT
GGCTTGTATTGGCGTTGCATTTCTTGTCTTTTTTTCAGAGCGCTACCAAA
ATAATTTGGCTCAATAAAAGATGGCAGCTAGCCCTTTATAAGGGCTTCCA
GTTCAATTTGCAATTTACTGCTGCTATTTGTTACCATCTATTACATCTTT
TGCTTGTGCTGCTGAGGATGGAGGCGCAATTTTTGTACCTCTATGCCTTG
ATATATTTTCTACAATGCATCAACGCATGTAGAAATATTATGAGATGTTG

FIGURE 11I

GCTTTGTTGGAAGTGCAAATCCAAGAACCATTACTTTATGATGCCAACT
ACTTTGTTTGC TGGCACACACATAACTATGACTACTGTATACCATATAAC
AGTGTACAGATACAATTGTCTTACTGAAGGTGACGGCATTTC AACACC
AAA ACTCAAAGAAGACTACCAAATGGTGGTTATTCTGAGGATAGGCACT
CAGGTGTTAAAGACTATGTCTGTTGTACATGGCTATTTACC CGAAGTTTAC
TACCAGCTTGAGTCTACACAAATTACTACAGACACTGGTATTGAAAATGC
TACATTCTTCATCTTTAACAAGCTTGTAAAGACCCACCGAATGTGCAAA
TACACACAATCGACGGCTCTTCAGGAGTTGCTAATCCAGCAATGGATCCA
ATTTATGATGAGCCGACGACGACTACTAGCGTGCCTTTGTAAGCACAAGA
AAGTGAGTACGAAC TTATGTACTCATTCGTTTCGGAAGAAACAGGTACGT
TAATAGTTAATAGCGTACTTCTTTTTCTTGC TTTTCGTGGTATTCTTGCTA
GTCACACTAGCCATCCTTACTGCGCTTCGATTGTGTGCGTACTGCTGCAA
TATTGTTAACGTGAGTTTAGTAAAACCAACGGTTACGTCTACTCGCGTG
TTAAAAATCTGAAC TCTTCTGAAGGAGTTCCTGATCTTCTGGTCTAAACG
AACTAACTATTATTATTATTCTGTTTGGAACTTTAACATTGCTTATCATG
GCAGACAACGGTACTATTACCGTTGAGGAGCTTAAACAACCTCCTGGAACA
ATGGAACTTAGTAATAGGTTTCC TATTCTTAGCCTGGATTATGTTACTAC
AATTTGCC TATTCTAATCGGAACAGGTTTTTGTACATAATAAAGCTTGTT
TTCCCTCGGCTCTTGTGGCCAGTAACTTGC TTTGTTTGTGCTTGCTG
TGCTACAGAATTAATTGGGTGACTGGCGGATTGCGATTGCAATGGCTT
GTATTGTAGGCTTGATGTGGCTTAGCTACTTCGTTGCTTTCCTTCAGGCTG
TTTGCTCGTACCCGCTCAATGTGGTCATTCAACCCAGAAAACAAACATTC
TCTCAATGTGCCTCTCCGGGGACAATTGTGACCAGACCGCTCATGGAAA
GTGAAC TTGTCATTGGTGC TGTGATCATTCGTGGTCACTTGC GAATGGCC
GGACACTCCCTAGGGCGCTGTGACATTAAGGACCTGCCAAAAGAGATCAC
TGTGGCTACATCACGAACGCTTCTTATTACAAATTAGGAGCGTCGCAGC
GTAGAGCACTGATT CAGGTTTTGCTGCATACAACCGCTACCGTATTGGA
AACTATAAATTAATAACAGACCACGCCGGTAGCAACGACAATATTGCTTT
GCTAGTACAGTAAGTGACAACAGATGTTTCATCTTGTGACTTCCAGGTT
ACAATAGCAGAGATATTGATTATCATTATGAGGACTTTCAGGATTGCTAT
TTGGAATCTTGACGTTATAATAAGTTCAATAGTGAGACAATATTTAAGC
CTCTAACTAAGAAGAATTATTCCGGAGTTAGATGATGAAGAACCTATGGAG
TTAGATTATCCATAAAAACGAACATGAAAATATTCTCTTCCTGACATTGA
TTGTATTACATCTTGCAGCTATATCACTATCAGGAGTGTGTAGAGGT
ACGACTGTACTACTAAAAGAACC TTGCCATCAGGAACATACGAGGGCAA
TTCACCA TTTACCCTCTTGCTGACAATAAATTTGC ACTAACTTGC ACTA
GCACACACTTTGCTTTTGCTTGTGCTGACGGTACTCGACATACCTATCAG
CTGCGTGCAAGATCAGTTTACC AAAACTTTTCATCAGACAAGAGGAGGT
TCAACAAGAGCTCTACTCGCCACTTTTTCTCATTGTTGCTGCTCTAGTAT
TTTTAATACTTTGCTTACCATTAAAGAGAAAAGACAGAATGAATGAGCTCA
CTTAATTGACTTCTATTGTGCTTTTTAGCCTTTCTGCTATTCTCTGT
TTAATAATGC TTAATTATATTTGGTTTTCACTCGAAATCCAGGATCTAGA
AGAACCTTGTACCAAAGTCTAAACGAACATGAAACTTCTCATTTGTTTGA
CTTGTATTTCTCTATGCAGTTGCATATGCACTGTAGTACAGCGCTGTGCA
TCTAATAAACCTCATGTGCTTGAAGATCCTTGTAAAGGTACAACACTAGGG
GTAATACTTATAGCACTGCTTGGCTTTGTGCTCTAGGAAAGGTTTTACCT
TTTCATAGATGGCACACTATGGTTCAAACATGCACACCTAATGTTACTAT
CAACTGTCAAGATCCAGCTGGTGGTGCCTTATAGCTAGGTGTTGGTACC
TTCATGAAGGTACCCAACTGCTGCATTTAGAGACGTACTTGTGTTTAA
AATAAACGAACAAATTA AAAATGTCTGATAATGGACCCCAATCAAACCAAC
GTAGTGCCCCCGCATTACATTTGGTGGACCCACAGATTCAACTGACAAT
AACCAGAATGGAGGACGCAATGGGGCAAGGCCAAAACAGCGCCGACCCCA
AGGTTTACCCAATAATAC TGCGTCTTGGTTACAGCTCTCACTCAGCATG
GCAAGGAGGAACCTTAGATTCCCTCGAGGCCAGGGCGTTCCAATCAACACC
AATAGTGGTCCAGATGACCAAA TTGGCTACTACCGAAGAGCTACCCGACG
AGTTCGTGGTGGTGACGGCAAAATGAAAGAGCTCAGCCCCAGATGGTACT
TCTATTACCTAGGAAC TGCCCCAGAAGCTTCACTTCCCTACGGCGCTAAC

FIGURE 11J

AAAGAAGGCATCGTATGGGTTGCAACTGAGGGAGCCTTGAATACACCCAA
AGACCACATTGGCACCCGCAATCCTAATAACAATGCTGCCACCGTGCTAC
AACTTCCTCAAGGAACAACATTGCCAAAAGGCTTCTACGCAGAGGGAAGC
AGAGGCGGCAGTCAAGCCTCTTCTCGCTCCTCATCACGTAGTCGCGGTAA
TTCAAGAAATTCAACTCCTGGCAGCAGTAGGGGAAATTCTCCTGCTCGAA
TGGCTAGCGGAGGTGGTGAAACTGCCCTCGCGCTATTGCTGCTAGACAGA
TTGAACCAGCTTGAGAGCAAAGTTTCTGGTAAAGGCCAACAAACAACAAAGG
CCAAACTGTCTACTAAGAAATCTGCTGCTGAGGCATCTAAAAGCCTCGCC
AAAAACGTACTGCCACAAAACAGTACAACGTCACTCAAGCATTGCGGAGA
CGTGGTCCAGAACAACCCAAGGAAATTCGGGGACCAAGACCTAATCAG
ACAAGGAAGTATTACAAACATTGGCCGCAAATGCACAATTTGCTCCAA
GTGCCTCTGCATTCTTTGGAATGTCACGCATTGGCATGGAAGTCACACCT
TCGGGAACATGGCTGACTTATCATGGAGCCATTAATTTGGATGACAAAGA
TCCACAATTCAAAGACAACGTCATACTGCTGAACAAGCACATTGACGCAT
ACAAAACATTCCCACCAACAGAGCCTAAAAGGACAAAAGAAAAGACT
GATGAAGCTCAGCCTTTGCCGCAGAGACAAAAGAAGCAGCCCACTGTGAC
TCTTCTTCCCTGCGGCTGACATGGATGATTTCTCCAGACAACCTCAAAATT
CCATGAGTGGAGCTTCTGCTGATTCAACTCAGGCATAAAACTCATGATG
ACCACACAAGGCAGATGGGCTATGTAAACGTTTTTCGCAATTCGGTTTACG
ATACATAGTCTACTCTTGTGCAGAATGAATTCGTAACATAACAGCACA
AGTAGGTTTAGTTAACTTTAATCTCACATAGCAATCTTTAATCAATGTGT
AACATTAGGGAGGACTTGAAAGAGCCACCACATTTTCATCGAGGCCACGC
GGAGTACGATCGAGGGTACAGTGAATAATGCTAGGGAGAGCTGCCTATAT
GGAGAGCCCTAATGTGTAAAATTAATTTTAGTAGTGTATCCCCATGTG
ATTTTAATAGCTTCTTAGGAGAATGACAAAAA

GenBank Accession No. AY274119.3, SEQ ID NO: 15

FIGURE 11K

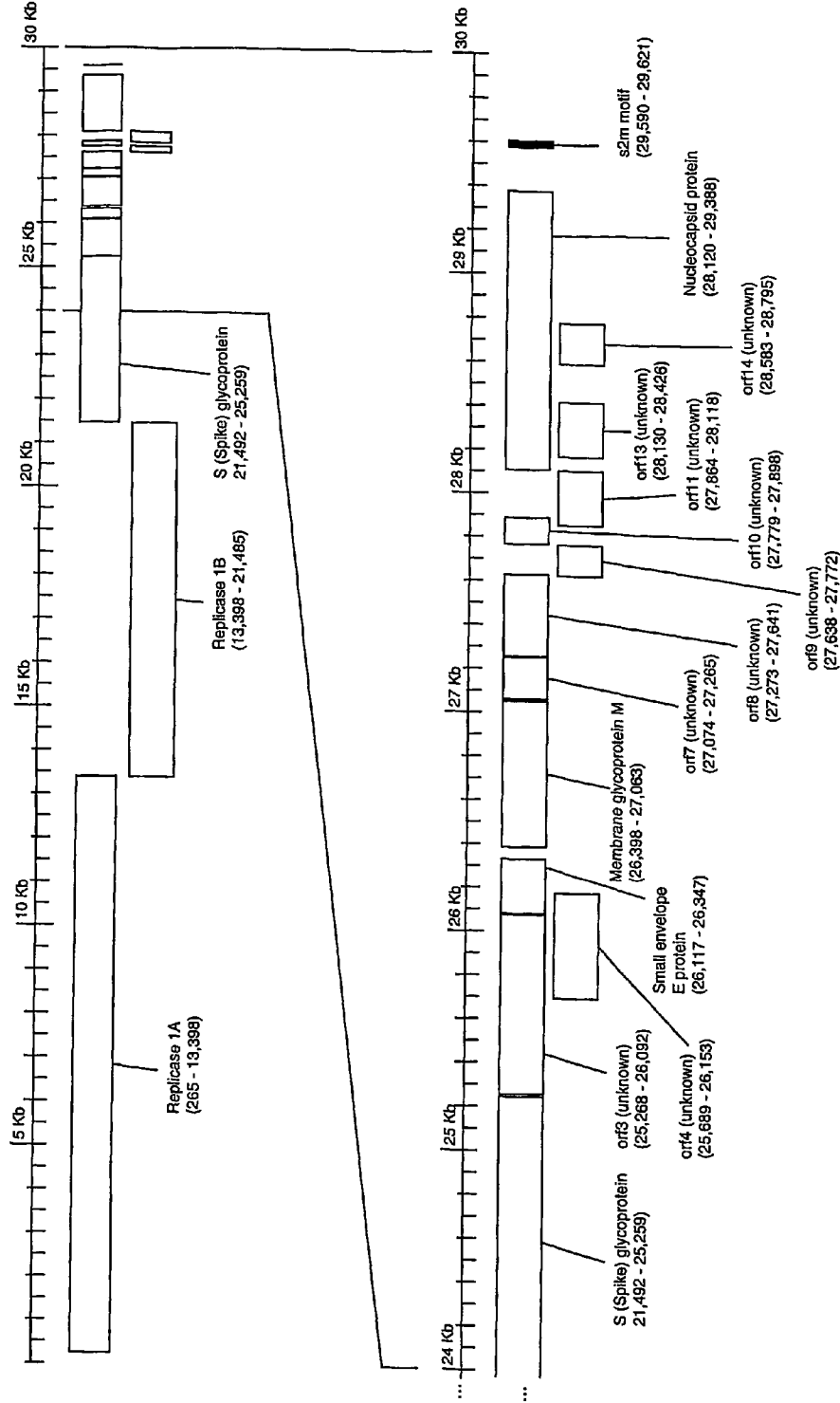


Figure 12

Replicase 1A

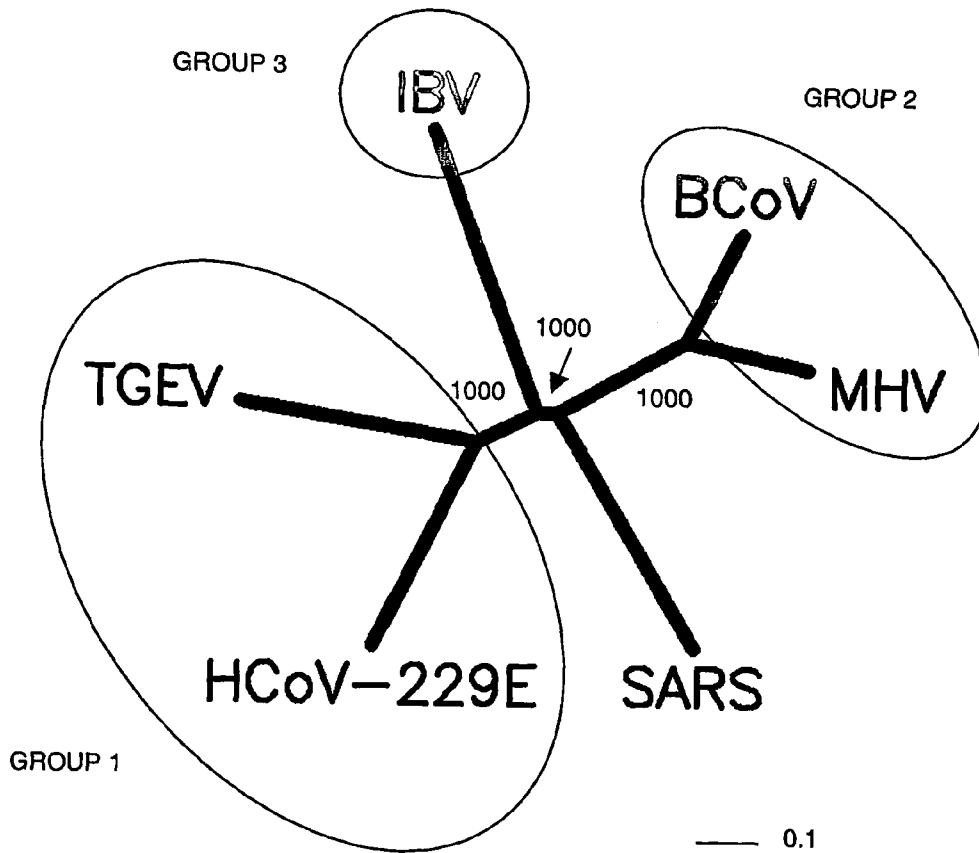


Figure 13A

Membrane Glycoprotein

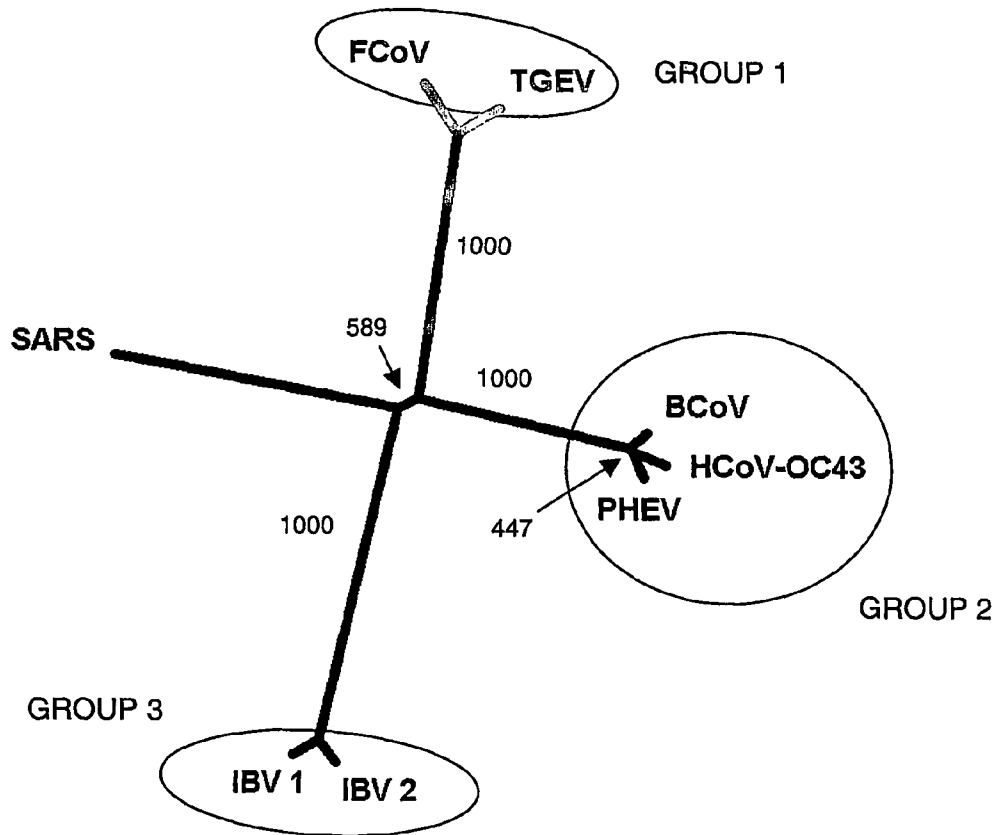


Figure 13B

Nucleocapsid

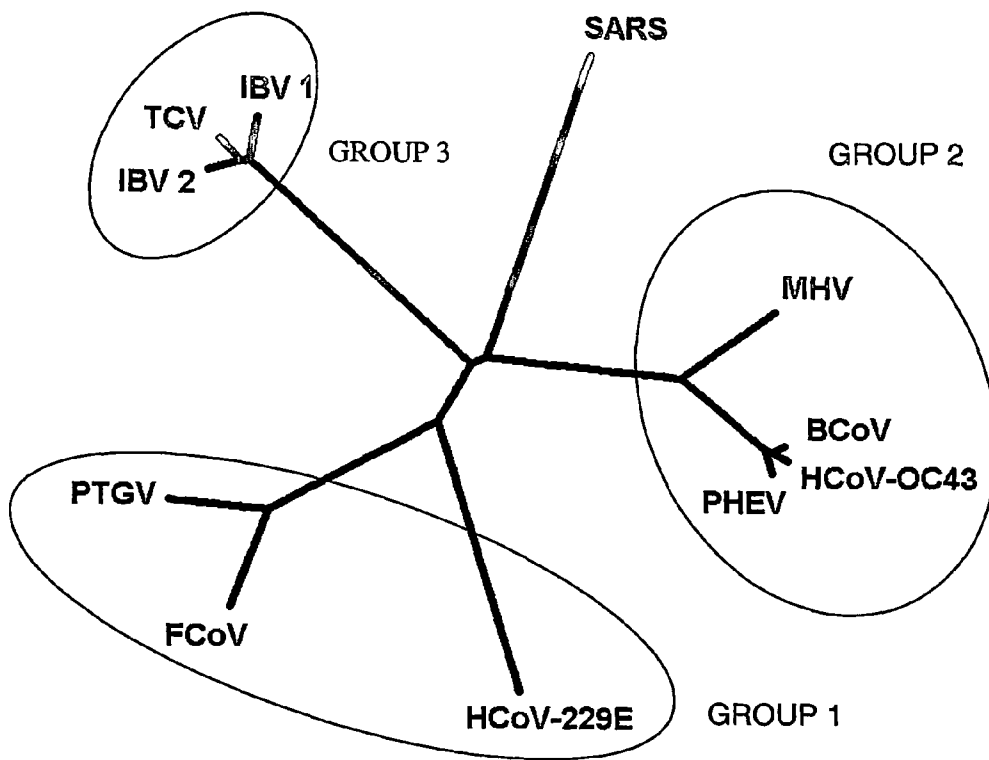


Figure 13C

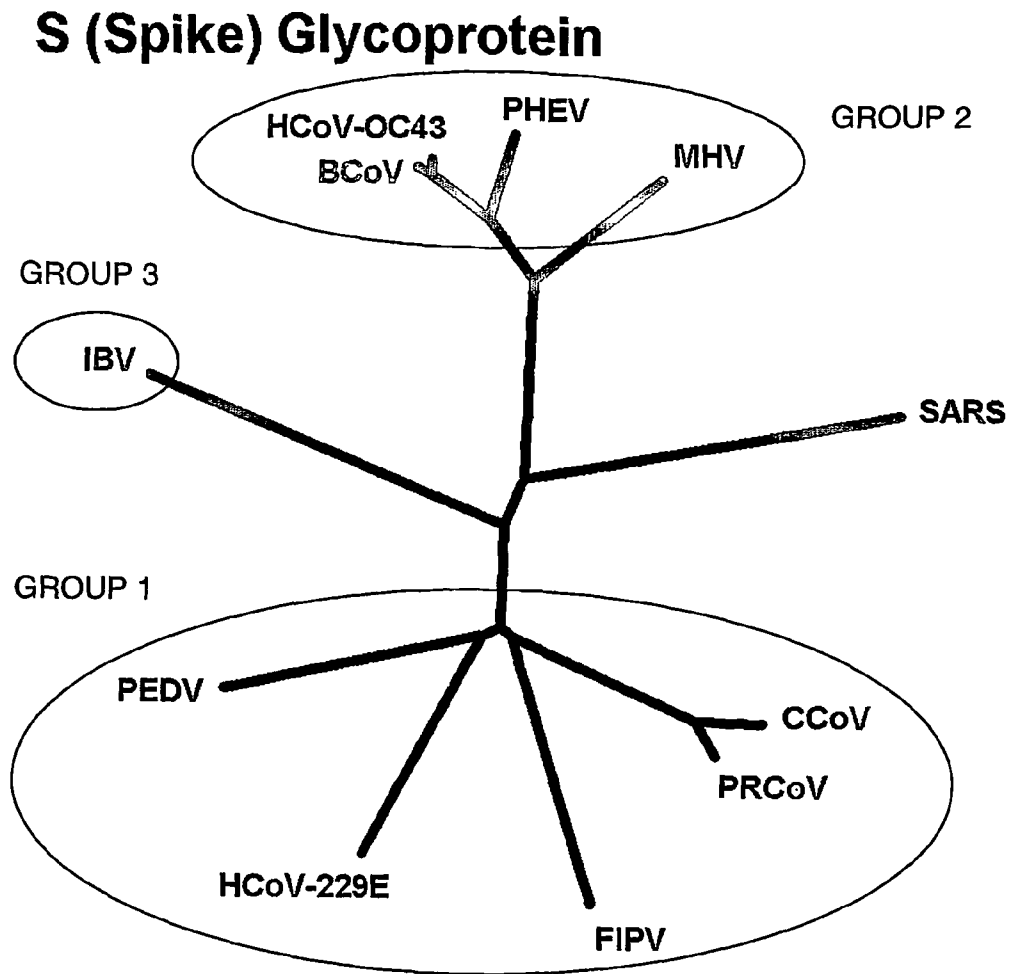


Figure 13D

229E
 PEDV -----MRSLIYFWLLLPVLP T L S L P Q D V T R C Q S T T-----NFRRF F S K F N V Q A P A
 CCov MIVLILCLLLFSYNSVICTSNND CV Q G N V T Q L P G N E-----N I K D F L F H T F K E E P
 PRC -----
 FICV -----MIF I I L T L L S V A K S E D A P H G V T L P Q F N T S H N N E R F E L N F Y N F L Q T W D I P P N T
 BoCov -----
 OC43 -----M F L I L L I S L P M A
 PHEV -----M F L I L L I S L P M A
 MHV -----M F F I L L I S L P S A
 TOR2_S -----M L F V F I L L L P S C
 AIBV -----

229E
 PEDV -----V V V L G G Y L P-----S M N S S S W Y C G T G I E T A S G V H G I F L S Y I D S G Q G F E
 CCov S V V V G G Y P T E-----V W Y N C S R S A T T A Y K D F S N I H A F Y F D M E A M E N S T G
 PRC -----
 FICV -----E T I L G G Y L P Y C G A G V N C G W Y N F S Q S V G Q N G K Y A Y I N T Q N L N I P N V H G V Y F D V R E H N D G E
 BoCov F A V I G D L K C T-----T V S I N D V D T G A P S I S T D I V D V T N G L G
 OC43 L A V I G D L K C T-----T V A I N D V D T G V P S T S T D I V D V T N G L G
 PHEV F A V I G D L K C T-----T S L I N D V D T G V P S I S S E V D V T N G L G
 MHV L G Y I G D F R C I Q-----T V M Y N G N N A S A P S I S T E A V D V S K G R G
 TOR2_S -----M F I F L L F L T L T S G S D L D R C T T F D D V Q A P N Y T Q
 AIBV -----

229E
 PEDV -----I G-----I S Q E P F D P S G Y Q L Y L H K A T N G N T N A T A R L R-----I C Q F P D N K T L G P T V N D V T G -
 CCov N A R G K P L L V H V H G D P V S I I I Y I S A Y R D D V Q P R P L L K H G L L C I T K N K I I D Y N T F T S A Q W S -
 PRC -----
 FICV -----W D D R D K V G L L I A I H G N S K Y S L L M V L Q D A V E A N Q P H V A V K I C H W K P G N I S S Y H A F S V N L G D
 BoCov T Y-----Y V L D R V Y L N T T L L L N G Y Y P T S G S T Y R N M A L K G T L L L S R L W F K P P F L S D F I N G -
 OC43 T Y-----Y V L D R V Y L N T T L L L N G Y Y P T S G S T Y R N M A L K G T L L L S R L W F K P P F L S D F I N G -
 PHEV T F-----Y V L D R V Y L N T T L L L N G Y Y P I S G A T F R N M A L K G T R L L S T L W F K P P F L S E F N D G -
 MHV T Y-----Y V L D R V Y L N A T L L L T G Y Y P V D G S N Y R N L A L T G T N T L S L T W F K P P F L S E F N D G -
 TOR2_S H T-----S S M R G V Y Y P D E I F R S D T L Y L T Q D L F L P F Y S N V T G F H T I N H T F G N P V I P F K D G -
 AIBV -----

229E
 PEDV -----R N C L F N K A I P---A Y M R D G K D I V V G I T W D N D R V T-V F A D K I Y H F Y L K N D W S R-----
 CCov -A I C L G D D R K I P F S V I P T D N G T K I F G L E W N D D Y V T A Y I S D R S H H L N I N N M W F N N V T I L Y S
 PRC -----
 FICV -----G G Q C V F N Q R F S---L D T V L T M N D F Y G F Q W E D T Y V D I Y L G G T I T K V W D N D W S I V E A S ---
 BoCov ---I F A K V K N---T K V I K K G V M Y S E F P A I T I G S T F V N T S Y S V V V Q P H T T N-----
 OC43 ---I F A K V K N---T K V I K H G V M Y S E F P A I T I G S T F V N T S Y S V V V Q P H T T N-----
 PHEV ---I F A K V K N---S R F S K D G V I Y S E F P A I T I G S T F V N T S Y S I V V E P H T S L-----
 MHV ---I F A K V Q N---L K T N T P T G A T S Y F P T I V I G S L F G N T S Y T V V L E P Y N N-----
 TOR2_S ---I Y F A A T E K---S N V V R G W V F G S T M N K S Q S V I I I N N S T N V V I R A C N F E L C D N---
 AIBV -----M L G K S L F L V T I L C A L C S A N L F D P A N Y V Y Y Y Q S A F R P-----

229E
 PEDV -----M F V L L V A Y A L L H I A G C Q T T N G L N--T S Y S V C N G---C V G Y S E N V F A V E S
 CCov --V A T R C Y N R R S C A M Q Y V Y T P T Y Y M L N V T S A G E D G- I Y Y E P C T A N--C T G Y A A N V F A T D S
 PRC R S S A T W Q K S A A Y V Y Q G V S N F T Y Y K L N N T N G L K S---Y E L C E D Y E Y C T G Y A T N V F A P T V
 FICV -----M K K L F V V L V V M P L I Y G D K F P T S V V S N-----C T D--Q C A S Y A N V P T T Q P
 BoCov -I S Y H W N R I N Y G Y M Q F V N R T T Y Y A Y N N T P G G A N Y T Q L Q L S E C H T D- Y C A G Y A K N V F V P- I
 OC43 -L D N K L Q G L L E I S V C Q Y T M C E Y P H T I C H E P K L- G N K R V E L W H W D T G V V S C L Y K R N F T Y D V N
 PHEV -L D N K L Q G L L E I S V C Q Y T M C E Y P N T I C H E P N L- G N R R V E L W H W D T G V V S C L Y K R N F T Y D V N
 MHV -I N G N L Q G L L Q I S V C Q Y T M C E Y P H T I C H E P N L- G N Q R I E L W H Y D T D V V S C L Y R R N F T Y D V N
 TOR2_S -----I I M A S V C T Y T I C Q L P Y T P C K P N T N G N R V I G F W H T D V K P P I C L L K R N F T F N V N
 AIBV ---P F F A V S K P M G T Q T H T M I F D N A F N C T F E Y I S D A F S L D V S E K S G N F K H L R E F V F K N K D G
 ---S N G W H L Q G G A Y A V V N S S N Y A N N A G S A S E C T---V G V I K D V Y N Q S A A S I A M T A P L Q G

FIGURE 14A

229E GGYIPSDFAFNN--WFLLTNTSSVVDGVVRSFQPLLLNCLWSVSGLRFTTGFVYFNGTGR
 PEDV NGHIPBGFSEFNN--WFLLSNDSTLLHGKVVSNQPLLNVNCLLAI PKIYGLGQFFSFNHTMD
 CCov GGYIPHGFSFNN--WFMRTNSSTFVSGRFVTNQPLLNVNCLWVPVSPFGVAAQQFCFEGAQF
 PRC GGFIPSDFSFNN--WFLLTNSSTLVSGKLVTKQPLLNVNCLWVPVSPFEAASTFCFEGADF
 FICV DGKIPEDFSFSN--WFLLSDKSTLVQGRVLSQPVFVQCLRPVPSWSNNTAVVHFKN-D
 BoCov ADYLYPHFYQEGGTFYAYFTDTGVVTKFLFNVLGTVLSHYVVLPLTCS----SAMTLEY
 OC43 ADYLYPHFYQEGGTFYAYFTDTGVVTKFLFNVLGTVLSHYVVMPLTCN----SAMTLEY
 PHEV ADYLYPHFYQEGGTFYAYFTDTGVVTKFLFKLYLGTVLSHYVVMPLTCN----SALSLEY
 MHV APWLYPHFYQGGTFYAYYADKPSATTFVSVYIGDILTQYFVLPFICTPTAGSTLAPLY
 TOR2_S FLYVYKGYQPIDVVRDLPSGFNTLKPFLKPLGINITNFRALLTAFSPAQDIWGTSAAY
 AIBV MAWSKSQFCSAHCDFSEITVVFVTHCYSSGSGSCPITGMIARGHIRISAMKNGSLFYNLTV

229E GDCKGFSDDVLSDVIRYNLN-FEENLRRGT-----ILFKTSYGV-VVFCYTNNT-----
 PEDV GVCNGAAVDRAPEALRFNINDTSVILAEGS-----IVLHTALGTNLSFVCSNSSD-----
 CCov SQCNGVSLNNTVDVIRFNLN-FTALVQSGMGATV-FSLNNTGGVILEISCYNDTVS---E
 PRC DQCNGAVLNNTVDVIRFNLN-FTTNVQSGKGATV-FSLNNTGGVILEISCYNDTVS---D
 FICV AFCP-----NVTADVLRFNLFSDTDVYTDSTNDEQLFFTFEDNTTASIACYSSANVTDFFQ
 BoCov WVTPLTSKQYLLAFNQDGVIFNAVDCKSDFMS---EIKCKTLSIAPSTGVYELNG-----
 OC43 WVTPLTSKQYLLAFNQDGVIFNAVDCKSDFMS---EIKCKTLSIAPSTGVYELNG-----
 PHEV WVTPLTRQFLLAFDQDGVLYHAVDCASDFMS---EIMCKTSSITPPTGVYELNG-----
 MHV WVTPLLRQYLFNFNEKGVITSAVDCASSYIS---EIKCKTQSLLPSTGVYDLG-----
 TOR2_S FVGYLKPTTFMLKYDENGTTITDAVDCSQNPLA---ELKCSVKSFEIDKGIYQTSN-----
 AIBV SVSKYPNEKSFQCVNNTSVVLNGDLVFTSNKTTDVT SAGVYFKAGGPFVNYSIMK-----

229E -LVSGDAHIPFGTVLGNFYCFVNTTIGTETTSAFVGLPKTVREFVISRTGHFYINGYRY
 PEDV -PHLAIFAIPLGATEVPYCYFLKVDTYNSTVYKFLAVLFPSTVREIVITKYGDVYVNGFY
 CCov SSFYSGEISFGVTDGPRYCF A---LYNGTALKYLGLTLPSPVKEIAISKWGHFYINGYF
 PRC SSFSSYGEIPFGVTNGPRYCYV---LYNGTALKYLGLTLPSPVKEIAISKWGHFYINGYF
 FICV PANNSVSHIPFGKT--AHFCFAN-FSHSIVSRQFLGILPPTVREFAFGRDGSIFVNGYKY
 BoCov -YTVQPIADVRRIPNLPDCNIEAWLNDKSVPSPLNWERKTF SNCNFMNSSLMSFIQADS
 OC43 -YTVQPIADVRRIPNLPDCNIEAWLNDKSVPSPLNWERKTF SNCNFMNSSLMSFIQADS
 PHEV -YTVQPVATVYRRIPDLFNCDIEAWLNSKTIVSSPLNWERKIF SNCNFMNMRMLMSFIQADS
 MHV -YTVQPVGVYRRVFNLPDCNIEAWLNTAKSVPSPLNWERRTFQNCNFMNSSLRYVQAES
 TOR2_S -FRVVPDGVVRRFPNITNLCPFGVEFNATKFPSPYAWERRKISNCVADYSVLYNSTFFST
 AIBV -EFKVLAYFVNGTAQDVLCDNSPKGLLACQYNTGNFSDGFYPTNSTLVREKFIYRES

229E FTLGNVEAVNFNVTTAETTD----FFTVALASYADVLVNVQSITIANI IYCNSVINRLRC
 PEDV LHLGLLDVAVTIYPTGHGTDDVSGFWTIASNTNFVDALIEVQGTSIQRILYCDPVSQKLC
 CCov PSTFPIDCISFNLTGDSGA----FWTIAYTSYTDALVQVENTAIKKTIVYCNSHINNICK
 PRC FSTFPIDCISFNLTGDSV----FWTIAYTSYTEALVQVENTAITNVYCN SYVMNICK
 FICV FSLPAIRSVNFSISSVEYEG----FWTIAYTNYTDMVDVNGTAITRLFYCDSPLNRIKC
 BoCov FTCNNIDAAKIYGMCFSSIT----IDKFAPNGRKVDLQGLNGLYLQSFNYRIDTTATSC
 OC43 FTCNNIDAAKIYGMCFSSIT----IDKFAPNGRKVDLQGLNGLYLQSFNYRIDTTATSC
 PHEV FGCNNIDASRLYGMCFGSIT----IDKFAPNSRKVDLQVGKSGYLQSFNYKIDTAVSSC
 MHV LSCNNIDASKVYGMCFGSVS----VDRFAPRSRQIDLQIGNSGFLQTANYKIDTAATSC
 TOR2_S FKCYGVSATKLNLCFNSVY----ADSFVVKGDDVRQIAPGQTGVADIADYNYKLEDDDFMGC
 AIBV SVNTTLALNTFTTNVSNQAQ----PNSGGVHTFHLYQTQTAQSGYFNFLSFLSQFVYKA

229E DQLSPYVPDGFYSTSP--IQSVLPVSVLSL-----VYHKHMFIVLYVDFKPKQ---
 PEDV SQVAFDLDDGFYPISSRNLLSHEQPI SFVTL-----SFNDHSFVNITVSA-----
 CCov SQLTANLQNGFYPVAS--SEVGLVNKSVVLLP-----SFYSHTSVNITIDLGMKR--
 PRC SQLTANLNGFYPVSS--SEVGSVNKSVVLLP-----SFLTHTIVNITIGLGMKR--
 FICV QQLKHELDPGFYSASM--LVKDKLPKTFVTMP-----QFYHWMNVTLHVVLNDTEKK
 BoCov -QLYYNLPAAVSVSRFNPSTWNRFRGFTEQSVFKPQPVGVFTHHDVVYAQHCFKAPKNF
 OC43 -QLYYNLPAAVSVSRFNPSTWNRFRGFTEQSVFKPQPVGVFTHHDVVYAQHCFKAPKNF
 PHEV -QLYYSLPAAVSVTHYNPSSWNRRYGFNNQS-----FGSRGLHDAVYSQQCFNTPTNY
 MHV -QLYYSLPKNNVTINNYNPSWNRRYGFKNVND-----
 TOR2_S -VLAWNTRNIDATSTG----NYNYKYRYLRHG-----
 AIBV SDYMYGSYHPICAFRP--ETINSGLWFNSLS-----

FIGURE 14B

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229E      ---SGGKCFNCYPAGVNITLANFNETKG---PLCVDTSHFT-----TKYVAVYAN
PEDV      ---FGGLSSANLVAS--DTTLNGFSS-----PCVDTRQFTI-----TLFYNVNTNS
CCov      ---SGYGQPIASTLS--NITLPMQDNNTD---VYCI RSNRF SVYF HSTCKSSLWDDVFNS
PRC       ---SGYGQPIASTLS--NITLPMQDNNTD---VYCV RSDQF SVVHSTCKSALWDNVFKR
FICV      YDILLAKAPELAALADVHFELAQANGSVTNVTSLCVQARQLA-----LFYKYTSL
BoCov     --CPCKLDGSLCVGNPGIDAGYKNSGIG---TCPAGTNYLT---CHNAA---QCDC
OC43      --CPCKLDGSLCVGNPGIDAGYKNSGIG---TCPAGTNYLT---CHNAV---QCNC
PHEV      --CPCRT--SQCIG---G-----AGTG---TCPVGTTRK---CFAAVTKATKCTC
MHV       -----
TOR2_S    -----
AIBV      -----

229E      ---VGRWSASINTGNCPFSFGKVN NFVKFSGVCFSLKDIPGG--CAMPIVANWAYS KYTT
PEDV      ---YGYVSKSQDS--NCPFTLQSVNDYLSF SKFCVSTSLLAGA--CTIDLFGYPAFGSGV
CCov      DCTDVL YATAVITKTGTCPFSFDKLN NYLTFNKFCLSLNFVGAN--CKFDVAARTRTNEQVV
PRC       NCTDVL DATAVITKTGTCPFSFDKLN NYLTFNKFCLSLSPVGAN--CKFDVAARTRTNEQVV
FICV      QGLYTYSNLVELQNYDCPFS PQQFN NYLQFETLCFDVNPVAVAG--CKWSLVHDVQWRQTQFA
BoCov     LCTPDPITSKSTGPKYKCPQTKYLVGIGEHCSGLAIKSDYCGGNPCTCQPQAFGLWSVDSC
OC43      LCTPDPITSKSTGPKYKCPQTKYLVGIGEHCSGLAIKSDYCGGNPCTCQPQAFGLWSVDSC
PHEV      WCQPD PSTYKGVNAWTC PQSKVSIQPGQHC PGLGLVEDDCSGNPCTCKPQAFIGWSSETC
MHV       -----
TOR2_S    -----
AIBV      -----

229E      IG---TLYVSWSDGDGITGVPQ--PVEGVSSFMNVTLDKCKTKYNIYDVSGVGVIRVSNDT
PEDV      LT---SLYFQFTKGELITGTPK--PLEGITDVSFMTLDVCTKYTIYGFKEGGIITLNTNS
CCov      R---SLYVIYEEGDNIYGVPS--DNSGLHDL SVLHLD SCTDYNIYGITGVGII RQTNST
PRC       R---SLYVIYEEGDSIVGVPS--DNSGLHDL SVLHLD SCTDYNIYGRTVGII RQTNST
FICV      T---ITVSYKHGSMITTHAKGHSWGFQDTSVLV KDECTDYNIYGFQGTGII RNTTSR
BoCov     LQGDRCNIFANFI PHDVNSGTTTC--STD LQKSN TD IILGVCVNYDLYGITGGIFVEV NAT
OC43      LQGDRCNIFANFI PHDVNSGTTTC--STD LQKSN TD IILGVCVNYDLYGITGGIFVEV NAT
PHEV      LQNGRCNIFANFI LNDVNSGTTTC--STD LQQNTLITDVCVNYDLYGITGGI LIEVNAT
MHV       ---RCQIFANILLNGINSGTTTC--STD LQLPNT EVATGVCVRYDLYGITGGV FKEVKAD
TOR2_S    G---YQPYRVVLSFELLNAPA--TVCGP KLS TD LKNQC VN FNGLTGTGVLTPSSKR
AIBV      L---LVVYTKSDGSRIQTRTEPLVLTQHNYNNTLDKCVAYNIYGRVGGFTINVTDS
          . . . * . . . * * . :

229E      FLN-----GITYTSTSGNLLGFKDVTKGTIYSITPCNP---PDQLVVYQAVVGAM
PEDV      ILA-----GVY YTS DSGQLLAFKNVTS GAVYSVTPCSF---SEQAAVND DVI GVI
CCov      LLS-----GLY YTS LSGDLLGFKNVDGVIY SVTPCDV---SAHAVIDGAI VGAM
PRC       LLS-----GLY YTS LSGDLLGFKNVDGVIY SVTPCDV---SAQAVIDGTTV GAI
FICV      LVA-----GLY YTS I SGDLLAFKNSTTGEI FTVVPCDL---TAQVAVINDEI VGAI
BoCov     YYNS-----WQNL LYDSNGNLYGFRDYL TNRTFMIRSCYSG--RVSAAFHANSSEPAL
OC43      YYNS-----WQNL LYDSNGNLYGFRDYL TNRTFMIRSCYSG--RVSAAFHANSSEPAL
PHEV      YYNS-----WQNL LYDSNGNLYGFRDYL TNRTFMIRSCYSG--RVSAVAFHANSSEPAL
MHV       YYNS-----WQAL LYDVNGNLNGFRDLT TNKTYTIRSCYSG--RVSAAYHKEAPEPAL
TOR2_S    FQP-----FQQFGRDVSDFTDSVRDPKTSEILDISP CAFGGVSVITPGTNASSEVAV
AIBV      VANFSYLADGGLALLDTS GAI DVFVVGSGYGLNYYKVNP CEDVN--QQFVVGSGNIVGIL
          . . . : . * :

229E      LSENFTSY-----GFSNVVELPKFFYASNGTYN-----
PEDV      SSLSNST-----FNNTRELPGFFYHSNDGSN-----
CCov      TSINSELL-----GLTHWTTTPNFYYSIYNYTNERTRGTAID--SND
PRC       TSINSELL-----GLTHWTTTPNFYYSIYNYTNDKTRGTPID--SND
FICV      TAVNQTDLFEFVNNTQARRSRSS TPNFVTSYTMPQFYIITKWNNDTS--S-----
BoCov     LFRNIKCN-----YVFNN TLSRQLQPINYFDSYLGCVVNADN-----STS
OC43      LFRNIKCN-----YVFNN TLSRQLQPINYFDSYLGCVVNADN-----STA
PHEV      MFRNLKCS-----HVFNN TILRQIQLVNYFDSYLGCVVNAYN-----NTA
MHV       LYRNINCS-----YVFNN ISREENPLNYFDSYLGCVVNADN-----RTD
TOR2_S    LYQDVNCT-----DVSTA IHADQLTPAWRIYSTGNVVFQTQAGCLIGA EHV
AIBV      TSRNETGS-----E-QVENQFYVKLTNSSHRRRRS-----IG

229E      -CTDAVLTYS SFGVCADGSI IAVQ----PRNWSYDSVSAIVTANLS-----
PEDV      -CTEPVLVYSNIGVCKSGSIGYV-----PSQYGVKIAPTVTGNIS-----
CCov      VDCEPIITYSNIGVCKNGALVFI-----NVTHSDGDVQPISTGNVT-----
PRC       VGCEPVITYSNIGVCKNGALVFI-----NVTHSDGDVQPISTGNVT-----
FICV      -NCTSAITYSSFAICNTGEIKYVNVTHVEIVDSDIGVIKPVSTGNIS-----
BoCov     SVVQTCDLTVGSGYCVDYSTKRRSR--RAITTG YRFNFEPFTVNSVND SLEPVGGLYEIQ

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FIGURE 14C

OC43 SAVQTCDLTVGSGYCVDYSTKRRSR-RAITTYGRPTNFEPFTVNSVNDLSLEHVGGLYEIQ
 PHEV SAVSTCDLTVGSGYCVDYVTALRSR-RSFTTYGRPTNFEPFAANLVNDSIEPVGGLYEIQ
 MHV EALEFNCNLRMGAGLCVDYKSRRAR-RSVSTGYRLTFPEYMPMLVNDVSVQSVGGLYEMQ
 TOR2_S DTSYECDIPIGAGICASYHTVSLRSTSQKSIVAYTMSLGADSSIAYSNN-----TIA
 AIBV QNVTSQCPYVSYGRFCTEPDGSGLMI----VPEELKQFVAPLLNITES-----VL

229E IPSNWTISVQVEYLQITSTPIVVDCASTYVCNGNVRVCELLKQYTSACKTIEDALRNSARL
 PEDV IPTNFSMSIRTEYLQLYNTPVSVDCATYVCNGNSRCKQLLTQYTAACKTIESALQLSARL
 CCov IPTNFTISVQVEYIQVYTTPVSIDCSRVCNGNPRCNKLLTQYVSACQTI EQALAMGARL
 PRC IPTNFTISVQVEYIQVYTTPVSIDCSRVCNGNPRCNKLLTQYVSACQTI EQALAMGARL
 FICV IPKNFTVAVQAEYIQVQKPVVDCATYVCNGNTHCLKLLTQYTSACQTIENALNLGARL
 BoCov IPSEFTIGNMEEFIQTSSPKVTIDCSAFVCGDYAACKSQLVEYGSFCDNINAILTEVNEL
 OC43 IPSEFTIGNMEEFIQTSSPKVTIDCSAFVCGDCAACKSQLVEYGSFCDNINAILTEVNEL
 PHEV IPSEFTIGNLEEFIQTRSPKVTIDCATFVCGDYAACRQQLAEYGSFCENINAILTEVNEL
 MHV IPTNFTIGHHEEFIQTRAPKVTIDCAAFVCGDNAACRQQLVEYGSFCDNVNAILNEVNNL
 TOR2_S IPTNFSISITTEVMPVSMATSVDCNMYICGDSTECANLLQYGSFCTQLNRLSGIAAE
 AIBV IPNSFNLTVTDEYIQTRMDKVINCLQVCGNSLECRKLFQQYGFVCDNLLSVNSVSVQK
 * : * : : * : * . . : * . * : : *

229E ESADVSEMLTFDKKAFILANVSSF-GD-----YNLSSVIPSP-----LPTSGSR--
 PEDV ESVEVNSMLTISEALQLATISSPFGDG-----YNFTNVLGASVY-----DPASGRV--
 CCov ENMEIDSMFLVSENAKLKASVEAFNSSTETLDPYIKEWENIGGSWLGGLKDLIPSHNSK--
 PRC ENMEVDSMLFVSENAKLKASVEAFNSSTETLDPYITQWENIGGFWEGLKYLPSDNSK--
 FICV ESLMLNDMITVSDRGLLELATVERFNATA-----LGGEKLGGLYFDG---LSSLPPK--
 BoCov LDTTQLQVANS LMNGVTLSTKIKDGVN-----FNVD DINFSPVLG---CLGSACNK--
 OC43 LDTTQLQVANS LMNGVTLSTKIKDGVN-----FNVD DINFSPVLG---CLGSECNK--
 PHEV LDTTQLQVANS LMNGVTLSTKIKDGIN-----FNVD DINFSPVLG---CLGSECNK--
 MHV LDNMQLQVASALMQGVTISSRLPDGIS-----GPIDDINFSPVLG---CIGSTCAEDG
 TOR2_S QDRNTREVFAQVKQMYKTPTLKYFGGF-----NFSQILPDPLKP-----
 AIBV EDMELLSFYSSSTKPKGYDTPVLSNVSTG-----EFNISLTLTTPSSP-----

229E -----VAGRSAIEDILFSKIVTSGLGTVDADYKNCCKGLS--IADLCAQYYNGIMVLP
 PEDV -----VQKRSVIEDLLFNKVVTVNGLGTVDEYKRCNNGRS--VADLVCAQYYSGVMVLP
 CCov -----RKYRSAIEDLLFDKVVTSGLGTVDEYKRCCTGGYD--IADLVCAQYYNGIMVLP
 PRC -----RKYRSAIEDLLFSKVVTSGLGTVDEYKRCCTGGYD--IADLVCAQYYNGIMVLP
 FICV -----IGKRSAVEDLLFNKVVTSGLGTVDDDYKCCSSGTD--VADLVCAQYYNGIMVLP
 BoCov -----VSSRSAIEDLLFSKVKLSDVG-FVEAYNNCTGGAE--IRDLCVQSYNGIKVLP
 OC43 -----VSSRSAIEDLLFSKVRKSDVG-FVEAYNNCTGGAG--IRDLCVQSYNGIKVLP
 PHEV -----ASTRSAIEDLLFDKVKLSDVG-FVQAYNNCTGGAE--IRDLCVQSYNGIKVLP
 MHV NGPSAIRGRSAIEDLLFDKVKLSDVG-FVEAYNNCTGGQE--VRDLVCVQSFNGIKVLP
 TOR2_S -----TKRSFIEDLLFNKVTADAG-FMKQYGECLGDIN--ARDLCAQKFNGLTVLPP
 AIBV -----SGRSFVEDLLFTSVEVTVGLP-TDAEYKCKTAGPLGTLKDLICAREYNGLVLP
 * : * : * . . : * . * : : *

229E VADAERMAMYTGSLIGGIALGGIT---SAVSIPFSLAIQARLNYVALQTDVQLQENQKIL
 PEDV VVDAEKLHMYGASLIGGMALGGIT---AAAALPFSYAVQARLNYLALQTDVQLQRNQQLL
 CCov VANDDKMYTASLAGGITLGLSGG---GAVSIPFAVAVQARLNYVALQTDVQLNKNQQYL
 PRC VANADKMTMYTASLAGGITLGFAGG---GAVSIPFAVAVQARLNYVALQTDVQLNKNQQYL
 FICV VVDGNKMSMYTASLIGGMALGSIT---SAVAVPFAMQVQARLNYVALQTDVQLQENQKIL
 BoCov LLSVNQISGYTLAATSASLFPPLS---AAGVVPFYLNVQYRINGIGVTMDVLSQNKLI
 OC43 LLSNDQISGYTLAATSANLFPFWS---AAGVVPFYLNVQYRINGIGVTMDVLSQNKLI
 PHEV LLSVNQISGYTLAATSASLFPFPT---AAGVVPFYLNVQYRINGIGVTMDVLSQNKLI
 MHV VLSESQISGYTAGATAAAMFPFPT---AAGVVPFYLNVQYRINGIGVTMDVLSQNKMI
 TOR2_S LLTDDMLAAATAALVSGTATAGWTFGAGAALQIPFAMQMYRFGNGIGVTQNVLYENQKQI
 AIBV IITADMQTMYTASLVGAMAFGGIT---SAAAIPTAQIARINHLGIAQSLLMKVQEKI
 : : * : * . . : * . * : : *

229E AASFNKAMTNIVDAFTGVNDAITQTSQALQTVATALNKIQDQVNVQGGNSLNHLTSQLRQN
 PEDV AESFNSAIGNITSAFESVKEAISQTSKGLNTVAHALTKVQEVVNSQGSALNQLTVQLQHN
 CCov ANAFNQAIGNITQAFGKVNDAIHQTSQGLATVAKVLAKVQDVVNTQGGALSHLTLQQLQNN
 PRC ASAFNQAIGNITQSPGKVNDAIHQTSRGLTTVAKALAKVQDVVNTQGGALRHLLTVQLQNN
 FICV ANAFNNAIGNITLALGKVSNAITTTSDGFNSMASALTKIQSVNVQGGALSQLTSQQLKN
 BoCov ANAFNNALDAIQEGFDATN-----S-ALVKIQAVVNANAEALNLLQQLSNR
 OC43 ANAFNNALDAIQEGFDATN-----S-ALVKIQAVVNANAEALNLLQQLSNR
 PHEV ASAFNNALDAIQEGFDATN-----S-ALVKIQAVVNANAEALNLLQQLSNR
 MHV ASAFNNALGAIQEGFDATN-----S-ALGKIQSVVNANAEALNLLNQLSNR
 TOR2_S ANQFNKAISQIQESLTTTS-----TALGKLQDVVNQNAQALNTLVKQLSSN
 AIBV AASFNKAIHGMQEGFRSTS-----LALQQVQDVVNQKQSAILTETMNSLNKN
 * : * : * . . : * . * : : *

FIGURE 14D

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229E      FQAISSSIQAIYDRDLTIQADQQVDRLITGRALNALNVFVSHLTLYTEVRSRQLAQQKV
PEDV      FQAISSSIDDIYSRLDILLADVQVDRLITGRLSALNAFVAQTLTKYTEVQASRQLAQQKV
CCov      FQAISSSIDDIYNRLDELSADAQVDRLITGRLTALNAFVSQTLTRQAEVRSRQLAKDKV
PRC       FQAISSSIDDIYNRLDELSADAQVDRLITGRLTALNAFVSQTLTRQAEVRSRQLAKDKV
PICV      FQAISSSIAETYNRLEKVEADAQVDRLITGRALNALNAVVSQTLTQYAEVKASRQIALEKV
BoCov     FQAISSSLQEILSRDLDALEAQAQIDRLINGRLTALNVVVSQQLSDSTLVKFSAAQAMEKV
OC43      FQAISSSLQEILSRDLDALEAQAQIDRLINGRLTALDAYVVSQQLSDSTLVKFSAAQAMEKV
PHEV      FQAISSSLQEILSRDLDALEAQAQIDRLINGRLTALNAVVSQQLSDSTLVKFSAAQAIIEKV
MHV       FQAISSSLQEILTRDAVEAQAQIDRLINGRLTALNAVYSKQLSDSTLIKFSAAQAIIEKV
TOR2_S    FQAISSVLNDILSRDLKVEAEVQIDRLITGRLQSLQTYVVTQQLIRAAEIRASANLAATKM
AIBV      >    FGAISSVIQDIYAQLDAIQADAQVDRLITGRLSLVLASAKQSEYIRVSQQRELATQKI
          * ***: : * :*: : * .*:****.*** :.. : : . * * :

229E      NECVKSQSKRYGFCG--NGTHIFSVIUNAPEGLVFLHTVLLPTQYKDVEAWSGLCV--DG--
PEDV      NECVKSQSQRVYGFCCGDEHIFSLVQAAPQGLLEFLHTVLPVGFVNVLAIALGLCV--NG--
CCov      NECVRSQSQRVYGFCCG--NGTHLFLSLANAAPNGMIFFFHTVLLPTAYETVTAWSGICASDGDR
PRC       NECVRSQSQRVYGFCCG--NGTHLFLSLANAAPNGMIFFFHTVLLPTAYETVTAWSGICALDGDR
PICV      NECVKSQSNRYGFCG--NGTHLFLSLVNSAPEGLLFFHTVLLPTWEVEVTAWSGICVNDT--
BoCov     NECVKSQSSRINFCCG--NGNHIISLVQNAFYGLYFIHFSYVPTKYVTAKVSPGLCI-----
OC43      NECVKSQSSRINFCCG--NGNHIISLVQNAFYGLYFIHFSYVPTKYVTAKVSPGLCI-----
PHEV      NECVKSQSSRINFCCG--NGNHIISLVQNAFYGLYFIHFSYVPTKYVTAKVSPGLCI-----
MHV       NECVKSQSTRINFCCG--NGNHIISLVQNAFYGLCFIHFYSVPTSFKTANVSPGLCI-----
TOR2_S    SECVLGGQSKRVDFCCG--KGYHLSMFPQAAPHGVVFLHVTVVPSQERNFTTAPAICH-----
AIBV      >    NECVKSQSNRYGFCG--SGRHVLSIQNAAPNGIYVFIHFTYTPETVFNVTATVGFVNLNA
          .*** .*: * .*** .* :*: : * * * : * : * * * * . : : *

229E      TNGYVLRQPNLALYK-----EGNYRITSRIMFEPRIPTMADFVQIENCNVTFVNI SRS
PEDV      EIALTLREPGLVLFTHLQTYTATEYFVSSRRMFEPKPTVSDVQIESCVVTVVNLTS
CCov      TFGLVVKDQVLTFRN-----LDDKFYLTPTMYQPIVATSSDFVQIEGCDVLFVNATVI
PRC       TFGLVVKDQVLTFRN-----LDDKFYLTPTMYQPRVATSSDFVQIEGCDVLFVNTVVS
PICV      -YAYVLKDFDHSIFS-----YNGTYMVTFRNMFQPRKQMSDFVQIITSCVTFLNMTYT
BoCov     -AGDRGIAPKSGYFVN-----VNNTWMFTGSGYYPPEPITGNVVVMSTCAVNYTKAPDV
OC43      -AGDRGIAPKSGYFVN-----VNNTWMFTGSRYYYPEPITGNVVVMSTCAVNYTKAPDV
PHEV      -AGDIGISPKSGYFIN-----VNNSWMFTGSSYYYPEPITQNNVVVMSTCAVNYTKAPDL
MHV       -SGDRGLAPKAGYFVQ-----DNGEWKFTGSNYYYPEPITDKNSVAMISCAVNYTKAPEV
TOR2_S    -EGKAYFPREGVVFVN-----GTSWFITQRNFFSQIITTDNTFVSGNCDVVIGIINNT
AIBV      >    SQYAIVPANGRGIFIQ-----VNGTYIITSRDMYMPRDYTAGDIVTLTSCQANYVNVNKT
          : : : * : : . * .

229E      ELQTIYP- EYIDVNKTLQELSYKL- PNYTVPDLV---VEQYNQTI LNLTSEISTLENKSA
PEDV      QLPDVIP- DYIDVNKTLDEILASL- PNRTGPSLP---LDVFNATV LNLTGEDIADLEQRSE
CCov      DLPSIIP- DYIDINQTVQDILENFRPNWTVPELP---LDIFNATV LNLTGEDIADLEQRSE
PRC       DLPSIIP- DYIDINQTVQDILENFRPNWTVPELT---LDVFNATV LNLTGEDIADLEQRSE
PICV      TFQEIIV- DYIDINKTIADMLEQYNPNYTTPELNL- LLDIFNQT KLNLTAEIDQLEQRAD
BoCov     MLNISTP- NLPDFKEELDQWFKNQ--TSVAPDLSL-DY-- INVTFDLQDEMNI-----
OC43      MLNISTP- NLPDFKEELDQWFKNQ--TSVAPDLSL-DY-- INVTFDLQDEMNI-----
PHEV      MLNISTP- NLPDFKEELQWFKNQ--SSVAPDLSL-DY-- INVTFDLQDEMNI-----
MHV       FLNNSIP- NLPDFKEELDKWFKNQ--TSIAPDLSL-DFEKLNVTFDLTYEMNI-----
TOR2_S    VYDPLQP- ELDSFKEELDKYFKNH---TSPDVLGDISGINASVNIQKEID-----
AIBV      >    VITTFVEDDDFNFDDELSKWNWNT--KHGLPDFD---DFNYTVPILNISGEID-----
          : . . . : . . * . . . : : * :

229E      ELNYTVQKLQTLIDNINSLVLDLKWLN RVETIYKWPWVWVLCISVVLIFVVSMLLCCCS
PEDV      SLRNTTEELRSLINNINNTLVLDLWLN RVETIYKWPWVWVLLIIVIVLIFVVSLLVFCFCCIS
CCov      KLHNTTVELAILIDNINNTLVNLEWLNRIETVYKWPWVWVLLIGLVVIFCIPILLFCCCS
PRC       KLHNTTVELAILIDNINNTLVNLEWLNRIETVYKWPWVWVLLIGLVVIFCIPILLFCCCS
PICV      NLTTIAHELQYIDNLNKTLVDLWLNRIETVYKWPWVWVLLIGLVVIFCIPILLFCCCS
BoCov     -----RLQEAIKVLNQSYINLKDIGTYEYVYKWPWVWVLLIGFAGVAMLVLLFFICCC
OC43      -----RLQEAIKVLNQSYINLKDIGTYEYVYKWPWVWVLLIGFAGVAMLVLLFFICCC
PHEV      -----RLQEAIKVLNQSYINLKDIGTYEYVYKWPWVWVLLIGLAGVAMLVLLFFICCC
MHV       -----RIQDAIKKLNESYINLKEVGYEYVYKWPWVWVLLIGLAGVAVCVLLFFICCC
TOR2_S    -----RLNEVAKNLNESLIDLQELGKYEYKWPWVWVLLGFIAGLTAIVMVTIILCCM
AIBV      >    -----NIQGVIQGLNDSLINLELSIIKTYIKWPWVWVWLAIGFAIIFILILGWVFFM
          . : . : * : : * . : : * : * * * : * : : :

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FIGURE 14E

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229E      TGCCG-FFSCFASSIRGCCSTKL-PYYDVEKIHIQ-----
PEDV      TGCCG-CCGCCGACFSGCCRGPRQLQPYEAFEKVVHVQ-----
CCOV      TGCCG-CIGCLGSCCHSICSRQFESYEPYIEKVVHVH-----
PRC       TGCCG-CIGCLGSCCHSIFSRQFENYEPYIEKVVHVH-----
FICV      TGFCG-CFGCVGSCCHSLCSRQFETYPYIEKVVHIIH-----
BoCov     TGCGTSCFKICGGCCD-DYTGHQELVIK---TSHDD-----
OC43      TGCGTSCFKKCGGCCD-DYTGHQELVIK---TSHDD-----
PHEV      TGCGTSCFKKCGGCCD-DYTGHQEFVIK---TSHDD-----
MHV       TGCGSCEFRCGSCCD-EYGGHQDSIVIHNI SAHED-----
TOR2_S    TSCCCLKGACSCGSCCKFDEDDSEPVKGVKLVHT-----
AIBV      > TGCCGCCCGCGFGLIPLISKCGKRSSYYTTFDNDVVTEQYRPKKSV
          *

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Key	Name	Genbank % ID*
229E	spike glycoprotein [Human coronavirus 229E].	AAK32191 28.6% (SEQ ID NO: 53)
AIBV	spike glycoprotein [Avian infectious bronchitis virus].	AAO34396 27.6% (SEQ ID NO: 54)
BoCov	E2 glycoprotein precursor (Spike glycoprotein)	F25193 30.5% (SEQ ID NO: 55)
CCOV	spike protein - canina coronavirus	S41453 26.1% (SEQ ID NO: 56)
FICV	peplomer protein [Feline infectious peritonitis virus].	BAA06805 25.4% (SEQ ID NO: 57)
MHV	E2 glycoprotein precursor (Spike glycoprotein)	F11225 31.9% (SEQ ID NO: 58)
OC43	surface protein - human coronavirus	S44241 30.7% (SEQ ID NO: 59)
PEDV	spike protein [Porcine epidemic diarrhea virus].	CAA80971 26.0% (SEQ ID NO: 60)
PHEV	spike glycoprotein [porcine hemagglutinating encephalomyelitis virus]	AAI80031 30.5% (SEQ ID NO: 61)
PRC	S protein [Porcine respiratory coronavirus].	AAA46905 27.5% (SEQ ID NO: 62)
TOR2_S	Sars associated virus S glycoprotein (SEQ ID NO: 33)	

FIGURE 14F

```

10          20          30          40          50
TOR2_E      MYSFVSEETGTLIVNSVLLFLAFVVFLVTLAILTALRLCAYCCNIVNVSLVKPTV
           : . . . : . . . : . . . : . . . : . . . : . . . : . . . : . . . :
PGV        MTFPRALTVIDDNG-MVINIIFWFLIIILILLSIALLNIIKLCMVCCNLGRTVVIIVPAQ
           10          20          30          40          50

           60          70
TOR2_E YVYSRVKLNLSSEGVPDLLV (SEQ ID NO: 35)
           . . . : . . .
PGV      HAYDAYKNFMRIKAYNPDGALLA (SEQ ID NO: 63)
           60          70          80

```

FIGURE 15

MESLVLGVNEKTHVQLSLPVLQVRDVLVVRGFGDSVEEALSEAREHLKNGT
 CGLVELEKGVLPQLEQPYVFLKRSDALSTNHGHKVVVELVAEMDGIQYGRS
 GITLGLVLPVHVGETPIAYRNVLLRKNKNGKAGGHSYIGIDLKSYDLGDELG
 TDPFIEDYEQNWNTHKSGALRELRELNNGGAVTRYVDNNTFCGPDGYFLDC
 IKDFLARAGKSMCTLSEQLDYIESKRGVYCCRDHEHEIAWFTEERSDKSYE
 HQTPPEIKSAKKFDTFKGCEPKFVFLNSRVKVIQPRVEKKTEGFMGRI
 RSVYFVASPQECNMHLSTLMKCNHCDEVSWQTCDFLKATCEHCGTENLV
 IEGPTTCGYLPTNAVVKMPCPACQDPEIGPEHVSADYHNHSNIETRLRKG
 GRTRCFGGCVFAYVGCYNKRAYVWPRASADIGSGHTGITGDNVETLNEDL
 LEILSRERVNINIVGDFHLNEEVAIILASFASSTSAFIDTIKSLDYKSFK
 TIVESCGNYKVTRGKPKVGAWNIGQQRSVLTPLCGFPQAAQVIRSI FAR
 TLDAANHSIPDLQRAAVTILDGISEQSLRLVDAMVYTSDLLTNSVIIMAY
 VTGGLVQQTSQLSNLGGTTVEKLRPIFEWIEAKLSAGVEFLKDAWEILK
 FLITGVFDIVKGGIQVASDNKDCVKCFIDVVKALEMCIQVTTIAGAKL
 RSLNLGEVFLAQSKGLYRQCIRGKEQLQLMLPLKAPKEVTFLEGDSHDTV
 LTSEEVVLKNGELEALETPVDSFTNGAIVGTPVCVNGMLLEIKKDEQYC
 ALSPGLLATMNVFRLKGGAPKIGVTFGEDTVWEVQGYKNVRIITFELDERV
 DKVLNEKCSVYTVESGTEVTEFACVVAEAVVKTLQFVSDLLTNMGI DLDE
 WSVATFYLFDDAGEENFSSRMYCSFYPPDEEEEDDAECEEEIIDEETCEHE
 YGTEDDYQGLPLEFGASAEVTRVEEEEEEDWLD DTTTEQSEIEPEPEPTPE
 EPVNOFTGYLKLTDNVAIKVDIVKEAQSANPMVIVNAANIHLKHGGGVA
 GALNKATNGAMQRESDDYIKLNGPLTVGGSCLLSGHNLAKKCLHVGPNL
 NAGEDIQLLKAAAYENFNSQDILLAPLLSAGIFGAKPLQSLQVCVQTVRTQ
 VYIAVNDKALYEQVMDYLDNLKPRVEAPKQEEPPNTEEDSKTEEKSVVQK
 PVDVKPKIKACIDEVTTLEETKFLTNKLLLFADINGKLYHDSQNMLRGE
 DMSFLEKDAFYMVGDVITSGDITCVVIPSCKAGGTTEMLSRALKKVPVDE
 YITTYPGQCGAGYTL EAKTALKKCKSAFYVLPSEAPNAKEEILGTVSWN
 LREMLAHAEBETRLMPCMDVRAIMATIQRKYKGIKIQEGIVDYGVRRFFF
 YTSKEPVASITTKLNSLNEPLVTMPIGYVTHGNLEEAARCMRSLKAPAV
 VSVSPDAVTTYNGYLTSSSKTSEHFVETVSLAGSYRDWSYSGQRTELG
 VEFKRGDKIVYHTLESFVEFHL DGEVLSL DKLKSLLSLREVKTTIKVFTT
 VDNTNLHTQLVDMSTYGGQFGPTVLDGADVTKIKPHVNHGKTFVFLPS
 DDLRSEAFEYHTLDESFLGRYMSALNHTKKKWKFPQVGLTSTIKWADNN
 CYLSSVLLALQQLVFNAPALQEAAYRARAGDAANFCALILAYSNKTVG
 ELGDVRETMTHLLQHANLES AKRVLNVVCKHCGKQTTTLTGVEAVMYMGT
 LSYDNLKTGVSIPCVCCRDAQYLVQQESSFVMSAPAEYKQQQTFCLC
 ANEYTGNYCCGHYHTITAKETLYRIDGAHLTKMSEYKGPVTDVFKETSY
 TTTIKPVSYKLDGVTYTEIEPKLDGYKKDNAYYTEQPIDLVPQTPLPNA
 SFDNFKLTCNFKFADDLNQMTGFTKASRELSVTFPDLNGDVVAIDYR
 HYSASFKKGAKLLHKPIVWHINQATTKTTFKPNFWCLRCLWSTKPVDTSN
 SFEVLAVEDTQGMNLCESQOPTSEEVENPTIQKEVIECDVKTEVVG
 NVILKPSDEGVKVTQELGHEDLMAAYVENTSITIKKPNELSLALGLKTI A
 THGIAAINSVPWSKILAYVKPFLGQAAITTSNCAKRLAQRVFNMYMPYVF
 TLLFQLCTFKSTNSRIRASLPTTIAKNSVKSVAKLCLDAGINYKSPKF
 SKLFTIAMWLLLSICLGLCVTAAFVLLSNF GAPSYCNGVRELYLNS
 SNVTMDFCESGFPCICLSGLDSLDSYPALETIQVTISSYKLDLTLGL
 AAEWVLAJMLFTKFFYLGLLSAIMQVFFGYFASHFISNSWLMWFIISIVQ
 MAPVSAMVRMYIFFASFYIWKSVVHIMDGTSSSTCMCYKRNRATRVEC
 TTI VNGMRSFYVYANGGRGPKTHNWNCLNCDTFFCTGSTFI SDEVARDL
 SLQFKRPI NPTDQSSYIVDSVAVKNGALHLYFDKAGQKTYERHPLSHFVN
 LDNLRANNFKGSLPINVIVFDGKSKCDESASKSASVYYSQLMCPILLLD
 QALVSDVGDSTEVSVKMFDAYVDTF SATPSVPMKALVATAHSELAKG
 VALDGLVSTFVSAARQGVVDTVDVTDKDVIECLKLSHHSDELVTDGSCNMF
 MLTYNKVENMTPRDLGACIDCNARHINAQVAKSHVNSLITWNVKDYMSLSE
 QLRKQIRSAAKKNNIPFRLTCATTRQVNVITTKISLKGKIVSTCFKLM
 LKATLLCVLAALVCIYVMPVHTLSIHGVTNEIIGYKAIQDGVTRDIIST
 DDCFANKHAGFDWFSQRGGSYKNDKSCPVAAIITREIGFIVPGLPCTV
 LRAINGDFLHFLPRVPSAVGNICYTF SKLIEYSDFATSACVLAECTIFK
 DAMGKPVYCYDTNLLBGSISYSELRPDTRYVLM DGSIIQFPNTYLEG SV
 RVVTTFDAEYCRHGTCESEVVICLSTSGRWVLMNEHYRALSGVFCGVDA
 MNLIANIFTPLVQPVGALDVSASV VAGGIIAILVTC AAYYFMKFRRVFGE
 YNHVVAANALLFLMSFTILCLVPAYSFLPGVYSVFYLYLTFYFTNDVSFL
 AHLQWFAMFSPVFPWITAIYVFCISLKHCHWFFNNYLRKRVMFNGVTF S
 TFEEAALCFFLLNKEMV LKLRSETLPLTQYNYR LALYNYKYFSGALDT
 TSYREAACCHLAKALNDFNSGADVLYQPPQTSITSAVLQSGFRKMAFPS
 GKVEGCMVQVTCGTTLNGLWLD DTVYCPRHVIC TAEDMLNPNYEDLLIR
 KSNHSFLVQAGNVQLRVIGHSMQNLRLKVDTSNPKTPKYKVFRIQPGQ
 TFSVLACYNGPSGVYQCAMPNHTIKGSFLNGSCG SVGFNIDYDCVSPC
 YMHMELPTGVHAGTDLEKFGYGFVDRQTAQAAGTDTTITLNLAWLYA
 AVINGDRWFLNRFTTTLNDFNLVAMKYNYEPLTQDHVDILGELSQTGIA
 VLDMCAALKELLQNGMNGRTILGSTILEDEFTPFDDVVRQCSGVTFOGKPK

FIGURE 16A

KIVKGTTHWMLLTFLTSLILVQSTQWSLFFFVYENAFLPFTLGIMAAIAA
CAMLVVKHKHAFCLFLLPSLATVAYFNMVYMPASWVMRIMTWLELADTS
LSGYRLKDCVMYASALVLLILMTARTVYDDAARRVWFLMNVITLVYKVVY
GNALDQAISMWALVISVTSNYSGVVPTIMFLARAIIVFCVEYYPLLFITG
NFIQCIMLVYCFLYCCCYFGLFCLLNRYFRLTLGVYDYLVSQEFRYM
NSQGLLPKSSIDAFKLNKLLGIGGKPCIKVATVQSKMSDVKCTSVVLL
SVLQQLRVESSSKLWAQCVQLHNDILLAKDTTEAFEKMSLSSVLLSMQG
AVDINRLCEEMLDNRATLQAIASEFSSLPYAAAYATAQEAYEQAVANGDS
EVLKLLKSLNVAKSEFDRDAAMQRKLEKMAQAMTQMYKQARSEDKRA
KVTSAMQTMLEFMLRKLNDALNNIINNARDGCVPLNIIPLTTAAKLMVV
VPDYGTYNKTCNFTFYASALWEIQVVDADSKIVQLSEINMDNSPNLA
WPLIVTALRANSVAVKLQNNELSFVALRQMSCAAGTTQTACTDDNALAYYN
NSKGGREFVALLSDHQDLKWARFPKSDGTGTIYTELEPPCFVTDTPKGP
KVKYLYFIKGLNNLNRGMVLSLAATVRLQAGNATEVPANSTVLSFCFAFA
VDFAKAYKDYLASGGQPIITNCVKMLCTHTGTGQAITVTPKANMDQESFGG
ASCCLYCRCHIDHPNPKGFCDLKGKYVQIPTTCANDPVGFTLRNVTCTVC
GMWKGVCSCDQLREPLMQSADASTF

(SEQ ID NO: 64)

FIGURE 16B

FKRVCG
VSAARLTPCGTGTSTDVVYRAFDIYNEKVAGFAKFLKTNCCRPFQEKDEEG
NLLDSYFVVKRHTMSNYQHEETIYNLVKDCPAVAVHDFKFRVDGDMVPH
ISRQRLTKYTMADLVYALRHFDEGNCDTLKEILVTYNCCDDDYFNKKDWY
DFVENPDILRVYANLGERVRQSLKTVQFCDAMRDAGIVGVLTLDNQDLN
GNWYDFGDFVQVAPGCGVPIVDSYSSLMPILTLTRALAAESHMDADLAK
PLIKWDLKDYDFTEERLCLFDRYFKYWDQTYHPNCINCLDDRCILHCANF
NVLFSVFPPTSFGPLVRKIFVDGVFVSTGYHFRELGVVHNQDVNLHS
SRLSFKELLVYAADPAMHAASGNLKKRFTCFVSAALTNVAFQTVKPG
NFNKDFYDFAVSKGFFKFGSSVELKHFFFAQDGNAAISDYDYRYNLP
CDIRQLLFVVEVDKYFDCYDGGCINANQVIVNNLDKSAGFPFNKWKAR
LYYDMSYEDQDALFAYTKRNVIPTITQMNLYAISAKNRARTVAGVSIC
STMTRQFHQKLLKSAATRGTAVVIGTSKFGGWHNMLKTVYSDVETPH
LMGWDPKCDRAMPNMLRIMASLVLARKHNTCCNLSHRFYRLANCAQVL
SEMVMCGGSLYVKPGGTSAGDATAYANSVFNICQAVTANVNALLSTDGN
KIADKYVRNLQHRLYECLYRNRDVEDHEFVDEFYAYLRKHFSMMILSDDAV
VCYNSNYAAQGLVASIKNFKAVLYYQNNVFMSEAKWTETDLTKGPHEFC
SQHTMLVKQGGDDYVLYPDPSPRILGAGCFVDDIVKTDGTLMIERFVSLA
IDAYPLTKHPNQEYADVFLYLQYIRKLHDELGHMLDMYSVMLTNDNTS
RYWEPFYEAMYTPHTVLQAVGACVLCNSQTSLRGACIRRPFLCCKCCY
DHVISTSHKLVLSVNPVVCNAPGCDVTDVTLQYLGGMSSYCKSHKPPISF
PLCANGQVFGLYKNTCVGSNDVTFDFAIATCDWTNAGDYILANTCTERLK
LFAAETLKATEETFKLSYGIATVREVLSRELHLSWEVKGKPRPPLNRNV
FTGYRVTKNSKVQIGEYTFEKGDYGDVAVYRGTTTYKLVNGDYFVLTSTHT
VMPLSAPTLVQEHYVRITGLYPTLNI SDEFSSNVANYQKVMQKYSTLQ
GPPGTGKSHFAIGLALYPSARIVYTACSHAVALCEKALKYLPIDKCS
RIIPARARVECFDKFKVNSTLEQYVFCVNALPETTADIVVDEISMATN
YDLSVVNARLRAKHVYVYIGDPAQLPAPRLLTKGTLPEYFNVSVCRLMKT
IGPDMFLGTCRRCPAEIVDTVSALVYDNKLKAHKDKSAQCFKMFYKGVIT
HDVSSAINRPQIGVVREFLTRNPAWRKAVFISPYNSQNAVASKILGLPTQ
TVDSSQGSEYDYVITQTTETAHSCNVNRFNVAITRAKIGILCIMS DRDL
YDKLQFTSLEIPRRNVATLQAENVTLGFKDCSKIIITGLHPTQAPTHLSVD
IKFKTEGLCVDIPGPKDMTYRRLISMMGFKMNYQVNGYPNMFITREEAI
RHVRAWIGFDVEGCHATRDAVGTNLPLQLGFSTGVNLVAVPTGYVDTENN
TEFTRVNAKPPPGDQFKHLIPLMYKGLPWNVVRKIVQMLSDTLKGLSDR
VVFLWAHGFELTSMKYFVKIGPERTCCLCDKRATCFSTSSDITYACWNHS
VGFYVYNPFMIDVQWQWFTGNLQSNHDQHCQVHGNAHVASCDAIMTRCL
AVHECFVKRVDWSVEYPIIGDELRVNSACRKYQHMMVKSALLADKFPVLH
DIGNPKAIKCVQAEVEWKFYDAQPCSDKAYKIEELFYSYATHHDKFTDG
VCLFWNCNVDRYPANAI VCRFDTRVLSNLNLP GCDGGS LYVNKHAFHTPA
FDKSAFTNLKQLPFFYSDSPCESHGKQVVSIDIDYVPLKSATCITRCNLG
GAVCRHHANEYRQYLDAYNMMISAGFSLWIYKQFDYTNLWNTFTRLQSL
NVAYNVNKGHFDGHAGEAPVSIINNAVYTKVDGIDVEIFENKTTLPVNV
AFELWAKRNKIPVPEIKILNNGVDIAANTVIWDYKREAPAHVSTIGVCT
MTDIAKKPTESACSSLTVLFDGRVEGQVDFRNARNGVLITEGSKGLTP
SKGPAQASVNGVTLIGESVKTFQFNYFKKVDGIIQQLPETYFTQSRDLEDF
KPRSQMETDFLELAMDEFIQRYKLEGYAFEHIVYGDVSHGQLGGLHLMIG
LAKRSQD SPLKLEDFIPMDSTVKNYFITDAQTGSSKVCVSDIDLLDDFV
EIIKSQDLSVISKVVKVTIDYAEISFMLWCKDGHVETFPKQLQASQAWQP
GVAMPNLYKMQRMLLEKCDLQNYGENAVIPKGIMMNAKYTQLCQYLNTL
TLAVPYNMVRVIFHGAGSDKGVAPGTAVLRQWLP TGTLVSDS LNDVSDA
DSTLIGDCATVHTANKWDLIISDMYDPRTKHVTKENDSKEGFFTYLTCGFI
KQKLAGGSIAVKITEHSWNADLYKLMGHFSWWTAFVTNVNASSSEAFLI
GANYLKPKPEQIDGYTMHANYIFWRNTNPIQLSSYSLFDMSKFLPKLRGT
AVMSLKENQINDMIYSLLEKGRLLIRENRRVVVSSDILVNN

(SEQ ID NO: 65)

FIGURE 17

MDLFMRFFTLRSITAQPVKIDNASPASTVHATATIPLQASLPFGWLIVIGV
AFLAVFQSATKIIALNKRWQLALYKGFQFICNLLLLFVVTIYSHLLLVAAG
MEAQFLYLYALIYFLQCINACRIIMRCWLCWKCKSKNPLLYDANYFVCWH
THNYDYCIPYNSVTDTIIVTEGDGISTPKLKEDYQIGGYSEDRHSGVKDY
VVVHGYFTEVYYQLESTQITTDGTGIENATFFIFNKLVKDPPNVQIHTIDG
SSGVANPAMDPIYDEPTTTTSVPL (SEQ ID NO: 66)

FIGURE 18

MMPTTLFAGTHITMTTVYHITVSQIQLSLLKVTAHQHNSKKTTLVIL
RIGTQVLKTMSTLYMAISPFFTSLSLHKLQTLVLKMLHSSSLTSLKTH
RMCKYTQSTALQELLIQQWIQFMMSRRLLACLCKHKKVSTNLCTHSFRK
KQVR (SEQ ID NO: 67)

FIGURE 19

MFHLVDFQVTIAEILIIIMRTFRIAIWNLDVLISSIVRQLFKPLTKKNYS
ELDDEEPMELDYP (SEQ ID NO: 68)

FIGURE 20

MKIILFLTIVFTSCELYHYQECVRGTTVLLKEPCPSGTYEGNSPFHPLA
DNKFALTCTSTHFAFACADGTRHTYQLRARSVSPKLFIRQEEVQQELYSP
LFLIVAALVFLILCFTIKRKTE (SEQ ID NO: 69)

FIGURE 21

MNELTLIDFYLCFLAFLFLVLIIMLIIFWFSLEIQDLEEPCTKV
(SEQ ID NO: 70)

FIGURE 22

MKLLIVLTCISLCSICTVVQRCASNKPHVLEDPCVKVQH
(SEQ ID NO: 71)

FIGURE 23

MCLKILVRYNTRGNTYSTAWLCALGKVLPHRWHTMVQTCTPNVTINCQD
PAGGALIARCWYLHEGHQTAAFRDVLVVLNKRTN (SEQ ID NO: 72)

FIGURE 24

MDPNQTNVPPALHLVDPQIQLTITRMEDAMGQGQNSADPKVYPIILRLG
SQLSLSMARRNLDSEARAFQSTPIVVQMTKLATTEELPDEFVVVTAK

(SEQ ID NO: 73)

FIGURE 25

MLPPCYNFLKEQHCQKASTQREAEAAVKPLLAPHHVAVIQEIQLLAAVG
EILLLEWLAEVVKLPSRYCC (SEQ ID NO: 74)

FIGURE 26

CIAVGQLCVFVNIGRPCCSGLCVFA--CTVKL conotoxin
CISLCS-CICTVVQRCASNKPHVLEDPCKVQH sars
**::. * : * ... * : *.*:

FIGURE 27

SARS VIRUS NUCLEOTIDE AND AMINO ACID SEQUENCES AND USES THEREOF

Sequence Listing

The instant application contains a Sequence Listing which has been submitted in ASCII format via EFS-Web and is hereby incorporated by reference in its entirety. Said ASCII copy, created on Jan. 18, 2010, is named 308410US.txt and is 462,420 bytes in size.

FIELD OF THE INVENTION

The invention is in the field of virology. More specifically, the invention is in the field of coronaviruses.

BACKGROUND OF THE INVENTION

Severe acute respiratory syndrome (SARS), a worldwide outbreak of atypical pneumonia with an overall mortality rate of about 3 to 6%, has been attributed to a coronavirus following tests of causation according to Koch's postulates, including monkey inoculation (R. Munch, *Microbes Infect* 5, 69-74, January 2003). The coronaviruses are members of a family of enveloped viruses that replicate in the cytoplasm of animal host cells (B. N. Fields et al., *Fields virology*, Lippincott Williams & Wilkins, Philadelphia, 4th ed., 2001). They are distinguished by the presence of a single-stranded plus sense RNA genome, approximately 30 kb in length, that has a 5' cap structure and 3' polyA tract. Hence the genome is essentially a very large mRNA. Upon infection of an appropriate host cell, the 5'-most open reading frame (ORF) of the viral genome is translated into a large polyprotein that is cleaved by viral-encoded proteases to release several nonstructural proteins including an RNA-dependent RNA polymerase (Pol) and an ATPase helicase (Hel). These proteins in turn are responsible for replicating the viral genome as well as generating nested transcripts that are used in the synthesis of the viral proteins. The mechanism by which these subgenomic mRNAs are made is not fully understood, however transcription regulating sequences (TRSs) at the 5' end of each gene may represent signals that regulate the discontinuous transcription of subgenomic mRNAs (sgmRNAs). The TRSs include a partially conserved core sequence (CS) that in some coronaviruses is 5'-CUAAAC-3'. Two major models have been proposed to explain the discontinuous transcription in coronaviruses and arterioviruses (M. M. C. Lai, D. Cavanagh, *Adv Virus Res.* 48, 1 (1997); S. G. Sawicki, D. L. Sawicki, *Adv. Exp. Med Biol.* 440, 215 (1998)). The discovery of transcriptionally active, subgenomic-size minus strands containing the antileader sequence and transcription intermediates active in the synthesis of mRNAs (D. L. Sawicki et al., *J. Gen Virol* 82, 386 (2001); S. G. Sawicki, D. L. Sawicki, *J. Virol.* 64, 1050 (1990); M. Schaad, R. S. J. Baric, *J. Virol.* 68, 8169 (1994); P. B. Sethna et al., *Proc. Natl. Acad. Sci. U.S.A.* 86, 5626 (1989)) favors the model of discontinuous transcription during the minus strand synthesis (S. G. Sawicki, D. L. Sawicki, *Adv. Exp. Med Biol.* 440, 215 (1998)).

The coronaviral membrane proteins, including the major proteins S (Spike) and M (Membrane), are inserted into the endoplasmic reticulum Golgi intermediate compartment (ERGIC) while full length replicated RNA (+ strands) assemble with the N (nucleocapsid) protein. This RNA-protein complex then associates with the M protein embedded in the membranes of the ER and virus particles form as the nucleocapsid complex buds into the ER. The virus then migrates through the Golgi complex and eventually exits the

cell, likely by exocytosis (B. N. Fields et al., *Fields virology*, Lippincott Williams & Wilkins, Philadelphia, 4th ed., 2001). The site of viral attachment to the host cell resides within the S protein.

The coronaviruses include a large number of viruses that infect different animal species. The predominant diseases associated with these viruses are respiratory and enteric infections, although hepatic and neurological diseases also occur with some viruses. Coronaviruses are divided into three serotypes, Types I, II and III. Phylogenetic analysis of coronavirus sequences also identifies three main classes of these viruses, corresponding to each of the three serotypes. Type II coronaviruses contain a hemagglutinin esterase (HE) gene homologous to that of Influenza C virus. It is presumed that the precursor of the Type II coronaviruses acquired HE as a result of a recombination event within a doubly infected host cell.

In view of the rapid worldwide dissemination of SARS, which has the potential of creating a pandemic, along with its alarming morbidity and mortality rates, it would be useful to have a better understanding of this coronavirus agent at the molecular level to provide diagnostics, vaccines, and therapeutics, and to support public health control measures.

SUMMARY OF THE INVENTION

In general, the invention provides the genomic sequence of a novel coronavirus, the SARS virus, and provides novel nucleic acid molecules encoding novel proteins that may be used, for example, for the diagnosis or therapy of a variety of SARS virus-related disorders.

In one aspect, the invention provides a substantially pure SARS virus nucleic acid molecule or fragment thereof, for example, a genomic RNA or DNA, cDNA, synthetic DNA, or mRNA molecule. In some embodiments, the nucleic acid molecule includes a sequence substantially identical to any of the sequences of SEQ ID NOs: 1-13, 15-18, 20-30, 90-159, 208, 209. In some embodiments, the nucleic acid molecule includes a sequence from SEQ ID NO: 1, SEQ ID NO: 2, or SEQ ID NO: 15 or a fragment of these sequences. In alternative embodiments, the nucleic acid molecule may include a sequence substantially identical to SEQ ID NO: 1, SEQ ID NO: 2, or SEQ ID NO: 15, or a fragment thereof. In alternative embodiments, the nucleic acid molecule may include a s2m motif (for example, a s2m sequence substantially identical to any of the sequence of SEQ ID NOs: 16, 17, and 18), a leader sequence (for example, a sequence substantially identical to the sequence of SEQ ID NO: 3), or a transcriptional regulatory sequence (for example, a sequence substantially identical to any of the sequence of SEQ ID NOs: 4-13 and 20-30). In alternative embodiments, the nucleic acid molecule includes a sequence substantially identical to any of the sequences of nucleotides 265-13,398; 13,398-21,485; 21,492-25,259; 25,268-26,092; 25,689-26,153; 26,117-26,347; 26,398-27,063; 27,074-27,265; 27,273-27,641; 27,638-27,772; 27,779-27,898; 27,864-28,118; 28,120-29,388; 28,130-28,426; 28,583-28,795; and 29,590-29,621 of SEQ ID NO: 15. In alternative embodiments, the nucleic acid molecule may encode a polyprotein or a polypeptide. In alternative embodiments, the invention provides a nucleic acid molecule including a sequence complementary to a SARS virus nucleotide sequence.

In an alternative aspect, the invention provides a substantially pure SARS virus polypeptide or fragment thereof, for example, a polyprotein, glycoprotein (for example, a matrix glycoprotein that may include a sequence substantially identical to the sequence of SEQ ID NO: 34), a transmembrane

protein (for example, a multitransmembrane protein, a type I transmembrane protein, or a type II transmembrane protein), a RNA binding protein, or a viral envelope protein. In alternative embodiments, the invention provides a replicase 1a protein, replicase 1b protein, a spike glycoprotein, a small envelope protein, a matrix glycoprotein, or a nucleocapsid protein. In alternative embodiments, the invention provides a nucleic acid molecule encoding a SARS virus polypeptide. In alternative embodiments, the SARS virus polypeptide includes an identifiable signal sequence (for example, a signal sequence substantially identical to the sequence of SEQ ID NOs: 76 or 85), a transmembrane domain (for example, a transmembrane domain substantially identical to any of the sequences of SEQ ID NOs: 77-86), a transmembrane anchor, a transmembrane helix, an ATP-binding domain, a nuclear localization signal, a hydrophilic domain, (for example, a hydrophilic domain substantially identical to the sequence of SEQ ID NOs: 87), or a lysine-rich sequence (for example, a sequence substantially identical to the sequence of SEQ ID NO: 14). In alternative embodiments, the SARS virus polypeptide may include a sequence substantially identical to any of the sequences of SEQ ID NOs: 14, 33-36, 64-74, and 76-87.

In alternative embodiments, the invention provides a vector (for example, a gene therapy vector or a cloning vector) including a SARS virus nucleic acid molecule (for example, a molecule including a sequence substantially identical to any of the sequences of SEQ ID NOs: 1-13, 15-18, 20-30, 90-159, 208, 209), or a host cell (for example, a mammalian cell, a yeast, a bacterium, or a nematode cell) including the vector.

In alternative embodiments, the invention provides a nucleic acid molecule having substantial nucleotide sequence identity (for example, 30%, 40%, 50%, 60%, 70%, 80%, 90% or 100% complementarity) to a sequence encoding a SARS virus polypeptide or fragment thereof, for example where the fragment includes at least six amino acids, and where the nucleic acid molecule hybridizes under high stringency conditions to at least a portion of a SARS virus nucleic acid molecule.

In alternative embodiments, the invention provides a nucleic acid molecule having substantial nucleotide sequence identity (for example, 30%, 40%, 50%, 60%, 70%, 80%, 90% or 100% complementarity) to a SARS virus nucleotide sequence, for example where the nucleic acid molecule includes at least ten nucleotides, and where the nucleic acid molecule hybridizes under high stringency conditions to at least a portion of a SARS virus nucleic acid molecule.

In alternative embodiments, the invention provides a nucleic acid molecule comprising a sequence that is antisense to a SARS virus nucleic acid molecule, or an antibody (for example, a neutralizing antibody) that specifically binds to a SARS virus polypeptide.

In alternative embodiments, the invention provides a method for detecting a SARS epitope, such as a virion or polypeptide in a sample, by contacting the sample with an antibody that specifically binds a SARS epitope, such as a virus polypeptide, and determining whether the antibody specifically binds to the polypeptide. In alternative embodiments, the invention provides a method for detecting a SARS virus genome, gene, or homolog or fragment thereof in a sample by contacting a SARS virus nucleic acid molecule, for example where the nucleic acid molecule includes at least ten nucleotides, with a preparation of genomic DNA from the sample, under hybridization conditions providing detection of DNA sequences having nucleotide sequence identity to a SARS virus nucleic acid molecule. In alternative embodiments, the invention provides a method of targeting a protein

for secretion from a cell, by attaching a signal sequence from a SARS virus polypeptide to the protein, such that the protein is secreted from the cell.

In alternative aspects, the invention provides a method for eliciting an immune response in an animal, by identifying an animal infected with or at risk for infection with a SARS virus and administering a SARS virus polypeptide or fragment thereof or fragment thereof, or administering a SARS virus nucleic acid molecule encoding a SARS virus polypeptide or fragment thereof to the animal. In alternative embodiments, the administering results in the production of an antibody in the mammal, or results in the generation of cytotoxic or helper T-lymphocytes in the mammal.

In alternative embodiments, the invention provides a kit for detecting the presence of a SARS virus nucleic acid molecule or polypeptide in a sample, where the kit includes a SARS virus nucleic acid molecule, or an antibody that specifically binds a SARS virus polypeptide.

In alternative aspects the invention provides a method for treating or preventing a SARS virus infection by identifying an animal (e.g., a human) infected with or at risk for infection with a SARS virus, and administering a SARS virus nucleic acid molecule or polypeptide, or administering a compound that inhibits pathogenicity or replication of a SARS virus, to the animal. In alternative embodiments, the invention provides the use of a SARS virus nucleic acid molecule or polypeptide for treating or preventing a SARS virus infection.

In alternative aspects the invention provides a method of identifying a compound for treating or preventing a SARS virus infection, by contacting sample including a SARS virus nucleic acid molecule or contacting a SARS virus polypeptide with the compound, where an increase or decrease in the expression or activity of the nucleic acid molecule or the polypeptide identifies a compound for treating or preventing a SARS virus infection.

In alternative aspects the invention provides a vaccine (e.g., a DNA vaccine) including a SARS virus nucleic acid molecule or polypeptide.

In alternative aspects the invention provides a microarray including a plurality of elements, wherein each element includes one or more distinct nucleic acid or amino acid sequences, and where the sequences are selected from a SARS virus nucleic acid molecule or polypeptide, or a antibody that specifically binds a SARS virus nucleic acid molecule or polypeptide.

In alternative aspects the invention provides a computer readable record (e.g., a database) including distinct SARS virus nucleic acid or amino acid sequences.

A "SARS virus" is a virus putatively belonging to the coronavirus family and identified as the causative agent for sudden acute respiratory syndrome (SARS). A SARS virus nucleic acid molecule may include a sequence substantially identical to the nucleotide sequences described herein or fragments thereof. A SARS virus polypeptide may include a sequence substantially identical to a sequence encoded by a SARS virus nucleic acid molecule, or may include a sequence substantially identical to the polypeptide sequences described herein, or fragments thereof.

A compound is "substantially pure" when it is separated from the components that naturally accompany it. Typically, a compound is substantially pure when it is at least 60%, more generally 75% or over 90%, by weight, of the total material in a sample. Thus, for example, a polypeptide that is chemically synthesized or produced by recombinant technology will be generally be substantially free from its naturally associated components. A nucleic acid molecule may be substantially pure when it is not immediately contiguous with (i.e.,

covalently linked to) the coding sequences with which it is normally contiguous in the naturally occurring genome of the organism from which the DNA of the invention is derived. A nucleic acid molecule may also be substantially pure when it is isolated from the organism in which it is normally found. A substantially pure compound can be obtained, for example, by extraction from a natural source; by expression of a recombinant nucleic acid molecule encoding a polypeptide compound; or by chemical synthesis. Purity can be measured using any appropriate method such as column chromatography, gel electrophoresis, HPLC, etc.

A "substantially identical" sequence is an amino acid or nucleotide sequence that differs from a reference sequence only by one or more conservative substitutions, as discussed herein, or by one or more non-conservative substitutions, deletions, or insertions located at positions of the sequence that do not destroy the biological function of the amino acid or nucleic acid molecule. Such a sequence can be at least 10%, 20%, 30%, 40%, 50%, 52.5%, 55% or 60% or 75%, or more generally at least 80%, 85%, 90%, or 95%, or as much as 99% or 100% identical at the amino acid or nucleotide level to the sequence used for comparison using, for example, the Align Program (Myers and Miller, CABIOS, 1989, 4:11-17) or FASTA. For polypeptides, the length of comparison sequences may be at least 4, 5, 10, or 15 amino acids, or at least 20, 25, or 30 amino acids. In alternate embodiments, the length of comparison sequences may be at least 35, 40, or 50 amino acids, or over 60, 80, or 100 amino acids. For nucleic acid molecules, the length of comparison sequences may be at least 15, 20, or 25 nucleotides, or at least 30, 40, or 50 nucleotides. In alternate embodiments, the length of comparison sequences may be at least 60, 70, 80, or 90 nucleotides, or over 100, 200, or 500 nucleotides. Sequence identity can be readily measured using publicly available sequence analysis software (e.g., Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, Wis. 53705, or BLAST software available from the National Library of Medicine, or as described herein). Examples of useful software include the programs Pile-up and PrettyBox. Such software matches similar sequences by assigning degrees of homology to various substitutions, deletions, insertions, and other modifications.

Alternatively, or additionally, two nucleic acid sequences may be "substantially identical" if they hybridize under high stringency conditions. In some embodiments, high stringency conditions are, for example, conditions that allow hybridization comparable with the hybridization that occurs using a DNA probe of at least 500 nucleotides in length, in a buffer containing 0.5 M NaHPO₄, pH 7.2, 7% SDS, 1 mM EDTA, and 1% BSA (fraction V), at a temperature of 65° C., or a buffer containing 48% formamide, 4.8×SSC, 0.2 M Tris-Cl, pH 7.6, 1× Denhardt's solution, 10% dextran sulfate, and 0.1% SDS, at a temperature of 42° C. (These are typical conditions for high stringency northern or Southern hybridizations.) Hybridizations may be carried out over a period of about 20 to 30 minutes, or about 2 to 6 hours, or about 10 to 15 hours, or over 24 hours or more. High stringency hybridization is also relied upon for the success of numerous techniques routinely performed by molecular biologists, such as high stringency PCR, DNA sequencing, single strand conformational polymorphism analysis, and in situ hybridization. In contrast to northern and Southern hybridizations, these techniques are usually performed with relatively short probes (e.g., usually about 16 nucleotides or longer for PCR or sequencing and about 40 nucleotides or longer for in situ hybridization). The high stringency conditions used in these

techniques are well known to those skilled in the art of molecular biology, and examples of them can be found, for example, in Ausubel et al., *Current Protocols in Molecular Biology*, John Wiley & Sons, New York, N.Y., 1998, which is hereby incorporated by reference.

The terms "nucleic acid" or "nucleic acid molecule" encompass both RNA (plus and minus strands) and DNA, including cDNA, genomic DNA, and synthetic (e.g., chemically synthesized) DNA. The nucleic acid may be double-stranded or single-stranded. Where single-stranded, the nucleic acid may be the sense strand or the antisense strand. A nucleic acid molecule may be any chain of two or more covalently bonded nucleotides, including naturally occurring or non-naturally occurring nucleotides, or nucleotide analogs or derivatives. By "RNA" is meant a sequence of two or more covalently bonded, naturally occurring or modified ribonucleotides. One example of a modified RNA included within this term is phosphorothioate RNA. By "DNA" is meant a sequence of two or more covalently bonded, naturally occurring or modified deoxyribonucleotides. By "cDNA" is meant complementary or copy DNA produced from an RNA template by the action of RNA-dependent DNA polymerase (reverse transcriptase). Thus a "cDNA clone" means a duplex DNA sequence complementary to an RNA molecule of interest, carried in a cloning vector.

An "isolated nucleic acid" is a nucleic acid molecule that is free of the nucleic acid molecules that normally flank it in the genome or that is free of the organism in which it is normally found. Therefore, an "isolated" gene or nucleic acid molecule is in some cases intended to mean a gene or nucleic acid molecule which is not flanked by nucleic acid molecules which normally (in nature) flank the gene or nucleic acid molecule (such as in genomic sequences) and/or has been completely or partially purified from other transcribed sequences (as in a cDNA or RNA library). In some cases, an isolated nucleic acid molecule is intended to mean the genome of an organism such as a virus. An isolated nucleic acid of the invention may be substantially isolated with respect to the complex cellular milieu in which it naturally occurs. In some instances, the isolated material will form part of a composition (for example, a crude extract containing other substances), buffer system or reagent mix. In other circumstances, the material may be purified to essential homogeneity, for example as determined by PAGE or column chromatography such as HPLC. The term therefore includes, e.g., a genome; a recombinant nucleic acid incorporated into a vector, such as an autonomously replicating plasmid or virus; or into the genomic DNA of a prokaryote or eukaryote, or which exists as a separate molecule (e.g., a cDNA or a genomic DNA fragment produced by PCR or restriction endonuclease treatment) independent of other sequences. It also includes a recombinant nucleic acid which is part of a hybrid gene encoding additional polypeptide sequences. Preferably, an isolated nucleic acid comprises at least about 50, 80 or 90 percent (on a molar basis) of all macromolecular species present. Thus, an isolated gene or nucleic acid molecule can include a gene or nucleic acid molecule which is synthesized chemically or by recombinant means. Recombinant DNA contained in a vector are included in the definition of "isolated" as used herein. Also, isolated nucleic acid molecules include recombinant DNA molecules in heterologous host cells, as well as partially or substantially purified DNA molecules in solution. In vivo and in vitro RNA transcripts of the DNA molecules of the present invention are also encompassed by "isolated" nucleic acid molecules. Such isolated nucleic acid molecules are useful in the manufacture of the encoded polypeptide, as probes for isolating homologous

sequences (e.g., from other species), for gene mapping (e.g., by in situ hybridization with chromosomes), or for detecting expression of the nucleic acid molecule in tissue (e.g., human tissue, such as peripheral blood), such as by Northern blot analysis.

Various genes and nucleic acid sequences of the invention may be recombinant sequences. The term "recombinant" means that something has been recombined, so that when made in reference to a nucleic acid construct the term refers to a molecule that is comprised of nucleic acid sequences that are joined together or produced by means of molecular biological techniques. The term "recombinant" when made in reference to a protein or a polypeptide refers to a protein or polypeptide molecule which is expressed using a recombinant nucleic acid construct created by means of molecular biological techniques. The term "recombinant" when made in reference to genetic composition refers to a gamete or progeny with new combinations of alleles that did not occur in the parental genomes. Recombinant nucleic acid constructs may include a nucleotide sequence which is ligated to, or is manipulated to become ligated to, a nucleic acid sequence to which it is not ligated in nature, or to which it is ligated at a different location in nature. Referring to a nucleic acid construct as "recombinant" therefore indicates that the nucleic acid molecule has been manipulated using genetic engineering, i.e. by human intervention. Recombinant nucleic acid constructs may for example be introduced into a host cell by transformation. Such recombinant nucleic acid constructs may include sequences derived from the same host cell species or from different host cell species, which have been isolated and reintroduced into cells of the host species. Recombinant nucleic acid construct sequences may become integrated into a host cell genome, either as a result of the original transformation of the host cells, or as the result of subsequent recombination and/or repair events.

As used herein, "heterologous" in reference to a nucleic acid or protein is a molecule that has been manipulated by human intervention so that it is located in a place other than the place in which it is naturally found. For example, a nucleic acid sequence from one species may be introduced into the genome of another species, or a nucleic acid sequence from one genomic locus may be moved to another genomic or extrachromosomal locus in the same species. A heterologous protein includes, for example, a protein expressed from a heterologous coding sequence or a protein expressed from a recombinant gene in a cell that would not naturally express the protein.

By "antisense," as used herein in reference to nucleic acids, is meant a nucleic acid sequence that is complementary to one strand of a nucleic acid molecule. In some embodiments, an antisense sequence is complementary to the coding strand of a gene, preferably, a SARS virus gene. The preferred antisense nucleic acid molecule is one which is capable of lowering the level of polypeptide encoded by the complementary gene when both are expressed in a cell. In some embodiments, the polypeptide level is lowered by at least 10%, or at least 25%, or at least 50%, as compared to the polypeptide level in a cell expressing only the gene, and not the complementary antisense nucleic acid molecule.

A "probe" or "primer" is a single-stranded DNA or RNA molecule of defined sequence that can base pair to a second DNA or RNA molecule that contains a complementary sequence (the target). The stability of the resulting hybrid molecule depends upon the extent of the base pairing that occurs, and is affected by parameters such as the degree of complementarity between the probe and target molecule, and the degree of stringency of the hybridization conditions. The

degree of hybridization stringency is affected by parameters such as the temperature, salt concentration, and concentration of organic molecules, such as formamide, and is determined by methods that are known to those skilled in the art. Probes or primers specific for SARS virus nucleic acid sequences or molecules may vary in length from at least 8 nucleotides to over 500 nucleotides, including any value in between, depending on the purpose for which, and conditions under which, the probe or primer is used. For example, a probe or primer may be 8, 10, 15, 20, or 25 nucleotides in length, or may be at least 30, 40, 50, or 60 nucleotides in length, or maybe over 100, 200, 500, or 1000 nucleotides in length. Probes or primers specific for SARS virus nucleic acid molecules may have greater than 20-30% sequence identity, or at least 55-75% sequence identity, or at least 75-85% sequence identity, or at least 85-99% sequence identity, or 100% sequence identity to the nucleic acid sequences described herein. In various embodiments of the invention, probes having the sequences: 5'-ATg AAT TAC CAA gTC AAT ggT TAC-3', SEQ ID NO: 160; 5'-gAA gCT ATT CgT CAC gTT Cg-3', SEQ ID NO: 161; 5'-CTg TAg AAA ATC CTA gCT ggA g-3', SEQ ID NO: 162; 5'-CAT AAC CAg TCg gTACAg CTA-3', SEQ ID NO: 163; 5'-TTA TCA CCC gCgAAg AAg CT-3', SEQ ID NO: 164; 5'-CTC TAg TTg CATGAC AgC CCT C-3', SEQ ID NO: 165; 5'-TCg TgC gTg gAT TggCTT TgA TgT-3', SEQ ID NO: 166; 5'-ggg TTg ggA CTA TCC TAA gTg TgA-3', SEQ ID NO: 167; 5'-TAA CAC ACA AAC ACC ATC ATC A-3', SEQ ID NO: 168; 5'-ggT Tgg gAC TAT CCT AAg TgT gA-3', SEQ ID NO: 169; 5'-CCA TCA TCA gAT AgA ATC ATC ATA-3', SEQ ID NO: 170; 5'-CCT CTC TTg TTC TTg CTC gCA-3', SEQ ID NO: 171; 5'-TAT AgT gAg CCg CCA CAC Atg-3', SEQ ID NO: 172; 5'-TAACA-CACAACICCATCATCA-3', SEQ ID NO: 173; 5'-CTAA-CATGCTTAGGATAATGG-3', SEQ ID NO: 174; 5'-GC-CTCTTTGTTCTTGCTCGC-3', SEQ ID NO: 175; 5'-CAGGTAAGCGTAAAACATC-3', SEQ ID NO: 176; 5'-TACACACCTCAGCGTTG-3', SEQ ID NO: 177; 5'-CACGAACGTGACGAAT-3', SEQ ID NO: 178; 5'-GC-CGGAGCTCTGCAGAAATC-3', SEQ ID NO: 179; 5'-CAG-GAAACAGCTATGAC TTGCATCACCAGTGTGTC-CACCAGGTT-3', SEQ ID NO: 180; 5'-TGTAACACGACGGCCAGTTGATGG-GATGGGACTATCCTAAGTGTGA-3', SEQ ID NO: 181; 5'-GCATAGGCAGTAGTTGCATC-3', SEQ ID NO: 182, as well as sequences amplified by specific combinations of these probes, may be excluded from specific uses according to the invention. Probes can be detectably-labeled, either radioactively or non-radioactively, by methods that are known to those skilled in the art. Probes can be used for methods involving nucleic acid hybridization, such as nucleic acid sequencing, nucleic acid amplification by the polymerase chain reaction, single stranded conformational polymorphism (SSCP) analysis, restriction fragment polymorphism (RFLP) analysis, Southern hybridization, northern hybridization, in situ hybridization, electrophoretic mobility shift assay (EMSA), and other methods that are known to those skilled in the art.

By "complementary" is meant that two nucleic acid molecules, e.g., DNA or RNA, contain a sufficient number of nucleotides that are capable of forming Watson-Crick base pairs to produce a region of double-strandedness between the two nucleic acids. Thus, adenine in one strand of DNA or RNA pairs with thymine in an opposing complementary DNA strand or with uracil in an opposing complementary RNA strand. It will be understood that each nucleotide in a

nucleic acid molecule need not form a matched Watson-Crick base pair with a nucleotide in an opposing complementary strand to form a duplex.

By "vector" is meant a DNA molecule derived, e.g., from a plasmid, bacteriophage, or mammalian or insect virus, or artificial chromosome, that may be used to introduce a polypeptide, for example a SARS virus polypeptide, into a host cell by means of replication or expression of an operably linked heterologous nucleic acid molecule. By "operably linked" is meant that a nucleic acid molecule such as a gene and one or more regulatory sequences (e.g., promoters, ribosomal binding sites, terminators in prokaryotes; promoters, terminators, enhancers in eukaryotes; leader sequences, etc.) are connected in such a way as to permit the desired function e.g. gene expression when the appropriate molecules (e.g., transcriptional activator proteins) are bound to the regulatory sequences. A vector may contain one or more unique restriction sites and may be capable of autonomous replication in a defined host or vehicle organism such that the cloned sequence is reproducible. By "DNA expression vector" is meant any autonomous element capable of directing the synthesis of a recombinant peptide. Such DNA expression vectors include bacterial plasmids and phages and mammalian and insect plasmids and viruses. A "shuttle vector" is understood as meaning a vector which can be propagated in at least two different cell types, or organisms, for example vectors which are first propagated or replicated in prokaryotes in order for, for example, subsequent transfection into eukaryotic cells. A "replicon" is a unit that is capable of autonomous replication in a cell and may include plasmids, chromosomes (e.g., mini-chromosomes), cosmids, viruses, etc. A replicon may be a vector.

A "host cell" is any cell, including a prokaryotic or eukaryotic cell, into which a replicon, such as a vector, has been introduced by for example transformation, transfection, or infection.

An "open reading frame" or "ORF" is a nucleic acid sequence that encodes a polypeptide. An ORF may include a coding sequence having i.e., a sequence that is capable of being transcribed into mRNA and/or translated into a protein when combined with the appropriate regulatory sequences. In general, a coding sequence includes a 5' translation start codon and a 3' translation stop codon.

A "leader sequence" is a relatively short nucleotide sequence located at the 5' end of an RNA molecule that acts as a primer for transcription.

A "transcriptional regulatory sequence" "TRS" or "intergenic sequence" is a nucleotide sequence that lies upstream of an open reading frame (ORF) and serves as a template for the reassociation of a nascent RNA strand-polymerase complex.

A "frameshift mutation" is caused by a shift in an open reading frame, generally due to a deletion or addition of at least one nucleotide, such that an alternative polypeptide is ultimately translated.

By "detectably labeled" is meant any means for marking and identifying the presence of a molecule, e.g., an oligonucleotide probe or primer, a gene or fragment thereof, a cDNA molecule, a polypeptide, or an antibody. Methods for detectably-labeling a molecule are well known in the art and include, without limitation, radioactive labeling (e.g., with an isotope such as ³²P or ³⁵S) and nonradioactive labeling such as, enzymatic labeling (for example, using horseradish peroxidase or alkaline phosphatase), chemiluminescent labeling, fluorescent labeling (for example, using fluorescein), bioluminescent labeling, antibody detection of a ligand attached to the probe, or detection of double-stranded nucleic acid. Also included in this definition is a molecule that is

detectably labeled by an indirect means, for example, a molecule that is bound with a first moiety (such as biotin) that is, in turn, bound to a second moiety that may be observed or assayed (such as fluorescein-labeled streptavidin). Labels also include digoxigenin, luciferases, and aequorin.

A "peptide," "protein," "polyprotein" or "polypeptide" is any chain of two or more amino acids, including naturally occurring or non-naturally occurring amino acids or amino acid analogues, regardless of post-translational modification (e.g., glycosylation or phosphorylation). An "polyprotein", "polypeptide", "peptide" or "protein" of the invention may include peptides or proteins that have abnormal linkages, cross links and end caps, non-peptidyl bonds or alternative modifying groups. Such modified peptides are also within the scope of the invention. The term "modifying group" is intended to include structures that are directly attached to the peptidic structure (e.g., by covalent coupling), as well as those that are indirectly attached to the peptidic structure (e.g., by a stable non-covalent association or by covalent coupling to additional amino acid residues, or mimetics, analogues or derivatives thereof, which may flank the core peptidic structure). For example, the modifying group can be coupled to the amino-terminus or carboxy-terminus of a peptidic structure, or to a peptidic or peptidomimetic region flanking the core domain. Alternatively, the modifying group can be coupled to a side chain of at least one amino acid residue of a peptidic structure, or to a peptidic or peptidomimetic region flanking the core domain (e.g., through the epsilon amino group of a lysyl residue(s), through the carboxyl group of an aspartic acid residue(s) or a glutamic acid residue(s), through a hydroxy group of a tyrosyl residue(s), a serine residue(s) or a threonine residue(s) or other suitable reactive group on an amino acid side chain). Modifying groups covalently coupled to the peptidic structure can be attached by means and using methods well known in the art for linking chemical structures, including, for example, amide, alkylamino, carbamate or urea bonds.

A "polyprotein" is the polypeptide that is initially translated from the genome of a plus-stranded RNA virus, for example, a SARS virus. Accordingly, a polyprotein has not been subjected to post-translational processing by proteolytic cleavage into its processed protein products, and therefore, retains its cleavage sites. In some embodiments of the invention, the protease cleavage sites of a polyprotein may be modified, for example, by amino acid substitution, to result in a polyprotein that is incapable of being cleaved into its processed protein products.

An antibody "specifically binds" or "selectively binds" an antigen when it recognizes and binds the antigen, but does not substantially recognize and bind other molecules in a sample, having for example an affinity for the antigen which is 10, 100, 1000 or 10000 times greater than the affinity of the antibody for another reference molecule in a sample. A "neutralizing antibody" is an antibody that selectively interferes with any of the biological activities of a SARS virus polypeptide or polyprotein, for example, replication of the SARS virus, or infection of host cells. A neutralizing antibody may reduce the ability of a SARS virus polypeptide to carry out its specific biological activity by about 50%, or by about 70%, or by about 90% or more, or may completely abolish the ability of a SARS virus polypeptide to carry out its specific biological activity. Any standard assay for the biological activity of any SARS virus polypeptide, for example, assays determining expression levels, ability to infect host cells, or ability to replicate DNA, including those assays described herein or

known to those of skill in the art, may be used to assess potentially neutralizing antibodies that are specific for SARS virus polypeptides.

A "signal sequence" is a sequence of amino acids that may be identified, for example by homology or biological activity to a peptide sequence with the known function of targeting a polypeptide to a particular region of the cell. A signal sequence or signal peptide may be a peptide of any length, that is capable of targeting a polypeptide to a particular region of the cell. In some embodiments, the signal sequence may direct the polypeptide to the cellular membrane so that the polypeptide may be secreted. In alternate embodiments, the signal sequence may direct the polypeptide to an intracellular compartment or organelle, such as the Golgi apparatus, or to the surface of a virus, such as the SARS virus. In alternate embodiments, a signal sequence may range from about 13 or 15 amino acids in length to about 60 amino acids in length.

A "transmembrane protein" is an amphipathic protein having a hydrophobic region ("transmembrane domain") that spans the lipid bilayer of the cell membrane from the cytoplasm to the cell surface, or spans the viral envelope, interspersed between hydrophilic regions on both sides of the membrane. The number of hydrophobic regions in an amphipathic protein is often proportional to the number of times that proteins spans the lipid bilayer. Thus, a single transmembrane protein spans the lipid bilayer once, and has a single transmembrane domain, while a multi-transmembrane protein spans the lipid bilayer multiple times. Multi-transmembrane proteins may enable virus entry into a host cell, or act to initiate transduction of a signal from the cell surface to the interior of the cell, for example, by a conformational change upon ligand binding. A "transmembrane anchor" is a transmembrane domain that maintains a polypeptide in its position in the cell membrane or viral envelope and is generally hydrophobic. A transmembrane anchor may generally be in the structure of an alpha helix, i.e., a "transmembrane helix". Multi-transmembrane proteins may have multiple transmembrane alpha-helices.

A "nuclear localization signal" is an amino acid sequence that permits the entry of a polypeptide into the nucleus of a cell through nuclear pores. A nuclear localization signal generally has a cluster of positively charged residues, for example, lysines. A "lysine-rich sequence" is a sequence having at least two contiguous lysine residues, or at least three contiguous lysine residues. In some embodiments, a lysine-rich sequence may be a nuclear localization signal.

An "ATP binding domain" is a consensus domain that is found in many ATP or GTP-binding proteins, and that forms a flexible loop (P-loop) between alpha-helical and beta pleated sheet domains. The general consensus for an ATP binding domain may be (A or G)-XXXXGK-(S or T).

A "RNA binding protein" is a protein that is capable of binding to a RNA molecule (see, for example, "RNA Binding Proteins: New Concepts in Gene Regulation" 1st ed, eds. K. Sandberg and S. E. Mulrone, Kluwers Academic Publishers, 2001). RNA binding proteins may contain common structural features such as arginine-rich tracts, for example, arginines alternating with aspartates, serines, or glycines, or zinc finger regions. RNA binding proteins may also have a common ribonucleotide sequence domain. RNA binding proteins are believed to play diverse roles in modulating post-transcriptional gene expression.

An "immune response" includes, but is not limited to, one or more of the following responses in a mammal: induction of antibodies, B cells, T cells (including helper T cells, suppressor T cells, cytotoxic T cells, $\gamma\delta$ T cells) directed specifically to the antigen(s) in a composition or vaccine, following

administration of the composition or vaccine. An immune response to a composition or vaccine thus generally includes the development in the host mammal of a cellular and/or antibody-mediated response to the composition or vaccine of interest. In general, the immune response will result in prevention or reduction of infection by a SARS virus.

An "immunogenic fragment" of a polypeptide or nucleic acid molecule refers to an amino acid or nucleotide sequence that elicits an immune response. Thus, an immunogenic fragment may include, without limitation, any portion of any of the SARS virus sequences described herein, or a sequence substantially identical thereto, that includes one or more epitopes (the antigenic determinant i.e., site recognized by a specific immune system cell, such as a T cell or a B cell). An "epitope" may include amino acids in a spatial orientation that they are non-contiguous in the amino acid sequence but are near each other due to the three dimensional conformation of the polypeptide. An epitope may include at least 3, 5, 8, or 10 or more amino acids. Immunogenic fragments or epitopes may be identified using standard methods known to those of skill in the art, such as epitope mapping techniques or antigenicity or hydrophathy plots using, for example, the Omega version 1.0 program from Oxford Molecular Group (see, for example, U.S. Pat. No. 4,708,871). Immunogenic fragments or epitopes may also be identified using methods for determining three dimensional molecule structure such as X-ray crystallography or nuclear magnetic resonance.

A "sample" may be a tissue biopsy, amniotic fluid, cell, blood, serum, plasma, urine, stool, sputum, conjunctiva, or any other specimen, or any extract thereof, obtained from a patient (human or animal), test subject, or experimental animal. A "sample" may also be a cell or cell line created under experimental conditions, and constituents thereof (such as cell culture supernatants, cell fractions, infected cells, etc.). The sample may be analyzed to detect the presence of a SARS virus gene, genome, polypeptide, nucleic acid molecule or virion, or to detect a mutation in a SARS virus gene, expression levels of a SARS virus gene or polypeptide, or the biological function of a SARS virus polypeptide, using methods that are known in the art. For example, methods such as sequencing, single-strand conformational polymorphism (SSCP) analysis, or restriction fragment length polymorphism (RFLP) analysis of PCR products derived from a sample can be used to detect a mutation in a SARS virus gene; ELISA or western blotting can be used to measure levels of SARS virus polypeptide or antibody affinity; northern blotting can be used to measure SARS mRNA levels, or PCR can be used to measure the level of a SARS virus nucleic acid molecule.

Other features and advantages of the invention will be apparent from the following description of the drawings and the invention, and from the claims.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1A-D show phylogenetic analyses of SARS proteins. Unrooted phylogenetic trees were generated by clustalw (Thompson, J. D. et al., *Nucleic Acids Res* 22, 4673-80, Nov. 11, 1994) bootstrap analysis using 1000 iterations. Genbank accessions for protein sequences are as follows: FIG. 1A: Replicase 1A: BoCov (Bovine Coronavirus): AAL40396, 229E (Human Coronavirus): NP_07355, MHV (Mouse Hepatitis Virus): NP_045298, AIBV (Avian Infectious bronchitis virus): CAC39113, TGEV (Transmissible Gastroenteritis Virus): NP_058423. FIG. 1B: Matrix Glycoprotein: PHEV (Porcine hemagglutinating encephalomyelitis virus): AAL80035, BoCov (Bovine Coronavirus):

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NP_150082, AIBV & AIBV2 (Avian infectious bronchitis virus): AAF35863 & AAK83027, MHV (Mouse hepatitis virus): AAF36439, TGEV (Transmissible gastroenteritis virus): NP_058427, 229E & OC43 (Human Coronavirus): NP_073555 & AAA45462, FCV (Feline coronavirus): BAC01160. FIG. 1C: Nucleocapsid: MHV (Mouse hepatitis virus): P18446, BoCov (Bovine coronavirus): NP_150083, AIBV (Avian infectious bronchitis virus): AAK27162, FCV (Feline coronavirus): CAA74230, PTGV (Porcine transmissible gastroenteritis virus): AAM97563, 229E & OC43 (Human coronavirus): NP_073555 & P33469, PHEV (porcine hemagglutinating encephalomyelitis virus): AAL80036, TCV (Turkey coronavirus): AAF23873. FIG. 1D: S (Spike) Protein: BoCov (Bovine coronavirus): AAL40400, MHV (Mouse hepatitis virus): P11225, OC43 & 229E (Human coronavirus): S44241 & AAK32191, PHEV (Porcine hemagglutinating encephalomyelitis virus): AAL80031, PRC (Porcine respiratory coronavirus): AAA46905, PEDV (Porcine epidemic diarrhea virus): CAA80971, CCov (Canine coronavirus): S41453, FICV (Feline infectious peritonitis virus): BAA06805, AIBV (Avian infectious bronchitis virus): AA034396.

FIG. 2 shows a schematic representation of the ORFs and s2m motif in the 29,736-base SARS virus genome.

FIGS. 3A-P show nucleotide sequences of the 29,736-base genome of the SARS virus (SEQ ID NOs: 1 and 2).

FIG. 4 shows an alignment of the s2m regions from Avian infectious bronchitis virus (AIBV; SEQ ID NO: 32) and equine rhinovirus serotype 2 (ERV-2; SEQ ID NO: 31) with the 3' untranslated region (UTR; SEQ ID NO: 18) of the SARS virus (TOR2). The conserved areas in the s2m region are indicated by asterisks.

FIG. 5 shows the amino acid sequence of the SARS virus S (Spike) Glycoprotein (SEQ ID NO: 33).

FIG. 6 shows the amino acid sequence of the SARS virus M (Matrix) Glycoprotein (residues 1-220 of SEQ ID NO: 34).

FIG. 7 shows the amino acid sequence of the SARS virus E (Small envelope) protein (SEQ ID NO: 35).

FIG. 8 shows the amino acid sequence of the SARS virus N (Nucleocapsid) Protein (SEQ ID NO: 36).

FIG. 9 shows an alignment of the matrix glycoprotein M from the SARS virus (Tor2_M or ORF5; SEQ ID NO: 34) and various other matrix glycoproteins (SEQ ID NOs: 37-43). Asterisks (*) indicate percentage identity to the SARS matrix protein as calculated by Align (Myers and Miller, CABIOS (1989) 4:11-17).

FIGS. 10A-B show an alignment of the nucleocapsid protein N from the SARS virus (Tor2_N; SEQ ID NO: 36) and various other nucleocapsid proteins (SEQ ID NOs: 44-52; and SEQ ID NO: 199 of AIBV2 nucleocapsid protein [Avian infectious bronchitis virus 2]). Asterisks (*) indicate percentage identity to the SARS nucleocapsid protein calculated by Align (Myers and Miller, CABIOS (1989) 4:11-17).

FIGS. 11A-K show the nucleotide sequence of the 29,751-base genome of the SARS virus (SEQ ID NO: 15).

FIG. 12 shows a schematic representation of the ORFs and s2m motif in the 29,751-base SARS virus genome.

FIGS. 13A-D show phylogenetic analyses of SARS proteins. Unrooted phylogenetic trees were generated by clustalw 1.74 (J. D. Thompson, D. G. Higgins, T. J. Gibson, Nucleic Acids Res 22, 4673-80 (Nov. 11, 1994) using the BLOSUM comparison matrix and a bootstrap analysis of 1000 iterations. Numbers indicate bootstrap replicates supporting each node. Phylogenetic trees were drawn with the Phylip Drawtree program 3.6a3 (Felsenstein, J. 1993. PHYLIP (Phylogeny Inference Package) version 3.5c. Distributed by the author. Department of Genetics, University of

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Washington, Seattle). Branch lengths indicate the number of substitutions per residue. Genbank accessions for protein sequences: A: Replicase 1A: BoCoV (Bovine Coronavirus): AAL40396, HCoV-229E (Human Coronavirus): NP_073555, MHV (Mouse Hepatitis Virus): NP_045298, IBV (Avian Infectious bronchitis virus): CAC39113, TGEV (Transmissible Gastroenteritis Virus): NP_058423. B: Membrane Glycoprotein: PHEV (Porcine hemagglutinating encephalomyelitis virus): AAL80035, BoCoV (Bovine Coronavirus): NP_150082, IBV & IBV2 (Avian infectious bronchitis virus): AAF35863 & AAK83027, MHV (Mouse hepatitis virus): AAF36439, TGEV (Transmissible gastroenteritis virus): NP_058427, HCoV-229E & HCoV-OC43 (Human Coronavirus): NP_073555 & AAA45462, FCov (Feline coronavirus): BAC01160. C: Nucleocapsid: MHV (Mouse hepatitis virus): P18446, BoCoV (Bovine coronavirus): NP_150083, IBV 1 & 2 (Avian infectious bronchitis virus): AAK27162 & NP_040838, FCov (Feline coronavirus): CAA74230, PTGV (Porcine transmissible gastroenteritis virus): AAM97563, HCoV-229E & HCoV-OC43 (Human coronavirus): NP_073555 & P33469, PHEV (porcine hemagglutinating encephalomyelitis virus): AAL80036, TCV (Turkey coronavirus): AAF23873. D: S (Spike) Protein: BoCoV (Bovine coronavirus): AAL40400, MHV (Mouse hepatitis virus): P11225, HCoV-OC43 & HCoV-229E (Human coronavirus): S44241 & AAK32191, PHEV (Porcine hemagglutinating encephalomyelitis virus): AAL80031, PRCOV (Porcine respiratory coronavirus): AAA46905, PEDV (Porcine epidemic diarrhea virus): CAA80971, CCov (Canine coronavirus): S41453, FIPV (Feline infectious peritonitis virus): BAA06805, IBV (Avian infectious bronchitis virus): AA034396.

FIGS. 14A-F show an alignment of the spike glycoprotein S from the SARS virus (Tor2_S; SEQ ID NO: 33) and various other spike glycoproteins (SEQ ID NOs: 53-62). Asterisks (*) indicate percentage identity to the SARS spike protein as calculated by Align (Myers and Miller, CABIOS (1989) 4:11-17).

FIG. 15 shows an alignment between the SARS virus Small envelope protein E (TOR2_E; SEQ ID NO: 35) and the Envelope protein (Protein 4) (X1 protein) (ORF 3) from Porcine transmissible gastroenteritis coronavirus (strain Purdue). Swissprot accession number P09048 (PGV; SEQ ID NO: 63), as calculated by FASTA (world wide web at ebi "dot" ac "dot" uk "forward slash" fasta33).

FIGS. 16A-B show the amino acid sequence of the SARS virus Replicase 1A protein (SEQ ID NO: 64).

FIG. 17 shows the amino acid sequence of the SARS virus Replicase 1B protein (SEQ ID NO: 65).

FIG. 18 shows the amino acid sequence of ORF3 of SARS virus (SEQ ID NO: 66).

FIG. 19 shows the amino acid sequence of ORF4 of SARS virus (SEQ ID NO: 67).

FIG. 20 shows the amino acid sequence (SEQ ID NO: 68) of ORF6 (nucleotides 27059-27247 of the 29,736-base genome sequence) or ORF 7 (nucleotides 27,074-27,265 of the 29,751-base genome sequence) of SARS virus.

FIG. 21 shows the amino acid sequence (SEQ ID NO: 69) of ORF7 (nucleotides 27258-27623 of the 29,736-base genome sequence) or ORF 8 (nucleotides 27,273-27,641 of the 29,751-base genome sequence), of SARS virus.

FIG. 22 shows the amino acid sequence (SEQ ID NO: 70) of ORF8 (nucleotides 27623-27754 of the 29,736-base genome sequence) or ORF9 8 (nucleotides 27,638-27,772 of the 29,751-base genome sequence) of SARS virus.

FIG. 23 shows the amino acid sequence (SEQ ID NO: 71) of ORF9 (nucleotides 27764-27880 of the 29,736-base

genome sequence) or ORF10 (nucleotides 27,779-27,898 of the 29,751-base genome sequence) of SARS virus.

FIG. 24 shows the amino acid sequence (SEQ ID NO: 72) of ORF10 (nucleotides 27849-28100 of the 29,736-base genome sequence) or ORF11 (nucleotides 27,864-28118 of the 29,751-base genome sequence) of SARS virus.

FIG. 25 shows the amino acid sequence of ORF13 of SARS virus (SEQ ID NO: 73).

FIG. 26 shows the amino acid sequence of ORF14 of SARS virus (SEQ ID NO: 74).

FIG. 27 shows an alignment of the secreted region of the SARS virus ORF 10 (SEQ ID NO: 201) of the 29,751-base genome sequence (sars) with the conotoxin from *Conus ventricosus* (conotoxin) (SEQ ID NO: 200). Sequence identity is indicated by asterisks and sequence homology is indicated by dots.

DETAILED DESCRIPTION OF THE INVENTION

In general, the invention provides nucleic acid molecules, polypeptides, and other reagents derived from a SARS virus, as well as methods of using such nucleic acid molecules, polypeptides, and other reagents.

The genome sequence (FIGS. 3A-P, 11A-K, SEQ ID NOs: 1, 2, and 15) reveals that the SARS coronavirus is only moderately related to other known coronaviruses, including two human coronaviruses, OC43 and 229E. Thus, the SARS virus is a previously unknown virus. The 5' end of the SARS genome contains a 5' leader sequence (Table 1; SEQ ID NO: 3) with sequence similarity to the highly conserved coronavirus core leader sequence, 5'-CUAAAC-3 (SEQ ID NO: 75;

Sawicki, S. G., et al., *Adv Exp Med Biol* 440, 215-9, 1998; Lai, M. M. and D. Cavanagh, *Adv Virus Res* 48, 1-100, 1997). Transcriptional regulatory sequences (TRSs) were identified upstream of all open reading frames (ORFs) (Tables 1 and 2; SEQ ID NOs: 3-13 and 20-30). ORF9 and ORF10 of the 29,736-base SARS genome (ORF 10 and ORF 11 of the 29,751 base genome) overlap by 12 amino acids, and have matches to the TRS consensus in close proximity to their respective initiating methionine codons.

The 3' UTR sequence (SEQ ID NO: 18) of SARS virus contains a s2m region having the sequence ACATTTTCATC-GAGGCCACGCGGAGTACGAT CGAGGGTACAGT-GAAT; SEQ ID NO: 16) that includes a conserved, discontinuous 32 base-pair s2m motif. The conserved 32 base-pair motif is a universal feature of astroviruses that has also been identified in avian coronavirus (AIBV) and the ERV-2 equine rhinovirus. This motif has been identified by Jonassen C. M. et al. (*J Gen Virol* 1998 April; 79 (Pt 4):715-8) as GCCGNG-GCCACGC(G/C)GAGTA(C/G)GANCGAGGGTACAG(G/C) (SEQ ID NO: 19), where N is generally not part of the conserved motif, and can be any nucleotide. The region corresponding to the 32 base-pair motif in SARS virus includes the sequence: CGAGGCCACGCGGAGTACGATC-GAGGGTACAG (SEQ ID NO: 17), and spans positions 29590-29621 of the 29,751 base genome. FIG. 4 shows an alignment of the s2m regions from Avian infectious bronchitis virus (AIBV; SEQ ID NO: 32) and equine rhinovirus serotype 2 (ERV-2; SEQ ID NO: 31), as defined in Jonassen C. M. et al. (*J Gen Virol* 1998 April; 79 (Pt 4):715-8), with the entire 3' untranslated region (UTR) of the SARS virus (TOR2) (SEQ ID NO: 18).

TABLE 1

Listing of the transcription regulatory sequences of the 29,736-base SARS genome, showing the nucleotide position (base) and associated open-reading frames (ORF). An asterisk (*) indicates consensus sequence.

Base	ORF	TRS Sequence	
45	Leader	TCTCTAAACGAACTTTAAAATCTGTG	(SEQ ID NO: 3)
21464	S	CAACTAAACGAACATG	(SEQ ID NO: 4)
25238	ORF3	CACATAAACGAACTTATG	(SEQ ID NO: 5)
26089	E	TGAGTACGAACTTATG	(SEQ ID NO: 6)
26326	M	GGTCTAAACGAACTAACT 40 ATG	(SEQ ID NO: 7)
26986	ORF6	AACTATAAATT 62 ATG	(SEQ ID NO: 8)
27244	ORF7	TCCATAAACGAACATG	(SEQ ID NO: 9)
27575	ORF8	TGCTCTA---GTATTTTTTAACTTTG 24 ATG	(SEQ ID NO: 10)
27751	ORF9	AGTCTAAACGAACATG	(SEQ ID NO: 11)
27837	ORF10	CTAATAAACCTCATG	(SEQ ID NO: 12)
28084	N	TAAATAAACGAACAAATTAAATG	(SEQ ID NO: 13)

TABLE 2

Listing of the transcription regulatory sequences of the 29,751-base SARS genome, showing the nucleotide position (base), associated open-reading frames (ORF), and identified transcription regulatory sequences. Numbers in parentheses within the alignment indicate distance to the putative initiating codon. The conserved core sequence is indicated in bold in the putative leader sequence. Contiguous sequences identical to region of the leader sequence containing the core sequence are shaded. No putative TRSs were detected for ORFs 4, 13 and 14, although ORF 13 may share the TRS associated with the N protein.

Base	ORF	TRS Sequence	
60	Leader	UCUCUAAACGAACUUUAAAUCUGUG	(SEQ ID NO: 20)
21479	S (Spike)	CAACU AAAACGAA CAUG	(SEQ ID NO: 21)
25252	ORF3	CACAU AAAACGAA CUUAUG	(SEQ ID NO: 22)
26104	Envelope	UGAGU ACGAA CUUAUG	(SEQ ID NO: 23)
26341	M	GGUCU AAAACGAA CUAACU (40) AUG	(SEQ ID NO: 24)
27001	ORF7	AGCU UAAAAU (62) AUG	(SEQ ID NO: 25)
27259	ORF8	UCCA UAAAACGAA CAUG	(SEQ ID NO: 26)
27590	ORF9	UGCU CUA--GUAUUUUUAUACUUUG (24) AUG	(SEQ ID NO: 27)
27766	ORF10	AGCU UAAAACGAA CAUG	(SEQ ID NO: 28)
27852	ORF11	CUA UAAAAC CUCAUG	(SEQ ID NO: 29)
28099	NUCLEOCAPSID	U AAAUAAAACGAA CAAAU UAAA AUG	(SEQ ID NO: 30)

The coding potentials of the 29,736-base and 29,751-base genomes are depicted in FIGS. 2 and 12, respectively. Open reading frames (ORFs) include the Replicase 1a and 1b translation products, the Spike glycoprotein, the small Envelope protein, the Membrane and the Nucleocapsid protein. Construction of unrooted phylogenetic trees using this set of known proteins from representatives of the three known coronavirus groups reveals that the proteins encoded by the SARS virus do not readily cluster more closely with any known group than with any other (FIGS. 1A-D and 13A-D). In addition, nine novel ORFs have been analyzed.

The Replicase 1a ORF located at nucleotides 250-13395 of the 29,736-base genome, and nucleotides 265-13,398 of the 29,751-base genome, and replicase 1b ORF located at nucleotides 13,398-21,485 of the 29,751-base genome, occupy 21.2 kb of the SARS virus genome (FIGS. 2 and 12). These genes encode a number of proteins that are produced by proteolytic cleavage of a large polyprotein (Ziebuhr, J. et al., *J Gen Virol* 81, 853-79, April, 2000). A frame shift mutation interrupts the protein-coding region, separating the 1a and 1b open-reading frames. The proteins encoded by the Replicase 1a and 1b ORFs are depicted in FIGS. 16A-B and 17, SEQ ID NOs: 64 and 65).

The Spike glycoprotein (S) (E2 glycoprotein gene; FIGS. 2 and 12; nucleotides 21477 to 25241 of the 29,736-base genome, and nucleotides 21,492 to 25,259 of the 29,751-base genome) encodes a surface projection glycoprotein precursor of about 1,255 amino acids in length (FIG. 5; SEQ ID NO: 33), which may be significant in the virulence of the SARS virus. Mutations in this gene are correlated with altered pathogenesis and virulence in other coronaviruses (B. N. Fields et al., *Fields virology* (Lippincott Williams & Wilkins, Philadelphia, ed. 4th, 2001). In other coronaviruses, the mature spike protein is inserted in the viral envelope with the majority of the protein exposed on the surface of the particles. Three molecules of the Spike protein form the characteristic peplomers or corona-like structures of this virus family. Analysis of the spike glycoprotein with SignalP (Nielson, H. et al., *Prot Engineer*. 10:1-6 (1997) indicates a signal peptide (MFIFLLFLTLTSG; SEQ ID NO: 76)(probability 0.996)

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with cleavage between residues 13 and 14. TMHMM (Sonhammer, E. L. et al., *Proc Int Conf Intell Syst Mol Biol* 6, 175-82 (1998)) indicates a transmembrane domain near the C-terminal end (WYVWLGFIAGLIAIVMVTILLCC; SEQ ID NO: 183). Together these data indicate a type I membrane protein with N-terminus and the majority of the protein (residues 14-1195) on the outside of the cell-surface or virus particle, which may be responsible for binding to a cellular receptor. The SARS virus Spike glycoprotein has limited sequence identity to other, known Spike glycoproteins (FIGS. 14A-F).

ORF 3 (FIGS. 2 and 12; nucleotides 25253-26074 of the 29,736-base genome and nucleotides 25,268-26,092 of the 29,751-base genome) encodes a protein of 274 amino acids (FIG. 18; SEQ ID NO: 66) that lacks significant similarities to any known protein when analyzed with BLAST (Altschul, S. F. et al., *Nucleic Acids Res* 25, 3389-402, Sep. 1, 1997), FASTA (Pearson, W. R. and D. J. Lipman, *Proc Natl Acad Sci USA* 85, 2444-8, April, 1988) or PFAM (Bateman, A. et al., *Nucleic Acids Res* 30, 276-80, Jan. 1, 2002). Analysis of the N-terminal 70 amino acids with SignalP indicates the existence of a signal peptide (MDLFMRFFTLRSITAQ; SEQ ID NO: 184) and a cleavage site (probability 0.540). Both TMpred (Hofinan, K. and W. Stoffel, *Biol. Chem. Hoopes-Seyler* 374, 166 (1993) and TMHMM indicate three transmembrane regions spanning approximately residues 34-56 (TIPLQASLPFGWLVIQVAFLAVF, SEQ ID NO: 77), 77-99 (FQFICNLLLLFVTTYSHLLLVAA, SEQ ID NO: 78), and 103-125 (AQFLYLYALIYFLQCINACRIIM, SEQ ID NO: 79). Both TMpred and TMHMM indicate that the C-terminus and a large 149 amino acid domain is located inside the viral or cellular membrane. The C-terminal (interior) region of the protein, corresponding to about amino acids 124-274 (MRCWLCWKCKSKNPLLYDANYFVCWHHT-NYDYCIPYNSVTDITIVVTEGDGI STPKLKEDYQIGGY-SEDRHSGVKDYVVVHGYFTEVYYQLEST-QITTDGTIENAT
FFIFNKLKVDPPNVQIHTIDGSSGVAN-PAMDPIYDEPTTTTSVPL; SEQ ID NO: 185) may encode a protein domain with ATP-binding properties (PD037277).

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ORF 4 (FIG. 12; nucleotides 25,689-26,153 of the 29,751-base genome) encodes a predicted protein of 154 amino acids (FIG. 19; SEQ ID NO: 67). This ORF overlaps entirely with ORF 3 and the E protein. ORF4 may be expressed from the ORF mRNA using an internal ribosomal entry site. BLAST analyses failed to identify matching sequences. Analysis with TMPred predicts a single transmembrane helix, amino acids 1-20 MMPTTLFAGTHITMTTVYHI, SEQ ID NO: 186.

The small envelope protein E (FIGS. 2 and 12; nucleotides 26102-26329 of the 29,736-base genome and nucleotides 26,117-26,347, ORF 5, of the 29,751-genome) encodes a protein of 76 amino acids (FIG. 7; SEQ ID NO: 35). BLAST and FASTA comparisons indicate that the protein, while novel, is homologous to multiple envelope proteins (alternatively known as small membrane proteins) from several coronaviruses. An alignment of the SARS virus E protein with the envelope protein of Porcine transmissible gastroenteritis coronavirus indicates approximately 28% sequence identity between the two proteins over a 61 amino acid overlap, as calculated by FASTA (FIG. 15). PFAM analysis of the protein indicates that the small envelope protein E is a member of the NS3_EnvE protein family. InterProScan (R. Apweiler et al., *Nucleic Acids Res* 29, 37-40, Jan. 1, 2001; Zdobnov, E. M. and R. Apweiler, *Bioinformatics* 17, 847-8, September, 2001) analysis indicates that the protein is a component of the viral envelope, and homologs of it are found in other viruses, including gastroenteritis virus and murine hepatitis virus. SignalP analysis indicates the presence of a transmembrane anchor (probability 0.939). TMPred analysis indicates a similar transmembrane anchor at positions 17-34 (VLLFLAFV-VLLVTLAIL, SEQ ID NO: 80), which is consistent with the known association of homologous proteins with the viral envelope. TMHMM indicates a type II membrane protein with the majority of the 46 residue C terminus hydrophilic domain (TALRLCAYCCNIVNVSLVKPTVYVYS-RVKNLNSSEGVPDLLV; SEQ ID NO: 187) located on the surface of the viral particle. The E protein may be important for viral replication.

The Matrix glycoprotein M (FIGS. 2 and 12; nucleotides 26383-27045 of the 29,736-base genome and nucleotides 26,398-27,063, ORF 6, of the 29,751-genome) encodes a protein of 221 amino acids (FIG. 6; SEQ ID NO: 34). BLAST and FASTA analysis of the protein, while novel, reveals homologies to coronaviral matrix glycoproteins (FIG. 9). The association of the spike glycoprotein (S) with the matrix glycoprotein (M) may be an essential step in the formation of the viral envelope and in the accumulation of both proteins at the site of virus assembly. Analysis of the amino acid sequence with SignalP indicates a signal sequence (probability 0.932), located at approximately residues 1-39 (MAD-NGTITVEELKQLLEQWNLVIGFLFLAWIMLLQFAYS; SEQ ID NO: 188) that is unlikely to be cleaved. TMHMM and TMPred analysis both indicate the presence of three transmembrane helices, located at approximately residues 15-37 (LLEQWNLVIGFLFLAWIMLLQFA; SEQ ID NO: 81), 50-72 (LVFLWLLWPVTLACFVLAAYRI; SEQ ID NO: 82) and 77-99 (GGIAIAMACIVGLMWLSYFVASF; SEQ ID NO: 83), with the 121 amino acid hydrophilic domain on the inside of the virus particle, where it may interact with nucleocapsid. The hydrophilic domain may run from approximately amino acids PLRGTIIVTRPLMESELVI-GAVIRGHLRMAGHSLGRCDIKDLPKEITVATSRITLS YYKLGASQVRVGTDSGFAAYN-RYRIGNYKLNTHAGSNDNIALLVQ (SEQ ID NO: 189) i.e. approximately amino acids 95 or 99 to 221 of SEQ ID NO: 34. PFAM analysis reveals a match to PFAM domain

PF01635, and alignments to 85 other sequences in the PFAM database bearing this domain, which is indicative of the coronavirus matrix glycoprotein.

ORF6 (FIG. 2; nucleotides 27059-27247 of the 29,736-base genome sequence) or ORF 7 (FIG. 12; nucleotides 27,074-27,265 of the 29,751-base genome sequence) encodes a protein of 63 amino acids (FIG. 20; SEQ ID NO: 68). TMpred analysis indicates a trans-membrane helix located between residues 3 or 4 and 22 (HLVDFQVTI-AEILIIIMRTF; SEQ ID NO: 84), with the N-terminus located outside the viral particle.

Similarly, the gene encoding ORF7 (FIG. 2; nucleotides 27258-27623 of the 29,736-base genome sequence) or ORF 8 (FIG. 12; nucleotides 27,273-27,641 of the 29,751-base genome sequence), encoding a protein of 122 amino acids (FIG. 21; SEQ ID NO: 69), has no significant BLAST or FASTA matches to known proteins. Analysis of this sequence with SignalP indicates a cleaved signal sequence (MKIIL-FLLIVFTSC; SEQ ID NO: 85) (probability 0.995), with the cleavage site located between residues 15 and 16. TMpred and TMHMM analysis also indicates a trans-membrane helix located approximately at residues 99-117 (SPLFLIVAALV-FLILCFTI; SEQ ID NO: 86). Together these data indicate that this protein is a type I membrane protein with the major hydrophilic domain of the protein (residues 16-98; ELY-HYQECVRGTTVLLKEPCP SGTYEGNSPFHPLADNK-FALCTCTSTHFAFACADGTRHTYQLRARSVSPKLFIRQ EEVQQELY; SEQ ID NO: 87) and the amino-terminus is oriented inside the lumen of the ER/Golgi, or on the surface of the cell membrane or virus particle, depending on the membrane localization of the protein.

ORF8 (FIG. 2; nucleotides 27623-27754 of the 29,736-base genome sequence) or ORF9 (FIG. 12; nucleotides 27,638-27,772 of the 29,751-base genome sequence), encodes a protein of 44 amino acids (FIG. 22; SEQ ID NO: 70). FASTA analysis of this sequence revealed some weak similarities (37% identity over a 35 amino acid overlap) to Swiss-Prot accession Q9M883, annotated as a putative sterol-C5 desaturase. A similarly weak match to a hypothetical *Clostridium perfringens* protein (Swiss-Prot accession CPE2366) was also detected. TMpred indicated a single strong trans-membrane helix FYLCFLAFLFLVLMILIIIF-WFS, SEQ ID NO: 190, with little preference for alternate models in which the N-terminus was located inside or outside the particle.

Similarly ORF9 (FIG. 2; nucleotides 27764-27880 of the 29,736-base genome sequence) or ORF10 (FIG. 12; nucleotides 27,779-27,898 of the 29,751-base genome sequence) encoding a protein of 39 amino acids (FIG. 23; SEQ ID NO: 71), exhibited no significant matches in BLAST and FASTA searches but encodes a trans-membrane helix LLIVLTCIS-LCSCICTVVQ (SEQ ID NO: 191) by TMPred, with the N-terminus located within the viral particle. The region immediately upstream of this protein exhibits a strong match to the TRS consensus (Table 2), indicating that a transcript initiates from this site. The large number of cysteine residues (6) may result in cross linking of the amino acids. Amino acids ICTVVQRCASNKPHVLEDPCKVQH (SEQ ID NO: 192) of this protein may be secreted. The secreted amino acids exhibit homology to toxin proteins, for example, to the conotoxin of *Conus ventricosus* (FIG. 27). Antigenic peptides from the hydrophilic (secreted) region, for example, CICTV-VQRCASNKPHVLEDPCK (SEQ ID NO: 193), were used to generate monoclonal antibodies using standard techniques. Furthermore, the C terminal amino acids form a sequence that shares homology to farnesylation sites (CKQH), which gen-

erally require C terminal location to be functional. This protein may act as a virulence factor and/or may facilitate transmission to humans.

ORF10 (FIG. 2; nucleotides 27849-28100 of the 29,736-base genome sequence) or ORF11 (FIG. 12; nucleotides 27,864-28118 of the 29,751-base genome sequence) encoding a protein of 84 amino acids (FIG. 24; SEQ ID NO: 72) exhibited only very short (9-10 residues) matches to a region of the human coronavirus E2 glycoprotein precursor (starting at residue 801). Analysis by SignalP and TMHMM predict a soluble protein. A detectable alignment to the TRS consensus sequence was also found (Table 2).

The protein (422 amino acids; FIG. 8; SEQ ID NO: 36) encoded by the Nucleocapsid gene (FIG. 2; nucleotides 28105-29370 of the 29,736-base genome sequence; FIG. 12, nucleotides 28,120-29,388 of the 29,751-base genome sequence) aligns well with nucleocapsid proteins from other representative coronaviruses (FIGS. 10A-B), although a short lysine rich region (KTFPPTEPKKDKKKKTDEAQ; SEQ ID NO: 14) is unique to SARS. This region is suggestive of a nuclear localization signal. Since some coronaviruses are able to replicate in enucleated cells, the SARS virus nucleocapsid protein may have evolved a novel nuclear function, which may play a role in pathogenesis. In addition, the basic nature of this peptide suggests it may assist in RNA binding. The SARS nucleocapsid protein is also a good candidate for diagnostic tests.

ORF13 (FIG. 12; nucleotides 28,130-28,426 of the 29,751-base genome sequence) encodes a novel protein of 98 amino acids (FIG. 25; SEQ ID NO: 73). ORF 14 (FIG. 12; nucleotides 28,583-28,795 of the 29,751-base genome sequence) encodes a novel protein of 70 amino acids (FIG. 26; SEQ ID NO: 74). TMPred predicts a single transmembrane helix VVAVIQEIQLLAAVGEILLLEW (SEQ ID NO: 194).

Various features of the SARS virus genome are summarized in Table 3. While Table 3 refers to the 29,751-base genome sequence, the features are also applicable to the 29,736-base genome sequence (SEQ ID NOs: 1 and 2).

TABLE 3

Features of the SARS virus 29,751-base genome sequence.

Feature	Start-End ¹	No. amino acids	No. bases	Frame	TRS
Orf 1a	265-13,398	4,382	13,149	+1	N/A
Orf 1b	13,398-21,485	2,628	7,887	+3	N/A
S protein	21,492-25,259	1,255	3,768	+3	Strong
Orf 3	25,268-26,092	274	825	+2	Strong
Orf 4	25,689-26,153	154	465	+3	Absent ²
E protein	26,117-26,347	76	231	+2	Weak
M protein	26,398-27,063	221	666	+1	Strong
Orf 7	27,074-27,265	63	192	+2	Weak
Orf 8	27,273-27,641	122	369	+3	Strong
Orf 9	27,638-27,772	44	135	+2	Weak
Orf 10	27,779-27,898	39	120	+2	Strong
Orf 11	27,864-28,118	84	255	+3	Weak
N protein	28,120-29,388	422	1,269	+1	Strong
Orf 13 ³	28,130-28,426	98	297	+2	Absent ²
Orf 14 ³	28,583-28,795	70	213	+2	Absent
s2m motif	29,590-29,621	N/A	30	N/A	N/A

¹End coordinates include the stop codon, except for ORF 1a and s2m.

²These ORFs overlap substantially or completely with other and may share TRSs.

N/A indicates not applicable.

Various polymorphisms may exist in the SARS virus. In the SARS 29,736-base genome sequences (SEQ ID NO: 1 or 2), for example, nucleotides 7904, 16607, 19168, 24857, or 26842 may be C or T; or nucleotides 19049, 23205, or 25283 may be G or A, and in the SARS 29,751-base genome

sequence (SEQ ID NO: 15), for example, nucleotides 7919, 16622, 19183, 24872, or 26857 may be C or T; or nucleotides 19064, 23220, or 25298 may be G or A. In some embodiments, the nucleotide changes may result in no change in the encoded amino acid, or in a conservative or non-conservative change in the encoded amino acid. In some embodiments, a nucleotide change, as described herein, at position 7904 or 7919, may result in a A to V amino acid substitution, in the Replicase 1A protein coding region; a change at position 19168 or 19183 may result in a V to A amino acid substitution, in the Replicase IB protein coding region; a change at position 23205 or 23220 may result in a A to S amino acid substitution (non-conservative change), affecting the Spike glycoprotein coding region; a change at position 25283 or 25298 may result in a R to G amino acid substitution (non-conservative change), affecting ORF3; or a change at position 26842 or 26857 may result in a S to P amino acid substitution (non-conservative change), affecting the Nucleocapsid protein coding region, in the SARS 29,736-base (SEQ ID NO: 1 or 2) and 29,751-base genome (SEQ ID NO: 15) sequences, respectively. In various embodiments, a nucleotide or amino acid sequence including a particular polymorphism may be selected, for example, for use in the methods of the invention, or may be excluded, for example, from a particular use according to the invention.

Various alternative embodiments of the invention are described below. These embodiments include, without limitation, identification and use of SARS virus nucleic acid and amino acid sequences for diagnostic or therapeutic uses.

Diagnosis of SARS Virus-Related Disorders

A SARS virus-related disorder is any disorder that is mediated by the SARS virus, or by a nucleic acid molecule or polypeptide derived from the SARS virus. Accordingly, SARS virus nucleic acid molecules and polypeptides may be used to diagnose and identify a SARS virus-related disorder in a mammal, for example, a human or a domestic, farm, wild, or experimental animal. In some embodiments, SARS virus nucleic acid molecules and polypeptides may be used to screen such animals, e.g., civet cats, for the presence of SARS virus. A SARS virus-related disorder may be a hepatic, enteric, respiratory, or neurological disorder, and may be accompanied by one or more symptoms or indications including, but not limited to, fever, cough, shortness of breath, headache, low blood oxygen concentration, liver damage, or reduced lymphocyte numbers. Accordingly, samples for diagnosis may be obtained from cells, blood, serum, plasma, urine, stool, conjunctiva, sputum, nasopharyngeal or oropharyngeal swabs, tracheal aspirates, bronchoalveolar lavage, pleural fluid, amniotic fluid, or any other specimen, or any extract thereof, or by tissue biopsy of for example lungs or major organs, obtained from a patient (human or animal), test subject, or experimental animal.

A SARS virus-related disorder may be diagnosed by amplifying a SARS nucleic acid molecule or fragment thereof from a sample. Probes or primers for use in amplification may be prepared using standard techniques. In some embodiments, probes or primers are selected from regions of a SARS virus genome as described herein that show limited sequence homology or identity (e.g., less than 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, or 100% identity) to other viruses or pathogens, or to host sequences.

Nucleic acid sequences can be amplified as needed by methods known in the art. For example, this can be accomplished by e.g., polymerase chain reaction "PCR" of DNA or of RNA by reverse transcriptase-PCR or "RT-PCR" (See generally PCR Technology: Principles and Applications for

DNA Amplification (ed. H. A. Erlich, Freeman Press, NY, N.Y., 1992); PCR Protocols: A Guide to Methods and Applications (eds. Innis, et al., Academic Press, San Diego, Calif., 1990); Mattila et al., *Nucleic Acids Res.* 19, 4967 (1991); Eckert et al., *PCR Methods and Applications* 1, 17 (1991); PCR (eds. McPherson et al., IRL Press, Oxford); and U.S. Pat. No. 4,683,202 issued Jul. 28, 1987 to Mullis) Variations of standard PCR techniques, such as for example real time RT-PCR using internal as well as amplification primers, resulting in increased sensitivity and speed, and reduction of risk of sample contamination (see for example Higuchi, R., et al., "Kinetic PCR Analysis: Real-time Monitoring of DNA Amplification Reactions," *Bio/Technology*, vol. 11, pp. 1026-1030 (1993); Heid et al., "Real Time Quantitative PCT", *Genome Research*, 1996, pp. 986-994; Gibson U E et al., "A novel method for real time quantitative RT-PCR," *Genome Res.* 1996 October; 6(10):995-1001), or the "Tacman" approach to PCR, described by for example Holland et al, *Proc. Natl. Acad. Sci.*, 88: 7276-7280 (1991), may be performed.

Other suitable amplification and analytical methods include the single base primer extension (see for example U.S. Pat. No. 6,004,744), mini-sequencing, ligase chain reaction (LCR) (see for example Wu and Wallace, *Genomics* 4, 560 (1989), Landegren et al., *Science* 241, 1077 (1988), transcription amplification (Kwoh et al., *Proc. Natl. Acad. Sci. USA* 86, 1173 (1989)), and self-sustained sequence replication (Guatelli et al., *Proc. Nat. Acad. Sci. USA*, 87, 1874 (1990)) and nucleic acid based sequence amplification (NASBA). The latter two amplification methods involve isothermal reactions based on isothermal transcription, which produce both single stranded RNA (ssRNA) and double stranded DNA (dsDNA) as the amplification products in a ratio of about 30 or 100 to 1, respectively.

A SARS virus-related disorder may also be diagnosed using an antibody directed against a SARS virus nucleic acid or amino acid sequence that specifically binds a nucleic acid molecule or polypeptide. In an alternative embodiment, the antibody may be directed against a SARS polypeptide, for example, the S polypeptide or fragment thereof that is located on the surface of the SARS virion. Methods for preparation of antibodies or for assaying antibody binding are well known in the art.

Serological diagnosis may included detection of antibodies against a SARS virus polypeptide or nucleic acid molecule, e.g., the Nucleocapsid protein, produced in response to infection using techniques such as indirect fluorescent antibody testing or enzyme-linked immunosorbent assays (ELISA). A SARS virus-related disorder may also be diagnosed by for example performing in situ probe hybridization studies on tissue specimens.

In some aspects, diagnostic tests as described herein or known to those of skill in the art may be performed for SARS virus variants that exhibit increased pathogenicity, such as strains having redundant sequences.

In some embodiments, reagents for diagnosis (e.g. probes, primers, antibodies, etc.) may be provided in kits which may optionally include instructions for using the reagent or may include other reagents for performing the appropriate assay e.g., controls, standards, buffers, etc.

Therapy or Prophylaxis for SARS Virus-Related Disorders

Compounds according to the invention may also be used to provide therapeutics or prophylactics for SARS virus-related disorders. Accordingly, such compounds may be used to treat a mammal, for example, a human or a domestic, farm, wild, or experimental animal that has or is at risk for a SARS virus-

related disorder. Such compounds may include, without limitation, compounds that interfere with SARS virus replication, expression of SARS virus proteins, or the ability of the SARS virus to infect a host cell. Accordingly, in some embodiments, compounds that act as antagonists to SARS virus polypeptides may be used as therapeutics or prophylactics for SARS virus related disorders. In some embodiments, purified SARS virus polypeptides may be used as for example competitive inhibitors to disrupt viral function. For example, a Spike protein lacking a functional domain, or having some other modification that maintains binding but reduces or eliminates pathogenicity, may be used to disrupt viral function. In some embodiments, antibodies that bind SARS virus polypeptides or nucleic acid molecules, for example, humanized antibodies, may be used as therapeutics or prophylactics.

In some embodiments, the SARS-virus compounds may be used as vaccines, or may be used to develop vaccines. For example, peptides derived from portions of SARS-virus proteins or polypeptides located on the outside of the virion or cell surface may be useful for vaccines or for generation of therapeutic or prophylactic antibodies.

A "vaccine" is a composition that includes materials that elicit a desired immune response. A vaccine may select, activate or expand memory B and T cells of the immune system to, for example, enable the elimination of infectious agents, such as a SARS virus, or a component thereof. In some embodiments, a vaccine includes a suitable carrier, such as an adjuvant, which is an agent that acts in a non-specific manner to increase the immune response to a specific antigen, or to a group of antigens, enabling the reduction of the quantity of antigen in any given vaccine dose, or the reduction of the frequency of dosage required to generate the desired immune response.

Vaccines according to the invention may include SARS virus polypeptides and nucleic acid molecules described herein, or immunogenic fragments thereof. In some embodiments, a SARS virus Spike polypeptide, Envelope polypeptide, or membrane glycoprotein or fragments thereof may be suitable for vaccine applications. In some embodiments, the vaccines may be multivalent and include one or more epitopes from a SARS virus polypeptide or fragment thereof.

In some embodiments of the invention, a vaccine may include a live or killed microorganism e.g., a SARS virus or a component thereof. If a live SARS virus is used, which may be administered in the form of an oral vaccine, it may contain non-reversible genetic alterations (for example, large deletions or insertions in the genomic sequence) that reduce or eliminate the virulence of the virus ("attenuated virus"), but not its induction of an immune response. In some embodiments, a live vaccine may include an attenuated non-SARS microorganism (e.g. bacteria or virus such as vaccinia virus) that is capable of expressing a SARS virus polypeptide or immunogenic fragment thereof as described herein. In some embodiments, a vaccine may include SARS virus polypeptides or nucleic acid molecules having modifications that facilitate ease of administration. For example, an indigestible SARS virus polypeptide or nucleic acid molecule may be used for oral administration, and a modification that is suitable for inhalation may be used for administration to the lung.

A "nucleic acid vaccine" or "DNA vaccine" as used herein, is a nucleic acid construct comprising a polynucleotide encoding a polypeptide antigen, particularly an antigenic amino acid subsequence identified by methods described herein or known in the art. The nucleic acid construct can also include transcriptional promoter elements, enhancer elements, splicing signals, termination and polyadenylation signals, and other nucleic acid sequences. Thus, a nucleic acid

vaccine is generally introduced into a subject animal using for example one or more DNA plasmids including one or more antigen-coding sequences (for example, a SARS virus Envelope polypeptide or membrane glycoprotein sequence) that are capable of transfecting cells in vivo and inducing an immune response (see for example Whalen RG et al. DNA-mediated immunization and the energetic immune response to hepatitis B surface antigen. Clin Immunol Immunopathol 1995; 75:1-12; Wolff JA et al. Direct gene transfer into mouse muscle in vivo. Science 1990; 247:1465-8; Fynan E F et al. DNA vaccines: protective immunizations by parental, mucosal, and genegun inoculations. Proc Natl Acad Sci USA 1993; 90:11478-82). In some embodiments, a library of nucleic acid fragments may be prepared by cloning SARS virus genomic DNA into a plasmid expression vector using known techniques and the library then used as a nucleic acid vaccine (see for example Barry MA, et al. Protection against mycoplasma infection using expression-library immunization. Nature 1995; 377:632-5).

The subject is administered the nucleic acid vaccine using standard methods. The vertebrate can be administered parenterally, subcutaneously, intravenously, intraperitoneally, intradermally, intramuscularly, topically, orally, rectally, nasally, buccally, vaginally, by inhalation spray, or via an implanted reservoir in dosage formulations containing conventional non-toxic, physiologically acceptable carriers or vehicles. Alternatively, the subject is administered the nucleic acid vaccine through the use of a particle acceleration or bombardment instrument (a "gene gun"). The form in which it is administered (e.g., capsule, tablet, solution, emulsion) will depend in part on the route by which it is administered. For example, for mucosal administration, nose drops, inhalants or suppositories can be used. The nucleic acid vaccine can be administered in conjunction with known adjuvants. The adjuvant is administered in a sufficient amount, which is that amount that is sufficient to generate an enhanced immune response to the nucleic acid vaccine. The adjuvant can be administered prior to (e.g., 1 or more days before) inoculation with the nucleic acid vaccine; concurrently with (e.g., within 24 hours of) inoculation with the nucleic acid vaccine; contemporaneously (simultaneously) with the nucleic acid vaccine (e.g., the adjuvant is mixed with the nucleic acid vaccine, and the mixture is administered to the vertebrate); or after (e.g., 1 or more days after) inoculation with the nucleic acid vaccine. The adjuvant can also be administered at more than one time (e.g., prior to inoculation with the nucleic acid vaccine and also after inoculation with the nucleic acid vaccine). As used herein, the term "in conjunction with" encompasses any time period, including those specifically described herein and combinations of the time periods specifically described herein, during which the adjuvant can be administered so as to generate an enhanced immune response to the nucleic acid vaccine (e.g., an increased antibody titer to the antigen encoded by the nucleic acid vaccine, or an increased antibody titer to the pathogenic agent). The adjuvant and the nucleic acid vaccine can be administered at approximately the same location on the vertebrate; for example, both the adjuvant and the nucleic acid vaccine are administered at a marked site on a limb of the subject.

In some embodiments, expression of a SARS virus gene or coding or non-coding region of interest may be inhibited or prevented using RNA interference (RNAi) technology, a type of post-transcriptional gene silencing. RNAi may be used to create a functional "knockout", i.e. a system in which the expression of a gene or coding or non-coding region of interest is reduced, resulting in an overall reduction of the encoded

product. As such, RNAi may be performed to target a nucleic acid of interest or fragment or variant thereof, to in turn reduce its expression and the level of activity of the product which it encodes. Such a system may be used for therapy or prophylaxis, as well as for functional studies. RNAi is described in for example published US patent applications 20020173478 (Gewirtz; published Nov. 21, 2002) and 20020132788 (Lewis et al.; published Nov. 7, 2002). Reagents and kits for performing RNAi are available commercially from for example Ambion Inc. (Austin, Tex., USA) and New England Biolabs Inc. (Beverly, Mass., USA).

The initial agent for RNAi in some systems is thought to be dsRNA molecule corresponding to a target nucleic acid. The dsRNA is then thought to be cleaved into short interfering RNAs (siRNAs) which are 21-23 nucleotides in length (19-21 bp duplexes, each with 2 nucleotide 3' overhangs). The enzyme thought to effect this first cleavage step has been referred to as "Dicer" and is categorized as a member of the Rnase III family of dsRNA-specific ribonucleases. Alternatively, RNAi may be effected via directly introducing into the cell, or generating within the cell by introducing into the cell a suitable precursor (e.g. vector, etc.) of such an siRNA or siRNA-like molecule. An siRNA may then associate with other intracellular components to form an RNA-induced silencing complex (RISC). The RISC thus formed may subsequently target a transcript of interest via base-pairing interactions between its siRNA component and the target transcript by virtue of homology, resulting in the cleavage of the target transcript approximately 12 nucleotides from the 3' end of the siRNA. Thus the target mRNA is cleaved and the level of protein product it encodes is reduced.

RNAi may be effected by the introduction of suitable in vitro synthesized siRNA or siRNA-like molecules into cells. RNAi may for example be performed using chemically-synthesized RNA, for which suitable RNA molecules may be chemically synthesized using known methods. Alternatively, suitable expression vectors may be used to transcribe such RNA either in vitro or in vivo. In vitro transcription of sense and antisense strands (encoded by sequences present on the same vector or on separate vectors) may be effected using for example T7 RNA polymerase, in which case the vector may comprise a suitable coding sequence operably-linked to a T7 promoter. The in vitro-transcribed RNA may in embodiments be processed (e.g. using *E. coli* RNase III) in vitro to a size conducive to RNAi. The sense and antisense transcripts combined to form an RNA duplex which is introduced into a target cell of interest. Other vectors may be used, which express small hairpin RNAs (shRNAs) which can be processed into siRNA-like molecules. Various vector-based methods are known in the art. Various methods for introducing such vectors into cells, either in vitro or in vivo (e.g. gene therapy) are known in the art.

Accordingly, in an embodiment, expression of a polypeptide including an amino acid sequence substantially identical to a SARS virus sequence may be inhibited by introducing into or generating within a cell an siRNA or siRNA-like molecule corresponding to a nucleic acid molecule encoding the polypeptide or fragment thereof, or to an nucleic acid homologous thereto. In various embodiments such a method may entail the direct administration of the siRNA or siRNA-like molecule into a cell, or use of the vector-based methods described above. In an embodiment, the siRNA or siRNA-like molecule is less than about 30 nucleotides in length. In a further embodiment, the siRNA or siRNA-like molecules are about 21-23 nucleotides in length. In an embodiment, siRNA or siRNA-like molecules comprise and 19-21 bp duplex portion, each strand having a 2 nucleotide 3' overhang. In

embodiments, the siRNA or siRNA-like molecule is substantially identical to a nucleic acid encoding the polypeptide or a fragment or variant (or a fragment of a variant) thereof. Such a variant is capable of encoding a protein having the activity of a SARS virus polypeptide. In embodiments, the sense strand of the siRNA or siRNA-like molecule is substantially identical to a SARS virus nucleic acid molecule or a fragment thereof (RNA having U in place of T residues of the DNA sequence).

SARS Virus Protein Expression

In general, SARS virus polypeptides according to the invention, may be produced by transformation of a suitable host cell with all or part of a SARS virus polypeptide-encoding genomic or cDNA molecule or fragment thereof (e.g., the genomic DNA or cDNAs described herein) in a suitable expression vehicle. Those skilled in the field of molecular biology will understand that any of a wide variety of expression systems may be used to provide the recombinant protein. The precise host cell used is not critical to the invention. The SARS virus polypeptide may be produced in a prokaryotic host (e.g., *E. coli* or a virus, for example, a coronavirus such as human OC43 or 229E, a bovine coronavirus, or a virus used for gene therapy, such as an adenovirus) or in a eukaryotic host (e.g., *Saccharomyces cerevisiae*, insect cells, e.g., Sf21 cells, or mammalian cells, e.g., COS 1, NIH 3T3, VeroE6, or HeLa cells). Such cells are available from a wide range of sources (e.g., the American Type Culture Collection, Rockland, Md.; also, see, e.g., Ausubel et al., Current Protocols in Molecular Biology, John Wiley & Sons, New York, 1994). The method of transformation or transfection and the choice of expression vehicle will depend on the host system selected. Transformation and transfection methods are described, e.g., in Ausubel et al. (supra); expression vehicles may be chosen from those provided, e.g., in Cloning Vectors: A Laboratory Manual, P. H. Pouwels et al, 1985, Supp. 1987), or from commercially available sources. Suitable animal models, e.g. a ferret animal model, or any other animal model suitable for analysis of SARS virus infection or expression of SARS virus nucleic acid molecules may be used.

In an alternative embodiment, the baculovirus expression system (using, for example, the vector pBacPAK9) available from Clontech (Pal Alto, Calif.) may be used. If desired, this system may be used in conjunction with other protein expression techniques, for example, the myc tag approach described by Evan et al. (Mol. Cell Biol. 5:3610-3616, 1985). In an alternative embodiment, a SARS virus polypeptide may be produced by a stably-transfected mammalian cell line. A number of vectors suitable for stable transfection of mammalian cells are available to the public, e.g., see Pouwels et al (supra); methods for constructing such cell lines are also publicly available, e.g., in Ausubel et al. (supra). In one example, cDNA encoding the SARS virus polypeptide is cloned into an expression vector which includes the dihydrofolate reductase (DHFR) gene. Integration of the plasmid and, therefore, the SARS virus polypeptide-encoding gene into the host cell chromosome is selected for by inclusion of 0.01-300 μ M methotrexate in the cell culture medium (as described in Ausubel et al., supra). This dominant selection can be accomplished in most cell types. Recombinant protein expression can be increased by DHFR-mediated amplification of the transfected gene. Methods for selecting cell lines bearing gene amplifications are described in Ausubel et al. (supra); such methods generally involve extended culture in medium containing gradually increasing levels of methotrexate. DHFR-containing expression vectors commonly used for this purpose include pCVSEII-DHFR and pAdD26SV(A)

(described in Ausubel et al., supra). Any of the host cells described above or, preferably, a DHFR-deficient CHO cell line (e.g., CHO DHFR.sup.—cells, ATCC Accession No. CRL 9096) are among the host cells preferred for DHFR selection of a stably-transfected cell line or DHFR-mediated gene amplification.

Once the recombinant SARS virus polypeptide is expressed, it is isolated, e.g., using affinity chromatography. In one example, an anti-SARS virus polypeptide antibody (e.g., produced as described herein) may be attached to a column and used to isolate the SARS virus polypeptide. Lysis and fractionation of SARS virus polypeptide-harboring cells prior to affinity chromatography may be performed by standard methods (see, e.g., Ausubel et al., supra). In another example, SARS virus polypeptides may be purified or substantially purified from a mixture of compounds such as an extract or supernatant obtained from cells (Ausubel et al., supra). Standard purification techniques can be used to progressively eliminate undesirable compounds from the mixture until a single compound or minimal number of effective compounds has been isolated.

Once isolated, the recombinant protein can, if desired, be further purified, e.g., by high performance liquid chromatography (see, e.g., Fisher, Laboratory Techniques In Biochemistry And Molecular Biology, eds., Work and Burdon, Elsevier, 1980).

Polypeptides of the invention, particularly short SARS virus peptide fragments, can also be produced by chemical synthesis (e.g., by the methods described in Solid Phase Peptide Synthesis, 2nd ed., 1984 The Pierce Chemical Co., Rockford, Ill.).

These general techniques of polypeptide expression and purification can also be used to produce and isolate useful SARS virus protein fragments or analogs (described herein).

In certain alternative embodiments, the SARS polypeptide might have attached any one of a variety of tags. Tags can be amino acid tags or chemical tags and can be added for the purpose of purification (for example a 6-histidine tag for purification over a nickel column). In other preferred embodiments, various labels can be used as means for detecting binding of a SARS polypeptide to another polypeptide, for example to a cell surface receptor. Alternatively, SARS DNA or RNA may be labeled for detection, for example in a hybridization assay. SARS virus nucleic acids or proteins, or derivatives thereof, may be directly or indirectly labeled, for example, with a radioscope, a fluorescent compound, a bioluminescent compound, a chemiluminescent compound, a metal chelator or an enzyme. Those of ordinary skill in the art will know of other suitable labels or will be able to ascertain such, using routine experimentation. In yet another embodiment of the invention, the polypeptides disclosed herein, or derivatives thereof, are linked to toxins.

Isolation and Identification of Additional SARS Virus Molecules

Based on the SARS virus sequences described herein, the isolation and identification of additional SARS virus-related sequences such as SARS virus genes and of additional SARS virus strains or isolates is made possible using standard techniques. In addition, the SARS virus sequences provided herein also provide the basis for identification of homologous sequences from other species and genera from both prokaryotes and eukaryotes such as viruses, bacteria, fungi, parasites, yeast, and/or mammals. In some embodiments, the nucleic acid sequences described herein may be used to design probes or primers, including degenerate oligonucleotide probes or primers, based upon the sequence of either DNA strand. The

probes or primers may then be used to screen genomic or cDNA libraries for sequences from for example naturally occurring variants or isolates of SARS viruses, using standard amplification or hybridization techniques.

In some embodiments, binding partners may be identified by tagging the polypeptides of the invention (e.g., those substantially identical to SARS virus polypeptides described herein) with an epitope sequence (e.g., FLAG or 2HA), and delivering it into host cells, either by transfection with a suitable vector containing a nucleic acid sequence encoding a polypeptide of the invention, followed by immunoprecipitation and identification of the binding partner. Cells may be infected with strains expressing the FLAG or 2HA fusions, followed by lysis and immunoprecipitation with anti-FLAG or anti-2HA antibodies. Binding partners may be identified by mass spectroscopy. If the polypeptide of the invention is not produced in sufficient quantities, such a method may not deliver enough tagged protein to identify its partner. As part of a complementary approach, each polypeptide of the invention may be cloned into a mammalian transfection vector fused to, for example, 2HA, GFP and/or FLAG. Following transfection, HeLa cells may be lysed and the tagged polypeptide immunoprecipitated. The binding partner may be identified by SDS PAGE followed by mass spectroscopy.

In some embodiments, polypeptides or antibodies of the invention may be tagged, produced, and used for example on affinity columns and/or in immunological assays to identify and/or confirm identified target compounds. FLAG, HA, and/or His tagged proteins can be used for such affinity columns to pull out host cell factors from cell extracts, and any hits may be validated by standard binding assays, saturation curves, and other methods as described herein or known to those of skill in the art.

In some embodiments, a two hybrid system may be used to study protein-protein interactions. The nucleic acid sequences described herein, or sequences substantially identical thereto, can be cloned into the pBT bait plasmid of the two hybrid system, and a commercially available murine spleen library of 5×10^6 independent clones, may be used as the target library for the baits. Potential hits may be further characterized by recovering the plasmids and retransforming to reduce false positives resulting from clonal bait variants and library target clones which activate the reporter genes independent of the cloned bait. Reproducible hits may be studied further as described herein.

Virulence may be assayed as described herein or as known to those of skill in the art. Once coding sequences have been identified, they may be isolated using standard cloning techniques, and inserted into any suitable vector or replicon for, for example, production of polypeptides. Such vectors and replicons include, without limitation, bacteriophage X (*E. coli*), pBR322 (*E. coli*), pACYC177 (*E. coli*), pKT230 (gram-negative bacteria), pGV1 106 (gram-negative bacteria), pLAFR1 (gram-negative bacteria), pME290 (non-*E. coli* gram-negative bacteria), pHV14 (*E. coli* and *Bacillus subtilis*), pBD9 (*Bacillus*), pIJ61 (*Streptomyces*), pUC6 (*Streptomyces*), YIp5 (*Saccharomyces*), YCp19 (*Saccharomyces*) or bovine papilloma virus (mammalian cells). In general, the polypeptides of the invention may be produced in any suitable host cell transformed or transfected with a suitable vector. The method of transformation or transfection and the choice of expression vehicle will depend on the host system selected. A wide variety of expression systems may be used, and the precise host cell used is not critical to the invention. For example, a polypeptide according to the invention may be produced in a prokaryotic host (e.g., *E. coli*) or in a eukaryotic host (e.g., *Saccharomyces cerevisiae*, insect cells, e.g., Sf21

cells, or mammalian cells, e.g., NIH 3T3, HeLa, or COS cells). Such cells are available from a wide range of sources (e.g., the American Type Culture Collection, Manassas, Va.). Bacterial expression systems for polypeptide production include the *E. coli* pET expression system (Novagen, Inc., Madison, Wis.), and the pGEX expression system (Pharmacia).

Compounds

In one aspect, compounds according to the invention include SARS virus nucleic acid molecules and polypeptides, such as the sequences disclosed in the Figures and Tables herein, and throughout the specification, and fragments thereof. In alternative embodiments, compounds according to the invention may be nucleic acid molecules that are at least 10 nucleotides in length, and that are derived from the sequences described herein. In alternative embodiments, compounds according to the invention may be peptides that are at least 5 amino acids in length, and that are derived from the sequences described herein.

In alternative embodiments, a compound according to the invention can be a non-peptide molecule as well as a peptide or peptide analogue. A peptide or peptide analogue will generally be as small as feasible while retaining full biological activity. A non-peptide molecule can be any molecule that exhibits biological activity as described herein or known in the art. Biological activity can, for example, be measured in terms of ability to elicit a cytotoxic response, to mediate DNA replication, or any other function of a SARS virus molecule.

Compounds can be prepared by, for example, replacing, deleting, or inserting an amino acid residue of SARS peptide or peptide analogue, as described herein, with other conservative amino acid residues, i.e., residues having similar physical, biological, or chemical properties, and screening for biological function.

It is well known in the art that some modifications and changes can be made in the structure of a polypeptide without substantially altering the biological function of that peptide, to obtain a biologically equivalent polypeptide. Such modifications may be made for the purpose of modifying function, or for facilitating administration or enhancing stability or inhibiting breakdown for, for example, therapeutic uses. For example, an indigestible SARS virus compound according to the invention may be used for oral administration; a modification that is suitable for inhalation may be used for administration to the lung; or addition of a leader sequence may increase protein expression levels.

In one aspect of the invention, SARS virus-derived peptides or epitopes may include peptides that differ from a portion of a native leader, protein or SARS virus sequence by conservative amino acid substitutions. The peptides and epitopes of the present invention also extend to biologically equivalent peptides that differ from a portion of the sequence of novel peptides of the present invention by conservative amino acid substitutions. As used herein, the term "conserved amino acid substitutions" refers to the substitution of one amino acid for another at a given location in the peptide, where the substitution can be made without substantial loss of the relevant function. In making such changes, substitutions of like amino acid residues can be made on the basis of relative similarity of side-chain substituents, for example, their size, charge, hydrophobicity, hydrophilicity, and the like, and such substitutions may be assayed for their effect on the function of the peptide by routine testing.

In some embodiments, conserved amino acid substitutions may be made where an amino acid residue is substituted for another having a similar hydrophilicity value (e.g., within a

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value of plus or minus 2.0), where the following may-be an amino acid having a hydrophobic index of about -1.6 such as Tyr (-1.3) or Pro (-1.6) are assigned to amino acid residues (as detailed in U.S. Pat. No. 4,554,101, incorporated herein by reference): Arg (+3.0); Lys (+3.0); Asp (+3.0); Glu (+3.0); Ser (+0.3); Asn (+0.2); Gln (+0.2); Gly (0); Pro (-0.5); Thr (-0.4); Ala (-0.5); His (-0.5); Cys (-1.0); Met (-1.3); Val (-1.5); Leu (-1.8); Ile (-1.8); Tyr (-2.3); Phe (-2.5); and Trp (-3.4).

In alternative embodiments, conserved amino acid substitutions may be made where an amino acid residue is substituted for another having a similar hydrophobic index (e.g., within a value of plus or minus 2.0). In such embodiments, each amino acid residue may be assigned a hydrophobic index on the basis of its hydrophobicity and charge characteristics, as follows: Ile (+4.5); Val (+4.2); Leu (+3.8); Phe (+2.8); Cys (+2.5); Met (+1.9); Ala (+1.8); Gly (-0.4); Thr (-0.7); Ser (-0.8); Trp (-0.9); Tyr (-1.3); Pro (-1.6); His (-3.2); Glu (-3.5); Gln (-3.5); Asp (-3.5); Asn (-3.5); Lys (-3.9); and Arg (-4.5).

In alternative embodiments, conserved amino acid substitutions may be made where an amino acid residue is substituted for another in the same class, where the amino acids are divided into non-polar, acidic, basic and neutral classes, as follows: non-polar: Ala, Val, Leu, Ile, Phe, Trp, Pro, Met; acidic: Asp, Glu; basic: Lys, Arg, His; neutral: Gly, Ser, Thr, Cys, Asn, Gln, Tyr.

Conservative amino acid changes can include the substitution of an L-amino acid by the corresponding D-amino acid, by a conservative D-amino acid, or by a naturally-occurring, non-genetically encoded form of amino acid, as well as a conservative substitution of an L-amino acid. Naturally-occurring non-genetically encoded amino acids include beta-alanine, 3-amino-propionic acid, 2,3-diamino propionic acid, alpha-aminoisobutyric acid, 4-amino-butyric acid, N-methylglycine (sarcosine), hydroxyproline, ornithine, citrulline, t-butylalanine, t-butylglycine, N-methylisoleucine, phenylglycine, cyclohexylalanine, norleucine, norvaline, 2-naphthylalanine, pyridylalanine, 3-benzothienyl alanine, 4-chlorophenylalanine, 2-fluorophenylalanine, 3-fluorophenylalanine, 4-fluorophenylalanine, penicillamine, 1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid, beta-2-thienylalanine, methionine sulfoxide, homoarginine, N-acetyl lysine, 2-amino butyric acid, 2-amino butyric acid, 2,4-diamino butyric acid, p-aminophenylalanine, N-methylvaline, homocysteine, homoserine, cysteine acid, epsilon-amino hexanoic acid, delta-amino valeric acid, or 2,3-diaminobutyric acid.

In alternative embodiments, conservative amino acid changes include changes based on considerations of hydrophilicity or hydrophobicity, size or volume, or charge. Amino acids can be generally characterized as hydrophobic or hydrophilic, depending primarily on the properties of the amino acid side chain. A hydrophobic amino acid exhibits a hydrophobicity of greater than zero, and a hydrophilic amino acid exhibits a hydrophilicity of less than zero, based on the normalized consensus hydrophobicity scale of Eisenberg et al. (*J. Mol. Bio.* 179:125-142, 184). Genetically encoded hydrophobic amino acids include Gly, Ala, Phe, Val, Leu, Ile, Pro, Met and Trp, and genetically encoded hydrophilic amino acids include Thr, His, Glu, Gln, Asp, Arg, Ser, and Lys. Non-genetically encoded hydrophobic amino acids include t-butylalanine, while non-genetically encoded hydrophilic amino acids include citrulline and homocysteine.

Hydrophobic or hydrophilic amino acids can be further subdivided based on the characteristics of their side chains. For example, an aromatic amino acid is a hydrophobic amino

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acid with a side chain containing at least one aromatic or heteroaromatic ring, which may contain one or more substituents such as -OH, -SH, -CN, -F, -Cl, -Br, -I, -NO₂, -NO, -NH₂, -NHR, -NRR, -C(O)R, -C(O)OH, -C(O)OR, -C(O)NH₂, -C(O)NHR, -C(O)NRR, etc., where R is independently (C₁-C₆) alkyl, substituted (C₁-C₆) alkyl, (C₁-C₆) alkenyl, substituted (C₁-C₆) alkenyl, (C₁-C₆) alkynyl, substituted (C₁-C₆) alkynyl, (C₅-C₂₀) aryl, substituted (C₅-C₂₀) aryl, (C₆-C₂₆) alkaryl, substituted (C₆-C₂₆) alkaryl, 5-20 membered heteroaryl, substituted 5-20 membered heteroaryl, 6-26 membered alkheteroaryl or substituted 6-26 membered alkheteroaryl. Genetically encoded aromatic amino acids include Phe, Tyr, and Trp, while non-genetically encoded aromatic amino acids include phenylglycine, 2-naphthylalanine, beta-2-thienylalanine, 1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid, 4-chlorophenylalanine, 2-fluorophenylalanine, 3-fluorophenylalanine, and 4-fluorophenylalanine.

An apolar amino acid is a hydrophobic amino acid with a side chain that is uncharged at physiological pH and which has bonds in which a pair of electrons shared in common by two atoms is generally held equally by each of the two atoms (i.e., the side chain is not polar). Genetically encoded apolar amino acids include Gly, Leu, Val, Ile, Ala, and Met, while non-genetically encoded apolar amino acids include cyclohexylalanine. Apolar amino acids can be further subdivided to include aliphatic amino acids, which is a hydrophobic amino acid having an aliphatic hydrocarbon side chain. Genetically encoded aliphatic amino acids include Ala, Leu, Val, and Ile, while non-genetically encoded aliphatic amino acids include norleucine.

A polar amino acid is a hydrophilic amino acid with a side chain that is uncharged at physiological pH, but which has one bond in which the pair of electrons shared in common by two atoms is held more closely by one of the atoms. Genetically encoded polar amino acids include Ser, Thr, Asn, and Gln, while non-genetically encoded polar amino acids include citrulline, N-acetyl lysine, and methionine sulfoxide.

An acidic amino acid is a hydrophilic amino acid with a side chain pKa value of less than 7. Acidic amino acids typically have negatively charged side chains at physiological pH due to loss of a hydrogen ion. Genetically encoded acidic amino acids include Asp and Glu. A basic amino acid is a hydrophilic amino acid with a side chain pKa value of greater than 7. Basic amino acids typically have positively charged side chains at physiological pH due to association with hydronium ion. Genetically encoded basic amino acids include Arg, Lys, and His, while non-genetically encoded basic amino acids include the non-cyclic amino acids ornithine, 2,3-diaminopropionic acid, 2,4-diaminobutyric acid, and homoarginine.

It will be appreciated by one skilled in the art that the above classifications are not absolute and that an amino acid may be classified in more than one category. In addition, amino acids can be classified based on known behaviour and or characteristic chemical, physical, or biological properties based on specified assays or as compared with previously identified amino acids. Amino acids can also include bifunctional moieties having amino acid-like side chains.

Conservative changes can also include the substitution of a chemically derivatised moiety for a non-derivatised residue, by for example, reaction of a functional side group of an amino acid. Thus, these substitutions can include compounds whose free amino groups have been derivatised to amine hydrochlorides, p-toluene sulfonyl groups, carbobenzoxy groups, t-butyloxycarbonyl groups, chloroacetyl groups or formyl groups. Similarly, free carboxyl groups can be deriva-

tized to form salts, methyl and ethyl esters or other types of esters or hydrazides, and side chains can be derivatized to form O-acyl or O-alkyl derivatives for free hydroxyl groups or N-im-benzylhistidine for the imidazole nitrogen of histidine. Peptide analogues also include amino acids that have been chemically altered, for example, by methylation, by amidation of the C-terminal amino acid by an alkylamine such as ethylamine, ethanolamine, or ethylene diamine, or acylation or methylation of an amino acid side chain (such as acylation of the epsilon amino group of lysine). Peptide analogues can also include replacement of the amide linkage in the peptide with a substituted amide (for example, groups of the formula $-\text{C}(\text{O})-\text{NR}$, where R is (C_1-C_6) alkyl, (C_1-C_6) alkenyl, (C_1-C_6) alkynyl, substituted (C_1-C_6) alkyl, substituted (C_1-C_6) alkenyl, or substituted (C_1-C_6) alkynyl) or isostere of an amide linkage (for example, $-\text{CH}_2\text{NH}-$, $-\text{CH}_2\text{S}$, $-\text{CH}_2\text{CH}_2-$, $-\text{CH}=\text{CH}-$ (cis and trans), $-\text{C}(\text{O})\text{CH}_2-$, $-\text{CH}(\text{OH})\text{CH}_2-$, or $-\text{CH}_2\text{SO}-$).

The compound can be covalently linked, for example, by polymerisation or conjugation, to form homopolymers or heteropolymers. Spacers and linkers, typically composed of small neutral molecules, such as amino acids that are uncharged under physiological conditions, can be used. Linkages can be achieved in a number of ways. For example, cysteine residues can be added at the peptide termini, and multiple peptides can be covalently bonded by controlled oxidation. Alternatively, heterobifunctional agents, such as disulfide/amide forming agents or thioether/amide forming agents can be used. The compound can also be constrained, for example, by having cyclic portions.

In some embodiments, three dimensional molecular modeling techniques may be used to identify or generate compounds that may be useful as therapeutics or diagnostics. Standard molecular modeling tools may be used, for example, those described in L-H Hung and R. Samudrala, PROTFINFO: secondary and tertiary protein structure prediction, *Nucleic Acids Research*, 2003, Vol. 31, No. 13 3296-3299; A. Yamaguchi, et al., Enlarged FAMSBASE: protein 3D structure models of genome sequences for 41 species, *Nucleic Acids Research*, 2003, Vol. 31, No. 1 463-468; J. Chen, et al., MMDB: Entrez's 3D-structure database, *Nucleic Acids Research*, 2003, Vol. 31, No. 1 474-477; R. A. Chiang, et al., The Structure Superposition Database, *Nucleic Acids Research*, 2003, Vol. 31, No. 1 505-510.

Peptides or peptide analogues can be synthesized by standard chemical techniques, for example, by automated synthesis using solution or solid phase synthesis methodology. Automated peptide synthesizers are commercially available and use techniques well known in the art. Peptides and peptide analogues can also be prepared using recombinant DNA technology using standard methods such as those described in, for example, Sambrook, et al. (*Molecular Cloning: A Laboratory Manual*, 2.sup.nd, ed., Cold Spring Harbor Laboratory, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y., 1989) or Ausubel et al. (*Current Protocols in Molecular Biology*, John Wiley & Sons, 1994).

Compounds, such as peptides (or analogues thereof) can be identified by routine experimentation by, for example, modifying residues within SARS peptides; introducing single or multiple amino acid substitutions, deletions, or insertions, and identifying those compounds that retain biological activity, e.g., those compounds that have cytotoxic ability.

In general, candidate compounds for prevention or treatment of SARS virus-mediated disorders are identified from large libraries of both natural product or synthetic (or semi-synthetic) extracts or chemical libraries according to methods known in the art. Candidate or test compounds may include,

without limitation, peptides, polypeptides, synthesised organic molecules, naturally occurring organic molecules, and nucleic acid molecules. In some embodiments, such compounds screen for the ability to inhibit SARS virus replication or pathogenicity, while maintaining the infected cell's ability to grow or survive.

Those skilled in the field of drug discovery and development will understand that the precise source of test extracts or compounds is not critical to the method(s) of the invention. Accordingly, virtually any number of chemical extracts or compounds can be screened using the exemplary methods described herein or using standard methods. Examples of such extracts or compounds include, but are not limited to, plant-, fungal-, prokaryotic- or animal-based extracts, fermentation broths, and synthetic compounds, as well as modification of existing compounds. Numerous methods are also available for generating random or directed synthesis (e.g., semi-synthesis or total synthesis) of any number of chemical compounds, including, but not limited to, saccharide-, lipid-, peptide-, and nucleic acid-based compounds. Synthetic compound libraries are commercially available. Alternatively, libraries of natural compounds in the form of bacterial, fungal, plant, and animal extracts are commercially available from a number of sources, including Biotics (Sussex, UK), Xenova (Slough, UK), Harbor Branch Oceanographic Institute (Ft. Pierce, Fla.), and PharmaMar, U.S.A. (Cambridge, Mass.). In addition, natural and synthetically produced libraries of, for example, SARS virus polypeptides containing leader sequences, are produced, if desired, according to methods known in the art, e.g., by standard extraction and fractionation methods. Furthermore, if desired, any library or compound is readily modified using standard chemical, physical, or biochemical methods.

When a crude extract is found to modulate cytotoxicity or viral infection, further fractionation of the positive lead extract is necessary to isolate chemical constituents responsible for the observed effect. Thus, the goal of the extraction, fractionation, and purification process is the careful characterization and identification of a chemical entity within the crude extract having, for example, anti-cytotoxicity or antiviral properties. The same assays described herein for the detection of activities in mixtures of compounds can be used to purify the active component and to test derivatives thereof. Methods of fractionation and purification of such heterogeneous extracts are known in the art. If desired, compounds shown to be useful agents for treatment are chemically modified according to methods known in the art. Compounds identified as being of therapeutic, prophylactic, diagnostic, or other value in for example cell culture systems, such as a Vero E6 culture system, may be subsequently analyzed using a ferret animal model, or any other animal model suitable for analysis of SARS.

Antibodies

The compounds of the invention can be used to prepare antibodies to SARS virus peptides, protein, polyproteins, or analogs thereof, or to SARS virus nucleic acid molecules or analogs thereof using standard techniques of preparation as, for example, described in Harlow and Lane (*Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y., 1988), or known to those skilled in the art. Antibodies may include polyclonal antibodies, monoclonal antibodies, hybrid antibodies (e.g., divalent antibodies having different pairs of heavy and light chains), chimeric antibodies (e.g., antibodies having constant and variable domains from different species and/or class), modified antibodies (e.g., antibodies in which the naturally occurring

sequence has been altered by for example recombinant techniques), Fab antibodies, anti-idiotypic antibodies, etc. Antibodies can be tailored to minimize adverse host immune response by, for example, using chimeric antibodies containing an antigen binding domain from one species and the Fc portion from another species, or by using antibodies made from hybridomas of the appropriate species. For example, "humanized" antibodies may be used for administration to humans.

To generate SARS virus polypeptide-specific antibodies, a SARS virus polypeptide coding sequence may be expressed, for example, as a C-terminal fusion with glutathione S-transferase (GST) (Smith et al., *Gene* 67:31-40, 1988). The fusion polypeptide may then be purified on glutathione-Sepharose beads, eluted with glutathione cleaved with thrombin (at the engineered cleavage site), and purified to the degree necessary for immunization of rabbits. Primary immunizations are carried out with Freud's complete adjuvant and subsequent immunizations with Freud's incomplete adjuvant. Antibody titres are monitored by Western blot and immunoprecipitation analyzes using the thrombin-cleaved SARS virus polypeptide fragment of the GST-SARS virus fusion polypeptide. Immune sera are affinity purified using CNBr-Sepharose-coupled SARS virus polypeptide. Antiserum specificity is determined using a panel of unrelated GST polypeptides.

As an alternate or adjunct immunogen to GST fusion polypeptides, peptides corresponding to relatively unique hydrophilic SARS virus polypeptides may be generated and coupled to keyhole limpet hemocyanin (KLH) through an introduced C-terminal lysine. Antiserum to each of these peptides is similarly affinity purified on peptides conjugated to BSA, and specificity tested in ELISA and Western blots using peptide conjugates, and by Western blot and immunoprecipitation using SARS virus polypeptide expressed as a GST fusion polypeptide.

Alternatively, monoclonal antibodies may be prepared using the SARS virus polypeptides described above and standard hybridoma technology (see, e.g., Kohler et al., *Nature*, 256:495, 1975; Kohler et al., *Eur. J Immunol.* 6:511, 1976; Kohler et al., *Eur. J. Immunol.* 6:292, 1976; Hammerling et al., *In Monoclonal Antibodies and T Cell Hybridomas*, Elsevier, NY, 1981; Ausubel et al., *supra*). Once produced, monoclonal antibodies are also tested for specific SARS virus polypeptide recognition by Western blot or immunoprecipitation analysis (by the methods described in Ausubel et al., *supra*). Antibodies which specifically recognize SARS virus polypeptides are considered to be useful in the invention; such antibodies may be used, e.g., in an immunoassay to monitor the level of SARS virus polypeptides produced by a mammal (for example, to determine the amount or location of a SARS virus polypeptide).

In an alternative embodiment, antibodies of the invention are not only produced using the whole SARS virus polypeptide, but using fragments of the SARS virus polypeptide which are unique or which lie outside highly conserved regions and appear likely to be antigenic, by criteria such as high frequency of charged residues may also be used. In one specific example, such fragments are generated by standard techniques of PCR and cloned into the pGEX expression vector (Ausubel et al., *supra*). Fusion polypeptides are expressed in *E. coli* and purified using a glutathione agarose affinity matrix as described in Ausubel et al. (*supra*). To attempt to minimize the potential problems of low affinity or specificity of antisera, two or three such fusions are generated for each polypeptide, and each fusion is injected into at least two rabbits. Antisera are raised by injections in a series,

preferably including at least three booster injections. SARS virus antibodies may also be prepared against SARS virus nucleic acid molecules.

Antibodies may be used as diagnostics, therapeutics, or prophylactics for SARS virus-related disorders. Antibodies may also be used to isolate SARS virus and compounds by for example affinity chromatography, or to identify SARS virus compounds isolated or generated by other techniques.

10 Arrays and Libraries

In some aspects, biological assays, such as diagnostic or other assays, using high density nucleic acid, polypeptide, or antibody arrays, for example high density miniaturized arrays or "microarrays," of SARS virus nucleic acid molecules or polypeptides, or antibodies capable of specifically binding such nucleic acid molecules or polypeptides, may be performed. Macroarrays, performed for example by manual spotting techniques, may also be used. Arrays generally require a solid support (for example, nylon, glass, ceramic, plastic, silicon, nitrocellulose or PVDF membranes, microwells, microbeads, e.g., magnetic microbeads, etc.) to which the nucleic acid molecules or polypeptides or antibodies are attached in a specified two-dimensional arrangement, such that the pattern of hybridization is easily determinable. Suspension arrays (particles in suspension) that are coded to facilitate identification may also be used. SARS virus nucleic acid molecules or polypeptide probes or targets may be compounds as described herein.

In some embodiments, high density nucleic acid arrays may for example be used to monitor the presence or level of expression of a large number of SARS virus nucleic acid molecules or genes or for detecting or identifying SARS virus nucleic acid sequence variations, mutations or polymorphisms. For the purpose of such arrays, "nucleic acids" may include any polymer or oligomer of nucleosides or nucleotides (polynucleotides or oligonucleotides), which include pyrimidine and purine bases, preferably cytosine, thymine, and uracil, and adenine and guanine, respectively, or may include peptide nucleic acids (PNA). In an alternative aspect, the invention provides nucleic acid microarrays including a number of distinct nucleic acid sequence arrays of the invention, thus providing specific "sets" of sequences. The number of distinct sequences may for example be any integer between 2 and 1×10^5 , such as at least 10^2 , 10^3 , 10^4 , or 10^5 .

The invention also provides gene knockout and expression libraries. Thus, nucleic acid molecules encoding SARS virus polypeptides or proteins (e.g., PCR products of ORF's or total mRNA) may for example be attached to a solid support, hybridized with single stranded detectably-labeled cDNAs (corresponding to an "antisense" orientation), and quantified using an appropriate method such that a signal is detected at each location at which hybridization has taken place. The intensity of the signal would then reflect the level of gene expression. Comparison of results from viruses, for example, of different strains or from different samples or subjects, would elucidate differing levels of expression of specified genes. Using similar techniques, homologous nucleic acids may be identified from different viruses if SARS virus nucleic acids are used in the microarray, and probed with nucleic acid molecules from different viruses or subjects. In some embodiments, this approach may involve constructing his-tagged ORP expression libraries of viral genomes in a bacterial host, similar to an expression library in yeast (Martzen M. R. et al., 1999. *Science*, 286:1153). ORF-encoded protein activities may for example be detected in purified his-tagged protein pools in cases where activities cannot be detected in extracts or cells. In one aspect of the invention, arrayed librar-

ies may be constructed of viral strains each of which bears a plasmid expressing a different SARS virus ORF under control of an inducible promoter. ORFs are amplified using PCR and cloned into a vector that enables their expression as N-terminal his-tagged polypeptides. These amplicons are also used to construct hybridization microarrays and enable targeted gene disruption, reducing expenses. A suitable expression host is selected, and genes encoding particular biochemical activities are identified by screening arrayed pools of his-tagged proteins as described previously (Martzen M. R., McCraith S. M., Spinelli S. L., Torres F. M., Fields S., Grayhack E. J., and Phizicky E. M., 1999. *Science*, 286: 1153).

In some embodiments, protein arrays (including antibody or antigen arrays) may be used for the analysis and identification of SARS virus polypeptides or host responses to such polypeptides. Thus, protein arrays may be used to detect SARS virus polypeptides in a patient; distinguish a SARS virus polypeptide from a host polypeptide; detect interactions between SARS virus polypeptides and for example host proteins; determine the efficacy of potential therapeutics, such as small molecules or ligands that may bind SARS virus polypeptides; determine protein-antibody interactions; and/or detect the interaction of enzyme-substrate interactions. Protein arrays may also be used to detect SARS virus antigens and antibodies in samples; to profile expression of SARS virus polypeptides; to identify suitable antibodies or map epitopes; or for a variety of protein function analyses.

A variety of methods are known for making and using microarrays, as for example disclosed in Cheung V. G., et al., 1999. *Nature Genetics Supplement*, 21:15-19; Lipshutz R. J., et al., 1999. *Nature Genetics Supplement*, 21:20-24; Bowtell D. D. L., 1999. *Nature Genetics Supplement*, 21:25-32; Singh-Gasson S., et al., 1999. *Nature Biotechnol.*, 17:974-978; and Schweitzer B., et al., 2002. *Nature Biotechnol.*, 20:359-365. Thus, for example, microarrays may be designed by synthesizing oligonucleotides with sequence variations based on a reference sequences, such as any SARS virus sequences described herein. Methods for storing, querying and analyzing microarray data have for example been disclosed in, for example, U.S. Pat. No. 6,484,183; U.S. Pat. No. 6,188,783; and Holloway A. J., et al., 2002. *Nature Genetics Supplement*, 32:481-489. Protein arrays may be constructed, detected, and analysed using methods known in the art for example mass spectrometric techniques, immunoassays such as ELISA and western (dot) blotting combined with for example fluorescence detection techniques, and adapted for high throughput analysis, as described in for example MacBeath, G. and Schreiber, S. L. *Science* 2000, 289, 1760-1763; Levit-Binnun N, et al. (2003) Quantitative detection of protein arrays. *Anal Chem* 75:1436-41; Kukar T, et al. (2002) Protein microarrays to detect protein-protein interactions using red and green fluorescent proteins. *Anal Biochem* 306: 50-4; Borrebaeck C A, et al. (2001) Protein chips based on recombinant antibody fragments: a highly sensitive approach as detected by mass spectrometry. *Biotechniques* 30:1126-1132; Huang R P (2001) Detection of multiple proteins in an antibody-based protein microarray system. *J Immunol Methods* 255:1-13; Emili A Q and Cagney G (2000) Large-scale functional analysis using peptide or protein arrays. *Nature Biotechnol* 18:393-397; Zhu H, et al. (2000) Analysis of yeast protein kinases using protein chips. *Nature Genet* 26:283-9; Lueking A, et al. (1999) Protein Microarrays for Gene Expression and Antibody Screening. *Anal. Biochem.* 270:103-111; or Templin M F, et al. (2002) Protein microarray technology. *Drug Discov Today* 7:815-822. Tools for microarray techniques are available commercially from

for example Affymetrix, Santa Clara, Calif.; Nanogen, San Diego, Calif.; or Sequenom, San Diego, Calif.

Computer Readable Records

Nucleic acid and polypeptide sequences, as described herein, or a fragment thereof, may be provided in a variety of media to facilitate access to these sequences and enable the use thereof. According, SARS virus nucleic acid and polypeptide sequences of the invention may be recorded or stored on computer readable media, using any technique and format that is appropriate for the particular medium.

In alternative embodiments, the invention provides computer readable media encoded with a number of distinct nucleic acid or amino acid data sequences of the invention. The number of distinct sequences may for example be any integer between 2 and 1×10^5 , such as at least 10^2 , 10^3 , 10^4 or 10^5 . In one embodiment, the invention features a computer medium having a plurality of digitally encoded data records. Each data record may include a value representing a nucleic acid or amino acid sequence of the invention. In some embodiments, the data record may further include values representing the level of expression, level or activity of a nucleic acid or amino acid sequence of the invention. The data record can be structured as a table, for example, a table that is part of a database such as a relational database (for example, a SQL database of the Oracle or Sybase database environments). The invention also includes a method of communicating information about a sample, for example by transmitting information, for example transmitting a computer readable record as described herein, for example over a computer network. The polypeptide and nucleic acid sequences of the invention, and sequence information pertaining thereto, may be routinely accessed by one of ordinary skill in the art for a variety of purposes, including for the purposes of comparing substantially identical sequences, etc. Such access may be facilitated using publicly available software as described herein. By "computer readable media" is meant any medium that can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media.

Pharmaceutical and Veterinary Compositions, Dosages, and Administration

Compounds of the invention can be provided alone or in combination with other compounds (for example, small molecules, peptides, or peptide analogues), in the presence of a liposome, an adjuvant, or any pharmaceutically acceptable carrier, in a form suitable for administration to humans or to animals.

Conventional pharmaceutical practice may be employed to provide suitable formulations or compositions to administer the compounds to patients suffering from or presymptomatic for SARS. Any appropriate route of administration may be employed, for example, parenteral, intravenous, subcutaneous, intramuscular, intracranial, intraorbital, ophthalmic, intraventricular, intracapsular, intraspinal, intracisternal, intraperitoneal, intranasal, aerosol, or oral administration. In some embodiments, compounds are delivered directly to the lung, by for example, formulations suitable for inhalation. In some embodiments, gene therapy techniques may be used for administration of SARS virus nucleic acid molecules, for example, as DNA vaccines. Formulations may be in the form of liquid solutions or suspensions; for oral administration,

formulations may be in the form of tablets or capsules; and for intranasal formulations, in the form of powders, nasal drops, or aerosols.

Methods well known in the art for making formulations are found in, for example, "Remington's Pharmaceutical Sciences" (18th edition), ed. A. Gennaro, 1990, Mack Publishing Company, Easton, Pa. Formulations for parenteral administration may, for example, contain excipients, sterile water, or saline, polyalkylene glycols such as polyethylene glycol, oils of vegetable origin, or hydrogenated naphthalenes. Biocompatible, biodegradable lactide polymer, lactide/glycolide copolymer, or polyoxyethylene-polyoxypropylene copolymers may be used to control the release of the compounds. Other potentially useful parenteral delivery systems for modulatory compounds include ethylene-vinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, and liposomes. Formulations for inhalation may contain excipients, for example, lactose, or may be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or may be oily solutions for administration in the form of nasal drops, or as a gel.

If desired, treatment with a compound according to the invention may be combined with more traditional therapies for the disease.

For therapeutic or prophylactic compositions, the compounds are administered to an individual in an amount sufficient to stop or slow the replication of the SARS virus, or to confer protective immunity against future SARS virus infection. Amounts considered sufficient will vary according to the specific compound used, the mode of administration, the stage and severity of the disease, the age, sex, and health of the individual being treated, and concurrent treatments. As a general rule, however, dosages can range from about 1 µg to about 100 mg per kg body weight of a patient for an initial dosage, with subsequent adjustments depending on the patient's response, which can be measured, for example by determining the presence of SARS nucleic acid molecules, polypeptides, or virions in the patient's peripheral blood.

In the case of vaccine formulations, an immunogenically effective amount of a compound of the invention can be provided, alone or in combination with other compounds, with an adjuvant, for example, Freund's incomplete adjuvant or aluminum hydroxide. The compound may also be linked with a carrier molecule, such as bovine serum albumin or keyhole limpet hemocyanin to enhance immunogenicity. In general, compounds of the invention should be used without causing substantial toxicity. Toxicity of the compounds of the invention can be determined using standard techniques, for example, by testing in cell cultures or experimental animals and determining the therapeutic index, i.e., the ratio between the LD50 (the dose lethal to 50% of the population) and the LD100 (the dose lethal to 100% of the population). In some circumstances however, such as in severe disease conditions, it may be necessary to administer substantial excesses of the compositions.

Virus Isolation

Virus isolation was performed on a bronchoalveolar lavage specimen of a fatal SARS case belonging to the original case cluster from Toronto, Canada. All work with the infectious agent was performed in a biosafety level 3 (BSL3) laboratory using a N100 mask for personal protection. Samples were removed from BSL3 after addition of the RNA extraction buffer. The virus isolate, named the "Tor2 isolate" was grown in African Green Monkey Kidney (Vero E6) cells, the viral particles were purified, and the genetic material (RNA) was extracted from the Tor2 isolate (Poutanen, S. M. et al., N Engl

J Med, Apr. 10, 2003). More specifically, one hundred microlitre specimens were used to inoculate Vero E6 cells (ATCC CRL 1586) on Dulbecco's Modified Eagle Medium supplemented with penicillin/streptomycin, glutamine and 2% fetal calf serum. The culture was incubated at 37° C. Cytopathogenic effect was observed 5 days post inoculation. The virus was passaged into newly seeded Vero E6 cells which showed a cytopathogenic effect as early as 2 days post infection (multiplicity of infection 10⁻²). A virus stock was prepared from passage 2 of these cells and preserved in liquid nitrogen. The titer of the virus stock was determined to be 1×10⁷ plaque forming units (p.f.u.) by plaque assay and 5×10⁶ by tissue culture infectious dose (TCID) 50.

For virus propagation, 10×T-162 flasks of Vero E6 cells were infected with a multiplicity of infection of 10⁻². When infected cells showed a cytopathogenic effect of '4+' (48 hours post infection), the cultures were then frozen and thawed to lyse the cells, and the supernatants were clarified from cell debris by centrifugation at 10,000 rpm in a Beckman high-speed centrifuge. The supernatants were treated with DNase and RNase for 3 hours at 37° C. to remove any cellular genomic nucleic acids and subsequently extracted with an equal volume of 1,1,2-trichloro-trifluoroethane. The top fraction was ultra-centrifuged through a 5%/40% glycerol step gradient at 151,000×g for 1 hour at 4° C. The virus pellet was resuspended in PBS. RNA was isolated using a commercial kit from QIAGEN and stored at -80° C. for further use.

cDNA Library Construction

The RNA and subsequent products were handled under biosafety level 2 (BSL2) conditions. The RNA sample was converted to a cDNA library, using a combined random-priming and oligo-dT priming strategy, and resultant subgenomic clones were processed under level 1 biosafety conditions. More specifically, purified viral RNA (55 ng) was used in the construction of a random primed and oligo-dT primed cDNA library, using the SuperScript Choice System for cDNA synthesis (Invitrogen). Linkers 5'-AATTCGCGGC-CGCGTCGAC-3', SEQ ID NO: 195, and 5'-pGTC-GACGCGGCCGCG-3', SEQ ID NO: 196, were ligated following cDNA synthesis. The cDNA synthesis products were visualized on agarose gels, revealing the anticipated low-yield smear. To produce sufficient cDNA for cloning, the cDNA product was size fractionated on a low-melting point preparative agarose gel, followed by PCR amplification using a single PCR primer 5'-AATTCGCGGCCGCGTCGAC-3', SEQ ID NO: 197, specific to the linkers. This yielded sufficient material for cloning.

Size-selected cDNA products were cloned and single sequence reads were generated from each end of the insert from randomly picked clones. A list of the SARS virus clones is provided in the accompanying sequence listing, which is incorporated by reference herein (SEQ ID NOs: 92-159, 208 and 209).

More specifically, size-selected cDNAs were ligated into the pCR4-TOPO TA cloning vector (Invitrogen, CA), or after digestion with the restriction nuclease Not I into the pBR194c vector (The Institute for Genomic Research, Rockville, Md., USA). Ligated clones were then transformed by electroporation into DH10B T1 cells (Invitrogen), plated on 22 cm agar plates with the appropriate antibiotic and grown for 16 hours at 37° C. Colonies were picked into 384-well Axygen culture blocks containing 2×YT media and grown in a shaking incubator for 18 hours at 37° C. Cells were lysed and DNA purified using standard laboratory procedures. Sequencing primers for the 194c clones were 5'-GGCCTCTTCGCTAT-

TACGC-3' (forward primer) (SEQ ID NO: 159) and 5' TGCAGGTCGACTCTAGAGGAT-3' (reverse primer) (SEQ ID NO: 198).

DNA Sequencing and Assembly of Reads

Sequences were assembled and the assembly edited to produce the genomic sequence of the SARS virus. More specifically, DNA sequencing of both ends of the plasmid templates was achieved using Applied Biosystems BigDye terminator reagent (version 3), with electrophoresis and data collection on AB 3700 and 3730 XL instruments DNA sequence reads were screened for non-viral contaminating sequences, trimmed for quality using PHRED (Ewing, B, and P. Green, *Genome Res* 8, 186-94, March, 1998) and assembled using PHRAP (Gordon, D. et al. *Genome Res* 8, 195-202, March, 1998). Simultaneously, sequences were used in BLAST searches of viral nucleotide and non-redundant protein datasets (NCBI, National Library of Medicine) to search for similarities. Sequence assemblies were visualized using CONSED (Gordon, D. et al. *Genome Res* 8, 195-202, March, 1998). Sequence mis-assemblies and contig joins were identified using Miropeats (Parsons, J. D., *Comput Appl Biosci* 11, 615-9 (December, 1995). As sequence data accrued, the additional sequences were assembled until it became apparent that the additional depth of sampling was increasing depth of coverage but not extending the length of the contig. At this point, 3,080 sequencing reads were generated, 2,634 of which were assembled into a single large contig.

The sequence information was imported into an ACEDB database (Durbin, J. Thierry-Mieg. 1991. A C. elegans Database. Documentation, code and data available from anonymous FTP servers at lirmm "dot" lirmm "dot" fr; cele "dot" mrc-1mb "dot" cam "dot" ac "dot" uk; and ncbi "dot" nlm "dot" nih "dot" gov) and subjected to biological analysis including the identification of open reading frames, detection of similar sequences by BLAST and searching for apparent frameshifts. When frameshifts were identified by this analysis, the sequence assembly was consulted for evidence of sequencing errors and if found, they were corrected. The sequences were also searched for any that could extend the 5' end of the sequence and these were incorporated when found. High quality sequence discrepancies between different

sequence reads were identified and resolved. Sequence reads classified as deleted or chimeric were identified through manual inspection and removed from the assembly. The resulting sequence has an average PHRED consensus quality score of 89.96. The lowest quality bases in the assembly are in the immediate vicinity of the 5' and 3' ends of the viral genome, with the lowest quality base having a PHRED score of 35. Most (29,694 of the 29,736 (99.86%)) of the bases have a consensus score of 90. Almost all regions of the genome are represented by reads derived from both strands of the plasmid sequencing templates, the exceptions being 50 bases at the 5' end represented by a single sequencing read, and 5 bases at the 3' end represented by a single read. The average base in the assembly is represented by 30 reads in the forward direction and 30 reads in the reverse direction, as determined by PHRED. RT-PCR products predicted from the sequence and spanning the entire genome yield PCR products of the anticipated size on agarose gels. To confirm the 5' end of the viral genome RACE was performed using the RLM-RACE kit from Ambion, and primers 5'-CAGGAAACAGCTATGACACCAAGAACAAGGCTCTCCA-3' (SEQ ID NO: 90) and 5'-CAGGAAACAGCTATGACGATAGGGCCTCTTCCACAGA-3' (SEQ ID NO: 91). Fourteen clones were recovered and sequenced. Analysis of these sequences confirmed the 5' end of the coronavirus genome. The SARS genomic sequences have been deposited into Genbank (Accession Nos. AY274119.1, AY274119.2, and AY274119.3).

While the invention has been described in connection with specific embodiments thereof, it will be understood that it is capable of further modifications and this application is intended to cover any variations, uses, or adaptations of the invention following, in general, the principles of the invention and including such departures from the present disclosure that come within known or customary practice within the art to which the invention pertains, and may be applied to the essential features set forth herein and in the scope of the appended claims.

All patents, patent applications, and publications referred to herein are hereby incorporated by reference in their entirety to the same extent as if each individual patent, patent application, or publication was specifically and individually indicated to be incorporated by reference in its entirety.

SEQUENCE LISTING

<160> NUMBER OF SEQ ID NOS: 206

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tcgggaacat ggctgactta tcatggagcc attaaattgg atgacaaaaga tccacaattc 29160
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gagcctaaaa aggacaaaaa gaaaaagact gatgaagctc agcctttgcc gcagagacaa 29280
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ctgctatat ggaagagccc taatgtgtaa aattaatfff agtagtgcta tccccatgtg 29700
attttaatag cttcttagga gaatgacaaa aaaaaaaaaa aaaaaaaaaa a 29751

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<210> SEQ ID NO 16
<211> LENGTH: 47
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 16
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acattttcat cgaggccacg cggagtacga tcgagggtac agtgaat 47
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<210> SEQ ID NO 17
<211> LENGTH: 32
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 17
```

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cgaggccacg cggagtacga tcgagggtac ag 32
```

```

<210> SEQ ID NO 18
<211> LENGTH: 339
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```
<400> SEQUENCE: 18
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acactcatga tgaccacaca aggcagatgg gctatgtaaa cgttttcgca attccgttta 60
cgatacatag tctactcttg tgcaaatga attctcgtaa ctaaacagca caagtagggt 120
tagttaactt taatctcaca tagcaatctt taatcaatgt gtaacattag ggaggacttg 180
aaagagccac cacattttca tcgaggccac gcgagtacg atcgagggta cagtgaataa 240
tgctagggag agctgctat atggaagagc cctaatgtgt aaaattaatt ttagtagtgc 300
tatccccatg tgattttaat agcttcttag gagaatgac 339

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<210> SEQ ID NO 19
<211> LENGTH: 35
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: s2m motif
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (5)..(5)
<223> OTHER INFORMATION: n is a, c, g, or t
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (23)..(23)
<223> OTHER INFORMATION: n is a, c, g, or t

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<400> SEQUENCE: 19
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gccgnggccca cgcsagtas gancgaggtt acags	35
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ucucuaaacg aacuuuaaaa ucugug	26
<210> SEQ ID NO 21 <211> LENGTH: 16 <212> TYPE: RNA <213> ORGANISM: Severe acute respiratory syndrome virus <400> SEQUENCE: 21	
caacuaaacg aacaug	16
<210> SEQ ID NO 22 <211> LENGTH: 18 <212> TYPE: RNA <213> ORGANISM: Severe acute respiratory syndrome virus <400> SEQUENCE: 22	
cacauaacg aacuuauug	18
<210> SEQ ID NO 23 <211> LENGTH: 16 <212> TYPE: RNA <213> ORGANISM: Severe acute respiratory syndrome virus <400> SEQUENCE: 23	
ugaguacgaa cuuauug	16
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ggucuaaacg aacuaacu	18
<210> SEQ ID NO 25 <211> LENGTH: 11 <212> TYPE: RNA <213> ORGANISM: Severe acute respiratory syndrome virus <400> SEQUENCE: 25	
aacuuaaaau u	11
<210> SEQ ID NO 26 <211> LENGTH: 17 <212> TYPE: RNA <213> ORGANISM: Severe acute respiratory syndrome virus <400> SEQUENCE: 26	
uccauaaaac gaacaug	17
<210> SEQ ID NO 27 <211> LENGTH: 24 <212> TYPE: RNA <213> ORGANISM: Severe acute respiratory syndrome virus	

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<400> SEQUENCE: 27
ugcucuagua uuuuuuuuac uuug 24

<210> SEQ ID NO 28
<211> LENGTH: 16
<212> TYPE: RNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 28
agucuaaacg aacaug 16

<210> SEQ ID NO 29
<211> LENGTH: 15
<212> TYPE: RNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 29
cuaaaaaacc ucaug 15

<210> SEQ ID NO 30
<211> LENGTH: 24
<212> TYPE: RNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 30
uaaaaaaacg aacaaaauaa aaug 24

<210> SEQ ID NO 31
<211> LENGTH: 136
<212> TYPE: DNA
<213> ORGANISM: Equine rhinovirus

<400> SEQUENCE: 31
accggttacc ctaaaattcc ctcccccttc tcttcaactcg ccgaggccac gccgagtagg 60
accgagggta cagcgagtct tttagtttaa ggtgtagat gtaaggtacg tgggctttct 120
tttggtttac ttcttc 136

<210> SEQ ID NO 32
<211> LENGTH: 178
<212> TYPE: DNA
<213> ORGANISM: Avian infectious bronchitis

<400> SEQUENCE: 32
tagtttagtt taagttagtt tagagtaggt ataaagatgc cagtgccggg gccacgcgga 60
gtacgatcga gggtagacga ctaggacgcc cattagggga agagctaaat tttagtttaa 120
gttaagttta attggctaag tatagttaaa atttataggc tagtatagag ttagagca 178

<210> SEQ ID NO 33
<211> LENGTH: 1255
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 33
Met Phe Ile Phe Leu Leu Phe Leu Thr Leu Thr Ser Gly Ser Asp Leu
1 5 10 15
Asp Arg Cys Thr Thr Phe Asp Asp Val Gln Ala Pro Asn Tyr Thr Gln
20 25 30
His Thr Ser Ser Met Arg Gly Val Tyr Tyr Pro Asp Glu Ile Phe Arg
35 40 45

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Ser Asp Thr Leu Tyr Leu Thr Gln Asp Leu Phe Leu Pro Phe Tyr Ser
 50 55 60
 Asn Val Thr Gly Phe His Thr Ile Asn His Thr Phe Gly Asn Pro Val
 65 70 75 80
 Ile Pro Phe Lys Asp Gly Ile Tyr Phe Ala Ala Thr Glu Lys Ser Asn
 85 90 95
 Val Val Arg Gly Trp Val Phe Gly Ser Thr Met Asn Asn Lys Ser Gln
 100 105 110
 Ser Val Ile Ile Ile Asn Asn Ser Thr Asn Val Val Ile Arg Ala Cys
 115 120 125
 Asn Phe Glu Leu Cys Asp Asn Pro Phe Phe Ala Val Ser Lys Pro Met
 130 135 140
 Gly Thr Gln Thr His Thr Met Ile Phe Asp Asn Ala Phe Asn Cys Thr
 145 150 155 160
 Phe Glu Tyr Ile Ser Asp Ala Phe Ser Leu Asp Val Ser Glu Lys Ser
 165 170 175
 Gly Asn Phe Lys His Leu Arg Glu Phe Val Phe Lys Asn Lys Asp Gly
 180 185 190
 Phe Leu Tyr Val Tyr Lys Gly Tyr Gln Pro Ile Asp Val Val Arg Asp
 195 200 205
 Leu Pro Ser Gly Phe Asn Thr Leu Lys Pro Ile Phe Lys Leu Pro Leu
 210 215 220
 Gly Ile Asn Ile Thr Asn Phe Arg Ala Ile Leu Thr Ala Phe Ser Pro
 225 230 235 240
 Ala Gln Asp Ile Trp Gly Thr Ser Ala Ala Ala Tyr Phe Val Gly Tyr
 245 250 255
 Leu Lys Pro Thr Thr Phe Met Leu Lys Tyr Asp Glu Asn Gly Thr Ile
 260 265 270
 Thr Asp Ala Val Asp Cys Ser Gln Asn Pro Leu Ala Glu Leu Lys Cys
 275 280 285
 Ser Val Lys Ser Phe Glu Ile Asp Lys Gly Ile Tyr Gln Thr Ser Asn
 290 295 300
 Phe Arg Val Val Pro Ser Gly Asp Val Val Arg Phe Pro Asn Ile Thr
 305 310 315 320
 Asn Leu Cys Pro Phe Gly Glu Val Phe Asn Ala Thr Lys Phe Pro Ser
 325 330 335
 Val Tyr Ala Trp Glu Arg Lys Lys Ile Ser Asn Cys Val Ala Asp Tyr
 340 345 350
 Ser Val Leu Tyr Asn Ser Thr Phe Phe Ser Thr Phe Lys Cys Tyr Gly
 355 360 365
 Val Ser Ala Thr Lys Leu Asn Asp Leu Cys Phe Ser Asn Val Tyr Ala
 370 375 380
 Asp Ser Phe Val Val Lys Gly Asp Asp Val Arg Gln Ile Ala Pro Gly
 385 390 395 400
 Gln Thr Gly Val Ile Ala Asp Tyr Asn Tyr Lys Leu Pro Asp Asp Phe
 405 410 415
 Met Gly Cys Val Leu Ala Trp Asn Thr Arg Asn Ile Asp Ala Thr Ser
 420 425 430
 Thr Gly Asn Tyr Asn Tyr Lys Tyr Arg Tyr Leu Arg His Gly Lys Leu
 435 440 445
 Arg Pro Phe Glu Arg Asp Ile Ser Asn Val Pro Phe Ser Pro Asp Gly
 450 455 460
 Lys Pro Cys Thr Pro Pro Ala Leu Asn Cys Tyr Trp Pro Leu Asn Asp

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465	470	475	480
Tyr Gly Phe Tyr Thr Thr Thr Gly Ile Gly Tyr Gln Pro Tyr Arg Val 485 490 495			
Val Val Leu Ser Phe Glu Leu Leu Asn Ala Pro Ala Thr Val Cys Gly 500 505 510			
Pro Lys Leu Ser Thr Asp Leu Ile Lys Asn Gln Cys Val Asn Phe Asn 515 520 525			
Phe Asn Gly Leu Thr Gly Thr Gly Val Leu Thr Pro Ser Ser Lys Arg 530 535 540			
Phe Gln Pro Phe Gln Gln Phe Gly Arg Asp Val Ser Asp Phe Thr Asp 545 550 555 560			
Ser Val Arg Asp Pro Lys Thr Ser Glu Ile Leu Asp Ile Ser Pro Cys 565 570 575			
Ala Phe Gly Gly Val Ser Val Ile Thr Pro Gly Thr Asn Ala Ser Ser 580 585 590			
Glu Val Ala Val Leu Tyr Gln Asp Val Asn Cys Thr Asp Val Ser Thr 595 600 605			
Ala Ile His Ala Asp Gln Leu Thr Pro Ala Trp Arg Ile Tyr Ser Thr 610 615 620			
Gly Asn Asn Val Phe Gln Thr Gln Ala Gly Cys Leu Ile Gly Ala Glu 625 630 635 640			
His Val Asp Thr Ser Tyr Glu Cys Asp Ile Pro Ile Gly Ala Gly Ile 645 650 655			
Cys Ala Ser Tyr His Thr Val Ser Leu Leu Arg Ser Thr Ser Gln Lys 660 665 670			
Ser Ile Val Ala Tyr Thr Met Ser Leu Gly Ala Asp Ser Ser Ile Ala 675 680 685			
Tyr Ser Asn Asn Thr Ile Ala Ile Pro Thr Asn Phe Ser Ile Ser Ile 690 695 700			
Thr Thr Glu Val Met Pro Val Ser Met Ala Lys Thr Ser Val Asp Cys 705 710 715 720			
Asn Met Tyr Ile Cys Gly Asp Ser Thr Glu Cys Ala Asn Leu Leu Leu 725 730 735			
Gln Tyr Gly Ser Phe Cys Thr Gln Leu Asn Arg Ala Leu Ser Gly Ile 740 745 750			
Ala Ala Glu Gln Asp Arg Asn Thr Arg Glu Val Phe Ala Gln Val Lys 755 760 765			
Gln Met Tyr Lys Thr Pro Thr Leu Lys Tyr Phe Gly Gly Phe Asn Phe 770 775 780			
Ser Gln Ile Leu Pro Asp Pro Leu Lys Pro Thr Lys Arg Ser Phe Ile 785 790 795 800			
Glu Asp Leu Leu Phe Asn Lys Val Thr Leu Ala Asp Ala Gly Phe Met 805 810 815			
Lys Gln Tyr Gly Glu Cys Leu Gly Asp Ile Asn Ala Arg Asp Leu Ile 820 825 830			
Cys Ala Gln Lys Phe Asn Gly Leu Thr Val Leu Pro Pro Leu Leu Thr 835 840 845			
Asp Asp Met Ile Ala Ala Tyr Thr Ala Ala Leu Val Ser Gly Thr Ala 850 855 860			
Thr Ala Gly Trp Thr Phe Gly Ala Gly Ala Ala Leu Gln Ile Pro Phe 865 870 875 880			
Ala Met Gln Met Ala Tyr Arg Phe Asn Gly Ile Gly Val Thr Gln Asn 885 890 895			

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Val Leu Tyr Glu Asn Gln Lys Gln Ile Ala Asn Gln Phe Asn Lys Ala
 900 905 910
 Ile Ser Gln Ile Gln Glu Ser Leu Thr Thr Thr Ser Thr Ala Leu Gly
 915 920 925
 Lys Leu Gln Asp Val Val Asn Gln Asn Ala Gln Ala Leu Asn Thr Leu
 930 935 940
 Val Lys Gln Leu Ser Ser Asn Phe Gly Ala Ile Ser Ser Val Leu Asn
 945 950 955 960
 Asp Ile Leu Ser Arg Leu Asp Lys Val Glu Ala Glu Val Gln Ile Asp
 965 970 975
 Arg Leu Ile Thr Gly Arg Leu Gln Ser Leu Gln Thr Tyr Val Thr Gln
 980 985 990
 Gln Leu Ile Arg Ala Ala Glu Ile Arg Ala Ser Ala Asn Leu Ala Ala
 995 1000 1005
 Thr Lys Met Ser Glu Cys Val Leu Gly Gln Ser Lys Arg Val Asp
 1010 1015 1020
 Phe Cys Gly Lys Gly Tyr His Leu Met Ser Phe Pro Gln Ala Ala
 1025 1030 1035
 Pro His Gly Val Val Phe Leu His Val Thr Tyr Val Pro Ser Gln
 1040 1045 1050
 Glu Arg Asn Phe Thr Thr Ala Pro Ala Ile Cys His Glu Gly Lys
 1055 1060 1065
 Ala Tyr Phe Pro Arg Glu Gly Val Phe Val Phe Asn Gly Thr Ser
 1070 1075 1080
 Trp Phe Ile Thr Gln Arg Asn Phe Phe Ser Pro Gln Ile Ile Thr
 1085 1090 1095
 Thr Asp Asn Thr Phe Val Ser Gly Asn Cys Asp Val Val Ile Gly
 1100 1105 1110
 Ile Ile Asn Asn Thr Val Tyr Asp Pro Leu Gln Pro Glu Leu Asp
 1115 1120 1125
 Ser Phe Lys Glu Glu Leu Asp Lys Tyr Phe Lys Asn His Thr Ser
 1130 1135 1140
 Pro Asp Val Asp Leu Gly Asp Ile Ser Gly Ile Asn Ala Ser Val
 1145 1150 1155
 Val Asn Ile Gln Lys Glu Ile Asp Arg Leu Asn Glu Val Ala Lys
 1160 1165 1170
 Asn Leu Asn Glu Ser Leu Ile Asp Leu Gln Glu Leu Gly Lys Tyr
 1175 1180 1185
 Glu Gln Tyr Ile Lys Trp Pro Trp Tyr Val Trp Leu Gly Phe Ile
 1190 1195 1200
 Ala Gly Leu Ile Ala Ile Val Met Val Thr Ile Leu Leu Cys Cys
 1205 1210 1215
 Met Thr Ser Cys Cys Ser Cys Leu Lys Gly Ala Cys Ser Cys Gly
 1220 1225 1230
 Ser Cys Cys Lys Phe Asp Glu Asp Asp Ser Glu Pro Val Leu Lys
 1235 1240 1245
 Gly Val Lys Leu His Tyr Thr
 1250 1255

<210> SEQ ID NO 34

<211> LENGTH: 220

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 34

Met Ala Asp Asn Gly Thr Ile Thr Val Glu Glu Leu Lys Gln Leu Leu
 1 5 10 15

Glu Gln Trp Asn Leu Val Ile Gly Phe Leu Phe Leu Ala Trp Ile Met
 20 25 30

Leu Leu Gln Phe Ala Tyr Ser Asn Arg Asn Arg Phe Leu Tyr Ile Ile
 35 40 45

Lys Leu Val Phe Leu Trp Leu Leu Trp Pro Val Thr Leu Ala Cys Phe
 50 55 60

Val Leu Ala Ala Val Tyr Arg Ile Asn Trp Val Thr Gly Gly Ile Ala
 65 70 75 80

Ile Ala Met Ala Cys Ile Val Gly Leu Met Trp Leu Ser Tyr Phe Val
 85 90 95

Ala Ser Phe Arg Leu Phe Ala Arg Thr Arg Ser Met Trp Ser Phe Asn
 100 105 110

Pro Glu Thr Asn Ile Leu Leu Asn Val Pro Leu Arg Gly Thr Ile Val
 115 120 125

Thr Arg Pro Leu Met Glu Ser Glu Leu Val Ile Gly Ala Val Ile Ile
 130 135 140

Arg Gly His Leu Arg Met Ala Gly His Ser Leu Gly Arg Cys Asp Ile
 145 150 155 160

Lys Asp Leu Pro Lys Glu Ile Thr Val Ala Thr Ser Arg Thr Leu Ser
 165 170 175

Tyr Tyr Lys Leu Gly Ala Ser Gln Arg Val Gly Thr Asp Ser Gly Phe
 180 185 190

Ala Ala Tyr Asn Arg Tyr Arg Ile Gly Asn Tyr Lys Leu Asn Thr Asp
 195 200 205

His Ala Gly Ser Asn Asp Asn Ile Ala Leu Leu Val
 210 215 220

<210> SEQ ID NO 35

<211> LENGTH: 76

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 35

Met Tyr Ser Phe Val Ser Glu Glu Thr Gly Thr Leu Ile Val Asn Ser
 1 5 10 15

Val Leu Leu Phe Leu Ala Phe Val Val Phe Leu Leu Val Thr Leu Ala
 20 25 30

Ile Leu Thr Ala Leu Arg Leu Cys Ala Tyr Cys Cys Asn Ile Val Asn
 35 40 45

Val Ser Leu Val Lys Pro Thr Val Tyr Val Tyr Ser Arg Val Lys Asn
 50 55 60

Leu Asn Ser Ser Glu Gly Val Pro Asp Leu Leu Val
 65 70 75

<210> SEQ ID NO 36

<211> LENGTH: 422

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 36

Met Ser Asp Asn Gly Pro Gln Ser Asn Gln Arg Ser Ala Pro Arg Ile
 1 5 10 15

Thr Phe Gly Gly Pro Thr Asp Ser Thr Asp Asn Asn Gln Asn Gly Gly

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20					25					30					
Arg	Asn	Gly	Ala	Arg	Pro	Lys	Gln	Arg	Arg	Pro	Gln	Gly	Leu	Pro	Asn
	35						40					45			
Asn	Thr	Ala	Ser	Trp	Phe	Thr	Ala	Leu	Thr	Gln	His	Gly	Lys	Glu	Glu
	50					55					60				
Leu	Arg	Phe	Pro	Arg	Gly	Gln	Gly	Val	Pro	Ile	Asn	Thr	Asn	Ser	Gly
	65				70					75				80	
Pro	Asp	Asp	Gln	Ile	Gly	Tyr	Tyr	Arg	Arg	Ala	Thr	Arg	Arg	Val	Arg
				85					90					95	
Gly	Gly	Asp	Gly	Lys	Met	Lys	Glu	Leu	Ser	Pro	Arg	Trp	Tyr	Phe	Tyr
			100					105					110		
Tyr	Leu	Gly	Thr	Gly	Pro	Glu	Ala	Ser	Leu	Pro	Tyr	Gly	Ala	Asn	Lys
		115					120					125			
Glu	Gly	Ile	Val	Trp	Val	Ala	Thr	Glu	Gly	Ala	Leu	Asn	Thr	Pro	Lys
	130					135					140				
Asp	His	Ile	Gly	Thr	Arg	Asn	Pro	Asn	Asn	Asn	Ala	Ala	Thr	Val	Leu
	145				150					155					160
Gln	Leu	Pro	Gln	Gly	Thr	Thr	Leu	Pro	Lys	Gly	Phe	Tyr	Ala	Glu	Gly
				165					170					175	
Ser	Arg	Gly	Gly	Ser	Gln	Ala	Ser	Ser	Arg	Ser	Ser	Ser	Arg	Ser	Arg
			180					185					190		
Gly	Asn	Ser	Arg	Asn	Ser	Thr	Pro	Gly	Ser	Ser	Arg	Gly	Asn	Ser	Pro
		195					200					205			
Ala	Arg	Met	Ala	Ser	Gly	Gly	Gly	Glu	Thr	Ala	Leu	Ala	Leu	Leu	Leu
	210					215					220				
Leu	Asp	Arg	Leu	Asn	Gln	Leu	Glu	Ser	Lys	Val	Ser	Gly	Lys	Gly	Gln
	225				230					235					240
Gln	Gln	Gln	Gly	Gln	Thr	Val	Thr	Lys	Lys	Ser	Ala	Ala	Glu	Ala	Ser
				245					250					255	
Lys	Lys	Pro	Arg	Gln	Lys	Arg	Thr	Ala	Thr	Lys	Gln	Tyr	Asn	Val	Thr
			260					265					270		
Gln	Ala	Phe	Gly	Arg	Arg	Gly	Pro	Glu	Gln	Thr	Gln	Gly	Asn	Phe	Gly
		275					280					285			
Asp	Gln	Asp	Leu	Ile	Arg	Gln	Gly	Thr	Asp	Tyr	Lys	His	Trp	Pro	Gln
	290					295					300				
Ile	Ala	Gln	Phe	Ala	Pro	Ser	Ala	Ser	Ala	Phe	Phe	Gly	Met	Ser	Arg
	305				310					315					320
Ile	Gly	Met	Glu	Val	Thr	Pro	Ser	Gly	Thr	Trp	Leu	Thr	Tyr	His	Gly
				325					330					335	
Ala	Ile	Lys	Leu	Asp	Asp	Lys	Asp	Pro	Gln	Phe	Lys	Asp	Asn	Val	Ile
			340					345					350		
Leu	Leu	Asn	Lys	His	Ile	Asp	Ala	Tyr	Lys	Thr	Phe	Pro	Pro	Thr	Glu
		355					360					365			
Pro	Lys	Lys	Asp	Lys	Lys	Lys	Lys	Thr	Asp	Glu	Ala	Gln	Pro	Leu	Pro
		370				375					380				
Gln	Arg	Gln	Lys	Lys	Gln	Pro	Thr	Val	Thr	Leu	Leu	Pro	Ala	Ala	Asp
	385				390					395					400
Met	Asp	Asp	Phe	Ser	Arg	Gln	Leu	Gln	Asn	Ser	Met	Ser	Gly	Ala	Ser
				405					410					415	
Ala	Asp	Ser	Thr	Gln	Ala										
			420												

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<211> LENGTH: 230
<212> TYPE: PRT
<213> ORGANISM: Bovine coronavirus

<400> SEQUENCE: 37

Met Ser Ser Val Thr Thr Pro Ala Pro Val Tyr Thr Trp Thr Ala Asp
 1           5           10           15
Glu Ala Ile Lys Phe Leu Lys Glu Trp Asn Phe Ser Leu Gly Ile Ile
 20           25           30
Leu Leu Phe Ile Thr Val Ile Leu Gln Phe Gly Tyr Thr Ser Arg Ser
 35           40           45
Met Phe Val Tyr Val Ile Lys Met Val Ile Leu Trp Leu Met Trp Pro
 50           55           60
Leu Thr Ile Ile Leu Thr Ile Phe Asn Cys Val Tyr Ala Leu Asn Asn
 65           70           75           80
Val Tyr Leu Gly Phe Ser Ile Val Phe Thr Ile Val Ala Ile Ile Met
 85           90           95
Trp Ile Val Tyr Phe Val Asn Ser Ile Arg Leu Phe Ile Arg Thr Gly
 100          105          110
Ser Trp Trp Ser Phe Asn Pro Glu Thr Asn Asn Leu Met Cys Ile Asp
 115          120          125
Met Lys Gly Arg Met Tyr Val Arg Pro Ile Ile Glu Asp Tyr His Thr
 130          135          140
Leu Thr Val Thr Ile Ile Arg Gly His Leu Tyr Met Gln Gly Ile Lys
 145          150          155          160
Leu Gly Thr Gly Tyr Ser Leu Ser Asp Leu Pro Ala Tyr Val Thr Val
 165          170          175
Ala Lys Val Ser His Leu Leu Thr Tyr Lys Arg Gly Phe Leu Asp Lys
 180          185          190
Ile Gly Asp Thr Ser Gly Phe Ala Val Tyr Val Lys Ser Lys Val Gly
 195          200          205
Asn Tyr Arg Leu Pro Ser Thr Gln Lys Gly Ser Gly Leu Asp Thr Ala
 210          215          220
Leu Leu Arg Asn Asn Ile
 225          230

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<210> SEQ ID NO 38
<211> LENGTH: 226
<212> TYPE: PRT
<213> ORGANISM: Avian infectious bronchitis virus

<400> SEQUENCE: 38

Met Ser Asn Gly Thr Glu Asn Cys Thr Leu Ser Thr Gln Gln Ala Ala
 1           5           10           15
Glu Leu Phe Lys Glu Tyr Asn Leu Phe Ile Thr Ala Phe Leu Leu Phe
 20           25           30
Leu Thr Ile Leu Leu Gln Tyr Gly Tyr Ala Thr Arg Ser Arg Phe Ile
 35           40           45
Tyr Ile Leu Lys Met Ile Val Leu Trp Cys Phe Trp Pro Leu Asn Ile
 50           55           60
Ala Val Gly Ile Ile Ser Cys Ile Tyr Pro Pro Asn Thr Gly Gly Leu
 65           70           75           80
Val Ala Ala Ile Ile Leu Thr Val Phe Ala Cys Leu Ser Phe Val Gly
 85           90           95
Tyr Trp Ile Gln Ser Phe Arg Leu Phe Lys Arg Cys Arg Ser Trp Trp
 100          105          110

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Ser Phe Asn Pro Glu Ser Asn Ala Val Gly Ser Ile Leu Leu Thr Asn
 115 120 125

Gly Gln Gln Cys Asn Phe Ala Ile Glu Ser Val Pro Met Val Leu Ser
 130 135 140

Pro Ile Ile Lys Asn Gly Ala Leu Tyr Cys Glu Gly Gln Trp Leu Ala
 145 150 155 160

Lys Cys Glu Pro Asp His Leu Pro Lys Asp Ile Phe Val Cys Thr Pro
 165 170 175

Asp Arg Arg Asn Ile Tyr Arg Met Val Gln Lys Tyr Thr Gly Asp Gln
 180 185 190

Ser Gly Asn Lys Lys Arg Phe Ala Thr Phe Val Tyr Ala Lys Gln Ser
 195 200 205

Val Asp Thr Gly Glu Leu Gly Ser Val Ala Thr Gly Gly Ser Ser Leu
 210 215 220

Tyr Thr
 225

<210> SEQ ID NO 39
 <211> LENGTH: 262
 <212> TYPE: PRT
 <213> ORGANISM: Transmissible gastroenteritis virus

<400> SEQUENCE: 39

Met Lys Ile Leu Leu Ile Leu Ala Cys Val Ile Ala Cys Ala Cys Gly
 1 5 10 15

Glu Arg Tyr Cys Ala Met Lys Ser Asp Thr Asp Leu Ser Cys Arg Asn
 20 25 30

Ser Thr Ala Ser Asp Cys Glu Ser Cys Phe Asn Gly Gly Asp Leu Ile
 35 40 45

Trp His Leu Ala Asn Trp Asn Phe Ser Trp Ser Ile Ile Leu Ile Val
 50 55 60

Phe Ile Thr Val Leu Gln Tyr Gly Arg Pro Gln Phe Ser Trp Phe Val
 65 70 75 80

Tyr Gly Ile Lys Met Leu Ile Met Trp Leu Leu Trp Pro Val Val Leu
 85 90 95

Ala Leu Thr Ile Phe Asn Ala Tyr Ser Glu Tyr Gln Val Ser Arg Tyr
 100 105 110

Val Met Phe Gly Phe Ser Ile Ala Gly Ala Ile Val Thr Phe Val Leu
 115 120 125

Trp Ile Met Tyr Phe Val Arg Ser Ile Gln Leu Tyr Arg Arg Thr Lys
 130 135 140

Ser Trp Trp Ser Phe Asn Pro Glu Thr Lys Ala Ile Leu Cys Val Ser
 145 150 155 160

Ala Leu Gly Arg Ser Tyr Val Leu Pro Leu Glu Gly Val Pro Thr Gly
 165 170 175

Val Thr Leu Thr Leu Leu Ser Gly Asn Leu Tyr Ala Glu Gly Phe Lys
 180 185 190

Ile Ala Gly Gly Met Asn Ile Asp Asn Leu Pro Lys Tyr Val Met Val
 195 200 205

Ala Leu Pro Ser Arg Thr Ile Val Tyr Thr Leu Val Gly Lys Lys Leu
 210 215 220

Lys Ala Ser Ser Ala Thr Gly Trp Ala Tyr Tyr Val Lys Ser Lys Ala
 225 230 235 240

Gly Asp Tyr Ser Thr Glu Ala Arg Thr Asp Asn Leu Ser Glu Gln Glu

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	245	250	255
Lys Leu Leu His Met Val			
	260		
<210> SEQ ID NO 40			
<211> LENGTH: 263			
<212> TYPE: PRT			
<213> ORGANISM: feline coronavirus			
<400> SEQUENCE: 40			
Met Lys Ile Leu Leu Ile Leu Ala Cys Ala Val Ala Cys Val Tyr Gly			
1	5	10	15
Glu Gln Ile Arg Tyr Cys Ala Met Gln Glu Thr Gly Leu Ser Cys Arg			
	20	25	30
Asn Gly Thr Ala Ser Asp Cys Glu Ser Cys Phe Asn Gly Gly Asp Leu			
	35	40	45
Ile Trp His Leu Ala Asn Trp Asn Phe Ser Trp Ser Ile Ile Leu Ile			
	50	55	60
Val Phe Ile Thr Val Leu Gln Tyr Gly Arg Pro Gln Phe Ser Trp Phe			
	65	70	75
Val Tyr Gly Ile Lys Met Leu Ile Met Trp Leu Leu Trp Pro Ile Val			
	85	90	95
Leu Ala Leu Thr Ile Phe Asn Ala Tyr Ser Glu Tyr Glu Val Ser Arg			
	100	105	110
Tyr Val Met Phe Gly Phe Ser Val Ala Gly Ala Val Val Thr Phe Ala			
	115	120	125
Leu Trp Met Met Tyr Phe Val Arg Ser Ile Gln Leu Tyr Arg Arg Thr			
	130	135	140
Lys Ser Trp Trp Ser Phe Asn Pro Glu Thr Asn Ala Ile Leu Cys Val			
	145	150	155
Asn Ala Leu Gly Arg Ser Tyr Val Leu Pro Leu Asp Gly Thr Pro Thr			
	165	170	175
Gly Val Thr Leu Thr Leu Leu Ser Gly Asn Leu Tyr Ala Glu Gly Phe			
	180	185	190
Lys Met Ala Gly Gly Leu Thr Ile Glu His Leu Pro Lys Tyr Val Met			
	195	200	205
Ile Arg Thr Pro Asn Arg Thr Ile Val Tyr Thr Leu Val Gly Lys Gln			
	210	215	220
Leu Lys Ala Thr Thr Ala Thr Gly Trp Ala Tyr Tyr Val Lys Ser Lys			
	225	230	235
Ala Gly Asp Tyr Ser Thr Glu Ala Arg Thr Asp Asn Leu Ser Glu His			
	245	250	255
Glu Lys Leu Leu His Met Val			
	260		

<210> SEQ ID NO 41
 <211> LENGTH: 231
 <212> TYPE: PRT
 <213> ORGANISM: Human coronavirus OC43
 MSSKTTTPAPVYIWTADEAIKFLKEWNFSLGIILLFITIILQPGYTSRSMFVYVIKMIILWLMNPLT
 IILTIFNCVYALNNVYLGSLIVFTIVAIIMWIVYFVNSIRLFI RTGSFWSFNPETNNLMCIDMKGT
 MYVRPIIEDYHTLTVTIIRGHLYIQGIKLGTGYSWADLPAYMTVAKVTHLCTYKRGFLDRISDTSG
 FAVVKS KVGNYRLPSTQKSGMDTALLRNNI
 <SEQ ID NO:37;prt;Porcine hemagglutinating encephalomyelitis virus
 <400> SEQUENCE: 41

Met Ser Ser Pro Thr Thr Pro Val Pro Val Ile Ser Trp Thr Ala Asp			
1	5	10	15

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Glu Ala Ile Lys Phe Leu Lys Glu Trp Asn Phe Ser Leu Gly Ile Ile
 20 25 30
 Val Leu Phe Ile Thr Ile Ile Leu Gln Phe Gly Tyr Thr Ser Arg Ser
 35 40 45
 Met Phe Val Tyr Val Ile Lys Met Val Ile Leu Trp Leu Met Trp Pro
 50 55 60
 Leu Thr Ile Ile Leu Thr Ile Phe Asn Cys Val Tyr Ala Leu Asn Asn
 65 70 75 80
 Val Tyr Leu Gly Phe Ser Ile Val Phe Thr Ile Val Ala Ile Ile Met
 85 90 95
 Trp Val Val Tyr Phe Val Asn Ser Ile Arg Leu Phe Ile Arg Thr Gly
 100 105 110
 Ser Trp Trp Ser Phe Asn Pro Glu Thr Asn Asn Leu Met Cys Ile Asp
 115 120 125
 Met Lys Gly Arg Met Tyr Val Arg Pro Ile Ile Glu Asp Tyr His Thr
 130 135 140
 Leu Thr Ala Thr Ile Ile Arg Gly His Leu Tyr Ile Gln Gly Ile Lys
 145 150 155 160
 Leu Gly Thr Gly Tyr Ser Leu Ser Asp Leu Pro Ala Tyr Val Thr Val
 165 170 175
 Ala Lys Val Thr His Leu Cys Thr Tyr Lys Arg Gly Phe Leu Asp Arg
 180 185 190
 Ile Gly Asp Thr Ser Gly Phe Ala Val Tyr Val Lys Ser Lys Val Gly
 195 200 205
 Asn Tyr Arg Leu Pro Ser Thr His Lys Gly Ser Gly Met Asp Thr Ala
 210 215 220
 Leu Leu Arg Asn Asn Ile Met
 225 230

<210> SEQ ID NO 42

<211> LENGTH: 223

<212> TYPE: PRT

<213> ORGANISM: Avian infectious bronchitis virus

<400> SEQUENCE: 42

Met Met Glu Asn Cys Thr Leu Asn Leu Glu Gln Ala Thr Leu Leu Phe
 1 5 10 15
 Lys Glu Tyr Asn Leu Phe Ile Thr Ala Phe Leu Leu Phe Leu Thr Ile
 20 25 30
 Leu Leu Gln Tyr Gly Tyr Ala Thr Arg Ser Arg Phe Ile Tyr Ile Leu
 35 40 45
 Lys Met Ile Val Leu Trp Cys Phe Trp Pro Leu Asn Ile Ala Val Gly
 50 55 60
 Val Ile Ser Cys Ile Tyr Pro Pro Asn Thr Gly Gly Leu Val Ala Ala
 65 70 75 80
 Ile Ile Leu Thr Val Phe Ala Cys Leu Ser Phe Val Gly Tyr Trp Ile
 85 90 95
 Gln Ser Cys Arg Leu Phe Lys Arg Cys Arg Ser Trp Trp Ser Phe Asn
 100 105 110
 Pro Glu Ser Asn Ala Val Gly Ser Ile Leu Leu Thr Asn Gly Gln Gln
 115 120 125
 Cys Asn Phe Ala Ile Glu Ser Val Pro Met Val Leu Ala Pro Ile Ile
 130 135 140
 Lys Asn Gly Val Leu Tyr Cys Glu Gly Gln Trp Leu Ala Lys Cys Glu

-continued

145	150	155	160
Pro Asp His Leu	Pro Lys Asp Ile Phe Val Cys Thr	Pro Asp Arg Arg	
	165	170	175
Asn Ile Tyr Arg	Met Val Gln Lys Tyr Thr Gly Asp Gln Ser Gly Asn		
	180	185	190
Lys Lys Arg Val Ala Thr Phe Val Tyr Ala Lys Gln Ser Val Asp Thr			
	195	200	205
Gly Glu Leu Glu Ser Val Pro Thr Gly Gly Ser Ser Leu Tyr Thr			
	210	215	220

<210> SEQ ID NO 43

<211> LENGTH: 455

<212> TYPE: PRT

<213> ORGANISM: Mouse Hepatitis Virus

<400> SEQUENCE: 43

Met Ser Phe Val	Pro Gly Gln Glu Asn Ala Gly Ser Arg Ser Ser Ser		
1	5	10	15
Val Asn Arg Ala	Gly Asn Gly Ile Leu Lys Lys Thr Thr Trp Ala Asp		
	20	25	30
Gln Thr Glu Arg	Gly Pro Asn Asn Gln Asn Arg Gly Arg Arg Asn Gln		
	35	40	45
Pro Lys Gln Thr	Ala Thr Thr Gln Pro Asn Ser Gly Ser Val Val Pro		
	50	55	60
His Tyr Ser Trp	Phe Ser Gly Ile Thr Gln Phe Gln Lys Gly Lys Glu		
	65	70	75
Phe Gln Phe Ala	Gln Gly Gln Gly Val Pro Ile Ala Asn Gly Ile Pro		
	85	90	95
Ala Ser Glu Gln	Lys Gly Tyr Trp Tyr Arg His Asn Arg Arg Ser Phe		
	100	105	110
Lys Thr Pro Asp	Gly Gln Gln Lys Gln Leu Leu Pro Arg Trp Tyr Phe		
	115	120	125
Tyr Tyr Leu Gly	Thr Gly Pro His Ala Gly Ala Glu Tyr Gly Asp Asp		
	130	135	140
Ile Asp Gly Val	Val Trp Val Ala Ser Gln Gln Ala Asp Thr Lys Thr		
	145	150	155
Thr Ala Asp Ile	Val Glu Arg Asp Pro Ser Ser His Glu Ala Ile Pro		
	165	170	175
Thr Arg Phe Ala	Pro Gly Thr Val Leu Pro Gln Gly Phe Tyr Val Glu		
	180	185	190
Gly Ser Gly Arg	Ser Ala Pro Ala Ser Arg Ser Gly Ser Arg Ser Gln		
	195	200	205
Ser Arg Gly Pro	Asn Asn Arg Ala Arg Ser Ser Ser Asn Gln Arg Gln		
	210	215	220
Pro Ala Ser Thr	Val Lys Pro Asp Met Ala Glu Glu Ile Ala Ala Leu		
	225	230	235
Val Leu Ala Lys	Leu Gly Lys Asp Ala Gly Gln Pro Lys Gln Val Thr		
	245	250	255
Lys Gln Ser Ala	Lys Glu Val Arg Gln Lys Ile Leu Asn Lys Pro Arg		
	260	265	270
Gln Lys Arg Thr	Pro Asn Lys Gln Cys Pro Val Gln Gln Cys Phe Gly		
	275	280	285
Lys Arg Gly Pro	Asn Gln Asn Phe Gly Gly Ser Glu Met Leu Lys Leu		
	290	295	300

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Gly Thr Ser Asp Pro Gln Phe Pro Ile Leu Ala Glu Leu Ala Pro Thr
 305 310 315 320

Pro Ser Ala Phe Phe Phe Gly Ser Lys Leu Glu Leu Val Lys Lys Asn
 325 330 335

Ser Gly Gly Ala Asp Asp Pro Thr Lys Asp Val Tyr Glu Leu Gln Tyr
 340 345 350

Ser Gly Ala Ile Arg Phe Asp Ser Thr Leu Pro Gly Phe Glu Thr Ile
 355 360 365

Met Lys Val Leu Asn Glu Asn Leu Asp Ala Tyr Gln Asp Gln Ala Gly
 370 375 380

Gly Ala Asp Val Val Ser Pro Lys Pro Gln Arg Lys Arg Gly Thr Lys
 385 390 395 400

Gln Lys Ala Leu Lys Gly Glu Val Asp Asn Val Ser Val Ala Lys Pro
 405 410 415

Lys Ser Ser Val Gln Arg Asn Val Ser Arg Glu Leu Thr Pro Glu Asp
 420 425 430

Arg Ser Leu Leu Ala Gln Ile Leu Asp Asp Gly Val Val Pro Asp Gly
 435 440 445

Leu Glu Asp Asp Ser Asn Val
 450 455

<210> SEQ ID NO 44

<211> LENGTH: 448

<212> TYPE: PRT

<213> ORGANISM: Bovine coronavirus

<400> SEQUENCE: 44

Met Ser Phe Thr Pro Gly Lys Gln Ser Ser Ser Arg Ala Ser Ser Gly
 1 5 10 15

Asn Arg Ser Gly Asn Gly Ile Leu Lys Trp Ala Asp Gln Ser Asp Gln
 20 25 30

Ser Arg Asn Val Gln Thr Arg Gly Arg Arg Ala Gln Pro Lys Gln Thr
 35 40 45

Ala Thr Ser Gln Gln Pro Ser Gly Gly Asn Val Val Pro Tyr Tyr Ser
 50 55 60

Trp Phe Ser Gly Ile Thr Gln Phe Gln Lys Gly Lys Glu Phe Glu Phe
 65 70 75 80

Ala Glu Gly Gln Gly Val Pro Ile Ala Pro Gly Val Pro Ala Thr Glu
 85 90 95

Ala Lys Gly Tyr Trp Tyr Arg His Asn Arg Arg Ser Phe Lys Thr Ala
 100 105 110

Asp Gly Asn Gln Arg Gln Leu Leu Pro Arg Trp Tyr Phe Tyr Tyr Leu
 115 120 125

Gly Thr Gly Pro His Ala Lys Asp Gln Tyr Gly Thr Asp Ile Asp Gly
 130 135 140

Val Tyr Trp Val Ala Ser Asn Gln Ala Asp Val Asn Thr Pro Ala Asp
 145 150 155 160

Ile Leu Asp Arg Asp Pro Ser Ser Asp Glu Ala Ile Pro Thr Arg Phe
 165 170 175

Pro Pro Gly Thr Val Leu Pro Gln Gly Tyr Tyr Ile Glu Gly Ser Gly
 180 185 190

Arg Ser Ala Pro Asn Ser Arg Ser Thr Ser Arg Ala Ser Ser Arg Ala
 195 200 205

Ser Ser Ala Gly Ser Arg Ser Arg Ala Asn Ser Gly Asn Arg Thr Pro
 210 215 220

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Thr Ser Gly Val Thr Pro Asp Met Ala Asp Gln Ile Ala Ser Leu Val
 225 230 235 240
 Leu Ala Lys Leu Gly Lys Asp Ala Ala Lys Pro Gln Gln Val Thr Lys
 245 250 255
 Gln Thr Ala Lys Glu Ile Arg Gln Lys Ile Leu Asn Lys Pro Arg Gln
 260 265 270
 Lys Arg Ser Pro Asn Lys Gln Cys Thr Val Gln Gln Cys Phe Gly Lys
 275 280 285
 Arg Gly Pro Asn Gln Asn Phe Gly Gly Gly Glu Met Leu Lys Leu Gly
 290 295 300
 Thr Ser Asp Pro Gln Phe Pro Ile Leu Ala Glu Leu Ala Pro Thr Ala
 305 310 315 320
 Gly Ala Phe Phe Phe Gly Ser Arg Leu Glu Leu Ala Lys Val Gln Asn
 325 330 335
 Leu Ser Gly Asn Leu Asp Glu Pro Gln Lys Asp Val Tyr Glu Leu Arg
 340 345 350
 Tyr Asn Gly Ala Ile Arg Phe Asp Ser Thr Leu Ser Gly Phe Glu Thr
 355 360 365
 Ile Met Lys Val Leu Asn Glu Asn Leu Asn Ala Tyr Gln Gln Gln Asp
 370 375 380
 Gly Thr Met Asn Met Ser Pro Lys Pro Gln Arg Gln Arg Gly Gln Lys
 385 390 395 400
 Asn Gly Gln Gly Glu Asn Asp Asn Ile Ser Val Ala Ala Pro Lys Ser
 405 410 415
 Arg Val Gln Gln Asn Lys Ile Arg Glu Leu Thr Ala Glu Asp Ile Ser
 420 425 430
 Leu Leu Lys Lys Met Asp Glu Pro Phe Thr Glu Asp Thr Ser Glu Ile
 435 440 445

<210> SEQ ID NO 45

<211> LENGTH: 409

<212> TYPE: PRT

<213> ORGANISM: Avian infectious bronchitis virus

<400> SEQUENCE: 45

Met Ala Ser Gly Lys Ala Ala Gly Lys Thr Asp Ala Pro Ala Pro Val
 1 5 10 15
 Ile Lys Leu Gly Gly Pro Lys Pro Pro Lys Val Gly Ser Ser Gly Asn
 20 25 30
 Ala Ser Trp Phe Gln Ala Leu Lys Ala Lys Lys Leu Asn Ala Pro Ala
 35 40 45
 Pro Lys Phe Glu Gly Ser Gly Val Pro Asp Asn Glu Asn Leu Lys Ile
 50 55 60
 Ser Gln Gln His Gly Tyr Trp Arg Arg Gln Ala Arg Tyr Lys Pro Gly
 65 70 75 80
 Lys Gly Gly Arg Lys Pro Val Pro Asp Ala Trp Tyr Phe Tyr Tyr Thr
 85 90 95
 Gly Thr Gly Pro Ala Ala Asp Leu Asn Trp Gly Asp Ser Gln Asp Gly
 100 105 110
 Ile Val Trp Val Ala Ala Lys Gly Ala Asp Val Lys Ser Arg Ser Asn
 115 120 125
 Gln Gly Thr Arg Asp Pro Asp Lys Phe Asp Gln Tyr Pro Leu Arg Phe
 130 135 140
 Ser Asp Gly Gly Pro Asp Gly Asn Phe Arg Trp Asp Phe Ile Pro Leu

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145		150		155		160									
Asn	Arg	Gly	Arg	Ser	Gly	Arg	Ser	Thr	Ala	Ala	Ser	Ser	Ala	Ala	Ser
				165					170					175	
Ser	Arg	Ala	Pro	Ser	Arg	Glu	Gly	Ser	Arg	Gly	Arg	Leu	Asn	Gly	Ala
			180					185					190		
Glu	Asp	Asp	Leu	Ile	Ala	Arg	Ala	Ala	Lys	Ile	Ile	Gln	Asp	Gln	Gln
	195						200					205			
Lys	Lys	Gly	Ser	Arg	Ile	Thr	Lys	Ala	Lys	Ala	Glu	Glu	Met	Ile	His
	210					215					220				
Arg	Arg	Tyr	Cys	Lys	Arg	Thr	Val	Pro	Pro	Gly	Val	Ser	Ile	Asp	Lys
225					230					235					240
Val	Phe	Gly	Pro	Arg	Thr	Lys	Gly	Lys	Glu	Gly	Asn	Phe	Gly	Asp	Asp
				245					250					255	
Lys	Met	Asn	Glu	Glu	Gly	Ile	Lys	Asp	Gly	Arg	Val	Thr	Ala	Met	Leu
			260					265					270		
Asn	Leu	Val	Pro	Ser	Ser	His	Ala	Cys	Leu	Phe	Gly	Ser	Gln	Val	Thr
		275					280					285			
Pro	Lys	Leu	Gln	Pro	Asp	Gly	Leu	His	Leu	Thr	Phe	Arg	Phe	Thr	Thr
	290					295					300				
Val	Val	Ser	Arg	Asp	Asp	Pro	Gln	Phe	Asp	Asn	Tyr	Val	Lys	Ile	Cys
305					310					315					320
Asp	Glu	Cys	Val	Asp	Gly	Val	Gly	Thr	Arg	Pro	Lys	Asp	Glu	Val	Val
				325					330					335	
Arg	Pro	Lys	Ser	Arg	Ser	Ser	Ser	Arg	Pro	Ala	Thr	Arg	Gly	Thr	Ser
			340					345					350		
Pro	Ala	Pro	Lys	Gln	Gln	Arg	Pro	Lys	Lys	Glu	Lys	Lys	Pro	Lys	Lys
		355					360					365			
Gln	Asp	Asp	Glu	Val	Asp	Lys	Ala	Leu	Thr	Ser	Asp	Glu	Glu	Arg	Asn
	370					375					380				
Asn	Ala	Gln	Leu	Glu	Phe	Asp	Asp	Glu	Pro	Lys	Val	Ile	Asn	Trp	Gly
385					390					395					400
Asp	Ser	Ala	Leu	Gly	Glu	Asn	Glu	Leu							
				405											

<210> SEQ ID NO 46

<211> LENGTH: 376

<212> TYPE: PRT

<213> ORGANISM: Feline coronavirus

<400> SEQUENCE: 46

Met	Ala	Thr	Gln	Gly	Gln	Arg	Val	Asn	Trp	Gly	Asp	Glu	Pro	Ser	Lys
1				5					10					15	
Arg	Arg	Gly	Arg	Ser	Asn	Ser	Arg	Gly	Arg	Lys	Asn	Asn	Asp	Ile	Pro
			20					25					30		
Leu	Ser	Tyr	Phe	Asn	Pro	Ile	Thr	Leu	Asp	Gln	Gly	Ser	Lys	Phe	Trp
		35					40					45			
Asn	Leu	Cys	Pro	Arg	Asp	Phe	Val	Pro	Lys	Gly	Ile	Gly	Asn	Lys	Asp
	50					55					60				
Gln	Gln	Ile	Gly	Tyr	Trp	Asn	Arg	Gln	Ala	Arg	Tyr	Arg	Ile	Val	Lys
65					70					75					80
Gly	Gln	Arg	Val	Glu	Leu	Pro	Glu	Arg	Trp	Phe	Phe	Tyr	Phe	Leu	Gly
				85					90					95	
Thr	Gly	Pro	His	Ala	Asp	Ala	Lys	Phe	Lys	Ala	Lys	Ile	Asp	Gly	Val
			100					105					110		

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Val Trp Val Ala Lys Asp Gly Ala Met Asn Lys Pro Thr Thr Leu Gly
 115 120 125
 Ser Arg Gly Ala Asn Asn Glu Ser Lys Ala Leu Lys Phe Asp Gly Lys
 130 135 140
 Val Pro Gly Glu Phe Gln Leu Glu Val Asn Gln Ser Arg Asp Asn Ser
 145 150 155 160
 Arg Leu Arg Ser Gln Ser Arg Ser Arg Ser Arg Asn Arg Ser Gln Ser
 165 170 175
 Arg Gly Arg Gln Gln Ser Asn Asn Lys Lys Asp Asp Ser Val Glu Gln
 180 185 190
 Ala Val Leu Ala Ala Leu Lys Lys Leu Gly Val Tyr Thr Glu Lys Gln
 195 200 205
 Gln Gln Arg Ser Arg Ser Lys Ser Lys Glu Arg Ser Asn Ser Lys Ile
 210 215 220
 Arg Asp Thr Thr Pro Lys Asn Glu Asn Lys His Thr Trp Lys Arg Thr
 225 230 235 240
 Ala Gly Lys Gly Asp Val Thr Arg Phe Tyr Gly Thr Arg Ser Asn Ser
 245 250 255
 Ala Asn Phe Gly Asp Ser Asp Leu Val Ala Asn Gly Ser Ser Ala Lys
 260 265 270
 His Tyr Pro Gln Leu Ala Glu Cys Val Pro Ser Val Ser Ser Ile Leu
 275 280 285
 Phe Gly Ser Tyr Trp Thr Ser Lys Glu Asp Gly Asp Gln Ile Glu Val
 290 295 300
 Thr Phe Thr His Lys Tyr His Leu Pro Lys Asp Asp Pro Lys Thr Gly
 305 310 315 320
 Gln Phe Leu Gln Gln Ile Asn Ala Tyr Ala Arg Pro Ser Glu Val Ala
 325 330 335
 Lys Glu Gln Arg Lys Arg Lys Ser Arg Ser Lys Ser Ala Glu Arg Ser
 340 345 350
 Glu Gln Glu Val Val Pro Asp Ala Leu Ile Glu Asn Tyr Thr Asp Val
 355 360 365
 Phe Asp Asp Thr Gln Val Glu Met Ile Asp Glu Val Thr Asn
 370 375 380

<210> SEQ ID NO 48

<211> LENGTH: 389

<212> TYPE: PRT

<213> ORGANISM: Human coronavirus 229E

<400> SEQUENCE: 48

Met Ala Thr Val Lys Trp Ala Asp Ala Ser Glu Pro Gln Arg Gly Arg
 1 5 10 15
 Gln Gly Arg Ile Pro Tyr Ser Leu Tyr Ser Pro Leu Leu Val Asp Ser
 20 25 30
 Glu Gln Pro Trp Lys Val Ile Pro Arg Asn Leu Val Pro Ile Asn Lys
 35 40 45
 Lys Asp Lys Asn Lys Leu Ile Gly Tyr Trp Asn Val Gln Lys Arg Phe
 50 55 60
 Arg Thr Arg Lys Gly Lys Arg Val Asp Leu Ser Pro Lys Leu His Phe
 65 70 75 80
 Tyr Tyr Leu Gly Thr Gly Pro His Lys Asp Ala Lys Phe Arg Glu Arg
 85 90 95
 Val Glu Gly Val Val Trp Val Ala Val Asp Gly Ala Lys Thr Glu Pro

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100					105					110					
Thr	Gly	Tyr	Gly	Val	Arg	Arg	Lys	Asn	Ser	Glu	Pro	Glu	Ile	Pro	His
		115					120					125			
Phe	Asn	Gln	Lys	Leu	Pro	Asn	Gly	Val	Thr	Val	Val	Glu	Glu	Pro	Asp
	130					135						140			
Ser	Arg	Ala	Pro	Ser	Arg	Ser	Gln	Ser	Arg	Ser	Gln	Ser	Arg	Gly	Arg
145					150					155					160
Gly	Glu	Ser	Lys	Pro	Gln	Ser	Arg	Asn	Pro	Ser	Ser	Asp	Arg	Asn	His
			165						170					175	
Asn	Ser	Gln	Asp	Asp	Ile	Met	Lys	Ala	Val	Ala	Ala	Ala	Leu	Lys	Ser
			180					185						190	
Leu	Gly	Phe	Asp	Lys	Pro	Gln	Glu	Lys	Asp	Lys	Lys	Ser	Ala	Lys	Thr
		195					200					205			
Gly	Thr	Pro	Lys	Pro	Ser	Arg	Asn	Gln	Ser	Pro	Ala	Ser	Ser	Gln	Thr
	210					215					220				
Ser	Ala	Lys	Ser	Leu	Ala	Arg	Ser	Gln	Ser	Ser	Glu	Thr	Lys	Glu	Gln
225					230					235					240
Lys	His	Glu	Met	Gln	Lys	Pro	Arg	Trp	Lys	Arg	Gln	Pro	Asn	Asp	Asp
			245						250					255	
Val	Thr	Ser	Asn	Val	Thr	Gln	Cys	Phe	Gly	Pro	Arg	Asp	Leu	Asp	His
			260					265					270		
Asn	Phe	Gly	Ser	Ala	Gly	Val	Val	Ala	Asn	Gly	Val	Lys	Ala	Lys	Gly
		275					280					285			
Tyr	Pro	Gln	Phe	Ala	Glu	Leu	Val	Pro	Ser	Thr	Ala	Ala	Met	Leu	Phe
	290					295					300				
Asp	Ser	His	Ile	Val	Ser	Lys	Glu	Ser	Gly	Asn	Thr	Val	Val	Leu	Thr
305					310					315					320
Phe	Thr	Thr	Arg	Val	Thr	Val	Pro	Lys	Asp	His	Pro	His	Leu	Gly	Lys
				325					330					335	
Phe	Leu	Glu	Glu	Leu	Asn	Ala	Phe	Thr	Arg	Glu	Met	Gln	Gln	His	Pro
			340					345					350		
Leu	Leu	Asn	Pro	Ser	Ala	Leu	Glu	Phe	Asn	Pro	Ser	Gln	Thr	Ser	Pro
		355					360					365			
Ala	Thr	Ala	Glu	Pro	Val	Arg	Asp	Glu	Val	Ser	Ile	Glu	Thr	Asp	Ile
	370					375					380				
Ile	Asp	Glu	Val	Asn											
385															

<210> SEQ ID NO 49

<211> LENGTH: 448

<212> TYPE: PRT

<213> ORGANISM: Human coronavirus

<400> SEQUENCE: 49

Met	Ser	Phe	Thr	Pro	Gly	Lys	Gln	Ser	Ser	Ser	Arg	Ala	Ser	Ser	Gly
1				5					10					15	
Asn	Arg	Ser	Gly	Asn	Gly	Ile	Leu	Lys	Trp	Ala	Asp	Gln	Ser	Asp	Gln
			20					25					30		
Val	Arg	Asn	Val	Gln	Thr	Arg	Gly	Arg	Arg	Ala	Gln	Pro	Lys	Gln	Thr
			35				40					45			
Ala	Thr	Ser	Gln	Gln	Pro	Ser	Gly	Gly	Asn	Val	Val	Pro	Tyr	Tyr	Ser
			50				55				60				
Trp	Phe	Ser	Gly	Ile	Thr	Gln	Phe	Gln	Lys	Gly	Lys	Glu	Phe	Glu	Phe
65					70					75					80

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Val Glu Gly Gln Gly Pro Pro Ile Ala Pro Gly Val Pro Ala Thr Glu
 85 90 95
 Ala Lys Gly Tyr Trp Tyr Arg His Asn Arg Gly Ser Phe Lys Thr Ala
 100 105 110
 Asp Gly Asn Gln Arg Gln Leu Leu Pro Arg Trp Tyr Phe Tyr Tyr Leu
 115 120 125
 Gly Thr Gly Pro His Ala Lys Asp Gln Tyr Gly Thr Asp Ile Asp Gly
 130 135 140
 Val Tyr Trp Val Ala Ser Asn Gln Ala Asp Val Asn Thr Pro Ala Asp
 145 150 155 160
 Ile Val Asp Arg Asp Pro Ser Ser Asp Glu Ala Ile Pro Thr Arg Phe
 165 170 175
 Pro Pro Gly Thr Val Leu Pro Gln Gly Tyr Tyr Ile Glu Gly Ser Gly
 180 185 190
 Arg Ser Ala Pro Asn Ser Arg Ser Thr Ser Arg Thr Ser Ser Arg Ala
 195 200 205
 Ser Ser Ala Gly Ser Arg Ser Arg Ala Asn Ser Gly Asn Arg Thr Pro
 210 215 220
 Thr Ser Gly Val Thr Pro Asp Met Ala Asp Gln Ile Ala Ser Leu Val
 225 230 235 240
 Leu Ala Lys Leu Gly Lys Asp Ala Thr Lys Pro Gln Gln Val Thr Lys
 245 250 255
 His Thr Ala Lys Glu Val Arg Gln Lys Ile Leu Asn Lys Pro Arg Gln
 260 265 270
 Lys Arg Ser Pro Asn Lys Gln Cys Thr Val Gln Gln Cys Phe Gly Lys
 275 280 285
 Arg Gly Pro Asn Gln Asn Phe Gly Gly Gly Glu Met Leu Lys Leu Gly
 290 295 300
 Thr Ser Asp Pro Gln Phe Pro Ile Leu Ala Glu Leu Ala Pro Thr Ala
 305 310 315 320
 Gly Ala Phe Phe Phe Gly Ser Arg Leu Glu Leu Ala Lys Val Gln Asn
 325 330 335
 Leu Ser Gly Asn Pro Asp Glu Pro Gln Lys Asp Val Tyr Glu Leu Arg
 340 345 350
 Tyr Asn Gly Ala Ile Arg Phe Asp Ser Thr Leu Ser Gly Phe Glu Thr
 355 360 365
 Ile Met Lys Val Leu Asn Glu Asn Leu Asn Ala Tyr Gln Gln Gln Asp
 370 375 380
 Gly Met Met Asn Met Ser Pro Lys Pro Gln Arg Gln Arg Gly His Lys
 385 390 395 400
 Asn Gly Gln Gly Glu Asn Asp Asn Ile Ser Val Ala Val Pro Lys Ser
 405 410 415
 Arg Val Gln Gln Asn Lys Ser Arg Glu Leu Thr Ala Glu Asp Ile Ser
 420 425 430
 Leu Leu Lys Lys Met Asp Glu Pro Tyr Thr Glu Asp Thr Ser Glu Ile
 435 440 445

<210> SEQ ID NO 50

<211> LENGTH: 449

<212> TYPE: PRT

<213> ORGANISM: porcine hemagglutinating encephalomyelitis

<400> SEQUENCE: 50

Met Ser Phe Thr Pro Gly Lys Gln Ser Ser Ser Arg Ala Ser Ser Gly
 1 5 10 15

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Asn Arg Ser Gly Asn Gly Ile Leu Lys Trp Ala Asp Gln Ser Asp Gln
 20 25 30
 Ser Arg Asn Val Gln Thr Arg Gly Arg Arg Val Gln Ser Lys Gln Thr
 35 40 45
 Ala Thr Ser Gln Gln Pro Ser Gly Gly Thr Val Val Pro Tyr Tyr Ser
 50 55 60
 Trp Phe Ser Gly Ile Thr Gln Phe Gln Lys Gly Lys Glu Phe Glu Phe
 65 70 75 80
 Ala Glu Gly Gln Gly Val Pro Ile Ala Pro Gly Val Pro Ser Thr Glu
 85 90 95
 Ala Lys Gly Tyr Trp Tyr Arg His Asn Arg Arg Ser Phe Lys Thr Ala
 100 105 110
 Asp Gly Asn Gln Arg Gln Leu Leu Pro Arg Trp Tyr Phe Tyr Tyr Leu
 115 120 125
 Gly Thr Gly Pro His Ala Lys Asp Gln Tyr Gly Thr Asp Ile Asp Gly
 130 135 140
 Val Phe Trp Val Ala Ser Asn Gln Ala Asp Ile Asn Thr Pro Ala Asp
 145 150 155 160
 Ile Val Asp Arg Asp Pro Ser Ser Asp Glu Ala Ile Pro Thr Arg Phe
 165 170 175
 Pro Pro Gly Thr Val Leu Pro Gln Gly Tyr Tyr Ile Glu Gly Ser Gly
 180 185 190
 Arg Ser Ala Pro Asn Ser Arg Ser Thr Ser Arg Ala Pro Asn Arg Ala
 195 200 205
 Pro Ser Ala Gly Ser Arg Ser Arg Ala Asn Ser Gly Asn Arg Thr Ser
 210 215 220
 Thr Pro Gly Val Thr Pro Asp Met Ala Asp Gln Ile Ala Ser Leu Val
 225 230 235 240
 Leu Ala Lys Leu Gly Lys Asp Ala Thr Lys Pro Gln Gln Val Thr Lys
 245 250 255
 Gln Thr Ala Lys Glu Val Arg Gln Lys Ile Leu Asn Lys Pro Arg Gln
 260 265 270
 Lys Arg Ser Pro Asn Lys Gln Cys Thr Val Gln Gln Cys Phe Gly Lys
 275 280 285
 Arg Gly Pro Asn Gln Asn Phe Gly Gly Gly Glu Met Leu Lys Leu Gly
 290 295 300
 Thr Ser Asp Pro Gln Phe Pro Ile Leu Ala Glu Leu Ala Pro Thr Ala
 305 310 315 320
 Gly Ala Phe Phe Phe Gly Ser Arg Leu Glu Leu Ala Lys Val Gln Asn
 325 330 335
 Leu Ser Gly Asn Pro Asp Glu Pro Gln Lys Asp Val Tyr Glu Leu Arg
 340 345 350
 Tyr Asn Gly Ala Ile Arg Phe Asp Ser Thr Leu Ser Gly Phe Glu Thr
 355 360 365
 Ile Met Lys Val Leu Asn Gln Asn Leu Asn Ala Tyr Gln His Gln Glu
 370 375 380
 Asp Gly Met Met Asn Ile Ser Pro Lys Pro Gln Arg Gln Arg Gly Gln
 385 390 395 400
 Lys Asn Gly Gln Val Glu Asn Asp Asn Val Ser Val Ala Ala Pro Lys
 405 410 415
 Ser Arg Val Gln Gln Asn Lys Ser Arg Glu Leu Thr Ala Glu Asp Ile
 420 425 430

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Ser Leu Leu Lys Lys Met Asp Glu Pro Tyr Thr Glu Asp Thr Ser Glu
 435 440 445

Ile

<210> SEQ ID NO 51

<211> LENGTH: 409

<212> TYPE: PRT

<213> ORGANISM: turkey coronavirus

<400> SEQUENCE: 51

Met Ala Ser Gly Lys Ala Thr Gly Lys Thr Asp Ala Pro Ala Pro Ile
 1 5 10 15

Ile Lys Leu Gly Gly Pro Lys Pro Lys Val Gly Ser Ser Gly Asn
 20 25 30

Ala Ser Trp Phe Gln Ser Ile Lys Ala Lys Lys Leu Asn Ser Pro Gln
 35 40 45

Pro Lys Phe Glu Gly Ser Gly Val Pro Asp Asn Glu Asn Ile Lys Thr
 50 55 60

Ser Gln Gln His Gly Tyr Trp Arg Arg Gln Ala Arg Phe Lys Pro Gly
 65 70 75 80

Lys Gly Gly Arg Lys Pro Val Pro Asp Ala Trp Tyr Phe Tyr Tyr Thr
 85 90 95

Gly Thr Gly Pro Ala Ala Asp Leu Asn Trp Gly Asp Thr Gln Asp Gly
 100 105 110

Ile Val Trp Val Ala Ala Lys Gly Ala Asp Val Lys Ser Arg Ser Asn
 115 120 125

Gln Gly Thr Arg Asp Pro Asp Lys Phe Asp Gln Tyr Pro Leu Arg Phe
 130 135 140

Ser Asp Gly Gly Pro Asp Ser Asn Phe Arg Trp Asp Phe Ile Pro Leu
 145 150 155 160

His Arg Gly Arg Ser Gly Arg Ser Thr Ala Ala Ser Ser Ala Ala Ser
 165 170 175

Ser Arg Ala Pro Ser Arg Asp Gly Ser Arg Gly Arg Arg Ser Gly Ser
 180 185 190

Glu Asp Asp Leu Ile Ala Arg Ala Lys Ile Ile Gln Asp Gln Gln
 195 200 205

Lys Lys Gly Ser Arg Ile Thr Lys Ala Lys Ala Asp Glu Met Ala His
 210 215 220

Arg Arg Tyr Cys Lys Arg Thr Val Pro Pro Gly Tyr Lys Val Asp Gln
 225 230 235 240

Val Phe Gly Pro Arg Thr Lys Gly Lys Glu Gly Asn Phe Gly Asp Asp
 245 250 255

Lys Met Asn Glu Gly Ile Lys Asp Gly Arg Val Thr Ala Met Leu
 260 265 270

Asn Leu Val Pro Ser Ser His Ala Cys Leu Phe Gly Ser Arg Val Thr
 275 280 285

Pro Lys Leu Gln Pro Asp Gly Leu His Leu Arg Phe Glu Phe Thr Thr
 290 295 300

Val Val Pro Arg Asp Asp Pro Gln Phe Asp Asn Tyr Val Thr Ile Cys
 305 310 315

Asp Gln Cys Val Asp Gly Ile Gly Thr Arg Pro Lys Asp Asn Glu Pro
 325 330 335

Arg Pro Lys Ser Arg Pro Ser Ser Arg Pro Ala Thr Arg Gly Asn Ser
 340 345 350

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Pro Ala Pro Arg Gln Gln Arg Pro Lys Lys Glu Lys Lys Pro Lys Lys
 355 360 365

Gln Asp Asp Glu Val Asp Lys Ala Leu Thr Ser Asp Glu Glu Arg Asn
 370 375 380

Asn Ala Gln Leu Glu Phe Asp Asp Glu Pro Lys Val Ile Asn Trp Gly
 385 390 395 400

Asp Ser Ala Leu Gly Glu Asn His Leu
 405

<210> SEQ ID NO 52
 <211> LENGTH: 1173
 <212> TYPE: PRT
 <213> ORGANISM: Human coronavirus 229E

<400> SEQUENCE: 52

Met Phe Val Leu Leu Val Ala Tyr Ala Leu Leu His Ile Ala Gly Cys
 1 5 10 15

Gln Thr Thr Asn Gly Leu Asn Thr Ser Tyr Ser Val Cys Asn Gly Cys
 20 25 30

Val Gly Tyr Ser Glu Asn Val Phe Ala Val Glu Ser Gly Gly Tyr Ile
 35 40 45

Pro Ser Asp Phe Ala Phe Asn Asn Trp Phe Leu Leu Thr Asn Thr Ser
 50 55 60

Ser Val Val Asp Gly Val Val Arg Ser Phe Gln Pro Leu Leu Leu Asn
 65 70 75 80

Cys Leu Trp Ser Val Ser Gly Leu Arg Phe Thr Thr Gly Phe Val Tyr
 85 90 95

Phe Asn Gly Thr Gly Arg Gly Asp Cys Lys Gly Phe Ser Ser Asp Val
 100 105 110

Leu Ser Asp Val Ile Arg Tyr Asn Leu Asn Phe Glu Glu Asn Leu Arg
 115 120 125

Arg Gly Thr Ile Leu Phe Lys Thr Ser Tyr Gly Val Val Val Phe Tyr
 130 135 140

Cys Thr Asn Asn Thr Leu Val Ser Gly Asp Ala His Ile Pro Phe Gly
 145 150 155 160

Thr Val Leu Gly Asn Phe Tyr Cys Phe Val Asn Thr Thr Ile Gly Thr
 165 170 175

Glu Thr Thr Ser Ala Phe Val Gly Ala Leu Pro Lys Thr Val Arg Glu
 180 185 190

Phe Val Ile Ser Arg Thr Gly His Phe Tyr Ile Asn Gly Tyr Arg Tyr
 195 200 205

Phe Thr Leu Gly Asn Val Glu Ala Val Asn Phe Asn Val Thr Thr Ala
 210 215 220

Glu Thr Thr Asp Phe Phe Thr Val Ala Leu Ala Ser Tyr Ala Asp Val
 225 230 235 240

Leu Val Asn Val Ser Gln Thr Ser Ile Ala Asn Ile Ile Tyr Cys Asn
 245 250 255

Ser Val Ile Asn Arg Leu Arg Cys Asp Gln Leu Ser Phe Tyr Val Pro
 260 265 270

Asp Gly Phe Tyr Ser Thr Ser Pro Ile Gln Ser Val Glu Leu Pro Val
 275 280 285

Ser Ile Val Ser Leu Pro Val Tyr His Lys His Met Phe Ile Val Leu
 290 295 300

Tyr Val Asp Phe Lys Pro Gln Ser Gly Gly Gly Lys Cys Phe Asn Cys
 305 310 315 320

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Tyr Pro Ala Gly Val Asn Ile Thr Leu Ala Asn Phe Asn Glu Thr Lys
 325 330 335

Gly Pro Leu Cys Val Asp Thr Ser His Phe Thr Thr Lys Tyr Val Ala
 340 345 350

Val Tyr Ala Asn Val Gly Arg Trp Ser Ala Ser Ile Asn Thr Gly Asn
 355 360 365

Cys Pro Phe Ser Phe Gly Lys Val Asn Asn Phe Val Lys Phe Gly Ser
 370 375 380

Val Cys Phe Ser Leu Lys Asp Ile Pro Gly Gly Cys Ala Met Pro Ile
 385 390 395 400

Val Ala Asn Trp Ala Tyr Ser Lys Tyr Tyr Thr Ile Gly Thr Leu Tyr
 405 410 415

Val Ser Trp Ser Asp Gly Asp Gly Ile Thr Gly Val Pro Gln Pro Val
 420 425 430

Glu Gly Val Ser Ser Phe Met Asn Val Thr Leu Asp Lys Cys Thr Lys
 435 440 445

Tyr Asn Ile Tyr Asp Val Ser Gly Val Gly Val Ile Arg Val Ser Asn
 450 455 460

Asp Thr Phe Leu Asn Gly Ile Thr Tyr Thr Ser Thr Ser Gly Asn Leu
 465 470 475 480

Leu Gly Phe Lys Asp Val Thr Lys Gly Thr Ile Tyr Ser Ile Thr Pro
 485 490 495

Cys Asn Pro Pro Asp Gln Leu Val Val Tyr Gln Gln Ala Val Val Gly
 500 505 510

Ala Met Leu Ser Glu Asn Phe Thr Ser Tyr Gly Phe Ser Asn Val Val
 515 520 525

Glu Leu Pro Lys Phe Phe Tyr Ala Ser Asn Gly Thr Tyr Asn Cys Thr
 530 535 540

Asp Ala Val Leu Thr Tyr Ser Ser Phe Gly Val Cys Ala Asp Gly Ser
 545 550 555 560

Ile Ile Ala Val Gln Pro Arg Asn Val Ser Tyr Asp Ser Val Ser Ala
 565 570 575

Ile Val Thr Ala Asn Leu Ser Ile Pro Ser Asn Trp Thr Ile Ser Val
 580 585 590

Gln Val Glu Tyr Leu Gln Ile Thr Ser Thr Pro Ile Val Val Asp Cys
 595 600 605

Ser Thr Tyr Val Cys Asn Gly Asn Val Arg Cys Val Glu Leu Leu Lys
 610 615 620

Gln Tyr Thr Ser Ala Cys Lys Thr Ile Glu Asp Ala Leu Arg Asn Ser
 625 630 635 640

Ala Arg Leu Glu Ser Ala Asp Val Ser Glu Met Leu Thr Phe Asp Lys
 645 650 655

Lys Ala Phe Thr Leu Ala Asn Val Ser Ser Phe Gly Asp Tyr Asn Leu
 660 665 670

Ser Ser Val Ile Pro Ser Leu Pro Thr Ser Gly Ser Arg Val Ala Gly
 675 680 685

Arg Ser Ala Ile Glu Asp Ile Leu Phe Ser Lys Ile Val Thr Ser Gly
 690 695 700

Leu Gly Thr Val Asp Ala Asp Tyr Lys Asn Cys Thr Lys Gly Leu Ser
 705 710 715 720

Ile Ala Asp Leu Ala Cys Ala Gln Tyr Tyr Asn Gly Ile Met Val Leu
 725 730 735

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Pro Gly Val Ala Asp Ala Glu Arg Met Ala Met Tyr Thr Gly Ser Leu
 740 745 750

 Ile Gly Gly Ile Ala Leu Gly Gly Leu Thr Ser Ala Val Ser Ile Pro
 755 760 765

 Phe Ser Leu Ala Ile Gln Ala Arg Leu Asn Tyr Val Ala Leu Gln Thr
 770 775 780

 Asp Val Leu Gln Glu Asn Gln Lys Ile Leu Ala Ala Ser Phe Asn Lys
 785 790 795 800

 Ala Met Thr Asn Ile Val Asp Ala Phe Thr Gly Val Asn Asp Ala Ile
 805 810 815

 Thr Gln Thr Ser Gln Ala Leu Gln Thr Val Ala Thr Ala Leu Asn Lys
 820 825 830

 Ile Gln Asp Val Val Asn Gln Gln Gly Asn Ser Leu Asn His Leu Thr
 835 840 845

 Ser Gln Leu Arg Gln Asn Phe Gln Ala Ile Ser Ser Ser Ile Gln Ala
 850 855 860

 Ile Tyr Asp Arg Leu Asp Thr Ile Gln Ala Asp Gln Gln Val Asp Arg
 865 870 875 880

 Leu Ile Thr Gly Arg Leu Ala Ala Leu Asn Val Phe Val Ser His Thr
 885 890 895

 Leu Thr Lys Tyr Thr Glu Val Arg Ala Ser Arg Gln Leu Ala Gln Gln
 900 905 910

 Lys Val Asn Glu Cys Val Lys Ser Gln Ser Lys Arg Tyr Gly Phe Cys
 915 920 925

 Gly Asn Gly Thr His Ile Phe Ser Ile Val Asn Ala Ala Pro Glu Gly
 930 935 940

 Leu Val Phe Leu His Thr Val Leu Leu Pro Thr Gln Tyr Lys Asp Val
 945 950 955 960

 Glu Ala Trp Ser Gly Leu Cys Val Asp Gly Thr Asn Gly Tyr Val Leu
 965 970 975

 Arg Gln Pro Asn Leu Ala Leu Tyr Lys Glu Gly Asn Tyr Tyr Arg Ile
 980 985 990

 Thr Ser Arg Ile Met Phe Glu Pro Arg Ile Pro Thr Met Ala Asp Phe
 995 1000 1005

 Val Gln Ile Glu Asn Cys Asn Val Thr Phe Val Asn Ile Ser Arg
 1010 1015 1020

 Ser Glu Leu Gln Thr Ile Val Pro Glu Tyr Ile Asp Val Asn Lys
 1025 1030 1035

 Thr Leu Gln Glu Leu Ser Tyr Lys Leu Pro Asn Tyr Thr Val Pro
 1040 1045 1050

 Asp Leu Val Val Glu Gln Tyr Asn Gln Thr Ile Leu Asn Leu Thr
 1055 1060 1065

 Ser Glu Ile Ser Thr Leu Glu Asn Lys Ser Ala Glu Leu Asn Tyr
 1070 1075 1080

 Thr Val Gln Lys Leu Gln Thr Leu Ile Asp Asn Ile Asn Ser Thr
 1085 1090 1095

 Leu Val Asp Leu Lys Trp Leu Asn Arg Val Glu Thr Tyr Ile Lys
 1100 1105 1110

 Trp Pro Trp Trp Val Trp Leu Cys Ile Ser Val Val Leu Ile Phe
 1115 1120 1125

 Val Val Ser Met Leu Leu Leu Cys Cys Cys Ser Thr Gly Cys Cys
 1130 1135 1140

 Gly Phe Phe Ser Cys Phe Ala Ser Ser Ile Arg Gly Cys Cys Glu

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1145 1150 1155
 Ser Thr Lys Leu Pro Tyr Tyr Asp Val Glu Lys Ile His Ile Gln
 1160 1165 1170

<210> SEQ ID NO 53
 <211> LENGTH: 1164
 <212> TYPE: PRT
 <213> ORGANISM: Avian infectious bronchitis virus

<400> SEQUENCE: 53

Met Leu Gly Lys Ser Leu Phe Leu Val Thr Ile Leu Cys Ala Leu Cys
 1 5 10 15

Ser Ala Asn Leu Phe Asp Pro Ala Asn Tyr Val Tyr Tyr Gln Ser
 20 25 30

Ala Phe Arg Pro Ser Asn Gly Trp His Leu Gln Gly Gly Ala Tyr Ala
 35 40 45

Val Val Asn Ser Ser Asn Tyr Ala Asn Asn Ala Gly Ser Ala Ser Glu
 50 55 60

Cys Thr Val Gly Val Ile Lys Asp Val Tyr Asn Gln Ser Ala Ala Ser
 65 70 75 80

Ile Ala Met Thr Ala Pro Leu Gln Gly Met Ala Trp Ser Lys Ser Gln
 85 90 95

Phe Cys Ser Ala His Cys Asp Phe Ser Glu Ile Thr Val Phe Val Thr
 100 105 110

His Cys Tyr Ser Ser Gly Ser Gly Ser Cys Pro Ile Thr Gly Met Ile
 115 120 125

Ala Arg Gly His Ile Arg Ile Ser Ala Met Lys Asn Gly Ser Leu Phe
 130 135 140

Tyr Asn Leu Thr Val Ser Val Ser Lys Tyr Pro Asn Phe Lys Ser Phe
 145 150 155 160

Gln Cys Val Asn Asn Phe Thr Ser Val Tyr Leu Asn Gly Asp Leu Val
 165 170 175

Phe Thr Ser Asn Lys Thr Thr Asp Val Thr Ser Ala Gly Val Tyr Phe
 180 185 190

Lys Ala Gly Gly Pro Val Asn Tyr Ser Ile Met Lys Glu Phe Lys Val
 195 200 205

Leu Ala Tyr Phe Val Asn Gly Thr Ala Gln Asp Val Ile Leu Cys Asp
 210 215 220

Asn Ser Pro Lys Gly Leu Leu Ala Cys Gln Tyr Asn Thr Gly Asn Phe
 225 230 235 240

Ser Asp Gly Phe Tyr Pro Phe Thr Asn Ser Thr Leu Val Arg Glu Lys
 245 250 255

Phe Ile Val Tyr Arg Glu Ser Ser Val Asn Thr Thr Leu Ala Leu Thr
 260 265 270

Asn Phe Thr Phe Thr Asn Val Ser Asn Ala Gln Pro Asn Ser Gly Gly
 275 280 285

Val His Thr Phe His Leu Tyr Gln Thr Gln Thr Ala Gln Ser Gly Tyr
 290 295 300

Tyr Asn Phe Asn Leu Ser Phe Leu Ser Gln Phe Val Tyr Lys Ala Ser
 305 310 315 320

Asp Tyr Met Tyr Gly Ser Tyr His Pro Ile Cys Ala Phe Arg Pro Glu
 325 330 335

Thr Ile Asn Ser Gly Leu Trp Phe Asn Ser Leu Ser Val Ser Leu Thr
 340 345 350

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Tyr	Gly	Pro	Leu	Gln	Gly	Gly	Tyr	Lys	Gln	Ser	Val	Phe	Ser	Gly	Lys
		355					360					365			
Ala	Thr	Cys	Cys	Tyr	Ala	Tyr	Ser	Tyr	Asn	Gly	Pro	Arg	Ala	Cys	Lys
	370					375					380				
Gly	Val	Tyr	Ser	Gly	Glu	Leu	Ser	Arg	Asp	Phe	Glu	Cys	Gly	Leu	Leu
385					390					395					400
Val	Tyr	Val	Thr	Lys	Ser	Asp	Gly	Ser	Arg	Ile	Gln	Thr	Arg	Thr	Glu
				405					410					415	
Pro	Leu	Val	Leu	Thr	Gln	His	Asn	Tyr	Asn	Asn	Ile	Thr	Leu	Asp	Lys
			420					425					430		
Cys	Val	Ala	Tyr	Asn	Ile	Tyr	Gly	Arg	Val	Gly	Gln	Gly	Phe	Ile	Thr
		435					440					445			
Asn	Val	Thr	Asp	Ser	Val	Ala	Asn	Phe	Ser	Tyr	Leu	Ala	Asp	Gly	Gly
	450					455					460				
Leu	Ala	Ile	Leu	Asp	Thr	Ser	Gly	Ala	Ile	Asp	Val	Phe	Val	Val	Gln
465					470					475					480
Gly	Ser	Tyr	Gly	Leu	Asn	Tyr	Tyr	Lys	Val	Asn	Pro	Cys	Glu	Asp	Val
				485					490					495	
Asn	Gln	Gln	Phe	Val	Val	Ser	Gly	Gly	Asn	Ile	Val	Gly	Ile	Leu	Thr
			500					505					510		
Ser	Arg	Asn	Glu	Thr	Gly	Ser	Glu	Gln	Val	Glu	Asn	Gln	Phe	Tyr	Val
		515					520					525			
Lys	Leu	Thr	Asn	Ser	Ser	His	Arg	Arg	Arg	Arg	Ser	Ile	Gly	Gln	Asn
	530					535					540				
Val	Thr	Ser	Cys	Pro	Tyr	Val	Ser	Tyr	Gly	Arg	Phe	Cys	Ile	Glu	Pro
545					550					555					560
Asp	Gly	Ser	Leu	Lys	Met	Ile	Val	Pro	Glu	Glu	Leu	Lys	Gln	Phe	Val
				565					570					575	
Ala	Pro	Leu	Leu	Asn	Ile	Thr	Glu	Ser	Val	Leu	Ile	Pro	Asn	Ser	Phe
			580					585					590		
Asn	Leu	Thr	Val	Thr	Asp	Glu	Tyr	Ile	Gln	Thr	Arg	Met	Asp	Lys	Val
		595					600					605			
Gln	Ile	Asn	Cys	Leu	Gln	Tyr	Val	Cys	Gly	Asn	Ser	Leu	Glu	Cys	Arg
	610					615					620				
Lys	Leu	Phe	Gln	Gln	Tyr	Gly	Pro	Val	Cys	Asp	Asn	Ile	Leu	Ser	Val
625					630					635					640
Val	Asn	Ser	Val	Ser	Gln	Lys	Glu	Asp	Met	Glu	Leu	Leu	Ser	Phe	Tyr
				645					650					655	
Ser	Ser	Thr	Lys	Pro	Lys	Gly	Tyr	Asp	Thr	Pro	Val	Leu	Ser	Asn	Val
			660					665					670		
Ser	Thr	Gly	Glu	Phe	Asn	Ile	Ser	Leu	Leu	Leu	Thr	Pro	Pro	Ser	Ser
		675					680					685			
Pro	Ser	Gly	Arg	Ser	Phe	Val	Glu	Asp	Leu	Leu	Phe	Thr	Ser	Val	Glu
		690				695					700				
Thr	Val	Gly	Leu	Pro	Thr	Asp	Ala	Glu	Tyr	Lys	Lys	Cys	Thr	Ala	Gly
705					710					715					720
Pro	Leu	Gly	Thr	Leu	Lys	Asp	Leu	Ile	Cys	Ala	Arg	Glu	Tyr	Asn	Gly
				725					730					735	
Leu	Leu	Val	Leu	Pro	Pro	Ile	Ile	Thr	Ala	Asp	Met	Gln	Thr	Met	Tyr
			740					745					750		
Thr	Ala	Ser	Leu	Val	Gly	Ala	Met	Ala	Phe	Gly	Gly	Ile	Thr	Ser	Ala
		755					760					765			
Ala	Ala	Ile	Pro	Phe	Ala	Thr	Gln	Ile	Gln	Ala	Arg	Ile	Asn	His	Leu

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770	775	780
Gly Ile Ala Gln Ser Leu Leu Met Lys Asn Gln Glu Lys Ile Ala Ala 785 790 795 800		
Ser Phe Asn Lys Ala Ile Gly His Met Gln Glu Gly Phe Arg Ser Thr 805 810 815		
Ser Leu Ala Leu Gln Gln Val Gln Asp Val Val Asn Lys Gln Ser Ala 820 825 830		
Ile Leu Thr Glu Thr Met Asn Ser Leu Asn Lys Asn Phe Gly Ala Ile 835 840 845		
Ser Ser Val Ile Gln Asp Ile Tyr Ala Gln Leu Asp Ala Ile Gln Ala 850 855 860		
Asp Ala Gln Val Asp Arg Leu Ile Thr Gly Arg Leu Ser Ser Leu Ser 865 870 875 880		
Val Leu Ala Ser Ala Lys Gln Ser Glu Tyr Ile Arg Val Ser Gln Gln 885 890 895		
Arg Glu Leu Ala Thr Gln Lys Ile Asn Glu Cys Val Lys Ser Gln Ser 900 905 910		
Asn Arg Tyr Gly Phe Cys Gly Ser Gly Arg His Val Leu Ser Ile Pro 915 920 925		
Gln Asn Ala Pro Asn Gly Ile Val Phe Ile His Phe Thr Tyr Thr Pro 930 935 940		
Glu Thr Phe Val Asn Val Thr Ala Ile Val Gly Phe Cys Val Asn Pro 945 950 955 960		
Leu Asn Ala Ser Gln Tyr Ala Ile Val Pro Ala Asn Gly Arg Gly Ile 965 970 975		
Phe Ile Gln Val Asn Gly Thr Tyr Tyr Ile Thr Ser Arg Asp Met Tyr 980 985 990		
Met Pro Arg Asp Ile Thr Ala Gly Asp Ile Val Thr Leu Thr Ser Cys 995 1000 1005		
Gln Ala Asn Tyr Val Asn Val Asn Lys Thr Val Ile Thr Thr Phe 1010 1015 1020		
Val Glu Asp Asp Asp Phe Asn Phe Asp Asp Glu Leu Ser Lys Trp 1025 1030 1035		
Trp Asn Asp Thr Lys His Gly Leu Pro Asp Phe Asp Asp Phe Asn 1040 1045 1050		
Tyr Thr Val Pro Ile Leu Asn Ile Ser Gly Glu Ile Asp Asn Ile 1055 1060 1065		
Gln Gly Val Ile Gln Gly Leu Asn Asp Ser Leu Ile Asn Leu Glu 1070 1075 1080		
Glu Leu Ser Ile Ile Lys Thr Tyr Ile Lys Trp Pro Trp Tyr Val 1085 1090 1095		
Trp Leu Ala Ile Gly Phe Ala Ile Ile Ile Phe Ile Leu Ile Leu 1100 1105 1110		
Gly Trp Val Phe Phe Met Thr Gly Cys Cys Gly Cys Cys Cys Gly 1115 1120 1125		
Cys Phe Gly Ile Ile Pro Leu Ile Ser Lys Cys Gly Lys Lys Ser 1130 1135 1140		
Ser Tyr Tyr Thr Thr Phe Asp Asn Asp Val Val Thr Glu Gln Tyr 1145 1150 1155		
Arg Pro Lys Lys Ser Val 1160		

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<211> LENGTH: 1363
<212> TYPE: PRT
<213> ORGANISM: Bovine coronavirus

<400> SEQUENCE: 54

Met Phe Leu Ile Leu Leu Ile Ser Leu Pro Met Ala Phe Ala Val Ile
 1           5           10          15

Gly Asp Leu Lys Cys Thr Thr Val Ser Ile Asn Asp Val Asp Thr Gly
      20           25           30

Ala Pro Ser Ile Ser Thr Asp Ile Val Asp Val Thr Asn Gly Leu Gly
      35           40           45

Thr Tyr Tyr Val Leu Asp Arg Val Tyr Leu Asn Thr Thr Leu Leu Leu
 50           55           60

Asn Gly Tyr Tyr Pro Thr Ser Gly Ser Thr Tyr Arg Asn Met Ala Leu
65           70           75           80

Lys Gly Thr Leu Leu Leu Ser Arg Leu Trp Phe Lys Pro Pro Phe Leu
      85           90           95

Ser Asp Phe Ile Asn Gly Ile Phe Ala Lys Val Lys Asn Thr Lys Val
      100          105          110

Ile Lys Lys Gly Val Met Tyr Ser Glu Phe Pro Ala Ile Thr Ile Gly
      115          120          125

Ser Thr Phe Val Asn Thr Ser Tyr Ser Val Val Val Gln Pro His Thr
      130          135          140

Thr Asn Leu Asp Asn Lys Leu Gln Gly Leu Leu Glu Ile Ser Val Cys
145          150          155          160

Gln Tyr Thr Met Cys Glu Tyr Pro His Thr Ile Cys His Pro Lys Leu
      165          170          175

Gly Asn Lys Arg Val Glu Leu Trp His Trp Asp Thr Gly Val Val Ser
      180          185          190

Cys Leu Tyr Lys Arg Asn Phe Thr Tyr Asp Val Asn Ala Asp Tyr Leu
      195          200          205

Tyr Phe His Phe Tyr Gln Glu Gly Gly Thr Phe Tyr Ala Tyr Phe Thr
      210          215          220

Asp Thr Gly Val Val Thr Lys Phe Leu Phe Asn Val Tyr Leu Gly Thr
225          230          235          240

Val Leu Ser His Tyr Tyr Val Leu Pro Leu Thr Cys Ser Ser Ala Met
      245          250          255

Thr Leu Glu Tyr Trp Val Thr Pro Leu Thr Ser Lys Gln Tyr Leu Leu
      260          265          270

Ala Phe Asn Gln Asp Gly Val Ile Phe Asn Ala Val Asp Cys Lys Ser
      275          280          285

Asp Phe Met Ser Glu Ile Lys Cys Lys Thr Leu Ser Ile Ala Pro Ser
      290          295          300

Thr Gly Val Tyr Glu Leu Asn Gly Tyr Thr Val Gln Pro Ile Ala Asp
305          310          315          320

Val Tyr Arg Arg Ile Pro Asn Leu Pro Asp Cys Asn Ile Glu Ala Trp
      325          330          335

Leu Asn Asp Lys Ser Val Pro Ser Pro Leu Asn Trp Glu Arg Lys Thr
      340          345          350

Phe Ser Asn Cys Asn Phe Asn Met Ser Ser Leu Met Ser Phe Ile Gln
      355          360          365

Ala Asp Ser Phe Thr Cys Asn Asn Ile Asp Ala Ala Lys Ile Tyr Gly
      370          375          380

Met Cys Phe Ser Ser Ile Thr Ile Asp Lys Phe Ala Ile Pro Asn Gly

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385	390					395					400				
Arg Lys Val Asp	Leu	Gln	Leu	Gly	Asn	Leu	Gly	Tyr	Leu	Gln	Ser	Phe			
	405						410				415				
Asn Tyr Arg Ile Asp	Thr	Thr	Ala	Thr	Ser	Cys	Gln	Leu	Tyr	Tyr	Asn				
	420				425					430					
Leu Pro Ala Ala Asn	Val	Ser	Val	Ser	Arg	Phe	Asn	Pro	Ser	Thr	Trp				
	435					440				445					
Asn Arg Arg Phe Gly	Phe	Thr	Glu	Gln	Phe	Val	Phe	Lys	Pro	Gln	Pro				
	450				455			460							
Val Gly Val Phe Thr	His	His	Asp	Val	Val	Tyr	Ala	Gln	His	Cys	Phe				
	465				470			475			480				
Lys Ala Pro Lys Asn	Phe	Cys	Pro	Cys	Lys	Leu	Asp	Gly	Ser	Leu	Cys				
		485				490					495				
Val Gly Asn Gly Pro	Gly	Ile	Asp	Ala	Gly	Tyr	Lys	Asn	Ser	Gly	Ile				
		500			505					510					
Gly Thr Cys Pro Ala	Gly	Thr	Asn	Tyr	Leu	Thr	Cys	His	Asn	Ala	Ala				
		515				520			525						
Gln Cys Asp Cys Leu	Cys	Thr	Pro	Asp	Pro	Ile	Thr	Ser	Lys	Ser	Thr				
		530				535			540						
Gly Pro Tyr Lys Cys	Pro	Gln	Thr	Lys	Tyr	Leu	Val	Gly	Ile	Gly	Glu				
		545			550			555			560				
His Cys Ser Gly Leu	Ala	Ile	Lys	Ser	Asp	Tyr	Cys	Gly	Gly	Asn	Pro				
		565			570					575					
Cys Thr Cys Gln Pro	Gln	Ala	Phe	Leu	Gly	Trp	Ser	Val	Asp	Ser	Cys				
		580				585				590					
Leu Gln Gly Asp Arg	Cys	Asn	Ile	Phe	Ala	Asn	Phe	Ile	Phe	His	Asp				
		595				600			605						
Val Asn Ser Gly Thr	Thr	Cys	Ser	Thr	Asp	Leu	Gln	Lys	Ser	Asn	Thr				
		610				615			620						
Asp Ile Ile Leu Gly	Val	Cys	Val	Asn	Tyr	Asp	Leu	Tyr	Gly	Ile	Thr				
		625			630			635			640				
Gly Gln Gly Ile Phe	Val	Glu	Val	Asn	Ala	Thr	Tyr	Tyr	Asn	Ser	Trp				
		645				650					655				
Gln Asn Leu Leu Tyr	Asp	Ser	Asn	Gly	Asn	Leu	Tyr	Gly	Phe	Arg	Asp				
		660				665				670					
Tyr Leu Thr Asn Arg	Thr	Phe	Met	Ile	Arg	Ser	Cys	Tyr	Ser	Gly	Arg				
		675			680				685						
Val Ser Ala Ala Phe	His	Ala	Asn	Ser	Ser	Glu	Pro	Ala	Leu	Leu	Phe				
		690				695			700						
Arg Asn Ile Lys Cys	Asn	Tyr	Val	Phe	Asn	Asn	Thr	Leu	Ser	Arg	Gln				
		705			710			715			720				
Leu Gln Pro Ile Asn	Tyr	Phe	Asp	Ser	Tyr	Leu	Gly	Cys	Val	Val	Asn				
		725				730					735				
Ala Asp Asn Ser Thr	Ser	Ser	Val	Val	Gln	Thr	Cys	Asp	Leu	Thr	Val				
		740				745				750					
Gly Ser Gly Tyr Cys	Val	Asp	Tyr	Ser	Thr	Lys	Arg	Arg	Ser	Arg	Arg				
		755				760			765						
Ala Ile Thr Thr Gly	Tyr	Arg	Phe	Thr	Asn	Phe	Glu	Pro	Phe	Thr	Val				
		770				775			780						
Asn Ser Val Asn Asp	Ser	Leu	Glu	Pro	Val	Gly	Gly	Leu	Tyr	Glu	Ile				
		785			790			795			800				
Gln Ile Pro Ser Glu	Phe	Thr	Ile	Gly	Asn	Met	Glu	Glu	Phe	Ile	Gln				
		805				810					815				

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Thr Ser Ser Pro Lys Val Thr Ile Asp Cys Ser Ala Phe Val Cys Gly
 820 825 830
 Asp Tyr Ala Ala Cys Lys Ser Gln Leu Val Glu Tyr Gly Ser Phe Cys
 835 840 845
 Asp Asn Ile Asn Ala Ile Leu Thr Glu Val Asn Glu Leu Leu Asp Thr
 850 855 860
 Thr Gln Leu Gln Val Ala Asn Ser Leu Met Asn Gly Val Thr Leu Ser
 865 870 875 880
 Thr Lys Leu Lys Asp Gly Val Asn Phe Asn Val Asp Asp Ile Asn Phe
 885 890 895
 Ser Pro Val Leu Gly Cys Leu Gly Ser Ala Cys Asn Lys Val Ser Ser
 900 905 910
 Arg Ser Ala Ile Glu Asp Leu Leu Phe Ser Lys Val Lys Leu Ser Asp
 915 920 925
 Val Gly Phe Val Glu Ala Tyr Asn Asn Cys Thr Gly Gly Ala Glu Ile
 930 935 940
 Arg Asp Leu Ile Cys Val Gln Ser Tyr Asn Gly Ile Lys Val Leu Pro
 945 950 955 960
 Pro Leu Leu Ser Val Asn Gln Ile Ser Gly Tyr Thr Leu Ala Ala Thr
 965 970 975
 Ser Ala Ser Leu Phe Pro Pro Leu Ser Ala Ala Val Gly Val Pro Phe
 980 985 990
 Tyr Leu Asn Val Gln Tyr Arg Ile Asn Gly Ile Gly Val Thr Met Asp
 995 1000 1005
 Val Leu Ser Gln Asn Gln Lys Leu Ile Ala Asn Ala Phe Asn Asn
 1010 1015 1020
 Ala Leu Asp Ala Ile Gln Glu Gly Phe Asp Ala Thr Asn Ser Ala
 1025 1030 1035
 Leu Val Lys Ile Gln Ala Val Val Asn Ala Asn Ala Glu Ala Leu
 1040 1045 1050
 Asn Asn Leu Leu Gln Gln Leu Ser Asn Arg Phe Gly Ala Ile Ser
 1055 1060 1065
 Ser Ser Leu Gln Glu Ile Leu Ser Arg Leu Asp Ala Leu Glu Ala
 1070 1075 1080
 Gln Ala Gln Ile Asp Arg Leu Ile Asn Gly Arg Leu Thr Ala Leu
 1085 1090 1095
 Asn Val Tyr Val Ser Gln Gln Leu Ser Asp Ser Thr Leu Val Lys
 1100 1105 1110
 Phe Ser Ala Ala Gln Ala Met Glu Lys Val Asn Glu Cys Val Lys
 1115 1120 1125
 Ser Gln Ser Ser Arg Ile Asn Phe Cys Gly Asn Gly Asn His Ile
 1130 1135 1140
 Ile Ser Leu Val Gln Asn Ala Pro Tyr Gly Leu Tyr Phe Ile His
 1145 1150 1155
 Phe Ser Tyr Val Pro Thr Lys Tyr Val Thr Ala Lys Val Ser Pro
 1160 1165 1170
 Gly Leu Cys Ile Ala Gly Asp Arg Gly Ile Ala Pro Lys Ser Gly
 1175 1180 1185
 Tyr Phe Val Asn Val Asn Asn Thr Trp Met Phe Thr Gly Ser Gly
 1190 1195 1200
 Tyr Tyr Tyr Pro Glu Pro Ile Thr Gly Asn Asn Val Val Val Met
 1205 1210 1215

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Ser Thr Cys Ala Val Asn Tyr Thr Lys Ala Pro Asp Val Met Leu
1220                               1225                1230

Asn Ile Ser Thr Pro Asn Leu His Asp Phe Lys Glu Glu Leu Asp
1235                               1240                1245

Gln Trp Phe Lys Asn Gln Thr Ser Val Ala Pro Asp Leu Ser Leu
1250                               1255                1260

Asp Tyr Ile Asn Val Thr Phe Leu Asp Leu Gln Asp Glu Met Asn
1265                               1270                1275

Arg Leu Gln Glu Ala Ile Lys Val Leu Asn Gln Ser Tyr Ile Asn
1280                               1285                1290

Leu Lys Asp Ile Gly Thr Tyr Glu Tyr Tyr Val Lys Trp Pro Trp
1295                               1300                1305

Tyr Val Trp Leu Leu Ile Gly Phe Ala Gly Val Ala Met Leu Val
1310                               1315                1320

Leu Leu Phe Phe Ile Cys Cys Cys Thr Gly Cys Gly Thr Ser Cys
1325                               1330                1335

Phe Lys Ile Cys Gly Gly Cys Cys Asp Asp Tyr Thr Gly His Gln
1340                               1345                1350

Glu Leu Val Ile Lys Thr Ser His Asp Asp
1355                               1360

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<210> SEQ ID NO 55
<211> LENGTH: 1453
<212> TYPE: PRT
<213> ORGANISM: canine coronavirus

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<400> SEQUENCE: 55

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Met Ile Val Leu Ile Leu Cys Leu Leu Leu Phe Ser Tyr Asn Ser Val
1           5           10           15

Ile Cys Thr Ser Asn Asn Asp Cys Val Gln Gly Asn Val Thr Gln Leu
20           25           30

Pro Gly Asn Glu Asn Ile Ile Lys Asp Phe Leu Phe His Thr Phe Lys
35           40           45

Glu Glu Pro Ser Val Val Val Gly Gly Tyr Tyr Pro Thr Glu Val Trp
50           55           60

Tyr Asn Cys Ser Arg Ser Ala Thr Thr Thr Ala Tyr Lys Asp Phe Ser
65           70           75           80

Asn Ile His Ala Phe Tyr Phe Asp Met Glu Ala Met Glu Asn Ser Thr
85           90           95

Gly Asn Ala Arg Gly Lys Pro Leu Leu Val His Val His Gly Asp Pro
100          105          110

Val Ser Ile Ile Ile Tyr Ile Ser Ala Tyr Arg Asp Asp Val Gln Pro
115          120          125

Arg Pro Leu Leu Lys His Gly Leu Leu Cys Ile Thr Lys Asn Lys Ile
130          135          140

Ile Asp Tyr Asn Thr Phe Thr Ser Ala Gln Trp Ser Ala Ile Cys Leu
145          150          155          160

Gly Asp Asp Arg Lys Ile Pro Phe Ser Val Ile Pro Thr Asp Asn Gly
165          170          175

Thr Lys Ile Phe Gly Leu Glu Trp Asn Asp Asp Tyr Val Thr Ala Tyr
180          185          190

Ile Ser Asp Arg Ser His His Leu Asn Ile Asn Asn Asn Trp Phe Asn
195          200          205

Asn Val Thr Ile Leu Tyr Ser Arg Ser Ser Ser Ala Thr Trp Gln Lys
210          215          220

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Ser Ala Ala Tyr Val Tyr Gln Gly Val Ser Asn Phe Thr Tyr Tyr Lys
 225 230 235 240
 Leu Asn Asn Thr Asn Gly Leu Lys Ser Tyr Glu Leu Cys Glu Asp Tyr
 245 250 255
 Glu Tyr Cys Thr Gly Tyr Ala Thr Asn Val Phe Ala Pro Thr Val Gly
 260 265 270
 Gly Tyr Ile Pro His Gly Phe Ser Phe Asn Asn Trp Phe Met Arg Thr
 275 280 285
 Asn Ser Ser Thr Phe Val Ser Gly Arg Phe Val Thr Asn Gln Pro Leu
 290 295 300
 Leu Val Asn Cys Leu Trp Pro Val Pro Ser Phe Gly Val Ala Ala Gln
 305 310 315 320
 Gln Phe Cys Phe Glu Gly Ala Gln Phe Ser Gln Cys Asn Gly Val Ser
 325 330 335
 Leu Asn Asn Thr Val Asp Val Ile Arg Phe Asn Leu Asn Phe Thr Ala
 340 345 350
 Leu Val Gln Ser Gly Met Gly Ala Thr Val Phe Ser Leu Asn Thr Thr
 355 360 365
 Gly Gly Val Ile Leu Glu Ile Ser Cys Tyr Asn Asp Thr Val Ser Glu
 370 375 380
 Ser Ser Phe Tyr Ser Tyr Gly Glu Ile Ser Phe Gly Val Thr Asp Gly
 385 390 395 400
 Pro Arg Tyr Cys Phe Ala Leu Tyr Asn Gly Thr Ala Leu Lys Tyr Leu
 405 410 415
 Gly Thr Leu Pro Ser Val Lys Glu Ile Ala Ile Ser Lys Trp Gly
 420 425 430
 His Phe Tyr Ile Asn Gly Tyr Asn Phe Phe Ser Thr Phe Pro Ile Asp
 435 440 445
 Cys Ile Ser Phe Asn Leu Thr Thr Gly Asp Ser Gly Ala Phe Trp Thr
 450 455 460
 Ile Ala Tyr Thr Ser Tyr Thr Asp Ala Leu Val Gln Val Glu Asn Thr
 465 470 475 480
 Ala Ile Lys Lys Val Thr Tyr Cys Asn Ser His Ile Asn Asn Ile Lys
 485 490 495
 Cys Ser Gln Leu Thr Ala Asn Leu Gln Asn Gly Phe Tyr Pro Val Ala
 500 505 510
 Ser Ser Glu Val Gly Leu Val Asn Lys Ser Val Val Leu Leu Pro Ser
 515 520 525
 Phe Tyr Ser His Thr Ser Val Asn Ile Thr Ile Asp Leu Gly Met Lys
 530 535 540
 Arg Ser Gly Tyr Gly Gln Pro Ile Ala Ser Thr Leu Ser Asn Ile Thr
 545 550 555 560
 Leu Pro Met Gln Asp Asn Asn Thr Asp Val Tyr Cys Ile Arg Ser Asn
 565 570 575
 Arg Phe Ser Val Tyr Phe His Ser Thr Cys Lys Ser Ser Leu Trp Asp
 580 585 590
 Asp Val Phe Asn Ser Asp Cys Thr Asp Val Leu Tyr Ala Thr Ala Val
 595 600 605
 Ile Lys Thr Gly Thr Cys Pro Phe Ser Phe Asp Lys Leu Asn Asn Tyr
 610 615 620
 Leu Thr Phe Asn Lys Phe Cys Leu Ser Leu Asn Pro Val Gly Ala Asn
 625 630 635 640

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Cys Lys Phe Asp Val Ala Ala Arg Thr Arg Thr Asn Glu Gln Val Val
 645 650 655
 Arg Ser Leu Tyr Val Ile Tyr Glu Glu Gly Asp Asn Ile Val Gly Val
 660 665 670
 Pro Ser Asp Asn Ser Gly Leu His Asp Leu Ser Val Leu His Leu Asp
 675 680 685
 Ser Cys Thr Asp Tyr Asn Ile Tyr Gly Ile Thr Gly Val Gly Ile Ile
 690 695 700
 Arg Gln Thr Asn Ser Thr Leu Leu Ser Gly Leu Tyr Tyr Thr Ser Leu
 705 710 715 720
 Ser Gly Asp Leu Leu Gly Phe Lys Asn Val Ser Asp Gly Val Ile Tyr
 725 730 735
 Ser Val Thr Pro Cys Asp Val Ser Ala His Ala Ala Val Ile Asp Gly
 740 745 750
 Ala Ile Val Gly Ala Met Thr Ser Ile Asn Ser Glu Leu Leu Gly Leu
 755 760 765
 Thr His Trp Thr Thr Thr Pro Asn Phe Tyr Tyr Tyr Ser Ile Tyr Asn
 770 775 780
 Tyr Thr Asn Glu Arg Thr Arg Gly Thr Ala Ile Asp Ser Asn Asp Val
 785 790 795 800
 Asp Cys Glu Pro Ile Ile Thr Tyr Ser Asn Ile Gly Val Cys Lys Asn
 805 810 815
 Gly Ala Leu Val Phe Ile Asn Val Thr His Ser Asp Gly Asp Val Gln
 820 825 830
 Pro Ile Ser Thr Gly Asn Val Thr Ile Pro Thr Asn Phe Thr Ile Ser
 835 840 845
 Val Gln Val Glu Tyr Ile Gln Val Tyr Thr Thr Pro Val Ser Ile Asp
 850 855 860
 Cys Ser Arg Tyr Val Cys Asn Gly Asn Pro Arg Cys Asn Lys Leu Leu
 865 870 875 880
 Thr Gln Tyr Val Ser Ala Cys Gln Thr Ile Glu Gln Ala Leu Ala Met
 885 890 895
 Gly Ala Arg Leu Glu Asn Met Glu Ile Asp Ser Met Leu Phe Val Ser
 900 905 910
 Glu Asn Ala Leu Lys Leu Ala Ser Val Glu Ala Phe Asn Ser Thr Glu
 915 920 925
 Thr Leu Asp Pro Ile Tyr Lys Glu Trp Pro Asn Ile Gly Gly Ser Trp
 930 935 940
 Leu Gly Gly Leu Lys Asp Ile Leu Pro Ser His Asn Ser Lys Arg Lys
 945 950 955 960
 Tyr Arg Ser Ala Ile Glu Asp Leu Leu Phe Asp Lys Val Val Thr Ser
 965 970 975
 Gly Leu Gly Thr Val Asp Glu Asp Tyr Lys Arg Cys Thr Gly Gly Tyr
 980 985 990
 Asp Ile Ala Asp Leu Val Cys Ala Gln Tyr Tyr Asn Gly Ile Met Val
 995 1000 1005
 Leu Pro Gly Val Ala Asn Asp Asp Lys Met Ala Met Tyr Thr Ala
 1010 1015 1020
 Ser Leu Ala Gly Gly Ile Thr Leu Gly Ser Leu Gly Gly Gly Ala
 1025 1030 1035
 Val Ser Ile Pro Phe Ala Ile Ala Val Gln Ala Arg Leu Asn Tyr
 1040 1045 1050
 Val Ala Leu Gln Thr Asp Val Leu Asn Lys Asn Gln Gln Ile Leu

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1055	1060	1065
Ala Asn 1070	Ala Phe Asn Gln 1075	Ile Gly Asn Ile Thr 1080
Gly Lys 1085	Val Asn Asp Ala 1090	Ile His Gln Thr Ser 1095
Thr Val 1100	Ala Lys Val Leu 1105	Ala Lys Val Gln Asp 1110
Gln Gly 1115	Gln Ala Leu Ser 1120	His Leu Thr Leu Gln 1125
Phe Gln 1130	Ala Ile Ser Ser 1135	Ser Ile Ser Asp Ile 1140
Asp Glu 1145	Leu Ser Ala Asp 1150	Ala Gln Val Asp Arg 1155
Arg Leu 1160	Thr Ala Leu Asn 1165	Ala Phe Val Ser Gln 1170
Gln Ala 1175	Glu Val Arg Ala 1180	Ser Arg Gln Leu Ala 1185
Asn Glu 1190	Cys Val Arg Ser 1195	Gln Ser Gln Arg Phe 1200
Asn Gly 1205	Thr His Leu Phe 1210	Ser Leu Ala Asn Ala 1215
Met Ile 1220	Phe Phe His Thr 1225	Val Leu Leu Pro Thr 1230
Val Thr 1235	Ala Trp Ser Gly 1240	Ile Cys Ala Ser Asp 1245
Phe Gly 1250	Leu Val Val Lys 1255	Asp Val Gln Leu Thr 1260
Leu Asp 1265	Asp Lys Phe Tyr 1270	Leu Thr Pro Arg Thr 1275
Ile Val 1280	Ala Thr Ser Ser 1285	Asp Phe Val Gln Ile 1290
Val Leu 1295	Phe Val Asn Ala 1300	Thr Val Ile Asp Leu 1305
Pro Asp 1310	Tyr Ile Asp Ile 1315	Asn Gln Thr Val Gln 1320
Asn Phe 1325	Arg Pro Asn Trp 1330	Thr Val Pro Glu Leu 1335
Phe Asn 1340	Ala Thr Tyr Leu 1345	Asn Leu Thr Gly Glu 1350
Glu Phe 1355	Arg Ser Glu Lys 1360	Leu His Asn Thr Thr 1365
Ile Leu 1370	Ile Asp Asn Ile 1375	Asn Asn Thr Leu Val 1380
Leu Asn 1385	Arg Ile Glu Thr 1390	Tyr Val Lys Trp Pro 1395
Leu Leu 1400	Ile Gly Leu Val 1405	Val Ile Phe Cys Ile 1410
Phe Cys 1415	Cys Cys Ser Thr 1420	Gly Cys Cys Gly Cys 1425
Gly Ser 1430	Cys Cys His Ser 1435	Ile Cys Ser Arg Arg 1440
Tyr Glu 1445	Pro Ile Glu Lys 1450	Val His Val His

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<210> SEQ ID NO 56
<211> LENGTH: 1464
<212> TYPE: PRT
<213> ORGANISM: Feline infectious peritonitis virus

<400> SEQUENCE: 56
Met Ile Phe Ile Ile Leu Thr Leu Leu Ser Val Ala Lys Ser Glu Asp
 1           5           10           15
Ala Pro His Gly Val Thr Leu Pro Gln Phe Asn Thr Ser His Asn Asn
 20           25           30
Glu Arg Phe Glu Leu Asn Phe Tyr Asn Phe Leu Gln Thr Trp Asp Ile
 35           40           45
Pro Pro Asn Thr Glu Thr Ile Leu Gly Gly Tyr Leu Pro Tyr Cys Gly
 50           55           60
Ala Gly Val Asn Cys Gly Trp Tyr Asn Phe Ser Gln Ser Val Gly Gln
 65           70           75           80
Asn Gly Lys Tyr Ala Tyr Ile Asn Thr Gln Asn Leu Asn Ile Pro Asn
 85           90           95
Val His Gly Val Tyr Phe Asp Val Arg Glu His Asn Asn Asp Gly Glu
 100          105          110
Trp Asp Asp Arg Asp Lys Val Gly Leu Leu Ile Ala Ile His Gly Asn
 115          120          125
Ser Lys Tyr Ser Leu Leu Met Val Leu Gln Asp Ala Val Glu Ala Asn
 130          135          140
Gln Pro His Val Ala Val Lys Ile Cys His Trp Lys Pro Gly Asn Ile
 145          150          155          160
Ser Ser Tyr His Ala Phe Ser Val Asn Leu Gly Asp Gly Gly Gln Cys
 165          170          175
Val Phe Asn Gln Arg Phe Ser Leu Asp Thr Val Leu Thr Thr Asn Asp
 180          185          190
Phe Tyr Gly Phe Gln Trp Thr Asp Thr Tyr Val Asp Ile Tyr Leu Gly
 195          200          205
Gly Thr Ile Thr Lys Val Trp Val Asp Asn Asp Trp Ser Ile Val Glu
 210          215          220
Ala Ser Ile Ser Tyr His Trp Asn Arg Ile Asn Tyr Gly Tyr Tyr Met
 225          230          235          240
Gln Phe Val Asn Arg Thr Thr Tyr Tyr Ala Tyr Asn Asn Thr Gly Gly
 245          250          255
Ala Asn Tyr Thr Gln Leu Gln Leu Ser Glu Cys His Thr Asp Tyr Cys
 260          265          270
Ala Gly Tyr Ala Lys Asn Val Phe Val Pro Ile Asp Gly Lys Ile Pro
 275          280          285
Glu Asp Phe Ser Phe Ser Asn Trp Phe Leu Leu Ser Asp Lys Ser Thr
 290          295          300
Leu Val Gln Gly Arg Val Leu Ser Ser Gln Pro Val Phe Val Gln Cys
 305          310          315          320
Leu Arg Pro Val Pro Ser Trp Ser Asn Asn Thr Ala Val Val His Phe
 325          330          335
Lys Asn Asp Ala Phe Cys Pro Asn Val Thr Ala Asp Val Leu Arg Phe
 340          345          350
Asn Leu Asn Phe Ser Asp Thr Asp Val Tyr Thr Asp Ser Thr Asn Asp
 355          360          365
Glu Gln Leu Phe Phe Thr Phe Glu Asp Asn Thr Thr Ala Ser Ile Ala

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370					375					380				
Cys	Tyr	Ser	Ser	Ala	Asn	Val	Thr	Asp	Phe	Gln	Pro	Ala	Asn	Ser
385					390					395				400
Val	Ser	His	Ile	Pro	Phe	Gly	Lys	Thr	Ala	His	Phe	Cys	Phe	Ala
				405					410					415
Phe	Ser	His	Ser	Ile	Val	Ser	Arg	Gln	Phe	Leu	Gly	Ile	Leu	Pro
			420					425					430	Pro
Thr	Val	Arg	Glu	Phe	Ala	Phe	Gly	Arg	Asp	Gly	Ser	Ile	Phe	Val
		435					440					445		Asn
Gly	Tyr	Lys	Tyr	Phe	Ser	Leu	Pro	Ala	Ile	Arg	Ser	Val	Asn	Phe
	450					455					460			Ser
Ile	Ser	Ser	Val	Glu	Glu	Tyr	Gly	Phe	Trp	Thr	Ile	Ala	Tyr	Thr
465					470					475				480
Tyr	Thr	Asp	Val	Met	Val	Asp	Val	Asn	Gly	Thr	Ala	Ile	Thr	Arg
				485					490					495
Phe	Tyr	Cys	Asp	Ser	Pro	Leu	Asn	Arg	Ile	Lys	Cys	Gln	Gln	Leu
			500					505					510	Lys
His	Glu	Leu	Pro	Asp	Gly	Phe	Tyr	Ser	Ala	Ser	Met	Leu	Val	Lys
		515					520					525		Lys
Asp	Leu	Pro	Lys	Thr	Phe	Val	Thr	Met	Pro	Gln	Phe	Tyr	His	Trp
	530					535					540			Met
Asn	Val	Thr	Leu	His	Val	Val	Leu	Asn	Asp	Thr	Glu	Lys	Lys	Tyr
545					550					555				560
Ile	Ile	Leu	Ala	Lys	Ala	Pro	Glu	Leu	Ala	Ala	Leu	Ala	Asp	Val
				565					570					575
Phe	Glu	Ile	Ala	Gln	Ala	Asn	Gly	Ser	Val	Thr	Asn	Val	Thr	Ser
			580					585					590	Leu
Cys	Val	Gln	Ala	Arg	Gln	Leu	Ala	Leu	Phe	Tyr	Lys	Tyr	Thr	Ser
		595					600					605		Leu
Gln	Gly	Leu	Tyr	Thr	Tyr	Ser	Asn	Leu	Val	Glu	Leu	Gln	Asn	Tyr
	610					615					620			Asp
Cys	Pro	Phe	Ser	Pro	Gln	Gln	Phe	Asn	Asn	Tyr	Leu	Gln	Phe	Glu
625					630					635				640
Leu	Cys	Phe	Asp	Val	Asn	Pro	Ala	Val	Ala	Gly	Cys	Lys	Trp	Ser
				645					650					655
Val	His	Asp	Val	Gln	Trp	Arg	Thr	Gln	Phe	Ala	Thr	Ile	Thr	Val
			660					665					670	Ser
Tyr	Lys	His	Gly	Ser	Met	Ile	Thr	Thr	His	Ala	Lys	Gly	His	Ser
		675					680					685		Trp
Gly	Phe	Gln	Asp	Thr	Ser	Val	Leu	Val	Lys	Asp	Glu	Cys	Thr	Asp
	690					695					700			Tyr
Asn	Ile	Tyr	Gly	Phe	Gln	Gly	Thr	Gly	Ile	Ile	Arg	Asn	Thr	Thr
705					710					715				720
Arg	Leu	Val	Ala	Gly	Leu	Tyr	Tyr	Thr	Ser	Ile	Ser	Gly	Asp	Leu
				725					730					735
Ala	Phe	Lys	Asn	Ser	Thr	Thr	Gly	Glu	Ile	Phe	Thr	Val	Val	Pro
			740					745					750	Cys
Asp	Leu	Thr	Ala	Gln	Val	Ala	Val	Ile	Asn	Asp	Glu	Ile	Val	Gly
		755					760					765		Ala
Ile	Thr	Ala	Val	Asn	Gln	Thr	Asp	Leu	Phe	Glu	Phe	Val	Asn	Asn
	770					775					780			Thr
Gln	Ala	Arg	Arg	Ser	Arg	Ser	Ser	Thr	Pro	Asn	Phe	Val	Thr	Ser
785					790					795				800

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Thr Met Pro Gln Phe Tyr Tyr Ile Thr Lys Trp Asn Asn Asp Thr Ser
 805 810 815

Ser Asn Cys Thr Ser Ala Ile Thr Tyr Ser Ser Phe Ala Ile Cys Asn
 820 825 830

Thr Gly Glu Ile Lys Tyr Val Asn Val Thr His Val Glu Ile Val Asp
 835 840 845

Asp Ser Ile Gly Val Ile Lys Pro Val Ser Thr Gly Asn Ile Ser Ile
 850 855 860

Pro Lys Asn Phe Thr Val Ala Val Gln Ala Glu Tyr Ile Gln Ile Gln
 865 870 875 880

Val Lys Pro Val Val Val Asp Cys Ala Thr Tyr Val Cys Asn Gly Asn
 885 890 895

Thr His Cys Leu Lys Leu Leu Thr Gln Tyr Thr Ser Ala Cys Gln Thr
 900 905 910

Ile Glu Asn Ala Leu Asn Leu Gly Ala Arg Leu Glu Ser Leu Met Leu
 915 920 925

Asn Asp Met Ile Thr Val Ser Asp Arg Gly Leu Glu Leu Ala Thr Val
 930 935 940

Glu Arg Phe Asn Ala Thr Ala Leu Gly Gly Glu Lys Leu Gly Gly Leu
 945 950 955 960

Tyr Phe Asp Gly Leu Ser Ser Leu Leu Pro Lys Ile Gly Lys Arg
 965 970 975

Ser Ala Val Glu Asp Leu Leu Phe Asn Lys Val Val Thr Ser Gly Leu
 980 985 990

Gly Thr Val Asp Asp Asp Tyr Lys Lys Cys Ser Ser Gly Thr Asp Val
 995 1000 1005

Ala Asp Leu Val Cys Ala Gln Tyr Tyr Asn Gly Ile Met Val Leu
 1010 1015 1020

Pro Gly Val Val Asp Gly Asn Lys Met Ser Met Tyr Thr Ala Ser
 1025 1030 1035

Leu Ile Gly Gly Met Ala Leu Gly Ser Ile Thr Ser Ala Val Ala
 1040 1045 1050

Val Pro Phe Ala Met Gln Val Gln Ala Arg Leu Asn Tyr Val Ala
 1055 1060 1065

Leu Gln Thr Asp Val Leu Gln Glu Asn Gln Lys Ile Leu Ala Asn
 1070 1075 1080

Ala Phe Asn Asn Ala Ile Gly Asn Ile Thr Leu Ala Leu Gly Lys
 1085 1090 1095

Val Ser Asn Ala Ile Thr Thr Thr Ser Asp Gly Phe Asn Ser Met
 1100 1105 1110

Ala Ser Ala Leu Thr Lys Ile Gln Ser Val Val Asn Gln Gln Gly
 1115 1120 1125

Glu Ala Leu Ser Gln Leu Thr Ser Gln Leu Gln Lys Asn Phe Gln
 1130 1135 1140

Ala Ile Ser Ser Ser Ile Ala Glu Ile Tyr Asn Arg Leu Glu Lys
 1145 1150 1155

Val Glu Ala Asp Ala Gln Val Asp Arg Leu Ile Thr Gly Arg Leu
 1160 1165 1170

Ala Ala Leu Asn Ala Tyr Val Ser Gln Thr Leu Thr Gln Tyr Ala
 1175 1180 1185

Glu Val Lys Ala Ser Arg Gln Ile Ala Leu Glu Lys Val Asn Glu
 1190 1195 1200

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Cys Val Lys Ser Gln Ser Asn Arg Tyr Gly Phe Cys Gly Asn Gly
 1205 1210 1215
 Thr His Leu Phe Ser Leu Val Asn Ser Ala Pro Glu Gly Leu Leu
 1220 1225 1230
 Phe Phe His Thr Val Leu Leu Pro Thr Glu Trp Glu Glu Val Thr
 1235 1240 1245
 Ala Trp Ser Gly Ile Cys Val Asn Asp Thr Tyr Ala Tyr Val Leu
 1250 1255 1260
 Lys Asp Phe Asp His Ser Ile Phe Ser Tyr Asn Gly Thr Tyr Met
 1265 1270 1275
 Val Thr Pro Arg Asn Met Phe Gln Pro Arg Lys Pro Gln Met Ser
 1280 1285 1290
 Asp Phe Val Gln Ile Thr Ser Cys Glu Val Thr Phe Leu Asn Met
 1295 1300 1305
 Thr Tyr Thr Thr Phe Gln Glu Ile Val Ile Asp Tyr Ile Asp Ile
 1310 1315 1320
 Asn Lys Thr Ile Ala Asp Met Leu Glu Gln Tyr Asn Pro Asn Tyr
 1325 1330 1335
 Thr Thr Pro Glu Leu Asn Leu Leu Leu Asp Ile Phe Asn Gln Thr
 1340 1345 1350
 Lys Leu Asn Leu Thr Ala Glu Ile Asp Gln Leu Glu Gln Arg Ala
 1355 1360 1365
 Asp Asn Leu Thr Thr Ile Ala His Glu Leu Gln Gln Tyr Ile Asp
 1370 1375 1380
 Asn Leu Asn Lys Thr Leu Val Asp Leu Asp Trp Leu Asn Arg Ile
 1385 1390 1395
 Glu Thr Tyr Val Lys Trp Pro Trp Tyr Val Trp Leu Leu Ile Gly
 1400 1405 1410
 Leu Val Val Val Phe Cys Ile Pro Leu Leu Leu Phe Cys Cys Leu
 1415 1420 1425
 Ser Thr Gly Phe Cys Gly Cys Phe Gly Cys Val Gly Ser Cys Cys
 1430 1435 1440
 His Ser Leu Cys Ser Arg Arg Gln Phe Glu Thr Tyr Glu Pro Ile
 1445 1450 1455
 Glu Lys Val His Ile His
 1460

<210> SEQ ID NO 57

<211> LENGTH: 1235

<212> TYPE: PRT

<213> ORGANISM: Mouse hepatitis virus

<400> SEQUENCE: 57

Met Leu Phe Val Phe Ile Leu Leu Leu Pro Ser Cys Leu Gly Tyr Ile
 1 5 10 15
 Gly Asp Phe Arg Cys Ile Gln Thr Val Asn Tyr Asn Gly Asn Asn Ala
 20 25 30
 Ser Ala Pro Ser Ile Ser Thr Glu Ala Val Asp Val Ser Lys Gly Arg
 35 40 45
 Gly Thr Tyr Tyr Val Leu Asp Arg Val Tyr Leu Asn Ala Thr Leu Leu
 50 55 60
 Leu Thr Gly Tyr Tyr Pro Val Asp Gly Ser Asn Tyr Arg Asn Leu Ala
 65 70 75 80
 Leu Thr Gly Thr Asn Thr Leu Ser Leu Thr Trp Phe Lys Pro Pro Phe
 85 90 95

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Leu Ser Glu Phe Asn Asp Gly Ile Phe Ala Lys Val Gln Asn Leu Lys
 100 105 110
 Thr Asn Thr Pro Thr Gly Ala Thr Ser Tyr Phe Pro Thr Ile Val Ile
 115 120 125
 Gly Ser Leu Phe Gly Asn Thr Ser Tyr Thr Val Val Leu Glu Pro Tyr
 130 135 140
 Asn Asn Ile Ile Met Ala Ser Val Cys Thr Tyr Thr Ile Cys Gln Leu
 145 150 155 160
 Pro Tyr Thr Pro Cys Lys Pro Asn Thr Asn Gly Asn Arg Val Ile Gly
 165 170 175
 Phe Trp His Thr Asp Val Lys Pro Pro Ile Cys Leu Leu Lys Arg Asn
 180 185 190
 Phe Thr Phe Asn Val Asn Ala Pro Trp Leu Tyr Phe His Phe Tyr Gln
 195 200 205
 Gln Gly Gly Thr Phe Tyr Ala Tyr Tyr Ala Asp Lys Pro Ser Ala Thr
 210 215 220
 Thr Phe Leu Phe Ser Val Tyr Ile Gly Asp Ile Leu Thr Gln Tyr Phe
 225 230 235 240
 Val Leu Pro Phe Ile Cys Thr Pro Thr Ala Gly Ser Thr Leu Ala Pro
 245 250 255
 Leu Tyr Trp Val Thr Pro Leu Leu Lys Arg Gln Tyr Leu Phe Asn Phe
 260 265 270
 Asn Glu Lys Gly Val Ile Thr Ser Ala Val Asp Cys Ala Ser Ser Tyr
 275 280 285
 Ile Ser Glu Ile Lys Cys Lys Thr Gln Ser Leu Leu Pro Ser Thr Gly
 290 295 300
 Val Tyr Asp Leu Ser Gly Tyr Thr Val Gln Pro Val Gly Val Val Tyr
 305 310 315 320
 Arg Arg Val Pro Asn Leu Pro Asp Cys Lys Ile Glu Glu Trp Leu Thr
 325 330 335
 Ala Lys Ser Val Pro Ser Pro Leu Asn Trp Glu Arg Arg Thr Phe Gln
 340 345 350
 Asn Cys Asn Phe Asn Leu Ser Ser Leu Leu Arg Tyr Val Gln Ala Glu
 355 360 365
 Ser Leu Ser Cys Asn Asn Ile Asp Ala Ser Lys Val Tyr Gly Met Cys
 370 375 380
 Phe Gly Ser Val Ser Val Asp Lys Phe Ala Ile Pro Arg Ser Arg Gln
 385 390 395 400
 Ile Asp Leu Gln Ile Gly Asn Ser Gly Phe Leu Gln Thr Ala Asn Tyr
 405 410 415
 Lys Ile Asp Thr Ala Ala Thr Ser Cys Gln Leu Tyr Tyr Ser Leu Pro
 420 425 430
 Lys Asn Asn Val Thr Ile Asn Asn Tyr Asn Pro Ser Ser Trp Asn Arg
 435 440 445
 Arg Tyr Gly Phe Lys Val Asn Asp Arg Cys Gln Ile Phe Ala Asn Ile
 450 455 460
 Leu Leu Asn Gly Ile Asn Ser Gly Thr Thr Cys Ser Thr Asp Leu Gln
 465 470 475 480
 Leu Pro Asn Thr Glu Val Ala Thr Gly Val Cys Val Arg Tyr Asp Leu
 485 490 495
 Tyr Gly Ile Thr Gly Gln Gly Val Phe Lys Glu Val Lys Ala Asp Tyr
 500 505 510

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Tyr Asn Ser Trp Gln Ala Leu Leu Tyr Asp Val Asn Gly Asn Leu Asn
515 520 525
Gly Phe Arg Asp Leu Thr Thr Asn Lys Thr Tyr Thr Ile Arg Ser Cys
530 535 540
Tyr Ser Gly Arg Val Ser Ala Ala Tyr His Lys Glu Ala Pro Glu Pro
545 550 555 560
Ala Leu Leu Tyr Arg Asn Ile Asn Cys Ser Tyr Val Phe Thr Asn Asn
565 570 575
Ile Ser Arg Glu Glu Asn Pro Leu Asn Tyr Phe Asp Ser Tyr Leu Gly
580 585 590
Cys Val Val Asn Ala Asp Asn Arg Thr Asp Glu Ala Leu Pro Asn Cys
595 600 605
Asn Leu Arg Met Gly Ala Gly Leu Cys Val Asp Tyr Ser Lys Ser Arg
610 615 620
Arg Ala Arg Arg Ser Val Ser Thr Gly Tyr Arg Leu Thr Thr Phe Glu
625 630 635 640
Pro Tyr Met Pro Met Leu Val Asn Asp Ser Val Gln Ser Val Gly Gly
645 650 655
Leu Tyr Glu Met Gln Ile Pro Thr Asn Phe Thr Ile Gly His His Glu
660 665 670
Glu Phe Ile Gln Ile Arg Ala Pro Lys Val Thr Ile Asp Cys Ala Ala
675 680 685
Phe Val Cys Gly Asp Asn Ala Ala Cys Arg Gln Gln Leu Val Glu Tyr
690 695 700
Gly Ser Phe Cys Asp Asn Val Asn Ala Ile Leu Asn Glu Val Asn Asn
705 710 715 720
Leu Leu Asp Asn Met Gln Leu Gln Val Ala Ser Ala Leu Met Gln Gly
725 730 735
Val Thr Ile Ser Ser Arg Leu Pro Asp Gly Ile Ser Gly Pro Ile Asp
740 745 750
Asp Ile Asn Phe Ser Pro Leu Leu Gly Cys Ile Gly Ser Thr Cys Ala
755 760 765
Glu Asp Gly Asn Gly Pro Ser Ala Ile Arg Gly Arg Ser Ala Ile Glu
770 775 780
Asp Leu Leu Phe Asp Lys Val Lys Leu Ser Asp Val Gly Phe Val Glu
785 790 795 800
Ala Tyr Asn Asn Cys Thr Gly Gly Gln Glu Val Arg Asp Leu Leu Cys
805 810 815
Val Gln Ser Phe Asn Gly Ile Lys Val Leu Pro Pro Val Leu Ser Glu
820 825 830
Ser Gln Ile Ser Gly Tyr Thr Ala Gly Ala Thr Ala Ala Ala Met Phe
835 840 845
Pro Pro Trp Thr Ala Ala Ala Gly Val Pro Phe Ser Leu Asn Val Gln
850 855 860
Tyr Arg Ile Asn Gly Leu Gly Val Thr Met Asn Val Leu Ser Glu Asn
865 870 875 880
Gln Lys Met Ile Ala Ser Ala Phe Asn Asn Ala Leu Gly Ala Ile Gln
885 890 895
Glu Gly Phe Asp Ala Thr Asn Ser Ala Leu Gly Lys Ile Gln Ser Val
900 905 910
Val Asn Ala Asn Ala Glu Ala Leu Asn Asn Leu Leu Asn Gln Leu Ser
915 920 925
Asn Arg Phe Gly Ala Ile Ser Ala Ser Leu Gln Glu Ile Leu Thr Arg

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930	935	940
Leu Asp Ala Val Glu 945	Ala Lys Ala Gln Ile 950	Asp Arg Leu Ile Asn Gly 955 960
Arg Leu Thr Ala 965	Leu Asn Ala Tyr Ile Ser 970	Lys Gln Leu Ser Asp Ser 975
Thr Leu Ile Lys Phe Ser Ala Ala 980	Gln Ala Ile Glu Lys Val Asn Glu 985	
Cys Val Lys Ser Gln Thr Thr Arg 995	Ile Asn Phe Cys Gly Asn Gly Asn 1000	
His Ile Leu Ser Leu Val Gln 1010	Asn Ala Pro Tyr Gly Leu Cys Phe 1015	
Ile His Phe Ser Tyr Val Pro Thr Ser Phe Lys Thr Ala Asn Val 1025		1030 1035
Ser Pro Gly Leu Cys Ile Ser Gly Asp Arg Gly Leu Ala Pro Lys 1040		1045 1050
Ala Gly Tyr Phe Val Gln Asp Asn Gly Glu Trp Lys Phe Thr Gly 1055		1060 1065
Ser Asn Tyr Tyr Tyr Pro Glu Pro Ile Thr Asp Lys Asn Ser Val 1070		1075 1080
Ala Met Ile Ser Cys Ala Val Asn Tyr Thr Lys Ala Pro Glu Val 1085		1090 1095
Phe Leu Asn Asn Ser Ile Pro Asn Leu Pro Asp Phe Lys Glu Glu 1100		1105 1110
Leu Asp Lys Trp Phe Lys Asn Gln Thr Ser Ile Ala Pro Asp Leu 1115		1120 1125
Ser Leu Asp Phe Glu Lys Leu Asn Val Thr Phe Leu Asp Leu Thr 1130		1135 1140
Tyr Glu Met Asn Arg Ile Gln Asp Ala Ile Lys Lys Leu Asn Glu 1145		1150 1155
Ser Tyr Ile Asn Leu Lys Glu Val Gly Thr Tyr Glu Met Tyr Val 1160		1165 1170
Lys Trp Pro Trp Tyr Val Trp Leu Leu Ile Gly Leu Ala Gly Val 1175		1180 1185
Ala Val Cys Val Leu Leu Phe Phe Ile Cys Cys Cys Thr Gly Cys 1190		1195 1200
Gly Ser Cys Cys Phe Arg Lys Cys Gly Ser Cys Cys Asp Glu Tyr 1205		1210 1215
Gly Gly His Gln Asp Ser Ile Val Ile His Asn Ile Ser Ala His 1220		1225 1230
Glu Asp 1235		

<210> SEQ ID NO 58

<211> LENGTH: 1363

<212> TYPE: PRT

<213> ORGANISM: human coronavirus

<400> SEQUENCE: 58

Met Phe Leu Ile Leu Leu Ile Ser Leu Pro Met Ala Leu Ala Val Ile
1 5 10 15

Gly Asp Leu Lys Cys Thr Thr Val Ala Ile Asn Asp Val Asp Thr Gly
20 25 30

Val Pro Ser Thr Ser Thr Asp Ile Val Asp Val Thr Asn Gly Leu Gly
35 40 45

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Thr	Tyr	Tyr	Val	Leu	Asp	Arg	Val	Tyr	Leu	Asn	Thr	Thr	Leu	Leu	Leu
50						55					60				
Asn	Gly	Tyr	Tyr	Pro	Thr	Ser	Gly	Ser	Thr	Tyr	Arg	Asn	Met	Ala	Leu
65					70					75					80
Lys	Gly	Thr	Leu	Leu	Leu	Ser	Arg	Leu	Trp	Phe	Lys	Pro	Pro	Phe	Leu
			85						90					95	
Ser	Asp	Phe	Ile	Asn	Gly	Ile	Phe	Ala	Lys	Val	Lys	Asn	Thr	Lys	Val
		100						105					110		
Ile	Lys	His	Gly	Val	Met	Tyr	Ser	Glu	Phe	Pro	Ala	Ile	Thr	Ile	Gly
		115						120					125		
Ser	Thr	Phe	Val	Asn	Thr	Ser	Tyr	Ser	Val	Val	Val	Gln	Pro	His	Thr
		130					135					140			
Thr	Asn	Leu	Asp	Asn	Lys	Leu	Gln	Gly	Leu	Leu	Glu	Ile	Ser	Val	Cys
145					150					155					160
Gln	Tyr	Thr	Met	Cys	Glu	Tyr	Pro	Asn	Thr	Ile	Cys	His	Pro	Asn	Leu
			165						170					175	
Gly	Asn	Arg	Arg	Val	Glu	Leu	Trp	His	Trp	Asp	Thr	Gly	Val	Val	Ser
		180						185					190		
Cys	Leu	Tyr	Lys	Arg	Asn	Phe	Thr	Tyr	Asp	Val	Asn	Ala	Asp	Tyr	Leu
		195					200					205			
Tyr	Phe	His	Phe	Tyr	Gln	Glu	Gly	Gly	Ile	Phe	Tyr	Ala	Tyr	Phe	Thr
		210				215					220				
Asp	Thr	Gly	Val	Val	Thr	Lys	Phe	Leu	Phe	Asn	Val	Tyr	Leu	Gly	Thr
225					230					235					240
Val	Leu	Ser	Tyr	Tyr	Tyr	Val	Met	Pro	Leu	Thr	Cys	Asn	Ser	Ala	Met
			245						250					255	
Thr	Leu	Glu	Tyr	Trp	Val	Thr	Pro	Leu	Thr	Ser	Lys	Gln	Tyr	Leu	Leu
			260					265					270		
Ala	Phe	Asn	Gln	Asp	Gly	Val	Ile	Phe	Asn	Ala	Val	Asp	Cys	Lys	Ser
		275					280					285			
Asp	Phe	Met	Ser	Glu	Ile	Lys	Cys	Lys	Thr	Leu	Ser	Ile	Ala	Pro	Ser
		290				295					300				
Thr	Gly	Val	Tyr	Glu	Leu	Asn	Gly	Tyr	Thr	Val	Gln	Pro	Ile	Ala	Asp
305					310					315					320
Val	Tyr	Arg	Arg	Ile	Pro	Asn	Leu	Pro	Asp	Cys	Asn	Ile	Glu	Ala	Trp
			325						330					335	
Leu	Asn	Asp	Lys	Ser	Val	Pro	Ser	Pro	Leu	Asn	Trp	Glu	Arg	Lys	Thr
			340					345					350		
Phe	Ser	Asn	Cys	Asn	Phe	Asn	Met	Ser	Ser	Leu	Met	Ser	Phe	Ile	Gln
		355					360					365			
Ala	Asp	Ser	Phe	Thr	Cys	Asn	Asn	Ile	Asp	Ala	Ala	Lys	Ile	Tyr	Gly
		370				375					380				
Met	Cys	Phe	Ser	Ser	Ile	Thr	Ile	Asp	Lys	Phe	Ala	Ile	Pro	Asn	Gly
385					390					395					400
Arg	Lys	Val	Asp	Leu	Gln	Leu	Gly	Asn	Leu	Gly	Tyr	Leu	Gln	Ser	Phe
			405						410					415	
Asn	Tyr	Arg	Ile	Asp	Thr	Thr	Ala	Thr	Ser	Cys	Gln	Leu	Tyr	Tyr	Asn
			420					425					430		
Leu	Pro	Ala	Ala	Asn	Val	Ser	Val	Ser	Arg	Phe	Asn	Pro	Ser	Ile	Trp
		435					440					445			
Asn	Arg	Arg	Phe	Gly	Phe	Thr	Glu	Gln	Ser	Val	Phe	Lys	Pro	Gln	Pro
		450				455					460				
Ala	Gly	Val	Phe	Thr	Asp	His	Asp	Val	Val	Tyr	Ala	Gln	His	Cys	Phe

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465	470	475	480
Lys Ala Pro Thr	Asn Phe Cys Pro Cys	Lys Leu Asp Gly Ser	Leu Cys
	485	490	495
Val Gly Asn Gly	Pro Gly Ile Asp	Ala Gly Tyr Lys	Asn Ser Gly Ile
	500	505	510
Gly Thr Cys Pro	Ala Gly Thr Asn Tyr	Leu Thr Cys His	Asn Ala Val
	515	520	525
Gln Cys Asn Cys	Leu Cys Thr Pro Asp	Pro Ile Thr Ser	Lys Ser Thr
	530	535	540
Gly Pro Tyr Lys	Cys Pro Gln Thr Lys Tyr	Leu Val Gly Ile	Gly Glu
	545	550	555
His Cys Ser Gly	Leu Ala Ile Lys Ser	Asp Tyr Cys Gly	Gly Asn Pro
	565	570	575
Cys Thr Cys Gln	Pro Gln Ala Phe	Leu Gly Trp Ser	Val Asp Ser Cys
	580	585	590
Leu Gln Gly Asp	Arg Cys Asn Ile Phe	Ala Asn Phe Ile	Leu His Asp
	595	600	605
Val Asn Ser Gly	Thr Thr Cys Ser Thr	Asp Leu Gln Lys	Ser Asn Thr
	610	615	620
Asp Ile Ile Leu	Gly Val Cys Val Asn Tyr	Asp Leu Tyr Gly	Ile Thr
	625	630	635
Gly Gln Gly Ile	Phe Val Glu Val Asn Ala	Pro Tyr Tyr Asn	Ser Trp
	645	650	655
Gln Asn Leu Leu	Tyr Asp Ser Asn Gly	Asn Leu Tyr Gly	Phe Arg Asp
	660	665	670
Tyr Leu Thr Asn	Arg Thr Phe Met Ile	Arg Ser Cys Tyr	Ser Gly Arg
	675	680	685
Val Ser Ala Ala	Phe His Ala Asn Ser Ser	Glu Pro Ala Leu	Leu Phe
	690	695	700
Arg Asn Ile Lys	Cys Asn Tyr Val Phe	Asn Asn Thr Leu	Ser Arg Gln
	705	710	715
Leu Gln Pro Ile	Asn Tyr Phe Asp Ser	Tyr Leu Gly Cys	Val Val Asn
	725	730	735
Ala Asp Asn Ser	Thr Ala Ser Ala Val	Gln Thr Cys Asp	Leu Thr Val
	740	745	750
Gly Ser Gly Tyr	Cys Val Asp Tyr Ser	Thr Lys Arg Arg	Ser Arg Arg
	755	760	765
Ala Ile Thr Thr	Gly Tyr Arg Phe Thr	Asn Phe Glu Pro	Phe Thr Val
	770	775	780
Asn Ser Val Asn	Asp Ser Leu Glu His Val	Gly Gly Leu Tyr	Glu Ile
	785	790	795
Gln Ile Pro Ser	Glu Phe Thr Ile Gly	Asn Met Glu Glu	Phe Ile Gln
	805	810	815
Thr Ser Ser Pro	Lys Val Thr Ile Asp	Cys Ser Ala Phe	Val Cys Gly
	820	825	830
Asp Cys Ala Ala	Cys Lys Ser Gln Leu	Val Glu Tyr Gly	Ser Phe Cys
	835	840	845
Asp Asn Ile Asn	Ala Ile Leu Thr Glu	Val Asn Glu Leu	Leu Asp Thr
	850	855	860
Thr Gln Leu Gln	Val Ala Asn Ser Leu	Met Asn Gly Val	Thr Leu Ser
	865	870	875
Thr Lys Leu Lys	Asp Gly Val Asn Phe	Asn Val Asp Asp	Val Asn Phe
	885	890	895

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Ser Pro Val Leu Gly Cys Leu Gly Ser Glu Cys Asn Lys Val Ser Ser
 900 905 910

Arg Ser Ala Ile Glu Asp Leu Leu Phe Ser Lys Val Arg Leu Ser Asp
 915 920 925

Val Gly Phe Val Glu Ala Tyr Asn Asn Cys Thr Gly Gly Ala Gly Ile
 930 935 940

Arg Asp Leu Ile Cys Val Gln Ser Tyr Asn Gly Ile Lys Val Leu Pro
 945 950 955 960

Pro Leu Leu Ser Asp Asn Gln Ile Ser Gly Tyr Thr Leu Ala Ala Thr
 965 970 975

Ser Ala Asn Leu Phe Pro Pro Trp Ser Ala Ala Ala Gly Val Pro Phe
 980 985 990

Tyr Leu Asn Val Gln Tyr Arg Ile Asn Gly Ile Gly Val Thr Met Asp
 995 1000 1005

Val Leu Ser Gln Asn Gln Lys Leu Ile Ala Asn Ala Phe Asn Asn
 1010 1015 1020

Ala Leu Asp Ala Ile Gln Glu Gly Phe Asp Ala Thr Asn Ser Ala
 1025 1030 1035

Leu Val Lys Ile Gln Ala Val Val Asn Ala Asp Ala Glu Ala Leu
 1040 1045 1050

Asn Asn Leu Leu Gln Gln Leu Ser Asn Arg Phe Gly Ala Ile Ser
 1055 1060 1065

Ser Ser Leu Gln Glu Ile Leu Ser Arg Leu Asp Ala Leu Glu Ala
 1070 1075 1080

Gln Ala Gln Ile Asp Arg Leu Ile Asn Gly Arg Leu Thr Ala Leu
 1085 1090 1095

Asp Ala Tyr Val Ser Gln Gln Leu Ser Asp Ser Thr Leu Val Lys
 1100 1105 1110

Phe Ser Ala Ala Gln Ala Met Glu Lys Val Asn Glu Cys Val Lys
 1115 1120 1125

Ser Gln Ser Ser Arg Ile Asn Phe Cys Gly Asn Gly Asn His Ile
 1130 1135 1140

Ile Ser Leu Val Gln Asn Ala Pro Tyr Gly Leu Tyr Phe Ile His
 1145 1150 1155

Phe Ser Tyr Val Pro Thr Lys Tyr Val Thr Ala Lys Val Ser Pro
 1160 1165 1170

Gly Leu Cys Ile Ala Gly Asp Arg Gly Ile Ala Pro Lys Ser Gly
 1175 1180 1185

Tyr Phe Val Asn Val Asn Asn Thr Trp Met Phe Thr Gly Ser Arg
 1190 1195 1200

Tyr Tyr Tyr Pro Glu Pro Ile Thr Gly Asn Asn Val Val Val Met
 1205 1210 1215

Ser Thr Cys Ala Val Asn Tyr Thr Lys Ala Pro Asp Val Met Leu
 1220 1225 1230

Asn Ile Ser Thr Pro Asn Leu Pro Asp Phe Lys Glu Glu Leu Asp
 1235 1240 1245

Gln Trp Phe Lys Asn Gln Thr Leu Val Ala Pro Asp Leu Ser Leu
 1250 1255 1260

Asp Tyr Ile Asn Val Thr Phe Leu Asp Leu Gln Asp Glu Met Asn
 1265 1270 1275

Arg Leu Gln Glu Ala Ile Lys Val Leu Asn Gln Ser Tyr Ile Asn
 1280 1285 1290

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Leu Lys Asp Ile Gly Thr Tyr Glu Tyr Tyr Val Lys Trp Pro Trp
 1295 1300 1305

Tyr Val Trp Leu Leu Ile Gly Phe Ala Gly Val Ala Met Leu Val
 1310 1315 1320

Leu Leu Phe Phe Ile Cys Cys Cys Thr Gly Cys Gly Thr Ser Cys
 1325 1330 1335

Phe Lys Lys Cys Gly Gly Cys Cys Asp Asp Tyr Thr Gly His Gln
 1340 1345 1350

Glu Leu Val Ile Lys Thr Ser His Glu Gly
 1355 1360

<210> SEQ ID NO 59

<211> LENGTH: 1383

<212> TYPE: PRT

<213> ORGANISM: Porcine epidemic diarrhea virus

<400> SEQUENCE: 59

Met Arg Ser Leu Ile Tyr Phe Trp Leu Leu Leu Pro Val Leu Pro Thr
 1 5 10 15

Leu Ser Leu Pro Gln Asp Val Thr Arg Cys Gln Ser Thr Thr Asn Phe
 20 25 30

Arg Arg Phe Phe Ser Lys Phe Asn Val Gln Ala Pro Ala Val Val Val
 35 40 45

Leu Gly Gly Tyr Leu Pro Ser Met Asn Ser Ser Ser Trp Tyr Cys Gly
 50 55 60

Thr Gly Ile Glu Thr Ala Ser Gly Val His Gly Ile Phe Leu Ser Tyr
 65 70 75 80

Ile Asp Ser Gly Gln Gly Phe Glu Ile Gly Ile Ser Gln Glu Pro Phe
 85 90 95

Asp Pro Ser Gly Tyr Gln Leu Tyr Leu His Lys Ala Thr Asn Gly Asn
 100 105 110

Thr Asn Ala Thr Ala Arg Leu Arg Ile Cys Gln Phe Pro Asp Asn Lys
 115 120 125

Thr Leu Gly Pro Thr Val Asn Asp Val Thr Thr Gly Arg Asn Cys Leu
 130 135 140

Phe Asn Lys Ala Ile Pro Ala Tyr Met Arg Asp Gly Lys Asp Ile Val
 145 150 155 160

Val Gly Ile Thr Trp Asp Asn Asp Arg Val Thr Val Phe Ala Asp Lys
 165 170 175

Ile Tyr His Phe Tyr Leu Lys Asn Asp Trp Ser Arg Val Ala Thr Arg
 180 185 190

Cys Tyr Asn Arg Arg Ser Cys Ala Met Gln Tyr Val Tyr Thr Pro Thr
 195 200 205

Tyr Tyr Met Leu Asn Val Thr Ser Ala Gly Glu Asp Gly Ile Tyr Tyr
 210 215 220

Glu Pro Cys Thr Ala Asn Cys Thr Gly Tyr Ala Ala Asn Val Phe Ala
 225 230 235 240

Thr Asp Ser Asn Gly His Ile Pro Glu Gly Phe Ser Phe Asn Asn Trp
 245 250 255

Phe Leu Leu Ser Asn Asp Ser Thr Leu Leu His Gly Lys Val Val Ser
 260 265 270

Asn Gln Pro Leu Leu Val Asn Cys Leu Leu Ala Ile Pro Lys Ile Tyr
 275 280 285

Gly Leu Gly Gln Phe Phe Ser Phe Asn His Thr Met Asp Gly Val Cys
 290 295 300

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Asn Gly Ala Ala Val Asp Arg Ala Pro Glu Ala Leu Arg Phe Asn Ile
 305 310 315 320
 Asn Asp Thr Ser Val Ile Leu Ala Glu Gly Ser Ile Val Leu His Thr
 325 330 335
 Ala Leu Gly Thr Asn Leu Ser Phe Val Cys Ser Asn Ser Ser Asp Pro
 340 345 350
 His Leu Ala Ile Phe Ala Ile Pro Leu Gly Ala Thr Glu Val Pro Tyr
 355 360 365
 Tyr Cys Phe Leu Lys Val Asp Thr Tyr Asn Ser Thr Val Tyr Lys Phe
 370 375 380
 Leu Ala Val Leu Pro Ser Thr Val Arg Glu Ile Val Ile Thr Lys Tyr
 385 390 395 400
 Gly Asp Val Tyr Val Asn Gly Phe Gly Tyr Leu His Leu Gly Leu Leu
 405 410 415
 Asp Ala Val Thr Ile Tyr Phe Thr Gly His Gly Thr Asp Asp Asp Val
 420 425 430
 Ser Gly Phe Trp Thr Ile Ala Ser Thr Asn Phe Val Asp Ala Leu Ile
 435 440 445
 Glu Val Gln Gly Thr Ser Ile Gln Arg Ile Leu Tyr Cys Asp Asp Pro
 450 455 460
 Val Ser Gln Leu Lys Cys Ser Gln Val Ala Phe Asp Leu Asp Asp Gly
 465 470 475 480
 Phe Tyr Pro Ile Ser Ser Arg Asn Leu Leu Ser His Glu Gln Pro Ile
 485 490 495
 Ser Phe Val Thr Leu Pro Ser Phe Asn Asp His Ser Phe Val Asn Ile
 500 505 510
 Thr Val Ser Ala Ala Phe Gly Gly Leu Ser Ser Ala Asn Leu Val Ala
 515 520 525
 Ser Asp Thr Thr Ile Asn Gly Phe Ser Ser Phe Cys Val Asp Thr Arg
 530 535 540
 Gln Phe Thr Ile Thr Leu Phe Tyr Asn Val Thr Asn Ser Tyr Gly Tyr
 545 550 555 560
 Val Ser Lys Ser Gln Asp Ser Asn Cys Pro Phe Thr Leu Gln Ser Val
 565 570 575
 Asn Asp Tyr Leu Ser Phe Ser Lys Phe Cys Val Ser Thr Ser Leu Leu
 580 585 590
 Ala Gly Ala Cys Thr Ile Asp Leu Phe Gly Tyr Pro Ala Phe Gly Ser
 595 600 605
 Gly Val Lys Leu Thr Ser Leu Tyr Phe Gln Phe Thr Lys Gly Glu Leu
 610 615 620
 Ile Thr Gly Thr Pro Lys Pro Leu Glu Gly Ile Thr Asp Val Ser Phe
 625 630 635 640
 Met Thr Leu Asp Val Cys Thr Lys Tyr Thr Ile Tyr Gly Phe Lys Gly
 645 650 655
 Glu Gly Ile Ile Thr Leu Thr Asn Ser Ser Ile Leu Ala Gly Val Tyr
 660 665 670
 Tyr Thr Ser Asp Ser Gly Gln Leu Leu Ala Phe Lys Asn Val Thr Ser
 675 680 685
 Gly Ala Val Tyr Ser Val Thr Pro Cys Ser Phe Ser Glu Gln Ala Ala
 690 695 700
 Tyr Val Asn Asp Asp Ile Val Gly Val Ile Ser Ser Leu Ser Asn Ser
 705 710 715 720

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Thr Phe Asn Asn Thr Arg Glu Leu Pro Gly Phe Phe Tyr His Ser Asn
725 730 735

Asp Gly Ser Asn Cys Thr Glu Pro Val Leu Val Tyr Ser Asn Ile Gly
740 745 750

Val Cys Lys Ser Gly Ser Ile Gly Tyr Val Pro Ser Gln Tyr Gly Gln
755 760 765

Val Lys Ile Ala Pro Thr Val Thr Gly Asn Ile Ser Ile Pro Thr Asn
770 775 780

Phe Ser Met Ser Ile Arg Thr Glu Tyr Leu Gln Leu Tyr Asn Thr Pro
785 790 795 800

Val Ser Val Asp Cys Ala Thr Tyr Val Cys Asn Gly Asn Ser Arg Cys
805 810 815

Lys Gln Leu Leu Thr Gln Tyr Thr Ala Ala Cys Lys Thr Ile Glu Ser
820 825 830

Ala Leu Gln Leu Ser Ala Arg Leu Glu Ser Val Glu Val Asn Ser Met
835 840 845

Leu Thr Ile Ser Glu Glu Ala Leu Gln Leu Ala Thr Ile Ser Ser Phe
850 855 860

Asn Gly Asp Gly Tyr Asn Phe Thr Asn Val Leu Gly Ala Ser Val Tyr
865 870 875 880

Asp Pro Ala Ser Gly Arg Val Val Gln Lys Arg Ser Val Ile Glu Asp
885 890 895

Leu Leu Phe Asn Lys Val Val Thr Asn Gly Leu Gly Thr Val Asp Glu
900 905 910

Asp Tyr Lys Arg Cys Ser Asn Gly Arg Ser Val Ala Asp Leu Val Cys
915 920 925

Ala Gln Tyr Tyr Ser Gly Val Met Val Leu Pro Gly Val Val Asp Ala
930 935 940

Glu Lys Leu His Met Tyr Ser Ala Ser Leu Ile Gly Gly Met Ala Leu
945 950 955 960

Gly Gly Ile Thr Ala Ala Ala Ala Leu Pro Phe Ser Tyr Ala Val Gln
965 970 975

Ala Arg Leu Asn Tyr Leu Ala Leu Gln Thr Asp Val Leu Gln Arg Asn
980 985 990

Gln Gln Leu Leu Ala Glu Ser Phe Asn Ser Ala Ile Gly Asn Ile Thr
995 1000 1005

Ser Ala Phe Glu Ser Val Lys Glu Ala Ile Ser Gln Thr Ser Lys
1010 1015 1020

Gly Leu Asn Thr Val Ala His Ala Leu Thr Lys Val Gln Glu Val
1025 1030 1035

Val Asn Ser Gln Gly Ser Ala Leu Asn Gln Leu Thr Val Gln Leu
1040 1045 1050

Gln His Asn Phe Gln Ala Ile Ser Ser Ser Ile Asp Asp Ile Tyr
1055 1060 1065

Ser Arg Leu Asp Ile Leu Leu Ala Asp Val Gln Val Asp Arg Leu
1070 1075 1080

Ile Thr Gly Arg Leu Ser Ala Leu Asn Ala Phe Val Ala Gln Thr
1085 1090 1095

Leu Thr Lys Tyr Thr Glu Val Gln Ala Ser Arg Lys Leu Ala Gln
1100 1105 1110

Gln Lys Val Asn Glu Cys Val Lys Ser Gln Ser Gln Arg Tyr Gly
1115 1120 1125

Phe Cys Gly Gly Asp Gly Glu His Ile Phe Ser Leu Val Gln Ala

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1130	1135	1140
Ala Pro Gln Gly Leu Leu Phe 1145	Leu His Thr Val Leu 1150	Val Pro Gly 1155
Asp Phe Val Asn Val Leu Ala 1160	Ile Ala Gly Leu Cys 1165	Val Asn Gly 1170
Glu Ile Ala Leu Thr Leu Arg 1175	Glu Pro Gly Leu Val 1180	Leu Phe Thr 1185
His Glu Leu Gln Thr Tyr Thr 1190	Ala Thr Glu Tyr 1195	Phe Val Ser Ser 1200
Arg Arg Met Phe Glu Pro Arg 1205	Lys Pro Thr Val Ser 1210	Asp Phe Val 1215
Gln Ile Glu Ser Cys Val Val 1220	Thr Tyr Val Asn Leu 1225	Thr Ser Asp 1230
Gln Leu Pro Asp Val Ile Pro 1235	Asp Tyr Ile Asp Val 1240	Asn Lys Thr 1245
Leu Asp Glu Ile Leu Ala Ser 1250	Leu Pro Asn Arg Thr 1255	Gly Pro Ser 1260
Leu Pro Leu Asp Val Phe Asn 1265	Ala Thr Tyr Leu Asn 1270	Leu Thr Gly 1275
Glu Ile Ala Asp Leu Glu Gln 1280	Arg Ser Glu Ser Leu 1285	Arg Asn Thr 1290
Thr Glu Glu Leu Arg Ser Leu 1295	Ile Asn Asn Ile Asn 1300	Asn Thr Leu 1305
Val Asp Leu Glu Trp Leu Asn 1310	Arg Val Glu Thr Tyr 1315	Ile Lys Trp 1320
Pro Trp Trp Val Trp Leu Ile 1325	Ile Val Ile Val Leu 1330	Ile Phe Val 1335
Val Ser Leu Leu Val Phe Cys 1340	Cys Ile Ser Thr Gly 1345	Cys Cys Gly 1350
Cys Cys Gly Cys Cys Gly Ala 1355	Cys Phe Ser Gly Cys 1360	Cys Arg Gly 1365
Pro Arg Leu Gln Pro Tyr Glu 1370	Ala Phe Glu Lys Val 1375	His Val Gln 1380

<210> SEQ ID NO 60

<211> LENGTH: 1349

<212> TYPE: PRT

<213> ORGANISM: porcine hemagglutinating encephalomyelitis virus

<400> SEQUENCE: 60

Met Phe Phe Ile Leu Leu Ile Ser Leu Pro Ser Ala Phe Ala Val Ile 1 5 10 15
Gly Asp Leu Lys Cys Thr Thr Ser Leu Ile Asn Asp Val Asp Thr Gly 20 25 30
Val Pro Ser Ile Ser Ser Glu Val Val Asp Val Thr Asn Gly Leu Gly 35 40 45
Thr Phe Tyr Val Leu Asp Arg Val Tyr Leu Asn Thr Thr Leu Leu Leu 50 55 60
Asn Gly Tyr Tyr Pro Ile Ser Gly Ala Thr Phe Arg Asn Met Ala Leu 65 70 75 80
Lys Gly Thr Arg Leu Leu Ser Thr Leu Trp Phe Lys Pro Pro Phe Leu 85 90 95
Ser Pro Phe Asn Asp Gly Ile Phe Ala Lys Val Lys Asn Ser Arg Phe 100 105 110

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Ser Lys Asp Gly Val Ile Tyr Ser Glu Phe Pro Ala Ile Thr Ile Gly
 115 120 125
 Ser Thr Phe Val Asn Thr Ser Tyr Ser Ile Val Val Glu Pro His Thr
 130 135 140
 Ser Leu Ile Asn Gly Asn Leu Gln Gly Leu Leu Gln Ile Ser Val Cys
 145 150 155 160
 Gln Tyr Thr Met Cys Glu Tyr Pro His Thr Ile Cys His Pro Asn Leu
 165 170 175
 Gly Asn Gln Arg Ile Glu Leu Trp His Tyr Asp Thr Asp Val Val Ser
 180 185 190
 Cys Leu Tyr Arg Arg Asn Phe Thr Tyr Asp Val Asn Ala Asp Tyr Leu
 195 200 205
 Tyr Phe His Phe Tyr Gln Glu Gly Gly Thr Phe Tyr Ala Tyr Phe Thr
 210 215 220
 Asp Thr Gly Phe Val Thr Lys Phe Leu Phe Lys Leu Tyr Leu Gly Thr
 225 230 235 240
 Val Leu Ser His Tyr Tyr Val Met Pro Leu Thr Cys Asn Ser Ala Leu
 245 250 255
 Ser Leu Glu Tyr Trp Val Thr Pro Leu Thr Thr Arg Gln Phe Leu Leu
 260 265 270
 Ala Phe Asp Gln Asp Gly Val Leu Tyr His Ala Val Asp Cys Ala Ser
 275 280 285
 Asp Phe Met Ser Glu Ile Met Cys Lys Thr Ser Ser Ile Thr Pro Pro
 290 295 300
 Thr Gly Val Tyr Glu Leu Asn Gly Tyr Thr Val Gln Pro Val Ala Thr
 305 310 315 320
 Val Tyr Arg Arg Ile Pro Asp Leu Pro Asn Cys Asp Ile Glu Ala Trp
 325 330 335
 Leu Asn Ser Lys Thr Val Ser Ser Pro Leu Asn Trp Glu Arg Lys Ile
 340 345 350
 Phe Ser Asn Cys Asn Phe Asn Met Gly Arg Leu Met Ser Phe Ile Gln
 355 360 365
 Ala Asp Ser Phe Gly Cys Asn Asn Ile Asp Ala Ser Arg Leu Tyr Gly
 370 375 380
 Met Cys Phe Gly Ser Ile Thr Ile Asp Lys Phe Ala Ile Pro Asn Ser
 385 390 395 400
 Arg Lys Val Asp Leu Gln Val Gly Lys Ser Gly Tyr Leu Gln Ser Phe
 405 410 415
 Asn Tyr Lys Ile Asp Thr Ala Val Ser Ser Cys Gln Leu Tyr Tyr Ser
 420 425 430
 Leu Pro Ala Ala Asn Val Ser Val Thr His Tyr Asn Pro Ser Ser Trp
 435 440 445
 Asn Arg Arg Tyr Gly Phe Asn Asn Gln Ser Phe Gly Ser Arg Gly Leu
 450 455 460
 His Asp Ala Val Tyr Ser Gln Gln Cys Phe Asn Thr Pro Asn Thr Tyr
 465 470 475 480
 Cys Pro Cys Arg Thr Ser Gln Cys Ile Gly Gly Ala Gly Thr Gly Thr
 485 490 495
 Cys Pro Val Gly Thr Thr Val Arg Lys Cys Phe Ala Ala Val Thr Lys
 500 505 510
 Ala Thr Lys Cys Thr Cys Trp Cys Gln Pro Asp Pro Ser Thr Tyr Lys
 515 520 525
 Gly Val Asn Ala Trp Thr Cys Pro Gln Ser Lys Val Ser Ile Gln Pro

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530					535					540					
Gly	Gln	His	Cys	Pro	Gly	Leu	Gly	Leu	Val	Glu	Asp	Asp	Cys	Ser	Gly
545					550					555					560
Asn	Pro	Cys	Thr	Cys	Lys	Pro	Gln	Ala	Phe	Ile	Gly	Trp	Ser	Ser	Glu
				565					570						575
Thr	Cys	Leu	Gln	Asn	Gly	Arg	Cys	Asn	Ile	Phe	Ala	Asn	Phe	Ile	Leu
			580					585					590		
Asn	Asp	Val	Asn	Ser	Gly	Thr	Thr	Cys	Ser	Thr	Asp	Leu	Gln	Gln	Gly
		595					600					605			
Asn	Thr	Ile	Ile	Thr	Thr	Asp	Val	Cys	Val	Asn	Tyr	Asp	Leu	Tyr	Gly
	610					615					620				
Ile	Thr	Gly	Gln	Gly	Ile	Leu	Ile	Glu	Val	Asn	Ala	Thr	Tyr	Tyr	Asn
625					630					635					640
Ser	Trp	Gln	Asn	Leu	Leu	Tyr	Asp	Ser	Ser	Gly	Asn	Leu	Tyr	Gly	Phe
				645					650					655	
Arg	Asp	Tyr	Leu	Ser	Asn	Arg	Thr	Phe	Leu	Ile	Arg	Ser	Cys	Tyr	Ser
			660					665					670		
Gly	Arg	Val	Ser	Ala	Val	Phe	His	Ala	Asn	Ser	Ser	Glu	Pro	Ala	Leu
		675					680					685			
Met	Phe	Arg	Asn	Leu	Lys	Cys	Ser	His	Val	Phe	Asn	Asn	Thr	Ile	Leu
	690					695					700				
Arg	Gln	Ile	Gln	Leu	Val	Asn	Tyr	Phe	Asp	Ser	Tyr	Leu	Gly	Cys	Val
705					710					715					720
Val	Asn	Ala	Tyr	Asn	Asn	Thr	Ala	Ser	Ala	Val	Ser	Thr	Cys	Asp	Leu
				725					730					735	
Thr	Val	Gly	Ser	Gly	Tyr	Cys	Val	Asp	Tyr	Val	Thr	Ala	Leu	Arg	Ser
			740					745					750		
Arg	Arg	Ser	Phe	Thr	Thr	Gly	Tyr	Arg	Phe	Thr	Asn	Phe	Glu	Pro	Phe
		755				760						765			
Ala	Ala	Asn	Leu	Val	Asn	Asp	Ser	Ile	Glu	Pro	Val	Gly	Gly	Leu	Tyr
		770				775					780				
Glu	Ile	Gln	Ile	Pro	Ser	Glu	Phe	Thr	Ile	Gly	Asn	Leu	Glu	Glu	Phe
785					790					795					800
Ile	Gln	Thr	Arg	Ser	Pro	Lys	Val	Thr	Ile	Asp	Cys	Ala	Thr	Phe	Val
				805					810					815	
Cys	Gly	Asp	Tyr	Ala	Ala	Cys	Arg	Gln	Gln	Leu	Ala	Glu	Tyr	Gly	Ser
			820					825					830		
Phe	Cys	Glu	Asn	Ile	Asn	Ala	Ile	Leu	Thr	Glu	Val	Asn	Glu	Leu	Leu
		835					840					845			
Asp	Thr	Thr	Gln	Leu	Gln	Val	Ala	Asn	Ser	Leu	Met	Asn	Gly	Val	Thr
	850					855					860				
Leu	Ser	Thr	Lys	Ile	Lys	Asp	Gly	Ile	Asn	Phe	Asn	Val	Asp	Asp	Ile
865					870					875					880
Asn	Phe	Ser	Pro	Val	Leu	Gly	Cys	Leu	Gly	Ser	Glu	Cys	Asn	Arg	Ala
				885					890					895	
Ser	Thr	Arg	Ser	Ala	Ile	Glu	Asp	Leu	Leu	Phe	Asp	Lys	Val	Lys	Leu
			900					905					910		
Ser	Asp	Val	Gly	Phe	Val	Gln	Ala	Tyr	Asn	Asn	Cys	Thr	Gly	Gly	Ala
		915					920					925			
Glu	Ile	Arg	Asp	Leu	Ile	Cys	Val	Gln	Ser	Tyr	Asn	Gly	Ile	Lys	Val
	930					935					940				
Leu	Pro	Pro	Leu	Leu	Ser	Glu	Asn	Gln	Ile	Ser	Gly	Tyr	Thr	Leu	Ala
945					950					955					960

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Ala Thr Ala Ala Ser Leu Phe Pro Pro Trp Thr Ala Ala Ala Gly Val
965 970 975

Pro Phe Tyr Leu Asn Val Gln Tyr Arg Ile Asn Gly Leu Gly Val Thr
980 985 990

Met Asp Val Leu Ser Gln Asn Gln Lys Leu Ile Ala Ser Ala Phe Asn
995 1000 1005

Asn Ala Leu Asp Ala Ile Gln Glu Gly Phe Asp Ala Thr Asn Ser
1010 1015 1020

Ala Leu Val Lys Ile Gln Ala Val Val Asn Ala Asn Ala Glu Ala
1025 1030 1035

Leu Asn Asn Leu Leu Gln Gln Leu Ser Asn Arg Phe Gly Ala Ile
1040 1045 1050

Ser Ala Ser Leu Gln Glu Ile Leu Ser Arg Leu Asp Ala Leu Glu
1055 1060 1065

Ala Lys Ala Gln Ile Asp Arg Leu Ile Asn Gly Arg Leu Thr Ala
1070 1075 1080

Leu Asn Ala Tyr Val Ser Gln Gln Leu Ser Asp Ser Thr Leu Val
1085 1090 1095

Lys Phe Ser Ala Ala Gln Ala Ile Glu Lys Val Asn Glu Cys Val
1100 1105 1110

Lys Ser Gln Ser Ser Arg Ile Asn Phe Cys Gly Asn Gly Asn His
1115 1120 1125

Ile Ile Ser Leu Val Gln Asn Ala Pro Tyr Gly Leu Tyr Phe Ile
1130 1135 1140

His Phe Ser Tyr Val Pro Thr Lys Tyr Val Thr Ala Lys Val Ser
1145 1150 1155

Pro Gly Leu Cys Ile Ala Gly Asp Ile Gly Ile Ser Pro Lys Ser
1160 1165 1170

Gly Tyr Phe Ile Asn Val Asn Asn Ser Trp Met Phe Thr Gly Ser
1175 1180 1185

Ser Tyr Tyr Tyr Pro Glu Pro Ile Thr Gln Asn Asn Val Val Val
1190 1195 1200

Met Ser Thr Cys Ala Val Asn Tyr Thr Lys Ala Pro Asp Leu Met
1205 1210 1215

Leu Asn Thr Ser Thr Pro Asn Leu Pro Asp Phe Lys Glu Glu Leu
1220 1225 1230

Tyr Gln Trp Phe Lys Asn Gln Ser Ser Val Ala Pro Asp Leu Ser
1235 1240 1245

Leu Asp Tyr Ile Asn Val Thr Phe Leu Asp Leu Gln Asp Glu Met
1250 1255 1260

Asn Arg Leu Gln Glu Ala Ile Lys Val Leu Asn Gln Ser Tyr Ile
1265 1270 1275

Asn Leu Lys Asp Ile Gly Thr Tyr Glu Tyr Tyr Val Lys Trp Pro
1280 1285 1290

Trp Tyr Val Trp Leu Leu Ile Gly Leu Ala Gly Val Ala Met Leu
1295 1300 1305

Val Leu Leu Phe Phe Ile Cys Cys Cys Thr Gly Cys Gly Thr Ser
1310 1315 1320

Cys Phe Lys Lys Cys Gly Gly Cys Cys Asp Asp Tyr Thr Gly His
1325 1330 1335

Gln Glu Phe Val Ile Lys Thr Ser His Asp Asp
1340 1345

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<210> SEQ ID NO 61
<211> LENGTH: 1225
<212> TYPE: PRT
<213> ORGANISM: Porcine respiratory coronavirus

<400> SEQUENCE: 61

Met Lys Lys Leu Phe Val Val Leu Val Val Met Pro Leu Ile Tyr Gly
 1          5          10          15
Asp Lys Phe Pro Thr Ser Val Val Ser Asn Cys Thr Asp Gln Cys Ala
 20          25          30
Ser Tyr Val Ala Asn Val Phe Thr Thr Gln Pro Gly Gly Phe Ile Pro
 35          40          45
Ser Asp Phe Ser Phe Asn Asn Trp Phe Leu Leu Thr Asn Ser Ser Thr
 50          55          60
Leu Val Ser Gly Lys Leu Val Thr Lys Gln Pro Leu Leu Val Asn Cys
 65          70          75          80
Leu Trp Pro Val Pro Ser Phe Glu Glu Ala Ala Ser Thr Phe Cys Phe
 85          90          95
Glu Gly Ala Asp Phe Asp Gln Cys Asn Gly Ala Val Leu Asn Asn Thr
100          105          110
Val Asp Val Ile Arg Phe Asn Leu Asn Phe Thr Thr Asn Val Gln Ser
115          120          125
Gly Lys Gly Ala Thr Val Phe Ser Leu Asn Thr Thr Gly Gly Val Thr
130          135          140
Leu Glu Ile Ser Cys Tyr Asn Asp Thr Val Ser Asp Ser Ser Phe Ser
145          150          155          160
Ser Tyr Gly Glu Ile Pro Phe Gly Val Thr Asn Gly Pro Arg Tyr Cys
165          170          175
Tyr Val Leu Tyr Asn Gly Thr Ala Leu Lys Tyr Leu Gly Thr Leu Pro
180          185          190
Pro Ser Val Lys Glu Ile Ala Ile Ser Lys Trp Gly His Phe Tyr Ile
195          200          205
Asn Gly Tyr Asn Phe Phe Ser Thr Phe Pro Ile Asp Cys Ile Ser Phe
210          215          220
Asn Leu Thr Thr Gly Asp Ser Asp Val Phe Trp Thr Ile Ala Tyr Thr
225          230          235          240
Ser Tyr Thr Glu Ala Leu Val Gln Val Glu Asn Thr Ala Ile Thr Asn
245          250          255
Val Thr Tyr Cys Asn Ser Tyr Val Asn Asn Ile Lys Cys Ser Gln Leu
260          265          270
Thr Ala Asn Leu Asn Asn Gly Phe Tyr Pro Val Ser Ser Ser Glu Val
275          280          285
Gly Ser Val Asn Lys Ser Val Val Leu Leu Pro Ser Phe Leu Thr His
290          295          300
Thr Ile Val Asn Ile Thr Ile Gly Leu Gly Met Lys Arg Ser Gly Tyr
305          310          315          320
Gly Gln Pro Ile Ala Ser Thr Leu Ser Asn Ile Thr Leu Pro Met Gln
325          330          335
Asp Asn Asn Thr Asp Val Tyr Cys Val Arg Ser Asp Gln Phe Ser Val
340          345          350
Tyr Val His Ser Thr Cys Lys Ser Ala Leu Trp Asp Asn Val Phe Lys
355          360          365
Arg Asn Cys Thr Asp Val Leu Asp Ala Thr Ala Val Ile Lys Thr Gly
370          375          380

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Thr Cys Pro Phe Ser Phe Asp Lys Leu Asn Asn Tyr Leu Thr Phe Asn
 385 390 395 400
 Lys Phe Cys Leu Ser Leu Ser Pro Val Gly Ala Asn Cys Lys Phe Asp
 405 410 415
 Val Ala Ala Arg Thr Arg Thr Asn Glu Gln Val Val Arg Ser Leu Tyr
 420 425 430
 Val Ile Tyr Glu Glu Gly Asp Ser Ile Val Gly Val Pro Ser Asp Asn
 435 440 445
 Ser Gly Leu His Asp Leu Ser Val Leu His Leu Asp Ser Cys Thr Asp
 450 455 460
 Tyr Asn Ile Tyr Gly Arg Thr Gly Val Gly Ile Ile Arg Gln Thr Asn
 465 470 475 480
 Arg Thr Leu Leu Ser Gly Leu Tyr Tyr Thr Ser Leu Ser Gly Asp Leu
 485 490 495
 Leu Gly Phe Lys Asn Val Ser Asp Gly Val Ile Tyr Ser Val Thr Pro
 500 505 510
 Cys Asp Val Ser Ala Gln Ala Val Ile Asp Gly Thr Ile Val Gly
 515 520 525
 Ala Ile Thr Ser Ile Asn Ser Glu Leu Leu Gly Leu Thr His Trp Thr
 530 535 540
 Ile Thr Pro Asn Phe Tyr Tyr Tyr Ser Ile Tyr Asn Tyr Thr Asn Asp
 545 550 555 560
 Lys Thr Arg Gly Thr Pro Ile Asp Ser Asn Asp Val Gly Cys Glu Pro
 565 570 575
 Val Ile Thr Tyr Ser Asn Ile Gly Val Cys Lys Asn Gly Ala Leu Val
 580 585 590
 Phe Ile Asn Val Thr His Ser Asp Gly Asp Val Gln Pro Ile Ser Thr
 595 600 605
 Gly Asn Val Thr Ile Pro Thr Asn Phe Thr Ile Ser Val Gln Val Glu
 610 615 620
 Tyr Ile Gln Val Tyr Thr Thr Pro Val Ser Ile Asp Cys Ser Arg Tyr
 625 630 635 640
 Val Cys Asn Gly Asn Pro Arg Cys Asn Lys Leu Leu Thr Gln Tyr Val
 645 650 655
 Ser Ala Cys Gln Thr Ile Glu Gln Ala Leu Ala Met Gly Ala Arg Leu
 660 665 670
 Glu Asn Met Glu Val Asp Ser Met Leu Phe Val Ser Glu Asn Ala Leu
 675 680 685
 Lys Leu Ala Ser Val Glu Ala Phe Asn Ser Ser Glu Thr Leu Asp Pro
 690 695 700
 Ile Tyr Thr Gln Trp Pro Asn Ile Gly Gly Phe Trp Leu Glu Gly Leu
 705 710 715 720
 Lys Tyr Ile Leu Pro Ser Asp Asn Ser Lys Arg Lys Tyr Arg Ser Ala
 725 730 735
 Ile Glu Asp Leu Leu Phe Ser Lys Val Val Thr Ser Gly Leu Gly Thr
 740 745 750
 Val Asp Glu Asp Tyr Lys Arg Cys Thr Gly Gly Tyr Asp Ile Ala Asp
 755 760 765
 Leu Val Cys Ala Gln Tyr Tyr Asn Gly Ile Met Val Leu Pro Gly Val
 770 775 780
 Ala Asn Ala Asp Lys Met Thr Met Tyr Thr Ala Ser Leu Ala Gly Gly
 785 790 795 800

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Ile Thr Leu Gly Ala Phe Gly Gly Gly Ala Val Ser Ile Pro Phe Ala
805 810 815

Val Ala Val Gln Ala Arg Leu Asn Tyr Val Ala Leu Gln Thr Asp Val
820 825 830

Leu Asn Lys Asn Gln Gln Ile Leu Ala Ser Ala Phe Asn Gln Ala Ile
835 840 845

Gly Asn Ile Thr Gln Ser Phe Gly Lys Val Asn Asp Ala Ile His Gln
850 855 860

Thr Ser Arg Gly Leu Thr Thr Val Ala Lys Ala Leu Ala Lys Val Gln
865 870 875 880

Asp Val Val Asn Thr Gln Gly Gln Ala Leu Arg His Leu Thr Val Gln
885 890 895

Leu Gln Asn Asn Phe Gln Ala Ile Ser Ser Ser Ile Ser Asp Ile Tyr
900 905 910

Asn Arg Leu Asp Glu Leu Ser Ala Asp Ala Gln Val Asp Arg Leu Ile
915 920 925

Thr Gly Arg Leu Thr Ala Leu Asn Ala Phe Val Ser Gln Thr Leu Thr
930 935 940

Arg Gln Ala Glu Val Arg Ala Ser Arg Gln Leu Ala Lys Asp Lys Val
945 950 955 960

Asn Glu Cys Val Arg Ser Gln Ser Gln Arg Phe Gly Phe Cys Gly Asn
965 970 975

Gly Thr His Leu Phe Ser Leu Ala Asn Ala Ala Pro Asn Gly Met Ile
980 985 990

Phe Phe His Thr Val Leu Leu Pro Thr Ala Tyr Glu Thr Val Thr Ala
995 1000 1005

Trp Ser Gly Ile Cys Ala Leu Asp Gly Asp Arg Thr Phe Gly Leu
1010 1015 1020

Val Val Lys Asp Val Gln Leu Thr Leu Phe Arg Asn Leu Asp Asp
1025 1030 1035

Lys Phe Tyr Leu Thr Pro Arg Thr Met Tyr Gln Pro Arg Val Ala
1040 1045 1050

Thr Ser Ser Asp Phe Val Gln Ile Glu Gly Cys Asp Val Leu Phe
1055 1060 1065

Val Asn Thr Thr Val Ser Asp Leu Pro Ser Ile Ile Pro Asp Tyr
1070 1075 1080

Ile Asp Ile Asn Gln Thr Val Gln Asp Ile Leu Glu Asn Phe Arg
1085 1090 1095

Pro Asn Trp Thr Val Pro Glu Leu Thr Leu Asp Val Phe Asn Ala
1100 1105 1110

Thr Tyr Leu Asn Leu Thr Gly Glu Ile Asp Asp Leu Glu Phe Arg
1115 1120 1125

Ser Glu Lys Leu His Asn Thr Thr Val Glu Leu Ala Ile Leu Ile
1130 1135 1140

Asp Asn Ile Asn Asn Thr Leu Val Asn Leu Glu Trp Leu Asn Arg
1145 1150 1155

Ile Glu Thr Tyr Val Lys Trp Pro Trp Tyr Val Trp Leu Leu Ile
1160 1165 1170

Gly Leu Val Val Ile Phe Cys Ile Pro Leu Leu Leu Phe Cys Cys
1175 1180 1185

Cys Ser Thr Gly Cys Cys Gly Cys Ile Gly Cys Leu Gly Ser Cys
1190 1195 1200

Cys His Ser Ile Phe Ser Arg Arg Gln Phe Glu Asn Tyr Glu Pro

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1205		1210		1215
Ile Glu Lys Val His Val His				
1220		1225		

<210> SEQ ID NO 62
 <211> LENGTH: 82
 <212> TYPE: PRT
 <213> ORGANISM: Porcine transmissible gastroenteritis coronavirus

<400> SEQUENCE: 62

Met Thr Phe Pro Arg Ala Leu Thr Val Ile Asp Asp Asn Gly Met Val				
1	5	10	15	
Ile Asn Ile Ile Phe Trp Phe Leu Leu Ile Ile Ile Leu Ile Leu Leu				
20	25	30		
Ser Ile Ala Leu Leu Asn Ile Ile Lys Leu Cys Met Val Val Cys Cys Asn				
35	40	45		
Leu Gly Arg Thr Val Ile Ile Val Pro Ala Gln His Ala Tyr Asp Ala				
50	55	60		
Tyr Lys Asn Phe Met Arg Ile Lys Ala Tyr Asn Pro Asp Gly Ala Leu				
65	70	75	80	

Leu Ala

<210> SEQ ID NO 63
 <211> LENGTH: 4376
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 63

Met Glu Ser Leu Val Leu Gly Val Asn Glu Lys Thr His Val Gln Leu				
1	5	10	15	
Ser Leu Pro Val Leu Gln Val Arg Asp Val Leu Val Arg Gly Phe Gly				
20	25	30		
Asp Ser Val Glu Glu Ala Leu Ser Glu Ala Arg Glu His Leu Lys Asn				
35	40	45		
Gly Thr Cys Gly Leu Val Glu Leu Glu Lys Gly Val Leu Pro Gln Leu				
50	55	60		
Glu Gln Pro Tyr Val Phe Ile Lys Arg Ser Asp Ala Leu Ser Thr Asn				
65	70	75	80	
His Gly His Lys Val Val Glu Leu Val Ala Glu Met Asp Gly Ile Gln				
85	90	95		
Tyr Gly Arg Ser Gly Ile Thr Leu Gly Val Leu Val Pro His Val Gly				
100	105	110		
Glu Thr Pro Ile Ala Tyr Arg Asn Val Leu Leu Arg Lys Asn Gly Asn				
115	120	125		
Lys Gly Ala Gly Gly His Ser Tyr Gly Ile Asp Leu Lys Ser Tyr Asp				
130	135	140		
Leu Gly Asp Glu Leu Gly Thr Asp Pro Ile Glu Asp Tyr Glu Gln Asn				
145	150	155	160	
Trp Asn Thr Lys His Gly Ser Gly Ala Leu Arg Glu Leu Thr Arg Glu				
165	170	175		
Leu Asn Gly Gly Ala Val Thr Arg Tyr Val Asp Asn Asn Phe Cys Gly				
180	185	190		
Pro Asp Gly Tyr Pro Leu Asp Cys Ile Lys Asp Phe Leu Ala Arg Ala				
195	200	205		
Gly Lys Ser Met Cys Thr Leu Ser Glu Gln Leu Asp Tyr Ile Glu Ser				
210	215	220		

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Lys Arg Gly Val Tyr Cys Cys Arg Asp His Glu His Glu Ile Ala Trp
 225 230 235 240
 Phe Thr Glu Arg Ser Asp Lys Ser Tyr Glu His Gln Thr Pro Phe Glu
 245 250 255
 Ile Lys Ser Ala Lys Lys Phe Asp Thr Phe Lys Gly Glu Cys Pro Lys
 260 265 270
 Phe Val Phe Pro Leu Asn Ser Lys Val Lys Val Ile Gln Pro Arg Val
 275 280 285
 Glu Lys Lys Lys Thr Glu Gly Phe Met Gly Arg Ile Arg Ser Val Tyr
 290 295 300
 Pro Val Ala Ser Pro Gln Glu Cys Asn Asn Met His Leu Ser Thr Leu
 305 310 315 320
 Met Lys Cys Asn His Cys Asp Glu Val Ser Trp Gln Thr Cys Asp Phe
 325 330 335
 Leu Lys Ala Thr Cys Glu His Cys Gly Thr Glu Asn Leu Val Ile Glu
 340 345 350
 Gly Pro Thr Thr Cys Gly Tyr Leu Pro Thr Asn Ala Val Val Lys Met
 355 360 365
 Pro Cys Pro Ala Cys Gln Asp Pro Glu Ile Gly Pro Glu His Ser Val
 370 375 380
 Ala Asp Tyr His Asn His Ser Asn Ile Glu Thr Arg Leu Arg Lys Gly
 385 390 395 400
 Gly Arg Thr Arg Cys Phe Gly Gly Cys Val Phe Ala Tyr Val Gly Cys
 405 410 415
 Tyr Asn Lys Arg Ala Tyr Trp Val Pro Arg Ala Ser Ala Asp Ile Gly
 420 425 430
 Ser Gly His Thr Gly Ile Thr Gly Asp Asn Val Glu Thr Leu Asn Glu
 435 440 445
 Asp Leu Leu Glu Ile Leu Ser Arg Glu Arg Val Asn Ile Asn Ile Val
 450 455 460
 Gly Asp Phe His Leu Asn Glu Glu Val Ala Ile Ile Leu Ala Ser Phe
 465 470 475 480
 Ser Ala Ser Thr Ser Ala Phe Ile Asp Thr Ile Lys Ser Leu Asp Tyr
 485 490 495
 Lys Ser Phe Lys Thr Ile Val Glu Ser Cys Gly Asn Tyr Lys Val Thr
 500 505 510
 Lys Gly Lys Pro Val Lys Gly Ala Trp Asn Ile Gly Gln Gln Arg Ser
 515 520 525
 Val Leu Thr Pro Leu Cys Gly Phe Pro Ser Gln Ala Ala Gly Val Ile
 530 535 540
 Arg Ser Ile Phe Ala Arg Thr Leu Asp Ala Ala Asn His Ser Ile Pro
 545 550 555 560
 Asp Leu Gln Arg Ala Ala Val Thr Ile Leu Asp Gly Ile Ser Glu Gln
 565 570 575
 Ser Leu Arg Leu Val Asp Ala Met Val Tyr Thr Ser Asp Leu Leu Thr
 580 585 590
 Asn Ser Val Ile Ile Met Ala Tyr Val Thr Gly Gly Leu Val Gln Gln
 595 600 605
 Thr Ser Gln Trp Leu Ser Asn Leu Leu Gly Thr Thr Val Glu Lys Leu
 610 615 620
 Arg Pro Ile Phe Glu Trp Ile Glu Ala Lys Leu Ser Ala Gly Val Glu
 625 630 635 640

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Phe Leu Lys Asp Ala Trp Glu Ile Leu Lys Phe Leu Ile Thr Gly Val
645 650 655
Phe Asp Ile Val Lys Gly Gln Ile Gln Val Ala Ser Asp Asn Ile Lys
660 665 670
Asp Cys Val Lys Cys Phe Ile Asp Val Val Asn Lys Ala Leu Glu Met
675 680 685
Cys Ile Asp Gln Val Thr Ile Ala Gly Ala Lys Leu Arg Ser Leu Asn
690 695 700
Leu Gly Glu Val Phe Ile Ala Gln Ser Lys Gly Leu Tyr Arg Gln Cys
705 710 715 720
Ile Arg Gly Lys Glu Gln Leu Gln Leu Leu Met Pro Leu Lys Ala Pro
725 730 735
Lys Glu Val Thr Phe Leu Glu Gly Asp Ser His Asp Thr Val Leu Thr
740 745 750
Ser Glu Glu Val Val Leu Lys Asn Gly Glu Leu Glu Ala Leu Glu Thr
755 760 765
Pro Val Asp Ser Phe Thr Asn Gly Ala Ile Val Gly Thr Pro Val Cys
770 775 780
Val Asn Gly Leu Met Leu Leu Glu Ile Lys Asp Lys Glu Gln Tyr Cys
785 790 795 800
Ala Leu Ser Pro Gly Leu Leu Ala Thr Asn Asn Val Phe Arg Leu Lys
805 810 815
Gly Gly Ala Pro Ile Lys Gly Val Thr Phe Gly Glu Asp Thr Val Trp
820 825 830
Glu Val Gln Gly Tyr Lys Asn Val Arg Ile Thr Phe Glu Leu Asp Glu
835 840 845
Arg Val Asp Lys Val Leu Asn Glu Lys Cys Ser Val Tyr Thr Val Glu
850 855 860
Ser Gly Thr Glu Val Thr Glu Phe Ala Cys Val Val Ala Glu Ala Val
865 870 875 880
Val Lys Thr Leu Gln Pro Val Ser Asp Leu Leu Thr Asn Met Gly Ile
885 890 895
Asp Leu Asp Glu Trp Ser Val Ala Thr Phe Tyr Leu Phe Asp Asp Ala
900 905 910
Gly Glu Glu Asn Phe Ser Ser Arg Met Tyr Cys Ser Phe Tyr Pro Pro
915 920 925
Asp Glu Glu Glu Glu Asp Asp Ala Glu Cys Glu Glu Glu Glu Ile Asp
930 935 940
Glu Thr Cys Glu His Glu Tyr Gly Thr Glu Asp Asp Tyr Gln Gly Leu
945 950 955 960
Pro Leu Glu Phe Gly Ala Ser Ala Glu Thr Val Arg Val Glu Glu Glu
965 970 975
Glu Glu Glu Asp Trp Leu Asp Asp Thr Thr Glu Gln Ser Glu Ile Glu
980 985 990
Pro Glu Pro Glu Pro Thr Pro Glu Glu Pro Val Asn Gln Phe Thr Gly
995 1000 1005
Tyr Leu Lys Leu Thr Asp Asn Val Ala Ile Lys Cys Val Asp Ile
1010 1015 1020
Val Lys Glu Ala Gln Ser Ala Asn Pro Met Val Ile Val Asn Ala
1025 1030 1035
Ala Asn Ile His Leu Lys His Gly Gly Gly Val Ala Gly Ala Leu
1040 1045 1050
Asn Lys Ala Thr Asn Gly Ala Met Gln Lys Glu Ser Asp Asp Tyr

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1055	1060	1065
Ile Lys Leu Asn Gly Pro 1070	Leu Thr Val Gly Gly 1075	Ser Cys Leu Leu 1080
Ser Gly His Asn Leu Ala 1085	Lys Lys Cys Leu His 1090	Val Val Gly Pro 1095
Asn Leu Asn Ala Gly Glu 1100	Asp Ile Gln Leu Leu 1105	Lys Ala Ala Tyr 1110
Glu Asn Phe Asn Ser Gln 1115	Asp Ile Leu Leu Ala 1120	Pro Leu Leu Ser 1125
Ala Gly Ile Phe Gly Ala 1130	Lys Pro Leu Gln Ser 1135	Leu Gln Val Cys 1140
Val Gln Thr Val Arg Thr 1145	Gln Val Tyr Ile Ala 1150	Val Asn Asp Lys 1155
Ala Leu Tyr Glu Gln Val 1160	Val Met Asp Tyr Leu 1165	Asp Asn Leu Lys 1170
Pro Arg Val Glu Ala Pro 1175	Lys Gln Glu Glu Pro 1180	Pro Asn Thr Glu 1185
Asp Ser Lys Thr Glu Glu 1190	Lys Ser Val Val Gln 1195	Lys Pro Val Asp 1200
Val Lys Pro Lys Ile Lys 1205	Ala Cys Ile Asp Glu 1210	Val Thr Thr Thr 1215
Leu Glu Glu Thr Lys Phe 1220	Leu Thr Asn Lys Leu 1225	Leu Leu Phe Ala 1230
Asp Ile Asn Gly Lys Leu 1235	Tyr His Asp Ser Gln 1240	Asn Met Leu Arg 1245
Gly Glu Asp Met Ser Phe 1250	Leu Glu Lys Asp Ala 1255	Pro Tyr Met Val 1260
Gly Asp Val Ile Thr Ser 1265	Gly Asp Ile Thr Cys 1270	Val Val Ile Pro 1275
Ser Lys Lys Ala Gly Gly 1280	Thr Thr Glu Met Leu 1285	Ser Arg Ala Leu 1290
Lys Lys Val Pro Val Asp 1295	Glu Tyr Ile Thr Thr 1300	Tyr Pro Gly Gln 1305
Gly Cys Ala Gly Tyr Thr 1310	Leu Glu Glu Ala Lys 1315	Thr Ala Leu Lys 1320
Lys Cys Lys Ser Ala Phe 1325	Tyr Val Leu Pro Ser 1330	Glu Ala Pro Asn 1335
Ala Lys Glu Glu Ile Leu 1340	Gly Thr Val Ser Trp 1345	Asn Leu Arg Glu 1350
Met Leu Ala His Ala Glu 1355	Glu Thr Arg Lys Leu 1360	Met Pro Ile Cys 1365
Met Asp Val Arg Ala Ile 1370	Met Ala Thr Ile Gln 1375	Arg Lys Tyr Lys 1380
Gly Ile Lys Ile Gln Glu 1385	Gly Ile Val Asp Tyr 1390	Gly Val Arg Phe 1395
Phe Phe Tyr Thr Ser Lys 1400	Glu Pro Val Ala Ser 1405	Ile Ile Thr Lys 1410
Leu Asn Ser Leu Asn Glu 1415	Pro Leu Val Thr Met 1420	Pro Ile Gly Tyr 1425
Val Thr His Gly Phe Asn 1430	Leu Glu Glu Ala Ala 1435	Arg Cys Met Arg 1440
Ser Leu Lys Ala Pro Ala 1445	Val Val Ser Val Ser 1450	Ser Pro Asp Ala 1455

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Val	Thr	Thr	Tyr	Asn	Gly	Tyr	Leu	Thr	Ser	Ser	Ser	Lys	Thr	Ser
1460						1465						1470		
Glu	Glu	His	Phe	Val	Glu	Thr	Val	Ser	Leu	Ala	Gly	Ser	Tyr	Arg
1475						1480						1485		
Asp	Trp	Ser	Tyr	Ser	Gly	Gln	Arg	Thr	Glu	Leu	Gly	Val	Glu	Phe
1490						1495						1500		
Leu	Lys	Arg	Gly	Asp	Lys	Ile	Val	Tyr	His	Thr	Leu	Glu	Ser	Pro
1505						1510						1515		
Val	Glu	Phe	His	Leu	Asp	Gly	Glu	Val	Leu	Ser	Leu	Asp	Lys	Leu
1520						1525						1530		
Lys	Ser	Leu	Leu	Ser	Leu	Arg	Glu	Val	Lys	Thr	Ile	Lys	Val	Phe
1535						1540						1545		
Thr	Thr	Val	Asp	Asn	Thr	Asn	Leu	His	Thr	Gln	Leu	Val	Asp	Met
1550						1555						1560		
Ser	Met	Thr	Tyr	Gly	Gln	Gln	Phe	Gly	Pro	Thr	Tyr	Leu	Asp	Gly
1565						1570						1575		
Ala	Asp	Val	Thr	Lys	Ile	Lys	Pro	His	Val	Asn	His	Glu	Gly	Lys
1580						1585						1590		
Thr	Phe	Phe	Val	Leu	Pro	Ser	Asp	Asp	Thr	Leu	Arg	Ser	Glu	Ala
1595						1600						1605		
Phe	Glu	Tyr	Tyr	His	Thr	Leu	Asp	Glu	Ser	Phe	Leu	Gly	Arg	Tyr
1610						1615						1620		
Met	Ser	Ala	Leu	Asn	His	Thr	Lys	Lys	Trp	Lys	Phe	Pro	Gln	Val
1625						1630						1635		
Gly	Gly	Leu	Thr	Ser	Ile	Lys	Trp	Ala	Asp	Asn	Asn	Cys	Tyr	Leu
1640						1645						1650		
Ser	Ser	Val	Leu	Leu	Ala	Leu	Gln	Gln	Leu	Glu	Val	Lys	Phe	Asn
1655						1660						1665		
Ala	Pro	Ala	Leu	Gln	Glu	Ala	Tyr	Tyr	Arg	Ala	Arg	Ala	Gly	Asp
1670						1675						1680		
Ala	Ala	Asn	Phe	Cys	Ala	Leu	Ile	Leu	Ala	Tyr	Ser	Asn	Lys	Thr
1685						1690						1695		
Val	Gly	Glu	Leu	Gly	Asp	Val	Arg	Glu	Thr	Met	Thr	His	Leu	Leu
1700						1705						1710		
Gln	His	Ala	Asn	Leu	Glu	Ser	Ala	Lys	Arg	Val	Leu	Asn	Val	Val
1715						1720						1725		
Cys	Lys	His	Cys	Gly	Gln	Lys	Thr	Thr	Thr	Leu	Thr	Gly	Val	Glu
1730						1735						1740		
Ala	Val	Met	Tyr	Met	Gly	Thr	Leu	Ser	Tyr	Asp	Asn	Leu	Lys	Thr
1745						1750						1755		
Gly	Val	Ser	Ile	Pro	Cys	Val	Cys	Gly	Arg	Asp	Ala	Thr	Gln	Tyr
1760						1765						1770		
Leu	Val	Gln	Gln	Glu	Ser	Ser	Phe	Val	Met	Met	Ser	Ala	Pro	Pro
1775						1780						1785		
Ala	Glu	Tyr	Lys	Leu	Gln	Gln	Gly	Thr	Phe	Leu	Cys	Ala	Asn	Glu
1790						1795						1800		
Tyr	Thr	Gly	Asn	Tyr	Gln	Cys	Gly	His	Tyr	Thr	His	Ile	Thr	Ala
1805						1810						1815		
Lys	Glu	Thr	Leu	Tyr	Arg	Ile	Asp	Gly	Ala	His	Leu	Thr	Lys	Met
1820						1825						1830		
Ser	Glu	Tyr	Lys	Gly	Pro	Val	Thr	Asp	Val	Phe	Tyr	Lys	Glu	Thr
1835						1840						1845		

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Ser	Tyr	Thr	Thr	Thr	Ile	Lys	Pro	Val	Ser	Tyr	Lys	Leu	Asp	Gly
1850						1855					1860			
Val	Thr	Tyr	Thr	Glu	Ile	Glu	Pro	Lys	Leu	Asp	Gly	Tyr	Tyr	Lys
1865						1870					1875			
Lys	Asp	Asn	Ala	Tyr	Tyr	Thr	Glu	Gln	Pro	Ile	Asp	Leu	Val	Pro
1880						1885					1890			
Thr	Gln	Pro	Leu	Pro	Asn	Ala	Ser	Phe	Asp	Asn	Phe	Lys	Leu	Thr
1895						1900					1905			
Cys	Ser	Asn	Thr	Lys	Phe	Ala	Asp	Asp	Leu	Asn	Gln	Met	Thr	Gly
1910						1915					1920			
Phe	Thr	Lys	Pro	Ala	Ser	Arg	Glu	Leu	Ser	Val	Thr	Phe	Phe	Pro
1925						1930					1935			
Asp	Leu	Asn	Gly	Asp	Val	Val	Ala	Ile	Asp	Tyr	Arg	His	Tyr	Ser
1940						1945					1950			
Ala	Ser	Phe	Lys	Lys	Gly	Ala	Lys	Leu	Leu	His	Lys	Pro	Ile	Val
1955						1960					1965			
Trp	His	Ile	Asn	Gln	Ala	Thr	Thr	Lys	Thr	Thr	Phe	Lys	Pro	Asn
1970						1975					1980			
Thr	Trp	Cys	Leu	Arg	Cys	Leu	Trp	Ser	Thr	Lys	Pro	Val	Asp	Thr
1985						1990					1995			
Ser	Asn	Ser	Phe	Glu	Val	Leu	Ala	Val	Glu	Asp	Thr	Gln	Gly	Met
2000						2005					2010			
Asp	Asn	Leu	Ala	Cys	Glu	Ser	Gln	Gln	Pro	Thr	Ser	Glu	Glu	Val
2015						2020					2025			
Val	Glu	Asn	Pro	Thr	Ile	Gln	Lys	Glu	Val	Ile	Glu	Cys	Asp	Val
2030						2035					2040			
Lys	Thr	Thr	Glu	Val	Val	Gly	Asn	Val	Ile	Leu	Lys	Pro	Ser	Asp
2045						2050					2055			
Glu	Gly	Val	Lys	Val	Thr	Gln	Glu	Leu	Gly	His	Glu	Asp	Leu	Met
2060						2065					2070			
Ala	Ala	Tyr	Val	Glu	Asn	Thr	Ser	Ile	Thr	Ile	Lys	Lys	Pro	Asn
2075						2080					2085			
Glu	Leu	Ser	Leu	Ala	Leu	Gly	Leu	Lys	Thr	Ile	Ala	Thr	His	Gly
2090						2095					2100			
Ile	Ala	Ala	Ile	Asn	Ser	Val	Pro	Trp	Ser	Lys	Ile	Leu	Ala	Tyr
2105						2110					2115			
Val	Lys	Pro	Phe	Leu	Gly	Gln	Ala	Ala	Ile	Thr	Thr	Ser	Asn	Cys
2120						2125					2130			
Ala	Lys	Arg	Leu	Ala	Gln	Arg	Val	Phe	Asn	Asn	Tyr	Met	Pro	Tyr
2135						2140					2145			
Val	Phe	Thr	Leu	Leu	Phe	Gln	Leu	Cys	Thr	Phe	Thr	Lys	Ser	Thr
2150						2155					2160			
Asn	Ser	Arg	Ile	Arg	Ala	Ser	Leu	Pro	Thr	Thr	Ile	Ala	Lys	Asn
2165						2170					2175			
Ser	Val	Lys	Ser	Val	Ala	Lys	Leu	Cys	Leu	Asp	Ala	Gly	Ile	Asn
2180						2185					2190			
Tyr	Val	Lys	Ser	Pro	Lys	Phe	Ser	Lys	Leu	Phe	Thr	Ile	Ala	Met
2195						2200					2205			
Trp	Leu	Leu	Leu	Leu	Ser	Ile	Cys	Leu	Gly	Ser	Leu	Ile	Cys	Val
2210						2215					2220			
Thr	Ala	Ala	Phe	Gly	Val	Leu	Leu	Ser	Asn	Phe	Gly	Ala	Pro	Ser
2225						2230					2235			
Tyr	Cys	Asn	Gly	Val	Arg	Glu	Leu	Tyr	Leu	Asn	Ser	Ser	Asn	Val

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Gly Asp	Ser Cys Asn Asn Phe	Met Leu Thr Tyr Asn	Lys Val Glu
2645	2650	2655	
Asn Met	Thr Pro Arg Asp Leu	Gly Ala Cys Ile Asp	Cys Asn Ala
2660	2665	2670	
Arg His	Ile Asn Ala Gln Val	Ala Lys Ser His Asn	Val Ser Leu
2675	2680	2685	
Ile Trp	Asn Val Lys Asp Tyr	Met Ser Leu Ser Glu	Gln Leu Arg
2690	2695	2700	
Lys Gln	Ile Arg Ser Ala Ala	Lys Lys Asn Asn Ile	Pro Phe Arg
2705	2710	2715	
Leu Thr	Cys Ala Thr Thr Arg	Gln Val Val Asn Val	Ile Thr Thr
2720	2725	2730	
Lys Ile	Ser Leu Lys Gly Gly	Lys Ile Val Ser Thr	Cys Phe Lys
2735	2740	2745	
Leu Met	Leu Lys Ala Thr Leu	Leu Cys Val Leu Ala	Ala Leu Val
2750	2755	2760	
Cys Tyr	Ile Val Met Pro Val	His Thr Leu Ser Ile	His Asp Gly
2765	2770	2775	
Tyr Thr	Asn Glu Ile Ile Gly	Tyr Lys Ala Ile Gln	Asp Gly Val
2780	2785	2790	
Thr Arg	Asp Ile Ile Ser Thr	Asp Asp Cys Phe Ala	Asn Lys His
2795	2800	2805	
Ala Gly	Phe Asp Ala Trp Phe	Ser Gln Arg Gly Gly	Ser Tyr Lys
2810	2815	2820	
Asn Asp	Lys Ser Cys Pro Val	Val Ala Ala Ile Ile	Thr Arg Glu
2825	2830	2835	
Ile Gly	Phe Ile Val Pro Gly	Leu Pro Gly Thr Val	Leu Arg Ala
2840	2845	2850	
Ile Asn	Gly Asp Phe Leu His	Phe Leu Pro Arg Val	Phe Ser Ala
2855	2860	2865	
Val Gly	Asn Ile Cys Tyr Thr	Pro Ser Lys Leu Ile	Glu Tyr Ser
2870	2875	2880	
Asp Phe	Ala Thr Ser Ala Cys	Val Leu Ala Ala Glu	Cys Thr Ile
2885	2890	2895	
Phe Lys	Asp Ala Met Gly Lys	Pro Val Pro Tyr Cys	Tyr Asp Thr
2900	2905	2910	
Asn Leu	Leu Glu Gly Ser Ile	Ser Tyr Ser Glu Leu	Arg Pro Asp
2915	2920	2925	
Thr Arg	Tyr Val Leu Met Asp	Gly Ser Ile Ile Gln	Phe Pro Asn
2930	2935	2940	
Thr Tyr	Leu Glu Gly Ser Val	Arg Val Val Thr Thr	Phe Asp Ala
2945	2950	2955	
Glu Tyr	Cys Arg His Gly Thr	Cys Glu Arg Ser Glu	Val Gly Ile
2960	2965	2970	
Cys Leu	Ser Thr Ser Gly Arg	Trp Val Leu Asn Asn	Glu His Tyr
2975	2980	2985	
Arg Ala	Leu Ser Gly Val Phe	Cys Gly Val Asp Ala	Met Asn Leu
2990	2995	3000	
Ile Ala	Asn Ile Phe Thr Pro	Leu Val Gln Pro Val	Gly Ala Leu
3005	3010	3015	
Asp Val	Ser Ala Ser Val Val	Ala Gly Gly Ile Ile	Ala Ile Leu
3020	3025	3030	

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Val Thr	Cys Ala Ala Tyr Tyr	Phe Met Lys Phe Arg	Arg Val Phe
3035	3040	3045	
Gly Glu	Tyr Asn His Val Val	Ala Ala Asn Ala Leu	Leu Phe Leu
3050	3055	3060	
Met Ser	Phe Thr Ile Leu Cys	Leu Val Pro Ala Tyr	Ser Phe Leu
3065	3070	3075	
Pro Gly	Val Tyr Ser Val Phe	Tyr Leu Tyr Leu Thr	Phe Tyr Phe
3080	3085	3090	
Thr Asn	Asp Val Ser Phe Leu	Ala His Leu Gln Trp	Phe Ala Met
3095	3100	3105	
Phe Ser	Pro Ile Val Pro Phe	Trp Ile Thr Ala Ile	Tyr Val Phe
3110	3115	3120	
Cys Ile	Ser Leu Lys His Cys	His Trp Phe Phe Asn	Asn Tyr Leu
3125	3130	3135	
Arg Lys	Arg Val Met Phe Asn	Gly Val Thr Phe Ser	Thr Phe Glu
3140	3145	3150	
Glu Ala	Ala Leu Cys Thr Phe	Leu Leu Asn Lys Glu	Met Tyr Leu
3155	3160	3165	
Lys Leu	Arg Ser Glu Thr Leu	Leu Pro Leu Thr Gln	Tyr Asn Arg
3170	3175	3180	
Tyr Leu	Ala Leu Tyr Asn Lys	Tyr Lys Tyr Phe Ser	Gly Ala Leu
3185	3190	3195	
Asp Thr	Thr Ser Tyr Arg Glu	Ala Ala Cys Cys His	Leu Ala Lys
3200	3205	3210	
Ala Leu	Asn Asp Phe Ser Asn	Ser Gly Ala Asp Val	Leu Tyr Gln
3215	3220	3225	
Pro Pro	Gln Thr Ser Ile Thr	Ser Ala Val Leu Gln	Ser Gly Phe
3230	3235	3240	
Arg Lys	Met Ala Phe Pro Ser	Gly Lys Val Glu Gly	Cys Met Val
3245	3250	3255	
Gln Val	Thr Cys Gly Thr Thr	Thr Leu Asn Gly Leu	Trp Leu Asp
3260	3265	3270	
Asp Thr	Val Tyr Cys Pro Arg	His Val Ile Cys Thr	Ala Glu Asp
3275	3280	3285	
Met Leu	Asn Pro Asn Tyr Glu	Asp Leu Leu Ile Arg	Lys Ser Asn
3290	3295	3300	
His Ser	Phe Leu Val Gln Ala	Gly Asn Val Gln Leu	Arg Val Ile
3305	3310	3315	
Gly His	Ser Met Gln Asn Cys	Leu Leu Arg Leu Lys	Val Asp Thr
3320	3325	3330	
Ser Asn	Pro Lys Thr Pro Lys	Tyr Lys Phe Val Arg	Ile Gln Pro
3335	3340	3345	
Gly Gln	Thr Phe Ser Val Leu	Ala Cys Tyr Asn Gly	Ser Pro Ser
3350	3355	3360	
Gly Val	Tyr Gln Cys Ala Met	Arg Pro Asn His Thr	Ile Lys Gly
3365	3370	3375	
Ser Phe	Leu Asn Gly Ser Cys	Gly Ser Val Gly Phe	Asn Ile Asp
3380	3385	3390	
Tyr Asp	Cys Val Ser Phe Cys	Tyr Met His His Met	Glu Leu Pro
3395	3400	3405	
Thr Gly	Val His Ala Gly Thr	Asp Leu Glu Gly Lys	Phe Tyr Gly
3410	3415	3420	
Pro Phe	Val Asp Arg Gln Thr	Ala Gln Ala Ala Gly	Thr Asp Thr

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3425	3430	3435
Thr Ile Thr Leu Asn Val Leu Ala Trp Leu Tyr Ala Ala Val Ile 3440 3445 3450		
Asn Gly Asp Arg Trp Phe Leu Asn Arg Phe Thr Thr Thr Leu Asn 3455 3460 3465		
Asp Phe Asn Leu Val Ala Met Lys Tyr Asn Tyr Glu Pro Leu Thr 3470 3475 3480		
Gln Asp His Val Asp Ile Leu Gly Pro Leu Ser Ala Gln Thr Gly 3485 3490 3495		
Ile Ala Val Leu Asp Met Cys Ala Ala Leu Lys Glu Leu Leu Gln 3500 3505 3510		
Asn Gly Met Asn Gly Arg Thr Ile Leu Gly Ser Thr Ile Leu Glu 3515 3520 3525		
Asp Glu Phe Thr Pro Phe Asp Val Val Arg Gln Cys Ser Gly Val 3530 3535 3540		
Thr Phe Gln Gly Lys Phe Lys Lys Ile Val Lys Gly Thr His His 3545 3550 3555		
Trp Met Leu Leu Thr Phe Leu Thr Ser Leu Leu Ile Leu Val Gln 3560 3565 3570		
Ser Thr Gln Trp Ser Leu Phe Phe Phe Val Tyr Glu Asn Ala Phe 3575 3580 3585		
Leu Pro Phe Thr Leu Gly Ile Met Ala Ile Ala Ala Cys Ala Met 3590 3595 3600		
Leu Leu Val Lys His Lys His Ala Phe Leu Cys Leu Phe Leu Leu 3605 3610 3615		
Pro Ser Leu Ala Thr Val Ala Tyr Phe Asn Met Val Tyr Met Pro 3620 3625 3630		
Ala Ser Trp Val Met Arg Ile Met Thr Trp Leu Glu Leu Ala Asp 3635 3640 3645		
Thr Ser Leu Ser Gly Tyr Arg Leu Lys Asp Cys Val Met Tyr Ala 3650 3655 3660		
Ser Ala Leu Val Leu Leu Ile Leu Met Thr Ala Arg Thr Val Tyr 3665 3670 3675		
Asp Asp Ala Ala Arg Arg Val Trp Thr Leu Met Asn Val Ile Thr 3680 3685 3690		
Leu Val Tyr Lys Val Tyr Tyr Gly Asn Ala Leu Asp Gln Ala Ile 3695 3700 3705		
Ser Met Trp Ala Leu Val Ile Ser Val Thr Ser Asn Tyr Ser Gly 3710 3715 3720		
Val Val Thr Thr Ile Met Phe Leu Ala Arg Ala Ile Val Phe Val 3725 3730 3735		
Cys Val Glu Tyr Tyr Pro Leu Leu Phe Ile Thr Gly Asn Thr Leu 3740 3745 3750		
Gln Cys Ile Met Leu Val Tyr Cys Phe Leu Gly Tyr Cys Cys Cys 3755 3760 3765		
Cys Tyr Phe Gly Leu Phe Cys Leu Leu Asn Arg Tyr Phe Arg Leu 3770 3775 3780		
Thr Leu Gly Val Tyr Asp Tyr Leu Val Ser Thr Gln Glu Phe Arg 3785 3790 3795		
Tyr Met Asn Ser Gln Gly Leu Leu Pro Pro Lys Ser Ser Ile Asp 3800 3805 3810		
Ala Phe Lys Leu Asn Ile Lys Leu Leu Gly Ile Gly Gly Lys Pro 3815 3820 3825		

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Cys Ile Lys Val Ala Thr Val Gln Ser Lys Met Ser Asp Val Lys
 3830 3835 3840
 Cys Thr Ser Val Val Leu Leu Ser Val Leu Gln Gln Leu Arg Val
 3845 3850 3855
 Glu Ser Ser Ser Lys Leu Trp Ala Gln Cys Val Gln Leu His Asn
 3860 3865 3870
 Asp Ile Leu Leu Ala Lys Asp Thr Thr Glu Ala Phe Glu Lys Met
 3875 3880 3885
 Val Ser Leu Leu Ser Val Leu Leu Ser Met Gln Gly Ala Val Asp
 3890 3895 3900
 Ile Asn Arg Leu Cys Glu Glu Met Leu Asp Asn Arg Ala Thr Leu
 3905 3910 3915
 Gln Ala Ile Ala Ser Glu Phe Ser Ser Leu Pro Ser Tyr Ala Ala
 3920 3925 3930
 Tyr Ala Thr Ala Gln Glu Ala Tyr Glu Gln Ala Val Ala Asn Gly
 3935 3940 3945
 Asp Ser Glu Val Val Leu Lys Lys Leu Lys Lys Ser Leu Asn Val
 3950 3955 3960
 Ala Lys Ser Glu Phe Asp Arg Asp Ala Ala Met Gln Arg Lys Leu
 3965 3970 3975
 Glu Lys Met Ala Asp Gln Ala Met Thr Gln Met Tyr Lys Gln Ala
 3980 3985 3990
 Arg Ser Glu Asp Lys Arg Ala Lys Val Thr Ser Ala Met Gln Thr
 3995 4000 4005
 Met Leu Phe Thr Met Leu Arg Lys Leu Asp Asn Asp Ala Leu Asn
 4010 4015 4020
 Asn Ile Ile Asn Asn Ala Arg Asp Gly Cys Val Pro Leu Asn Ile
 4025 4030 4035
 Ile Pro Leu Thr Thr Ala Ala Lys Leu Met Val Val Val Pro Asp
 4040 4045 4050
 Tyr Gly Thr Tyr Lys Asn Thr Cys Asp Gly Asn Thr Phe Thr Tyr
 4055 4060 4065
 Ala Ser Ala Leu Trp Glu Ile Gln Gln Val Val Asp Ala Asp Ser
 4070 4075 4080
 Lys Ile Val Gln Leu Ser Glu Ile Asn Met Asp Asn Ser Pro Asn
 4085 4090 4095
 Leu Ala Trp Pro Leu Ile Val Thr Ala Leu Arg Ala Asn Ser Ala
 4100 4105 4110
 Val Lys Leu Gln Asn Asn Glu Leu Ser Pro Val Ala Leu Arg Gln
 4115 4120 4125
 Met Ser Cys Ala Ala Gly Thr Thr Gln Thr Ala Cys Thr Asp Asp
 4130 4135 4140
 Asn Ala Leu Ala Tyr Tyr Asn Asn Ser Lys Gly Gly Arg Phe Val
 4145 4150 4155
 Leu Ala Leu Leu Ser Asp His Gln Asp Leu Lys Trp Ala Arg Phe
 4160 4165 4170
 Pro Lys Ser Asp Gly Thr Gly Thr Ile Tyr Thr Glu Leu Glu Pro
 4175 4180 4185
 Pro Cys Arg Phe Val Thr Asp Thr Pro Lys Gly Pro Lys Val Lys
 4190 4195 4200
 Tyr Leu Tyr Phe Ile Lys Gly Leu Asn Asn Leu Asn Arg Gly Met
 4205 4210 4215

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Val Leu Gly Ser Leu Ala Ala Thr Val Arg Leu Gln Ala Gly Asn
 4220 4225 4230

Ala Thr Glu Val Pro Ala Asn Ser Thr Val Leu Ser Phe Cys Ala
 4235 4240 4245

Phe Ala Val Asp Pro Ala Lys Ala Tyr Lys Asp Tyr Leu Ala Ser
 4250 4255 4260

Gly Gly Gln Pro Ile Thr Asn Cys Val Lys Met Leu Cys Thr His
 4265 4270 4275

Thr Gly Thr Gly Gln Ala Ile Thr Val Thr Pro Glu Ala Asn Met
 4280 4285 4290

Asp Gln Glu Ser Phe Gly Gly Ala Ser Cys Cys Leu Tyr Cys Arg
 4295 4300 4305

Cys His Ile Asp His Pro Asn Pro Lys Gly Phe Cys Asp Leu Lys
 4310 4315 4320

Gly Lys Tyr Val Gln Ile Pro Thr Thr Cys Ala Asn Asp Pro Val
 4325 4330 4335

Gly Phe Thr Leu Arg Asn Thr Val Cys Thr Val Cys Gly Met Trp
 4340 4345 4350

Lys Gly Tyr Gly Cys Ser Cys Asp Gln Leu Arg Glu Pro Leu Met
 4355 4360 4365

Gln Ser Ala Asp Ala Ser Thr Phe
 4370 4375

<210> SEQ ID NO 64

<211> LENGTH: 2697

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 64

Phe Lys Arg Val Cys Gly Val Ser Ala Ala Arg Leu Thr Pro Cys Gly
 1 5 10 15

Thr Gly Thr Ser Thr Asp Val Val Tyr Arg Ala Phe Asp Ile Tyr Asn
 20 25 30

Glu Lys Val Ala Gly Phe Ala Lys Phe Leu Lys Thr Asn Cys Cys Arg
 35 40 45

Phe Gln Glu Lys Asp Glu Glu Gly Asn Leu Leu Asp Ser Tyr Phe Val
 50 55 60

Val Lys Arg His Thr Met Ser Asn Tyr Gln His Glu Glu Thr Ile Tyr
 65 70 75 80

Asn Leu Val Lys Asp Cys Pro Ala Val Ala Val His Asp Phe Phe Lys
 85 90 95

Phe Arg Val Asp Gly Asp Met Val Pro His Ile Ser Arg Gln Arg Leu
 100 105 110

Thr Lys Tyr Thr Met Ala Asp Leu Val Tyr Ala Leu Arg His Phe Asp
 115 120 125

Glu Gly Asn Cys Asp Thr Leu Lys Glu Ile Leu Val Thr Tyr Asn Cys
 130 135 140

Cys Asp Asp Asp Tyr Phe Asn Lys Lys Asp Trp Tyr Asp Phe Val Glu
 145 150 155 160

Asn Pro Asp Ile Leu Arg Val Tyr Ala Asn Leu Gly Glu Arg Val Arg
 165 170 175

Gln Ser Leu Leu Lys Thr Val Gln Phe Cys Asp Ala Met Arg Asp Ala
 180 185 190

Gly Ile Val Gly Val Leu Thr Leu Asp Asn Gln Asp Leu Asn Gly Asn
 195 200 205

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Trp Tyr Asp Phe Gly Asp Phe Val Gln Val Ala Pro Gly Cys Gly Val
 210 215 220
 Pro Ile Val Asp Ser Tyr Tyr Ser Leu Leu Met Pro Ile Leu Thr Leu
 225 230 235 240
 Thr Arg Ala Leu Ala Ala Glu Ser His Met Asp Ala Asp Leu Ala Lys
 245 250 255
 Pro Leu Ile Lys Trp Asp Leu Leu Lys Tyr Asp Phe Thr Glu Glu Arg
 260 265 270
 Leu Cys Leu Phe Asp Arg Tyr Phe Lys Tyr Trp Asp Gln Thr Tyr His
 275 280 285
 Pro Asn Cys Ile Asn Cys Leu Asp Asp Arg Cys Ile Leu His Cys Ala
 290 295 300
 Asn Phe Asn Val Leu Phe Ser Thr Val Phe Pro Thr Ser Phe Gly
 305 310 315 320
 Pro Leu Val Arg Lys Ile Phe Val Asp Gly Val Pro Phe Val Val Ser
 325 330 335
 Thr Gly Tyr His Phe Arg Glu Leu Gly Val Val His Asn Gln Asp Val
 340 345 350
 Asn Leu His Ser Ser Arg Leu Ser Phe Lys Glu Leu Leu Val Tyr Ala
 355 360 365
 Ala Asp Pro Ala Met His Ala Ala Ser Gly Asn Leu Leu Leu Asp Lys
 370 375 380
 Arg Thr Thr Cys Phe Ser Val Ala Ala Leu Thr Asn Asn Val Ala Phe
 385 390 395 400
 Gln Thr Val Lys Pro Gly Asn Phe Asn Lys Asp Phe Tyr Asp Phe Ala
 405 410 415
 Val Ser Lys Gly Phe Phe Lys Glu Gly Ser Ser Val Glu Leu Lys His
 420 425 430
 Phe Phe Phe Ala Gln Asp Gly Asn Ala Ala Ile Ser Asp Tyr Asp Tyr
 435 440 445
 Tyr Arg Tyr Asn Leu Pro Thr Met Cys Asp Ile Arg Gln Leu Leu Phe
 450 455 460
 Val Val Glu Val Val Asp Lys Tyr Phe Asp Cys Tyr Asp Gly Gly Cys
 465 470 475 480
 Ile Asn Ala Asn Gln Val Ile Val Asn Asn Leu Asp Lys Ser Ala Gly
 485 490 495
 Phe Pro Phe Asn Lys Trp Gly Lys Ala Arg Leu Tyr Tyr Asp Ser Met
 500 505 510
 Ser Tyr Glu Asp Gln Asp Ala Leu Phe Ala Tyr Thr Lys Arg Asn Val
 515 520 525
 Ile Pro Thr Ile Thr Gln Met Asn Leu Lys Tyr Ala Ile Ser Ala Lys
 530 535 540
 Asn Arg Ala Arg Thr Val Ala Gly Val Ser Ile Cys Ser Thr Met Thr
 545 550 555 560
 Asn Arg Gln Phe His Gln Lys Leu Leu Lys Ser Ile Ala Ala Thr Arg
 565 570 575
 Gly Ala Thr Val Val Ile Gly Thr Ser Lys Phe Tyr Gly Gly Trp His
 580 585 590
 Asn Met Leu Lys Thr Val Tyr Ser Asp Val Glu Thr Pro His Leu Met
 595 600 605
 Gly Trp Asp Tyr Pro Lys Cys Asp Arg Ala Met Pro Asn Met Leu Arg
 610 615 620

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Ile	Met	Ala	Ser	Leu	Val	Leu	Ala	Arg	Lys	His	Asn	Thr	Cys	Cys	Asn	625	630	635	640
Leu	Ser	His	Arg	Phe	Tyr	Arg	Leu	Ala	Asn	Glu	Cys	Ala	Gln	Val	Leu	645	650	655	
Ser	Glu	Met	Val	Met	Cys	Gly	Gly	Ser	Leu	Tyr	Val	Lys	Pro	Gly	Gly	660	665	670	
Thr	Ser	Ser	Gly	Asp	Ala	Thr	Thr	Ala	Tyr	Ala	Asn	Ser	Val	Phe	Asn	675	680	685	
Ile	Cys	Gln	Ala	Val	Thr	Ala	Asn	Val	Asn	Ala	Leu	Leu	Ser	Thr	Asp	690	695	700	
Gly	Asn	Lys	Ile	Ala	Asp	Lys	Tyr	Val	Arg	Asn	Leu	Gln	His	Arg	Leu	705	710	715	720
Tyr	Glu	Cys	Leu	Tyr	Arg	Asn	Arg	Asp	Val	Asp	His	Glu	Phe	Val	Asp	725	730	735	
Glu	Phe	Tyr	Ala	Tyr	Leu	Arg	Lys	His	Phe	Ser	Met	Met	Ile	Leu	Ser	740	745	750	
Asp	Asp	Ala	Val	Val	Cys	Tyr	Asn	Ser	Asn	Tyr	Ala	Ala	Gln	Gly	Leu	755	760	765	
Val	Ala	Ser	Ile	Lys	Asn	Phe	Lys	Ala	Val	Leu	Tyr	Tyr	Gln	Asn	Asn	770	775	780	
Val	Phe	Met	Ser	Glu	Ala	Lys	Cys	Trp	Thr	Glu	Thr	Asp	Leu	Thr	Lys	785	790	795	800
Gly	Pro	His	Glu	Phe	Cys	Ser	Gln	His	Thr	Met	Leu	Val	Lys	Gln	Gly	805	810	815	
Asp	Asp	Tyr	Val	Tyr	Leu	Pro	Tyr	Pro	Asp	Pro	Ser	Arg	Ile	Leu	Gly	820	825	830	
Ala	Gly	Cys	Phe	Val	Asp	Asp	Ile	Val	Lys	Thr	Asp	Gly	Thr	Leu	Met	835	840	845	
Ile	Glu	Arg	Phe	Val	Ser	Leu	Ala	Ile	Asp	Ala	Tyr	Pro	Leu	Thr	Lys	850	855	860	
His	Pro	Asn	Gln	Glu	Tyr	Ala	Asp	Val	Phe	His	Leu	Tyr	Leu	Gln	Tyr	865	870	875	880
Ile	Arg	Lys	Leu	His	Asp	Glu	Leu	Thr	Gly	His	Met	Leu	Asp	Met	Tyr	885	890	895	
Ser	Val	Met	Leu	Thr	Asn	Asp	Asn	Thr	Ser	Arg	Tyr	Trp	Glu	Pro	Glu	900	905	910	
Phe	Tyr	Glu	Ala	Met	Tyr	Thr	Pro	His	Thr	Val	Leu	Gln	Ala	Val	Gly	915	920	925	
Ala	Cys	Val	Leu	Cys	Asn	Ser	Gln	Thr	Ser	Leu	Arg	Cys	Gly	Ala	Cys	930	935	940	
Ile	Arg	Arg	Pro	Phe	Leu	Cys	Cys	Lys	Cys	Cys	Tyr	Asp	His	Val	Ile	945	950	955	960
Ser	Thr	Ser	His	Lys	Leu	Val	Leu	Ser	Val	Asn	Pro	Tyr	Val	Cys	Asn	965	970	975	
Ala	Pro	Gly	Cys	Asp	Val	Thr	Asp	Val	Thr	Gln	Leu	Tyr	Leu	Gly	Gly	980	985	990	
Met	Ser	Tyr	Tyr	Cys	Lys	Ser	His	Lys	Pro	Pro	Ile	Ser	Phe	Pro	Leu	995	1000	1005	
Cys	Ala	Asn	Gly	Gln	Val	Phe	Gly	Leu	Tyr	Lys	Asn	Thr	Cys	Val	1010	1015	1020		
Gly	Ser	Asp	Asn	Val	Thr	Asp	Phe	Asn	Ala	Ile	Ala	Thr	Cys	Asp	1025	1030	1035		
Trp	Thr	Asn	Ala	Gly	Asp	Tyr	Ile	Leu	Ala	Asn	Thr	Cys	Thr	Glu					

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1040	1045	1050
Arg Leu Lys Leu Phe Ala Ala 1055	Glu Thr Leu Lys Ala 1060	Thr Glu Glu 1065
Thr Phe Lys Leu Ser Tyr Gly 1070	Ile Ala Thr Val Arg 1075	Glu Val Leu 1080
Ser Asp Arg Glu Leu His Leu 1085	Ser Trp Glu Val Gly 1090	Lys Pro Arg 1095
Pro Pro Leu Asn Arg Asn Tyr 1100	Val Phe Thr Gly Tyr 1105	Arg Val Thr 1110
Lys Asn Ser Lys Val Gln Ile 1115	Gly Glu Tyr Thr Phe 1120	Glu Lys Gly 1125
Asp Tyr Gly Asp Ala Val Val 1130	Tyr Arg Gly Thr Thr 1135	Thr Tyr Lys 1140
Leu Asn Val Gly Asp Tyr Phe 1145	Val Leu Thr Ser His 1150	Thr Val Met 1155
Pro Leu Ser Ala Pro Thr Leu 1160	Val Pro Gln Glu His 1165	Tyr Val Arg 1170
Ile Thr Gly Leu Tyr Pro Thr 1175	Leu Asn Ile Ser Asp 1180	Glu Phe Ser 1185
Ser Asn Val Ala Asn Tyr Gln 1190	Lys Val Gly Met Gln 1195	Lys Tyr Ser 1200
Thr Leu Gln Gly Pro Pro Gly 1205	Thr Gly Lys Ser His 1210	Phe Ala Ile 1215
Gly Leu Ala Leu Tyr Tyr Pro 1220	Ser Ala Arg Ile Val 1225	Tyr Thr Ala 1230
Cys Ser His Ala Ala Val Asp 1235	Ala Leu Cys Glu Lys 1240	Ala Leu Lys 1245
Tyr Leu Pro Ile Asp Lys Cys 1250	Ser Arg Ile Ile Pro 1255	Ala Arg Ala 1260
Arg Val Glu Cys Phe Asp Lys 1265	Phe Lys Val Asn Ser 1270	Thr Leu Glu 1275
Gln Tyr Val Phe Cys Thr Val 1280	Asn Ala Leu Pro Glu 1285	Thr Thr Ala 1290
Asp Ile Val Val Phe Asp Glu 1295	Ile Ser Met Ala Thr 1300	Asn Tyr Asp 1305
Leu Ser Val Val Asn Ala Arg 1310	Leu Arg Ala Lys His 1315	Tyr Val Tyr 1320
Ile Gly Asp Pro Ala Gln Leu 1325	Pro Ala Pro Arg Thr 1330	Leu Leu Thr 1335
Lys Gly Thr Leu Glu Pro Glu 1340	Tyr Phe Asn Ser Val 1345	Cys Arg Leu 1350
Met Lys Thr Ile Gly Pro Asp 1355	Met Phe Leu Gly Thr 1360	Cys Arg Arg 1365
Cys Pro Ala Glu Ile Val Asp 1370	Thr Val Ser Ala Leu 1375	Val Tyr Asp 1380
Asn Lys Leu Lys Ala His Lys 1385	Asp Lys Ser Ala Gln 1390	Cys Phe Lys 1395
Met Phe Tyr Lys Gly Val Ile 1400	Thr His Asp Val Ser 1405	Ser Ala Ile 1410
Asn Arg Pro Gln Ile Gly Val 1415	Val Arg Glu Phe Leu 1420	Thr Arg Asn 1425
Pro Ala Trp Arg Lys Ala Val 1430	Phe Ile Ser Pro Tyr 1435	Asn Ser Gln 1440

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Asn	Ala	Val	Ala	Ser	Lys	Ile	Leu	Gly	Leu	Pro	Thr	Gln	Thr	Val
1445						1450						1455		
Asp	Ser	Ser	Gln	Gly	Ser	Glu	Tyr	Asp	Tyr	Val	Ile	Phe	Thr	Gln
1460						1465						1470		
Thr	Thr	Glu	Thr	Ala	His	Ser	Cys	Asn	Val	Asn	Arg	Phe	Asn	Val
1475						1480						1485		
Ala	Ile	Thr	Arg	Ala	Lys	Ile	Gly	Ile	Leu	Cys	Ile	Met	Ser	Asp
1490						1495						1500		
Arg	Asp	Leu	Tyr	Asp	Lys	Leu	Gln	Phe	Thr	Ser	Leu	Glu	Ile	Pro
1505						1510						1515		
Arg	Arg	Asn	Val	Ala	Thr	Leu	Gln	Ala	Glu	Asn	Val	Thr	Gly	Leu
1520						1525						1530		
Phe	Lys	Asp	Cys	Ser	Lys	Ile	Ile	Thr	Gly	Leu	His	Pro	Thr	Gln
1535						1540						1545		
Ala	Pro	Thr	His	Leu	Ser	Val	Asp	Ile	Lys	Phe	Lys	Thr	Glu	Gly
1550						1555						1560		
Leu	Cys	Val	Asp	Ile	Pro	Gly	Ile	Pro	Lys	Asp	Met	Thr	Tyr	Arg
1565						1570						1575		
Arg	Leu	Ile	Ser	Met	Met	Gly	Phe	Lys	Met	Asn	Tyr	Gln	Val	Asn
1580						1585						1590		
Gly	Tyr	Pro	Asn	Met	Phe	Ile	Thr	Arg	Glu	Glu	Ala	Ile	Arg	His
1595						1600						1605		
Val	Arg	Ala	Trp	Ile	Gly	Phe	Asp	Val	Glu	Gly	Cys	His	Ala	Thr
1610						1615						1620		
Arg	Asp	Ala	Val	Gly	Thr	Asn	Leu	Pro	Leu	Gln	Leu	Gly	Phe	Ser
1625						1630						1635		
Thr	Gly	Val	Asn	Leu	Val	Ala	Val	Pro	Thr	Gly	Tyr	Val	Asp	Thr
1640						1645						1650		
Glu	Asn	Asn	Thr	Glu	Phe	Thr	Arg	Val	Asn	Ala	Lys	Pro	Pro	Pro
1655						1660						1665		
Gly	Asp	Gln	Phe	Lys	His	Leu	Ile	Pro	Leu	Met	Tyr	Lys	Gly	Leu
1670						1675						1680		
Pro	Trp	Asn	Val	Val	Arg	Ile	Lys	Ile	Val	Gln	Met	Leu	Ser	Asp
1685						1690						1695		
Thr	Leu	Lys	Gly	Leu	Ser	Asp	Arg	Val	Val	Phe	Val	Leu	Trp	Ala
1700						1705						1710		
His	Gly	Phe	Glu	Leu	Thr	Ser	Met	Lys	Tyr	Phe	Val	Lys	Ile	Gly
1715						1720						1725		
Pro	Glu	Arg	Thr	Cys	Cys	Leu	Cys	Asp	Lys	Arg	Ala	Thr	Cys	Phe
1730						1735						1740		
Ser	Thr	Ser	Ser	Asp	Thr	Tyr	Ala	Cys	Trp	Asn	His	Ser	Val	Gly
1745						1750						1755		
Phe	Asp	Tyr	Val	Tyr	Asn	Pro	Phe	Met	Ile	Asp	Val	Gln	Gln	Trp
1760						1765						1770		
Gly	Phe	Thr	Gly	Asn	Leu	Gln	Ser	Asn	His	Asp	Gln	His	Cys	Gln
1775						1780						1785		
Val	His	Gly	Asn	Ala	His	Val	Ala	Ser	Cys	Asp	Ala	Ile	Met	Thr
1790						1795						1800		
Arg	Cys	Leu	Ala	Val	His	Glu	Cys	Phe	Val	Lys	Arg	Val	Asp	Trp
1805						1810						1815		
Ser	Val	Glu	Tyr	Pro	Ile	Ile	Gly	Asp	Glu	Leu	Arg	Val	Asn	Ser
1820						1825						1830		

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Ala Cys Arg Lys Val Gln His Met Val Val Lys Ser Ala Leu Leu 1835 1840 1845
Ala Asp Lys Phe Pro Val Leu His Asp Ile Gly Asn Pro Lys Ala 1850 1855 1860
Ile Lys Cys Val Pro Gln Ala Glu Val Glu Trp Lys Phe Tyr Asp 1865 1870 1875
Ala Gln Pro Cys Ser Asp Lys Ala Tyr Lys Ile Glu Glu Leu Phe 1880 1885 1890
Tyr Ser Tyr Ala Thr His His Asp Lys Phe Thr Asp Gly Val Cys 1895 1900 1905
Leu Phe Trp Asn Cys Asn Val Asp Arg Tyr Pro Ala Asn Ala Ile 1910 1915 1920
Val Cys Arg Phe Asp Thr Arg Val Leu Ser Asn Leu Asn Leu Pro 1925 1930 1935
Gly Cys Asp Gly Gly Ser Leu Tyr Val Asn Lys His Ala Phe His 1940 1945 1950
Thr Pro Ala Phe Asp Lys Ser Ala Phe Thr Asn Leu Lys Gln Leu 1955 1960 1965
Pro Phe Phe Tyr Tyr Ser Asp Ser Pro Cys Glu Ser His Gly Lys 1970 1975 1980
Gln Val Val Ser Asp Ile Asp Tyr Val Pro Leu Lys Ser Ala Thr 1985 1990 1995
Cys Ile Thr Arg Cys Asn Leu Gly Gly Ala Val Cys Arg His His 2000 2005 2010
Ala Asn Glu Tyr Arg Gln Tyr Leu Asp Ala Tyr Asn Met Met Ile 2015 2020 2025
Ser Ala Gly Phe Ser Leu Trp Ile Tyr Lys Gln Phe Asp Thr Tyr 2030 2035 2040
Asn Leu Trp Asn Thr Phe Thr Arg Leu Gln Ser Leu Glu Asn Val 2045 2050 2055
Ala Tyr Asn Val Val Asn Lys Gly His Phe Asp Gly His Ala Gly 2060 2065 2070
Glu Ala Pro Val Ser Ile Ile Asn Asn Ala Val Tyr Thr Lys Val 2075 2080 2085
Asp Gly Ile Asp Val Glu Ile Phe Glu Asn Lys Thr Thr Leu Pro 2090 2095 2100
Val Asn Val Ala Phe Glu Leu Trp Ala Lys Arg Asn Ile Lys Pro 2105 2110 2115
Val Pro Glu Ile Lys Ile Leu Asn Asn Leu Gly Val Asp Ile Ala 2120 2125 2130
Ala Asn Thr Val Ile Trp Asp Tyr Lys Arg Glu Ala Pro Ala His 2135 2140 2145
Val Ser Thr Ile Gly Val Cys Thr Met Thr Asp Ile Ala Lys Lys 2150 2155 2160
Pro Thr Glu Ser Ala Cys Ser Ser Leu Thr Val Leu Phe Asp Gly 2165 2170 2175
Arg Val Glu Gly Gln Val Asp Leu Phe Arg Asn Ala Arg Asn Gly 2180 2185 2190
Val Leu Ile Thr Glu Gly Ser Val Lys Gly Leu Thr Pro Ser Lys 2195 2200 2205
Gly Pro Ala Gln Ala Ser Val Asn Gly Val Thr Leu Ile Gly Glu 2210 2215 2220
Ser Val Lys Thr Gln Phe Asn Tyr Phe Lys Lys Val Asp Gly Ile

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2225	2230	2235
Ile Gln Gln Leu Pro Glu Thr Tyr Phe Thr Gln Ser Arg Asp Leu 2240 2245 2250		
Glu Asp Phe Lys Pro Arg Ser Gln Met Glu Thr Asp Phe Leu Glu 2255 2260 2265		
Leu Ala Met Asp Glu Phe Ile Gln Arg Tyr Lys Leu Glu Gly Tyr 2270 2275 2280		
Ala Phe Glu His Ile Val Tyr Gly Asp Phe Ser His Gly Gln Leu 2285 2290 2295		
Gly Gly Leu His Leu Met Ile Gly Leu Ala Lys Arg Ser Gln Asp 2300 2305 2310		
Ser Pro Leu Lys Leu Glu Asp Phe Ile Pro Met Asp Ser Thr Val 2315 2320 2325		
Lys Asn Tyr Phe Ile Thr Asp Ala Gln Thr Gly Ser Ser Lys Cys 2330 2335 2340		
Val Cys Ser Val Ile Asp Leu Leu Leu Asp Asp Phe Val Glu Ile 2345 2350 2355		
Ile Lys Ser Gln Asp Leu Ser Val Ile Ser Lys Val Val Lys Val 2360 2365 2370		
Thr Ile Asp Tyr Ala Glu Ile Ser Phe Met Leu Trp Cys Lys Asp 2375 2380 2385		
Gly His Val Glu Thr Phe Tyr Pro Lys Leu Gln Ala Ser Gln Ala 2390 2395 2400		
Trp Gln Pro Gly Val Ala Met Pro Asn Leu Tyr Lys Met Gln Arg 2405 2410 2415		
Met Leu Leu Glu Lys Cys Asp Leu Gln Asn Tyr Gly Glu Asn Ala 2420 2425 2430		
Val Ile Pro Lys Gly Ile Met Met Asn Val Ala Lys Tyr Thr Gln 2435 2440 2445		
Leu Cys Gln Tyr Leu Asn Thr Leu Thr Leu Ala Val Pro Tyr Asn 2450 2455 2460		
Met Arg Val Ile His Phe Gly Ala Gly Ser Asp Lys Gly Val Ala 2465 2470 2475		
Pro Gly Thr Ala Val Leu Arg Gln Trp Leu Pro Thr Gly Thr Leu 2480 2485 2490		
Leu Val Asp Ser Asp Leu Asn Asp Phe Val Ser Asp Ala Asp Ser 2495 2500 2505		
Thr Leu Ile Gly Asp Cys Ala Thr Val His Thr Ala Asn Lys Trp 2510 2515 2520		
Asp Leu Ile Ile Ser Asp Met Tyr Asp Pro Arg Thr Lys His Val 2525 2530 2535		
Thr Lys Glu Asn Asp Ser Lys Glu Gly Phe Phe Thr Tyr Leu Cys 2540 2545 2550		
Gly Phe Ile Lys Gln Lys Leu Ala Leu Gly Gly Ser Ile Ala Val 2555 2560 2565		
Lys Ile Thr Glu His Ser Trp Asn Ala Asp Leu Tyr Lys Leu Met 2570 2575 2580		
Gly His Phe Ser Trp Trp Thr Ala Phe Val Thr Asn Val Asn Ala 2585 2590 2595		
Ser Ser Ser Glu Ala Phe Leu Ile Gly Ala Asn Tyr Leu Gly Lys 2600 2605 2610		
Pro Lys Glu Gln Ile Asp Gly Tyr Thr Met His Ala Asn Tyr Ile 2615 2620 2625		

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Phe Trp Arg Asn Thr Asn Pro Ile Gln Leu Ser Ser Tyr Ser Leu
 2630 2635 2640

Phe Asp Met Ser Lys Phe Pro Leu Lys Leu Arg Gly Thr Ala Val
 2645 2650 2655

Met Ser Leu Lys Glu Asn Gln Ile Asn Asp Met Ile Tyr Ser Leu
 2660 2665 2670

Leu Glu Lys Gly Arg Leu Ile Ile Arg Glu Asn Asn Arg Val Val
 2675 2680 2685

Val Ser Ser Asp Ile Leu Val Asn Asn
 2690 2695

<210> SEQ ID NO 65
 <211> LENGTH: 274
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 65

Met Asp Leu Phe Met Arg Phe Phe Thr Leu Arg Ser Ile Thr Ala Gln
 1 5 10 15

Pro Val Lys Ile Asp Asn Ala Ser Pro Ala Ser Thr Val His Ala Thr
 20 25 30

Ala Thr Ile Pro Leu Gln Ala Ser Leu Pro Phe Gly Trp Leu Val Ile
 35 40 45

Gly Val Ala Phe Leu Ala Val Phe Gln Ser Ala Thr Lys Ile Ile Ala
 50 55 60

Leu Asn Lys Arg Trp Gln Leu Ala Leu Tyr Lys Gly Phe Gln Phe Ile
 65 70 75 80

Cys Asn Leu Leu Leu Leu Phe Val Thr Ile Tyr Ser His Leu Leu Leu
 85 90 95

Val Ala Ala Gly Met Glu Ala Gln Phe Leu Tyr Leu Tyr Ala Leu Ile
 100 105 110

Tyr Phe Leu Gln Cys Ile Asn Ala Cys Arg Ile Ile Met Arg Cys Trp
 115 120 125

Leu Cys Trp Lys Cys Lys Ser Lys Asn Pro Leu Leu Tyr Asp Ala Asn
 130 135 140

Tyr Phe Val Cys Trp His Thr His Asn Tyr Asp Tyr Cys Ile Pro Tyr
 145 150 155 160

Asn Ser Val Thr Asp Thr Ile Val Val Thr Glu Gly Asp Gly Ile Ser
 165 170 175

Thr Pro Lys Leu Lys Glu Asp Tyr Gln Ile Gly Gly Tyr Ser Glu Asp
 180 185 190

Arg His Ser Gly Val Lys Asp Tyr Val Val Val His Gly Tyr Phe Thr
 195 200 205

Glu Val Tyr Tyr Gln Leu Glu Ser Thr Gln Ile Thr Thr Asp Thr Gly
 210 215 220

Ile Glu Asn Ala Thr Phe Phe Ile Phe Asn Lys Leu Val Lys Asp Pro
 225 230 235 240

Pro Asn Val Gln Ile His Thr Ile Asp Gly Ser Ser Gly Val Ala Asn
 245 250 255

Pro Ala Met Asp Pro Ile Tyr Asp Glu Pro Thr Thr Thr Ser Val
 260 265 270

Pro Leu

<210> SEQ ID NO 66

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<211> LENGTH: 154
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 66

Met Met Pro Thr Thr Leu Phe Ala Gly Thr His Ile Thr Met Thr Thr
 1 5 10 15
 Val Tyr His Ile Thr Val Ser Gln Ile Gln Leu Ser Leu Leu Lys Val
 20 25 30
 Thr Ala Phe Gln His Gln Asn Ser Lys Lys Thr Thr Lys Leu Val Val
 35 40 45
 Ile Leu Arg Ile Gly Thr Gln Val Leu Lys Thr Met Ser Leu Tyr Met
 50 55 60
 Ala Ile Ser Pro Lys Phe Thr Thr Ser Leu Ser Leu His Lys Leu Leu
 65 70 75 80
 Gln Thr Leu Val Leu Lys Met Leu His Ser Ser Ser Leu Thr Ser Leu
 85 90 95
 Leu Lys Thr His Arg Met Cys Lys Tyr Thr Gln Ser Thr Ala Leu Gln
 100 105 110
 Glu Leu Leu Ile Gln Gln Trp Ile Gln Phe Met Met Ser Arg Arg Arg
 115 120 125
 Leu Leu Ala Cys Leu Cys Lys His Lys Lys Val Ser Thr Asn Leu Cys
 130 135 140
 Thr His Ser Phe Arg Lys Lys Gln Val Arg
 145 150

<210> SEQ ID NO 67
 <211> LENGTH: 63
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 67

Met Phe His Leu Val Asp Phe Gln Val Thr Ile Ala Glu Ile Leu Ile
 1 5 10 15
 Ile Ile Met Arg Thr Phe Arg Ile Ala Ile Trp Asn Leu Asp Val Ile
 20 25 30
 Ile Ser Ser Ile Val Arg Gln Leu Phe Lys Pro Leu Thr Lys Lys Asn
 35 40 45
 Tyr Ser Glu Leu Asp Asp Glu Glu Pro Met Glu Leu Asp Tyr Pro
 50 55 60

<210> SEQ ID NO 68
 <211> LENGTH: 122
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 68

Met Lys Ile Ile Leu Phe Leu Thr Leu Ile Val Phe Thr Ser Cys Glu
 1 5 10 15
 Leu Tyr His Tyr Gln Glu Cys Val Arg Gly Thr Thr Val Leu Leu Lys
 20 25 30
 Glu Pro Cys Pro Ser Gly Thr Tyr Glu Gly Asn Ser Pro Phe His Pro
 35 40 45
 Leu Ala Asp Asn Lys Phe Ala Leu Thr Cys Thr Ser Thr His Phe Ala
 50 55 60
 Phe Ala Cys Ala Asp Gly Thr Arg His Thr Tyr Gln Leu Arg Ala Arg
 65 70 75 80

-continued

Asp Pro Gln Ile Gln Leu Thr Ile Thr Arg Met Glu Asp Ala Met Gly
 20 25 30
 Gln Gly Gln Asn Ser Ala Asp Pro Lys Val Tyr Pro Ile Ile Leu Arg
 35 40 45
 Leu Gly Ser Gln Leu Ser Leu Ser Met Ala Arg Arg Asn Leu Asp Ser
 50 55 60
 Leu Glu Ala Arg Ala Phe Gln Ser Thr Pro Ile Val Val Gln Met Thr
 65 70 75 80
 Lys Leu Ala Thr Thr Glu Glu Leu Pro Asp Glu Phe Val Val Val Thr
 85 90 95
 Ala Lys

<210> SEQ ID NO 73
 <211> LENGTH: 70
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 73

Met Leu Pro Pro Cys Tyr Asn Phe Leu Lys Glu Gln His Cys Gln Lys
 1 5 10 15
 Ala Ser Thr Gln Arg Glu Ala Glu Ala Val Lys Pro Leu Leu Ala
 20 25 30
 Pro His His Val Val Ala Val Ile Gln Glu Ile Gln Leu Leu Ala Ala
 35 40 45
 Val Gly Glu Ile Leu Leu Leu Glu Trp Leu Ala Glu Val Val Lys Leu
 50 55 60
 Pro Ser Arg Tyr Cys Cys
 65 70

<210> SEQ ID NO 74
 <211> LENGTH: 6
 <212> TYPE: RNA
 <213> ORGANISM: Coronavirus

<400> SEQUENCE: 74

cuaaac

6

<210> SEQ ID NO 75
 <211> LENGTH: 13
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 75

Met Phe Ile Phe Leu Leu Phe Leu Thr Leu Thr Ser Gly
 1 5 10

<210> SEQ ID NO 76
 <211> LENGTH: 23
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 76

Thr Ile Pro Leu Gln Ala Ser Leu Pro Phe Gly Trp Leu Val Ile Gly
 1 5 10 15
 Val Ala Phe Leu Ala Val Phe
 20

<210> SEQ ID NO 77
 <211> LENGTH: 23
 <212> TYPE: PRT

-continued

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 77

Phe Gln Phe Ile Cys Asn Leu Leu Leu Leu Phe Val Thr Ile Tyr Ser
 1 5 10 15
 His Leu Leu Leu Val Ala Ala
 20

<210> SEQ ID NO 78

<211> LENGTH: 23

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 78

Ala Gln Phe Leu Tyr Leu Tyr Ala Leu Ile Tyr Phe Leu Gln Cys Ile
 1 5 10 15
 Asn Ala Cys Arg Ile Ile Met
 20

<210> SEQ ID NO 79

<211> LENGTH: 18

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 79

Val Leu Leu Phe Leu Ala Phe Val Val Phe Leu Leu Val Thr Leu Ala
 1 5 10 15
 Ile Leu

<210> SEQ ID NO 80

<211> LENGTH: 23

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 80

Leu Leu Glu Gln Trp Asn Leu Val Ile Gly Phe Leu Phe Leu Ala Trp
 1 5 10 15
 Ile Met Leu Leu Gln Phe Ala
 20

<210> SEQ ID NO 81

<211> LENGTH: 23

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 81

Leu Val Phe Leu Trp Leu Leu Trp Pro Val Thr Leu Ala Cys Phe Val
 1 5 10 15
 Leu Ala Ala Val Tyr Arg Ile
 20

<210> SEQ ID NO 82

<211> LENGTH: 23

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 82

Gly Gly Ile Ala Ile Ala Met Ala Cys Ile Val Gly Leu Met Trp Leu
 1 5 10 15
 Ser Tyr Phe Val Ala Ser Phe
 20

-continued

<210> SEQ ID NO 83
 <211> LENGTH: 20
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 83

His Leu Val Asp Phe Gln Val Thr Ile Ala Glu Ile Leu Ile Ile Ile
 1 5 10 15

Met Arg Thr Phe
 20

<210> SEQ ID NO 84
 <211> LENGTH: 15
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 84

Met Lys Ile Ile Leu Phe Leu Thr Leu Ile Val Phe Thr Ser Cys
 1 5 10 15

<210> SEQ ID NO 85
 <211> LENGTH: 19
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 85

Ser Pro Leu Phe Leu Ile Val Ala Ala Leu Val Phe Leu Ile Leu Cys
 1 5 10 15

Phe Thr Ile

<210> SEQ ID NO 86
 <211> LENGTH: 83
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 86

Glu Leu Tyr His Tyr Gln Glu Cys Val Arg Gly Thr Thr Val Leu Leu
 1 5 10 15

Lys Glu Pro Cys Pro Ser Gly Thr Tyr Glu Gly Asn Ser Pro Phe His
 20 25 30

Pro Leu Ala Asp Asn Lys Phe Ala Leu Thr Cys Thr Ser Thr His Phe
 35 40 45

Ala Phe Ala Cys Ala Asp Gly Thr Arg His Thr Tyr Gln Leu Arg Ala
 50 55 60

Arg Ser Val Ser Pro Lys Leu Phe Ile Arg Gln Glu Glu Val Gln Gln
 65 70 75 80

Glu Leu Tyr

<210> SEQ ID NO 87
 <211> LENGTH: 37
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

<400> SEQUENCE: 87

caggaaacag ctatgacacc aagaacaagg ctctcca

37

<210> SEQ ID NO 88
 <211> LENGTH: 37
 <212> TYPE: DNA

-continued

<213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

<400> SEQUENCE: 88

caggaaacag ctatgacgat agggcctctt ccacaga 37

<210> SEQ ID NO 89
 <211> LENGTH: 496
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus
 <220> FEATURE:
 <221> NAME/KEY: misc_feature
 <222> LOCATION: (11)..(11)
 <223> OTHER INFORMATION: n is a, c, g, or t

<400> SEQUENCE: 89

acctacccag ngaaaagcca accaacctcg atctcttgta gatctgttct ctaaacgaac 60
 tttaaaatct gtgtagctgt cgctcggctg catgcctagt gcacctacgc agtataaaca 120
 ataataaatt ttactgtcgt tgacaagaaa cgagtaactc gtcctcttcc tgcagactgc 180
 ttacggtttc gtccgtgttg cagtcgatca tcagcatacc taggtttcgt cggggtgga 240
 ccgaaaggta agatggagag ccttgttctt ggtgtcaacg agaaaacaca cgtccaactc 300
 agtttgctg tccttcaggt tagagacgtg ctagtgcgtg gcttcgggga ctctgtggaa 360
 gaggcctat cggaggcacg tgaacacctc aaaaatggca cttgtggtct agtagagctg 420
 gaaaaaggcg tactgcccc gcttgaacag ccctatgtgt tcattaacg ttctgatgcc 480
 ttaagcacca atcacg 496

<210> SEQ ID NO 90
 <211> LENGTH: 523
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 90

gtcgacaaca atttctgtgg ccagatggg tacctcttg attgcatcaa agattttctc 60
 gcacgcgagg gcaagtcaat gtgactctt tccgaacaac ttgattacat cgagtcgaag 120
 agaggtgtct actgctgccg tgacatgag catgaaattg cctgggtcac tgagcgtct 180
 gataagagct acgagcacca gacaccttc gaaattaaga gtgccaagaa atttgacct 240
 ttcaaagggg aatgcccmaa gtttgtgtt cctcttaact caaaagtcaa agtcattcaa 300
 ccacgtgttg aaaagaaaa gactgagggt ttcattgggc gtatacgtc tgtgtaccct 360
 gttgcatctc cacaggagtg taacaatatg cacttgctc ccttgatgaa atgtaatcat 420
 tgcgatgaag tttcatggca gacgtgcgac tttctgaaag ccacttgta acattgtggc 480
 actgaaaatt tagttattga aggacctact acatgtgggt acc 523

<210> SEQ ID NO 91
 <211> LENGTH: 324
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 91

cttaggtgac gagcttgga ctgatccat tgaagattat gaacaaaact ggaacactaa 60
 gcatggcagt ggtgactcc gtgaactcac tcgtgagctc aatggagggt cagtcactcg 120
 ctatgtcgac aacaatttct gtggcccaga tgggtacct cttgattgca tcaaagattt 180
 tctgcacgc ggggcaagt caatgtgcac ttttccgaa caacttgatt acatcgagtc 240

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gaagagaggt gtctactgct gccgtgacca tgagcatgaa attgcctggt tcaactgagcg 300
ctcctgataa gagctacgag cacc 324

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<210> SEQ ID NO 92
<211> LENGTH: 495
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 92

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tgctataata agcgtgccta ctgggttccct cgtgctagtg ctgatattgg gctcaggcca 60
tactggcatt actggtgaca atgtggagac cttgaatgag gatctccttg agatactgag 120
tcgtgaacgt gttaacatta acattgttgg cgattttcat ttgaatgaag aggttgccat 180
cattttggca tctttctctg cttctacaag tgcctttatt gacactataa agagtcttga 240
ttacaagtct ttcaaaacca ttgttgagtc ctgcggtaac tataaagtta ccaagggaaa 300
gcccgtaaaa ggtgcttggg acattggaca acagagatca gttttaacac cactgtgtgg 360
ttttccctca caggctgctg gtgttatcag atcaattttt gcgcgcacac ttgatgcagc 420
aaaccactca attcctgatt tgcaaagagc agctgtcacc atacttgatg gtatttctga 480
acagtcatta cgtct 495

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<210> SEQ ID NO 93
<211> LENGTH: 486
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 93

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gccactcaaa cattgaaact cgactccgca agggaggtag gactagatgt tttggaggct 60
gtgtgtttgc ctatgttggc tgctataata agcgtgccta ctgggttccct cgtgctagtg 120
ctgatattgg ctccagccat actggcatta ctggtgacaa tgtggagacc ttgaatgagg 180
atctccttga gatactgagt cgtgaacgtg ttaacattaa cattgttggc gattttcatt 240
tgaatgaaga ggttgccatc attttggcat ctttctctgc ttctacaagt gcctttattg 300
acactataaa gagtcttgat tacaagtctt tcaaaacat tgttgagtcc tgcggtaact 360
ataaagttac caagggaaag cccgtaaaag gtgcttgaa cattggacaa cagagatcag 420
ttttaacacc actgtgtggt tttccctcac aggctgctgg tgttatcaga tcaatttttg 480
cgcgca 486

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<210> SEQ ID NO 94
<211> LENGTH: 567
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 94

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cactactgtg gaaaaactca ggcctatctt tgaatggatt gaggcgaaac ttagtgcagg 60
agttgaattt ctcaaggatg cttgggagat totcaaattt ctcattacag gtgtttttga 120
catcgtcaag ggtcaaaatac aggttgcttc agataacatc aaggattgtg taaaatgctt 180
cattgatggt gttacaagg cactcgaaat gtgcattgat caagtcacta tcgctggcgc 240
aaagttgcca tcaactcaact taggtgaagt cttcatcgct caaagcaagg gactttaccg 300
tcagtgata cgtggcaagg agcagctgca actactcatg cctcttaagg caccaaaga 360
agtaaccttt cttgaagggt attcacatga cacagtactt acctctgagg aggttgttct 420

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caagaacggt gaactcgaag cactcgagac gcccgttgat agcttcacaa atggagctat 480
cgttggcaca ccagctctgtg taaatggcct catgctctta gagattaagg acaaagaaca 540
atactgcgca ttgtctcctg gtttact 567

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<210> SEQ ID NO 95
<211> LENGTH: 516
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 95

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gggagattct caaatttctc attacaggtg tttttgacat cgtcaagggt caaatacagg 60
ttgcttcaga taacatcaag gattgtgtaa aatgcttcat tgatgttggt aacaaggcac 120
tcgaaatgtg cattgatcaa gtcactatcg ctggcgcaaa gttcggatca ctcaacttag 180
gtgaagtctt catcgctcaa agcaagggac tttaccgtca gtgtatacgt ggcaaggagc 240
agctgcaact actcatgcct cttaggcac caaaagaagt aacctttctt gaaggtgatt 300
cacatgacac agtacttacc tctgaggagg ttgttctcaa gaacggtgaa ctggaagcac 360
tcgagacgcc cgttgatagc ttcacaaatg gagctatcgt tggcacacca gtctgtgtaa 420
atggcctcat gctcttagag attaggaca aagaacaata ctgcgcattg tctcctgggt 480
tactggctac aaacaatgtc tttcgcttaa aagggg 516

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<210> SEQ ID NO 96
<211> LENGTH: 448
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 96

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agttcgagtt gaggaagaag aagaggaaga ctggctggat gatactactg agcaatcaga 60
gattgagcca gaaccagaac ctacacctga agaaccagtt aatcagttta ctggttattt 120
aaaacttact gacaatggtg ccattaaatg tgttgacatc gttaaggagg cacaaagtgc 180
taatcctatg gtgattgtaa atgctgctaa catacacctg aaacatggtg gtggtgtagc 240
aggtgcactc aacaaggcaa ccaatggtgc catgcaaaag gagagtgatg attacattaa 300
gctaaatggc cctcttacag taggagggtc ttgtttgctt tctggacata atcttgctaa 360
gaagtgtctg catgttggtg gacctaacct aaatgcaggt gaggacatcc agcttcttaa 420
ggcagcatat gaaaatttca attcacag 448

```

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<210> SEQ ID NO 97
<211> LENGTH: 333
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 97

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agaggatgat tatcaaggtc tccctctgga atttgggtgcc tcagctgaaa cagttcgagt 60
tgaggaagaa gaagaggaag actggctgga tgatactact gagcaatcag agattgagcc 120
agaaccagaa cctacacctg aagaaccagt taatcagttt actggttatt taaaacttac 180
tgacaatggt gccattaaat gtggtgacat cgtaaggag gcacaaagtg ctaatcctat 240
ggtgattgta aatgctgcta acatacacct gaaacatggt ggtggtgtag caggtgcact 300
caacaaggca accaatggtg ccatgcaaaa gga 333

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<210> SEQ ID NO 98
<211> LENGTH: 399

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<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 98

gagatgctct caagagcttt gaagaaagtg ccagttgatg agtatataac cacgtaccct    60
ggacaaggat gtgctgggta tacacttgag gaagctaaga ctgctcttaa gaaatgcaaa    120
tctgcatttt atgtactacc ttcagaagca cctaattgcta aggaagagat tctaggaact    180
gtatcctgga atttgagaga aatgcttgct catgctgaag agacaagaaa attaatgcct    240
atatgcatgg atggttagagc cataatggca accatccaac gtaagtataa aggaattaaa    300
attcaagagg gcatcgttga ctatgggtgc cgattcttct tttatactag taaagagcct    360
gtagcttcta ttattacgaa gctgaactct ctaaagtag                                399

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<210> SEQ ID NO 99
<211> LENGTH: 437
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 99

agaaatctgt cgtacagaag cctgtcgatg tgaagccaaa aattaaggcc tgcattgatg    60
aggttaccac aacctggaa gaaactaagt ttcttaccac taagttaact ttgtttgctg    120
atatcaatgg taagctttac catgattctc agaacatgct tagagggtgaa gatatgtctt    180
tccttgagaa ggatgcacct tacatggtag gtgatggtat cactagtggg gatatacctt    240
gtgttgtaat accctccaaa aaggctgggt gcactactga gatgctctca agagctttga    300
agaaagtgcc agttgatgag tatataacca cgtaccctgg acaaggatgt gctggttata    360
cacttgagga agctaagact gctcttaaga aatgcaaact tgcattttat gtactacctt    420
cagaagcacc taatgct                                437

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<210> SEQ ID NO 100
<211> LENGTH: 569
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 100

cctctatcgt attgacggag ctcaccttac aaagatgtca gagtacaaag gaccagtgac    60
tgatgttttc tacaaggaaa catcttacac tacaaccatc aagcctgtgt cgtataaaact    120
cgatggagtt acttacacag agattgaacc aaaattggat gggtattata aaaaggataa    180
tgcttactat acagagcagc ctatagacct tgtaccaact caaccattac caaatgagag    240
ttttgataat ttcaaaactca catgttctaa cacaaaattt gctgatgatt taaatcaaat    300
gacaggcttc acaaagccag cttaacgaga gctatctgtc acattcttcc cagacttgaa    360
tggogatgta gtggctattg actatagaca ctattcagcg agtttcaaga aagggtgctaa    420
attactgcat aagccaattg tttggccatc taaccaggct acaaccaaga caacgttcaa    480
accaaact tgggtgttac gttgtctttg gagtacaaag ccagtagata cttcaaatc     540
attgaagtt ctggcagtag aagacacat                                569

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<210> SEQ ID NO 101
<211> LENGTH: 187
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 101

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tcagcagata cttcaaatc atttgaagt ctggcagtag aagacacaca aggaatggac 60
aatcttgctt gtgaaagtca acaaccacc tctgaagaag tagtggaaaa tcctaccata 120
cagaaggaag tcatagagcg tgacgtgaaa actaccgaag ttgtaggcaa tgtcactt 180
aaacat 187

```

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<210> SEQ ID NO 102
<211> LENGTH: 271
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 102

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aaatgcgacg agtctgcttc taagtctgct tctgtgtact acagtcagct gatgtgccaa 60
cctattctgt tgcttgacca agctcttgta tcagacgttg gagatagtac tgaagtttcc 120
gttaagatgt ttgatgctta tgcgacacc ttttcagcaa cttttagtgt tcctatggaa 180
aaacttaagg cacttggttc tacagctcac agcgagttag caaagggtgt agcttttagat 240
ggtgtccttt ctacattcgt gtcagctgcc c 271

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<210> SEQ ID NO 103
<211> LENGTH: 363
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 103

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catttcatca gcaattcttg gctcatgtgg tttatcatta gtattgtaca aatggcacc 60
gtttctgcaa tggtaggat gtacatcttc tttgcttctt tctactacat atggaagagc 120
tatgttcata tcatggatgg ttgcacctct tcgacttgca tgatgtgcta taagcgcaat 180
cgtgccacac gcgttgatgg tacaactatt gttaatggca tgaagagatc tttctatgtc 240
tatgcaaatg gaggccgtgg cttctgcaag actcacaatt ggaattgtct caattgtgac 300
acattttgca ctggtagtag attcattagt gatgaagtgg ctcgagatth gtcactccag 360
ttt 363

```

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<210> SEQ ID NO 104
<211> LENGTH: 500
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 104

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agagatcttg gcgcatgtat tgactgtaat gcaaggcata tcaatgcca aggtagcaaa 60
aagtcaaat gtttcaactca tctggaatgt aaaagactac atgtctttat ctgaacagct 120
gcgtaaaaca attcgtatgt ctgccaagaa gaacaacata ctttttagac taactgtgtc 180
tacaactaga caggttgctca atgtcataac tactaaaatc tcaactcaagg gtgtaagat 240
tgtagtact tgttttaaac ttatgcttaa ggccacatta ttgtgcgttc ttgctgcatt 300
ggtttgttat atcgttatgc cagtacatc attgtcaatc catgatggtt acacaaatga 360
aatcattggt tacaagcca ttcaggatgg tgcactcgt gacatcattt ctactgatga 420
ttgttttgca aataaacatg ctggttttga cgcattggtt agccagcgtg gtggttcata 480
caaaaatgac aaaagctgcc 500

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<210> SEQ ID NO 105
<211> LENGTH: 537
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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-continued

<400> SEQUENCE: 105

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cattgtcaat ccatgatggt tacacaaatg aaatcattgg ttacaaagcc attcaggatg    60
gtgtcactcg tgacatcatt tctactgatg attgttttgc aaataaacat gctggttttg   120
acgcatgggt tagccagcgt ggtggttcat acaaaaatga caaaagctgc cctgtagtag   180
ctgctatcat tacaagagag attggtttca tagtgcctgg cttaccgggt actgtgctga   240
gagcaatcaa tggtgacttc ttgcatttcc tacctcgtgt ttttagtgct gttggcaaca   300
tttgctacac accttccaaa ctcatgagt atagtgattt tgctacctct gcttgcggtc   360
ttgtgctga gtgtacaatt ttaaggatg ctatgggcaa acctgtgcca tattgttatg   420
acactaattt gctagagggt tctatttctt atagtgagct tcgtccagac actcgttatg   480
tgcttatgga tggttccatc atacagtttc ctaacactta cctggagggg tctgtta    537

```

<210> SEQ ID NO 106

<211> LENGTH: 427

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 106

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cacttttggg tttgatgtct ttcactatac tctgtctggg accagcttac agctttctgc    60
cgggagtcta ctcaagtctt tacttgtact tgacattcta tttaccaat gatgtttcat   120
tcttggtcga ccttcaatgg tttgccatgt tttctcctat tgtgcctttt tggataacag   180
caatctatgt attctgtatt tctctgaagc actgccattg gttctttaac aactatctta   240
ggaaaagagt catgtttaat ggagttacat ttagtacctt cgaggaggct gctttgtgta   300
cctttttgct caacaaggaa atgtacctaa aattgcgtag cgagacactg ttgccactta   360
cacagataaa caggatctct gctctatata acaagtacaa gtatttcagt ggagccttag   420
atactac                                           427

```

<210> SEQ ID NO 107

<211> LENGTH: 537

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 107

```

agtaacaact tttgatgctg agtactgtag acatggtaca tgcgaaaggc cagaagtagg    60
tatttgecta tctaccagtg gtagatgggt tottaataat gagcattaca gagctctatc   120
aggagttttc tgtggtgttg atgcatgaa tctcatagct aacatcttta ctctctttgt   180
gcaacctgtg ggtgctttag atgtgtctgc ttcagtagtg gctggtggtta ttattgccat   240
attggtgact tgtgtgcctt actactttat gaaattcaga cgtgtttttg gtgagtacaa   300
ccatgtttgt gctgctaatt cacttttggg tttgatgtct ttcactatac tctgtctggg   360
accagcttac agctttctgc cgggagtcta ctcaagtctt tacttgtact tgacattcta   420
tttcaccaat gatgtttcat tcttggtcga ccttcaatgg tttgccatgt tttctcctat   480
tgtgcctttt tggataacag caatctatgt attctgtatt tctctgaagc actgcca    537

```

<210> SEQ ID NO 108

<211> LENGTH: 551

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 108

-continued

```

agtatactgt ccaagacatg tcatttgac agcagaagac atgcttaatc ctaactatga 60
agatctgctc attcgc aaat ccaaccatag ctttcttggt caggctggca atgttcaact 120
tcgtgttatt ggccattcta tgcaaaattg tctgcttagg cttaaagttg atacttctaa 180
ccctaagaca cccaagtata aatttgtccg tatccaacct ggtcaaacat tttcagttct 240
agcatgctac aatggttcac catctgggtt ttatcagttg gccatgagac ctaatcatac 300
cattaaaggt tctttcctta atggatcatg tggtagtggt ggttttaaca ttgattatga 360
ttgctgtctc ttctgctata tgcacatata ggagcttcca acaggagtac acgctggtag 420
tgacttagaa ggtaaattct atggctcatt tgttgacaga caaactgcac aggctgcagg 480
tacagacaca accataacat taaatgtttt ggcatggctg tatgctgctg ttatcaatgg 540
tgataggtgg t 551

```

```

<210> SEQ ID NO 109
<211> LENGTH: 593
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 109

```

```

acttagcaaa ggctcctaaat gacttttagca actcaggtgc tgatgttctc taccaaccac 60
cacagacatc aatcacttct gctgttctgc agagtggttt taggaaaatg gcattcccgt 120
caggcaaaat tgaaggggtc atggtacaag taacctgtgg aactacaact cttaatggat 180
tgtggttga tgacacagta tactgtccaa gacatgtcat ttgcacagca gaagacatgc 240
ttaatcctaa ctatgaagat ctgctcattc gcaaatccaa ccatagcttt cttgttcagg 300
ctggcaatgt tcaacttctg gttattggcc attctatgca aaattgtctg cttaggetta 360
aagttgatac ttctaaccct aagacaccca agtataaatt tgcctgtatc caacctggtc 420
aaacattttc agttctagca tgctacaatg gttcaccatc tgggttttat cagtgtgcca 480
tgagacctaa tcataccatt aaaggttctt tccttaatgg atcatgtggt agtgttggtt 540
ttaaacttga ttatgattgc gtgtctttct gctatatgca tcatatggag ctt 593

```

```

<210> SEQ ID NO 110
<211> LENGTH: 504
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 110

```

```

tgtgctgctt tgaagagact gctgcagaat gggatgaat ggtcgtacta tccttggtag 60
cactatttta gaagatgagt ttacaccatt tgatgttgtt agacaatgct ctgggtgttac 120
cttcaagggt taagtccaag aaaattgtta agggcactca tcattggatg cttttaactt 180
tcttgacatc actattgatt cttgttcaaa gtacacagtg gtcactgttt ttctttgttt 240
acgagaatgc tttcttgcca tttactcttg gtattatggc aattgctgca tgtgctatgc 300
tgcttgtaa gcataagcac gcattcttct gcttgtttct gttacctctc cttgcaacag 360
ttgcttactt taatatggtc tacatgcctg ctatgctggg gatgcgtatc atgacatggc 420
ttgaattggc tgacactagc ttgtctggtt ataggcttaa ggattgtggt atgtatgctt 480
cagctttagt tttgcttatt ctca 504

```

```

<210> SEQ ID NO 111
<211> LENGTH: 298
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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-continued

<400> SEQUENCE: 111

```

taggcttaag gattgtggtta tgtatgcttc agcttttagtt ttgettattc tcatgacagc   60
tcgcactggt tatgatgatg ctgctagacg tgtttggaca ctgatgaatg tcattacact   120
tgtttacaaa gtctactatg gtaatgcttt agatcaagct atttccatgt gggccttagt   180
tatttctgta acctctaact attctggtgt cgttacgact atcatgtttt tagctagagc   240
tatagtgttt gtgtgtgttg agtattaccc attgttattt attacctggc aacacctt    298

```

<210> SEQ ID NO 112

<211> LENGTH: 530

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 112

```

aaacaggcaa gatctgagga caagagggca aaagtaacta gtgctatgca aacaatgctc   60
ttcactatgc ttaggaagct tgataatgat gcacttaaca acattatcaa caatgcgcggt  120
gatggttggt ttccactcaa catcatacca ttgactacag cagccaaact catggttggt  180
gtccctgatt atggtaccta caagaacact tgtgatggta acacctttac atatgcatct  240
gcactctggg aatccagca agttgttgat gcggatagca agattgttca acttagtgaa  300
attaacatgg acaattcacc aaatttggtc tggcctctta ttgttacagc tctaagagcc  360
aactcagctg ttaaactaca gaataatgaa ctgagtccag tagcactacg acagatgtcc  420
tgtgcggtct gtaccacaca aacagcttgt actgatgaca atgcacttgc ctactataac  480
aattcgaagg gaggtagggt tgtgtcggca ttactatcag accaccaagc    530

```

<210> SEQ ID NO 113

<211> LENGTH: 605

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 113

```

gaagtcgttc tcaaaaagt aaagaaatct ttgaatgtgg ctaaatctga gtttgaccgt   60
gatgtgcca tgcaacgcaa gttggaaaag atggcagatc aggctatgac ccaaatgtac   120
aaacaggcaa gatctgagga caagagggca aaagtaacta gtgctatgca aacaatgctc   180
ttcactatgc ttaggaagct tgataatgat gcacttaaca acattatcaa caatgcgcggt  240
gatggttggt ttccactcaa catcatacca ttgactacag cagccaaact catggttggt  300
gtccctgatt atggtaccta caagaacact tgtgatggta acacctttac atatgcatct  360
gcactctggg aatccagca agttgttgat gcggatagca agattgttca acttagtgaa  420
attaacatgg acaattcacc aaatttggtc tggcctctta ttgttacagc tctaagagcc  480
aactcagctg ttaaactaca gaataatgaa ctgagtccag tagcactacg acagatgtcc  540
tgtgcggtct gtaccacaca aacagcttgt actgatgaca atgcacttgc ctactataac  600
aatcc                                           605

```

<210> SEQ ID NO 114

<211> LENGTH: 176

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 114

```

acactggtag aggacaggca attactgtaa caccagaagc taacatggac caagagtcct   60

```

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 ttggtgggtgc ttcattgtgt ctgtattgta gatgccacat tgaccatcca aatcctaaag 120

gattctgtga cttgaaaggt aagtagctcc aaatacctac cacttgtgct aatgat 176

<210> SEQ ID NO 115

<211> LENGTH: 516

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 115

actgtaaac cagaagctaa catggaccaa gagtcctttg gtggtgcttc atgttgtctg 60

tattgtagat gccacattga ccatccaaat cctaaaggat tctgtgactt gaaaggtgag 120

tacgtccaaa tacctaccac ttgtgctaata gaccagtggt gttttacact tagaaacaca 180

gtctgtaccg tctgcggaat gtggaaggt tatggctgta gttgtgacca actccgcgaa 240

cccttgatgc agtctgcgga tgcacaaacg tttttaaagc ggtttgcggt gtaagtgcag 300

cccgcttac accgtgcggc acaggcacta gtactgatgt cgtctacagg gcttttgata 360

tttacaacga aaaagttgct ggttttgcaa agttcctaaa aactaattgc tgcgcttcc 420

aggagaagga tgaggaagc aatttattag actcttactt ttagttagg aggcatacta 480

tgtctaccta ccaacatgaa gagactattt ataact 516

<210> SEQ ID NO 116

<211> LENGTH: 366

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 116

accacttatt aagtgggatt tgctgaaata tgattttacg gaagagagac tttgtctctt 60

cgaccgttat tttaaatatt gggaccagac ataccatccc aattgtatta actgtttgga 120

tgataggtgt atccttcatt gtgcaaacgt taatgtgta tttctgctg tgtttccacg 180

tacaagtttt ggaccactag taagaaaaat attttagat ggtgttcctt ttgttgtttc 240

aactggatac cattttcgtg agttaggagt cgtacataat caggatgtaa acttacatag 300

ctcgcgtctc agtttcaagg aacttttagt gtatgctgct gatccagcta tgcacgcagc 360

ttctgg 366

<210> SEQ ID NO 117

<211> LENGTH: 291

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 117

tgaaaaagt gctggttttg caaagttcct aaaaactaat tgctgtcgt tccaggagaa 60

ggatgaggaa ggcaatttat tagactctta cttttagatt aagaggcata ctatgtctaa 120

ctaccaacat gaagagacta tttataaact ggttaaagat tgtccagcgg ttgctgtcca 180

tgactttttc aagtttagag tagatggtga catggtacca catatatcac gtcagcgtct 240

aactaaatac acaatggctg atttagtcta tgctctacgt cattttgatg a 291

<210> SEQ ID NO 118

<211> LENGTH: 480

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 118

gagtcacata tggatgctga tctcgaaaa ccaacttatta agtgggattt gctgaaatat 60

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gattttacgg aagagagact ttgtctcttc gaccgttatt ttaaattattg ggaccagaca 120
taccatccca attgtattaa ctgtttgat gataggtgta tccttcattg tgcaaaacttt 180
aatgtgttat tttctactgt gtttccacct acaagttttg gaccactagt aagaaaaata 240
ttttagatg gtgttccctt tgttgtttca actggatacc attttcgtga gttaggagtc 300
gtacataate aggatgtaaa cttacatagc tcgctgtcca gtttcaagga acttttagtg 360
tatgctgctg atccagctat gcatgcagct tctggcaatt tattgctaga taaacgcact 420
acatgctttt cagtagctgc actaacaac aatgttgctt tcaaaactgt caaacccggt 480

```

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<210> SEQ ID NO 119
<211> LENGTH: 405
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 119

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aatgggaact ggtacgattt cggtgatttc gtacaagtag caccaggctg cggagttcct 60
attgtggatt catattactc attgtgatg cccatcctca ctttgactag ggcattggct 120
gctgagtccc atatggatgc tgatctcgca aaaccactta ttaagtgaga tttgctgaaa 180
tatgatttta cggaagagag actttgtctc ttcgaccggt attttaata ttgggaccag 240
acataccatc ccaattgat taactgtttg gatgataggt gtatccttca ttgtgcaaac 300
tttaatgtgt tattttctac tgtgtttcca cctacaagct ttggaccact agtaagaaaa 360
atattgtag atgggtgttc tttgttgtt tcaactggat accat 405

```

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<210> SEQ ID NO 120
<211> LENGTH: 562
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (67)..(67)
<223> OTHER INFORMATION: n is a, c, g, or t

```

```

<400> SEQUENCE: 120

```

```

ctattgatgc ttaccactt acaaaacatc ctaatcagga gtatgctgat gtctttcact 60
tgtattnaca atacattaga aagttacatg atgagcttac tggccacatg ttggacatgt 120
attccgtaat gctaactaat gataacacct cacggtactg ggaacctgag ttttatgagg 180
ctatgtacac accacataca gtcttgacag ctgtaggtgc ttgtgtattg tgcaattcac 240
agacttcact tcggtgcggt gcctgtatta ggagaccatt cctatgttgc aagtgtctgt 300
atgacctgt catttcaaca tcacacaaat tagtgttgtc tgtaatccc tatgtttgca 360
atgcccagg ttgtgatgtc actgatgtga cacaactgta tctaggaggt atgagctatt 420
attgcaagtc acataagcct cccattagtt ttccattatg tgctaaggt caggtttttg 480
gtttatacaa aaacacatgt gtaggcagtg acaatgtcac tgacttcaat gcgatagcaa 540
catgtgattg gactaatgct gg 562

```

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<210> SEQ ID NO 121
<211> LENGTH: 580
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 121

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```

gctatgtaca caccacatac agtcttcgag gctgtaggtg cttgtgtatt gtgcaattca 60

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-continued

cagacttcac ttcggtgcgg tgcctgtatt aggagacat tcctatgttg caagtgcctgc	120
tatgaccatg tcatttcaac atcacacaaa ttagtggttg ctgttaatcc ctatgtttgc	180
aatgccccag gttgtgatgt cactgatgtg acacaactgt atctaggagg tatgagctat	240
tattgcaagt cacataagcc tcccattagt tttccattat gtgctaattg tcaggttttt	300
ggtttataca aaaacacatg tgtaggcagt gacaatgtca ctgacttcaa tgcgatagca	360
acatgtgatt ggactaatgc tggcgattac atacttgcca acacttgtae tgagagactc	420
aaagcttttcg cagcagaaaac gctcaaagcc actgaggaaa catttaagct gtcatatggt	480
attgccactg tacgcgaagt actctctgac agagaattgc atctttcatg ggaggttgga	540
aaacctagac caccattgaa cagaaactat gtctttactg	580

<210> SEQ ID NO 122
 <211> LENGTH: 610
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 122

tggtgatgct gttgtgtaca gaggtactac gacatacaag tgaatgttg gtgattactt	60
tgtgttgaca tctcacactg taatgccact tagtgcaect actctagtgc cacaagagca	120
ctatgtgaga attactggct tgtacccaac actcaacatc tcagatgagt tttctagcaa	180
tgttgcaaat tatcaaaagg tcggcatgca aaagtactct aactccaag gaccacctgg	240
tactgtaag agtcattttg ccatcggact tgctctctat taccatctg ctgcgatagt	300
gtatacggca tgctctcatg cagctgttga tgcctatgt gaaaaggcat taaaatattt	360
gcccatagat aatgtagta gaatcatacc tgcgcgtgcg cgcgtagagt gttttgataa	420
attcaaaagt aattcaacac tagaacagta tgttttctgc actgtaaatg cattgccaga	480
aaacaactgct gacattgtag tctttgatga aatctctatg gctactaatt atgacttgag	540
tgttgtcaat gctagacttc gtgcaaaaaca ctacgtctat attggcgatc ctgctcaatt	600
accagccct	610

<210> SEQ ID NO 123
 <211> LENGTH: 429
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 123

ccaacactca acatctcaga tgagttttct agcaatgttg caaattatca aaaggtcggc	60
atgcaaaagt actctacact ccaaggacca cctggtagtg gtaagagtca ttttgccatc	120
ggacttgctc tctattacc atctgctcgc atagtgtata cggcatgctc tcatgcagct	180
gttgatgccc tatgtgaaaa ggcattaaaa tatttgccca tagataaatg tagtagaatc	240
atacctgcgc gtgcgcgcgt agagtgtttt gataaattca aagtgaattc aacactagaa	300
cagtatgttt tctgcactgt aatgcattg ccagaaacaa ctgctgacat tgtagtcttt	360
gatgaaatct ctatggctac taattatgac ttgagtgttg tcaatgctag acttcgtgca	420
aaacactac	429

<210> SEQ ID NO 124
 <211> LENGTH: 486
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 124

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caatgtggct atcacaaggg caaaaattgg cttttgtgc ataagtctg atagagatct 60
ttatgacaaa ctgcaattta caagtctaga aataccacgt cgcaatgtgg ctacattaca 120
agcagaaaaat gtaactggac tttttaagga ctgtagtaag atcattactg gtcttcatcc 180
tacacaggca cctacacacc tcagcgttga tataaagttc aagactgaag gattatgtgt 240
tgacatacca ggcataccaa aggacatgac ctaccgtaga ctcatctcta tgatggggtt 300
caaatgaat taccaagtca atggttacc taatatgttt atcaccgcg aagaagctat 360
tcgtcacgtt cgtgcgtgga ttggcttga tgtagagggc tgcatgcaa ctagagatgc 420
tgtgggtact aacctacctc tccagctagg attttctaca ggtgttaact tagtagctgt 480
accgac 486

```

```

<210> SEQ ID NO 125
<211> LENGTH: 427
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 125

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```

aaaggacatg acctaccgta gactcatctc tatgatgggt tcaaaatga attaccaagt 60
caatggttac cctaatatgt ttatcaccgc cgaagaagct atctgacacg ttcgtgcgtg 120
gattggcttt gatgtagagg gctgtcatgc aactagagat gctgtgggta ctaacctacc 180
tctccagcta ggattttcta caggtgttaa cttagtagct gtaccgactg gttatgttga 240
cactgaaaaa aacacagaat tcaccagagt taatgcaaaa cctccaccag gtgaccagtt 300
taaacatctt ataccactca tgtataaagg cttgccctgg aatgtagtgc gtattaagat 360
agtacaaatg ctcagtata cactgaaagg attgtcagac agagtctgtg tcgtcctttg 420
ggcgcat 427

```

```

<210> SEQ ID NO 126
<211> LENGTH: 392
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 126

```

```

atggaaatgc acatgtggct agttgtgatg ctatcatgac tagatgttta gcagtccatg 60
agtgctttgt taagcgcgtt gattggctctg ttgaataccc tattatagga gatgaactga 120
gggttaatte tgcttcgaga aaagtacaac acatggttgt gaagtctgca ttgcttgctg 180
ataagtttcc agttcttcat gacattggaa atccaaaggc tatcaagtgt gtgctcagg 240
ctgaagtaga atggaagtcc tacgatgctc agccatgtag tgacaaagct tacaaaatag 300
aggaactctt ctattcttat gctacacatc acgataaatt cactgatggg gtttgtttgt 360
tttgaattg taacgttgat cgtaaccag cc 392

```

```

<210> SEQ ID NO 127
<211> LENGTH: 483
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 127

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```

gcttcatcag atacttatgc ctgctggaat cattctgtgg gttttgacta tgtctataac 60
ccatztatga ttgatgttca gcagtggggc tttacgggta accttcagag taacctgac 120
caacattgcc aggtacatgg aatgcacat gtggctagtt gtgatgctat catgactaga 180

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tgtttagcag tccatgagtg ctttgtaag cgcgttgatt ggtctgttga ataccctatt 240
ataggagatg aactgagggg taattctgct tgcagaaaag tacaacacat ggttgtgaag 300
tctgcattgc ttgctgataa gtttccagtt cttcatgaca ttggaaatcc aaaggctatc 360
aagtgtgtgc ctcagcgtga agtagaatgg aagttctacg atgctcagcc atgtagtgc 420
aaagcttaca aaatagagga actcttctat tcttatgcta cacatcacga taaattcact 480
gat 483

```

```

<210> SEQ ID NO 128
<211> LENGTH: 326
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 128

```

```

tcaaagggac cagcacaagc tagcgtcaat ggagtcacat taattggaga atcagtaaaa 60
acacagttta actactttaa gaaagtagac ggcattatc aacagttgcc tgaacctac 120
tttactcaga gcagagactt agaggathtt aagcccagat cacaatgga aactgacttt 180
ctcgagctcg ctatggatga attcatacag cgatataagc tcgagggcta tgccttcgaa 240
cacatcgttt atggagatgt cagtcatgga caacttggcg gtcttcattt aatgataggg 300
ttagccaagc gctcacaaga ttcact 326

```

```

<210> SEQ ID NO 129
<211> LENGTH: 457
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 129

```

```

acaccttcaa agggaccagc acaagctagc gtcaatggag tcacattaat tggagaatca 60
gtaaaaaaac agtttaacta ctttaagaaa gtagacggca ttattcaaca gttgcctgaa 120
acctacttta ctcagagcag agacttagag gattttaagc ccagatcaca aatggaaact 180
gactttctcg agctcgtat ggatgaattc atacagcagc ataagctcga gggctatgcc 240
ttcgaacaca tcgtttatgg agatttcagt catggacaac ttggcgggtct tcatttaatg 300
ataggcttag ccaagcgcct acaagattca ccaacttaat tagaggatgt tatccctatg 360
gacagcacag tgaaaaaatta cttcataaca gatgcgcaaa caggttcacg aaaatgtgtg 420
tgttctgtga ttgatctttt acttgatgac tttgtcg 457

```

```

<210> SEQ ID NO 130
<211> LENGTH: 493
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 130

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```

cgcaaagtat actcaactgt gtcaataact aaatacactt actttagctg taccctaaa 60
catgagagtt attcactttg gtgctggctc tgataaagga gttgcaccag gtacagctgt 120
gctcagacaa tggttgcca ctggcacact acttgcgat tcagatctta atgacttcgt 180
ctccgacgca gattctactt taattggaga ctgtgcaaca gtacatacgg ctaataaatg 240
ggactttatt attagcgata tgtatgaccc taggacccaa catgtgacaa aagagaatga 300
ctctaaagaa gggtttttca cttatctgtg tggatttata aagcaaaaac tagccctggg 360
tggttctata gctgtaaaaga taacagagca ttcttggaat gctgaccttt acaagcttat 420
gggcatttc tcatgggtga cagcttttgt tacaatgta aatgcatcat catcggaagc 480

```

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atttttaatt ggg 493

<210> SEQ ID NO 131
 <211> LENGTH: 490
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 131

acttaaatac acttacttta gctgtaccct acaacatgag agttattcac tttggtgctg 60
 gctctgataa aggagttgca ccaggtagag ctgtgctcag acaatggttg ccaactggca 120
 cactacttgt cgattcagat cttaatgact tegtctccga cgcagattct actttaattg 180
 gagactgtgc aacagtagat acggctaata aatgggacct tattattagc gatatgtatg 240
 accctaggac caaacatgtg acaaaagaga atgactctaa agaagggttt ttcacttate 300
 tgtgtggatt tataaagcaa aaactagccc tgggtgggtc tatagctgta aagataacag 360
 agcattcttg gaatgctgac ctttacaagc ttatgggcca tttctcatgg tggacagcct 420
 ttgttcaaaa tgtaaatgca tcatcatcgg aagcattttt aattggggct aactatcttg 480
 gcaagccgaa 490

<210> SEQ ID NO 132
 <211> LENGTH: 550
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 132

taaggagaat caaatcaatg atatgattta ttctcttctg gaaaaaggta ggcttatcat 60
 tagagaaaaa aacagagttg tggtttcaag tgatattctt gttacaact aaacgaacat 120
 gtttattttc ttattatttc ttactctcac tagtggtagt gaccttgacc ggtgcaccac 180
 ttttgatgat gttcaagctc ctaattacac tcaacatact tcatctatga ggggggttta 240
 ctatcctgat gaaattttta gatcagacac tctttattta actcaggatt tatttcttcc 300
 attttattct aatggtacag ggtttcatac tattaatcat acgtttggca accctgtcat 360
 accttttaag gatggtatct attttgctgc cacagagaaa tcaaatgttg tccgtgggtg 420
 ggtttttggt tctaccatga acaacaagtc acagtcggtg attattatta acaattctac 480
 taatgttgtt atacgagcat gtaactttga attgtgtgac aacctttctt ttgctgtttc 540
 taaaccata 550

<210> SEQ ID NO 133
 <211> LENGTH: 490
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 133

acttaaatac acttacttta gctgtaccct acaacatgag agttattcac tttggtgctg 60
 gctctgataa aggagttgca ccaggtagag ctgtgctcag acaatggttg ccaactggca 120
 cactacttgt cgattcagat cttaatgact tegtctccga cgcagattct actttaattg 180
 gagactgtgc aacagtagat acggctaata aatgggacct tattattagc gatatgtatg 240
 accctaggac caaacatgtg acaaaagaga atgactctaa agaagggttt ttcacttate 300
 tgtgtggatt tataaagcaa aaactagccc tgggtgggtc tatagctgta aagataacag 360
 agcattcttg gaatgctgac ctttacaagc ttatgggcca tttctcatgg tggacagcct 420

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ttgttacaaa tgtaaatgca tcatcatcgg aagcattttt aattggggct aactatcttg 480
gcaagccgaa 490

```

```

<210> SEQ ID NO 134
<211> LENGTH: 550
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 134
taaggagaat caaatcaatg atatgattta ttctcttctg gaaaaaggta ggcttatcat 60
tagagaaaac aacagagttg tggtttcaag tgatattctt gttaacaact aacgaacat 120
gtttattttc ttattatttc ttactctcac tagtggtagt gaccttgacc ggtgcaccac 180
ttttgatgat gttcaagctc ctaattacac tcaacatact tcatctatga ggggggttta 240
ctatcctgat gaaattttta gatcagacac tctttattta actcaggatt tattttcttc 300
attttattct aatgttacag ggtttcatal tattaatcat acgtttggca accctgtcat 360
accttttaag gatggatttt attttgctgc cacagagaaa tcaaatgttg tccgtgggtg 420
ggtttttggg tctaccatga acaacaagtc acagtcgggtg attattatta acaattctac 480
taatgtgttt atacgagcat gtaactttga attgtgtgac aacctttctt ttgctgtttc 540
taaaccata 550

```

```

<210> SEQ ID NO 135
<211> LENGTH: 400
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 135
atcaatgata tgatttattc tcttctggaa aaaggtaggc ttatcattag agaaaacaac 60
agagttgtgg tttcaagtga tattcttggt aacaactaaa cgaacatggt tattttctta 120
ttattttcta ctctcactag tggtagtgac cttgaccggg gcaccacttt tgatgatggt 180
caagtcctca attacactca acatacttca tctatgaggg gggtttacta tctgatgaa 240
atttttagat cagacactct ttatttaact caggatttat ttcttcatt ttattctaat 300
gttacagggg tctactat taatcatacg tttggcaacc ctgtcatacc ttttaaggat 360
ggtatttatt ttgctgccac agagaaatca aatgttgtcc 400

```

```

<210> SEQ ID NO 136
<211> LENGTH: 288
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 136
tgatctttgc ttctccaatg tctatgcaga ttctttggta gtcaaggag atgatgtaag 60
acaaatagcg ccaggacaaa ctgggtgtat tgctgattat aattataaat tgccagatga 120
ttctatgggt tgtgtccttg cttggaatac taggaacatt gatgctactt caactggtaa 180
ttataattat aatataggtt atcttagaca tggcaagctt aggcctttg agagagacat 240
atctaagtgt cctttctcca cctgatggca aacettgcac cccacctg 288

```

```

<210> SEQ ID NO 137
<211> LENGTH: 411
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 137

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ctttgagaga gacatatcta atgtgccttt ctcccctgat ggcaaacctt gcaccccacc 60
tgctcttaat tgttattggc cattaaatga ttatggtttt tacaccacta ctggcattgg 120
ctaccaacct tacagagttg tagtactttc ttttgaactt taaatgcac cggccacggt 180
ttgtggacca aaattatcca ctgaccttat taagaaccag tgtgtcaatt ttaattttaa 240
tggactcact ggtactgggt tgtaactcc ttcttcaaag agatttcaac catttcaaca 300
aattttgccg tgatgtttct gatttcaactg attccggtcg agatcctaaa acatctgaaa 360
tattagacat ttcacctgc gcttttgggg gtgtaagtgt aattacacct g 411

```

```

<210> SEQ ID NO 138
<211> LENGTH: 357
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 138

```

```

tggaaatatt ttggtggttt taatttttca caaatattac ctgaccctct aaagccaact 60
aagaggtcct ttattgagga ctgtctcttt aataaggtga cactcgtga tgctggcttc 120
atgaagcaat atggcgaatg cctaggtgat attaagtcta gagatctcat ttgtgcgcag 180
aagttcaatg gacttacagt gttgccacct ctgctcactg atgatatgat tgctgcctac 240
actgctgctc tagtttagtg tactgccact gctggatgga catttggtgc tggcgtgct 300
cttcaaatc cttttgctat gcaaatggca tataggttca atggcattgg agttact 357

```

```

<210> SEQ ID NO 139
<211> LENGTH: 434
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 139

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```

caatatggcg aatgcttagg tgatattaat gctagagatc tcatttgtgc gcagaagttc 60
aatggactta cagtgttgcc acctctgctc actgatgata tgattgctgc ctacactgct 120
gctctagtta gtggtactgc cactgctgga tggacatttg gtgctggcgc tgctcttcaa 180
ataccttttg ctatgcaaat ggcatatagg ttcaatggca ttggagtac ccaaaatggt 240
ctctatgaga accaaaaaca aatcgccaac caatttaaca aggcgattag tcaaatcaa 300
gaatcactta caacaacatc aactgcattg ggcaagctgc aagacgttgt taaccagaat 360
gctcaagcat taaacacact tgtaaacaa cttagcteta attttggtgc aatttcaagt 420
gtgctaatag atat 434

```

```

<210> SEQ ID NO 140
<211> LENGTH: 557
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 140

```

```

acagacaata ctttgtctc aggaaattgt gatgtcgtta ttggcatcat taacaacaca 60
gtttatgac ctctgcaacc tgagcttgac tcattcaaag aagagctgga caagtacttc 120
aaaaatcata catcaccaga tgttgatctt ggcgacattt caggcattaa cgcttctgtc 180
gtcaacatte aaaaagaaat tgaccgcctc aatgaggtcg ctaaaaattt aaatgaatca 240
ctcattgacc ttcaagaatt gggaaaatat gagcaatata taaatggcc ttggtatggt 300
tggctcggct tcattgctgg actaattgcc atcgtcatgg ttacaatctt gctttgttgc 360

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atgactagtt gttgcagttg cctcaagggg gcatgctctt gtggttcttg ctgcaagttt 420
gatgaggatg actctgagcc agttctcaag ggtgtcaaat tacattacac ataaacgaac 480
ttatggattt gtttatgaga ttttttactc ttagatcaat tactgcacag ccagtaaaaa 540
ttgacaatgc ttctcct 557

```

```

<210> SEQ ID NO 141
<211> LENGTH: 530
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 141

```

```

atgtttggtt cggcttcatt gctggactaa ttgccatcgt catggttaca atcttgcttt 60
gttgcagatg tagttgttgc agttgctca aggggtgatg ctcttggtgt tcttgctgca 120
agtttgatga ggatgactct gagccagttc tcaaggggtg caaattacat tacacataaa 180
cgaacttatg gatttgttta tgagattttt tactcttaga tcaattactg cacagccagt 240
aaaaattgac aatgcttctc ctgcaagtac tgttcagctc acagcaacga taccgctaca 300
agcctcactc cctttcggat ggcttggtat tggcgttgca tttcttgctg tttttcagag 360
cgctacaaa ataattgcgc tcaataaaaag atggcagcta gccctttata agggcttcca 420
gttcatttgc aatttactgc tgctatttgt taccatctat tcacatcttt tgcttgctgc 480
tgcaggtatg gaggcgcaat ttttgtacct ctatgccttg atatattttc 530

```

```

<210> SEQ ID NO 142
<211> LENGTH: 320
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 142

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```

ttgctcgtac ccgctcaatg tggtcattca acccagaaac aaacattctt ctcaatgtgc 60
ctctccgggg gacaattgtg accagaccgc tcatgaaag tgaacttgtc attggtgctg 120
tgatcattcg tggtcacttg cgaatggcgc gacctcctt agggcgctgt gacattaagg 180
acctgcaaaa agagatcact gtggctacat cacgaacgct ttcttattac aaattaggag 240
cgtcgcagcg ttaggcact gattcaggtt ttgctgcata caaccgctac cgtattggaa 300
actataaatt aaatacagac 320

```

```

<210> SEQ ID NO 143
<211> LENGTH: 417
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

```

```

<400> SEQUENCE: 143

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```

cgaacttatg tactcattcg tttcggaga aacaggtacg ttaatagtta atagcgtact 60
tctttttctt gctttcgtgg tattcttctt agtcacacta gccatcetta ctgcgcttcg 120
attgtgtgcg tactgctgca atattgttaa cgtgagttta gtaaaaccaa cggtttacgt 180
ctactcgcgt gttaaaaatc tgaactcttc tgaaggagtt cctgatcttc tggctctaac 240
gaaactaacta ttattattat tctgtttgga actttaacat tgcttatcat ggcagacaac 300
ggtactatta ccggtgagga gcttaacaa ctctggaac aatggaacct agtaaatagg 360
ttctatttcc tagcctggat tatgttacta caatttgctt attctaactg gaacagg 417

```

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<210> SEQ ID NO 144
<211> LENGTH: 516

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<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 144

cttgtcattg gtgctgtgat cattcgtggt cacttgcgaa tggccgaca ctccttagg 60
cgctgtgaca ttaaggacct gccaaaagag atcactgtgg ctacatcacg aacgctttct 120
tattacaaat taggagcgtc gcagcgtgta ggcactgatt caggttttgc tgcatacaac 180
cgctaccgta ttggaaacta taaattaaat acagaccacg ccggtagcaa cgacaatatt 240
gctttgctag tacagtaagt gacaacagat gtttcatctt gttgacttcc aggttacaat 300
agcagagata ttgattatca ttatgaggac tttcaggatt gctatttggga atcttgacgt 360
tataataagt tcaatagtga gacaattatt taagcctcta actaagaaga attattcgga 420
gttagatgat gaagaaccta tggagttaga ttatccataa aacgaacatg aaaattattc 480
tcttctgac attgatttta tttacatctt gcgagc 516

```

```

<210> SEQ ID NO 145
<211> LENGTH: 310
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 145

cgatgtttca tcttgttgac ttccaggta caatagcaga gatattgatt atcattatga 60
ggactttcag gattgctatt tggatcttg acgttataat aagttcaata gtgagacaat 120
tatttaagcc tctaactaag aagaattatt cggagttaga tgatgaagaa cctatggagt 180
tagattatcc ataaaacgaa catgaaaatt attctcttcc tgacattgat tgtatttaca 240
tcttgcgagc tatatcacta tcaggagtgt gtttagaggta cgactgtact actaaaagaa 300
ccttgcccat 310

```

```

<210> SEQ ID NO 146
<211> LENGTH: 556
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 146

agaaagacag aatgaatgag ctcactttaa ttgacttcta tttgtgcttt ttagcctttc 60
tgctattcct tgttttaata atgcttatta tattttggtt ttcactcgaa atccaggatc 120
tagaagaacc ttgtacaaa gtctaaacga acatgaaact tctcattggt ttgacttgta 180
tttctctatg cagttgcata tgcactgtag tacagcgtcg tgcattctaat aaacctcatg 240
tgcttgaaga tccttgaag gtacaacact aggggtaata cttatagcac tgcttggctt 300
tgtgctctag gaaaggtttt accttttcat agatggcaca ctatggttca aacatgcaca 360
cctaattgta ctatcaactg tcaagatcca gctggtggtg cgcttatagc taggtgttgg 420
taccttcatg aaggtcacca aactgctgca tttagagacg tacttgttgt tttaaataaa 480
cgaacaaatt aaaatgtctg ataatggacc ccaatcaaac caacgtagtg cccccgcac 540
tacatttggg ggacct 556

```

```

<210> SEQ ID NO 147
<211> LENGTH: 110
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 147

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acgaacatga aaattattct cttcctgaca ttgattgtat ttacatcttg cgagctatat 60

cactatcagg agtgtgtag aggtacgact gtactactaa aagaacctg 110

<210> SEQ ID NO 148

<211> LENGTH: 363

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 148

gcatttagag acgtacttgt tgttttaaat aaacgaacaa attaaaatgt ctgataatgg 60

acctcaatca agccaacgta gtgcccccg cattacattt ggtggacca cagattcaac 120

tgacaataac cagaatggag gacgcaatgg ggcaaggcca aaacagcgcc gacccaagg 180

tttaccat aatactgcgt cttggttcac agctctcact cagcatggca aggaggaact 240

tagattccct cgaggccagg gcgttccaat caacaccaat agtggccag atgaccaaat 300

tggctactac cgaagagcta cccgacgagt tcgtggtggt gacggcaaaa tgaagagct 360

cag 363

<210> SEQ ID NO 149

<211> LENGTH: 294

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 149

ctatcagctg cgtgcaagat cagtttcacc aaaacttttc atcagacaag aggaggttca 60

acaagagctc tactcgccac tttttctcat tgttctgct ctagtatttt taatactttg 120

cttcaccatt aagagaaaga cagaatgaat gagctcactt taattgactt ctatttgtgc 180

tttttagcct ttctgctatt ccttgtttta ataatgctta ttatattttg gttttcactc 240

gaaatccagg atctagaaaa acctgttacc aaaggctaaa cgaacatgaa actt 294

<210> SEQ ID NO 150

<211> LENGTH: 504

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 150

caaaactgctg catttagaga cgtacttgtt gtttaataa acgaacaaat taaaatgtct 60

gataatggac cccaatcaaa ccaacgtagt gcccccgca ttacatttgg tggaccaca 120

gattcaactg acaataacca gaatggagga cgcaatgggg caaggccaaa acagcgccga 180

ccccaaaggtt taccacaata tactgctct tggttcacag ctctcactca gcatggcaag 240

gaggaactta gattccctcg aggccagggc gttccaatca acaccaatag tggccagat 300

gaccaaattg gctactaccg aagagctacc cgacgagttc gtggtggtga cggcaaaatg 360

aaagagctca gcccagatg gtacttctat tacctaggaa ctggcccaga agcttcactt 420

ccctacggcg ctaacaaaga aggcacgta tgggttgcaa ctgagggagc cttgaataca 480

cccaaagacc acattggcac ccgt 504

<210> SEQ ID NO 151

<211> LENGTH: 474

<212> TYPE: DNA

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 151

ctcgccactt tttctcattg ttgctgctct agtattttta atactttgct tcaccattaa 60

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gagaaagaca gaatgaatga gctcacttta attgacttct atttgtgctt tttagccttt 120
ctgctattcc ttgttttaaat aatgcttatt atatttttggg tttcactcga aatccaggat 180
ctagaagaac ctgttaccaa agtctaaacg aacatgaaac ttctcattgt tttgacttgt 240
atcttctctat gcagttgcat atgcactgta gtacagcgct gtgcatctaa taaacctcat 300
gtgcttgaag atccttgtaa ggtacaacac taggggtaat acttatagca ctgcttggct 360
ttgtgctcta ggaaaggttt taccttttca tagatggcac actatggttc aaacatgcac 420
acctaagtgt actatcaact gtcaagatcc agctgggtggc gcgcttatag ctg 474

```

```

<210> SEQ ID NO 152
<211> LENGTH: 516
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 152

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```

cattaagaga aagacagaat gaatgagctc actttaattg acttctatct gtgcttttta 60
gcctttctgc tattccttgt ttaataatg cttattatat tttggttttc actcgaaatc 120
caggatctag aagaaccttg taccaaaagt taaacgaaca tgaaccttct cattgttttg 180
acttgtatct ctctatgcag ttgcatatgc actgtagtac agcgctgtgc atctaataaa 240
cctcatgtgc ttgaagatcc ttgtaaggta caacactagg ggtaataact atagcactgc 300
ttggctttgt gctctaggaa aggttttacc ttttcataga tggcacacta tggttcaaac 360
atgcacacct aatgttacta tcaactgtca agatccagct ggtgggtgccc ttatagctag 420
gtgttggtac cttcatgaag gtcaccaaac tgctgcattt agagacgtac ttgttgtttt 480
aaataaacga acaaattaaa atgtctgata atggac 516

```

```

<210> SEQ ID NO 153
<211> LENGTH: 451
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 153

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```

ccaaggttta cccaataata ctgcttcttg gttcacagct ctactcagc atggcaagga 60
ggaacttaga ttccctcgag gccagggcgt tccaatcaac accaatagtg gtccagatga 120
ccaaattggc tactaccgaa gagctaccgg acgagttcgt ggtgggtgacg gcaaaatgaa 180
agagctcagc cccagatggt acttctatta cctaggaact ggcccagaag cttcacttcc 240
ctacggcgct aacaaagaag gcatcgtatg ggttgcaact gagggagcct tgaatacacc 300
caaagaccac attggcaccg gcaatcctaa taacaatgct gccaccgtgc tacaacttcc 360
tcaaggaaca acattgcca aaggcttcta cgcagaggga agcagaggcg gcagtcaagc 420
ctcttctcgc tcctcatcac gtagtgcggg t 451

```

```

<210> SEQ ID NO 154
<211> LENGTH: 495
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 154

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```

gatgaagctc agcctttgcc gcagagacaa aagaagcagc ccaactgtgac tcttcttct 60
gctgctgaca tggatgattt ctccagacaa cttcaaaatt ccatgagtgg agcttctgct 120
gattcaactc aggcataaac actcatgatg accacacaag gcagatgggc tatgtaaacy 180

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ttttcgcaat tccggtttacg atacatagtc tactcttgtg cagaatgaat tctcgtaact 240
aacagcaca agtaggttta gttaacttta atctcacata gcaatcttta atcaatgtgt 300
aacattaggg aggacttgaa agagccacca cattttcatc gaggccacgc ggagtacgat 360
cgagggtaca gtgaataatg ctaggggagag ctgcctatat ggaagagccc taatgtgtaa 420
aattaatfff agtagtgcta tccccatgtg attttaatag cttcttagga gaatgacaaa 480
aaaaaaaaaa aaaaa 495

```

```

<210> SEQ ID NO 155
<211> LENGTH: 512
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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```

<400> SEQUENCE: 155

```

```

acaaggccaa actgtcacta agaaatctgc tgctgaggca tctaaaagc ctgcgcaaaa 60
acgtactgcc acaaaacagt acaacgtcac tcaagcattt gggagacgtg gtccagaaca 120
aacccaagga aatttcgggg accaagacct aatcagacaa ggaactgatt acaaacattg 180
gccgcaaatt gcacaatttg ctccaagtgc ctctgcattc tttggaatgt cacgcattgg 240
catggaagtc acaccttcgg gaacatggct gacttatcat ggagccatta aattggatga 300
caaagatcca caattcaaag acaacgtcat actgctgaac aagcacattg acgcatacaa 360
aacattccca ccaacagagc ctaaaaagga caaaaagaaa aagactgatg aagctcagcc 420
tttgccgag agacaaaaga agcagcccac tgtgactctt cttcctgcgg ctgatatgga 480
tgatttctcc agacaacttc aaaattccat ga 512

```

```

<210> SEQ ID NO 156
<211> LENGTH: 442
<212> TYPE: DNA
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 156

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tgtgactctt cttcctgcgg ctgatatgga tgtttctcca gacaacttca aaattccatg 60
agtggagctt ctgctgattc aactcaggca taaacactca tgatgaccac acaaggcaga 120
tgggctatgt aaacgttttc gcaattccgt ttacgatata tagtctactc ttgtgcagaa 180
tgaattctcg taactaaaca gcacaagtag gtttagttaa ctttaatctc acatagcaat 240
ctttaatcaa tgtgtaacat tagggaggac ttgaaagagc caccacattt tcatcgaggc 300
cacgcgaggt acgatcgagg gtacagtga taatgctagg gagagctgcc tatatggaag 360
agccctaagt tgtaaaaatta atttttagtag tgctatcccc atgtgatfff aatagcttct 420
taggagaatg acaaaaaaaaa aa 442

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<210> SEQ ID NO 157
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

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<400> SEQUENCE: 157

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atgaattacc aagtcaatgg ttac 24

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<210> SEQ ID NO 158
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 158

gaagctattc gtcacgttcg                20

<210> SEQ ID NO 159
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 159

ctgtagaaaa tcctagctgg ag            22

<210> SEQ ID NO 160
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 160

cataaccagt cggtacagct a             21

<210> SEQ ID NO 161
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 161

ttatcaccgc cgaagaagct              20

<210> SEQ ID NO 162
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 162

ctctagttgc atgacagccc tc           22

<210> SEQ ID NO 163
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 163

tcgtgcgtgg attggctttg atgt        24

<210> SEQ ID NO 164
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 164

gggttgggac taccctaagt gtga        24

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<210> SEQ ID NO 165
 <211> LENGTH: 22
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

 <400> SEQUENCE: 165

 taacacacaa acaccatcat ca 22

<210> SEQ ID NO 166
 <211> LENGTH: 23
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

 <400> SEQUENCE: 166

 gggtgggact atcctaagtg tga 23

<210> SEQ ID NO 167
 <211> LENGTH: 24
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

 <400> SEQUENCE: 167

 ccatcatcag atagaatcat cata 24

<210> SEQ ID NO 168
 <211> LENGTH: 21
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

 <400> SEQUENCE: 168

 cctctcttgt tcttgctgc a 21

<210> SEQ ID NO 169
 <211> LENGTH: 21
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

 <400> SEQUENCE: 169

 tatagtgagc cgccacacat g 21

<210> SEQ ID NO 170
 <211> LENGTH: 21
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer
 <220> FEATURE:
 <221> NAME/KEY: misc_feature
 <222> LOCATION: (12)..(12)
 <223> OTHER INFORMATION: n is a, c, g, or t

 <400> SEQUENCE: 170

 taacacacaa cncatcatc a 21

<210> SEQ ID NO 171
 <211> LENGTH: 21

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 171

ctaacatgct taggataatg g 21

<210> SEQ ID NO 172
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 172

gcctctcttg ttcttgcteg c 21

<210> SEQ ID NO 173
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 173

caggtaagcg taaaactcat c 21

<210> SEQ ID NO 174
<211> LENGTH: 17
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 174

tacacacctc agcggtg 17

<210> SEQ ID NO 175
<211> LENGTH: 16
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 175

cacgaacgtg acgaat 16

<210> SEQ ID NO 176
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 176

gccggagctc tgcagaattc 20

<210> SEQ ID NO 177
<211> LENGTH: 47
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 177

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caggaaacag ctatgacttg catcaccact agttgtgccca ccaggtt 47

<210> SEQ ID NO 178
 <211> LENGTH: 46
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

<400> SEQUENCE: 178

tgtaaaaacga cggccagttg atgggatggg actatcctaa gtgtga 46

<210> SEQ ID NO 179
 <211> LENGTH: 20
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Primer

<400> SEQUENCE: 179

gcataggcag tagttgcatc 20

<210> SEQ ID NO 180
 <211> LENGTH: 8
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: ATP Binding Domain
 <220> FEATURE:
 <221> NAME/KEY: MISC_FEATURE
 <222> LOCATION: (1)..(1)
 <223> OTHER INFORMATION: Xaa = A or G
 <220> FEATURE:
 <221> NAME/KEY: misc_feature
 <222> LOCATION: (2)..(5)
 <223> OTHER INFORMATION: Xaa can be any naturally occurring amino acid
 <220> FEATURE:
 <221> NAME/KEY: MISC_FEATURE
 <222> LOCATION: (8)..(8)
 <223> OTHER INFORMATION: Xaa = S or T

<400> SEQUENCE: 180

Xaa Xaa Xaa Xaa Xaa Gly Lys Xaa
 1 5

<210> SEQ ID NO 181
 <211> LENGTH: 23
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 181

Trp Tyr Val Trp Leu Gly Phe Ile Ala Gly Leu Ile Ala Ile Val Met
 1 5 10 15

Val Thr Ile Leu Leu Cys Cys
 20

<210> SEQ ID NO 182
 <211> LENGTH: 16
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 182

Met Asp Leu Phe Met Arg Phe Phe Thr Leu Arg Ser Ile Thr Ala Gln
 1 5 10 15

<210> SEQ ID NO 183
 <211> LENGTH: 150

-continued

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<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 183

Met Arg Cys Trp Leu Cys Trp Lys Cys Lys Ser Lys Asn Pro Leu Leu
1          5          10          15

Tyr Asp Ala Asn Tyr Phe Val Cys Trp His Thr His Asn Tyr Asp Tyr
          20          25          30

Cys Ile Pro Tyr Asn Ser Val Thr Asp Thr Ile Val Val Thr Glu Gly
          35          40          45

Asp Gly Ile Ser Thr Pro Lys Leu Lys Glu Asp Tyr Gln Ile Gly Gly
          50          55          60

Tyr Ser Glu Asp Arg His Ser Gly Val Lys Asp Tyr Val Val Val His
65          70          75          80

Gly Tyr Phe Thr Glu Val Tyr Tyr Gln Leu Glu Ser Thr Gln Ile Thr
          85          90          95

Thr Asp Thr Gly Ile Glu Asn Ala Thr Phe Phe Ile Phe Asn Lys Leu
          100          105          110

Val Lys Asp Pro Pro Asn Val Gln Ile His Thr Ile Asp Gly Ser Ser
          115          120          125

Gly Val Ala Asn Pro Ala Met Asp Pro Ile Tyr Asp Glu Pro Thr Thr
          130          135          140

Thr Thr Ser Val Pro Leu
145          150

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<210> SEQ ID NO 184
<211> LENGTH: 20
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 184

Met Met Pro Thr Thr Leu Phe Ala Gly Thr His Ile Thr Met Thr Thr
1          5          10          15

Val Tyr His Ile
          20

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<210> SEQ ID NO 185
<211> LENGTH: 42
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 185

Thr Ala Leu Arg Leu Cys Ala Tyr Cys Cys Asn Ile Val Asn Val Ser
1          5          10          15

Leu Val Lys Pro Thr Val Tyr Val Tyr Ser Arg Val Lys Asn Leu Asn
          20          25          30

Ser Ser Glu Gly Val Pro Asp Leu Leu Val
          35          40

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<210> SEQ ID NO 186
<211> LENGTH: 39
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 186

Met Ala Asp Asn Gly Thr Ile Thr Val Glu Glu Leu Lys Gln Leu Leu
1          5          10          15

Glu Gln Trp Asn Leu Val Ile Gly Phe Leu Phe Leu Ala Trp Ile Met
          20          25          30

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Leu Leu Gln Phe Ala Tyr Ser
35

<210> SEQ ID NO 187
<211> LENGTH: 100
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 187

Pro Leu Arg Gly Thr Ile Val Thr Arg Pro Leu Met Glu Ser Glu Leu
1 5 10 15
Val Ile Gly Ala Val Ile Ile Arg Gly His Leu Arg Met Ala Gly His
20 25 30
Ser Leu Gly Arg Cys Asp Ile Lys Asp Leu Pro Lys Glu Ile Thr Val
35 40 45
Ala Thr Ser Arg Thr Leu Ser Tyr Tyr Lys Leu Gly Ala Ser Gln Arg
50 55 60
Val Gly Thr Asp Ser Gly Phe Ala Ala Tyr Asn Arg Tyr Arg Ile Gly
65 70 75 80
Asn Tyr Lys Leu Asn Thr Asp His Ala Gly Ser Asn Asp Asn Ile Ala
85 90 95

Leu Leu Val Gln
100

<210> SEQ ID NO 188
<211> LENGTH: 23
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 188

Phe Tyr Leu Cys Phe Leu Ala Phe Leu Leu Phe Leu Val Leu Ile Met
1 5 10 15
Leu Ile Ile Phe Trp Phe Ser
20

<210> SEQ ID NO 189
<211> LENGTH: 19
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 189

Leu Leu Ile Val Leu Thr Cys Ile Ser Leu Cys Ser Cys Ile Cys Thr
1 5 10 15
Val Val Gln

<210> SEQ ID NO 190
<211> LENGTH: 24
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 190

Ile Cys Thr Val Val Gln Arg Cys Ala Ser Asn Lys Pro His Val Leu
1 5 10 15
Glu Asp Pro Cys Lys Val Gln His
20

<210> SEQ ID NO 191
<211> LENGTH: 22
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

-continued

<400> SEQUENCE: 191

Cys Ile Cys Thr Val Val Gln Arg Cys Ala Ser Asn Lys Pro His Val
 1 5 10 15

Leu Glu Asp Pro Cys Lys
 20

<210> SEQ ID NO 192

<211> LENGTH: 22

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 192

Val Val Ala Val Ile Gln Glu Ile Gln Leu Leu Ala Ala Val Gly Glu
 1 5 10 15

Ile Leu Leu Leu Glu Trp
 20

<210> SEQ ID NO 193

<211> LENGTH: 19

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Linker

<400> SEQUENCE: 193

aattcgcggc cgcgtcgac

19

<210> SEQ ID NO 194

<211> LENGTH: 15

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Linker

<400> SEQUENCE: 194

gtcgcgcggg ccgcg

15

<210> SEQ ID NO 195

<211> LENGTH: 19

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 195

aattcgcggc cgcgtcgac

19

<210> SEQ ID NO 196

<211> LENGTH: 19

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Primer

<400> SEQUENCE: 196

ggcctcttcg ctattacgc

19

<210> SEQ ID NO 197

<211> LENGTH: 21

<212> TYPE: DNA

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Primer

-continued

<400> SEQUENCE: 197

tgcaggtcga ctctagagga t

21

<210> SEQ ID NO 198

<211> LENGTH: 410

<212> TYPE: PRT

<213> ORGANISM: Avian infectious bronchitis virus

<400> SEQUENCE: 198

Met Ala Ser Gly Lys Ala Ala Gly Lys Thr Asp Ala Pro Ala Pro Val
1 5 10 15

Ile Lys Leu Gly Gly Pro Lys Pro Pro Lys Val Gly Ser Ser Gly Asn
20 25 30

Ala Ser Trp Phe Gln Ala Ile Lys Ala Lys Lys Leu Asn Thr Pro Pro
35 40 45

Pro Lys Phe Glu Gly Ser Gly Val Pro Asp Asn Glu Asn Ile Lys Pro
50 55 60

Ser Gln Gln His Gly Tyr Trp Arg Arg Gln Ala Arg Phe Lys Pro Gly
65 70 75 80

Lys Gly Gly Arg Lys Pro Val Pro Asp Ala Trp Tyr Phe Tyr Tyr Thr
85 90 95

Gly Thr Gly Pro Ala Ala Asp Leu Asn Trp Gly Asp Thr Gln Asp Gly
100 105 110

Ile Val Trp Val Ala Ala Lys Gly Ala Asp Thr Lys Ser Arg Ser Asn
115 120 125

Gln Gly Thr Arg Asp Pro Asp Lys Phe Asp Gln Tyr Pro Leu Arg Phe
130 135 140

Ser Asp Gly Gly Pro Asp Gly Asn Phe Arg Trp Asp Phe Ile Pro Leu
145 150 155 160

Lys Asn Arg Gly Arg Ser Gly Arg Ser Thr Ala Ala Ser Ser Ala Ala
165 170 175

Ala Ser Arg Ala Pro Ser Arg Glu Gly Ser Arg Gly Arg Arg Ser Asp
180 185 190

Ser Gly Asp Asp Leu Ile Ala Arg Ala Ala Lys Ile Ile Gln Asp Gln
195 200 205

Gln Lys Lys Gly Ser Arg Ile Thr Lys Ala Lys Ala Asp Glu Met Ala
210 215 220

His Arg Arg Tyr Cys Lys Arg Thr Ile Pro Pro Asn Tyr Arg Val Asp
225 230 235 240

Gln Val Phe Gly Pro Arg Thr Lys Gly Lys Glu Gly Asn Phe Gly Asp
245 250 255

Asp Lys Met Asn Glu Glu Gly Ile Lys Asp Gly Arg Val Thr Ala Met
260 265 270

Leu Asn Leu Val Pro Ser Ser His Ala Cys Leu Phe Gly Ser Arg Val
275 280 285

Thr Pro Lys Leu Gln Leu Asp Gly Leu His Leu Arg Phe Glu Phe Thr
290 295 300

Thr Val Val Pro Cys Asp Asp Pro Gln Phe Asp Asn Tyr Val Lys Ile
305 310 315 320

Cys Asp Gln Cys Val Asp Gly Val Gly Thr Arg Pro Lys Asp Asp Glu
325 330 335

Pro Lys Pro Lys Ser Arg Ser Ser Ser Arg Pro Ala Thr Arg Gly Asn
340 345 350

Ser Pro Ala Pro Arg Gln Gln Arg Pro Lys Lys Glu Lys Lys Leu Lys

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355	360	365
Lys Gln Asp Asp Glu Ala Asp Lys Ala Leu Thr Ser Asp Glu Glu Arg		
370	375	380
Asn Asn Ala Gln Leu Glu Phe Tyr Asp Glu Pro Lys Val Ile Asn Trp		
385	390	395
Gly Asp Ala Ala Leu Gly Glu Asn Glu Leu		
405	410	

<210> SEQ ID NO 199
 <211> LENGTH: 30
 <212> TYPE: PRT
 <213> ORGANISM: conotoxin

<400> SEQUENCE: 199

Cys Ile Ala Val Gly Gln Leu Cys Val Phe Trp Asn Ile Gly Arg Pro		
1	5	10
Cys Cys Ser Gly Leu Cys Val Phe Ala Cys Thr Val Lys Leu		
20	25	30

<210> SEQ ID NO 200
 <211> LENGTH: 31
 <212> TYPE: PRT
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 200

Cys Ile Ser Leu Cys Ser Cys Ile Cys Thr Val Val Gln Arg Cys Ala		
1	5	10
Ser Asn Lys Pro His Val Leu Glu Asp Pro Cys Lys Val Gln His		
20	25	30

<210> SEQ ID NO 201
 <211> LENGTH: 310
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 201

cgatgtttca tcttgttgac ttccaggta caatagcaga gatattgatt atcattatga	60
ggactttcag gattgctatt tggaatcttg acgttataat aagttcaata gtgagacaat	120
tatttaagcc tctaactaag aagaattatt cggagttaga tgatgaagaa cctatggagt	180
tagattatcc ataaaacgaa catgaaaatt attctcttcc tgacattgat tgtatttaca	240
tcttgcgagc tatatcacta tcaggagtgt gtttagaggta cgactgtact actaaaagaa	300
ccttgcccat	310

<210> SEQ ID NO 202
 <211> LENGTH: 556
 <212> TYPE: DNA
 <213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 202

agaaagacag aatgaatgag ctcactttaa ttgactteta tttgtgcttt ttagcctttc	60
tgctattcct tgttttaata atgcttatta tattttgggt ttcactcgaa atccaggatc	120
tagaagaacc ttgtacaaa gtctaaacga acatgaaact tctcattggt ttgacttgta	180
tttctctatg cagttgcata tgcaactgtag tacagcgtg tgcacttaac aaacctcatg	240
tgcttgaaga tccttgtaag gtacaacact aggggtaata cttatagcac tgcttggtt	300
tgtgctctag gaaaggtttt accttttcat agatggcaca ctatggttca aacatgcaca	360

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cctaagtta ctatcaactg tcaagatcca gctgggtggtg cgcttatagc taggtgttg 420
taccttcacg aaggtcacca aactgctgca tttagagacg tacttgtgtt ttaaataaa 480
cgaacaaatt aaatgtctg ataatggacc ccaatcaaac caacgtagtg cccccgcac 540
tacatttggg ggaccc 556

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<210> SEQ ID NO 203
<211> LENGTH: 1255
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

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<400> SEQUENCE: 203

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Met Phe Ile Phe Leu Leu Phe Leu Thr Leu Thr Ser Gly Ser Asp Leu
1 5 10 15
Asp Arg Cys Thr Thr Phe Asp Asp Val Gln Ala Pro Asn Tyr Thr Gln
20 25 30
His Thr Ser Ser Met Arg Gly Val Tyr Tyr Pro Asp Glu Ile Phe Arg
35 40 45
Ser Asp Thr Leu Tyr Leu Thr Gln Asp Leu Phe Leu Pro Phe Tyr Ser
50 55 60
Asn Val Thr Gly Phe His Thr Ile Asn His Thr Phe Gly Asn Pro Val
65 70 75 80
Ile Pro Phe Lys Asp Gly Ile Tyr Phe Ala Ala Thr Glu Lys Ser Asn
85 90 95
Val Val Arg Gly Trp Val Phe Gly Ser Thr Met Asn Asn Lys Ser Gln
100 105 110
Ser Val Ile Ile Ile Asn Asn Ser Thr Asn Val Val Ile Arg Ala Cys
115 120 125
Asn Phe Glu Leu Cys Asp Asn Pro Phe Phe Ala Val Ser Lys Pro Met
130 135 140
Gly Thr Gln Thr His Thr Met Ile Phe Asp Asn Ala Phe Asn Cys Thr
145 150 155 160
Phe Glu Tyr Ile Ser Asp Ala Phe Ser Leu Asp Val Ser Glu Lys Ser
165 170 175
Gly Asn Phe Lys His Leu Arg Glu Phe Val Phe Lys Asn Lys Asp Gly
180 185 190
Phe Leu Tyr Val Tyr Lys Gly Tyr Gln Pro Ile Asp Val Val Arg Asp
195 200 205
Leu Pro Ser Gly Phe Asn Thr Leu Lys Pro Ile Phe Lys Leu Pro Leu
210 215 220
Gly Ile Asn Ile Thr Asn Phe Arg Ala Ile Leu Thr Ala Phe Ser Pro
225 230 235 240
Ala Gln Asp Ile Trp Gly Thr Ser Ala Ala Ala Tyr Phe Val Gly Tyr
245 250 255
Leu Lys Pro Thr Thr Phe Met Leu Lys Tyr Asp Glu Asn Gly Thr Ile
260 265 270
Thr Asp Ala Val Asp Cys Ser Gln Asn Pro Leu Ala Glu Leu Lys Cys
275 280 285
Ser Val Lys Ser Phe Glu Ile Asp Lys Gly Ile Tyr Gln Thr Ser Asn
290 295 300
Phe Arg Val Val Pro Ser Gly Asp Val Val Arg Phe Pro Asn Ile Thr
305 310 315 320
Asn Leu Cys Pro Phe Gly Glu Val Phe Asn Ala Thr Lys Phe Pro Ser
325 330 335

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Val	Tyr	Ala	Trp	Glu	Arg	Lys	Lys	Ile	Ser	Asn	Cys	Val	Ala	Asp	Tyr
			340					345						350	
Ser	Val	Leu	Tyr	Asn	Ser	Thr	Phe	Phe	Ser	Thr	Phe	Lys	Cys	Tyr	Gly
		355					360					365			
Val	Ser	Ala	Thr	Lys	Leu	Asn	Asp	Leu	Cys	Phe	Ser	Asn	Val	Tyr	Ala
		370				375					380				
Asp	Ser	Phe	Val	Val	Lys	Gly	Asp	Asp	Val	Arg	Gln	Ile	Ala	Pro	Gly
385					390					395					400
Gln	Thr	Gly	Val	Ile	Ala	Asp	Tyr	Asn	Tyr	Lys	Leu	Pro	Asp	Asp	Phe
			405						410					415	
Met	Gly	Cys	Val	Leu	Ala	Trp	Asn	Thr	Arg	Asn	Ile	Asp	Ala	Thr	Ser
			420					425					430		
Thr	Gly	Asn	Tyr	Asn	Tyr	Lys	Tyr	Arg	Tyr	Leu	Arg	His	Gly	Lys	Leu
		435					440					445			
Arg	Pro	Phe	Glu	Arg	Asp	Ile	Ser	Asn	Val	Pro	Phe	Ser	Pro	Asp	Gly
		450				455					460				
Lys	Pro	Cys	Thr	Pro	Pro	Ala	Leu	Asn	Cys	Tyr	Trp	Pro	Leu	Asn	Asp
465					470					475					480
Tyr	Gly	Phe	Tyr	Thr	Thr	Thr	Gly	Ile	Gly	Tyr	Gln	Pro	Tyr	Arg	Val
				485					490					495	
Val	Val	Leu	Ser	Phe	Glu	Leu	Leu	Asn	Ala	Pro	Ala	Thr	Val	Cys	Gly
			500					505					510		
Pro	Lys	Leu	Ser	Thr	Asp	Leu	Ile	Lys	Asn	Gln	Cys	Val	Asn	Phe	Asn
		515					520					525			
Phe	Asn	Gly	Leu	Thr	Gly	Thr	Gly	Val	Leu	Thr	Pro	Ser	Ser	Lys	Arg
		530				535					540				
Phe	Gln	Pro	Phe	Gln	Gln	Phe	Gly	Arg	Asp	Val	Ser	Asp	Phe	Thr	Asp
545					550					555					560
Ser	Val	Arg	Asp	Pro	Lys	Thr	Ser	Glu	Ile	Leu	Asp	Ile	Ser	Pro	Cys
				565					570					575	
Ala	Phe	Gly	Gly	Val	Ser	Val	Ile	Thr	Pro	Gly	Thr	Asn	Ala	Ser	Ser
			580					585					590		
Glu	Val	Ala	Val	Leu	Tyr	Gln	Asp	Val	Asn	Cys	Thr	Asp	Val	Ser	Thr
		595					600					605			
Ala	Ile	His	Ala	Asp	Gln	Leu	Thr	Pro	Ala	Trp	Arg	Ile	Tyr	Ser	Thr
		610				615					620				
Gly	Asn	Asn	Val	Phe	Gln	Thr	Gln	Ala	Gly	Cys	Leu	Ile	Gly	Ala	Glu
625					630					635					640
His	Val	Asp	Thr	Ser	Tyr	Glu	Cys	Asp	Ile	Pro	Ile	Gly	Ala	Gly	Ile
				645					650					655	
Cys	Ala	Ser	Tyr	His	Thr	Val	Ser	Leu	Leu	Arg	Ser	Thr	Ser	Gln	Lys
		660						665					670		
Ser	Ile	Val	Ala	Tyr	Thr	Met	Ser	Leu	Gly	Ala	Asp	Ser	Ser	Ile	Ala
		675					680					685			
Tyr	Ser	Asn	Asn	Thr	Ile	Ala	Ile	Pro	Thr	Asn	Phe	Ser	Ile	Ser	Ile
		690				695					700				
Thr	Thr	Glu	Val	Met	Pro	Val	Ser	Met	Ala	Lys	Thr	Ser	Val	Asp	Cys
705					710					715					720
Asn	Met	Tyr	Ile	Cys	Gly	Asp	Ser	Thr	Glu	Cys	Ala	Asn	Leu	Leu	Leu
				725					730					735	
Gln	Tyr	Gly	Ser	Phe	Cys	Thr	Gln	Leu	Asn	Arg	Ala	Leu	Ser	Gly	Ile
			740					745					750		
Ala	Ala	Glu	Gln	Asp	Arg	Asn	Thr	Arg	Glu	Val	Phe	Ala	Gln	Val	Lys

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755					760					765					
Gln	Met	Tyr	Lys	Thr	Pro	Thr	Leu	Lys	Tyr	Phe	Gly	Gly	Phe	Asn	Phe
770					775					780					
Ser	Gln	Ile	Leu	Pro	Asp	Pro	Leu	Lys	Pro	Thr	Lys	Arg	Ser	Phe	Ile
785					790					795					800
Glu	Asp	Leu	Leu	Phe	Asn	Lys	Val	Thr	Leu	Ala	Asp	Ala	Gly	Phe	Met
				805					810					815	
Lys	Gln	Tyr	Gly	Glu	Cys	Leu	Gly	Asp	Ile	Asn	Ala	Arg	Asp	Leu	Ile
			820					825					830		
Cys	Ala	Gln	Lys	Phe	Asn	Gly	Leu	Thr	Val	Leu	Pro	Pro	Leu	Leu	Thr
		835					840					845			
Asp	Asp	Met	Ile	Ala	Ala	Tyr	Thr	Ala	Ala	Leu	Val	Ser	Gly	Thr	Ala
		850				855					860				
Thr	Ala	Gly	Trp	Thr	Phe	Gly	Ala	Gly	Ala	Ala	Leu	Gln	Ile	Pro	Phe
		865				870					875				880
Ala	Met	Gln	Met	Ala	Tyr	Arg	Phe	Asn	Gly	Ile	Gly	Val	Thr	Gln	Asn
				885					890						895
Val	Leu	Tyr	Glu	Asn	Gln	Lys	Gln	Ile	Ala	Asn	Gln	Phe	Asn	Lys	Ala
			900					905						910	
Ile	Ser	Gln	Ile	Gln	Glu	Ser	Leu	Thr	Thr	Thr	Ser	Thr	Ala	Leu	Gly
		915					920					925			
Lys	Leu	Gln	Asp	Val	Val	Asn	Gln	Asn	Ala	Gln	Ala	Leu	Asn	Thr	Leu
		930				935					940				
Val	Lys	Gln	Leu	Ser	Ser	Asn	Phe	Gly	Ala	Ile	Ser	Ser	Val	Leu	Asn
		945				950					955				960
Asp	Ile	Leu	Ser	Arg	Leu	Asp	Lys	Val	Glu	Ala	Glu	Val	Gln	Ile	Asp
				965					970						975
Arg	Leu	Ile	Thr	Gly	Arg	Leu	Gln	Ser	Leu	Gln	Thr	Tyr	Val	Thr	Gln
			980					985						990	
Gln	Leu	Ile	Arg	Ala	Ala	Glu	Ile	Arg	Ala	Ser	Ala	Asn	Leu	Ala	Ala
		995					1000					1005			
Thr	Lys	Met	Ser	Glu	Cys	Val	Leu	Gly	Gln	Ser	Lys	Arg	Val	Asp	
		1010				1015					1020				
Phe	Cys	Gly	Lys	Gly	Tyr	His	Leu	Met	Ser	Phe	Pro	Gln	Ala	Ala	
		1025				1030					1035				
Pro	His	Gly	Val	Val	Phe	Leu	His	Val	Thr	Tyr	Val	Pro	Ser	Gln	
		1040				1045					1050				
Glu	Arg	Asn	Phe	Thr	Thr	Ala	Pro	Ala	Ile	Cys	His	Glu	Gly	Lys	
		1055				1060					1065				
Ala	Tyr	Phe	Pro	Arg	Glu	Gly	Val	Phe	Val	Phe	Asn	Gly	Thr	Ser	
		1070				1075					1080				
Trp	Phe	Ile	Thr	Gln	Arg	Asn	Phe	Phe	Ser	Pro	Gln	Ile	Ile	Thr	
		1085				1090					1095				
Thr	Asp	Asn	Thr	Phe	Val	Ser	Gly	Asn	Cys	Asp	Val	Val	Ile	Gly	
		1100				1105					1110				
Ile	Ile	Asn	Asn	Thr	Val	Tyr	Asp	Pro	Leu	Gln	Pro	Glu	Leu	Asp	
		1115				1120					1125				
Ser	Phe	Lys	Glu	Glu	Leu	Asp	Lys	Tyr	Phe	Lys	Asn	His	Thr	Ser	
		1130				1135					1140				
Pro	Asp	Val	Asp	Leu	Gly	Asp	Ile	Ser	Gly	Ile	Asn	Ala	Ser	Val	
		1145				1150					1155				
Val	Asn	Ile	Gln	Lys	Glu	Ile	Asp	Arg	Leu	Asn	Glu	Val	Ala	Lys	
		1160				1165					1170				

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Asn Leu Asn Glu Ser Leu Ile Asp Leu Gln Glu Leu Gly Lys Tyr
1175 1180 1185

Glu Gln Tyr Ile Lys Trp Pro Trp Tyr Val Trp Leu Gly Phe Ile
1190 1195 1200

Ala Gly Leu Ile Ala Ile Val Met Val Thr Ile Leu Leu Cys Cys
1205 1210 1215

Met Thr Ser Cys Cys Ser Cys Leu Lys Gly Ala Cys Ser Cys Gly
1220 1225 1230

Ser Cys Cys Lys Phe Asp Glu Asp Asp Ser Glu Pro Val Leu Lys
1235 1240 1245

Gly Val Lys Leu His Tyr Thr
1250 1255

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<210> SEQ ID NO 204

<211> LENGTH: 422

<212> TYPE: PRT

<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 204

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Met Ser Asp Asn Gly Pro Gln Ser Asn Gln Arg Ser Ala Pro Arg Ile
1 5 10 15

Thr Phe Gly Gly Pro Thr Asp Ser Thr Asp Asn Asn Gln Asn Gly Gly
20 25 30

Arg Asn Gly Ala Arg Pro Lys Gln Arg Arg Pro Gln Gly Leu Pro Asn
35 40 45

Asn Thr Ala Ser Trp Phe Thr Ala Leu Thr Gln His Gly Lys Glu Glu
50 55 60

Leu Arg Phe Pro Arg Gly Gln Gly Val Pro Ile Asn Thr Asn Ser Gly
65 70 75 80

Pro Asp Asp Gln Ile Gly Tyr Tyr Arg Arg Ala Thr Arg Arg Val Arg
85 90 95

Gly Gly Asp Gly Lys Met Lys Glu Leu Ser Pro Arg Trp Tyr Phe Tyr
100 105 110

Tyr Leu Gly Thr Gly Pro Glu Ala Ser Leu Pro Tyr Gly Ala Asn Lys
115 120 125

Glu Gly Ile Val Trp Val Ala Thr Glu Gly Ala Leu Asn Thr Pro Lys
130 135 140

Asp His Ile Gly Thr Arg Asn Pro Asn Asn Asn Ala Ala Thr Val Leu
145 150 155 160

Gln Leu Pro Gln Gly Thr Thr Leu Pro Lys Gly Phe Tyr Ala Glu Gly
165 170 175

Ser Arg Gly Gly Ser Gln Ala Ser Ser Arg Ser Ser Ser Arg Ser Arg
180 185 190

Gly Asn Ser Arg Asn Ser Thr Pro Gly Ser Ser Arg Gly Asn Ser Pro
195 200 205

Ala Arg Met Ala Ser Gly Gly Gly Glu Thr Ala Leu Ala Leu Leu Leu
210 215 220

Leu Asp Arg Leu Asn Gln Leu Glu Ser Lys Val Ser Gly Lys Gly Gln
225 230 235 240

Gln Gln Gln Gly Gln Thr Val Thr Lys Lys Ser Ala Ala Glu Ala Ser
245 250 255

Lys Lys Pro Arg Gln Lys Arg Thr Ala Thr Lys Gln Tyr Asn Val Thr
260 265 270

Gln Ala Phe Gly Arg Arg Gly Pro Glu Gln Thr Gln Gly Asn Phe Gly

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275					280					285					
Asp	Gln	Asp	Leu	Ile	Arg	Gln	Gly	Thr	Asp	Tyr	Lys	His	Trp	Pro	Gln
	290					295					300				
Ile	Ala	Gln	Phe	Ala	Pro	Ser	Ala	Ser	Ala	Phe	Phe	Gly	Met	Ser	Arg
305					310					315					320
Ile	Gly	Met	Glu	Val	Thr	Pro	Ser	Gly	Thr	Trp	Leu	Thr	Tyr	His	Gly
				325					330					335	
Ala	Ile	Lys	Leu	Asp	Asp	Lys	Asp	Pro	Gln	Phe	Lys	Asp	Asn	Val	Ile
			340					345					350		
Leu	Leu	Asn	Lys	His	Ile	Asp	Ala	Tyr	Lys	Thr	Phe	Pro	Pro	Thr	Glu
		355					360					365			
Pro	Lys	Lys	Asp	Lys	Lys	Lys	Lys	Thr	Asp	Glu	Ala	Gln	Pro	Leu	Pro
	370					375					380				
Gln	Arg	Gln	Lys	Lys	Gln	Pro	Thr	Val	Thr	Leu	Leu	Pro	Ala	Ala	Asp
385					390					395					400
Met	Asp	Asp	Phe	Ser	Arg	Gln	Leu	Gln	Asn	Ser	Met	Ser	Gly	Ala	Ser
				405					410					415	
Ala	Asp	Ser	Thr	Gln	Ala										
			420												
<210> SEQ ID NO 205															
<211> LENGTH: 221															
<212> TYPE: PRT															
<213> ORGANISM: Sars associated coronavirus															
<400> SEQUENCE: 205															
Met	Ala	Asp	Asn	Gly	Thr	Ile	Thr	Val	Glu	Glu	Leu	Lys	Gln	Leu	Leu
1			5						10					15	
Glu	Gln	Trp	Asn	Leu	Val	Ile	Gly	Phe	Leu	Phe	Leu	Ala	Trp	Ile	Met
		20					25						30		
Leu	Leu	Gln	Phe	Ala	Tyr	Ser	Asn	Arg	Asn	Arg	Phe	Leu	Tyr	Ile	Ile
		35					40					45			
Lys	Leu	Val	Phe	Leu	Trp	Leu	Leu	Trp	Pro	Val	Thr	Leu	Ala	Cys	Phe
	50					55					60				
Val	Leu	Ala	Ala	Val	Tyr	Arg	Ile	Asn	Trp	Val	Thr	Gly	Gly	Ile	Ala
65				70					75					80	
Ile	Ala	Met	Ala	Cys	Ile	Val	Gly	Leu	Met	Trp	Leu	Ser	Tyr	Phe	Val
			85					90						95	
Ala	Ser	Phe	Arg	Leu	Phe	Ala	Arg	Thr	Arg	Ser	Met	Trp	Ser	Phe	Asn
			100				105						110		
Pro	Glu	Thr	Asn	Ile	Leu	Leu	Asn	Val	Pro	Leu	Arg	Gly	Thr	Ile	Val
			115				120					125			
Thr	Arg	Pro	Leu	Met	Glu	Ser	Glu	Leu	Val	Ile	Gly	Ala	Val	Ile	Ile
		130				135					140				
Arg	Gly	His	Leu	Arg	Met	Ala	Gly	His	Ser	Leu	Gly	Arg	Cys	Asp	Ile
145				150					155					160	
Lys	Asp	Leu	Pro	Lys	Glu	Ile	Thr	Val	Ala	Thr	Ser	Arg	Thr	Leu	Ser
				165					170					175	
Tyr	Tyr	Lys	Leu	Gly	Ala	Ser	Gln	Arg	Val	Gly	Thr	Asp	Ser	Gly	Phe
			180				185						190		
Ala	Ala	Tyr	Asn	Arg	Tyr	Arg	Ile	Gly	Asn	Tyr	Lys	Leu	Asn	Thr	Asp
			195			200					205				
His	Ala	Gly	Ser	Asn	Asp	Asn	Ile	Ala	Leu	Leu	Val	Gln			
			210			215					220				

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<210> SEQ ID NO 206
<211> LENGTH: 76
<212> TYPE: PRT
<213> ORGANISM: Severe acute respiratory syndrome virus

<400> SEQUENCE: 206

Met Tyr Ser Phe Val Ser Glu Glu Thr Gly Thr Leu Ile Val Asn Ser
1           5           10           15

Val Leu Leu Phe Leu Ala Phe Val Val Phe Leu Leu Val Thr Leu Ala
20           25           30

Ile Leu Thr Ala Leu Arg Leu Cys Ala Tyr Cys Cys Asn Ile Val Asn
35           40           45

Val Ser Leu Val Lys Pro Thr Val Tyr Val Tyr Ser Arg Val Lys Asn
50           55           60

Leu Asn Ser Ser Glu Gly Val Pro Asp Leu Leu Val
65           70           75
    
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What is claimed is:

1. An isolated SARS virus nucleic acid molecule comprising a nucleic acid sequence selected from the group consisting of SEQ ID NOs: 1, 2 and 15.
2. The molecule of claim 1, wherein said molecule is selected from the group consisting of genomic RNA, DNA, cDNA, synthetic DNA and mRNA.
3. The molecule of claim 1, wherein said molecule comprises a s2m motif.
4. The molecule of claim 1, wherein said molecule comprises a leader sequence.
5. The molecule of claim 1, wherein said molecule comprises a transcriptional regulatory sequence.
6. The molecule of claim 1, wherein said molecule encodes a polyprotein.
7. The molecule of claim 1, wherein said molecule encodes a polypeptide.

8. A vector comprising the nucleic acid molecule of claim 1.
9. An isolated host cell comprising the vector of claim 8.
10. The host cell of claim 9, wherein said cell is selected from the group consisting of a mammalian cell, a yeast, a bacterium, and a nematode cell.
11. An isolated nucleic acid molecule comprising a sequence complementary to the entire sequence of SEQ ID NOs: 1, 2 or 15.
12. A kit for detecting the presence of a SARS virus in a sample, wherein said kit comprises the isolated SARS virus nucleic acid molecule of claim 1 or 11.
13. A microarray comprising a plurality of elements, wherein the microarray comprises the nucleic acid of claim 1 or 11.
14. A composition comprising the nucleic acid of claim 1 or 11.

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