

(12) **United States Patent**
Kapil

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(54) **IMMUNOGENIC COMPOSITIONS, VACCINES AND DIAGNOSTICS BASED ON CANINE DISTEMPER VIRUSES CIRCULATING IN NORTH AMERICAN DOGS**

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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 635 days.

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Related U.S. Application Data

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(51) **Int. Cl.**

A61K 39/12	(2006.01)
A61K 39/175	(2006.01)
C12Q 1/70	(2006.01)
C12P 19/38	(2006.01)
C12P 19/34	(2006.01)

(52) **U.S. Cl.**

USPC **424/204.1**; 424/213.1; 435/5; 435/87; 435/91.1; 435/91.3; 435/91.33

(58) **Field of Classification Search**

None
See application file for complete search history.

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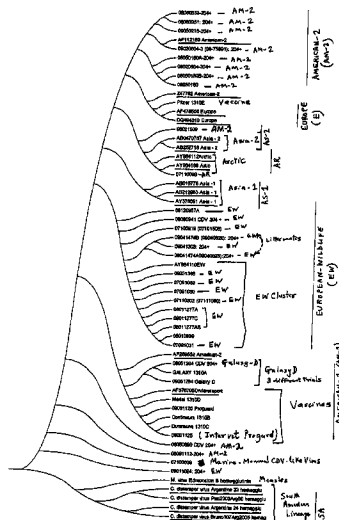
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(57) **ABSTRACT**

Immunogenic compositions and broad-spectrum vaccines containing newly identified isolates of canine distemper virus (CDV) collected from a geographic area are provided. The newly identified isolates exhibit attributes of both European wildlife lineage CDV and one or both of Arctic and American-2 lineage CDV. Therefore, the vaccines are broadly protective against infection with European wildlife lineage CDV and either Arctic lineage CDV or American-2 lineage CDV, or both Arctic and American-2 lineage CDV.

16 Claims, 24 Drawing Sheets



(56)

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07091030

AAGGTGAATTTTACTAACTACTGCGATAACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCCATCCT
CCTGTCAGCACTCTCTGGGGCAGAGGTGACATATTTCCACCATACAGATGCAGTGGAGCTGCTACCTCAGTAG
GCAGAGTTTTCCCCTATCAGTGTCAATTGTCCATGTCTTTGATCTCAAGAAAATCAGAGATAATCAATATGCTA
ACCGCTATCTCAAACGGAGTGTATGGTAAAACCTATTTACTAGTGCCTGATTATATTGAAGAGGAGTTTCGACAC
ACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCA
ACTATATGGTCTCCAGAGAATTCCAAAGCTAAGGTATGTACTATAGCAGTGGCGAGTTGACACTGGCTTCC
TTGTGTAGGTGAGAGCACCGTGTGTTATATCATGACAGCAATGGTTTCGCAAGATAGTATCCTAGCAGTGAC
GCTGGGAATATTTGGGGCAACAACCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAA
AAATACATATAACAAATCACCGTGGGTTCATAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATGGTCTCT
GAGAAAACAGGAAGAGCAAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCATATGTGCAACCAAAC
GTCATGGGAACCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCACTG
ACCTTCAACTTAACATATCGTTTACATACGGTCCGGTTTACTGAATGGAGA (SEQ ID NO:1)

FIG. 1

07091031

GGTGAATTTTACTAACTACTGCGATAACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCCATCCTCC
TGTCAGCACTCTCTGGGGCAGAGGTGACATATTTCCACCATACAGATGCAGTGGAGCTGCTACCTCAGTAGGC
AGAGTTTTCCCCTATCAGTGTCAATTGTCCATGTCTTTGATCTCAAGAAAATCAGAGATAATCAATATGCTAAC
CGCTATCTCAAACGGAGTGTATGGTAAAACCTATTTACTAGTGCCTGATTATATTGAAGAGGAGTTTCGACACAC
AAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAAC
TATATGGTCTCCAGAGAATTCCAAAGCTAAGGTATGTACTATAGGAGTGGCGAGTTGACACTGGCTTCCCTT
GTGTGTAGGTGAGAGCACCGTGTGTTATATCATGACAGCAATGGTTTCGCAAGATAGTATCCTAGCGGTGACGG
TGGGAATATTTGGGGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAA
ATACATATAACAAATCACCGTGGGTTCATAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGA
GAAACAGGAAGAGCAAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCATATGTGCAACCAAACGT
CATGGGAACCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCACTGAC
CTTCAACTTAACATATCGTTTACATACGGTCCGGTTTACTGAATGGAGACGGTATGGATTATTATGAAAGCCC
ACTGTCGGACTCCGGATGGCTTACCATTCTCCAAAAACGGAACAGTCCTTGGATTGATAAACAAAGCAAGTA
GAGGAGACCAGTTATTGTAATCCCCATGTGTTGACATTTGCGCCCAGGGAATCAAGTGGGAATTGTTATTTA
CCTATTCAAACATCCCAG (SEQ ID NO:2)

FIG. 2

07091032

GGTGAATTTTACTAACTACTGCGATAACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCCATC
CTCCTGTCAGCACTCTCTGGGGCAGAGGTGACATATTTCCACCATACAGATGCAGTGGAGCTGCTACCT
CAGTAGGCAGAGTTTTCCCCTATCAGTGTCAATTGTCCATGTCTTTGACCTCAAGAAAATCAGAGATAAT
CAATATGCTAACCGCTATCTCAAACGGAGTGTATGGTAAAACCTATTTACTAGTGCCTGATTATATTGAA
GAGGAGTTTCGACACACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATGACATGC
CATTACTCCAGACAACCTAATATATGGTCTCCAGAGAATTCCAAAGCTAAGGTATGTACTATAGCAGT
GGCGAGTTGACACTGGCTTCCCTTGTGTGTAGGTGAGAGCACCGTGTGTTATATCATGACAGCAATGGT
TCGCAAGATAGTATCCTAGCAGTGACCTGGGAATATTTGGGGCAACATCTATGGATCAAGTTGAAGAGG
TGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATATAACAAATCACCGTGGGTTCATAAAAGATTC
AATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAACAGGAAGAGCAAAAAAATTGTCTGGAGTCG
GCTTGTCAAAGAAAATCCTACCCATATGTGCAACCAAACGTCATGGGAACCTTCGGAGGAGGACAGTTG
CCATCTTATGGGCGG (SEQ ID NO:3)

FIG. 3

07101508

CTAGTAAGATCAGGTGAATTTTACTAATTACTGCTATACAATTGGGATCAGAAAATCTATTGCATCGGCA
GCAAATCCCATCCTCCAGTCAGCACTCTCTGGGGGCAGAGGTGACATATCCCACCATACAGATGCAGTG
GAGCTGCTACCTCAGTAGGCAGAGTTTTCCCCCTATCAGTGTCAATTGTCCATGTCTTTGATCTCAAGAAA
ATCAGAGATAATCAATATGCTAACCCTATCTCAAACGGAGTGTATGGTAAAACCTATTTACTAGTGCCT
GATTATATTGAAGAGGAGTTCGACACACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGC
TGAATGACATGCCATTAATCCAGACAACCAACTATATGGTCTCCAGAGAATTCCAAAGCTAAGGTATG
TACTATAGCAGTGGCGAGTTGACACTGGCTTCTTGTGTGTAGGTGAGAGCACCGTGTGTTATATCAT
GACAGCAATGGTTCGCAAGATAATATCCTAGTAGTGACGCTGGGAATATTTGGGGCAACATCTATGGATC
AAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATATAACAAATCACCGTGGGTT
CATAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAACAGGAAGAGCAAAAAATT
TGCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCAACGTCATGGGAACCTTCCGGAG
GAGGACAGTTGCCATCTTATGGGCGTTGACATTACCTTAGATNCAAGCACTGACCTTCAACTTAACAT
ATCGTTTACATACCGTCCGGTTACTGAATGGAGACGGNATGGATTATTATGAAAGCCCCTGTCCG
(SEQ ID NO:4)

FIG. 4

07100609

AACTTGTATCCGGCTCTTGGGTTGCATGAGTTTTCCGGGGAGTTAAACAACCATGAATCCCTTATGATGC
TATATCAACAGATGGGTGAAACAGCACCGTACATGGTTATTCTGAAAATCTGTCCAGAACAAATTTAG
TGAGGATCCTACCATTGCTCTGGAGTTATGCTATGGGAGTTGGTGTGAACTTGAAAACCTCCATGGGA
GGGTTAAATTTCCGTAGATCCTACTTTGACCCAGCTTATTTCAAGGCTCGGGCAAGAAAATGGTTAGAGAT
CGGCCGGTAAGGTAAGCTCTGCACCTGCCGCCGAGCTTGGCATCACCAAGGAAGAGGCTCAGCTAGTGTG
AGAAAATAGCATCCAAGACAACAGAGGACCA (SEQ ID NO:5)

FIG. 5

07110098

AAGTGAATTTTACTAGTTACTGTGATACAATTGGGATCAGAAAATCCATTGCATGGCAGCAAATCCCGT
CCTTTTGTGCACTCTCCGGAGGCAGAGGTGACATATCCCACCATACAGATGCAGTGGAGCTACTACT
TCAGTTGGCAAATCTTTCCCCCTATCAGTATCATTATCCATGTCTTTGATCTCAAGAACATCAGAGATAA
TCAATATGCTGACCTCTATCTCAGACGGAGTGTATGGTAAAACCTATTTGCTAGTGCCTGATTATATGA
AGGGGAGTTTCGACACGCAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAAGGTGGCTGAATGACATG
CCATTATTCAGACAACCAACTATATGATCCTCCCGGAGAATTCTAAAACCAAGGTATGTAATATAGCAG
TGGGCGAGTTGACACTGGCTTCTTGTGTGTAGATGAGAGCACTGTATTATTATATCATGACAGCAATGG
TTCACAAGATGGTATCTAGTAGTGACGCTGGGAATCTTTGGGGCAACACCTATGGATCAAGTCGAAGAG
GTGATACCTGTGCTCACCCATCAGTCGAAAAAATACATATAACAAATCACCGTGGTTTCATAAAAGATT
CAGTAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAACCCTAGAGGAACAAGAAAATTTGTCTGGAGTC
GGCTTGTGAGAAAAATCCTACCCTATGTGCAATCAAACATCATGGGAACCTTTGGAGGAGGACAGTTG
CCATCTTATGGGCGTTGACGTTACATCTAGATGCAAGCATTGACCGTCAACTTAACATATCATTACAT
ACGGTCC (SEQ ID NO:6)

FIG. 6

07111080

TCAAGAAAATCAGAGATAATCAATATGCTACCCGCTATCTCAAACGGAGTGTATGGTAAAACCTTATTTAC
TAGTGCCTGATTATATGAAGAGGAGTTCGACACACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAA
CGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCCTCCCAGAGAATTCCAAAGCTA
AGGTATGTACTATAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGAGCACCGTGTGT
ATATCATGACAGCAATGGTTCGCAAGATAGTATCCTAGCAGTGACGCTGGGAATATTTGGGGCAACATCT
ATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACTCATCAGTAGAAAAAATACATATAACAAATCACC
GTGGGTTTATAAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAACAGGAAGAGCA
AAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCAAACGTCATGGGAACCC
TTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCACTGACCTTCAAC
TTAACATATCGTTTACATACGGTCCGGTTATACTGAATGGAGACGGTATGGATTATTATGAAAGCCCACT
GTCGGACTCCGGATGGCTTACCATTCTCCCAAAAAACGGAACAGTCCTTGGATTGATAAATAAAGCAAGT
AGAGGAGACCAGTTCATTGTAATCCCCCATGTGTTGACATTTGCGCCAGGGAATCAAGTGGGAATTGTT
ATTTACCTATTCAAACATCCCAGATTATAGA (SEQ ID NO:7)

FIG. 7

08010939

GAATTTTACTAATTACTGCGATACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCCATCCTC
CTGTGACACTCTCTGGGGGAGAGGTGACATATTTCCACCATAACAGATGCAGTGGAGCTGCTACCTCAG
TAGGCAGAGTTTTCCCTATCAGTGTCAATTGTCCATGTCTTTGATCTCAAGAAAATCAGAGATAATCAA
TATGCTAACCGCTATCTCAAACGGAGTGTATGGTAAAACCTTATTTACTAGTGCCTGATTATATTGAAGAG
GAGTTCGACACACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATGACATGCCAT
TACTCCAGACAACCAACTATATGGTCCTCCCAGAGAATTCCAAAGCTAAGGTATGTACTATAGCAGTGGG
CGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGAGCACCGTGTGTATATCATGACAGCAATGGTTCG
CAAGATAATATCCTAGTAGTGACGCTGGGAATATTTGGGGCAACATCTATGGATCAAGTTGAAGAGGTGA
TACCTGTTGCTACCCATCAGTAGAAAAAATACATATAACAAATCACCGTGGGTTCAATAAAAAGATTCAAT
AGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAACAGGAAGAGCAAAAAAATTTGTCTGGAGTCGGCT
TGTCAAAGAAAATCCTACCCTATGTGCAACCAAACGTCATGGGAACCCCTTCGGAGGAGGACAGTTGCCAT
CTTATGGGCGGTTGACATTACCTCTAGATCCAAGCACTGACCTTCACTTAACATATCGTTTACATACGG
TCCGGTTATACTGAATGGAGACGGTATGGATTATTATGAAAGCCCACTGTCCGACTCCG (SEQ ID
NO: 8)

FIG. 8

08011277A

GCCGGGCTGCATCACCCCCTAGTAAGACAGGTGAATTTTACTTATTACTGCGATACAATTGGGATCAGAA
AATCTATTGCATCGGCAGCAAATCCCATCCTCCTGTGAGCACTCTCTGGGGGCAGAGGTGACATATTCCC
ACCATACAGATGCAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTTCCCCCTATCAGTGTCAATGTCCATG
TCTTTGATCTCAAGAAAATCAGAGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGTATGGTAAAA
CTTATTTACTAGTGCCTGATTATATTGAAGAGGAGTTCGACACACAAAAGATTGAGTCTTTGAGATAGG
GTTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAGAAT
TCCAAAGCTAAGGTATGTAATAGCAGTGGGCGAGTTGACACTGGCTTCTTGTGTGTAGGTGAGAGCA
CCGTGTTGTTATATCATGACAGCAATGGTTGCAAGATAATATCCTAGTAGTGACGCTGGGAATATTTGG
GGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATATA
ACAAATCACCGTGGGTTCAAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAAC
AGGAAGAGCAAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCATGTGCAACCAAACGTC
ATGGGAACCCCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCACT
GACCCCTCCAACCTAACATATCGTTTACATACCGTCCGGTTATACTTGAATGGAGACGGTATGGATAATT
ATGAAAGCCCACTGTCCGACTCGGATGGCTTACCATTTCCTTCCAAAACGGAACAGTCCTTGGATTGATA
AACAAACCAGTAGGGGAGACCAGTTCAATTGTATCCCCCATGTGTTGACCATTGCCCCAGGGAATCAAGGG
GAATGTATTTACCTATTCAACCTTCCCAAATAATGGGATAAAGGATGGCCCTCCTGAATCCAAATTACGG
TGTTGCCCTAAC (SEQ ID NO:9)

FIG. 9

08011277B

TTGGTTAAGGCCATCCTTTTTCCCTAATCTGGGCTGTTTGAATAGGTAAATAACAATTCCCCACTTGATT
CCCTGGGCGCAAATGTCAACACATGGGGGATTACAATGAACTGGTCTCCCCCTACTTGCTTTGTTTATCAA
TCCAAGGACTGTTCCGTTTTTGGGAGGAATGGTAAGCCATCCGGAGTCCGACAGTGGGCTTTCATAATAA
TCCATACCGTCTCCATTCAGTATAACCGGACCGTATGTAAACGATATGTTAAGTTGAAGGTCAGTGCTTG
GATCTAGAGGTAATGTCAACCGCCATAAGATGGCAACTGTCTCCTCCGAAGGGTTCCCATGACGTTTG
GTTGCACATAGGGTAGGATTTTCTTTGACAAGCCGACTCCAGACAATTTTTTTGCTCTTCTGTTTCTCA
GAGACCAATGCAGGCACCATCCAGGTTGCTATTGAATCTTTTATGAACCCACGGTGATTTGTTATATGTA
TTTTTTCTACTGATGGGTGAGCAACAGGTATCACCTCTTCAACTTGATCCATAGATGTTGCCCAAATAT
TCCCAGCGTCACTACTAGGATATTATCTTTCGAACCATTGCTGTGATGATATAACAACACGGTGCTCTCA
CCTACACACAAGGAAGCCAGTGTCAACTCGCCCACTGCTATAGTACATACCTTAGCTTTGGAATTCTCTG
GGAGGACCATATAGTTGGTTGTCTGGAGTAATGGCATGTCAATTCAGCCACCGTTTTGATGAACCTATCTC
AAAGACTCGAATCTTTTTGTGTGTGCAACTCCTCTTCAATATAAATCAGGCACCTAGTAAATAAAGTTTA
CCATACACCTCCGTTTGGAGATAGCCGTTAGCATATTGATTATCTCTGATCCTCTTGGAGATCAAAGACAT
GGACAATGACACTGATAGGCGGGAAAACCTCTGCCTACTGAGGTAGCAGCTCTACTGCTTTTGTGGGTGG
GAAATATTTAACCCCTTTGCCCCGAAAGTGCTTACAGGAGGATGGGATTTGCTGCCGATCCAATAAATTT
TCTGATCCCAATTGTATCGAAGAACTAATAAATTACCTGGACCTTACTTGGGGGGGTGATGAACCAGCGC
(SEQ ID NO:10)

FIG. 10

08011277C

AGATCAAGGTGAATTTTACTAATTACTGCGATACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAA
TCCCATCCTCCTGTCAGCACTCTCTGGGGCAGAGGTGACATATTCCCACCATACAGATGCAGTGGAGCT
GCTACCTCAGTAGGCAGAGTTTTCCCCTATCAGTGTCAATTGTCCATGTCTTTGATCTCAAGAAAATCAG
AGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGTATGGTAAAATCTATTTACTAGTGCCTGATTA
TATTGAAGAGGAGTTCGACACACAAAAGATTGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAAT
GACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAGAATTCCAAAGCTAAGGTATGTACTA
TAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGAGCACCGTGTGTTATATCATGACAG
CAATGGTTCGCAAGATAATATCCTAGTAGTGACGCTGGGAATATTTGGGGCAACATCTATGGATCAAGTT
GAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATATAACAAATCACCGTGGGTTCATAA
AAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAAAGGAAAGAGCAAAAAAATGTCT
GGAGTCGGCTTGTCAAAGAAAATCCTACCTATGTGCAACCAAACGTGATGGGAACCTTTTCGGAGGAGG
ACAGTTGCCATCTTATGGGCGGTTGAC (SEQ ID NO: 11)

FIG. 11

08011277D

CCCCAATTGGCATTGAACCATGTATCCGGCTCTTGGGTTGCATGAGTTTTCCGGGGAGTTAAACAACCAT
GAATCCCTTATGATGCTATATCAACAGATGGGTGAAACAGCACCGTACATGGTTATTCTGGAAAATCTG
TCCAGAACAAATTTAGTGCAGGCTCCTACCCATTGCTCTGGAGTTATGCTATGGGAGTTGGTGTGAACT
TGAAAATCCATGGGAGGGTTAAATTTCCGGTAGATCCTACTTTGACCCAGCTTATTTCAGGCTCGGGCAA
GAAATGGTTAGAAGATCTGCCGGTAAGGTAAGCTCTGCACCTTGCCGCCGAGCTCGGCATCACCAGGAAG
AGGCTCAGCTAGTGTGAGAAATAGCATCCAAGACAACAGAGGACCTCCATTTGGCATTGAAACTATGTA
TCCGGCTCTTGGGTTGCATGAGTTTTCCGGGGAGTTAAACAACCTTGAATCTTAATGACCTTTTTCCGCA
GGGAACAAACCCACAATCGCTGAATTTCTGTGAAATATGGCTCACACATTGTGGCAGCTCGACACCGACT
TTAACCTTACCTATGGAATTTGGCGTTGAAACTGTAAATCCCTCTTCGGGTTACCACCTCTTTTGATCAC
TTTAACCGTTATTTACGCCGGCAGCCACGTTAGAACAATATCCGCCTTCGCAAGTTTTCCTGCCTCCTCCT
TCCACCCAATTAGAGGGCCCCCTCCTTTGTTATGAACCCCTTA (SEQ ID NO: 12)

FIG. 12

08011671

CCTGGGCGCCTTACCCCCCTAGTAAGCTCAGGTGAATTTTACTAACTACTGCGATAACCTTGGGATCA
GAAAATCTATTGCATCGGCAGCAAAATCCCATCCTCCTGTGAGCACTCTCTGGGGCAGAGGTGACATATT
CCCACCATAACCGATGAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTTCCCCTGTGAGTGTCAATTGTCC
ATGTCTTTGATCTCAAGAAAATCAGAGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGTATGGTA
AACTTATTTACTAGTGCCTGATTATATTGAAGAGGAGTTCGACACACAAAAGATTGAGTCTTTGAGAT
AGGGTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAG
AATTCCAAAGCTAAGGTATGTACTATAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGA
GCACCGTGTGTTATATCATGACAGCAATGGTTCGCAAGATAGTATCCTAGCAGTGACGCTGGGAATATT
TGGGGCAACATCTATGGATCAAGTTGAAGAGGCGATACCTGTTGCTCACCCATCAGTAGAAAAAATACAT
ATAACAAATCACCGTGGGTTATAAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGA
AACAGGAAGAGCAAAAACAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCAAAC
GTCATGGGAACCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTTAGATCCAAGCA
CTGACCTTCAACTCAACATATCGCTTACATACCGTCCGGCTATACTGAATGGGAGACGGTATGGATTTTA
TGACAAGCCCCCTGTGGGACTCCCGGATGGCTTACCACCCCTCCCAAAACCGGAACAGCTCCTTCGAT
TGATAAACCAAACCGTACGAGGAGACTCAGTTTCAATTGTTATTTCCCCTACGTGTTGACATTTCCGCCCC
AGGCCATCCATGTGCGATTGCTCTTTACCCAATAACCCACCCACATCATGGATACAGCTCTCCTTACTG
ACTCCACACTACCGCTGTTGCCTACCCTCCCGCTCTCCCTTCCCCTA (SEQ ID NO: 13)

FIG. 13

08021509

GTGAATTTTACTAATTACTGCGATACTATGGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCCATCC
TTTTATCAGCACTCTCCGGAGGTAGAGGTGACATATCCCACCATACAGATGCAATGGAGCTACTATTTCC
AGTAGGCAAGATTTTCCCCNATCAGTATCATTATCTATGTCTTTGATCTCAAGAACATCAGAGATAATC
AATATGCTAACCGCTATCTCAGACGGAGTGATGG (SEQ ID NO:14)

FIG. 14

08030074

CAAGGTGAATTTTACTAATTACTGCGATAACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCC
ATCCTCCTGTGAGCACTCTCTGGGGCAGAGGTGACATATCCCACCATACAGATGAGTGGAGCTGCTA
CCTCAGTAGGCAGAGTTTCCCCCTATCAGTGACATTGTCCATGTCTTTGATCTCAAGAAAATCAGAGAT
AATCAATATGCTAACCGCTATCTCAAACGGAGTGATGGTAAGACTTATTTACTAGTGCCTGATTATATT
GAAGAGGAGTTCGACACACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATGACA
TGCCATTAATCCAGACAACCAACTATATGGTCCTCCAGAGAATTCCAAAGCTAAGGTATGTACTATAGC
AGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGAGCACCGTGTGTATATCATGACAGCAAT
GGTTCGCAAGATAATATCCTAGTAGTGACGCTGGGAATATTGGGGCAACATCTATGGATCAAGTTGAAG
AGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATATAACAAATCACCGTGGGTTCCATAAAGA
TTCAATAGCAACCTGGATGGTGCCTGCATGGTCTCTGAGAAACAGGAAGAGCAAAAAAATTGTCTGGAG
TCGGCTGTCAAAGAAAATCCTACCCTATGTGCAACCAACGTATGGGAACCCCTTCGGAGGANGGACAG
TTGCCATCTTATGGGCGGTTGACATTACCTTAGATCCAAGCACTGACCTTCAACTTAACATATC
(SEQ ID NO:15)

FIG. 15

08030776

TCCTGTTTGCCTTTCCCCCCTAGTAAGATCAGGTGAATTTTACTAACAACCTGCGATAACAATTGGGATC
AGAAAATCTATTGCATCGGCAGCAAATCCCATCCTCCTGTGAGCACTCTCTGGGGCAGAGGTGACATAT
TCCCACCATACAGATGAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTCCCCCTATCAGTGTCAATTGTC
CATGTCTTTGATCTCAAGAAAATCAGAGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGATGGT
AAAATTTACTAGTGCCTGATTATATTGAAGAGGAGTTCGACACACAAAAGATTCGAGTCTTTGAGA
TAGGGTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCCCTCCCAGA
GAATTCCAAAGCTAAGGTATGTACTATAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAG
AGCACCGTGTGTATATCATGACAGCAATGGTTCGCAAGATAGTATCCTAGCAGTGACGCTGGGAATAT
TTGGGGCAACAACCTATGGATCAAGTTGAAGAGGTGATACCTGTGCTCACCCATCAGTAGAAAAAATACA
TATAACAAATCACCGTGGGTTCCATAAAGATTCAAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAG
AAACAGGAAGAGCAAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCAAA
CGTTATGGGAACCCCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAG
CACTGACCTTCAACTTAACATATCGTTTACATACGGTCCGGTTATCCTGAATGGAGACGGTATGGATAT
TATGAAAGCCCACTGTCCGACTCCCAGTGGCTTACCATTCTCCAAAACGGAACAGTCTTGGATTGATA
AACAAACAAGTAGAGGAGACCAGTTCATTGAATCCCATGTGTGACTTTTCGCCAGGGAATCAAGTGG
AATTGTATTTACTATCAACTTCCAGATTATGGATAAGATGTCTTCTGATTCCAATACGGTGTGCCTTA
(SEQ ID NO:16)

FIG. 16

08030777

TCGTGGTGCTTAACCCCCCTAGTAAGATCAGGTGAATTTTACTAATTACTGCGATACTATTGGGATCAG
AAAATCTATTGCATCGGCAGCAAATCCCATCCTTTTATCAGCACTCTCCGGAGGTAGAGGTGACATATTC
CCACCATA CAGATGCAATGGAGCTACTATTT CAGTAGGCAAGATTTTCCCCCTATCAGTATCATTATCTA
TGTCTTTGATCTCAAGAACATCAGAGATAATCAATATGCTAACCGCTATCTCAGACGGAGTGATGGTAA
AACTTATTTACTAATGCCTGATTATATTGAAGGGGAGTTCGACACGCAAAAGATTCGAGTCTTTGAGATA
GGGTT CATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAGA
ATTCCAAAGCCAAGGTATGTACTATAGCAGTGGGCGAGTTGACACTGGCTTCTTTGTGTGTAGATGAGAG
CACCGTATTGTTATATCATGACAGCAATGGTTCACAAGATGGTGTCTAGTAGTGACGCTGGGAATATTC
GGGGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTGCGTGACCCATTAGCAGAAAAAATACATA
TAACAAATCACCGTGGGATCATAAAAGACTCAATAGCAACCTGGATGGTGCCTGCATTAGTTTCTGAGAA
ACAAGAGGAACAAACAAATTTGTCTGGAGTCAGCTTGTCAAAGAAAATCCTACCCTATGTGCAATCAAACG
TCATGGGAACCTTTGGAGGAGGACAGTTGCCATCTTATGGGCGGCTGACATTACCTCTACATCCAAGCA
TTGACCTCCACTAACATATCATTTACATACGGTCCGACTATACTGAATGGAGACGGATGGCTATTATGA
GAGCCCCCTGCGGACTCCGGATGGCTTACCTTTCCCTCCAGCACGGCACAGCTGGATTGATAAACAAG
AGTAGAGGACGACCAGTTATTGTCAATCCCCCTGTGTTGACATTTCCGCCCCGGCATCCACCCGAAATTGC
TATTACCCTATCCACATTCCCCTTCGCGCTCAAGATCCCCCTCCTGCTCCCCACCACGGCGCGCTCCCT
ATCTCC (SEQ ID NO: 17)

FIG. 17

08031346

CCTAGTAGATCAAGGTGAATTTTACTAATTACTGCGATACAATTGGGATCAGAAAATCTATTGCATCGGC
AGCAAATCCAATCCTCCTGT CAGCACTCTCTGGGGCAGAGGTGACATATTCACCACATACAGATGCAGT
GGAGCTGCTACCTCAGTAGGCAGAGTTTCCCCCTATCAGTGTCTATTGTCCATGTCTTTGATCTCAAGAA
AATCAGAGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGATGGTAAAATCTATTTACTAGTGCC
TGATTATATTGAAGAGGAGTTCGACACACAAAAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGG
CTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAGAATTCCAAAGCTAAGGTAT
GTACTATAGCAGTGGGCGAGTTGACACTGGCTTCTTGTGTGTAGGTGAGAGCACCGTGTGTTATATCA
TGACAGCAATGGTTCGCAAGATAGTATCCTAGCAGTGACGCTGGGAATATTTGGGGCAACATCTATGGAT
CAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATATAACAAATCAC
(SEQ ID NO:18)

FIG. 18

08040383

GTTTGATAGGTAAATAACAGTTTCCACTTGATTCCCTGGGTGCAAATGACAACACATGAGGGACCACAGT
GAACTGGTCTCCTCTACTTGCTTTGTTTATCAATCCAAGAAATGTTCCATTCTTAGGAGGAATGGTAAGC
CATCCGGATTCCAAAAGTGGGCTTTCATAATAATCCATACCATCTCCATT CAGTATAACCGGACCGTATG
TAAATGATATGTTAAGTTTACGGTCAATGCTTGCACTAGATGTAACGTCAACCGCCATAAGATGGCAA
CTGTCCTCCTCCAAAGGGTCCCATGATGTTGATTGCACATGGGGTAGGATTTTCTCTGACAAGCCGAC
TCCAGACAATTTCTGTTTCTCTAGGTTCTCAGAGACCAATGCAGGCACCATCCAGGTTGCTACTGAAT
CTTTTATGAAACCACGGTGAATTTGTTATATGATTTTTTTCGACTGATGGGTGAGCGACAGGTATCACCTC
TTGCTGTGATGATATAATAATACAGTGTCTCTCATACACAAGGAAGCCAGTGTCAACTCGCCACTG
CTATAGTACATACCTTGGTTTTAGAAATCTCCGGGAGGATCATATAGTTGGTTGTCTGGAATAATGGCAT
GTCATT CAGCCACCTTTTGATGAACCC TATCTCAAAGACTCGAATCTTTTTCGCTGTGCAACTCCCCTTCA
ATATAATCAGGCACTAGCAAATAAGTTTTACCATACTCCGTCTGAGATAGAGGTGAGCATATTGATTA
TCTCTGATGTTCTTGAGATCAAAGAC (SEQ ID NO:19)

FIG. 19

08050180A

AATGCTTCCTTTACCCACCCTAGTAAGATCAAGTAAATTTTACGGTAAATAAATAGCGATACAATTGGGA
TCAGAAAATCTATTGCATCGGCAGCAAATCCTATCCTTTTATCAGCACTCTCCGGAGGTAGAGGTGACAT
ATCCACCATACAGGTGCAGTGGAGCTACTACTTCAGTAGGCAGAGTCTTCCCCCTATCAGTATCATTG
TCCATGTCTTTGGTCTCAAGAACATCTGAAATAATCAATATGCTAACCGCTATCTCAGACGGTGTGTATG
GTAAAACCTTATTTGCTAGTTCTTGATTATCTTGAAGGGGAGTTTCGACACGCAAAGATTTCGAGTCTTTGA
GATAGGGTTTCATCAAACGGTGGCTGAACAACATGCCATTACTCCAGACAACCAACTATATGGTCTCCCG
GAGGATTCCAAAGCCAAGGTATGTACTATAGCGGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGATG
AGAGCACCGTATTGTTATATCATGACAGCAGTGGTTCACAAGATGGTATTCTAGTGGTGACGCTGGGAAT
ATTTGGGGCAACACCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATA
CATATAGCAAACCACCGTGGGTTTCATCAAAGATTCAATAGCAAACCTGGATGGTGCCTGCATTGGTCTCTG
AGAAACAAGAGGAACAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCA
AACGTCATGGGAACCTTTGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCAA
AGCATTGACCTCCAGCTTAACATCTCATTACATATGGTCCGGTTATACTGAATGGAGACGGTATGGATT
ATTATGAAAGTCCGCTTTTGAACCTCCGGATGGCTTACCATTCTCCCAAGAACGGAACAGTCTTTGGATT
GATAAACAAGCAAGTAGAGGAGACCAGTTCAGTGTATCCCCATGTGTGACATTTGCGCCAGGGAATCA
AGTGGAAATTGTATTTACCTATTCAAACATCCCAGATATGGATAAAGATGTCCTTACTGAATCCAAATTAG
TGGTGTTCCTAAC (SEQ ID NO:20)

FIG. 20

08060351

ACCGGGGTGCTTACCCCCCTAGTAAGATCAAGTGAATTTTACGAAAACTGCGATCCAATTGGGATCAG
GAAATCTATTGCAACGGCAGCAAATCCTATCCTTTTATCAGCACCTCCGGAGGTAGAGGTGACATATTC
CCATCATACAGATGCAGTGGAGCTACTACTTCAGTAGGCAGAGTCTTCCCCCTATCAGTATCATTGTCCA
TGTCTTTGATCTCAAGAACATCTGAAATAATCAATATGCTAACCGCTATCTCAGACGGAGTGTATGGTAA
AACTTATCTGCTAGTTCTTGATTATCTTGAAGGGGAGTTTCGACACGCAAAGATTTCGAGTCTTTGAGATA
GGGTTTCATCAAACGGTGGCTGAACAACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAGG
ATTCCAAAGCCAAGGTATGTACTATAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGATGAGAG
CACCATATTGTTATATCATGACAGCAATGGTTCACAAGATGGTATTCTAGTGGTGACGCTGGGAATATTT
GGGGCAACACCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATA
TAGCAAACCATCGTGGGTTTATCAAAGATTCAATAGCAAACCTGGATGGTGCCTGCATTGGTCTCTGAGAA
ACAAGAGGAACAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCAAACG
TCATGGGAACCTTTGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCA
TTGACCTTCAGCTTACATCTCATTACATACGGCCCGTTATACTGAATGGAGACGGTATGGATACTATGA
AAGCCACTTTTACTCCGGATGGCTTACCATTCTCCAAGAACGGAACAGTCTTTGGATTGATAAACA
AAGCAAGTAGAGGAGACCAGTTCAGTGTATCCCCATGTGTTGACATTTGCGCCAGGAATCAGTGGAAATT
GTTATTTACCTATTCAAACCTTCCAATTATGGATAAAGATGTCCTTACTGGATCCAAATTATGGTGTTCCT
AACC (SEQ ID NO:21)

FIG. 21

08060352

CATTGGTGCATTAACCCACCTAGTAAGACAAGTGAATTTTACTAATATACTGCGATACAATTGGGATCAG
GAAATCTATTGCATCGGCAGCAAATCCTATCCTTTTATCAGCACCTCCGGAGGTAGAGGTGACATATTC
CCATCATAAGATGCAGTGGAGCTACTACTTCAGTAGGCAGAGTCTCCCCCTATCAGTATCATTGTCCA
TGTCTTTGATCTCAAGAACATCTGAAATAATCAATATGCTAACCGCTATCTCAGACGGAGTGTATGGTAA
AACTTATCTGCTAGTTCCTGATTATCTTGAAGGGGAGTTCGACACGCAAAAGATTGAGTCTTTGAGATA
GGGTTTCATCAAACGGTGGCTGAACAACATGCCATTACTCCAGACAACCAACTATATGGTCCTCCCAGAGG
ATTCCAAAGCCAAGGTATGTAATACTAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGATGAGAG
CACCATATTGTTATATCATGACAGCAATGGTTCACAAGATGGTATTCTAGTGGTGACGCTGGGAATATTT
GGGGCAACACCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACATA
TAGCAAACCATCGTGGGTTTATCAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAA
ACAAGAGGAACAAAAAATTTGCTGGAGTCGGCTTGTCAAAGAAAATCCTACCCTATGTGCAACCAAACG
TCATGGGAACCTTTGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCA
TTGACCTTCAGCTTAACATCTCATTTACATACGGTCCGGTTATACTGAATGGAGACGGTATGGATTACTA
TGAAAGCCCACTTTTAGACTCCGGATGGCTTACCATTCTCCCAAGAACGGAACAGTCTTGGATTGATA
AACAAAGCAAGTAGAGGAGACCAGTTCAGTGAATCCCCATGTGTTGACATTTGCGCCCAGGGAATCAA
GTGAAATTTGTTATTTACCTATTCCAAACATCCCAGATTATGGATAAAGGATGTCCTTACTGAAATTTCTA
AATTAGTGGGGGTTTGGCCCTAAGAC (SEQ ID NO:22)

FIG. 22

08080696

GCCTCCCAGGGGCACCTTCCCCCCCCAGTAGCTCAGGTGAATCTCACTTAAAACCTGCGCCCCCTTGGGA
TCTTACAATCTATTGCATCGGCAGCAAATCCCCTCCTTTTATCAGCACTCTCCCAGGTAGAGGTGACAT
ATTCCCACCATACCGATGCAATGGAGCTACTATTTCACTAGGCAAGATTTCCCCCTATCAGTATCATT
TCTATGTCTTTGATCTCACGAACATCAGAGATAATCAATATGCTAACCGCTATCTCATACGGAGTGTATG
GTAAAACCTTATTTACTAATGCCCGACTATATTGAAGGGGAG (SEQ ID NO:23)

FIG. 23

08080941

TTGATTTGACTCCCCGATTTTCCACTGTGCATTAACCACCTAGTAAGATCAAGGTGAATTTTACTGACT
CTGGAACAAATGGGATCAAGAAATTTATTGCATGGCAGCAAATCCCATCTCCTGTCAGCACTCTATGGGG
GCAGAGGTGACATATTTCCACCATACAAGATGCAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTCCCCC
TATCAGTGTCAATTGGCCATGTCTTTGACCTCAAGAAAATCAGAGGATAATCAATATGCTAACCGCTATCT
CAAAACGGAGTGTATGGTAAAACCTATTTACTAGTGCCTGATTATATTGAAGAGGAGTTCGACACACAAA
AAGATTCGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATAACATGCCATTACTCCAGACAACCTA
ACTATATGGTCTCCAGAGAATTCCAAAGCTAAGGTATGTAATACTAGCAGTGGGCGAGTTGACACTGGC
TTCCTTGTGTGTAGGTGAGAGCACCGTGTGTTATATCATGACAGCAATGGTTCGCAAGATAGTATCCTA
GCAGTGACGCTGGGAATATTTGGGGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACC
CATCAGTAGAAAAATACATATAACAAATCACCGTGGGTTCAAAAAGATTCAATAGCAACCTGGATGGT
GCCTGCATTGGTCTCTGAGAAAACAGGAAGAGCAAAAAAATTTGCTGGAGTCGGCTTGTCAAAGAAAATCC
TACCCTATGTGCAACCAAACGTCATGGGAACCTTCCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGA
CATTACCTCTAGATCCAAGCACTGACCTTCAACTTAACATATCGTTTACGTACGGTCCGGTTATACTGAA
TGGAGACGGTATGGATTATTATGAAAGCCCACTGTCGGACTCCGGATGGCTTACCATTCTCCAAAAAC
GGAACAGTCTTGGATTGATAAACAAAGCAAGTAGAGGAGATCAGTTCATTGTAATCCCCATGTGTTGA
CATTTGCGCCAGAGAATCAAGTGGGAATGTTATTTACCTATTCAAACATCCCATATTAGGAAAAAGGG
AGGCCTACCCGGGA (SEQ ID NO:24)

FIG. 24

08081112

TATGGTTTCATTACCCCCGGCGTAAGTGAATTTGAATCGTAGTAATTGCTGTGATAAAAATTGGGATTGGA
AATGTATTGCATTGTTATGAAATTTCTACCTTTTCAGCACTTGCCTCCGTTGGTTGAGGGGACTTATTCCC
ATCATACATATGCAGTGGAGCTACTACCTCATCCGGCAGAGTTATATTTGATCATCATTATTGCACATGT
TTGTGACCTAAAAACATCTGGCATATGCAATCTGCTAACC CGGATCTCATGTGGAGTGTATGGCAAAC
TTATCTGCTACTTCCCTGATTTCTTGAAGGGGAGTCCGACACTCTGCCGATGTCCGACAAGCTGATCGGG
TTCATCAAACCTCTGGCTGAACAACATGTTGCGCTCTGACAACCTCCGATTGGCCTGCCAGAGGATTT
TACAGCCAAGGTATGTACCATATCCCAGGGAACTTACACTGCCTTCCTTGTGTGTTAGCCAGAGCCCC
ATATTGTCCCATAATGATATGAATGTCTACAAGAGGTCATTTCCATGTGACCCCGGTTCAATTTGTGG
CAATGGCGGTGGTTCAATTGGAACAGGGTATATCTGACCCTATCTTCACTAGAGAAATTACATATGACA
AACCATCATGGCTTGATCAAAGAATAACTTCCCTTTCTGGCTGACGCTTGACTTGCCCTTATATATACCAT
ATTTTCTTAATAAATCGCGGTCAATTGCCTGTGGAGCCAATTTTACCCTCTTCCAACCTTATGTTACG
GGCTTTCCTTGCCGGAGGACCGTTGC (SEQ ID NO:25)

FIG. 25

08120827

GCGATTTTGCCCTGTGCATTAACCCACCTAGTAAGATCAAGGTAAATTTTACTAAATTTCTGCGAAACATG
TGGATCAGAAAATCTATGGCATCGGCAGCAATCCCATCCTCCTGCAGCCCTCTTGGGGCAGAGGTGACAT
ATTCCCACCATACAGATGCAGTGAAGCTGCTACCTCAGTAGGCCAGAGTTTTCCCTATCAGGGTCATTG
TGCATGTCTTTGACCTCAAGAAAGTCAGAGATAATCAAATATGCTAACCCGCTATCTCAAACGAGTGTGTA
TGGGAAAAACTTATTTACTAGTGCCTGGATTATATTGAAGAGGAGTTCGACACACAAAAGATTCGAGTCT
TTGAGATAGGGTTTCAATCAAACGGTGGCTGAATAACATGCCATTACTCCAGACAACCTAATATATGGTCTT
CCCAGAGAATTCAAAGCTAAGGTATGTACTATAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTA
GGTGAGAGCACCGTGTGTTATATCATGACAGCAATGGTTCGCAAGATAGTATCCTAGCAGTGACGCTGG
GAATATTTGGGGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAA
AATACATATAACAAATCACCGTGGGTTCTATAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTC
TCTGAGAAACAGGAAGAGCAAAAAAATGTCTGGAGTCCGCTTGTCAAAGAAAATCCTACCCATATGTGCA
ACCAAACGTCATGGGAACCCCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGA
TCCAAGTACTGAATGGAGACGGTATCGTTTACATACGGTCCGGTTATACTGAATGGAGACGGTATG
GATTATTATGAAAGCCCACTGTCCGACTCCGATGGCTTACCATTCCCTCCCAAAAACGGAACAGTCCCTTG
GATTGATAAACAAAGCAAGTAGAGGAGATCAGTTCATTGTAATCCCCATGTGTTAACATTTGCGCCCAG
AGAATCAAGTGGGGATGTTATTTTCTATTCAAACATGCCCATATTATGATAAAGGATGGCCTTAACC
CG (SEQ ID NO:26)

FIG. 26

08120857

AGTTCGACGCACAAAAGATTTCGAGTGTGAGATAGGGTTGATCGGACGAGGAGGTGAAGGACATGCCATT
ACTCCAGACAGCTAACTATATGGTCCGCCAGAGAATTCCAAAGCTAAGGTATGTAATAGCAGTGGGC
GAGGTGGCACTGGCTTCCTTGTGTGTAGGGGAGAGCGCCGTGTTGTTATATCATGGCAGCAATGGTTCCG
AAGATAGTATCGTAGCAGTGACGCTGGGAATATTTGGGGCAACATCTATGGATCAAGTTGAAGAGGTGAT
ACCTGTTGCTCACCCATCAGTAGAGAAAATACATATAGCAAATCACCGTGGGTTTCATAAAGATTCAATA
GCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAACAGGAAGAGCAAAAAAATGTTCTGGAGTCCGCTT
GTCAAAGAAAATCCTACCGTATGTGCAGCCAAACGGCATGGGAACCCCTTCGGAGGAGGACAGTTGCCATC
TTATGGGCGGTTGACATTACCTCTAGATCCAAGCGCTGCCTTCAACTAACATATCGTTTACATACGGTC
CGTTATACTGAATGGAGACGGTATGGATTATATGAAAGCCCACTGTCCGGCTCCGGATGGCTTGCCAT
TCCTCCCAAAAACGGAACAGTCCCTTGGATTGATAAACAAAGCAAGTAGAGGAGATCAGTTCATTGTAATC
CCCATGTGTGGACATTTGCGCCAGAGAATCAAGTGGGGATGTTTTTTAACTATGCAAACGGCGCA
TATGAGGGGGGAGGGGGGGCGGGAGGCT (SEQ ID NO:27)

FIG. 27

09011024

CAGTGAGAGCAAAAATGTAGGAAAGGGCAGGAATTCATGCTCAAGGAGCGGATGTGGGGAGAGGTTCG
AGTCCCGCCAGCAGTGCAGGAAGGGGTACTCAGTAGCGGGTTTCCCCCTAGGAGGGGGATTGTCCAGTC
TTTGATATCAGAAAAGAAGGATATCAATATGCTAACCGCTATCGCCAAAGGAGGGTATGGTAAGAGCTTA
TTGGGAGTGCCTGATTAGAGGGAGGGAAGTTCTACAGGAGAGAGATTGGAGTGGTGAGATGGGGGTTCGT
CAAGCGTGGATGAATGACATACCATTACTCCAGACAACCAAGTATAGGGGCCCTCCAGAGAATGCCAAA
GCTAAGGTATGTAATAGCAGTGGGCGAGTTACGCTGGCTTCCCTTGTGTGTAGGTGAGAGCGCCGTGTT
GTTATATCATGACAGCAATGGTTCGCAAGATAGTATCCTAGCTGTGACGCTGGGAATATTTGGGGCAGCA
TCTATGGATCAAGTTGAAGAGGTGATGCCTGTTGCTCACCCATCAGTAGAAAAAATACATATAACAAATC
GCCGTGGGTTTCATAAAAAGATTCAATAGCAGCATGGATGGTGCCTGCATTGGTCTCTGAGAAGCAGGAAGA
GCAAAAAAATTTGT CAGGAGT CCGGTTGTCAAAGAAAATCCTACCCGATGTGCAACCAAACGTCATGGGAA
CCCTTCGGAGGAGGACAGGTGCCATCTTATGGGCGGTTGGCATTACCTCTAGAGCCAAGCACTGGCCCTC
AACTTGACATATCGTTTACATACGGGCGGTTATACTGAATGGAGACGGTATGGATTATTATGAAAAGCCC
ACTGTCCGACGCCGGATGGCTTACCATTCCCTCCCAAAAACGGAACAGTCCGTGGATTGATAAAACAAAGCA
AGTAGAGGAGGCCAGTTCATTGTAATCCCCATGTGTTGACATTTGCGCCAGGGAATCAAGTGGGAATT
GCTATTTTCCCTATTCAGAACACCCAGATTAGGATAGAAGGAGGGGCCCTGGGCCG (SEQ ID NO:28)

FIG. 28

09020504-3

CTTGTGGGCTTAAACCCTAGTAATACAAAGTGAATTTTACTAATTACTGCGATACAATTGGGATCAA
AAATCTATTGCATCGGCAGCAAATCCTATCCTTTTATCAGCACTCTCCGGAGGCAGAGGTGACATATTCC
CACCATACAGATGCAGTGGAGCTACTACTTCAGTAGGCAGAGTCTTCCCCTTATCAGTATCATTGTCCAT
GTCTTTGATCTCAAGAACATCTGAAATAATCAATATGCTAACCGCTATCTCAGACGGAGTGTATGGTAAA
ACTTATTTGCTAGTTCCTGATTATCTTGAAGGGGAGTTTCGACACGCCGAAGATTTCGAGTCTTTGAGATAG
GGTTCATCAAACGGTGGCTGAACAACATGCCATTAATCCAGACAACCAACTATATGGTCCCTCCGGAGGA
TTCCAAAGCTAAGGTATGTAATAGCAGTGGGCGAGTTGACACTGGCTTCTTATGTGTAGATGAGAGC
ACCGTATTTGTTATATCATGACAGCAATGGTTCACAAGATGGTATTCTAGTGGTGACGCTGGGAATATTTG
GGGCAACACCTATGGATCGAGTTGAAGAGGTGATACCTGTTGCTCACCCGTCAGTAGAAAAAATACATAT
GGCAAACCACCGTGGGTTTCATCAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGAAA
CAAGAGGAACAAAAAATTTGTCTGGAGTCCGGCTTGTCAAAGAAAATCCCTACCCATGTGCAACCAAACG
TCATGGGAAACCCCTTTGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGC
ATTGACCTTACCTTAAACATCTCATTTACATACGGCCCAGTTATACTGAATGGGACGGTATGGATTATT
ATGAAAGCCACTTTTGGACTCCGGATGGCTTACCATTCCCTCCCAAGAACGGAACAGTCCCTGGATTGAT
AAACAGAGCAGTAGAGGAGAACAGTTCACTGTAATCCCCATGTGTTGACTTGCGCAAGGGGATCAAGTGG
AAATGTATTTACCTATTCAAACATCTTAAATTATGGATAAAGATGCCCTCACCGAGCCCAAATTAGTGG
TGTTGCCTCAT (SEQ ID NO:29)

FIG. 29

09041289

CTCCCTTTCGGCTTGAACATGTATCCGGCTCTTGGGTTGCATGAGTTTTCCGGGGAGTTAACAACCATTG
AATCCCTTATGATGCTATATCAACAGATGGGTGAAACAGCACCGTACATGGTTATTCTGGAAAATTCTGT
CCAGAACAAATTTAGTGCAGGATCCTACCCATTGCTCTGGAGTTATGCTATGGGAGTTGGTGTGAACTT
GAAAACCTATGGGAGGGTTAAATTTCCGGTAGATCCTACTTTGACCCAGCTTATTTTCAGGCTCGGGCAAG
AAATGGTTAGAAGATCGGCCGGTAAGGTAAGCTCTGCACTTGCCGCCGAGCTTGGCATCACCAAGGAAGA
GGCTCAGCTAGTGTGAGAAATAGCATCCAAGACAACAGAGGACCCGCATTTGGCATTGAAACTATGT TC
CGGCTCTTGGGTTGCATGAGTTTTCCGGGGAGTTAACAACCATTGAATCCCTTGTGATGCTTTACCACCA
AATGGGTGAAGACCCCCCATGGTTATTCTTGGAAAATTTGTCCGACAAAATTAGTGCAGGATCTACCAT
TGCTCTGGAGTTATGCTATGGGAGTTGGTGGTGAACCTGAAAACCCCATGGGGGGGTTAAATTTCCGGCAG
ATTCTTCTTTGACAGTTAATTTTAGGCTCGGCCAGAAAATGGTTAGAAAACCTCGGCCGGTTAGGGG AG
CTTTGTCTTTGCCCGCTTGGGTTCCCCCCCCGAAAGGTTTCCCCCTTTTCTATATATT
(SEQ ID NO:30)

FIG. 30

09041303

TGTGAATGTGAACTTCCGCGATCTCCACTGGTGCATTAACCCACTAGTAAGATCAAGGTGAATTTACTAA
CTACGCGATACAATTGGGATCAGAAAATCTATTGCATCGGCAGCAAATCCCATCCTCCTGTGAGCACTCT
CTGGGGGCAGAGGTGACATATTCCACCATAACCGATGCAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTT
CCCCCTGTCAGTGTCAATTGTCCATGTCTTTGATCTCAAGAAAATCAGAGATAATCAATATGCTAACCGCT
ATCTCAAACGGAGTGTATGGTAAACTTATTTACTAGTGCCTGATTATATTGAAGAGGAGTTGACACAC
AAAAGATTGAGTCTTTGAGATAGGGTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAAC
CAACTATATGGTCTCCAGAGAAATCCAAAGCTAAGGTATGTACTATAGCAGTGGGCGAGTTGACACTG
GCTTCTTGTGTGTAGGTGAGAGCACCGTGTCAATTATATCATGACAGCAATGGTTCGCAAGATAGTATCC
TAGCAGTGACGCTGGGAAATTTGGGGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCA
CCCATCAGTAGAAAAAATACATATAACAAATCACCGTGGGTTCATAAAAAGATTCAATAGCAACCTGGATG
GTGCCTGCATTGGTCTCTGAGAAACAGGAAGAGCAAAAAAATGTCTGGAGTCCGGCTTGTCAAAGAAAAT
CCTACCCTATGTGCAACCAAACGTCATGGGAACCTTCCGAGGAGGACAGTTGCCATCTTATGGGCGGTT
GACATTACCTCTAGATCCAAGCACTGACCTTCAACTTAACATATCGTTTACATACGGTCCGGTTATACTG
AATGGAGACGGTATGGATTATTATGAAAGCCCACTGTCGGACTCCGGATGGCTTACCATTCTCCCCAAA
ACGGAACAGTCTTGGATTGATAAACAAAGCAAGTAGAGGAGACCAGTTTCATTGTAATCCCCCATGTGTT
GACATTTGCGCCCAGGGAATCAAGTGGGAATTGTTATTTACCTATTCAAACATCCCAGATTATGAAAAGA
TGCTTAACCCG (SEQ ID NO: 31)

FIG. 31

09041474A

TCTGCTGCTTAACCACCTAGTAAGATCAGGTGAATTTTACTAACTACTGCGATAACAATTGGGATCAGAAA
ATCTATTGCATCGGCAGCAAATCCCATCCTCCTGTGAGCACTCTCTGGGGGCAGAGGTGACATATCCCA
CCATAACCGATGCAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTTTCCCCTGTGAGTGTATTGTCCATGT
CTTTGATCTCAAGAAAAATCAGAGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGTATGGTAAAAAC
TTATTTACTAGTGCTGATTATATTGAAGAGGAGTTTCGACACACAAAAGATTTCGAGTCTTTGAGATAGGG
TTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCCTCCCAGAGAATT
CCAAAGCTAAGGTATGTAATAAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGAGCAC
CGTGTCAATTATATCATGACAGCAATGGTTTCGCAAGATAGTATCCTAGCAGTGACGCTGGGAATATTTGGG
GCAACATCTATGGATCAAGTTGAAGAGGTGAACCTGTTGCTCACCCATCAGTAGAAAAAATACATATAAC
AAATCACCGTGGGTTTCATAAAAAGATTCAATAGCAACTGGATGGTGCCTGCATTGGTCTCTGAGAAACAGG
AAGAGCAAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAAATCCTACCCTATGTGCAACCAAACGTCATG
GGAACCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGCACTGAC
CTTCAACTTAACATATCGTTTACATACGGTCCGGTTATACTGAATGGAGACGGTATGGATTATTATGAAA
GCCCACTGTCCGACTCCGGATGGCTTACCATTCCCTCCAAAAACGGAACAGTCCTTTGGATTGATAAACAA
AGCAGTAGAGGAGACCAGTTTCAATTGTAATCCCCCATGTGTTGACATTTGCGCCAGGGAATCAAGTGGGA
ATTGTTATTTACTTATTCAAACATCCAGATTATGGATAAAGATGTCCTTACTGAGTCCAAATTAGTGTGT
GTGCCTA (SEQ ID NO:32)

FIG. 32

09040826

ATTGGTTGCCCTTAACCACCTAGTAAGATCAGGTGAATTTTACTAACTACTGCGATAACAATTGGGATCA
GAAAATCTATTGCATCGGCAGCAAATCCCATCCTCCTGTGAGCACTCTCTGGGGGCAGAGGTGACATATT
CCCACCATAACCGATGCAGTGGAGCTGCTACCTCAGTAGGCAGAGTTTTTCCCCTGTGAGTGTATTGTCC
ATGTCCTTTGATCTCAAGAAAAATCAGAGATAATCAATATGCTAACCGCTATCTCAAACGGAGTGTATGGTA
AAACTTATTTACTAGTGCTGATTATATTGAAGAGGAGTTTCGACACACAAAAGATTTCGAGTCTTTGAGAT
AGGGTTTCATCAAACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCCTCCCAGAG
AATTCCAAAGCTAAGGTATGTAATAAGCAGTGGGCGAGTTGACACTGGCTTCCTTGTGTGTAGGTGAGA
GCACCGTGTCAATTATATCATGACAGCAATGGTTTCGCAAGATAGTATCCTAGCAGTGACGCTGGGAATATT
TGGGGCAACATCTATGGATCAAGTTGAAGAGGTGATACCTGTTGCTCACCCATCAGTAGAAAAAATACAT
ATAACAAATCACCGTGGGTTTCATAAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTGGTCTCTGAGA
AACAGGAAGAGCAAAAAAATTGTCTGGAGTCGGCTTGTCAAAGAAAAATCCTACCCTATGTGCAACCAAAC
GTCATGGGAACCTTCGGAGGAGGACAGTTGCCATCTTATGGGCGGTTGACATTACCTCTAGATCCAAGC
ACTGACCTTCAACTTAACATATCGTTTACATACGGTCCGGTTATACTGAATGGAGACGGTATGGATTATT
ATGAAAGCCCACTGTCCGACTCCGGATGGCTTACCATTCCCTCCAAAAACGGAACAGTCCTTGAATGATA
AACAAAGCAAGTAGAGGAGACCAGTTTATTGTAATCCCTCTGTGTTTACATTTGCGCCAGGATCAAGT
GGCATTGTTTACTTATTCAAACCTCCGAATTATGGATAAAGATGTCCTTACTGATCCAAACTAGTGCG
TTGCTCAA (SEQ ID NO:33)

FIG. 33

	CDV isolate	155	156	157	158	159	160	161	162	165	169	170	171	172	173	174	175	176	177	178	179	180	185	186	187	189
Reference	Ondersteport	GAG	TCA	ATT	GGG	ATC	AGA	AAA	GCT	TCG	CCT	ATC	CTT	TTA	TCA	GCC	CTA	TCT	GGG	GGC	AGA	AGT	CCA	CAC	AGA	AGT
	AY964110(EW)	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	AF112189(AM-2)	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCT	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	AY962112(AR)	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCC	TTG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGC	AGA	GGT	CCA	TAC	AGA	AGT
European Wildlife	9041474	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	OGA	AGT
	8120857	CAA	ACC	CTT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCC	CCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	OGA	AGT
	9041303	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	OGA	AGT
	8031346	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCA	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	7091032	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	7091030	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	7111080	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	08011277A	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	8080941	GAA	ACA	CTT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	08011277C	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	7101508	TAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CAG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	8010939	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	7091031	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTC	CTG	TCA	GCA	CTC	TCT	GGG	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	American-2	09020504-3(08-7589)	GAT	ACA	ATT	GGG	ATC	AAA	AAA	TCT	TCG	CCT	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGC	AGA	GGT	CCA	TAC	AGA
8060351		GAT	ACA	ATT	GGG	ATC	AGG	AAA	TCT	ACG	CCT	ATC	CTT	TTA	TCA	GCA	CCC	TCC	GGA	GGT	AGA	GGT	TCA	TAC	AGA	AGT
8021509		GAT	ACT	ATG	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGT	AGA	GGT	CCA	TAC	AGA	AAT
9050216		GAC	ACA	ATT	GGG	ATC	ACG	AAA	TCT	TCG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGC	ACA	GGT	TCA	TAC	AGA	CGT
8050180		GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGT	AGA	GGT	CCA	TAC	AGA	AGT
7100609		GGC	ACA	ATT	GGG	TAG	CAT	GAG	TTT	TCG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGC	AGA	GGT	CCA	TAC	AGA	AGT
8080696		GCC	CCC	ATT	GGG	ATC	TTA	CAA	TCT	TCG	CCC	ATC	CTC	TTA	TCA	GCA	CTC	TCC	CGA	GGC	AGA	GGT	CCA	TAC	AGA	AAT
9011024		GAT	TCC	CTG	GGG	GCA	CAT	GTC	AAC	TCG	CCC	ATC	CTC	TTA	TCA	GCA	CTC	TCC	GGA	GGC	AGA	GGT	CCA	TAC	AGA	AGT
8030777	GAT	ACT	ATT	GGC	ATC	AGA	AAA	TCT	TCG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGT	AGA	GGT	CCA	TAC	AGA	AAT	
Arcic	7110098	GAT	ACA	ATT	GGG	ATC	AGA	AAA	TCC	TTG	CCC	ATC	CTT	TTA	TCA	GCA	CTC	TCC	GGA	GGC	AGA	GGT	CCA	TAC	AGA	AGT
	gagE	tra5	atd	ggg	atd		aaak	gctA	acgT	cccP		cttL	ctgI	tra5	gcaA	ctal	tcc5	gggG	ggcG	agaR	O	ccaP	O	agaR	agT5	
	gatD	acaT	cttL	ggcG	tagterm		gagE	tcc5	tcc5	ctcP		ctcL	ttal	tcc5	gcaA	ctcL	tcc5	gggG	ggcG	acaT		cca5		agaR	aatN	
	taty	accT	atgM		gcaA		caaQ	tcc5	ttgI	ccaP		ctaI	cagQ		ccaP			ggaG	ggT6					ccaR		
	ggcD	actT	ctgI			agaR	gctV											ggaR								
	gccA	accP				aggI												ccaP								
						acgT	caH																			

Figure 34A

	CDV Isolate	191	192	193	195	197	198	199	201	203	205	206	209	210	212	213	214	215	220	222	225	226	228	229	232	233	235
Reference	Ordersteport	GCT	ACT	ACT	GTA	AAA	GTT	TTC	CTA	GTC	TTA	TCC	CTG	ATC	AGA	ACC	TCA	GAG	CTG	GCT	GAC	GGC	TAT	GGC	TAC	TTG	GTG
	AY964110(EW)	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	AF112189(AM-2)	GCT	ACT	ACT	GTA	AGA	GTT	TTC	CTA	GTA	TTG	TCC	TCG	ATC	AGA	ACA	TCA	GAG	CTA	GCT	GAC	GGA	TAT	GGT	TAT	TTA	GTG
	AY962112(AR)	GCT	ACT	ACT	GTT	AAA	TCT	TTC	CTA	GTA	TTA	TCC	TCG	ATC	AGA	ACA	TCA	GAG	CTG	TCT	GAC	GGA	TAT	GGT	TAT	TTA	GTG
European Wildlife	9041474	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTG	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	8120857	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	CAT	GGT	TAT	TTA	GTG
	9041303	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTG	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	8031346	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	7091032	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	7091030	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	7111080	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	08011277A	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	8080941	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	08011277C	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	7101508	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	8010939	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	7091031	GCT	GCT	ACC	GTA	AGA	GTT	TTC	CTA	GTG	TTG	TCC	TCG	ATC	AGA	AAA	TCA	GAG	CTA	GCT	AAC	GGA	TAT	GGT	TAT	TTA	GTG
	American-2	09020504-3(08-7589)	GCT	ACT	ACT	GTA	AGA	GTC	TTC	CTA	GTA	TTG	TCC	TCG	ATC	AGA	ACA	TCT	GAA	CTA	GCT	GAC	GGA	TAT	GGT	TAT	TTG
8060351		GCT	ACT	ACT	GTA	AGA	GTC	TTC	CTA	GTA	TTG	TCC	TCG	ATC	AGA	ACA	TCT	GAA	CTA	GCT	GAC	GGA	TAT	GGT	TAT	CTG	GTT
8021509		GCT	ACT	ATC	GTA	AAG	ATT	TTC	CTA	GTA	TTA	TCT	TCG	ATC	AGA	ACA	TCA	GAG	CTA	GCT	GAC	GGA	TAT	GGT	TAT	TTA	GTG
9050216		GCT	ACT	ACT	GTA	AGA	GTC	TTC	CTA	GTA	TTG	TCC	TCG	ATC	AGA	ACA	TCT	GAA	CTA	GCT	CAC	GGA	TAT	GGT	TAT	CTG	GTT
8050180		GCT	ACT	ACT	GTA	AGA	GTC	TTC	CTA	GTA	TTG	TCC	TCG	GTC	AGA	ACA	TCT	GAA	CTA	GCT	GAC	GGA	TAT	GGT	TAT	TTG	GTT
8030777		GCT	ACT	ATC	GTA	AAG	ATT	TTC	CTA	GTA	TTA	TCT	TCG	ATC	AGA	ACA	TCA	GAG	CTA	GCT	GAC	GGA	TAT	GGT	TAT	TTA	GTG
Arctic	8080696	GCT	ACT	ATC	CTA	AAG	ATT	TCC	CTA	GTA	TTA	TCT	TCG	ATC	CGA	ACA	TCA	GAG	CTA	GCT	TAC	GGA	TAT	GGT	TAT	TTA	GTG
	7110098	GCT	ACT	ACT	GTT	AGA	TCT	TTC	CTA	GTA	TTA	TCC	TCG	ATC	AGA	ACA	TCT	GAG	CTG	TCT	GAC	GGT	TAT	GGT	TAT	TTG	GTG
			actT	actI	gtvV	aaak	gttV	tteF	ctal	gtcV	ttal	tccS	ctgL	atcl	agar	acaT	tcaS	gagE	ctgL	gctA	gacD	ggcG	tatY	ggcG	0	ttgL	gtgV
			gcta	accT	gttV	agal	gtcV	tccS	ctgL	gtgV	ttgL	tctS	tccS	accT	cgaR	accT	tctS	gaaE	ctal	tctS	aacN	gggG	catH	gggG		ttal	gtgV
			atcl	atcl	ctal	aagK	attI			gtvV				gtcV	aaaK						cacH	gggG				ctgL	

Figure 34B

	CDV isolate	236	237	238	239	241	242	245	246	247	252	255	258	261	262	267	269	273	275	277	278	280	282	286	288	291
Reference	Ondersteport	CCT	GAT	GAT	ATA	AGA	GAG	ACT	CAA	GAG	GAA	TTC	AGG	AAT	GAC	CAA	ACC	GTA	CCG	AAT	TCC	GCC	GTA	GCA	GGT	ACA
	AY964110(EW)	CCA	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	AF112189(AM-2)	CAT	GAT	TAT	ATT	GGG	GGG	ACG	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCG	AAT	TCC	GCC	GTA	GCG	GGC	ACA
European Wildlife	AY962112(AR)	CCT	GAT	TAT	ATT	GGG	GAG	ACG	CAA	AAG	GAG	TTC	AGG	AAT	GAC	CAG	ACC	ATC	CCG	AAT	TCT	ACC	GTA	GCA	GGC	ACA
	9041474	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	8120857	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAC	AAC	CAG	ACT	GTC	CCA	AAT	TCC	GCT	GCA	GCA	GGC	CCC
	9041303	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	8031346	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	7091032	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACT	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	7091030	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	7111080	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	08011277A	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	8080941	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	AAC	CAG	ACT	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	08011277C	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	7101508	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	8010939	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GCA	GGC	ACA
	7091031	CCT	GAT	TAT	ATT	GAG	GAG	ACA	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCT	GTA	GGA	GGC	ACA
	American-2	09020504-3(08-7589)	CCT	GAT	TAT	ATT	GGG	GAG	ACG	CCG	AAG	GAG	TTC	CGG	AAC	AAC	CAG	ACC	GTC	CCG	GAT	TCC	GCT	GTA	GCA	GGC
8060351		CCT	GAT	TAT	ATT	GGG	GAG	ACG	CAA	AAG	GAG	TTC	CGG	AAC	AAC	CAG	ACC	GTC	CCA	GAT	TCC	GCC	GTA	GCA	GGC	ACA
8021509		CCT	GAT	TAT	ATT	GGG	GAG	ACG	CAA	AAG	GAG	TTC	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ACA
9050216		CCC	GAT	TAT	ATT	GGG	GAG	ACC	CAA	AAG	GAG	TCC	CGG	AAC	AAC	CAG	ACC	GTC	CCG	GAT	TCC	GCC	GTA	GCA	ND	ACA
8050180		CCT	GAT	TAT	ATT	GGG	GAG	ACG	CAA	AAG	GAG	TTC	CGG	AAC	AAC	CAG	ACC	GTC	CCG	GAT	TCC	GCC	GTA	GCG	GGC	ACA
8030777		CCT	GAT	TAT	ATT	GGG	GAG	ACG	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	GTC	CCA	AAT	TCC	GCC	GTA	GCA	GGC	ACA
Arctic	8080696	CCC	GAC	TAT	ATT	GGG	GAG	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
	7110098	CCT	GAT	TAT	ATT	GGG	GAG	ACG	CAA	AAG	GAG	TTC	CGG	AAT	GAC	CAG	ACC	ATC	CCG	AAT	TCT	ACC	GTA	GCA	GGC	ACA
	cctP	gatD	o	ataM	agaR	gagE	actT	caaQ	o	o	ttf	o	aatH	gacD	o	accT	gatV	ccgP	aatH	ttcS	gcaA	gatA	gcaA	ggcG	accT	cccP
	ccaP	gacD		attI	gagE	gggG	acaT	ccgP			ttcS		aacN	aacN		actT	gtcV	ccaP	gatD	ttcS	gcaA	gcaA	ggcG			
	catH			o	gggG		acgT										atcI					accT				
	cccP																									

Figure 34C

	CDV isolate	294	295	296	298	301	302	303	309	311	314	315	317	319	321	322	323	324	327	330	331	332	333	337	340	341
Reference	Ondersteport	TCC	TTG	TGT	GAA	ACT	GTA	TTA	AGT	TCA	GGT	ATT	GTA	ACA	GGG	ATA	TTT	TGG	CCT	CAC	ATT	GAG	GAA	GTC	CCA	TCA
	AY964110(EW)	TCC	GTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	AF112189(AM-2)	TCC	TTG	TGT	GAT	ACC	GTA	TTG	GAT	TCA	GGT	ATT	GTG	ACG	GGG	ATA	TTT	GGG	CCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	AY962112(AR)	TCC	TTG	TGT	GAT	ACT	GTA	TTA	AAT	TCA	GGT	ATT	GTA	ACG	GGG	ATA	TTT	GGG	CCT	CAA	GTC	GAA	GAG	GTC	CCA	TCA
European Wildlife	9041474	TCC	TTG	TGT	GGT	ACC	GTG	TCA	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	8120857	TCC	TCG	CGT	GGT	ACC	GTG	CTG	CCT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGC	ND	ND	ND	ND	ND	ND	ND	ND
	9041303	TCC	TTG	TGT	GGT	ACC	GTG	TCA	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	8031346	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	7091032	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	7091030	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	ACT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	7111080	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	08011277A	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AAT	ATC	GTA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	8080941	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AGT	ATC	GCA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	08011277C	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AAT	ATC	GTA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	7101508	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AAT	ATC	GTA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	8010939	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	AAT	ATC	GTA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
	7091031	TCC	TTG	TGT	GGT	ACC	GTG	TTG	AAT	TCG	ND	ATC	GCG	ACG	GGG	ATA	TTT	GGG	ND	ND	ND	ND	ND	ND	ND	ND
	American-2	09020504-3(08-7589)	TCC	TTA	TGT	GAT	ACC	GTA	TTG	AAT	TCA	GGT	ATT	GTG	ACG	GGG	ATA	TTT	GGG	CCT	CGA	GTT	GAA	GAG	GTT	CCG
8060351		TCC	TTG	TGT	GAT	ACC	ATA	TTG	AAT	TCA	GGT	ATT	GTG	ACG	GGG	ATA	TTT	GGG	CCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
8021509		ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	GGG	ND	ND	ND	ND	ND	ND	ND	ND
9050216		ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	GGG	ND	ND	ND	ND	ND	ND	ND	ND
8050180		TCC	TTG	TGT	GAT	ACC	GTA	TTG	AAT	TCA	GGT	ATT	GTA	ACG	GGG	ATA	TTT	GGG	CCT	CAA	GTT	GAA	GAG	GTT	CCA	TCA
8030777		TCT	TTG	TGT	GAT	ACC	GTA	TTG	AAT	TCA	GGT	GTT	GTA	ACG	GGG	ATA	TTT	GGG	TCT	CAA	GTT	GAA	GAG	GTC	CCA	TTA
8080696		ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
Afzdc	7110098	TCC	TTG	TGT	GAT	ACT	GTA	TTA	AAT	TCA	GGT	ATT	GTG	ACG	GGG	ATA	TTT	GGG	CCT	CAA	GTC	GAA	GAG	GTC	CCA	TCA
		tcc5	ttg	tgt	gat	act	gta	tta	aat	tca	ggg	att	gtg	acg	ggg	ata	ttt	ggg	cct	caa	gtc	gaa	gag	gtc	cca	tca
		tct5	gtgV	cgtR	gatG	accT	gtgV	ttgL	aatN	tcg5	gggG	atcl	gcaA			atcl			tct5	caaQ	gtrV			gtrV	ccgP	ttal
			ttal		gatD			tca5	gatD		aatN	gtrV	gtrV						actT	qgaR	gtrV					

Figure 34D

	Isolate	342	343	347	348	349	352	353	354	356	358	365	366	367	370	371	372	373	375	376	380	386	388	391	393	396	
Reference	Ondersteport	ATG	GAG	ATA	ACA	AAC	GGT	TTT	ATA	GAT	ATT	GCC	CTG	GCC	AAA	CAA	GAA	GAA	AAA	GGT	TCA	ACC	CCC	AAC	ACG	GAA	
	AY964110(EW)	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	AF112189(AM-2)	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTA	AAA	CAA	GAG	GAA	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	AY962112(AR)	GTC	GAA	ATA	ACA	AAT	GGT	TTC	ATA	GAT	GTA	GCA	TTG	GTC	AAC	CTA	GAG	GAA	GAA	AAT	TCG	TCC	CCT	AAT	ACA	GAA	
European Wildlife	9041474	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	8120857	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	
	9041303	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	8031346	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	7091032	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	7091030	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	7111080	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAA	TCG	TCC	CCT	AAC	ACG	GAA	
	08011277A	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GGA	
	8080941	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	08011277C	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	7101508	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	ATT	TCG	TCC	CCT	AAC	ACG	GAA	
	8010939	GTA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAT	ATA	GCA	TTG	GTC	AAA	CAG	GAA	GAG	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	7091031	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	
American-2	09020504-3(D8-7589)	GTA	GAA	ATG	GCA	AAT	GGG	TTC	ATC	GAT	ATA	GCA	TTG	GTC	AAA	CAA	GAG	GAA	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	8060351	GTA	GAA	ATA	GCA	AAT	GGG	TTC	ATC	GAT	ATA	GCA	TTG	GTC	AAA	CAA	GAG	GAA	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	8021509	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	
	9050216	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	
	8050180	GTA	GAA	ATA	GCA	AAT	GGG	TTC	ATC	GAT	ATA	GCA	TTG	GTC	AAA	CAA	GAG	GAA	AAA	AAT	TCG	TCC	CCT	AAC	ACG	GAA	
	8030777	GCA	GAA	ATA	ACA	AAT	GGG	TTC	ATA	GAC	ATA	GCA	TTG	GTT	AAA	CAA	GAG	GAA	GCA	AAT	TCG	TCC	CCT	AAT	ACG	GAA	
Arctic	7110098	GTC	GAA	ATA	ACA	AAT	GGT	TTC	ATA	GAT	GTA	GCA	TTG	GTC	AAC	CTA	GAG	GAA	GAA	AAT	TCG	TCC	CCT	AAT	ACA	GAA	
		atgM gtaV gtcV	0	atal atgM	acaT gcaA	0	gggG gggG	0	atal atcd	0	attl atal gtaV	0	0	gcaA gtcV gtaV gttV	aaaK aacN	caaQ cagQ ctal	gaaE gagE	gaaE gagE	gcaA gaaE aaaK	0	0	0	0	0	aaaN aatN	acaT acgT	

Figure 34 E

	CDV isolate	398	401	410	411	412	414	415	416	417	418	419	420	422	423	424	425	427	428
Reference	Ondersteport	TTC	AGA	ACA	TTA	CCT	GAT	GCA	AGT	GTT	GAC	CTT	CAA	AAC	CTA	TCG	TTC	TAC	GGT
	AY964110(EW)	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
	AF112189(AM-2)	TTT	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ATT	GAC	CTT	CAA	AAC	CTC	TCG	TTT	TAC	GGT
	AY962112(AR)	TTT	GGA	ACG	TTA	CAT	GAT	GCA	AGC	ATT	GAC	CGT	CAA	AAC	CTA	TCA	TTT	TAC	GGT
European Wildlife	9041474	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
	8120857	TTC	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
	9041303	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
	8031346	TTC	GGA	GAC	ATT	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	TAC	CTA	TCG	TTT	TAC	ND
	7091032	TTC	GGA	ACA	TTT	CCT	GAT	CCA	AGC	ACT	GAA	CTT	CAA	AAC	CTA	TCG	TTT	TAC	CGG
	7091030	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
	7111080	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND
	08011277A	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CCT	CAA	AAC	CTA	TCG	TTT	TAC	CGT
	8080941	TTT	GGA	ACT	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	ND	ND	ND	ND	ND
	08011277C	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
	7101508	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
	8010939	TTC	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ACT	GAC	CTT	CAA	AAC	CTA	TCG	TTT	TAC	GGT
7091031	TTC	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	TAC	GGT
American-2	09020504-3(08-7589)	TTT	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ATT	GAC	CTT	CAC	AAC	ATC	TCA	TTT	TAC	GGC
	8060351	TTT	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ATT	GAC	CTT	CAG	AAC	ATC	TCA	TTT	TAC	GGC
	8021509	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	TAC	GGT
	9050216	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	TAC	GGT
	8050180	TTT	GGA	ACA	TTA	CCT	GAT	CCA	AGC	ATT	GAC	CTC	CAG	AAC	ATC	TCA	TTT	TAT	GGT
	8030777	TTT	GGA	ACA	TTA	CCT	CAT	CCA	AGC	ATT	GAC	CTC	CAA	AAC	CTA	TCA	TTT	TAC	GGT
Arctic	7110098	TTT	GGA	ACG	TTA	CAT	GAT	CCA	AGC	ATT	GAC	CTT	CAA	AAC	CTA	TCA	TTT	TAC	GGT
		ttrP	O	acaT	ttal	cctP	gatD	O	O	gtrV	gacD	cttL	caaQ	aacN	ctal	trgS	ttcF	tacC	ggtG
		ttcP		acgT	tttF	cath	cath			actT	gaeE	cgtR	cach	tacY	ctcl	tcaS	tttF	tatY	ggoG
				gacD						attI	ctcl	cagQ		atd					

Figure 34 F

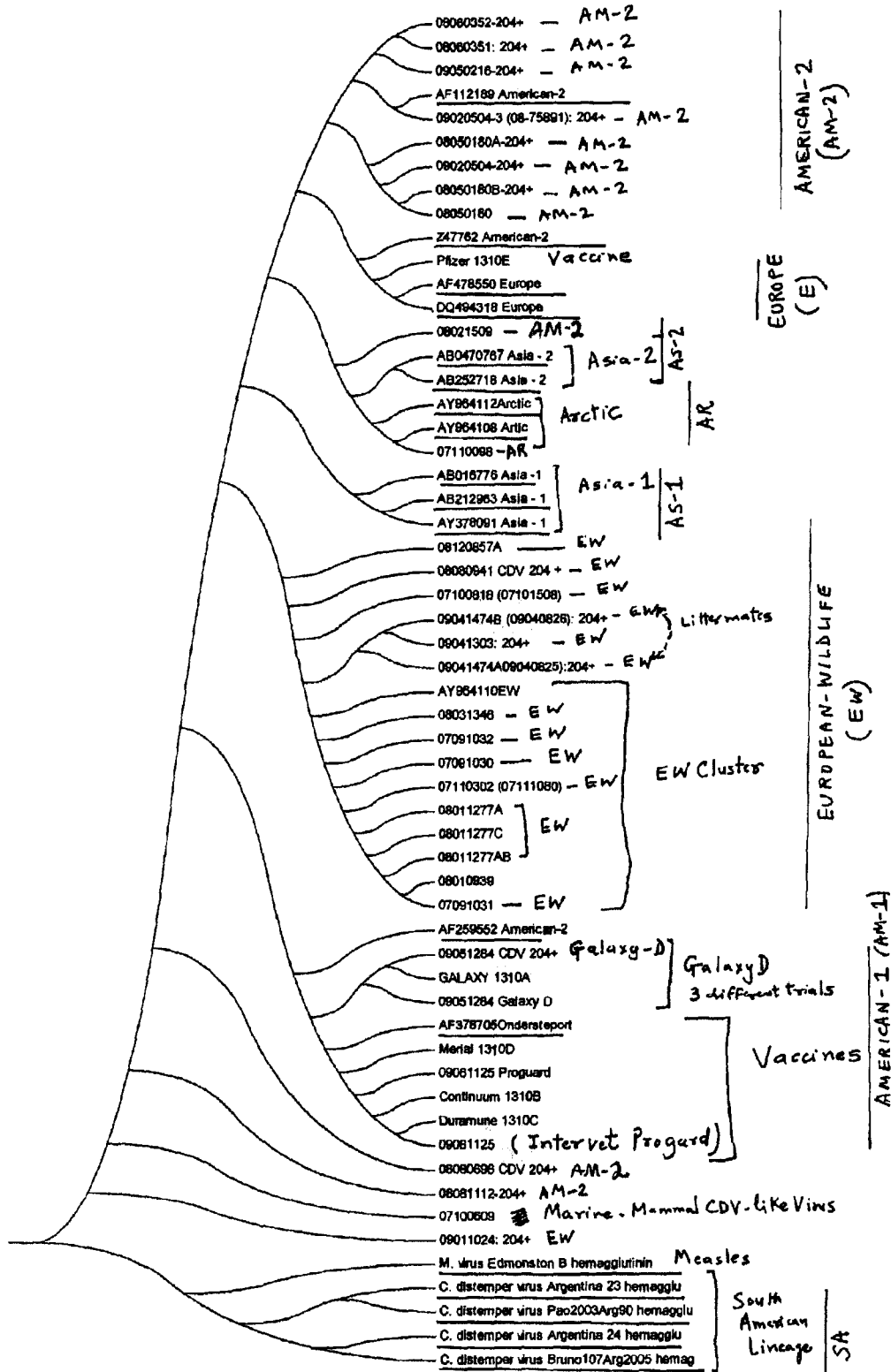


Figure 35

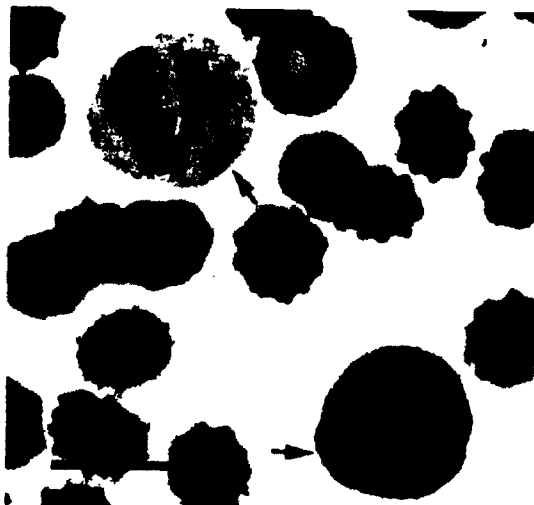


Figure 36

09041474B

atgctctcctaccaagacaaggtgggtgccttctataaggataatgcaagagctaattcatccaagctgt
ccctagtgcagagaagagcaagggggcaggagaccacctatctgctgtttgtccttctcactcctactggt
tggaatcctggccttgccttgcctatcactggagttcgatttcaccaagatcaactagcaacgtggaattt
agcagattgctaaaagaggatattggagaaatcagaggctgtacatcaccaagtcataagatgttttgacgc
cgctcttcaaaattattggagatgagattgggttacggctgccacaaaactaaacgagatcaaacaatt
catccttcaaaagacaaaacttcttcaatcctaacaggggaattcgacttccgtgatctccactgggtgcatt
aaccacctagtaagatcaaggtgaattttactaactactgcgatacaattgggatcagaaaatctattg
catcggcagcaaatcccatcctcctgtcagcactctctgggggcagaggtgacatattcccaccataccg
atgcagtgaggctgctacctcagtaggcagagttttccccctgtcagtgctattgtccatgtctttgatc
tcaagaaaatcagagataatcaatattgctaaccgctatctcaaacggagtgtatggtaaaacttatttac
tagtgctgattatattgaagaggagttcgacacacaaaagattcgagtcctttgagatagggttcatcaa
acggtggctgaatgacatgccattactccagacaaccaactatatggtcctccagagaattccaaagct
aaagtatgtactatagcagtgggcgagttgacactggcttccttgtgtgtaggtgagagcacgctgtcat
tatatcatgacagcaatgggttcgcaagatagtatcctagcagtgacgctgggaatattggggcaacatc
tatggatcaagttgaagaggtgatacctgttgctcaccocatcagtagaaaaatacatatacaaatcac
cgtgggttcataaaagattcaatagcaacctggatgggtgcctgcattggtctctgagaaacaggaagagc
aaaaaaattgtctggagtgggttgcaagaaaatcctaccctatgtgcaaccaaactcatgggaacc
cttcggaggaggacagttgccatcttatgggcgggtgacattacctctagatccaagcactgacctcaa
cttaacatatcgtttacatacgggtccgggttatactgaatggagacgggatggatattatgaaagcccac
tgtcggactccggatggcttaccattcctccaaaaacgggaacagtccttggattgataaacaagcaag
tagaggagaccagttcatttgtaatccccatgtgttgacatttgccgccagggaatcaagtgggaattgt
tatttacctattcaaacatcccagattatggataaagatgtccttactgagtcctaatttagtgggtgtgc
ctacacagaattttagatatgtcatagcaacatatgatatatcccgggacaatcatgcgatcgtttacta
tgtctatgacccaattcggacgatttcttatacgtacctttagactaactaccaaaggttagacctgat
ttcctaaggattgaaatgttttgggttgatgatattgtgggtgtcaccagttctaccgattcgaggctg
acatcactaactctaccaccagtggttgagaatttagtccgtataagattctcatgtaaccggttcaagacc
ttga (SEQ ID NO: 42)

Fig. 37

08021509

ATGCTCTCCTACCGAGACAAGGTGGGTGCCTTCTATAAGGACAATGCTAGAGCTAATTCATCCAAGCTGT
CCTTAGTGACAGAAGAGCAAGGGGGCAGGAGACCACCTATTTGCTGTTTGTCTTCTCATCCTACTGGT
TGGAATCATGGCCTTGCTTGCTATCACTGGAGTTCGATTTACCAAGTATCAACTAGCAATATGGAGTTT
AGCAGATTGCTGAAAGAGGATCTGGAGAAATCAGAGGCCGTACATACCAAGTCATAGATGTCTTGACGC
CGCTCTTCAAAATATTGGAGATGAGATTGGGTTACGGTTGCCACAAAACTAAACGAGATCAAAACAATT
TATCCTTCAAAAGACAAACTTCTTCAATCCGAACAGGGAATTCGACTCCGCGATCTCCACTGGTGCATT
AACCCACCTAGTAAGATCAAGGTGAATTTACTAATTACTGCGATACTATGGGGATCAGAAAATCTATTG
CATCGGCAGCAAATCCCATCCTTTTATCAGCACTCTCCGAGGTAGAGGTGACATATCCACCATACAG
ATGCAATGGAGCTACTATTTCAAGTAGGCAAGATTTTCCCCTATCAGTATCATTATCTATGTCTTTGATC
TCAAGAACATCAGAGATAATCAATATGCTAACCCTATCTCAGACGGAGTGATGGTAAACTTATTTAC
TAATGCCTGATTATATTGAAGGGGAGTTCGACACGCAAAAGATTGAGTCTTTGAGATAGGGTTCATCAA
ACGGTGGCTGAATGACATGCCATTACTCCAGACAACCAACTATATGGTCTCCAGAGAAATCCAAAGCT
AAGGTATGTACTATAGCAGTGGGCGAGTtGACACTGGCTTCTTTGTGTGAGGTGAGAGCACCGTATTGT
TATATCATGACAGCAATGGTTCACAAGATGGTATTCTAGTAGTGACGCTGGGAATATTCGGGGCAACATC
TATGGATCAAGTTGAAGAGGTGATACCTGTCGCTGACCCATTAGTAGAAAAAATACATATAACAATCAC
CGCGGATCATAAAAGATTCAATAGCAACCTGGATGGTGCCTGCATTAGTTTCTGAGAAACAAGAGGAAC
AAAAAATTGTCTGGAGTCAGCTTGTCAAAGAAAATCCTACCCTATGTGCAATCAAACGTCATGGGAACC
CTTTGGAGGAGGACAGTTGCCATCTTATGGGCGTTCGACATTACCTCTAGATCCAAGCATTGACCTTCAA
CTTAACATATCATTACATACGGTCCGATTATACTGAATGGGGACGGTATGGATTATTATGAGAGCCCAC
TGTTGGACTCCGGATGGCTTACCATTCTCCCAAGAACGGAACAGTCCTTGGATTGATAAACAAGCAAG
TAGAGGAGACCAGTTCAGTGAATCCCCATGTGTTGACATTTGCGCCAGGGAATCAAGTGGAAATTGT
TATTTACCTATTCAAACATCCAGATTATGGATAAAGATGTCCTTACTGAGTCCAATTTAGTGGTGTTC
CTACACAGAATTTTAGATATGTCTGAGCAACATATGATATATCTCGGGACGATCATGCCGATGTTTTATTA
TGTTTATGACCCAATACGGACGATTTCTTATACGTACCCATTTAGACTAACTACTAAGGGTAGACCTGAT
TTCTTAAGGATTGAGTGTTTTGTGTGGGATGACGATTTGTGGTGTACCAGTTTTACCGATTCCAGGCCG
ACATCACAACCTCTACAACCAGTGTGAGAATTTAGTCCGTATGAGATTCTCATGTACCCTTCCAGACC
TTGA (SEQ ID NO: 43)

Figure 38

MLSYQDKVGAFYKDNARANSSKLSLVTEEQGRRPPYLLFVLLILLVIGILALLAITGVRFHQVSS
NVEFSRLLKEDMEKSEAVHHQVIDVLTPLFKIIGDEIGLRLPQKLNEIKQFILQKTNFFNPNREF
DFRDLHWCINPPSKIKNFTNYCDTIGIRKSIASAANPILLSALSNGGRDIFPPYRCGAATSVG
RVFPLSVLSMSLISRKSEIINMLTAISNGVYKTYLLVPDYIEEEFDTQKIRVFEIGFIKRWLN
DMPLLQTTNYMVLPENSKAKVCTIAVGELTLASLCVGESTVSLYHDSNGSQDSILAVTLGIFGAT
SMDQVEEVIIPVAHPSVEKIHI TNHRGFIKDSIATWMVPALVSEKQEEQKNCLESACQRKSYPMC
QTSWEPFGGGQLPSYGRLLPLDPSTDLQLNISFTYGPVILNGDGMYYESPLSDSGWLTIPPKN
GTVLGLINKASRGDQFIVIPHVLTFA PRESSGNCYLPIQTSQIMDKDVLTESNLVVLPTQNFYV
IATYDISRDNHAI VYVYDPIRTISYTYPFRLTTKGRPDFLRIECFVWDDDLWCHQFYRFEADIN
STTSVENLVIRIRFSCNRSRP (SEQ ID NO: 44)

Fig. 39

MLSYRDKVGAFYKDNARANSSKLSLVTEEQGRRPPYLLFVLLILLVIGIMALLAITGVRFHQVST
SNMEFSRLLKEDLEKSEAVHHQVIDVLTPLFKIIGDEIGLRLPQKLNEIKQFILQKTNFFNPNRE
FDFRDLHWCINPPSKIKNFTNYCDTMGIRKSIASAANPILLSALSNGGRDIFPPYRCNGATISV
GKIFPLSVLSMSLISRSEIINMLTAISDGVYKTYLLMPDYIEGEFDTQKIRVFEIGFIKRWL
NDMPLLQTTNYMVLPENSKAKVCTIAVGELTLASLCVGESTVLLYHDSNGSQDGILVVTLGIFGA
TSMDQVEEVIIPVADPLVEKIHI TNHRGIIKDSIATWMVPALVSEKQEEQKNCLESACQRKSYPMC
NQTSWEPFGGGQLPSYGRLLPLDPSIDLQLNISFTYGPVILNGDGMYYESPLSDSGWLTIPPKN
NGTVLGLINKASRGDQFTVIPHVLTFA PRESSGNCYLPIQTSQIMDKDVLTESNLVVLPTQNFYV
VVATYDISRDDHAI VYVYDPIRTISYTYPFRLTTKGRPDFLRIECFVWDDDLWCHQFYRFEADI
TNSTTSVENLVRRMRFSCNRSRP (SEQ ID NO: 45)

Fig. 40

1

**IMMUNOGENIC COMPOSITIONS,
VACCINES AND DIAGNOSTICS BASED ON
CANINE DISTEMPER VIRUSES
CIRCULATING IN NORTH AMERICAN DOGS**

CROSS-REFERENCE TO RELATED
APPLICATIONS

This application claims benefit of U.S. provisional patent application 61/148,791, filed Jan. 30, 2009, the complete contents of which is hereby incorporated by reference.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The invention generally relates to newly identified isolates of canine distemper virus (CDV). In particular, the invention provides improved CDV immunogenic compositions, vaccines and diagnostics that contain or take into account these newly discovered isolates, and describes a systematic protocol for selecting, based on genetic makeup, broad spectrum isolates for use in immunogenic compositions, vaccines and diagnostics.

2. Background of the Invention

Canine distemper virus (CDV) is a single-stranded RNA Morbillivirus that affects dogs of all ages. CDV causes a multi-systemic infection that may involve the ocular, respiratory, gastrointestinal, integument and nervous systems, and is usually rapidly fatal. While the disease is a devastating problem for dogs, other species are also susceptible to the virus, for example, raccoons, foxes, coyotes, wolves, various fur-producing animals, and large non-domestic cats such as lions, leopards, cheetahs, and tigers. In the past, vaccines have proven to be effective in reducing the incidence of CDV infection. However, there appears to be a resurgence of the incidence of CDV, even in fully vaccinated animals.

The hemagglutinin (H) protein of CDV is a viral surface protein that is involved in host cell-virus binding, and mutations in the protein affect host cell-virus interactions. H protein is considered to be a virulence factor for CDV. The H protein displays significant (e.g. about 10%) variation in amino acid sequence among CDV isolates, and phylogenetic analysis of this variation serves as the basis for the division of viral isolates into seven lineages: American-1, American-2, Arctic-like, Asia-1, Asia-2, Europe, and European wildlife (McCarthy, A. J., M. A. Shaw, and S. J. Goodman. 2007. *Proc. Biol. Sci.* 274:3165-3174). Antibodies to H protein provide protection against infection, and are thus the likely basis for vaccine efficacy. However, antibodies do not necessarily cross-react between lineages. Hence, vaccines based on a particular isolate may or may not provide the vaccine recipient with protection against infection with other isolates. This is particularly problematic given 1) the high rate of mutation exhibited by RNA viruses such as CDV and 2) the increase in the global transport of dogs from one country to another, which fosters the introduction of new lineages into territories where they were previously unknown. Further, for dogs vaccinated with a particular CDV isolate, exposure to a genetically distant CDV may lead to sequestration of the incoming CDV virus in immunologically privileged sites (e.g. brain, ganglion, spinal cord, central, autonomic nervous systems, nasal plenum and bladder epithelium), allowing the propagation and spread of the genetically distant CDV without detection, since neurological symptoms may be overlooked by veterinary practitioners due to lack of sensitivity of the diagnostic tests and expense of long term treatment of a neurological patient.

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Unfortunately, CDV vaccines currently in use have not been updated for about 60 years (Woma et al., 2010. Phylogenetic analysis of the hemagglutinin gene of the current wild-type canine distemper viruses from South Africa: Lineage Africa.Vet. Microbiol. doi:10.1016/j.vetmic.2009.11.013) and have not kept pace with these changes. The use of these outdated vaccines is the likely cause of recent outbreaks of CDV infection, since these vaccines may not provide protection against infection with newly emerging lineages of CDV. Moreover, PCR sequencing has revealed that the vaccine isolate used in one commercial vaccine was misidentified (Demeter et al., 2009: Controversial results of the genetic analysis of a canine distemper vaccine strain. *Vet. Microbiol.* Published Online), further complicating the problem of determining how to best detect, monitor, and prevent CDV infection and transmission.

Clearly, epidemiological studies to investigate the rise in CDV clinical cases are warranted, as is the development of new immunological and vaccine compositions and diagnostic methods that take into account emerging isolates of CDV.

SUMMARY OF THE INVENTION

In one aspect, the present invention provides newly identified isolates of canine distemper virus (CDV). Accordingly, the present invention further provides updated immunogenic compositions and vaccine compositions. The vaccine compositions of the present invention are designed to provide broad-spectrum protection against emerging forms of CDV. In addition, the present invention provides updated diagnostic methods and kits for detecting CDV infection. The diagnostic methods and kits provide the ability to detect the newly evolved forms of the virus. The compositions and diagnostic methods and kits of the present invention are based, at least in part, on the discovery of previously unknown CDV variants, and take into account the emergence of mutant forms of the virus for which prior vaccine formulations and diagnostics are inadequate.

In a further aspect, the present invention provides a systematic method for selecting an antigen, e.g., a pathogenic isolate or portion thereof, that correspond to the genetic makeup of a broad spectrum of the source of the antigen, e.g. pathogen isolates or portion thereof, for use in such compositions and diagnostics.

The present invention further provides an isolated canine distemper virus (CDV) of European wildlife (EW) lineage comprising the characteristics of CDV 9041474B CDV-EW (ATCC Deposit No. PTA-10596). In another embodiment, the invention provides an attenuated strain of CDV isolated in cell cultures in which CDV strain CDV 9041474B CDV-EW (ATCC Deposit No. PTA-10596), or a progeny strain thereof, has been propagated. In a particular embodiment of this type, the attenuated strain of CDV may be plaque-purified. In yet another embodiment, the invention provides an immunogenic composition or vaccine, comprising the isolated CDV comprising the characteristics of CDV 9041474B CDV-EW (ATCC Deposit No. PTA-10596), or progeny thereof.

In still another embodiment, the invention provides an isolated canine distemper virus (CDV) of American-2 (AM-2) lineage having the characteristics of CDV 08021509 CDV-AM-2 (ATCC Deposit No. PTA-10597). In yet another embodiment, the invention provides an attenuated strain of CDV isolated in cell cultures in which CDV strain CDV 08021509 CDV-AM-2 (ATCC Deposit No. PTA-10597), or a progeny strain thereof, has been propagated. In a particular embodiment of this type, the attenuated strain of CDV may be plaque-purified. In a further embodiment, the invention pro-

vides an immunogenic composition or vaccine comprising the isolated CDV having the characteristics of CDV 08021509 CDV-AM-2 (ATCC Deposit No. PTA-10597), or progeny thereof.

The present invention also provides methods of eliciting an immune response to canine distemper virus in a subject in need thereof. One such method comprises administering to the subject an immunogenic composition or vaccine comprising an isolated CDV comprising the characteristics of CDV 9041474B CDV-EW (ATCC Deposit No. PTA-10596), or progeny thereof. In another such embodiment, the method comprises administering to said subject the immunogenic composition or vaccine comprising the isolated CDV having the characteristics of CDV 08021509 CDV-AM-2 (ATCC Deposit No. PTA-10597), or progeny thereof.

The present invention also provides diagnostic kits. One such embodiment comprises oligonucleotide primers specific for amplifying a nucleotide sequence as set forth in SEQ ID NO: 42. In another embodiment, the diagnostic kit comprising oligonucleotide primers specific for amplifying a nucleotide sequence as set forth in SEQ ID NO: 43. Those of skill in the art will recognize that such primers are based on the nucleotide sequence of a nucleotide sequence of interest that is to be amplified (e.g. a sequence that is targeted). In some embodiments, primers are homologous to or complementary to unique sequences of the target sequence.

The present invention further provides the recombinant and/or isolated nucleic acid molecules of the present invention. One such nucleic acid encodes the amino acid sequence of SEQ ID NO: 44, and/or the nucleic acid complement thereof. In another such embodiment, the nucleic acid molecule comprises the nucleotide sequence of SEQ ID NO: 42, and/or the complement thereof. In still another such embodiment, the isolated nucleic acid molecule comprises a nucleotide sequence that has greater than 99.5% identity to that of SEQ ID NO: 42, and/or to the complement thereof.

In still another embodiment, the recombinant and/or isolated nucleic acid molecule encodes the amino acid sequence of SEQ ID NO: 45, and/or the nucleic acid complement thereof. In yet another embodiment, the nucleic acid molecule comprises the nucleotide sequence of SEQ ID NO: 43, and/or the complement thereof. In still another such embodiment, the isolated nucleic acid molecule comprises a nucleotide sequence that has greater than 95% identity to that of SEQ ID NO: 43, and/or to the complement thereof.

The present invention further provides expression vectors that can comprise any of the isolated nucleic acid molecules of the present invention. In one such embodiment the expression vector comprises the isolated nucleic acid molecule encoding the amino acid sequence of SEQ ID NO: 44, and/or the complement thereof. In yet another embodiment, the invention provides an expression vector that comprises an isolated nucleic acid molecule encoding the amino acid sequence of SEQ ID NO: 45, or the complement thereof.

The present invention further provides immunogenic compositions and vaccines that can comprise any of the expression vectors of the present invention. In one such embodiment the immunogenic composition or vaccine comprises an expression vector that comprises the isolated nucleic acid molecule encoding the amino acid sequence of SEQ ID NO: 44, and/or the complement thereof. In another embodiment the immunogenic composition or vaccine comprises an expression vector that comprises the isolated nucleic acid molecule encoding the amino acid sequence of SEQ ID NO: 43, and/or the complement thereof.

The present invention further provides all of the isolated and/or recombinant proteins, polypeptides, peptides, fusion

proteins and chimeric proteins of the present invention. In one such embodiment the polypeptide comprises the amino acid sequence of SEQ ID NO: 44. In another such embodiment that polypeptide is encoded by the nucleotide sequence of SEQ ID NO: 42.

In a further embodiment, the present invention provides immunogenic compositions and vaccines comprising CDV virions that encode a hemagglutinin protein. In related embodiments, the hemagglutinin can be partially encoded by a nucleic acid that comprises a nucleotide sequence as set forth in SEQ ID NOS: 1-33. In still other related embodiments, the present invention provides methods of eliciting an immune response to canine distemper virus in a subject in need thereof by administering to the subject one or more of such immunogenic compositions or vaccines.

The present invention further provides a method of selecting one or more isolates of a pathogen for use in immunogenic compositions, wherein the isolate(s) utilize(s) one or more of a most frequently used codon to encode a selected immunogenic protein, polypeptide or peptide of interest. One such method comprises the steps of 1) determining, for each isolate in a plurality of pathogen isolates, a nucleotide sequence encoding said selected immunogenic protein, polypeptide or peptide of interest; 2) for nucleotide sequences obtained in the determining step, obtaining codon usage data for one or more amino acid residues of interest in said immunogenic protein, polypeptide or peptide of interest, whereby data for frequency of codon usage is obtained; 3) identifying, from said data for frequency of codon usage, a most frequently used codon for each of said amino acid residues of interest in the immunogenic protein, polypeptide or peptide of interest; and 4) selecting, from among the plurality of pathogen isolates, one or more isolates that utilize(s) one or more of the most frequently used codons to encode the protein, polypeptide or peptide of interest. In one embodiment of the invention, the pathogen is a canine distemper virus.

In yet another embodiment, the invention provides a method of selecting one or more nucleotide sequences for a nucleic acid (which may be from an isolate of a pathogen) for use in immunogenic compositions. The nucleic acid utilizes one or more of a most frequently used codon to encode a selected immunogenic protein, polypeptide or peptide of interest. The method comprises the steps of 1) determining (e.g. from a plurality of pathogen isolates) a plurality of nucleotide sequences which encode the selected immunogenic protein, polypeptide or peptide of interest; 2) for nucleotide sequences obtained in the determining step, obtaining codon usage data for one or more amino acid residues of interest in said immunogenic protein, polypeptide or peptide of interest, whereby data for frequency of codon usage is obtained; 3) identifying, from the data for frequency of codon usage, a most frequently used codon for each of the amino acid residues of interest in the immunogenic protein, polypeptide or peptide of interest; and 4) selecting, from among the plurality of nucleotide sequences, the nucleotide sequence(s) that utilize(s) one or more of the most frequently used codons to encode the protein, polypeptide or peptide of interest.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1. Isolate 07091030 (SEQ ID NO: 1, nucleotides (nt) 438-1302 of H gene of CDV). This isolate is from a dog with a history of seizures and exhibiting neutrophils with multiple, intra-cytoplasmic inclusions. This CDV isolate formed large, multi-nucleated, syncytia in a Vero cell line expressing canine signaling lymphocyte-activation molecule (Vero+SLAM).

Based on the H protein sequence, this is a European wild life lineage isolate. This CDV isolate may be particularly useful, for example, as a challenge virus.

FIG. 2. Isolate 07091031 (SEQ ID NO: 2, nt 440-1494 of H gene).

FIG. 3. Isolate 07091032 (SEQ ID NO: 3, nt 440-1225 of H gene).

FIG. 4. Isolate 07101508 (SEQ ID NO: 4, nt 427-1335 of H gene). This isolate has high identity with European wildlife (EW) genetic lineage but is from a dog from Southern California with a history of Ondersteport vaccination. This isolate produces large syncytia in Vero+SLAM cells. The isolate has high identity with CDV from the lesser panda and Danish mink but is not closely related to the Ondersteport vaccine isolate.

FIG. 5. Isolate 07100609 (SEQ ID NO: 5, nt 833-1213 of N gene). The sequence of the nucleocapsid encoding genes of this CDV-like virus from a marine mammal (seal), matches canine isolate 164071 from the US (EU716337).

FIG. 6. Isolate 07110098 (SEQ ID NO: 6, nt 439-1286 of H gene).

FIG. 7. Isolate 07111080 (SEQ ID NO: 7, nt 630-1501 of H gene).

FIG. 8. Isolate 08010939 (SEQ ID NO: 8, nt 443-1343 of H gene).

FIG. 9. Isolate 08011277A (SEQ ID NO: 9, nt 423-1556 of H gene).

FIG. 10. Isolate 08011277B (SEQ ID NO: 10, nt 1530-410 of H gene).

FIG. 11. Isolate 08011277C (SEQ ID NO: 11, nt 433-1230 of H gene).

FIG. 12. Isolate 08011277D (SEQ ID NO: 12, nt 833-1578 of nucleocapsid gene). The H gene of this isolate could not be amplified. Lack of amplification indicates genetic variation (and hence, lack of homology) in the primer binding sequences. The nucleocapsid gene sequence matches that of a CDV isolate.

FIG. 13. Isolate 08011671 (SEQ ID NO: 13, nt 422-1589 of H gene).

FIG. 14. Isolate 08021509 (SEQ ID NO: 14, nt 441-686 of H gene).

FIG. 15. Isolate 08030074 (SEQ ID NO: 15, nt 447-1282 of H gene).

FIG. 16. Isolate 08030776 (SEQ ID NO: 16, nt 422-1541 of H gene).

FIG. 17. Isolate 08030777 (SEQ ID NO: 17, nt 411-1537 of H gene).

FIG. 18. Isolate 08031346 (SEQ ID NO: 18, nt 436-1059 of H gene).

FIG. 19. Isolate 08040383 (SEQ ID NO: 19, nt 1486-620 of H gene).

FIG. 20. Isolate 08050180A (SEQ ID NO: 20, nt 418-1552 of H gene).

FIG. 21. Isolate 08060351 (SEQ ID NO: 21, nt 412-1536 of H gene).

FIG. 22. Isolate 08060352 (SEQ ID NO: 22, nt 408-1553 of H gene).

FIG. 23. Isolate 08080696 (SEQ ID NO: 23, nt 423-726 of H gene).

FIG. 24. Isolate 08080941 (SEQ ID NO: 24, nt 387-1522 of H gene).

FIG. 25. Isolate 08081112 (SEQ ID NO: 25, nt 411-1207 of H gene).

FIG. 26. Isolate 08120827 (SEQ ID NO: 26, nt 413-1535 of H gene).

FIG. 27. Isolate 08120857 (SEQ ID NO: 27, nt 724-1522 of H gene).

FIG. 28. Isolate 09011024 (SEQ ID NO: 28, nt 578-1613 of H gene).

FIG. 29. Isolate 09020504-3 (SEQ ID NO: 29, nt 418-1549 of H gene).

FIG. 30. Isolate 09041289 (SEQ ID NO: 30, nt 889-1646 of H gene).

FIG. 31. Isolate 09041303 (SEQ ID NO: 31, nt 394-1526 of H gene).

FIG. 32. Isolate 09041474A (SEQ ID NO: 32, nt 410-1539 of H gene). Fully vaccinated dogs (two vaccinations with commercial Ondersteport CDV vaccine) in a large shelter in Tennessee developed upper respiratory tract disease, high fevers, green nasal discharge, cough, and eventually neurological symptoms, e.g. twitching. About 20 out of 55 dogs died, including the 4 month old female from which isolates 09041474A and 09041474B (see FIG. 37) were obtained.

FIG. 33. Isolate 09040826 (SEQ ID NO: 33, nt 418-1546 of H gene).

FIGS. 34A-F. Codon table showing canine distemper virus hemagglutinin (H) codon sequences from field isolates aligned with CDV-H vaccine and reference strain codon sequences using BioEdit program. Sequences: Ondersteport=CDV vaccine sequence; AY964110=reference European wildlife (EW) strain; AF112189=reference American-2 (AM-2) strain; AY962112=reference Arctic (Ar) strain. Differences in codons are shaded.

FIG. 35. Phylogenetic tree showing genetic relatedness of the many recent United States CDV isolates in the United States and GenBank reference sequences (underlined). The phylogenetic tree was constructed using MEGA4.1 program (available free of charge at the website located at megasoftware.net; Tamura K, Dudley J, Nei M & Kumar S (2007) MEGA4: Molecular Evolutionary Genetics Analysis (MEGA) software version 4.0. Molecular Biology and Evolution 24: 1596-1599).

FIG. 36. Blood film showing erythrocyte inclusions of CDV from a CDV infected dog (OADDL:07091030). CDV inclusions (arrows) are visible within a neutrophil and a lymphocyte. (Aqueous Romanowsky stain; bar=10 microns).

FIG. 37. Complete sequence of H gene from 09041474B (SEQ ID NO: 42), including stop codons.

FIG. 38. Complete sequence of H gene from 08021509 (SEQ ID NO: 43), including stop codons.

FIG. 39. Complete amino acid sequence of H protein from 09041474B (SEQ ID NO: 44).

FIG. 40. Complete amino acid sequence of H protein from 08021509 (SEQ ID NO: 45).

DETAILED DESCRIPTION

The present invention provides compositions that elicit an immunogenic response to CDV, vaccine compositions designed to provide protection against CDV infection, and CDV diagnostics. The compositions contain newly discovered CDV variants. The new variants reflect the evolutionary trends of CDV, and provide an indication of the predominant CDV strains currently circulating in the United States. The variants were, in part, isolated from dogs that had already been vaccinated for CDV, but which nevertheless contracted CDV and became ill, i.e. the dogs were the victims of vaccine failure. In order to stop or curtail the spread of CDV and to prevent vaccine failure, one or more of these new variants should be incorporated into new vaccine protocols. The invention also provides new diagnostic methods for detecting the new CDV isolates, and for differentiating between canine distemper caused by vaccine administration and canine distemper caused by an emerging virus which was not included

in the vaccine, and against which the vaccine did not provide protection. In addition, the invention provides a method of analyzing RNA virus field isolates and emerging pathogens in order to determine which isolates are likely to be useful for inclusion in broad-spectrum immunogenic and vaccine compositions.

Several of the new isolates belong to the European wildlife CDV lineage; others belong to the American-2 (AM-2) and Arctic genetic lineages. 34 CDV viruses have been isolated, propagated in cell culture and the H genes of the viruses have been partially or fully sequenced. These viruses were from various states in the US (e.g. Oklahoma, Florida, Georgia, California, Missouri, Texas, Kansas and Tennessee). FIGS. 1-33 show partial sequences of cDNA complementary to the H gene from the isolates; FIGS. 37 and 38 show the complete H gene sequence for isolates 09041474B and 08021509, respectively, including stop sequences (TGA for both); and FIGS. 39 and 40 show the corresponding amino acid sequences of the H protein from isolates 09041474B and 08021509, respectively. In the amino acid sequences, the amino acid as position 5 in the sequence from 09041474B is glutamine, and amino acid at position 5 in the sequence from 08021509 is arginine.

The present invention also provides recombinant and/or isolated nucleic acids that encode any of the H-proteins of the present invention, but which do not include the signal sequence (the first 12 amino acid residues at the amino terminus of the H protein) and/or the stop codon and/or comprise an alternative signal sequence and/or an alternative stop codon. Expression vectors comprising such nucleic acids are also included in the present invention, as are the expressed recombinant polypeptides (including isolated recombinant polypeptides) that these nucleic acids encode.

FIGS. 34A-F show CDV hemagglutinin (H) sequences from several of the field isolates aligned with CDV-H vaccine sequences and reference strains. An analysis of this data served as the basis for the development of the Relative Preferred Codon Usage (RPCU) method/concept of analysis and selection of broad spectrum pathogen isolates for use in vaccines. The method is based on the newly recognized patterns of RPCU in pathogens, as disclosed herein. While the method is widely applicable to many pathogenic organisms (discussed in detail below), herein, an exemplary use for the analysis of RNA virus isolates is described. As used herein, an "isolate" (e.g. which may be a pathogen, such as, for example an RNA virus) has been substantially isolated and purified from a biological sample, and may have been subjected to passage and/or propagation in a suitable cell culture. Alternatively, the isolate may have been obtained from a pure culture, e.g. from an ATCC deposit.

In order to practice the method of RPCU, a representative number of isolates of interest are isolated from biological samples of animal subjects (including humans) obtained in one or more populations of interest. The isolates can be obtained within a given geographical region or area of interest, and/or from animal subjects suspected of harboring the pathogen. Suitable biological samples are those with the highest concentrations of the virus collected during the acute stage of pathogen infection, and include various tissue samples, bodily fluids or excreted substances (phlegm, saliva, blood, urine, stool, swabs [a generic term for many types of samples], etc.). Examples of suitable pathogens (e.g. emerging pathogens), which can be analyzed by RPCU include but are not limited to: various viruses such as RNA viruses, single-stranded DNA viruses, influenza viruses, HIV virus, etc.; various bacterial pathogens such as *Mycobacteria*, *Yersinia*, *Rickettsia* and *Bartonella* species; various fungi; and

various protozoan pathogens such as Malaria, *Trypanosoma*, *Toxoplasma*, *Entamoeba*, *Giardia*, and *Cryptosporidia* species, etc. Those of skill in the art will recognize that a "representative number" of isolates of a pathogen may vary, but will generally be in the range of at least 20-25, usually at least 30-35, and may be any number of isolates (or sequences) without limit, depending on the availability of biological samples, the availability of resources that can be directed to the effort, etc.

A "population of interest" will generally be a population of individuals that are susceptible to a pathogen of interest, and may include individuals that exhibit disease symptoms when infected with the virus, or may be "carriers" who harbor the virus with few or no symptoms, but which are nevertheless infected with the pathogen.

A geographical area or region of interest may be, for example, a region or area bounded by geographical and/or legal boundaries, for example, a country, continent, state, county, etc.; or regions separated by geographical barriers such as bodies of water (rivers, lakes, oceans, etc.), mountain ranges, deserts, etc.; or regions/areas with a common climate or weather pattern, e.g. similar average temperature or rainfall, presence or absence of snow and ice, etc.

At least one protein, polypeptide or peptide of interest common to all isolates is selected for detailed gene sequence analysis. Generally, such a protein, polypeptide or peptide of interest is one that is known to be immunogenic. By immunogenic, it is meant that the protein, polypeptide or peptide elicits an immune response (e.g. the production of antibodies) in a host when the protein, polypeptide or peptide is present in the host (e.g. when a pathogen comprising the protein, polypeptide or peptide infects a host). Major immunogens can be determined by any of several methods known to those of skill in the art, including by Western blot analysis of serum from a convalescent patient that has recovered and is protected from further infection with the pathogen. Those of skill in the art will recognize that one or more than one such proteins, polypeptides or peptides may be analyzed using RPCU, but at least one is selected. Otherwise, more can be systematically selected for analysis but the one that is predominant based on, for example, Western blot analysis can be a good starting point. An unknown immunogenic protein can be sequenced by MALDI-TOFF. This helps to design the primers to amplify the sequences.

After selection of a suitable protein, polypeptide or peptide, the nucleotide sequence encoding the protein, polypeptide or peptide is determined using techniques that are well established, e.g. polymerase chain reaction (PCR) sequencing, etc. Alternatively, the sequences that are compared using this method may be obtained directly from biological samples without isolation of the virus, or they may be known sequences obtained from a database (e.g. GenBank or others).

The nucleotide sequences are then subjected to analysis to identify triplet codons and to align the codons in the correct translation (reading) frame. Sequence analysis may be conducted with any of the many nucleotide analysis programs, including but not limited to CLUSTAL W analysis using BioEdit software, the Multiple Sequence Alignment Program (MAP) provided by the Baylor School of Medicine, etc. Usually, the sequences that are analyzed are cDNA sequences, although the method is not limited to the use of cDNA, e.g. DNA, RNA, etc. may also be analyzed. Optionally, the corresponding sequence from one or more reference sequences (e.g. RNA virus reference strains) is included in the analysis. Generally, a reference strain will represent, for example, a pathogen type (e.g. a virus lineage) which was previously dominant (e.g. present at a high frequency) in the population

and/or region under consideration. Such a reference strain may be, for example, an RNA virus that has been used in a vaccine against infection with the RNA virus. The preferred candidates for reference sequences are those which display the highest levels of homology and identity with both nucleotide and protein sequences of the pathogens (e.g. emerging pathogens), when analyzed using, for example, BLASTn and BLASTp programs. The level of homology and/or identity will be at least about 90% or 95%, or even at least about 99% or greater.

The triplet codons specifying the amino acids from the isolates (and, optionally, of one or more reference sequences) are aligned in frame in a format that may be readily compared, including but not limited to in tabular form, for example, in an Excel table. Those of skill in the art are well aware that the triplet code is redundant, and that more than one codon can encode the same amino acid. For each position corresponding to an amino acid residue in the protein, polypeptide or peptide of interest, the identity of the three nucleotides encoding the residue from each isolate is noted and compared across all isolates. For example, in a hypothetical protein of interest, if position 50 is a leucine, possible codons for this residue include tta and ttg. The actual codon at position 50 of all isolates is noted and compared to the codon present at all other isolates. From this comparison, the Relative Preferred Codon Usage (RPCU) can be determined. For example, if 75% of the isolates use "tta" to encode the Leu residue and 25% of the isolates use "ttg", then the RPCU value of tta is 75% and that of ttg is only 25%. Thus, tta is the preferred (i.e. the most frequently occurring or used) codon at that residue. In this manner, the most frequently used or preferred codon for each residue of interest of the sequence of interest is determined. The analysis may be carried out for all residues of a sequence, or for only a subset of residues, e.g. residues that are known to be involved in crucial pathogen activities or which are known to be part of an epitope or antigenic region, or which associated with virulence, etc.

The goal of the RPCU method is to identify isolates with nucleotide sequences which possess a high percentage of preferred codons for use in broad-spectrum vaccine preparations. This can be accomplished by any of several means. For example, a theoretical "ideal" sequence comprising only the most preferred codons can be determined and the actual sequences of the isolates can be compared to the theoretical sequence. The level of homology between each isolate and the ideal sequence is calculated. The isolate that displays the highest level (e.g. amount, percentage, etc.) of homology to the ideal sequence will be the isolate that utilizes the highest number of preferred codons. This isolate is the best "fit" to the ideal, and is selected as a vaccine component. Alternatively, an ideal sequence may not be determined but codon usage at each position is tabulated or calculated as described above, and a comparison is made among sequences by other methods that will occur to those of skill in the art, e.g. by simple visual inspection, to identify a sequence from an isolate that utilizes a very high level, or the highest level, number of preferred codons.

Without being bound by theory, this selection is consistent with the understanding that when a mutation occurs in the three-nucleotide sequence that encodes a residue, that mutation is likely to be perpetuated or to become widespread only if pathogens (e.g. RNA viruses) containing the mutation have some selective advantage over pathogens which do not contain the mutation. For example, some codons are translated more rapidly or with greater accuracy than others, and pathogens with such mutations may reproduce and infect new hosts more successfully than non-mutant pathogens. Therefore,

these codons are eventually present more frequently in a population of pathogens due to natural selection. An isolate with a high percentage of frequently used codons likely possesses the cumulative advantages associated with the codons, and is likely to display the favorable characteristics of afforded by the codons, and thus, when included in a vaccine preparation, is likely to provide broad-spectrum protection. RPCU takes into consideration the internal protein epitopes of pathogens (such as viruses) that interact with the genomic nucleic acids (Pepin K M, J Domsic, and R McKenna. 2008. Genomic evolution in a virus under specific selection for host recognition. *Infection, genetics, and evolution*: 825-834).

RPCU impacts biological functions of RNA viruses such as CDV. The codons (triplet of nucleotides) are the most basic biological unit because they encode amino acids, which form the functional units of protein, including epitopes (e.g. about 6-7 amino-acids) and antigenic regions, or sequence motifs, which are directly involved in eliciting a host immune response to antigenic proteins, such as the H protein of CDV. Thus, the selection of CDV isolates based on RPCU reflects a phenotype/function of the virus, such as gene expression, H-protein expression and titers of the virus. Codon usage can also affect the breadth of protein expression and hence influence the tissues in which the virus or a protein is expressed. In a preliminary analysis, a gel-based PCR analysis of H protein expression by CDV showed that most American-2 isolates had uniformly lower (about 8-10 fold lower) H-gene PCR product expression, compared to most EW isolates. This result likely reflects the robust and biologically favorable H-gene codon composition in most EW CDV isolates, which leads to a higher frequency of EW CDV isolates in canine populations. An application of this method to CDV RNA viruses is presented in Example 4 below.

Those of skill in the art will recognize that computer implemented software (a computer program) may be developed to implement the RPCU method. Such software includes or encodes instructions for causing a computer to carry out the RPCU method, and may include, for example, means for entering sequences and other relevant data (e.g. name of isolate, codon alignment features, etc.), means for displaying entered data, means for representing the results of the analysis (e.g. a display on a screen, or a printout ["hard copy"] of the results in a suitable form, e.g. as a sequence, as one or more numerical indicators (such as "sequence #4" or "#4" as the best result), in graphical form, tabular form, etc.). Means for statistical analyses may also be included in the computer program, and the analysis may provide gradations of results, i.e. the program may rank the candidate sequences in terms of those that are likely to be the most suitable to those that are the least likely, and/or may simply provide one or more highest ranking (most suitable) sequences. The computer program may include instructions for carrying out an algorithm that is used to carry out the analysis.

The RPFU method is thus a method of selecting and/or obtaining an isolate of a pathogen for use in immunogenic compositions and vaccines. Such an isolate can be selected or otherwise obtained (e.g., through modification by standard genetic engineering techniques), which has preferred codons that encode a protein, polypeptide or peptide (usually an immunogen) of interest.

The RPFU method may include steps of: obtaining a plurality of isolates of the pathogen (e.g. an RNA virus) from biological samples from a plurality of different animals infected with the pathogen; selecting an immunogenic protein, polypeptide or peptide of interest associated with the RNA virus; determining a nucleotide sequence encoding the immunogenic protein, polypeptide or peptide of interest for

each isolate of the plurality of isolates; identifying, in each of the nucleotide sequences encoding the immunogenic protein, polypeptide or peptide of interest, codons encoding amino acid residues of interest in the immunogenic protein, polypeptide or peptide of interest; determining, by comparing the nucleotide sequences encoding the immunogenic protein, polypeptide or peptide of interest, frequency of codon usage data for each of the amino acid residues of interest in the immunogenic protein, polypeptide or peptide of interest; from said frequency of codon usage data, identifying a most frequently used codon for each of the amino acid residues of interest in the immunogenic protein, polypeptide or peptide of interest; and selecting, from among the plurality of isolates of the pathogen, an isolate that utilizes one or more of the most frequently used codons to encode the protein, polypeptide or peptide of interest. In some RNA viruses, quasi-populations of the immunogen may be analyzed by RPCU software.

FIG. 35 shows a phylogenetic tree of the genetic relatedness of the isolates, based on the sequence of the H gene. In the tree, a very close related population of CDV isolates has emerged, a "predominant CDV population" that belongs to the European wild-life lineage. These European wild-life viruses can be checked by challenge with other minor CDV viruses of two other lineages, Arctic and American, i.e. the "minor CDV population". One or more broadly reactive, predominant isolates that are protective against both the European-Wildlife and one or the other (or both) of Arctic and American-2 lineage viruses can be used to make a CDV immunogenic composition for use as a vaccine that is effective against all CDV lineages currently circulating in the United States, or other suitable locations. In other words, the immunogenic compositions should elicit an immune reaction (e.g. antibody production) against European wild-life lineage viruses and against one or both of Arctic and American-2 lineage viruses. Vaccines of the invention should be protective against European wild-life lineage viruses and against one or both of Arctic and American-2 lineage viruses. The CDV used in such an immunogenic composition or vaccine contains nucleic acid sequences encoding antigens (antigenic regions, antigenic determinants, etc.) previously found only in and believed to be characteristic of European wild-life viruses, together with antigens previously found only in and believed to be characteristic of Arctic or American lineage viruses, or preferably both Arctic and American-2 lineage viruses.

One example of such a virus is isolate 09041474B, the complete hemagglutinin gene sequence of which is set forth in SEQ ID NO: 42. Several criteria were used for selection of this CDV isolate as a vaccine candidate. First, a panel of current CDV isolates was developed. (Historically, only a few isolates have been available based on published reports. Moreover, the reports to investigate the issue of CDV vaccine failure have been few.) Second, hemagglutinin sequencing and CDV genotyping were performed. Global nucleotide analysis using BALSTn, CLUSTAL-W, and phylogenetic analysis allowed clustering and characterization of CDV isolates in lineages, and 09041474B was determined to be of EW lineage. Then, codon usage tables (shown in FIGS. 34A-F) were used to select CDV vaccine isolate 09041474B according to the Relative Preferred Codon Usage (RPCU) analysis method described above.

The H gene sequence from isolate 09041474B (complete sequence, FIG. 37), differs from reference EW sequence (AY964110 in FIGS. 34A-F) in several respects. Firstly, the nucleotide sequence for the H gene of AY964110 was obtained using a tissue sample, i.e. a CDV virus with this

sequence was not isolated. In contrast, the H gene sequence from 09041474B as described herein was obtained from an isolated CDV that had been grown and propagated in cell culture. The two sequences also differ in codon usage. For 09041474B, the codon at position 187 is CGA, the codon at position 201 is CTG, the codon at position 236 is CCT and the codon at position 303 is TCA. In further contrast to AY964110, in the 09041474B sequence, the codon at position 303 encodes serine rather than leucine (see FIGS. 34A-F). Isolate 09041474B displays robust growth in cell culture and it is possible that serine at position 303 confers advantages with respect to eliciting an immune response in subjects to whom virions with this sequence are administered. A deposit of the 09041474B CDV isolate, labeled 09041474B CDV-EW (whole, live viruses at a low passage of 2-3) was made at the American Type Culture Collection (ATCC) in Manassas, Va., with deposition #PTA-10596, deposit date Jan. 21, 2010. A second CDV isolate of interest, 08021509 (American-2, labeled 08021509 CDV-AM-2) was also deposited at ATCC with deposition #PTA-10597, deposit date Jan. 21, 2010. Both deposits were of whole viruses. The invention includes viruses that have the characteristics of the isolates that were deposited, for example: nucleotide sequences as disclosed herein; virulence and propagation attributes; attributes of syncytia (e.g. size, shape, number, appearance (e.g. clearly demarcated, fuzzy, etc.)), number of nuclei in the syncytia, etc.; among others.

In some embodiments of the invention, a multivalent CDV immunogenic composition and/or vaccine (e.g. European wildlife and American-2) will be employed. These two CDV lineages can be combined in a single preparation or administered separately as two separate dosages, for example, if they interfere with induction of CDV immunity.

The present invention provides all of the isolates disclosed herein, as well as vaccines and immunogenic compositions made from the isolates, and/or from antigenic portions of the isolates as described herein, e.g. nucleic acids comprising the nucleotide sequences as set forth in SEQ ID NOS: 1-33 and 42-43 and/or the proteins, polypeptides, or peptides encoded therefrom. The vaccines of the invention may be formulated in any suitable manner, including but not limited to using the whole virus (e.g. killed or attenuated, as described in detail below). In this embodiment, any of the novel CD viruses disclosed herein may be used to prepare a vaccine. Generally, such viruses may be identified by isolating the virus from tissue samples from dogs with symptoms of CDV infection, especially dogs that have been previously vaccinated against CDV, and sequencing and comparing the viral genome to known sequences (i.e. compared to CDVs isolated prior to the present invention, especially to CDV isolates that are currently used in vaccines). In particular, such new virus isolates may have an H gene sequence that contains a region that is identical to or homologous to that of isolate 09041474B (an exemplary isolate).

Generally, such viruses will have an H gene (or portion thereof) that is at least about 75%, preferably about 80%, more preferably about 85%, most preferably about 90%, or even 95, 96, 97, 98, 99, or 100% homologous (and/or identical) to the nucleic acid sequences disclosed herein, or complements thereof. In one particular embodiment of the present invention, the CDV isolate comprises an H-gene with a nucleotide sequence, comprising greater than 99% homology (and/or identity) with SEQ ID NO: 42 (i.e., that of isolate 09041474B). In a related embodiment, the CDV isolate comprises an H-gene with a nucleotide sequence comprising greater than 99.5% homology (and/or identity) with that of SEQ ID NO: 42 (i.e., that of isolate 09041474B).

In yet another embodiment, the CDV isolate comprises an H-gene comprising a nucleotide sequence with greater than 95% homology (and/or identity) with SEQ ID NO: 43 (i.e., that of isolate 08021509). In a related embodiment, the CDV isolate comprises an H-gene comprising a nucleotide sequence with greater than 99% homology (and/or identity) with that of SEQ ID NO: 43 (i.e., that of isolate 08021509).

Alternatively, such viruses may encode H proteins containing amino acid sequences that are at least about 75%, preferably about 80%, more preferably about 85%, most preferably about 90%, or even 95, 96, 97, 98, 99, or 100% identical to the amino acid sequence encoded by the nucleic acid sequence disclosed herein. Those of skill in the art are familiar with methods to calculate % homology or % identity. Such variant viruses may have H gene coding sequences that differ from those disclosed herein because of natural variations among isolates, or due to changes that are introduced deliberately e.g. by genetic engineering techniques. In other words, the viruses may be recombinant.

In other embodiments, only antigenic portions of the viruses described herein are present in the immunogenic or vaccine compositions of the invention. Such compositions may be formulated using, for example, nucleic acids encoding antigenic peptides or proteins as presented in SEQ ID NOS: 1-33 and 42-43, or antigenic epitopes or regions of peptides or proteins, from those sequences. For example, the vaccine preparations of the invention may comprise nucleic acid sequences that include the sequences set forth herein, complements thereof, and/or proteins, polypeptides or peptides encoded by such sequences. In addition, vaccines with certain variations of such sequences are also encompassed. While the sequences represent cDNA, the invention also includes corresponding ssRNA, ssDNA, double-strand (ds) DNA, dsRNA, complementary DNA, and RNA of any form (e.g. mRNA, RNA/DNA hybrids, etc.) that is based on, derived from or that complements these sequences. Such sequences may be either sense or antisense sequences. Further, sequences which display at least about 90% homology, or even about 95, 96, 97, 98 or 99% or greater homology to nucleic acid sequences of the CDVs disclosed herein, are also contemplated for use in the vaccines. Such sequences may differ, for example, by containing alternate codons that encode the same amino acid at one or more positions in order to maximize expression. In addition, portions of these sequences which encode epitopes or antigenic regions of e.g. the H protein are also contemplated, as are sequences which display 70%, or even more preferably about 80, 90, or 95% or even greater identity (e.g. 96, 97, 98 or 99% identity) to such amino acid sequences. Generally, about 6-8 amino acids constitute an epitope. Such sequences may vary, for example, by containing conservative or non-conservative amino acid substitutions, or deletions (especially amino or carboxy terminal deletions), or various insertions, etc., so long as the resulting protein/peptide is antigenic as described herein. Such antigenic regions are preferably at least about 10 amino acids in length, but may be much longer, e.g. encompassing an entire protein such as the H protein.

Further, nucleic acid sequences which hybridize to sequences disclosed herein (or to portions of those sequences) under stringent conditions (especially conditions of high stringency) are also contemplated. Stringent conditions refer to hybridization conditions which allow a nucleic acid sequence to hybridize to a particular sequence. In general, high stringent conditions refer to the hybridization conditions which allow a nucleic acid sequence of at least 50 nucleotides and preferably about 200 or more nucleotides to hybridize to a particular sequence at about 65° C. in a solution

comprising about 1 M salt, preferably 6×SSC or any other solution having a comparable ionic strength, and washing at 65° C. in a solution comprising about 0.1 M salt, or less, preferably 0.2×SSC or any other solution having a comparable ionic strength. These conditions allow the detection of sequences having about 90% or more sequence identity.

Nucleic acids encompassed by the present invention, e.g. those with nucleotide sequences set forth in SEQ ID NOS: 1-33 and 42-43 (and variants thereof as described herein) may be obtained in various ways. For example, they may be obtained from natural sources such as from a viral isolate; alternatively, they may be produced synthetically. Those of skill in the art will understand that the capability exists in the art to synthetically produce very large sequences, e.g. entire viral or bacterial genomes (e.g. *Mycoplasma*), and the present invention encompasses sequences of any origin or manufacture that comprise the sequences disclosed herein, as well as the proteins, polypeptides and/or peptides expressed from the sequences.

The invention also provides recombinant constructs such as recombinant viruses, vectors, and expression vectors which express the proteins/polypeptides/peptides described herein (i.e. the amino acid sequences encoded by the nucleic acid sequence set forth in SEQ ID NOS: 1-33 and 42-43, or variants thereof). Such constructs include those which have been produced, for example, by cloning one or more of the sequences disclosed herein into a vector or host (e.g. plasmids, cosmids, viral vectors such as adenoviral and poxviral vectors, or bacterial vectors, etc.).

In one embodiment, the construct is an expression vector that includes the previously noted nucleic acids and/or fragments thereof. Recombinant expression vectors used in this invention are typically self-replicating DNA or RNA constructs comprising nucleic acids encoding a CDV hemagglutinin of the present invention and/or an antigenic fragment thereof, usually operably linked to suitable genetic control elements that are capable of regulating expression of the nucleic acids in compatible host cells. Genetic control elements may include a prokaryotic promoter system or a eukaryotic promoter expression control system, and typically include a transcriptional promoter, an optional operator to control the onset of transcription, transcription enhancers to elevate the level of mRNA expression, a sequence that encodes a suitable ribosome binding site, and a sequence that terminates transcription and translation. Expression vectors may also contain an origin of replication that allows the vector to replicate independently of the host cell. Recombinant expression vectors may be constructed by any of several means known to those of skill in the art. For example, genetic engineering techniques are known by which sequences of interest are removed e.g. from an isolate of origin such as a virus and ligated into a suitable expression vector. Alternatively, portions of an expression vector or an entire expression vector may be made synthetically; or a combination of ligation and synthesis protocols may be employed.

In addition, other useful elements may be included in the constructs described herein. For example, the constructs may encode various sequences such as histidine tags or other tags that are used to facilitate protein isolation, such as glutathione-S transferase (GST), and maltose binding protein; various linker or spacer sequences; various adjuvants and sequences that increase the antigenicity of the protein (e.g. haptens); sequences which introduce a desired/convenient restriction enzyme cleavage site or which encode a desired protease cleavage site; sequences encoding fluorescent or other detectable labels, or tagging or marking sequences (e.g. Green Fluorescent Protein (GFP), or portions thereof); vari-

ous sequences that direct the location, export or processing of the encoded protein (e.g. leader sequences); heterologous signal sequences (i.e. signal sequences not normally associated with CDV H protein in nature); etc. Other possibilities will occur to those of skill in the art and are also intended to be encompassed by the present invention. When such sequences are included in the constructs, if they are contiguous with the viral sequences described herein, the entire coding sequence may be translated as a fusion or chimeric protein/polypeptide/peptide, and may or may not (depending on the sequence) be susceptible to post-translational modification. The expressed recombinant proteins/polypeptides/peptides of the present invention and their corresponding fusion or chimeric proteins/polypeptides/peptides are also provided by the present invention. In particular embodiments such recombinant proteins/polypeptides/peptides and their corresponding fusion or chimeric proteins/polypeptides/peptides are also isolated.

In addition, the present invention provides host cells that comprise such expression vectors. The host cell is optionally a prokaryote or a eukaryote host cell. Expression of nucleic acids encoding a CDV hemagglutinin of the present invention can be carried out by conventional methods in either prokaryotic or eukaryotic cells.

The vaccines and immunogenic compositions of the invention may comprise any of the sources of the sequences described herein, e.g. a virus isolate, an attenuated virus, a recombinant construct, etc. Several methods of making vaccines suitable for vaccination against CDV are known in the art. See, for example, U.S. Pat. Nos. 4,193,990 and 4,193,991 to Appel et al., U.S. Pat. No. 4,303,645 to Carmichael et al., U.S. Pat. No. 4,971,793 to Wood et al.; U.S. Pat. No. 5,882,652 to Valdes et al., and U.S. Pat. Nos. 5,885,585 and 5,814,510 to Parrish et al., each of which offers variations of suitable vaccine-formulating strategies. The complete contents of each of these patents are hereby incorporated by reference. Generally, to manufacture a vaccine, a viral or other vector containing genetic sequences of the invention (either naturally, or due to genetic engineering) is employed. Examples of such viral vectors include viruses and virions (e.g. CDV) that are "killed", inactivated or otherwise attenuated so as to not cause severe disease symptoms in the animal to which it is administered, together with a suitable physiological carrier. The CDV virus can be inactivated (rendered unable to replicate) using chemicals such as formalin, binary ethylene amine, beta propriolactone, by using gamma irradiation or heat, or by other methods known in the art. Attenuation may be carried out, e.g. by repeated passage of the viral isolate in suitable host cells, and subsequent isolation of the resulting clonal isolate. In some embodiments, the attenuated virus retains the ability to replicate within the host, although this is not strictly necessary. Preferably, no disease symptoms will occur as a result of administration. However, those of skill in the art will recognize that many effective vaccine compositions cause some discomfort or relatively minor distress upon or after administration. However, the benefits of being protected against full-blown disease far outweigh this possibility. The attenuated virus may be a virus that naturally contains the nucleic acid sequence(s) of the invention (e.g. a CDV), or the virus may be recombinant in that the nucleic acid sequence is inserted into the virus by genetic engineering. In the case of recombinant vaccines, the nucleic acid sequences may be incorporated into viruses other than CDV to form heterotypic recombinant vaccines. Examples of such viruses include but are not limited to various herpesviruses, adenoviruses, poxviruses, non-pathogenic "orphan viruses", enteric viruses such as enterovirus, and others well known in the art. In

addition, expression of the H gene could be accomplished in bacterial, yeast or parasite recombinant systems. In a preferred embodiment, the virus is a live, attenuated (modified) high titer CDV, and the nucleic acid is ssRNA. In addition, other forms of the vaccine are also contemplated. For example, "empty" virion particle vaccines (without nucleic acid) are also contemplated, as are vaccines comprising antigenic virion or other CDV proteins that are not assembled into a capsid. In addition, the vaccines of the invention may be multivalent and include multiple viruses. Alternatively, a single virus genetically engineered to contain nucleic acids encoding proteins from two or more of the novel CDVs can be constructed by recombinant technology by exchanging coding regions, as is known by those of skill in the art.

The CDV that is used in the compositions described herein is generally attenuated and safe, i.e. produces no or few symptoms of disease when administered to a suitable host animal. A CDV vaccine should not elicit antibody production in cerebrospinal fluid of a host. However, administration of the attenuated CDV still results in an immune response (e.g. a protective immune response) to CDV immunogens such as the H protein. The most frequently used method for producing an attenuated live-virus vaccine is to serially passage the virus in cell culture. For example, the virus may be passaged in a primary canine cell culture or canine cell line that does not harbor an oncogene, although other cell lines may also be used (e.g. chick embryo or fibroblast, VERO-SLAM cells, baby hamster kidney cell lines, as well as other hamster cell lines (Sultan S, N T Lan, T Ueda, R Yamaguchi, K Maeda, and K Kai. 2009. Propagation of Asian isolates of canine distemper virus (CDV) in hamster cell lines. *Acta Veterinaria Scandinavica* 51:38 doi: 10.1186/1751-0147-51-38), etc. Typically, for the first passage, a cell culture is infected with the selected inoculum of CDV. After obtaining clear evidence of virus replication (for example, virus-induced cytopathic effects [CPE] in the infected cells), an aliquot of the cell culture medium, or infected cells, or both, of the first passage are used to infect a second cell culture. The process is repeated until one or more mutations in the viral genome cause sufficient attenuation so that the virus can be safely used as a vaccine. The number of passages may vary somewhat e.g. at least about 20 and usually about 50 passages are used, but as many as e.g. 150 passages may be used. By then, the virus is sufficiently attenuated (i.e., reduced in virulence or diseases-producing ability) to be used in a vaccine formulation. The degree of attenuation is usually determined empirically by exposing the natural host to progressively greater passage levels of the virus.

It is also possible to attenuate the CDV viruses by repeat passages at decreasing incubation temperatures with or without mutagenic chemicals. Normally, CDV viruses are propagated at 37° C. However, over e.g. 50 passages at successively decreasing incubation temperatures for example, clonal strains of the virus are produced which no longer have the ability to replicate at core body temperature (37° C.) or above. Such viruses retain the ability to multiply in areas of the body that typically exhibit lower temperature, e.g. the nasal cavity, but do not replicate at the core body temperature. For example, in one embodiment, a cold-adapted, temperature CDV propagates in tissue culture cells at temperatures from about 26° C. to about 34° C., but does not do so at a non-permissive temperature of about 37° C. (US patent application 2006121521, Dowling and Younger, the complete contents of which is hereby incorporated by reference). These viruses are therefore completely safe for use in CDV vaccines for animals, including wildlife and highly susceptible species such as large cats, mink and ferrets.

Other suitable vaccine components, e.g. pharmacologically acceptable carriers, are well-known to those of skill in the art, as is the preparation of such compositions for use as vaccines. Typically, such compositions are prepared either as liquid solutions or suspensions, however solid forms such as tablets, pills, powders and the like are also contemplated. Solid forms suitable for solution in, or suspension in, liquids prior to administration may also be prepared. The preparation may also be emulsified. The active ingredients may be mixed with excipients which are pharmaceutically acceptable and compatible with the active ingredients. Suitable excipients are, for example, water, saline, dextrose, glycerol, and the like, or combinations thereof. In addition, the composition may contain minor amounts of auxiliary substances such as wetting or emulsifying agents, pH buffering agents, and the like. If it is desired to administer an oral form of the composition, various thickeners, flavorings, diluents, emulsifiers, dispersing aids or binders and the like may be added. The composition of the present invention may contain any such additional ingredients so as to provide the composition in a form suitable for administration. The final amount of the translatable nucleic acid in the formulations may vary. However, in general, the amount will be from about 1-99%. The compositions may further comprise an adjuvant, suitable examples of which include but are not limited to Seppic, Quil A, Alhydrogel, oil-in-water emulsions, aluminum phosphate, carbolol, Emulsigen, and the like.

The immunogenic/vaccine preparations of the present invention may be administered by any of many suitable means which are well known to those of skill in the art, including but not limited to by injection, orally, intranasally, intratracheal, by ingestion of a food product containing the antigen, by intramuscular, subcutaneous, intravenous, transdermal, and intradermal routes, by eyedrops, or with a nebulizer or a needle-free instrument, etc. However, in a preferred embodiment, the mode of administration is by injection. In addition, the compositions may be administered alone or in combination with other medicaments or immunogenic compositions, e.g. as part of a multi-component vaccine. In particular, the immunogenic CDV could be combined with rabies virus, *Borrelia burgdorferi*, *Ehrlichia canis*, canine parvovirus, canine adenovirus, canine parainfluenza virus, canine coronavirus, *Babesia canis*, *Anaplasma phagocytophilum*, *Giardia* species, *Leishmania* species, *Leptospira* species or any combination thereof. etc. Further, administration may be a single event, or multiple booster doses (of the same or a different strain) may be administered at various timed intervals to augment the immune response. In addition, administration may be prophylactic, i.e. before exposure to the virus has occurred, or is suspected to have occurred, or after the fact, i.e. after a known or suspected exposure, or therapeutically, e.g. after the occurrence of disease symptoms associated with viral infection.

The invention also provides various types of recombinant vectors and/or expression vectors that contain and express the nucleic acid sequences disclosed herein (or portions thereof that encode antigenic peptides and/or polypeptides). Examples of such vectors and expression systems include but are not limited to: various bacterial (e.g. *Escherichia coli*) or probiotic-based (e.g. *Lactobacillus*) expression vectors; adenoviral vectors, baculovirus, *Pichia*, and other yeast expression systems; pox vectors such as raccoon pox vectors; etc. Such recombinant vectors and expression systems may also be utilized in vaccine preparations. Alternatively, they may be employed for other purposes such as for laboratory manipulation of the sequences, or for research or diagnostic purposes.

The invention provides methods of immunizing or preventing the symptoms of CDV infection in a subject (e.g. a mammal) in need thereof by administering to the subject a composition of the invention. Generally, the CDV vaccines are administered in an amount sufficient to provide active immunity in puppies and/or adult dogs. Preferably, the immune response is protective against future exposure to CDV, i.e. administration of the composition prevents the symptoms of disease associated with CDV infection, when compared to non-vaccinated controls. However, much benefit may also accrue if the immune response simply lessens or decreases the severity of disease symptoms, even if all symptoms are not eliminated.

In a preferred embodiment of the invention, the animals that are vaccinated using the vaccines of the invention are domestic dogs, including both adult dogs and puppies. However, the vaccination of other potential CDV hosts is also contemplated. Other potential hosts include other canids such as wild canids (e.g. wolves, wild dog species, etc.), larger species of cats (whether domesticated or wild), mink, red panda, foxes, lion and tigers, ferrets, rabbits, goats, etc. as well as other carnivores in general. Ferrets are highly susceptible to CDV. Thus a highly attenuated, modified live virus vaccine or recombinant CDV vaccine can be used. According to the American Ferret Association, MD, three CDV vaccines can be administered to healthy kits at 8, 11, and 14 weeks of age. While the vaccines will of course be used in domestic animals, wild or partially domesticated animals may also benefit from such vaccination, e.g. animals in zoos or protected areas, parks, in research facilities, etc. Wildlife can in particular be vaccinated with killed CDV vaccine because they are more susceptible to modified live virus vaccines, e.g. by use of edible bait which contains vaccine components. Any animal that can host the CDV variants, whether or not the virus causes disease symptoms in the host, may benefit by being vaccinated by the vaccine preparations provided herein. Vaccination of animals that are asymptomatic upon infection by the virus (i.e. silent carriers) would be beneficial in order to curtail the spread of the virus to more susceptible populations.

The invention also provides antibodies that bind specifically or selectively to antigenic determinants or antigenic regions of the CDV disclosed herein. In some embodiments, the antibodies are neutralizing antibodies that can neutralize the virus and thus prevent infection. Such differential antibodies may be polyclonal or monoclonal, although monoclonal antibodies are generally preferred. The antibodies may be of canine origin. Monoclonal antibodies will be prepared by injecting the viruses (e.g. killed viruses, proteins, or nucleic acids encoding the proteins) in mice or another suitable host such as rabbit or canine host. After 3 boosters, the spleens will be harvested and fused with myeloma cells. The monoclonal antibodies producing clones will be screened by ELISA, HA-HI, and indirect fluorescent antibody test. The clones that react with the viruses described herein, or with proteins isolated from the same, will be saved for development of CDV diagnostic assays. Polyclonal antibodies may be prepared by injecting one or more peptides that span amino acid codons that are preferred antigenic targets e.g. the H protein, into rabbits.

The invention also provides diagnostic methods and kits for the detection of the CDV variants described herein. Such kits include, for example, oligonucleotide primers specific for amplifying (e.g. by polymerase chain reaction, PCR) the nucleic acid sequences disclosed herein. Alternatively, such kits may include antibodies (e.g. monoclonal or polyclonal) that bind selectively or specifically to unique antigenic determinants displayed by the novel CDV variants. The kits are

useful in monitoring the CDV status of, for example, any animal that is susceptible to CDV, especially canines. The kits are especially useful to monitor the CDV status of puppies and dogs that are exported or transported from one jurisdiction to another. In one embodiment, the diagnostic tests and methods of the invention are used to detect the presence of CDV in dogs (or other animals) that have been fully vaccinated but have nevertheless developed symptoms of CDV infection. Using the methods of the invention, it is possible to determine the genotype of the etiological agent of disease, and to ascertain whether the disease symptoms are caused by the vaccine strain, or by superinfection with a genetic variant that was not neutralized by vaccination, i.e. the vaccine did not provide protection against the genetic variant.

The invention is further illustrated in the following Examples, which should not be construed so as to limit the invention in any way.

EXAMPLES

Example 1

Preliminary Studies of Seven CDV Isolates

Canine distemper virus (CDV) is a highly contagious virus that causes multi-systemic disease in dogs. Seven cases of CDV in dogs from the USA were received. These CDV isolates formed large, multi-nucleated, syncytia in a Vero cell line expressing canine signaling lymphocyte-activation molecule (SLAM) (described below). Based on the hemagglutinin gene sequences, the CDV isolates from 3 states (CA, MO, and OK) formed two CDV genetic groups: Group I (major, 6/7) consisted of CDV isolates closely related to the European wildlife lineage of CDV. The group II (minor, 1/7) was genetically related to the Arctic-like lineage of CDV. However, both the CDV groups were genetically different from the current vaccine strains that belong to American-I lineage of the old (1930-1950) CDV isolates.

In this study, an evolutionary and genetic analysis of 7 CDV isolates from the United States was performed using the H gene sequences. The biological effects of the 1-1 gene sequence variation were investigated using an in vitro cell culture system. Ante-mortem samples included ocular swabs, nasal swabs, and peripheral blood anticoagulated with EDTA. The swabs were received in 1 to 2 ml of cold normal saline sent on ice by overnight delivery within 24 h of collection. Urine samples were not tested. Post-mortem samples were from tonsils, brains, bladders, and lungs (Kubo, T., Y. Kagawa, H. Taniyama, and A. Hasegawa. 2007. Distribution of inclusion bodies in tissues from 100 dogs infected with canine distemper virus. *J. Vet. Med. Sci.* 69:527-529). Approximately 2 to 5 g of each tissue was received in tubes sent on ice by overnight delivery for virological examination. The specimens were obtained from seven suspected cases of CD from three states in the United States (Oklahoma, four; Missouri, one; and California, two).

For direct fluorescent antibody testing, tissues were sectioned at 8- μ m thickness and fixed with an acetone (75%)-methanol (25%) mixture at room temperature. Veterinary Medical Research and Development (VMRD), Pullman, Wash., USA supplied pretitrated, lot-to-lot certified conjugates for veterinary diagnostic applications. As part of quality control/quality assurance, the conjugates were tested before use on negative and known positive CDV controls. After addition of ready-to-use, prediluted, fluorescein isothiocyanate-labeled, anti-CDV monoclonal antibody (VMRD, Pullman, Wash.) or polyclonal antibody conjugates (VMRD,

Pullman, Wash.), the sections were incubated for 30 min at 37° C. After the unbound antibody conjugates were washed, the sections were counterstained with Evans blue for 15 min. After being mounted in buffered glycerol (pH 9.4), the sections were examined by fluorescent microscopy. Positive cells showed apple-green fluorescence in the cytoplasm and negative cells were brick-red.

For isolation, the tissues from CDV-infected samples were finely chopped, freeze-thawed twice to release the virus, and centrifuged at 8,000 \times g. The clear supernatant was filtered through a 0.22 μ m syringe filter. The Vero cell line was derived from the kidney of a normal, adult African green monkey (*Ceropithecus*) in Japan. The recombinant cell line was derived by transfection of the Vero cells with canine signaling lymphocyte activation molecule (SLAM, also known as CD150) as described before by Seki et al. (Seki, F., N. Ono, R. Yamaguchi, and Y. Yanagi. 2003. Efficient isolation of wild strains of canine distemper virus in Vero cells expressing canine SLAM (CD 150) and their adaptability to marmoset B95a cells. *J. Virol.* 77:9943-9950). The inoculums (about 1 ml per 25-cm² flask) were incubated for 1 h at 37° C. with rocking every 20 minutes. After inoculation on a recombinant Vero cell line expressing canine SLAM, about 3.5 ml of Dulbecco's modified of Eagle's medium (Cellgro, Hendron, Va.) with 5% fetal calf serum was added. The cells were examined daily for cytopathic effects (multinucleated-syncytium formation) (Seki, supra). Vero cells expressing canine SLAM have been found to be useful for the primary isolation of CDV (Lan, N. T., R. Yamaguchi, K. Uchida, S. Sugano, and S. Tateyama. 2005. Growth profiles of recent canine distemper isolates on Vero cells expressing canine signaling lymphocyte activation molecule (SLAM). *J. Comp. Path.* 133:77-81).

For total RNA extraction (host and viral RNAs) from specimens, QIAmp viral RNA extraction kits were used (Qiagen Inc., CA). The quality and quantity of the RNA were checked by A₂₆₀/A₂₈₀ using a Nonodrop spectrophotometer (Nonodrop Technologies, CA).

For detection of CDV RNA, reverse transcriptase (RT)-PCR based on the nucleocapsid (N) gene was targeted (Kim, Y. H., K. W. Cho, H. Y. Youn, H. S. Yoo, and H. R. Han. 2001. Detection of canine distemper virus (CDV) through one step RT-PCR combined with nested PCR. *J. Vet. Sci.* 2:59-63). This protocol provides high sensitivity due to the nested amplification of the target gene, high copy number of the N gene, and the conserved sequence of the N-gene among CDV isolates. Briefly, the first-round product was amplified by the forward primer (Primer 1: 5'-ATTGGGATTGCTTAGGA-3', SEQ ID NO: 34) and reverse primer (Primer 2: 5'-GGCGCTCATCTTGGACAT-3', SEQ ID NO: 35). The protocol was reverse transcription at 45° C. for 1 hour, 95° C. for 3 min; 30 cycles of PCR with denaturation at 94° C. for 30 s, annealing at 54° C. for 30 s, and an extension at 72° C. for 1 min; and a final extension at 72° C. for 7 min, with the reaction mixture held at 4° C. The small-portion (1-microliter) product of the first reaction was subjected to a second round of amplification using primer 3 (5'-GT-TAGCTAGTTTCATCCT-3', SEQ ID NO: 36) and primer 4 (5'-GGTCCTCTGTTGTCTTGG-3', SEQ ID NO: 37). The protocol for the second round was denaturation at 95° C. for 3 min; 30 cycles of denaturation at 94° C. for 30 s and annealing at 54° C. for 30 s; with an extension at 72° C. for 1 min. The final extension was performed at 72° C. for 7 min, and the reaction mixture was held at 4° C. before electrophoresis. The size of the second-round amplicon was 419 base-pairs, verified by including molecular size standards in agarose gel analysis.

For CDV genotyping, the H gene was used as the target (Martella, V., G. Elia, M. S. Lucente, N. Decaro, E. Lorusso, K. Banyai, M. Blixenkroner-Moller, N. T. Lan, R. Yamaguchi, F. Cirone, L. E. Carmichael, and C. Buonavoglia. 2007. Canine distemper virus (CDV) by hemi-nested multiplex PCR provides a rapid approach for investigation of CDV outbreaks. *Vet. Microbiol.* 122:32-42). The forward primer (primer 204+, nucleotides 388 to 409, 5'-GAATTCGACT-TCCGCGATCTCC-3', SEQ ID NO: 38) and reverse primer (primer 232b-, nucleotides 1543 to 1519, 5'-TAGGCAA-CACCACTAATTTRGACTC-3', SEQ ID NO: 39) yield an amplicon of 1160 base-pairs. The H-gene RT-PCR protocol was RT at 50° C. for 30 min and 94° C. for 2 min. The PCR protocol was 35 cycles of 94° C. for 1 min, 50° C. for 1 min, and 72° C. for 3 min and a final extension at 72° C. for 10 min, with the reaction mixture held at 4° C. The positive and negative CDV controls were included in each run of both detection (N gene) and genotyping (H gene) RT-PCR protocols. For phylogenetic analysis of the H gene sequences, the amplicons were sequenced at the Oklahoma Medical Research Foundation, Oklahoma City, Okla. The sequences were subjected to Basic Local Alignment Search Tool for Nucleotides (BLASTN) analysis (Altschul, S. F., T. L. Madden, A. A. Schaffer, J. Zhang, Z. Zhang, W. Miller, and D. J. Lipman. 1997. Gapped BLAST and PSI-BLAST: a new generation of protein database search programs. *Nuc. Acid Res.* 25:3389-3402) and compared to GenBank H gene sequences for CDV isolates from different species and geographic areas around the world. The percentage identities of the H gene sequences were recorded. Further, the H gene sequences were

phocytes, consistent with CDV inclusions (FIG. 36). The presence of CDV inclusions was confirmed by the direct fluorescent-antibody test in both cases.

Six of the seven CDV positive samples were successfully isolated in the Vero cell line with canine SLAM/CD150. The cytopathic effects of CDV isolates were characterized by multinucleated syncytia that formed 1 to 2 days after inoculation. The presence of CDV was further detected by RT-PCR for the hemagglutinin gene. One CDV sample (OADDL 07061535) was tested only by RT-PCR and sequencing of the H gene; there was insufficient sample for virus isolation.

Based on RT-PCR for the H gene followed by sequencing, the level of identity among the CDV isolates (OK-1, OK-2, OK-3, OK-4, CA-1 and CA-2; major group I) was highest with a canine CDV isolate 19876 from Missouri, that is genetically most related to a Danish mink CDV isolate (Pardo, I. D. R., G. C. Johnson, and S. B. Kleiboeker. 2005. Phylogenetic characterization of canine distemper viruses detected in naturally infected dogs in North America. *J. Clin. Microbiol.* 43:5009-5017). Thus, it was the predominant CDV variant (six of seven isolates) in this study. These six CDV isolates belonged to the European wildlife lineage of CDV isolates. However, one isolate (MO-1; minor group II) was found that was most genetically similar to the canine CDV isolate 21260 from Missouri (Pardo, supra) that is closely related to a lesser-panda CDV isolate. This CDV isolate belongs to the Arctic-like lineage of CDV isolates. The information on the 2007 OADDL CDV isolates is summarized in Table 1, and partial nucleotide sequences of the H gene of these isolates are provided in FIGS. 1-7

TABLE 1

OADDL Canine Distemper Virus (CDV) Isolates								
OADDL No./ Designation	State of Origin	Vaccination Status ¹	Age (weeks)	Breed	% H Gene Identity			
					Homology to Current Vaccine Virus ²	Homology to MO 19876 ³	Virus Isolation	CDV Lineage ⁵
07061535	OK-1	I	12	Mixed	89	98	nd ⁶	EW
07091030	OK-1	NV	44	Siberian husky	89	98	yes	EW
07091031	OK-1	NV	na	na	89	99	yes	EW
07091032	OK-1	NV	na	na	84	96	yes	EW
07101508	CA-1	V	10	American bulldog	89	98	yes	EW
07110098	MO-1	V	10	Weimaraner	89	90	yes	A
07111080	CA-2	V	136 ⁴	Border collie	97	96	yes	EW

¹I - vaccination incomplete; NV = not vaccinated; V = vaccinated

²All isolates had less than 90% identity with the current vaccine isolates (Ondersteport, Lederle and Convac)

³All isolates had less than 90% identity with the USA MO 19876 CDV isolate except MO-1, which belongs to the Arctic lineage

⁴Animal recovered completely after supportive therapy

⁵EW = European wildlife; A = Arctic

⁶nd = not done.

subjected to phylogenetic analysis and sequence comparison with H gene sequences of the vaccine CDV isolates (Ondersteport, Convac, Lederle, and Snyder Hill CDV isolates) deposited in GenBank. Alignments of the top 100 matches with known sequences were used to perform phylogenetic analysis by neighbor-joining using Jukes-Cantor method (NCBI, MD).

Peripheral blood films from two of the CD case samples (OADDL 07091030 and OADDL 07091031) were stained with an aqueous Romanowsky stain and examined by light microscopy. Both blood films revealed numerous eosinophilic structures within the cytoplasm of neutrophils and lym-

The hemagglutinin glycoprotein varies approximately 10% among the CDV isolates and envelope protein H determines the cytopathology and tropism of the virus (von Messling, V. G. Zimmer, G. Herrler, L. Haas, and R. Cattaneo. 2001. In a preliminary analysis, the OADDL CDV isolates were compared with all the H sequences in the GenBank and it was found that CDV isolates cluster in geographically distinct lineages. For example, all the Argentine CDV isolates formed one distinct cluster. The South American CDV isolates were not included in the recent analysis of CDV isolates based on geography and H gene phylogeny (McCarthy, supra). However, they form a distinct South American cluster.

In recent papers, the terms genotype, cluster, and lineage have been used interchangeably by different investigators, but the results on CDV phylogeny were similar in all the studies (Martella, V., F. Cirone, G. Elia, E. Lorusso, N. Decaro, M. Campolo, C. Desario, M. S. Lucente, A. L. Bellacicco, M. Blixenkrone-Moller, L. E. Carmichael, and C. Buonavoglia. 2006. Heterogeneity within the hemagglutinin genes of canine distemper virus (CDV) strains in Italy. *Vet. Microbiol.* 116:301-309; McCarthy, supra; Mochizuki, M., M. Hashimoto, S. Hagiwara, Y. Yoshida, and S. Ishiguro. 1999. Genotypes of canine distemper virus determined by analysis of the hemagglutinin genes of recent isolates from dogs in Japan. *J. Clin. Microbiol.* 37:2936-2942), including this analysis, because all the investigators used the GenBank accession sequences. A member of a particular genotype of CDV has been proposed to have a more than 95% identity in the nucleotides of the H gene sequences (Mochizuki, supra) and, thus, the intragenotypic variation is less than 5% (Martella, supra).

The CDV isolate OK-1 (OADDL 07061535) was obtained from a 3-month-old, female, mixed breed, vaccinated dog from Oklahoma with history of conjunctivitis, nasal discharge and weight loss. The dog had not finished the complete course of vaccination, and had a history of roaming and eating garbage. This CDV isolate had maximum identity (98%) with CDV isolate 19876 (GenBank accession number AY964110.1). Based on the H gene analysis, CDV isolate 19876 belongs to the European wildlife lineage of CDV isolates along with OK-1.

The CDV isolate OK-2 (OADDL 07091030) was obtained from a tissue pool of an 11-month-old unvaccinated Siberian husky from Oklahoma. On necropsy, the conjunctival and tracheal epithelium contained intracytoplasmic, eosinophilic inclusions surrounded by clear halos. In the tonsils, there were marked lymphoid depletion and numerous inclusion bodies in the epithelium. This isolate had maximum identity (98%) with CDV isolate 19876 (canine origin, Missouri), and 94% identity with CDV isolates from Hungary (GenBank accession number EF095750.1), a Danish mink (Z47759.1), and a lesser panda (AF178039.1), CDV strain A75/17 (AF164967.1), and morbillivirus from a German ferret isolate (X84999.1). The CDV isolate A75/17 from the United States is regarded as a virulent prototype of field CDV isolates (Simon-Martinez, supra). The level of identity of the H gene with the vaccine isolates (Convac, Lederle, and Ondersteport) was 89%.

The CDV isolate OK-3 (OADDL 07091031) was obtained from a dog in a shelter in Oklahoma. A blood tube was obtained but no other history was available on this case. Inclusions consistent with CDV were observed in leukocytes on a peripheral blood film and further confirmed by direct fluorescent-antibody test. The blood sample was positive for CDV by virus isolation. The H gene was sequenced and had 99% identity with CDV canine isolate 19876 (GenBank accession number AY964110.1), and 95% identity with CDV isolates from Hungary (EF095750.1), a Danish mink (Z47759.1), and a lesser panda (AF178039.1), CDV virus strain A75/17; CDV isolate 01-2641 and a German ferret morbillivirus strain (X84999.1).

The CDV isolate OK-4 (OADDL 07091032) was obtained from a tissue pool (bladder and lungs) from a dog adopted from an animal shelter in Oklahoma. This CDV isolate had maximum identity (96%) with CDV isolate 19876 (GenBank accession number AY964110.1). In descending order, it had 93% identity with CDV isolates from Hungary (EF095750.1), a lesser panda (AF178039.1), and a Danish mink (Z47759.1); 84% identity with the vaccine isolates; and 70% identity with the phocine distemper virus.

The CDV isolate CA-1 (OADDL 07101508) was obtained from a tissue pool from a 10-week-old male vaccinated American bull dog from California that died of CD. Three out of 4 littermates died of CD with respiratory signs, hyperkeratosis, and seizures. Of the three dead littermates, the necropsy report was available for one littermate. Its lungs were firm and congested on necropsy. The necropsy results of one of the four littermates were completely normal. The H gene sequence was 98% identical to a canine origin CDV isolate 19876 (GenBank accession number AY964110.1). The CDV isolate was 94% identical to the Hungarian CDV isolate, the lesser panda isolate (AF178039.1), and CDV strain A75/17 (AF164967). The CDV H gene sequence was 93% identical to CDV isolate 01-2641 (AY526496.1). The H gene of this CDV isolate lacked the Pst I site present in all vaccine CDV isolates (Demeter, Z., B. Lakatos, E. A. Palade, T. Kozma, P. Forgach, and M. Rusvai. 2007. Genetic diversity of Hungarian canine distemper virus strains. *Vet. Microbiol.* 122:258-269).

The CDV isolate MO-1 (OADDL 07110098) was obtained from nasal and conjunctival swabs of a 10-week-old, CDV vaccinated Weimaraner dog that had clinical signs compatible with CD. The dog developed 'chewing-gum' seizures, thickened footpads, coughing, nasal discharge, and congested lungs. The swabs were collected before euthanasia, and CDV was isolated in cell culture. The CDV isolate H gene had maximum identity (98%) with CDV isolates 21261 and 18133 from Missouri, and 97% identity with CDV isolates from Italy (48/05 and 179/94) and Hungary (H06Bp10S, H06 Bp8F, H05 Bp7F, H05 Bp6F, and H05 BpBp5F). The H gene sequence of this CDV isolate had 95% identity with a CDV isolate from a Greenlandic dog and only 90% identity with CDV 19876. It had 89% identity with the vaccine CDV isolates and 70% identity with the phocine distemper virus H gene. Moreover, this CDV isolate lacks the Pst I restriction site present in all vaccine CDV isolates (Demeter, supra). Based on phylogenetic analysis this isolate belongs to the Arctic-like lineage of the CDV isolates.

The CDV isolate CA-2 (OADDL 07111080) was obtained from a combination of nasal, pharyngeal, tonsil, and conjunctival swabs of a 32-month-old neutered male, vaccinated Border collie with a history of vomiting, diarrhea and lymphopenia. The H gene sequence had maximum identity (96%) with CDV isolate 19876. This isolate had 93% identity with the Hungarian CDV isolate, the lesser Panda CDV isolate, and the Danish mink CDV isolate; 92% identity with CDV strain A75/17 (GenBank accession number AF164967.1); and 92% identity with the German ferret CDV isolate. Based on phylogenetic analysis, this CDV isolate clusters with CDV isolates of the European wildlife lineage. This dog recovered after treatment and has been clinically healthy for the last 3 months. The survival of this dog after a natural exposure to a CDV isolate of European wildlife lineage is probably due to resistance based on age, genetic resistance, and immunity after complete vaccination with a commercial CDV vaccine. This dog had a CDV titer of 1:16 by CDV serum neutralization 3 months after recovery from CDV infection.

Five out of six OADDL 2007 CDV isolates were found to produce multinucleated, syncytia in a Vero cell line expressing the canine SLAM receptor. It has been proposed that syncytial size is a correlate of the degree of virulence of the CDV isolates (Cosby, S. L., C. Lyons, S. P. Fitzgerald, S. J. Martin, S. Pressdee, I. V. Allen. 1981. *J. Gen. Virol.* 52:345-353) because it correlates with the ability of the CDV to spread from cell-to-cell. The aggressive spread in cell culture, the ability to produce large numbers of inclusions in canine lymphocytes that naturally express SLAM/CD150, and the

ability to produce fatal infections in vaccinated dogs indicate that these canine isolates of European wildlife lineage are virulent for dogs.

This Example shows that the EW lineage is emerging as the predominant CDV isolate in the US.

Example 2

Additional CDV Isolates

Using the methods described in Example 1, additional CDV isolates were identified and are listed in Table 2.

TABLE 2

OADDL Canine Distemper Virus (CDV) Isolates							
OADDL No./ Designation	State of Origin ¹	Clinical Sign ²	CPE ³	Vaccination Status ⁴	Age (weeks)	Breed	CDV Lineage ⁵
8010939	OK	R, N	na	na	20	Miniature Schnauzer	EW
08011277-A	OK	R	+	na	12	Small breed	EW
08011277-B	OK	R	+	na	12	Small breed	EW
08011277-C	OK	R	+	na	12	Small breed	EW
8011671	GA	R	na	V	10	Mix	EW
8021509	FL	R, N	+	V	12	Mix	AM-2
8030674	CA	R, N	na	V	8	Golden Retriever Mix	EW
8030776	OK	R, N	+	V	16	Mix	EW
8030777	FL	N	+	V	12	Mix	AM-2
8031346	CA	na	na	V	12	Pitbull	EW
8040383	MO	R	+/-	na	6	Weimaraner	AR
8050180A	OK	R	na	na	14	Pitbull Mix	AM-2
8060351	MO	R, N	na	V	9	Shih Tzu	AM-2
8060352	MO	R	na	V	8	Welsh Terrier	AM-2
8080696	FL	R	+	V	24	Mix	EW
8080941	OK	N	+	na	12	Rat Terrier	EW
8081112	MO	N	+/-	V	11	Irish Terrier	AM-2
8120827	OK	N	+	na	0.3	Dachshund	EW
8120857	OK	N	+	na	5	Yorkshire Terrier	EW
9011024	na	N	+	V	156	Akita Mix	EW
09020504-3 (08-75891)	KS	na	+	na	na	na	AM-2
09020504-2 (56928)	KS	na	na	na	na	na	AM-2
09020504 (58829B)	KS	na	+	na	na	Leopard	AM-2
9041303	na	R	+/-	V	24	Cattle Dog Mix	EW
09041474A	TN	R	+	V	16	Border Collie Mix	EW
0904147B	TN	R	+	V	16	Border Collie Mix	EW

¹State of origin; na = not available

²Clinical Signs: R = Respiratory (coughing, ocular and nasal discharge, sneezing); N = nervous (tremors, twitching, prurition change, exterior rigidity)

³CPE = Cytopathic Effect: + = positive; - = negative; +/- = suspect; na = not available

⁴Vaccination Status: V = vaccinated; NV = not vaccinated; na = not available

⁵CDV lineage: EW = European wildlife; Am-2 = American-2

Example 3

Continuing Investigations of Emerging CDV Isolates: Differences in Cytopathology (CPE) Among Recent USA Canine Distemper Viruses

The results obtained in Examples 1 and 2 prompted a continued effort to isolate and characterize additional CDV isolates from the USA. Studies were carried out as described for Example 1. CDV samples were inoculated in Vero+SLAM cell line. Most CDV isolates produced large syncytia with large number of nuclei. A smaller number of CDV isolates produced smaller sized syncytia with few cell nuclei. In several CDV isolates, multiple syncytia appeared after virus

sample inoculation 18-24 hours after inoculation (Fast-growing CDV isolates). In some CDV isolates, the margins of the syncytia were well defined with almost circular margins. In other CDV isolates, the margins of the syncytia were not that clearly demarcated. These “fuzzy” CPE CDV isolates tended to spread rapidly with daughter syncytia next to the mother syncytium. In other CDV isolates, daughter syncytia appeared far away indicating another colony (colonies) of virus growth. In short, CDV isolates showed variable cytopathology. CDV isolates from USA differ in the speed (Fast and Slow), spread (Large and Small), size, shape (Round and Irregular) of the syncytia formation and invasiveness of the

host cells by CDV isolates. These biological properties may have bearing on the protection offered by the current vaccines. For example, codon usage of critical viral genes can affect the replication efficiency of CDV, as discussed in Example 11.

Example 4

Comparative Genetic Analysis of CDV Isolate Sequences: Relative Preferred Codon Usage (RPCU)

Ten hemagglutinin residues (29, 178, 180, 225, 386, 412, 475, 530, 549, and 603) are known to be under positive selection among CDV lineages (McCarthy, supra). The

hemagglutinin (H) gene of the isolates were sequenced or partially sequenced and the resulting sequences are shown in FIGS. 1 to 33. The partial H-gene sequenced (about nucleotide 534-1236 hemagglutinin gene fragment) from the wild type CDV isolates were aligned with reference CDV sequences from GenBank using CLUSTAL W provided with Bio-edit program). Of note, a viral isolate containing the reference sequence for the EW strain has never been isolated, propagated in cell lines, or characterized. The reference EW sequence was obtained by sequencing carried out on tissue extracts.

The results of the RPCU analysis, depicted in FIGS. 34A-F, showed the following:

At residue 180, the codon is AGT in Onderstepoort-like vaccines such as Galaxy, Proguard, Continuum, and Vanguard. However, this codon is GGT in European wildlife (EW), and in all wild type CDV isolates. This codon can be useful in designing a Taqman RT-PCR to distinguish the Onderstepoort-like vaccines from wild type CDV isolates circulating in the USA.

At residue 225, the codon is GAC encoding aspartic acid (D) in all vaccines. However, it is AAC in American-2 CDV and in raccoon distemper virus (RDV) 09050216 it is CAC (encoding histidine, H).

At residue 386, the codon is ACC (threonine, T) in all Onderstepoort-like vaccines (all except the Pfizer vaccine). However, it is TAC in EW, AM-2 and AR lineage CDV wild type viruses and in the Pfizer vaccine. This codon is thus useful in developing a differential Taqman RT-PCR for distinguishing most of the commercial CDV vaccines and wild type CDV isolates.

At residue 412, the codon is CCT, which encodes proline (P) in vaccines and wild type CDV viruses. However, in isolate 07110098 it is CAT (encoding histidine, H) as is also the case for the Arctic-lineage of CDV.

Relative Preferred Codon Usage (RPCU) has been developed from analysis of the codon usage comparisons. One goal of an RPCU analysis is to identify RNA viral isolates (e.g. CDV isolates) suitable for making a broad-spectrum vaccine capable of providing protection against most isolates circulating in a particular geographic area (e.g. the United States). RFCU also allows evaluation of the genetic distance of isolates that are outside an area of interest based on codon usage. RPCU is based in part on the observation that, in addition to amino acid residues, codons themselves are under evolutionary selection pressure (Gustavo et al., Lost in Translation: Codon Usage and HIV-1 Evolution, AIDS Reviews 2004; 6:54-60).

To develop and implement RPCU analysis, a codon usage Table depicted in FIGS. 34A-F was created in which each entry presents the three nucleotides of a single triplet codon. The codon usage Table can also be created using a concatenation approach, e.g. using an Excel program with MEGA4.1 software. To create this Table, first all the sequences were subjected to CLUSTAL W analysis using BioEdit software. Then, triplets of nucleotides in frame with the coding sequence were manually entered into an Excel table. In this manner, residue positions 155-428 of the hemagglutinin protein were analyzed, and the codons for residues of interest were included in the Table. For each residue position, at the bottom of the column, the codons used were retyped in lower case letters and the encoded amino acid was indicated using a single capital letter. The alternative codons that did not lead to change of amino acid (substitutive mutations, S/-) were also noted. For example, residue position 176 uses two codons (tct and tcc, both encoding serine, S). The residues positions for which only the Onderstepoort sequence differs from all other

codons in the Table (i.e. is an outlier) were also noted by an "O" at the bottom of the column) e.g. residue positions 180 and 186. Some residue positions were identical in all CDV isolates. These identical residue positions are not shown in the codon usage Table because they did not affect the selection of the CDV vaccine isolate.

To identify the relatively preferred codon at each residue, the entire Table was examined residue by residue, i.e. column by column. For example, in the column representing codons at residue position 185, either CCA or TCA is used to encode the amino acid at this position. However, CCA is the preferred codon compared to TCA, since CCA is present in the majority of isolates. Similarly, preferred codons were determined at each of the residue positions. For some residues, the least preferred codons were also identified. As can be seen, most of the least preferred codons occur in American-2 and Arctic isolates. This pattern of relative preferred codon usage could be one of the major reasons for the biological advantage of EW over AM-2 and AR CDV isolates. Canine distemper virus is labile in the environment but highly contagious, similar to the measles virus. A CDV isolate in a geographical area that has a replication advantage due to more biologically fit codon usage and higher replication titers and shedding (e.g. in nasal secretions and other portals of delivery in a dog population) has the ability to spread to and affect even vaccinated dogs. Thus, codon usage is the minimum functional unit of virulence factors (such as H protein of CDV) with effects on epidemiological, biological and disease outcomes in host populations.

As noted above, a major goal of the present RPCU analysis is to identify one or more CDV isolates for use in a broad-spectrum CDV vaccine development. Preferably, a vaccine preparation containing or based on such CDV isolates would provide protection against infection by most CDV isolates currently circulating in the US, or at least lessen deleterious symptoms associated with such infection. EW isolate 09041474B was identified as using a preferred codon at most residue positions (e.g. at positions 185, 192, 193, 203, 205, etc.) and was thus selected for further vaccine development.

The approach of RPCU was developed in part because the residues that are critical for the immunogenicity of the CDV H protein have not been determined. RPCU analysis provides a method to identify robust CDV vaccine candidates in the absence of detailed knowledge of the antigenic characteristics of H protein residues. RPCU can be used to analyze and select vaccine candidates from among isolates of other types of RNA viruses as well, and will be useful in cases where the newly emerged virus has not been well studied but an emergency vaccination is called for to stop a growing outbreak.

Further, the new variant AM-2 isolates described herein have higher isolate specific codon usage (as determined by RPCU) and, while this makes them less suitable for a broadly reactive CDV vaccine, these genetically unique isolates will make excellent challenge viruses to check the efficacy of the improved CDV vaccines. RPCU is consistent with self-optimization for new host adaptation being one of the fundamental reasons for evolution of emerging pathogens of animals and humans.

Example 5

Phylogenetic Analysis of Recent CDV Sequences, Reference CDV Sequences from Gen-Bank for Each CDV Genetic Lineage, and all Commercial CDV Vaccines

FIG. 35 shows the phylogenetic analysis of several CDV hemagglutinin partial sequences from recent US CDV

samples compared to commercial CDV vaccines strains and GenBank reference sequences for all known CDV genetic lineages, including South American sequences. Observations that can be made are as follows:

American-1 (AM-1) genetic lineage: reference sequence=AF378705, Onderstepoort strain from 1950's; none of the currently circulating CDV isolates from the US cases were of this CDV lineage. Most of the current commercial CDV vaccines (Continuum DAP, Intervet (n=1 lot); Duramune Max 5, Fort Dodge (n=1 lot); Galaxy DA2PPv, Schering Plough (n=3 lots); Merial (recombinant canary pox vectored CDV-H gene vaccine (n=1 lot) are all based on the AM-1 lineage.

American-2 (AM-2) genetic lineage: reference sequences=AF112189; Z47762; AF259552. Based on the phylogenetic analysis, the 3 reference sequences were dispersed in three locations on the CDV tree shown by the underlined sequences. All but one AM-2 CDV isolate clustered with and around the AF112189 reference sequence. A total of 8 AM-2 CDV isolates were isolated in this study. America-2 is the second largest cluster of CDV circulating in the USA now. An isolate from this lineage should be included in the updated CDV vaccine and has thus been deposited with ATCC (Manassas, Va.).

Arctic (AR) genetic lineage: Two reference sequences (AY964112 and AY964108) constituted this cluster. We identified a few USA samples in this cluster. Both (07110098; 08040383) the samples were from Missouri, USA. This is minor CDV genetic cluster. This CDV lineage will be suitable for some parts of the USA.

Asia-1 (AS-1) genetic Lineage: Three reference sequences (AB016776; AB 212963; AY378091) constituted this cluster. None of the USA CDV isolate is related to this lineage. This type of CDV lineage has been reported in Japan, China, and Korea. CDV vaccines for these Asian countries may include this lineage.

Asia-2 (AS-2) Genetic Lineage: Two reference sequences (AB0470767 and AB 252718) constitute this genetic cluster. None of the USA CDV isolate is related to this lineage. This type of CDV lineage has been reported in Japan, China, and Korea. CDV vaccines for these Asian countries should include this lineage.

European-Wildlife (EW) Genetic Lineage: The reference sequence for this lineage was AY964110. Most of the EW CDV isolates (n=14) from the US clustered around the reference strain of CDV. However, two EW were branched separately. This major cluster of CDV isolates should be included in the updated CDV vaccines for use in USA dogs.

European (E) CDV Lineage: This lineage cluster contained only the two reference sequences (AF478550, DQ494318). Pfizer CDV vaccine branched close to this cluster. South American (SA) CDV Lineage: Four reference CDV sequences clustered in this lineage. These isolates separated as a new branch from the tree indicating that they are unique from USA isolates and current vaccines.

Measles virus Edmonton B sequence was used as an outlier sequence for comparison with US vaccines and current isolates for phylogenetic analysis.

A potential new lineage of CDV has been described in South Africa (Woma, supra).

Example 6

Vaccine Development

Many limitations in the current art of CDV vaccine development had to be overcome to select the isolates for improved

CDV vaccine development. One of the current limitations of CDV vaccines is the availability of CDV isolates from cases of vaccine failure. The problem of CDV vaccine failure has not been fully appreciated and there is not much published data to support these observations. Moreover, current veterinary diagnostic techniques have not been extensively applied to the problem of vaccine failure due to a lack of available methods and the cost of diagnostic testing of ante-mortem samples showing neurological symptoms. Thus, prior to the present invention, the data on current CDV isolates with respect to improving the quality of CDV vaccines was very limited, consisting mainly of isolated reports that were largely overlooked and reports of single dogs. This is undoubtedly because most owners of the deceased dogs do not wish to pay for further medical investigations and simply dispose of the animal carcasses. Therefore, no further scientific information is obtained.

The 34 CDV isolates described herein are compared using hyper-immune serum against the American-1 CDV isolate utilized in the commercially available Onderstepoort canine distemper vaccine. Hyper-immune serum is a useful reagent because it is prepared by administering multiple vaccines to adult dogs that are immunocompetent. Hyper-immune sera offer the best case scenario. However, in a field situation, most dogs will receive two CDV vaccines as puppies. Thus, although hyper-immune serum is a useful reagent, it has limitations for designing and selecting CDV isolates for vaccine preparation. Here, we propose a novel bio-informatics approach (RPCU, described above) for broad-spectrum CDV vaccine development. All available genotypes of CDV currently circulating in the US are included in the analysis. Several (e.g. at least 3-5 isolates of each CDV genotype) are selected (using phylogenetic analysis) for antigenic comparison based on sequence alignment using Bio-Edit. This allows CDV isolates that have the maximum antigenic distance to be selected and compared to the current CDV vaccines. Any isolate that is at least 4-fold lower in SN test using either hyper-immune serum or a serum from a dog that has received only two vaccines is selected for further testing in vivo.

Both sero-negative and low sero-positive puppies (at least 5 puppies in each group) are injected with each of the selected (e.g. 3) lineages of CDV isolates either as a single injection multivalent CDV vaccine or in separate, back-to-back injections of each of the 2-3 different genetic variants 3-4 weeks apart. This experiment identifies one or more broadly reacting CDV isolates that will elicit a higher level of titers against all the 34 CDV isolates. For example, isolates identified by RPCU analysis such as 09041474B are confirmed to be broadly reacting isolates due to shared codons of H protein. This translates into higher vaccine titers, better protection in challenge experiments, no clinical evidence of disease in vaccinated dogs (or alternatively, mild clinical symptoms), and longer duration of immunity. Moreover, CDV isolates selected using the RFCU will provide broad protection against other genetic and antigenic variants that are present in other continents.

Additional criteria are also used to select CDV isolates for use in a modified live virus vaccine, including the following:

A). CDV isolates should grow to high titers (10^6 or more) on an approved non-recombinant cell line such as Vero and canine kidney cell line. It is expected that CDV isolates will grow to higher titer (2-3 log higher titers) in the recombinant Vero+SLAM (Signaling Lymphocyte Activation Molecule) cell line. Although recombinant Vero SLAM is suitable for primary isolation of morbilliviruses, it is expensive to propagate CDV isolates therein due to the required addition of the selection antibiotic, gentamycin. CDV isolates that grow in

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one or more non-recombinant Vero cell lines that are approved by the USDA for animal vaccine production are selected for further propagation.

B). The speed of growth of CDV isolates is another criterion for evaluation. Most CDV isolates can grow to high titers in 3 days. However, the growth is slightly slower in conventional Vero cell line or dog kidney cells.

C). A few CDV isolates (n=5) from the major EW-branch are selected for further evaluation as vaccine antigens. A few isolates (n=5) from the second largest cluster (AM-2) are also selected as vaccine antigens. A few Arctic CDV isolates are also evaluated.

Example 7

Preparation of a Broad-Spectrum CDV Vaccine

A broadly reactive and predominant CDV isolate (e.g. a candidate identified by RPCU) is selected and confirmed by CDV-serum neutralization (CDV-SN) and/or plaque reduction tests. In some embodiments, this is a wild-type CDV isolate that is obtained from a fully vaccinated adult dog (above 5-6 months of age) that died in spite of complete vaccination (specifically 2 CDV vaccines).

Dogs are vaccinated subcutaneously or intranasally with the vaccine. The antibody titer against the vaccine strain is checked by CDV-SN or plaque reduction assays using serum from the vaccinated dog. The selected CDV strain shows high cross-reactivity with a panel of recent CDV isolates from the US belonging to all the CDV lineages circulating in the USA. All isolates are checked for cross-reactivity with sera from the vaccinated dog or ferrets (a laboratory model animal for CDV). A titer of $\geq 1:8$, preferably $\geq 1:16$, more preferably $\geq 1:32$, and most preferably $\geq 1:64$ after one vaccination of a naïve puppy 6-8 weeks of age is sufficient using a CDV-SN assay. In addition, the vaccine will not induce any cerebrospinal fluid (CSF) titers against CDV. Lack of CDV titers in the CSF indicates that the vaccine virus has not crossed the blood-brain barrier and is safe for use in puppies. A vaccine that is safe in puppies is very likely safe in dogs. In summary, all the guidelines of the Code of Federal Regulations (CFR) will be followed to develop an effective broad spectrum and safe CDV vaccine that will be approved by the USDA for use in dogs and other species susceptible to CDV.

Challenge studies are performed in which dogs are vaccinated with the broad spectrum CDV vaccine and then exposed to circulating, wild type CDV (e.g. 08080696 EW; 08081112 AM-2; 09011024 EW). These CDV viruses are genetically distinct and so are suitable as challenge viruses rather than as vaccine components. Dogs vaccinated with the vaccine preparation of the invention develop few or no symptoms of disease. Low-passage CDV isolates from the US as described herein can be used as the challenge virus. Unvaccinated controls will develop symptoms of CDV.

With respect to evaluating the results of challenge studies, diagnostic laboratories typically use relatively insensitive tests that often may not detect weakly CDV positive cases, such as those involving the nervous system. CDV may be detected using immunohistochemistry of brain samples after an animal dies or is sacrificed or euthanized. However, CSF and brain biopsy are expensive and invasive procedures and are not used routinely. In a recent study, urine has been described as a sensitive sample for detection of CDV in live dogs (Amude, A. M., A. A. Alfieri, and A. F. Alfieri. 2006. Antemortem diagnosis of CDV infection by RT-PCR in distemper dogs with neurological deficits without the typical clinical presentation. *Vet. Res. Comm.* 30:679-687). Thus,

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viruria (the presence of CDV virus or RNA in urine) may be an important parameter to include in vaccine-protection evaluation studies, and may be used non-invasively to detect the residual virus in CDV vaccinated dogs. Moreover, after extensive in depth diagnostic investigation, it has been found that CDV can cause residual CDV infections that were not evaluated in past CDV vaccine approval processes. Highly sensitive PCR assays are used and safety data is provided on the improved broad-spectrum CDV vaccines. Viruria is used as one of the parameters of CDV vaccine efficacy.

An ideal candidate CDV vaccine should protect against all genetically diverse CDV isolates. CDV isolates from other continents should be included to check global coverage of a broad-spectrum CDV vaccine. American-1 is not checked because this virus has not been found in the US in nature for the last 20 years, existing only in vaccines that have not been updated for 6 decades.

Example 8

Evaluation of CDV Vaccines in Ferret Models

Prior to testing the vaccines in a large animal model such as dogs, they can be evaluated in a ferret model. Ferrets are known to be suitable models of CDV infection and evaluation of CDV vaccines (Pillet et al., 2009: Ferrets as a model for morbillivirus pathogenesis, complications, and vaccines. *Curr. Top. Microbiol. Immunol.* 330:73-87). Ferrets are used to screen a large number of CDV vaccine candidates. The ferrets are vaccinated with attenuated modified live CDV, preferably a European-wildlife type (e.g. 09041474B). European-wildlife is preferred because a closely related cluster of these CDV viruses is causing vaccine failure in dogs in the US. Ferrets are checked for serum antibody titers against CDV after vaccination. The bleed dates are 0, 7, 14 and 21 days after vaccination. Low passage CDV viruses (e.g. European wildlife, or Arctic, or American-2) are then administered to the ferrets as challenge viruses.

Ferrets also have been documented to show CDV vaccine failure based on a recent case report (Zehnder et al., 2008: An unusual presentation of canine distemper virus infection in a domestic ferret (*Mustela putorius furo*) DOI: 10.1111/j). This domestic ferret, from the US, was repeatedly vaccinated using chick-embryo modified live virus vaccine 18 months prior to the onset of clinical CDV problems and annually thereafter. This vaccinated ferret developed a systemic CDV infection manifesting in skin lesions, with a prolonged course of disease yet with complete absence of respiratory and neurologic signs. Thus, CDV should be suspected in vaccinated ferrets with skin lesions (Zehnder, supra).

Example 9

Critical H-Protein Residues Undergoing Positive Selection Among CDV Lineages: Application to Diagnostics

Depending on the specific needs of diagnostic clients, differential RT-PCR experiments and kits (including primers) are designed around critical H-residues (see Example 3) to differentially detect CDV wild types; to differentiate the Onderstepoort-like vaccines from the Pfizer vaccine; and to differentiate CDV viruses down to the level of major CDV lineages using rapid assays with a 1 hour turnaround time. Ongoing monitoring of CDV viruses by complete H-gene

sequencing further refines these fast CDV differential assays by identifying new CDV variants that arise in the future.

Example 10

Comparative Growth Characteristics of Three Selected CDV Isolates in Cell Culture

From a panel of current CDV isolates at OADDL, two CDV isolates (09041474B and 08021509) were selected for depositing at the American Type Culture Collection (ATCC). The isolate 09041474B has been selected for developing a broad-spectrum CDV vaccine against current CDV isolates.

Three CDV isolates: 09041474B (European-Wildlife); 08021509 (American-2); and 07110098 (Arctic), were propagated in cell culture and observed for their speed of growth based on cytopathology, and the flasks were frozen when most of the monolayer (over 80%) was exhibiting cytopathology. The results showed that the speed of growth of these selected isolates was as follows: EW>>>AM-2>AR. In other words, the EW 09041474B isolate grew significantly faster than AM-2 and AR. The individual plaques of 09041474 were very large. At 23 hours, the entire flask of cells was covered with very large syncytia that touched each other (were fused) leaving almost no space between syncytia.

Isolate 08021509 (AM-2) displayed medium size plaques and grew as isolated plaques (non-fused plaques) initially. The speed of AM-2 isolate growth was at least half or less that of 09041474B. This isolate was harvested at about 96 hours after inoculation.

Arctic isolate 07110098 grew very slowly and the plaque size was small. Only a few isolated small plaques were detected. This isolate was not deposited at ATCC. This isolate was harvested at 7 days post infection. Even at one week, the monolayer showed only about 25% cytopathology. Based on these growth characteristics, this isolate is not suitable for vaccine preparation.

The type of active replication displayed by 09041474B is indicative of optimum (robust) growth that is expected from a CDV isolate that exhibits high "Relative Preferred Codon Usage (RPCU)". This isolate has been chosen for vaccine preparation because it will replicate to higher titers after inoculation and express relatively higher amounts of hemagglutinin protein, the major CDV immunogen. The ATCC deposit number for 09041474B is PTA-10596, deposited Jan. 21, 2010. The ATCC deposit number for 08021509 is PTA-10597, deposited Jan. 21, 2010.

Example 11

Full-Length Hemagglutinin Sequences of Two Exemplary/Selected CDV Isolates Deposited at ATCC, MD

To derive the full-length sequences of the two CDV selected isolates, new primers were designed:

5'-TCGAAATCCTATGTGAGATCACT-3' (forward primer, CDVff1, SEQ IS NO: 40) and 5'-ATGCTGGAGATGGTT-TAATCAATCG-3' (reverse primer, CDVHS-2, SEQ IS NO: 41). The RNA was extracted from the same batch of CDV isolates that were deposited at ATCC on Jan. 21, 2010. A QIAGEN viral RNA extraction kit was used according to the manufacturer's instructions. The primers for the full-length H-protein have been published (Lan N T, Yamaguchi R, Inomata A, Furuya Y, Uchida K, Sugano S, and S Tateyama. 2006. Comparative analyses of canine distemper viral isolates from clinical cases of canine distemper in vaccinated dogs. *Vet. Microbiol.* 115:32-42) but the RT-PCR protocol was not described. Thus, a new protocol was developed based on the properties of the primers.

A one step RT-PCR protocol was as follows: reverse transcription at 45° C. for 1 hour, denature at 95° C. for 3 minutes, followed by 30 cycles of 94° C. for 30 seconds, anneal at 50° C. for 30 seconds, extend at 72° C. for 2 minutes, final extension at 72° C. for 7 minutes, and hold the reaction at 4° C.

The reaction set up was as follows for each PCR reaction: 12.5 ul of 2x reaction buffer (Invitrogen, Cat#10928-034), both primers 1.7 ul each at (15 uM), MgSO₄ (50 mM), dNTPs (10 mM) 0.5 ul, and RT/platinum-Taq (0.5 ul). The PCR amplicons were purified by electrophoresis on 1.5% agarose gel. Correct full length amplicons about 2100 bp were observed. The amplicons were purified on a Promega Wizard column. Sequencing was performed at Noble Research Center, Stillwater, Okla. The forward and reverse sequences of both CDV isolates were subjected to sequence analysis (FIGS. 37 and 38). The CDV isolate 09041474B had the highest match with European-Wildlife CDV isolates. The CDV isolate 08021509 had the highest match with American-2 CDV isolates.

While the invention has been described in terms of its preferred embodiments, those skilled in the art will recognize that the invention can be practiced with modification within the spirit and scope of the appended claims. Accordingly, the present invention should not be limited to the embodiments as described above, but should further include all modifications and equivalents thereof within the spirit and scope of the description provided herein.

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tatgtgcaac caaacgtcat gggaaacctt cggaggagga cagttgccat cttatgggag	780
gttgacatta cctctagatc caagcactga ccttcaactt aacatctcgt ttacatacgg	840
tccgggttat actgaatgga gacggtatgg attattatga aagcccactg tcggactccg	900

<210> SEQ ID NO 9
 <211> LENGTH: 1133
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 9

gccgggctgc atcaccocct agtaagacag gtgaatttta cttattactg cgatacaatt	60
gggatcagaa aatctattgc atcggcagca aatcccaccc tcctgtcagc actctctggg	120
ggcagagggtg acatattccc accatacaga tgcagtgagg ctgtacctc agtaggcaga	180
gttttcccc tatcagtgtc attgtccatg tctttgatct caagaaaatc agagataatc	240
aatatgctaa ccgctatctc aaacggagtg tatggtaaaa cttatttact agtgcctgat	300
tatattgaag aggagttoga cacacaaaag attcagatct ttgagatagg gttcatcaaa	360
cgggtggctga atgacatgcc attactccag acaaccaact atatggctct cccagagaat	420
tccaaagcta aggtatgtac tatagcagtg gccgagttga cactggcttc cttgtgtgta	480
ggtgagagca ccgtgttgtt atatcatgac agcaatggtt cgcaagataa taccctagta	540
gtgacgctgg gaatatttgg ggcaacatct atggatcaag ttgaagaggt gatacctgtt	600
gctcaccat cagtagaaaa aatacatata acaaatcacc gtgggttcat aaaagattca	660
atagcaacct ggatgggtgc tgcattgttc tctgagaaac aggaagagca aaaaaattgt	720
ctggagtcgg cttgtcaaag aaaatcctac cctatgtgca accaaacgctc atgggaaccc	780

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ttcggaggag gacagttgcc atcttatggg cggttgacat tacctctaga tccaagcact	840
gacccttcca acttaacata tcgtttacat accgtccggt tatacttgaa tggagacggt	900
atggataatt atgaaagccc actgtcggac teggatggct taccatttcc ttccaaaacg	960
gaacagtcct tggattgata aacaaaccag taggggagac cagttcattg tatccccat	1020
gtgttgacca ttgccccagg gaatcaaggg gaatgtattt acctattcaa ccttcccaaa	1080
taatgggata aaggatggcc ctectgaatc caaattacgg tgttgccta aac	1133

<210> SEQ ID NO 10
 <211> LENGTH: 1120
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 10

ttggttaagg ccctcctttt tccctaactc gggctgttg aataggtaaa taacaattcc	60
ccacttgatt ccctgggggc aaatgtcaac acatggggga ttacaatgaa ctggtctccc	120
ctacttgctt tgtttatcaa tccaaggact gttccgttt tgggaggaat ggtaagccat	180
cggagtcgc acagtgggct ttcataataa tccataccgt ctccattcag tataaccgga	240
cogtatgtaa acgatatggt aagttgaagg tcagtgcttg gatctagagg taatgtcaac	300
cgcccataag atggcaactg tctcctccg aagggtccc atgacgtttg gttgcacata	360
gggtaggatt ttctttgaca agccgactcc agacaatttt tttgctctc ctgtttctca	420
gagaccaatg caggcaccat ccaggttgc attgaatctt ttatgaacct acggtgattt	480
gttatatgta tttttctac tgatgggtga gcaacaggtc tcacctctc aacttgatcc	540
atagatggtg ccccaaatat tcccagcgtc actactagga tattatcttg cgaaccattg	600
ctgtcatgat ataacaacac ggtgctctca cctacacaca aggaagccag tgtcaactcg	660
cccactgcta tagtacatag cttagctttg gaattctctg ggaggaccat atagttggtt	720
gtctggagta atggcatgct attcagccac cgtttgatga accctatctc aaagactcga	780
atctttttgt gtgtcgaact cctcttcaat ataaatcagg cacctagtaa ataaagtta	840
ccatacacct ccgtttgaga tagccgggta gcatattgat tatctctgat cctcttgaga	900
tcaaagacat ggacaatgac actgataggc gggaaaactc tgcctactga ggtagcagct	960
ctactgcttt tgttgggtgg gaaatattha accctttgcc cccgaaagtg cttacaggag	1020
gatgggattt gctgccgac caataaattt tctgatccca attgtatcga agaactaata	1080
aattacctgg accttacttg ggggggtgat gaaccagcgc	1120

<210> SEQ ID NO 11
 <211> LENGTH: 797
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 11

agatcaaggt gaattttact aattactgcg atacaattgg gatcagaaaa tctattgcat	60
cggcagcaaa tcccactctc ctgtcagcac tctctggggg cagaggtgac atattccac	120
catacagatg cagtggagct gctacctcag taggcagagt tttccccta tcagtgtcat	180
tgtccatgct tttgatctca agaaaatcag agataatcaa tatgctaacc gctatctcaa	240
acggagtgta tggtaaaact tatttactag tgcctgatta tattgaagag gagttcgaca	300
cacaaaagat tccagctctt gagatagggt tcatcaaacg gtggctgaat gacatgccat	360
tactccagac aaccaactat atggctctcc cagagaattc caaagctaag gtatgtacta	420

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tagcagtggg cgagttgaca ctggcttct tgtgtgtagg tgagagcacc gtgttggtat	480
atcatgacag caatggttcg caagataata tcttagtagt gacgctggga atatttgggg	540
caacatctat ggatcaagtt gaagaggtga tacctgttgc tcaccatca gtagaaaaa	600
tacatataac aaatcacogt gggttcataa aagattcaat agcaacctgg atggtgctg	660
cattggctctc tgagaaacag gaagagcaaa aaaattgtct ggagtcggct tgtcaaagaa	720
aatcctaccc tatgtgcaac caaacgtcat gggaaacctt tcggaggagg acagttgcca	780
tcttatgggc ggttgac	797

<210> SEQ ID NO 12
 <211> LENGTH: 745
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 12

cccccaattgg cattgaacca tgtatccggc tcttgggttg catgagtttt ceggggagtt	60
aacaaccatt gaatccctta tgatgctata tcaacagatg ggtgaaacag cacctacat	120
ggttattctg gaaaattctg tccagaacaa atttagtgc ggctcctacc cattgctctg	180
gagttatgct atgggagttg gtgttgaact tgaaaactcc atgggagggt taaatttcgg	240
tagatcctac tttgaccogt cttatttcag gctcgggcaa gaaatggta gaagatctgc	300
cggtaaggta agctctgcac ttgcccgcga gctcggcacc accaaggaag aggctcagct	360
agtgtcagaa atagcatcca agacaacaga ggacctcca tttggcattg aaactatgta	420
tccggctctt gggttgcatg agttttccgg ggagttaaca acccttgaat cttaatgacc	480
ttttccgca gggaaacaac ccacaatgc tgaattctgt gaaatatggc tcaccacatt	540
gtggcagctc gacaccgact ttaaccttac ctatggaatt tggcggtgaa actgtaaatc	600
cctcttcggg ttaccacctc tttgatcac ttaaacggtt atttacgccg gcagccacgt	660
tagaacatat ccgctctgc aagtttctct gcctcctctc tcacccaat tagagggccc	720
ccctcctttg ttatgaacc cctta	745

<210> SEQ ID NO 13
 <211> LENGTH: 1167
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 13

cctgggcgcc ttacccccct ctagtaagct caggtgaatt ttactaacta ctgcgatacc	60
cttgggatca gaaaatctat tgcacggca gcaaatccca tcctcctgtc agcaactctct	120
gggggcagag gtgacatatt cccaccatac cgatgcagtg gagctgtac ctgagtaggc	180
agagttttcc ccctgtcagt gtcattgtcc atgtcttga tctcaagaaa atcagagata	240
atcaatatgc taaccgctat ctcaaacgga gtgtatggta aaacttattt actagtgcct	300
gattatatg aagaggagtt cgacacacaa aagattcgag tctttgagat agggttcatc	360
aaacgggtggc tgaatgacat gccattactc cagacaacca actatatggt cctcccagag	420
aattccaaag ctaaggtagt tactatagca gtgggagagt tgacactggc ttccttgtgt	480
gtaggtgaga gcaccgtgtc gttatatcat gacagcaatg gttecgcaaga tagtataccta	540
gcagtgacgc tgggaatatt tggggcaaca tctatggatc aagttgaaga ggcgatacct	600
gttgcctacc catcagtaga aaaaatacat ataacaaatc accgtgggtt cataaaagat	660
tcaatagcaa cctggatggt gcctgcattg gtctctgaga aacaggaaga gcaaaacaat	720

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tgtctggagt cggcttgta aagaaaatcc taccctatgt gcaaccaaac gtcacgggaa 780
cccttcggag gaggacagtt gccatcttat gggcggttga cattacetta gatccaagca 840
ctgaccttca actcaacata tcgcttacat accgtccggc tatactgaat gggagacggt 900
atggatttta tgacaagccc cctgtcgga ctcccggatg gcttaccacc ccctccaaa 960
accggaacag ctccctcgat tgataacca aaccagtacg aggagactca gtttcattgt 1020
tattccccta cgtgttgaca tttccgcccc aggccatcca tgteggattg ctetttacc 1080
aataaccac cccacatcat ggatacagct ctccctactg actccacact accgctgttg 1140
cctaccctcc cgctctccct tccccta 1167

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<210> SEQ ID NO 14
<211> LENGTH: 245
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (161)..(161)
<223> OTHER INFORMATION: n is a, c, g, or t

```

```
<400> SEQUENCE: 14
```

```

gtgaatttta ctaattactg cgatactatg gggatcagaa aatctattgc atcggcagca 60
aatcccatcc ttttatcagc actctccgga ggtagagggtg acatattccc accatacaga 120
tgcaatggag ctactatttc agtaggcaag attttcccc natcagatc attatctatg 180
tctttgatct caagaacatc agagataatc aatatgctaa ccgctatctc agacggagtg 240
tatgg 245

```

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<210> SEQ ID NO 15
<211> LENGTH: 835
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus
<220> FEATURE:
<221> NAME/KEY: misc_feature
<222> LOCATION: (764)..(764)
<223> OTHER INFORMATION: n is a, c, g, or t

```

```
<400> SEQUENCE: 15
```

```

caagtgtaat tttactaatt actgcgatac aattgggac agaaaatcta ttgcacggc 60
agcaaatccc atcctcctgt cagcactctc tgggggcaga ggtgacatat tcccaccata 120
cagatgcagt ggagctgcta cctcagtagg cagagttttc cccctacag tgacattgtc 180
catgtctttg atctcaagaa aatcagagat aatcaatatg ctaaccgcta tctcaaacgg 240
agtgtatggt aagacttatt tactagtgcc tgattatatt gaagaggagt tcgacacaca 300
aaagattcga gtctttgaga tagggttcat caaacggtgg ctgaatgaca tgccattact 360
ccagacaacc aactatatgg tctcccaga gaattccaaa gctaaggatg gtactatagc 420
agtggcgag ttgacactgg cttccttgty tgtaggtagg agcacctgtg tggtatatca 480
tgacagcaat ggttcgcaag ataatacct agtagtgacg ctgggaatat ttggggcaac 540
atctatggat caagttgaag aggtgatacc tgttgctcac ccatcagtag aaaaaataca 600
tataacaaat caccgtgggt tcataaaaga ttcaatagca acctggatgg tgccctgcatt 660
ggtctctgag aaacaggaag agcaaaaaa ttgtctggag tcggcttgctc aaagaaaatc 720
ctaccctatg tgcaacaaa cgctcatggga acccttcgga gganggacag ttgccatctt 780
atggcggtt gacattacct ctgatccaa gcactgacct tcaacttaac atacc 835

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<210> SEQ ID NO 16
<211> LENGTH: 1119
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 16
tcctgtttgc cttcccccc cctagtaaga tcaggtgaat tttactaaca actgcgatac   60
aattgggatc agaaaatcta ttgcatcggc agcaaatccc atcctcctgt cagcactctc   120
tgggggcaga ggtgacatat tcccaccata cagatgcagt ggagctgcta cctcagtagg   180
cagagttttc cccctatcag tgcattgtc catgtctttg atctcaagaa aatcagagat   240
aatcaatatg ctaaccgcta tctcaaacgg agtgtatggt aaaacttatt tactagtgcc   300
tgattatatt gaagaggagt tcgacacaca aaagattcga gtctttgaga tagggttcat   360
caaacggtgg ctgaatgaca tgccattact ccagacaacc aactatatgg tcctcccaga   420
gaattccaaa gctaaggtag gtactatagc agtgggcgag ttgacactgg ctctcctgtg   480
tgtaggtgag agcacctgtg tgttatatca tgacagcaat ggttcgcaag atagtatcct   540
agcagtgaoc ctgggaatat ttggggcaac aactatggat caagttgaag aggtgatacc   600
tgttgctcac ccatcagtag aaaaaataca tataacaaat caccgtgggt tcataaaaga   660
ttcaatagca acctggatgg tgctgcatt ggtctctgag aaacaggaag agcaaaaaaa   720
ttgtctggag tcggctgtc aaagaaaatc ctaccctatg tgcaacccaa cgttatggga   780
acccttcgga ggaggacagt tgccatctta tgggcgggtg acattacctc tagatccaag   840
cactgacett caacttaaca tatcgtttac atacgggctcg gttatcctga atggagacgg   900
tatggattat tatgaaagcc cactgtogga ctcccgatgg cttaccattc ctccaaaacg   960
gaacagtcct tggattgata aacaaaacaag tagaggagac cagttcattg aatccccatg  1020
tgttgacttt tcgcccaggg aatcaagtgg aattgtattt actatcaact tccagattat  1080
ggataagatg tccttctgat tccaatcgg tgtgcctta                               1119

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<210> SEQ ID NO 17
<211> LENGTH: 1126
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 17
tcgtgggtgct taaccccccc tagtaagatc aggtgaattt tactaattac tgcgatacta   60
ttgggatcag aaaatctatt gcacggcag caaatcccat ctttttatca gcaactctccg   120
gaggtagagg tgacatattc ccaccataca gatgcaatgg agctactatt tcagtaggca   180
agattttccc cctatcagta tcattatcta tgtctttgat ctcaagaaca tcagagataa   240
tcaatatgct aaccgctatc tcagacggag tgtatggtaa aacttattta ctaatgectg   300
attatattga aggggagttc gacacgcaaa agattcgagt ctttgagata gggttcatca   360
aacggtggct gaatgacatg ccattactcc agacaaccaa ctatatggct ctcccagaga   420
attccaaagc caaggtatgt actatagcag tgggcgagtt gacactggct tctttgtgtg   480
tagatgagag caccgtattg ttatatcatg acagcaatgg ttcacaagat ggtgttctag   540
tagtgacgct gggaaatattc ggggcaacat ctatggatca agttgaagag gtgatacctg   600
tcgctgaccc attagcagaa aaaatacata taacaaatca ccgtgggatc ataaaagact   660
caatagcaac ctggatgggt cctgcattag tttctgagaa acaagaggaa caaacaattt   720
gtctggagtc agcttgtcaa agaaaatcct accctatgtg caatcaaacg tcatgggaac   780
cctttggagg aggacagttg ccattctatg ggcggctgac attacctcta catccaagca   840

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ttgacctcca cttaacatat catttacata cggctccgact atactgaatg gagacggatg   900
gctattatga gagccccctg cggactccgg atggcttacc ttccctcca gcacggcaca   960
gcctggattg ataacaag agtagaggac gaccagttat tgtcattccc ctgtgttgac  1020
atctgcctcc cggcatccac ccgaaattgc tattacccta tcccacattc cccttcgcgc  1080
tcaagatccc cctcctgctc cccaccacgg cgcgctccct atctcc                1126

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<210> SEQ ID NO 18
<211> LENGTH: 623
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

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<400> SEQUENCE: 18

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```

cctagtagat caaggtgaat ttactaatt actgcgatac aattgggatc agaaaatcta   60
ttgcatcggc agcaaatcca atcctcctgt cagcactctc tgggggcaga ggtgacatat  120
tcccaccata cagatgcagt ggagctgcta cctcagtagg cagagtttcc ccctatcag  180
tgtcattgtc catgtctttg atctcaagaa aatcagagat aatcaatag ctaaccgcta  240
tctcaaacgg agtgtatggt aaaacttatt tactagtgcc tgattatatt gaagaggagt  300
tcgacacaca aaagattcga gtctttgaga tagggttcat caaacggtgg ctgaatgaca  360
tgccattact ccagacaacc aactatatgg tcctcccaga gaattccaaa gctaagggat  420
gtactatagc agtgggcgag ttgacactgg ctctcctgtg tgtaggtgag agcaccgtgt  480
tgttatatca tgacagcaat ggttcgcaag atagtatcct agcagtgacg ctgggaatat  540
ttggggcaac atctatggat caagttgaag aggtgatacc tgttgctcac ccatcagtag  600
aaaaaataca tataacaat cac                                           623

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<210> SEQ ID NO 19
<211> LENGTH: 866
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

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<400> SEQUENCE: 19

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gtttgatagg taaataacag ttccacttg atccctcggg tgcaaatgac aacacatgag   60
ggaccacagt gaactgtgct cctctacttg cttgtttat caatccaaga attgttccat  120
tcttaggagg aatggtaagc catccggatt ccaaaagtgg gctttcataa taatccatac  180
catctccatt cagtataacc ggaccgtatg taaatgatat gttaagtta cggatcaatgc  240
ttgcatctag atgtaacgtc aaccgcccac aagatggcaa ctgtcctcct ccaaagggtt  300
cccatgatgt ttgattgac atggggtagg atttctctcy acaagccgac tccagacaat  360
tttctgttcc ctctaggttc tcagagacca atgcaggcac catccaggtt gctactgaat  420
cttttatgaa accacgtgta tttgttatat gtatttttcc gactgatggg tgagcgacag  480
gtatcacctc ttcgacttga tccataggtg ttgccccaaa gattcccagc gtcactacta  540
gaataccatc ttgtgaacca ttgctgtcat gatataataa tacagtgtct tcatctacac  600
acaaggaagc cagtgtcaac tcgcccactg ctatagtaca taccttgggt ttagaattct  660
ccgggaggat catatagttg gttgtctgga ataatggcat gtcattcagc caccttttga  720
tgaaccctat ctcaaagact cgaatctttt gcgtgtcgaa ctcccctca atataatcag  780
gcactagcaa ataagtttta ccatacactc cgtctgagat agaggtcagc atattgatta  840
tctctgatgt tcttgagatc aaagac                                           866

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<210> SEQ ID NO 20
<211> LENGTH: 1134
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 20
aatgcttcct ttaccacacc tagtaagatc aagtaaattt tacggtaaat aaatagcgat    60
acaattggga tcagaaaaatc tattgcatcg gcagcaaatc ctatcctttt atcagcactc    120
tccggaggta gaggtgacat attcccacca tacagggtgca gtggagctac tacttcagta    180
ggcagagtct tccccctatc agtatcattg tccatgtctt tggctcctcaag aacatctgaa    240
ataatcaata tgctaaccgc tatctcagac ggtgtgtatg gtaaaaactta tttgctagtt    300
cctgattatc ttgaagggga gttcgacacg caaaagattc gagtctttga gataggggtc    360
atcaaacggt ggctgaacaa catgccatta ctccagacaa ccaactatat ggtcctcccg    420
gaggattcca aagccaaggt atgtactata gcggtgggcg agttgacact ggcttctctg    480
tgtgtagatg agagcaccgt attgttatat catgacagca gtggttcaca agatggtatt    540
ctagtgggtg cgctgggaat atttggggca acacctatgg atcaagtga agagggtgata    600
cctgttgctc acccatcagt agaaaaaata catatagcaa accaccgtgg gttcatcaaa    660
gattcaatag caacctggat ggtgcctgca ttggtctctg agaaacaaga ggaacaaaaa    720
aattgtctgg agtcggcttg tcaaagaaaa tcctacoccta tgtgcaacca aacgtcatgg    780
gaaccctttg gaggaggaca gttgccatct tatgggcggt tgacattacc tctagatcaa    840
agcattgacc tccagcttaa catctcattt acatatggtc cggttatact gaatggagac    900
ggtatggatt attatgaaag tccgcttttg aactccggat ggcttaccat tcctcccaag    960
aacggaacag tccttgatt gataaaciaa gcaagtagag gagaccagtt cactgtatcc    1020
ccatgtgtga catttgccc cagggaaatca agtgaattg tatttaccta tcaaacatc    1080
ccagatatgg ataagatgt ccttactgaa tccaaattag tgggtgtgac taac          1134

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<210> SEQ ID NO 21
<211> LENGTH: 1124
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 21
accgggggtgc ttaccccccc tagtaagatc aagtgaattt tacgaaaaac tgcgatccaa    60
ttgggatcag gaaatctatt gcaacggcag caaatcctat ccttttatca gcaccctccg    120
gaggttagagg tgacatattc ccatcataca gatgcagtgg agctactact tcagtaggca    180
gagtcttccc cctatcagta tcattgtcca tgtctttgat ctcaagaaca tctgaaataa    240
tcaatatgct aaccgctatc tcagacggag tgtatggtaa aacttatctg ctagtctctg    300
attatcttga aggggagttc gacacgcaaa agattcgagt ctttgagata gggttcatca    360
aacggtggct gaacaacatg ccattactcc agacaaccaa ctatatggtc ctcccagagg    420
attccaaagc caaggtatgt actatagcag tgggcgagtt gacactggct tccttgtgtg    480
tagatgagag caccatattg ttatatcatg acagcaatgg ttcacaagat ggtattctag    540
tggtgacgct gggaaatatt ggggcaacac ctatggatca agttgaagag gtgatacctg    600
ttgctcacc c atcagtagaa aaaatacata tagcaaacca tcgtgggttt atcaaagatt    660
caatagcaac ctggatgggt cctgcattgg tctctgagaa acaagaggaa caaaaaaatt    720
gtctggagtc ggcttgtcaa agaaaatcct accctatgtg caaccaaaacg tcatgggaac    780
cctttggagg aggacagttg ccatcttatg ggcggttgac attacctcta gatccaagca    840

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ttgaccttca gcttacatct catttacata cggcccggtta tactgaatgg agacgggatg 900
gatactatga aagcccactt ttagactccg gatggcttac cattcctcca agaacggaac 960
agtccttggg ttgataaaca aagcaagtag aggagaccag ttcactgtat ccccatgtgt 1020
tgacatttgc gccaggaatc agtggaatg gttatttacc tattcaaac tcccaattat 1080
ggataagagt cctactggat ccaaatatg gtgttccct aacc 1124

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<210> SEQ ID NO 22
<211> LENGTH: 1145
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

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<400> SEQUENCE: 22

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```

cattggtgca ttaaccacc tagtaagaca agtgaatgtt actaatatac tgcgatacaa 60
ttgggatcag gaaatctatt gcactcggcag caaatcctat ccttttatca gcaccctccg 120
gaggttagagg tgacatattc ccatcataka gatgcagtgg agctactact tcagtaggca 180
gagtcttccc cctatcagta tcattgtcca tgtctttgat ctcaagaaca tctgaaataa 240
tcaatatgct aaccgctatc tcagacggag tgtatggtaa aacttatctg ctagtctctg 300
attatcttga aggggagttc gacacgcaaa agattcagat ctttgagata gggttcatca 360
aacggtggct gaacaacatg ccattactcc agacaaccaa ctatatggtc ctcccagagg 420
attccaaagc caaggtatgt actatagcag tgggcgagtt gacactggct tccttgtgtg 480
tagatgagag caccatattg ttatatcatg acagcaatgg ttcacaagat ggtattctag 540
tggtgacgct ggggaatatt ggggcaacac ctatggatca agttgaagag gtgatactg 600
ttgctcacc atcagtagaa aaaatacata tagcaaacca tcgtgggttt atcaaagatt 660
caatagcaac ctggatggg cctgcattgg tctctgagaa acaagaggaa caaaaaaatt 720
gtctggagtc ggcttgcaca agaaaatcct accctatgtg caaccaaagc tcatgggaac 780
cctttggagg aggacagttg ccatcttatg ggcggttgac attacctcta gatccaagca 840
ttgaccttca gcttaacatc tcatttacat acggtcoggt tatactgaat ggagacggta 900
tggattacta tgaagccca ctttagact ccggatggct taccattcct cccaagaacg 960
gaacagtcct tggattgata aacaaagcaa gtagaggaga ccagttcact gtaatcccc 1020
atgtgttgac atttgcgcc agggaatcaa gtggaatgt ttatttacct attccaaaca 1080
tcccagatta tggataaagg atgtccttac tgaagtctta aattagtggg ggtttgcct 1140
aagac 1145

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<210> SEQ ID NO 23
<211> LENGTH: 321
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

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<400> SEQUENCE: 23

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gcctcccagg ggcaccttcc cccccagta gctcagggtg atctcactta aaactgcgcc 60
ccccttggga tcttacaatc tattgcatcg gcagcaaatc ccctcctttt atcageactc 120
tcccagagta gaggtgacat attcccacca taccgatgca atggagctac tatttacta 180
ggcaagatgt cccccctatc agtatcatta tctatgtctt tgatctcacy aacatcagag 240
ataatcaata tgctaaccgc tatctcatak ggagtgtatg gtaaaactta tttactaatg 300
cccgactata ttgaagggga g 321

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<210> SEQ ID NO 24
<211> LENGTH: 1135
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 24
ttgatttcga ctccccgatt ttccactgtg cattaaccac ctagtaagat caaggtgaat    60
tttactgact ctggaacaaa tgggatcaag aaatttattg catggcagca aatcccatct    120
cctgtcagca ctctatgggg gcagaggtga catattccca ccatacaaga tgcagtggag    180
ctgctacctc agtaggcaga gttttccccc tatcagtgtc attggccatg tctttgacct    240
caagaaaatc agaggataat caatatgcta accgctatct caaaacggag tgtatggtaa    300
aacctattta ctagtgcctg atttatattga agaggagttc gacacacaaa aagattcgag    360
tctttgagat aggggttcac aaacgggtgc tgaataacat gccattactc cagacaacta    420
actatatggt cctcccagag aattccaaag ctaaggtagt tactatagca gtgggcgagt    480
tgacactggc ttccttgtgt gtaggtgaga gcaccgtgtt gttatatcat gacagcaatg    540
gttcgcaaga tagtataccta gcagtacgc tgggaatatt tggggcaaca tctatggatc    600
aagttgaaga ggtgatacct gttgctcacc catcagtaga aaaaatacat ataacaaatc    660
accgtggggt cataaaagat tcaatagcaa cctggatggt gcctgcattg gtctctgaga    720
aacaggaaga gcaaaaaaat tgtctggagt cggcttgta aagaaaatcc taccctatgt    780
gcaaccaaac gtcatgggaa cccttcggag gaggacagt gccatcttat gggcggttga    840
cattacctct agatccaagc actgaccttc aacttaacat atcgtttacg tacggtcagg    900
ttatactgaa tggagacggt atggattatt atgaaagccc actgtcggac tccggatggc    960
ttaccattcc tccccaaaac ggaacagtcc ttggattgat aaacaaagca agtagaggag   1020
atcagttcat tgtaatcccc catgtgttga catttgccgc cagagaatca agtgggaatt   1080
gttatttacc tattcaaaca tcccatatta ggaaaaaggg aggcctaccc gggga      1135

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<210> SEQ ID NO 25
<211> LENGTH: 796
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 25
tatggttcat tccccccgg cgtaagtga tttgaatcgt agtaattgct gtgataaaat    60
tgggattgga aatgtattgc attgttatga aattctacct ttccagcact tgccctcgtt    120
ggttgagggg acttattccc atcacacata tgcagtggag ctactacctc atccggcaga    180
gttatatttg atcatcatta ttgcacatgt ttgtgacctt aaaaacatct ggcatatgca    240
atctgctaac cgcgatctca tgtggagtgt atggcaaac ttatctgcta ctctcgtatt    300
ttcttgaagg ggagtccgac actctgccga tgtccgacaa gctgatcggg ttcataaac    360
tctggctgaa caacatgttg cgcgtctgac aacctccgat ttggcctgcc cagaggattt    420
tacagccaag gtatgtacca tatcccaggg gaacttcaca ctgccttcc tgtgtgttag    480
ccagagcccc atattgtccc ataatgatat gaatgtccta caagaggta ttttccatgt    540
gacccccgct tcatttgtgg caatggcggg ggttcaattg gaacagggtg tatctgacce    600
tatctttcac tagagaaatt acatatgaca aaccatcatg gcttgatcaa agaataactt    660
cctttctggc tgacgcttga cttgccctta tatataccat attttcttaa taaatcgagg    720
tcaattgctt gtggagccaa attttaccac tcttccaacc ttatgttacg ggccttccct    780
gccggaggac cgttgc      796

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<210> SEQ ID NO 26
 <211> LENGTH: 1122
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 26

```

gcgattttgc cctgtgcatt aaccaccta gtaagatcaa ggtaaat ttt actaaattct    60
gcgaaacatg tggatcagaa aatctatggc atcggcagca atcccatcct cctgcagccc    120
tcttggggca gaggtgacat attcccacca tacagatgca gtgaggctgc tacctcagta    180
ggccagagtt tccccctatc agggtcattg tgcattgtct tgacctcaag aaagtcagag    240
ataatcaaat atgctaacc gctatctcaa acggagtgta tgggaaaaac ttatttacta    300
gtgcctggat tatattgaag aggagtcca cacacaaaag attcagatct ttgagatagg    360
gttcatcaaa cgggtggctga ataacatgcc attactccag acaactaact atatggctct    420
cccagagaat tccaaagcta aggtatgtac tatagcagtg ggcgagttga cactggcttc    480
cttgtgtgta ggtgagagca cctgtgtgtt atatcatgac agcaatggtt cgcaagatag    540
tatcctagca gtgacgtgga gaatatttgg ggcaacatct atggatcaag ttgaagaggt    600
gatacctggt gctcaccat cagtagaaaa aatacatata acaaatcacc gtgggttcat    660
aaaagattca atagcaacct ggatggtgcc tgcattggtc tctgagaac aggaagagca    720
aaaaaattgt ctggagtcgg cttgtcaaaag aaaatcctac cctatgtgca accaaacgtc    780
atgggaaccc ttcggaggag gacagttgcc atcttatggg cggttgacat tacctctaga    840
tccaagcact gacctaac ttaacatc gtttacatc ggtccggtta tactgaatgg    900
agacgggatg gattattatg aaagccact gtcggactcc ggatggctta ccattctccc    960
caaaaacgga acagtccttg gattgataaa caaagcaagt agaggagatc agttcattgt   1020
aatcccccat gtgttaacat ttgcgcccag agaatcaagt gggggattgt tattttccta   1080
ttcaaacatg cccatattat gataaaggat ggccttaacc cg                               1122

```

<210> SEQ ID NO 27
 <211> LENGTH: 798
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 27

```

agttcgacgc aaaaagatt cgagtgtgta gatagggttg atcggacgag gaggtgaagg    60
acatgccatt actccagaca gctaactata tggtcgcccc agagaattcc aaagctaagg    120
tatgtactat agcagtgggc gaggtggcac tggcttctct gtgtgtaggg gagagcgccg    180
tgttgttata tcattggcagc aatggttcgc aagatagtat cgtagcagtg acgctgggaa    240
tatttggggc aacatctatg gatcaagttg aagagggtgat acctgttgcct caccatcag    300
tagagaaaat acatatagca aatcacctg ggttcataaa agattcaata gcaacctgga    360
tgggtcctgc attggtctct gagaaacagg aagagcaaaa aaattgtctg gagtcggctt    420
gtcaaaagaaa atcctaccgt atgtgcagcc aaacggcatg ggaacccttc ggaggaggac    480
agttgccatc ttatggggcg ttgacattac ctctagatcc aagcgtgcc ttcaacttaa    540
catatcgttt acatacggtc cggttatact gaatggagac ggtatggatt attatgaaag    600
cccactgtcg ggctccggat ggcttgccat tctcccaaaa aacggaacag tccttgatt    660
gataaaca aaa gcaagtagag gagatcagtt cattgtaatc ccccatgtgt ggacatttgc    720
gcccagagaa tcaagtgggg gattgttttt taaactatgc aaacggcgca tatgaggggg    780

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gagggggggc gggaggct 798

<210> SEQ ID NO 28
 <211> LENGTH: 1035
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 28

cagtgagagc aaaaatgtag gaaagggcag gaattccatg ctcaaggagc ggatgtgggg 60
 agaggttgag agtcccgccg gcagtgacag aaggggtact cagtagcggg gtttcccctc 120
 aggaggggga ttgtccagtc tttgatatac gaaaagaagg atatcaatat gctaaccgct 180
 atcgccaaag gagggtagtg taagagctta ttgggagtg ctagtagagc ggagggaagt 240
 tctacaggag agagattgga gtggtgagat gggggttcgt caagcggtag atgaatgaca 300
 taccattact ccagacaacc aagtataggg gcctcccaga gaatgccaaa gctaaggtag 360
 gtactatagc agtgggcgag ttacgctggc ttccttgtgt gtaggtgaga gcgccgtgtt 420
 gttatatcat gacagcaatg gttcgcaaga tagtacctc gctgtgacgc tgggaatatt 480
 tggggcgaca tctatggatc aagttgaaga ggtgatgcct gttgctcacc catcagtaga 540
 aaaaatacat ataacaatc gccgtgggtt cataaaagat tcaatagcag catggatggt 600
 gcctgcattg gtctctgaga agcaggaaga gcaaaaaat tgtcaggagc cgggtgttca 660
 aagaaaatcc taccgatgt gcaaccaaac gtcattggaa cccttcggag gaggacaggt 720
 gccatcttat gggcggttgg cattacctct agagccaagc actggccttc aacttgacat 780
 atcgtttaca tacgggcgag ttatactgaa tggagacggt atggattatt atgaaagccc 840
 actgtcggac gccggatggc ttaccattcc tccccaaaac ggaacagtc gtggattgat 900
 aaacaaagca agtagaggag gccagttcat tgtaatcccc catgtgttga cttttgcgcc 960
 cagggaatca agtgggaatt gctattttcc tattcagaac accccagatt aggatagaag 1020
 gaggggcctg ggccg 1035

<210> SEQ ID NO 29
 <211> LENGTH: 1131
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 29

cttgtgggct taaaccacct agtaatacaa agtgaatttt actaattact gcgatacaat 60
 tgggatcaaa aaatctattg catcggcagc aaatcctatc cttttatcag cactctccgg 120
 aggcagaggt gacatattcc caccatacag atgcagtgga gctactactt cagtaggcag 180
 agtcttcccc ttatcagtat cattgtccat gtctttgatc tcaagaacat ctgaaataat 240
 caatatgcta accgctatct cagacggagt gtatggtaaa acttatttgc tagttcctga 300
 ttatcttgaa ggggagttcg acacgccgaa gattcagatc tttgagatag ggttcatcaa 360
 acggtggctg aacaacatgc cattaatcca gacaaccaac tatatggctc tcccggagga 420
 ttccaaagct aaggtatgta ctatagcagt gggcgagttg aactggctt ccttatgtgt 480
 agatgagagc accgtattgt tatatcatga cagcaatggt tcacaagatg gtattctagt 540
 ggtgacgctg ggaatatttg gggcaacacc tatggatcga gttgaagagg tgatacctgt 600
 tgctcaccgg tcagtagaaa aaatacatat ggcaaacacc cggtgggttca tcaaagattc 660
 aatagcaacc tggatggtgc ctgcattggt ctctgagaaa caagaggaac aaaaaattg 720
 tctggagtcg gcttgcataa gaaaatccct accctatgtg caaccaaacc tcatgggaaa 780

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cccttggag gaggacagtt gccatcttat gggcggttga cattacctct agatccaagc 840
attgaccttc accttaacat ctcatttaca tacggcccag ttatactgaa tggggacggt 900
atggattatt atgaaagccc acttttggac tccggatggc ttaccattcc tcccaagaac 960
ggaacagtcc ttggattgat aaacagagca gtagaggaga acagttcact gtaatcccca 1020
tgtgttgact tgcgcaaggg gatcaagtgg aaattgtatt tacctattca aacatcttaa 1080
attatggata aagatgcctt caccgagccc aaattagtgg tgttgcctca t 1131

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<210> SEQ ID NO 30
<211> LENGTH: 759
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

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```
<400> SEQUENCE: 30
```

```

ctccccttgc gcttgaacat gtatccggct cttgggttgc atgagtttcc cggggagtta 60
acaaccattg aatcccttat gatgctatat caacagatgg gtgaaacagc accgtacatg 120
gttattcttg aaaattctgt ccagaacaaa tttagtgcag gatcctaccc attgctctgg 180
agttatgcta tgggagtgtg tgttgaactt gaaaactcta tgggagggtt aaatttcggt 240
agatcctact ttgaccagc ttatttcagg ctccggcaag aaatggttag aagatcggcc 300
ggtaaggtaa gctctgcact tgcgcccag cttggcatca ccaaggaaga ggctcagcta 360
gtgtcagaaa tagcatccaa gacaacagag gacccgcatt tggcattgaa actatgtatc 420
cggtccttgg gttgcatgag ttttccgggg agttaacaac cattgaaatc cttgtgatgc 480
tttaccacca aatgggtgaa ggacccccca tggttattct tggaaaattt gtcgcacaaa 540
attagtgcag gatctacat tgctctggag ttatgctatg ggagtgtgtg gtgaacttga 600
aaacccccat gggggggtta atttcggcag attcttcttt gacagttaat tttaggctcg 660
gccagaaaat ggtagaaaa ctccgcccgt taggggaaag ctttgccttt gcccgcttgg 720
gttccccccc cgaaagggtt ccccccttt ctatatatt 759

```

```

<210> SEQ ID NO 31
<211> LENGTH: 1132
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

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```
<400> SEQUENCE: 31
```

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tgtgaatgtg aacttccgcg atctccactg gtgcattaac cactagtaa gatcaaggtg 60
aatttactaa ctacgcgata caattgggat cagaaaatct attgcatcgg cagcaaatcc 120
catctcctg tcagcactct ctgggggcag aggtgacata ttcccacct accgatgcag 180
tggagctgct acctcagtag gcagagtttt ccccctgtca gtgtcaattgt ccatgtcttt 240
gatctcaaga aatcagaga taatcaatat gctaaccgct atctcaaacg gagtgtatgg 300
taaaacttat ttactagtgc ctgattatat tgaagaggag ttcgacacac aaaagattcg 360
agtctttgag ataggggttca tcaaacggtg gctgaaatgac atgccattac tccagacaac 420
caactatatg gtctcccag agaattccaa agctaaggtg tgtactatag cagtgggcca 480
gttgacactg gcttccttgt gtgtaggtga gagcaccgtg tcattatata atgacagcaa 540
tgggttcgaa gatagatcc tagcagtgc gctgggaata tttggggcaa catctatgga 600
tcaagttgaa gaggtgatac ctgttctca cccatcagta gaaaaaatac atataacaaa 660
tcaccgtggg ttcataaaag attcaatagc aacctggatg gtgcctgcat tggctctctga 720
gaaacaggaa gagcaaaaaa attgtctgga gtcggcttgt caaagaaaat cctaccctat 780

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gtgcaaccaa acgcatgagg aacccttcgg aggaggacag ttgccatctt atgggcggtt	840
gacattacct ctatagccaa gcactgacct tcaacttaac atatcgttta catacgggcc	900
ggttatactg aatggagacg gtatggatta ttatgaaagc ccaactgctg actccggatg	960
gcttaccatt cctcccaaaa acggaacagt ccttggattg ataaacaaag caagtagagg	1020
agaccagttc attgtaatcc cccatgtggt gacatttgcg ccaggggaat caagtgaggaa	1080
ttgttattta cctattcaaa catcccagat tatgaaaaga tgccttaacc cg	1132

<210> SEQ ID NO 32
 <211> LENGTH: 1127
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 32

tctgctgctt aaccacctag taagatcagg tgaattttac taactactgc gatacaattg	60
ggatcagaaa atctattgca tcggcagcaa atcccacct cctgtcagca ctctctgggg	120
gcagaggtga catattccca ccataccgat gcagtggagc tgctacctca gtaggcagag	180
ttttcccctc gtcagtgtca ttgtccatgt ctttgatctc aagaaaatca gagataatca	240
atatgctaac cgctatctca aacggagtgt atggtaaac ttatttacta gtgcctgatt	300
atattgaaga ggagtctgac acacaaaaga ttcgagtctt tgagataggg ttcataaac	360
ggtggctgaa tgacatgcca ttactccaga caaccaacta tatggctctc ccagagaatt	420
ccaaagctaa ggtatgtact atagcagtgg gcgagttgac actggcttcc ttgtgtgtag	480
gtgagagcac cgtgtcatta tatcatgaca gcaatggttc gcaagatagt atcctagcag	540
tgacgctggg aatatttggg gcaacatcta tggatcaagt tgaagaggtg aacctgttgc	600
tcacccatca gtgaaaaaaa tacatataac aaatcaccgt gggttcataa aagattcaat	660
agcaactgga tgggtcctgc attggtctct gagaaacagg aagagcaaaa aaattgtctg	720
gagtcggctt gtcaaaagaaa atcctaccct atgtgcaacc aaacgtcatg ggaaccctc	780
ggaggaggac agttgccatc ttatggggcg ttgacattac ctctagatcc aagcactgac	840
cttcaactta acatattcgtt tacatacggc cgggttatac tgaatggaga cggtatggat	900
tattatgaaa gcccaactgc ggactccgga tggcttacca ttccctccaa aaacggaaca	960
gtccttggat tgataaacia agcagtagag gagaccagtt cattgtaatc ccccatgtgt	1020
tgacatttgc gcccagggaa tcaagtggga attgttattt acctattcaa acatccagat	1080
tatggataaa gatgtcctta ctgagtccaa attagtgtgt gtgccta	1127

<210> SEQ ID NO 33
 <211> LENGTH: 1128
 <212> TYPE: DNA
 <213> ORGANISM: canine distemper virus

<400> SEQUENCE: 33

attggttgc cttaacccac ctatgaagat caggtgaatt ttactaacta ctgcgataca	60
attgggatca gaaaaatctat tgcateggca gcaaatccca tcctcctgctc agcaactctc	120
gggggcagag gtgacatatt cccaccatac cgatgcagtg gagctgctac ctccagtaggc	180
agagttttcc ccctgtcagt gtcattgtcc atgtcttga tctcaagaaa atcagagata	240
atcaatatgc taaccgctat ctcaaacgga gtgtatggta aaacttattt actagtgcct	300
gattatattg aagaggagtt cgacacacia aagattcgag tctttgagat agggttcatc	360
aaacgggtgc tgaatgacat gccattactc cagacaacca actatatggt cctccagag	420

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aattccaaag ctaaggtatg tactatagca gtgggagag tgacactggc ttccttgtgt 480
gtaggtgaga gcaccgtgtc attatatcat gacagcaatg gttcgcaaga tagtatecta 540
gcagtgacgc tgggaatatt tggggcaaca tctatggatc aagttgaaga ggtgatacct 600
gttgctcacc catcagtaga aaaaatacat ataacaaatc accgtggggt cataaaagat 660
tcaatagcaa cctggatggt gcctgcattg gtctctgaga aacaggaaga gcaaaaaaat 720
tgtctggagt cggtttgca aagaaaatcc taccctatgt gcaaccaaac gtcattggaa 780
cccttcggag gaggacagtt gccatcttat gggcggttga cattacctct agatccaagc 840
actgaccttc aacttaacat atcgtttaca tacgggtccgg ttatactgaa tggagacggt 900
atggattatt atgaagccc actgtcggac tccggatggc ttaccattcc tccccaaac 960
ggaacagtcc ttgaatgata aacaaagcaa gtagaggaga ccagtttatt gtactcctc 1020
tgtgtttgac atttgcgccc aggatcaagt ggcattggtt ctacctatcc aaacttcga 1080
attatggata aagatgtcct tactgatcca aactagtggg ttgctcaa 1128

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<210> SEQ ID NO 34
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

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```
<400> SEQUENCE: 34
```

```
atgtgggatt gcttagga 18
```

```

<210> SEQ ID NO 35
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

```

```
<400> SEQUENCE: 35
```

```
ggcgtcctc ttggacat 18
```

```

<210> SEQ ID NO 36
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

```

```
<400> SEQUENCE: 36
```

```
gttagctagt ttcactct 18
```

```

<210> SEQ ID NO 37
<211> LENGTH: 18
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

```

```
<400> SEQUENCE: 37
```

```
ggctcctctgt tgtcttgg 18
```

```

<210> SEQ ID NO 38
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

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<400> SEQUENCE: 38
gaattcgact tccgcatct cc 22

<210> SEQ ID NO 39
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

<400> SEQUENCE: 39
taggcaacac cactaattr gactc 25

<210> SEQ ID NO 40
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

<400> SEQUENCE: 40
tcgaaatcct atgtgagatc act 23

<210> SEQ ID NO 41
<211> LENGTH: 26
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide primer

<400> SEQUENCE: 41
atgctggaga tggtttaatt caatcg 26

<210> SEQ ID NO 42
<211> LENGTH: 1824
<212> TYPE: DNA
<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 42
atgctctcct accaagacaa ggtgggtgcc ttctataagg ataatgcaag agctaattca 60
tccaagctgt ccctagtgc agaagacaa gggggcagga gaccaccta tttgctgttt 120
gtcctttca tcctactggt tggaaactcg gccttgcttg ctatcactgg agttcgattt 180
caccaagtat caactagcaa cgtggaattt agcagattgc taaaagagga tatggagaaa 240
tcagaggctg tacatcacca agtcatagat gttttgacgc cgctcttcaa aattattgga 300
gatgagattg gggttacggct gccacaaaaa ctaaacgaga tcaacaatt catccttcaa 360
aagacaaact tcttcaatcc taacagggaa ttcgacttcc gtgatctcca ctggtgcatt 420
aaccaccta gtaagatcaa ggtgaatctt actaactact gcgatacaat tgggatcaga 480
aaatctattg catcggcagc aaatccatc ctccctgtcag cactctctgg gggcagaggt 540
gacatattcc caccataacc atgcagtgga gctgctacct cagtaggcag agttttcccc 600
ctgtcagtgt cattgtccat gtctttgatc tcaagaaaat cagagataat caatattgcta 660
accgctatct caaacggagt gtatggtaaa acttatttac tagtgcctga ttatattgaa 720
gaggagttcg acacacaaaa gattcgagtc tttgagatag ggttcatcaa acggtggctg 780
aatgacatgc cattaactca gacaaccaac tatatggtcc tcccagagaa ttccaaagct 840
aaagtatgta ctatagcagt gggcgagttg acactggctt ccttgtgtgt aggtgagagc 900
accgtgtcat tataatcatga cagcaatggt tcgcaagata gtatcctagc agtgacgctg 960

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ggaatatttg gggcaacatc tatggatcaa gttgaagagg tgatacctgt tgctcaccca 1020
tcagtagaaa aaatacatat aacaaatcac cgtgggttca taaaagattc aatagcaacc 1080
tggatggtgc ctgcattggt ctctgagaaa caggaagagc aaaaaaattg tctggagtcg 1140
gcttgtaaaa gaaaaatccta ccctatgtgc aaccaaacgt catgggaacc ctteggagga 1200
ggacagttgc catcttatgg gcggttgaca ttacctctag atccaagcac tgacctcaa 1260
cttaacatat cgtttacata cggtcgggtt atactgaatg gagacggtat ggattattat 1320
gaaagcccac tgtcggactc cggatggctt accattcctc caaaaaacgg aacagtcctt 1380
ggattgataa acaaaagcaag tagaggagac cagttcattg taatccccc tgtggtgaca 1440
tttgcgccc gggaaatcaag tgggaattgt tattcaccta tccaacatc ccagattatg 1500
gataaagatg tccttactga gtccaattta gtggtgttgc ctacacagaa ttttagatat 1560
gtcatagcaa catatgatat atcccgggac aatcatgcga tcgtttacta tgtctatgac 1620
ccaattcgga cgatttctta tacgtacca tttagactaa ctaccaaagg tagacctgat 1680
ttcctaagga ttgaatgttt tgtttgggat gatgatttgt ggtgtcacca gttctaccga 1740
ttcagagctg acatcactaa ctctaccacc agtgttgaga atttagtccg tataagattc 1800
tcatgtaacc gtccaagacc ttga 1824

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<210> SEQ ID NO 43

<211> LENGTH: 1824

<212> TYPE: DNA

<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 43

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atgctctcct accgagacaa ggtgggtgcc ttctataagg acaatgctag agctaattca 60
tccaagctgt ccttagtgac agaagagcaa gggggcagga gaccacccta tttgctgttt 120
gtccttctca tcctactggt tggaaatcatg gccttgcttg ctatcaactgg agttcgattt 180
caccaagtat caactagcaa tatggagtgt agcagattgc tgaaagagga tctggagaaa 240
tcagaggcgg tacatcacca agtcatagat gtcttgacgc cgctcttcaa aattattgga 300
gatgagatg ggttacggtt gccacaaaa ctaaacgaga tcaaacatt tatcctcaa 360
aagacaaact tcttcaatcc gaacagggaa ttcgacttcc gcgatctcca ctggtgcatt 420
aaccaccta gtaagatcaa ggtgaatttt actaattact gcgatactat ggggatcaga 480
aatctatg catcggcagc aaatcccac cttttatcag cactctccgg aggtagaggt 540
gacatatcc caccatagc atgcaatgga gctactattt cagtaggcaa gattttcccc 600
ctatcagat cattatctat gtctttgatc tcaagaacat cagagataat caatatgcta 660
accgctatct cagacggagt gtatggtaaa acttatttac taatgcctga ttatattgaa 720
ggggagtctg acacgcaaaa gattcgagtc tttgagatag ggttcatcaa acggtggtctg 780
aatgacatgc cactactcca gacaaccaac tatatggtcc tcccagagaa ttccaaagct 840
aaggtatgta ctatagcagt gggcgagttg acactggctt ctttgtgtgt aggtgagagc 900
accgtattgt tatatcatga cagcaatggt tcacaagatg gtattctagt agtgacgctg 960
ggaatattcg gggcaacatc tatggatcaa gttgaagagg tgatacctgt cgctgaccca 1020
ttagtagaaa aaatacatat aacaaatcac cgcgggatca taaaagattc aatagcaacc 1080
tggatggtgc ctgcattagt ttctgagaaa caagaggaac aaaaaaattg tctggagtca 1140
gcttgtaaaa gaaaaatccta ccctatgtgc aatcaaacgt catgggaacc ctttggagga 1200
ggacagttgc catcttatgg gcggttgaca ttacctctag atccaagcat tgacctcaa 1260

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cttaacatat cattacata cggtcgatt atactgaatg gggacggtat ggattattat 1320
gagagcccac tgttggaactc cggatggcctt accattcctc ccaagaacgg aacagtcctt 1380
ggattgataa acaaagcaag tagaggagac cagttcactg taatccccc tgtgttgaca 1440
tttgcgccc a gggaaatcaag tggaaattgt tatttaccta ttcaaacatc ccagattatg 1500
gataaagatg tccttactga gtccaattta gtggtggtgc ctacacagaa ttttagatat 1560
gtcgtagcaa catatgat atctcgggac gatcatgcca ttgtttatta tgtttatgac 1620
ccaatacgg a cgatttctta tacgtacca tttagactaa ctactaagg tagacctgat 1680
ttcttaagga ttgagtgttt tgtgtgggat gacgatttgt ggtgtcacca gttttaccga 1740
ttcgaggccg acatcaccaa ctctacaacc agtgtcgaga atttagtccg tatgagattc 1800
tcatgtaacc gttccagacc ttga 1824

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<210> SEQ ID NO 44
<211> LENGTH: 607
<212> TYPE: PRT
<213> ORGANISM: canine distemper virus

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```

<400> SEQUENCE: 44

```

```

Met Leu Ser Tyr Gln Asp Lys Val Gly Ala Phe Tyr Lys Asp Asn Ala
1 5 10 15
Arg Ala Asn Ser Ser Lys Leu Ser Leu Val Thr Glu Glu Gln Gly Gly
20 25 30
Arg Arg Pro Pro Tyr Leu Leu Phe Val Leu Leu Ile Leu Leu Val Gly
35 40 45
Ile Leu Ala Leu Leu Ala Ile Thr Gly Val Arg Phe His Gln Val Ser
50 55 60
Thr Ser Asn Val Glu Phe Ser Arg Leu Leu Lys Glu Asp Met Glu Lys
65 70 75 80
Ser Glu Ala Val His His Gln Val Ile Asp Val Leu Thr Pro Leu Phe
85 90 95
Lys Ile Ile Gly Asp Glu Ile Gly Leu Arg Leu Pro Gln Lys Leu Asn
100 105 110
Glu Ile Lys Gln Phe Ile Leu Gln Lys Thr Asn Phe Phe Asn Pro Asn
115 120 125
Arg Glu Phe Asp Phe Arg Asp Leu His Trp Cys Ile Asn Pro Pro Ser
130 135 140
Lys Ile Lys Val Asn Phe Thr Asn Tyr Cys Asp Thr Ile Gly Ile Arg
145 150 155 160
Lys Ser Ile Ala Ser Ala Ala Asn Pro Ile Leu Leu Ser Ala Leu Ser
165 170 175
Gly Gly Arg Gly Asp Ile Phe Pro Pro Tyr Arg Cys Ser Gly Ala Ala
180 185 190
Thr Ser Val Gly Arg Val Phe Pro Leu Ser Val Ser Leu Ser Met Ser
195 200 205
Leu Ile Ser Arg Lys Ser Glu Ile Ile Asn Met Leu Thr Ala Ile Ser
210 215 220
Asn Gly Val Tyr Gly Lys Thr Tyr Leu Leu Val Pro Asp Tyr Ile Glu
225 230 235 240
Glu Glu Phe Asp Thr Gln Lys Ile Arg Val Phe Glu Ile Gly Phe Ile
245 250 255
Lys Arg Trp Leu Asn Asp Met Pro Leu Leu Gln Thr Thr Asn Tyr Met
260 265 270

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Val Leu Pro Glu Asn Ser Lys Ala Lys Val Cys Thr Ile Ala Val Gly
    275                                280                                285

Glu Leu Thr Leu Ala Ser Leu Cys Val Gly Glu Ser Thr Val Ser Leu
    290                                295                                300

Tyr His Asp Ser Asn Gly Ser Gln Asp Ser Ile Leu Ala Val Thr Leu
    305                                310                                315                                320

Gly Ile Phe Gly Ala Thr Ser Met Asp Gln Val Glu Glu Val Ile Pro
    325                                330                                335

Val Ala His Pro Ser Val Glu Lys Ile His Ile Thr Asn His Arg Gly
    340                                345                                350

Phe Ile Lys Asp Ser Ile Ala Thr Trp Met Val Pro Ala Leu Val Ser
    355                                360                                365

Glu Lys Gln Glu Glu Gln Lys Asn Cys Leu Glu Ser Ala Cys Gln Arg
    370                                375                                380

Lys Ser Tyr Pro Met Cys Asn Gln Thr Ser Trp Glu Pro Phe Gly Gly
    385                                390                                395                                400

Gly Gln Leu Pro Ser Tyr Gly Arg Leu Thr Leu Pro Leu Asp Pro Ser
    405                                410                                415

Thr Asp Leu Gln Leu Asn Ile Ser Phe Thr Tyr Gly Pro Val Ile Leu
    420                                425                                430

Asn Gly Asp Gly Met Asp Tyr Tyr Glu Ser Pro Leu Ser Asp Ser Gly
    435                                440                                445

Trp Leu Thr Ile Pro Pro Lys Asn Gly Thr Val Leu Gly Leu Ile Asn
    450                                455                                460

Lys Ala Ser Arg Gly Asp Gln Phe Ile Val Ile Pro His Val Leu Thr
    465                                470                                475                                480

Phe Ala Pro Arg Glu Ser Ser Gly Asn Cys Tyr Leu Pro Ile Gln Thr
    485                                490                                495

Ser Gln Ile Met Asp Lys Asp Val Leu Thr Glu Ser Asn Leu Val Val
    500                                505                                510

Leu Pro Thr Gln Asn Phe Arg Tyr Val Ile Ala Thr Tyr Asp Ile Ser
    515                                520                                525

Arg Asp Asn His Ala Ile Val Tyr Tyr Val Tyr Asp Pro Ile Arg Thr
    530                                535                                540

Ile Ser Tyr Thr Tyr Pro Phe Arg Leu Thr Thr Lys Gly Arg Pro Asp
    545                                550                                555                                560

Phe Leu Arg Ile Glu Cys Phe Val Trp Asp Asp Asp Leu Trp Cys His
    565                                570                                575

Gln Phe Tyr Arg Phe Glu Ala Asp Ile Thr Asn Ser Thr Thr Ser Val
    580                                585                                590

Glu Asn Leu Val Arg Ile Arg Phe Ser Cys Asn Arg Ser Arg Pro
    595                                600                                605

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<210> SEQ ID NO 45

<211> LENGTH: 607

<212> TYPE: PRT

<213> ORGANISM: canine distemper virus

<400> SEQUENCE: 45

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Met Leu Ser Tyr Arg Asp Lys Val Gly Ala Phe Tyr Lys Asp Asn Ala
1      5      10      15

Arg Ala Asn Ser Ser Lys Leu Ser Leu Val Thr Glu Glu Gln Gly Gly
20     25     30

Arg Arg Pro Pro Tyr Leu Leu Phe Val Leu Leu Ile Leu Leu Val Gly
35     40     45

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Ile	Met	Ala	Leu	Leu	Ala	Ile	Thr	Gly	Val	Arg	Phe	His	Gln	Val	Ser
	50					55					60				
Thr	Ser	Asn	Met	Glu	Phe	Ser	Arg	Leu	Leu	Lys	Glu	Asp	Leu	Glu	Lys
65				70						75					80
Ser	Glu	Ala	Val	His	His	Gln	Val	Ile	Asp	Val	Leu	Thr	Pro	Leu	Phe
				85					90					95	
Lys	Ile	Ile	Gly	Asp	Glu	Ile	Gly	Leu	Arg	Leu	Pro	Gln	Lys	Leu	Asn
			100					105					110		
Glu	Ile	Lys	Gln	Phe	Ile	Leu	Gln	Lys	Thr	Asn	Phe	Phe	Asn	Pro	Asn
		115					120					125			
Arg	Glu	Phe	Asp	Phe	Arg	Asp	Leu	His	Trp	Cys	Ile	Asn	Pro	Pro	Ser
	130					135					140				
Lys	Ile	Lys	Val	Asn	Phe	Thr	Asn	Tyr	Cys	Asp	Thr	Met	Gly	Ile	Arg
145				150						155					160
Lys	Ser	Ile	Ala	Ser	Ala	Ala	Asn	Pro	Ile	Leu	Leu	Ser	Ala	Leu	Ser
				165					170					175	
Gly	Gly	Arg	Gly	Asp	Ile	Phe	Pro	Pro	Tyr	Arg	Cys	Asn	Gly	Ala	Thr
			180						185					190	
Ile	Ser	Val	Gly	Lys	Ile	Phe	Pro	Leu	Ser	Val	Ser	Leu	Ser	Met	Ser
			195				200					205			
Leu	Ile	Ser	Arg	Thr	Ser	Glu	Ile	Ile	Asn	Met	Leu	Thr	Ala	Ile	Ser
	210					215					220				
Asp	Gly	Val	Tyr	Gly	Lys	Thr	Tyr	Leu	Leu	Met	Pro	Asp	Tyr	Ile	Glu
225					230					235					240
Gly	Glu	Phe	Asp	Thr	Gln	Lys	Ile	Arg	Val	Phe	Glu	Ile	Gly	Phe	Ile
				245					250					255	
Lys	Arg	Trp	Leu	Asn	Asp	Met	Pro	Leu	Leu	Gln	Thr	Thr	Asn	Tyr	Met
			260					265						270	
Val	Leu	Pro	Glu	Asn	Ser	Lys	Ala	Lys	Val	Cys	Thr	Ile	Ala	Val	Gly
			275				280						285		
Glu	Leu	Thr	Leu	Ala	Ser	Leu	Cys	Val	Gly	Glu	Ser	Thr	Val	Leu	Leu
	290					295					300				
Tyr	His	Asp	Ser	Asn	Gly	Ser	Gln	Asp	Gly	Ile	Leu	Val	Val	Thr	Leu
305				310						315					320
Gly	Ile	Phe	Gly	Ala	Thr	Ser	Met	Asp	Gln	Val	Glu	Glu	Val	Ile	Pro
				325					330					335	
Val	Ala	Asp	Pro	Leu	Val	Glu	Lys	Ile	His	Ile	Thr	Asn	His	Arg	Gly
				340				345						350	
Ile	Ile	Lys	Asp	Ser	Ile	Ala	Thr	Trp	Met	Val	Pro	Ala	Leu	Val	Ser
		355					360						365		
Glu	Lys	Gln	Glu	Glu	Gln	Lys	Asn	Cys	Leu	Glu	Ser	Ala	Cys	Gln	Arg
	370					375						380			
Lys	Ser	Tyr	Pro	Met	Cys	Asn	Gln	Thr	Ser	Trp	Glu	Pro	Phe	Gly	Gly
385					390					395					400
Gly	Gln	Leu	Pro	Ser	Tyr	Gly	Arg	Leu	Thr	Leu	Pro	Leu	Asp	Pro	Ser
				405					410					415	
Ile	Asp	Leu	Gln	Leu	Asn	Ile	Ser	Phe	Thr	Tyr	Gly	Pro	Ile	Ile	Leu
			420					425					430		
Asn	Gly	Asp	Gly	Met	Asp	Tyr	Tyr	Glu	Ser	Pro	Leu	Leu	Asp	Ser	Gly
		435					440						445		
Trp	Leu	Thr	Ile	Pro	Pro	Lys	Asn	Gly	Thr	Val	Leu	Gly	Leu	Ile	Asn
	450					455					460				
Lys	Ala	Ser	Arg	Gly	Asp	Gln	Phe	Thr	Val	Ile	Pro	His	Val	Leu	Thr
465					470					475					480

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Phe	Ala	Pro	Arg	Glu	Ser	Ser	Gly	Asn	Cys	Tyr	Leu	Pro	Ile	Gln	Thr
				485					490					495	
Ser	Gln	Ile	Met	Asp	Lys	Asp	Val	Leu	Thr	Glu	Ser	Asn	Leu	Val	Val
			500					505					510		
Leu	Pro	Thr	Gln	Asn	Phe	Arg	Tyr	Val	Val	Ala	Thr	Tyr	Asp	Ile	Ser
		515					520					525			
Arg	Asp	Asp	His	Ala	Ile	Val	Tyr	Tyr	Val	Tyr	Asp	Pro	Ile	Arg	Thr
	530					535					540				
Ile	Ser	Tyr	Thr	Tyr	Pro	Phe	Arg	Leu	Thr	Thr	Lys	Gly	Arg	Pro	Asp
545					550					555					560
Phe	Leu	Arg	Ile	Glu	Cys	Phe	Val	Trp	Asp	Asp	Asp	Leu	Trp	Cys	His
				565					570					575	
Gln	Phe	Tyr	Arg	Phe	Glu	Ala	Asp	Ile	Thr	Asn	Ser	Thr	Thr	Ser	Val
			580					585						590	
Glu	Asn	Leu	Val	Arg	Met	Arg	Phe	Ser	Cys	Asn	Arg	Ser	Arg	Pro	
		595					600					605			

The invention claimed is:

1. An isolated canine distemper virus (CDV) of European wildlife (EW) lineage comprising the characteristics of CDV 9041474B CDV-EW (ATCC Deposit No. PTA-10596).

2. An attenuated strain of CDV isolated in cell culture in which the CDV strain of claim 1 or a progeny strain thereof has been propagated.

3. An immunogenic composition, comprising the isolated CDV of claim 1, or progeny thereof.

4. A method of eliciting an immune response to canine distemper virus in a subject in need thereof, comprising the step of

administering to said subject the immunogenic composition of claim 3.

5. An isolated nucleic acid molecule encoding the amino acid sequence of SEQ ID NO: 44, or the complement thereof.

6. The isolated nucleic acid molecule of claim 5 that comprises the nucleotide sequence of SEQ ID NO: 42, or the complement thereof.

7. An isolated polypeptide comprising the amino acid sequence of SEQ ID NO: 44.

8. An isolated canine distemper virus (CDV) of European wildlife (EW) lineage encoding a polypeptide comprising the amino acid sequence of SEQ ID NO: 44.

9. An isolated canine distemper virus (CDV) of American-2 (AM-2) lineage having the characteristics of CDV 08021509 CDV-AM-2 (ATCC Deposit No. PTA-10597).

10. An attenuated strain of CDV isolated in cell culture in which the CDV strain of claim 9 or a progeny strain thereof has been propagated.

11. An immunogenic composition, comprising the isolated CDV of claim 9, or progeny thereof.

12. A method of eliciting an immune response to canine distemper virus in a subject in need thereof, comprising the step of

administering to said subject the immunogenic composition of claim 11.

13. An isolated nucleic acid molecule encoding the amino acid sequence of SEQ ID NO: 45, or the complement thereof.

14. The isolated nucleic acid molecule of claim 13 that comprises the nucleotide sequence of SEQ ID NO: 43, or the complement thereof.

15. An isolated polypeptide comprising the amino acid sequence of SEQ ID NO: 45.

16. An isolated canine distemper virus (CDV) of European wildlife (EW) lineage encoding a polypeptide comprising the amino acid sequence of SEQ ID NO: 45.

* * * * *