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(54) **Title:** EUKARYOTIC CELLS COMPRISING ADENOVIRUS-ASSOCIATED VIRUS POLYNUCLEOTIDES

(57) **Abstract:** The present inventions provide eukaryotic cells, such as mammalian cells, that comprise adeno-associated virus (AAV) polynucleotides, including AAV capsid proteins (Cap), and are capable of expressing the polypeptides encoded by the AAV polynucleotides, and thereby are capable of producing AAV, including recombinant AAV. The eukaryotic cells also may comprise adenovirus (Ad) polynucleotides. The present inventions also provide methods of expressing AAV polynucleotides, as well as Ad polynucleotides, in eukaryotic cells, such as CHO cells, HEK 293 and BHK cells. The present inventions further provides other products and methods described herein.

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EUKARYOTIC CELLS COMPRISING ADENOVIRUS-ASSOCIATED VIRUS POLYNUCLEOTIDES

This Application claims priority to U.S. Application Serial No. 63/256,730, filed October 18, 2021, which is hereby incorporated by reference in its entirety.

FIELD OF THE INVENTIONS

[0001] The present inventions provide eukaryotic cells that comprise adeno-associated virus (AAV) polynucleotides, including AAV capsid proteins (Cap). The cells are capable of expressing the polypeptides encoded by the AAV polynucleotides, and thereby are capable of producing AAV, including recombinant AAV. The eukaryotic cells also may comprise adenovirus (Ad) polynucleotides. The present inventions also provide methods of expressing AAV polynucleotides, as well as Ad polynucleotides, in eukaryotic cells. The present inventions also provide methods for producing recombinant adeno-associated virus utilizing eukaryotic cells that express AAV and Ad polypeptides encoded by polynucleotides, as well as recombinant AAV produced by these inventive methods. The present inventions further provide other products and methods described herein.

REFERENCE TO ELECTRONIC SEQUENCE LISTING

[0002] The application contains a Sequence Listing, which has been submitted electronically in .XML format and is hereby incorporated by reference in its entirety. Said .XML copy, created on October 5, 2022 , is named “135975-61702.xml” and is 229,879 bytes in size. The sequence listing contained in this .XML file is part of the specification and is hereby incorporated by reference herein in its entirety.

BACKGROUND OF THE INVENTIONS

[0003] Adeno-associated virus (AAV) is a non-enveloped, single-stranded DNA virus and is used as a gene delivery vector for both research and therapeutics. Weitzman and Linden, Adeno-Associated Virus Biology (chapter 1), *Meth. Molec. Biol.* 807: 1-23 (2011). Gene transfer vectors based on AAV have demonstrated promise for human gene therapy based on their safety profile and potential to achieve long-term efficacy in animal models. Wang *et al., Nature*, 18: 358-78 (2019). A major challenge for advancing AAV-based therapies into clinical development is the difficulty and cost of producing sufficient quantities of AAV through transient methodologies.

[0004] AAV has been produced in HEK 293, BHK, human amniotic (for example, epithelial cells such as HAEpiC) and SF9 lines. However, expression is transient due to the use of plasmid vectors containing the necessary AAV and helper virus genes. For example, recombinant AAV production in HEK 293 cells using adenovirus helper gene products utilizes adenovirus E2A, E4, VA RNA and AAV Rep and Cap, along with AAV inverted terminal repeats (ITR) flanking the polynucleotide of interest. The reliance on non-integrated plasmid vectors means that the requisite gene products will be lost over time and need to be continually re-established.

[0005] The AAV genome includes a capsid gene referred to as “Cap” or “CAP”. Cap in nature is translated to produce, via alternative start codons and transcript splicing, three size-variant structural proteins referred to as VP1 (about 90kDa), VP2 (about 72kDa) and VP3 (about 60kDa). An AAV capsid contains 60 subunits total of the VP proteins. A ratio of 1:1:10 is considered the most typical

ratio for VP1:VP2:VP3, which is a stoichiometry of 5 VP1 subunits:5 VP2 subunits:50 VP3 subunits. However, there can be variation. Wörner *et al., Nature Communications* 12:1642 (2021). AAV polynucleotides and proteins, including CAP, can be selected from any serotype.

[0006] Thus, there exists the need to develop improved cells and production methods that avoid the transient nature of non-integrated plasmid vectors.

SUMMARY OF THE INVENTIONS

[0007] The present inventions provide stable eukaryotic cells, such as mammalian cells (for example, primate, rodent and canine cells), comprising integrated AAV polynucleotides and Ad polynucleotides. The protein and VA RNA products of the Ad polynucleotides act in a helper capacity. All AAV and Ad types are amenable for use according to the present inventions. The present inventions advantageously can employ site-specific integration into the cell genome, which refers to pre-selected genomic sites for exogenous DNA to be inserted into a cellular genome. Random insertion can be employed as well.

[0008] Herein described are polynucleotides, where each can comprise (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding Adeno-associated virus (AAV) Cap protein, and (v) a polyadenylation site. For example, the polynucleotide can be in a CHO cell and have the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a CHO cell genome, such as a CHO chromosome. Alternatively, the polynucleotide can be

in a HEK 293 cell and have the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein operably linked. The polynucleotide can be integrated into a HEK 293 cell genome, such as a HEK 293 cell chromosome. Another alternative is the polynucleotide can be in a BHK cell and have the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein operably linked. The polynucleotide can be integrated into a BHK cell genome, such as a BHK cell chromosome. In yet another alternative is the polynucleotide can be in a human amniotic cell and have the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein operably linked. The polynucleotide can be integrated into a human amniotic cell genome, such as a human amniotic cell chromosome. Additionally, the polynucleotide can be integrated into non-chromosomal locations as known by the person skilled in the art, such as episomes.

[0009] The polynucleotide can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. AAV proteins and polynucleotides, including CAP, can be selected from any serotype. When Cap is from serotype 5 (“Cap5”) and expressed in a CHO cell, the polynucleotide allows for production of AAV Cap5 VP2 and VP3 protein, wherein the amount of VP3 produced is greater than the amount of VP2 produced. Production of VP1 can be less than 1% the level of VP2 production. Production ratios can vary based upon experimental conditions and analytical techniques.

[0010] Additionally, polynucleotides are described, wherein each can comprise (i) a promoter, (ii) an intron, (iii) a first internal ribosome entry site, (iv) a first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry

site, (vi) a second polynucleotide encoding AAV Cap protein, and (vii) a polyadenylation site. For example, the polynucleotide can be in a CHO cell and the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a CHO cell genome. Alternatively, the polynucleotide can be in a HEK 293 cell and the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked. The polynucleotide can be integrated into a HEK 293 cell genome. In another alternative, the polynucleotide can be in a BHK cell and the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked. The polynucleotide can be integrated into a BHK cell genome. In another alternative, the polynucleotide can be in a human amniotic cell and the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked. The polynucleotide can be integrated into a human amniotic cell genome. The polynucleotide can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. AAV proteins and polynucleotides, including CAP, can be selected from any serotype. When Cap is from serotype 5 (“Cap5”) and expressed in a CHO cell, the polynucleotide allows for production of AAV Cap5 proteins VP1, VP2 and VP3. The

amount of VP3 production can be greater than the amount of VP1 production and the amount of VP2 production. The amount of VP2 production can be greater than the amount of VP1 production. Production ratios can vary based upon experimental conditions and analytical techniques.

[0011] Moreover, there are described polynucleotides, wherein each polynucleotide can comprise (i) a promoter, (ii) an intron, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site, wherein the polynucleotide allows for production of AAV Cap VP1 protein when expressed. For example, the polynucleotide can be in a CHO cell and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a CHO cell genome. Alternatively, the polynucleotide can be in a HEK 293 cell and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a HEK 293 cell genome. In another alternative, the polynucleotide can be in a BHK cell and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a BHK cell genome. In still another alternative, the polynucleotide can be in a human amniotic cell and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a human amniotic cell genome. The polynucleotide can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator.

[0012] Furthermore, there are described polynucleotides, wherein each can comprise (i) a promoter, (ii) an internal ribosome binding site, (iii) a

polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site, wherein the polynucleotide allows for production of AAV VP1 Cap protein when expressed. The polynucleotide can be in a CHO cell and the (i) promoter, (ii) internal ribosome binding site, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a CHO genome. Alternatively, the polynucleotide can be in a HEK 293 cell and the (i) promoter, (ii) internal ribosome binding site, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a HEK 293 cell genome. In another alternative, the polynucleotide can be in a BHK cell and the (i) promoter, (ii) internal ribosome binding site, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a BHK cell genome. In yet another alternative, the polynucleotide can be in a human amniotic cell and the (i) promoter, (ii) internal ribosome binding site, and (iii) polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a human amniotic cell genome. The polynucleotide can comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator.

[0013] Additionally, there are described eukaryotic cells, where each cell can comprise a polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a first internal ribosome entry site, (iv) a first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, (vi) a second polynucleotide encoding AAV Cap protein, and (vii) a polyadenylation site. The (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV

Cap protein can be operably linked. The polynucleotide can be integrated into a cell genome. The cell can be a CHO cell, a HEK 293 cell, a BHK cell, a Human Amniotic Cell or other eukaryotic cell. The cell can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. The cell can further comprise: a polynucleotide encoding AAV Rep, a polynucleotide encoding Ad E1A, a polynucleotide encoding Ad E1B, a polynucleotide encoding Ad E2A or E2A orf, a polynucleotide encoding Ad E4 or E4 orf 6, a polynucleotide encoding VA RNA, and a polynucleotide encoding AAV ITRs and a protein of interest.

[0014] There are also described eukaryotic cells, where each cell can comprise (A) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and (B) a second polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site. The (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (A) first polynucleotide can be operably linked, and wherein the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein of (B) second polynucleotide can be operably linked. The cell can have at least one polynucleotide integrated into a cell genome. The cell can be a CHO cell, a HEK 293 cell, a BHK cell, a human amniotic cell or other eukaryotic cell. The cell can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. The cell can further comprise a polynucleotide encoding AAV Rep, a polynucleotide encoding Ad E1A, a polynucleotide encoding Ad E1B, a polynucleotide encoding Ad E2A or E2A orf, a

polynucleotide encoding E4 or E4 orf 6, a polynucleotide encoding VA RNA, and a polynucleotide encoding AAV ITRs and a protein of interest.

[0015] Additionally, there are described eukaryotic cells, where each cell can comprise: (A) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and (B) a second polynucleotide comprising (i) a promoter, (ii) an internal ribosome entry site, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site. The cells can have the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (A) first polynucleotide operably linked, and the (i) promoter, (ii) internal ribosome entry site, and (iii) polynucleotide encoding AAV Cap protein of (B) second polynucleotide operably linked. At least one polynucleotide can be integrated into a cell genome. The cell can be a CHO cell, a HEK 293 cell, a BHK cell, a human amniotic cell or other eukaryotic cell. The cell can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. The cell can further comprise: a polynucleotide encoding AAV Rep, a polynucleotide encoding Ad E1A, a polynucleotide encoding Ad E1B, a polynucleotide encoding Ad E2A or E2A orf, a polynucleotide encoding Ad E4 or E4 orf 6, a polynucleotide encoding VA RNA, and a polynucleotide encoding AAV ITRs and a protein of interest.

[0016] There also are described cell cultures comprising any of the above cells in any type of media, including growth media and maintenance media. Additionally, there are described methods of producing AAV proteins, including Cap proteins, and methods that can result in the production of recombinant AAV.

[0017] There are described methods of producing adeno-associated virus (AAV) Cap protein in cell culture, wherein a method comprises the steps of: providing eukaryotic cells, wherein a cell comprises a polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a first internal ribosome entry site, (iv) a first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, (vi) a second polynucleotide encoding AAV Cap protein, and (vii) a polyadenylation site; and culturing the cells in a culture medium to allow the cells to produce AAV Cap protein, wherein the polynucleotide allows for production of AAV Cap proteins VP1, VP2 and VP3. The (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein can be operably linked. The polynucleotide can be integrated into a cell genome. The cell can be a CHO cell, a HEK 293 cell, a BHK cell, a human amniotic cell or other eukaryotic cell. The cell can further comprise an operator. The cell can further comprise: a polynucleotide encoding AAV Rep, a polynucleotide encoding Ad E1A, a polynucleotide encoding Ad E1B, a polynucleotide encoding Ad E2A or E2A orf, a polynucleotide encoding Ad E4 or E4 orf 6, a polynucleotide encoding VA RNA, and a polynucleotide encoding AAV ITRs and a protein of interest, wherein the cell can produce recombinant AAV.

[0018] Also described are methods of producing adeno-associated virus (AAV) Cap protein in cell culture, wherein a method comprises the steps of providing eukaryotic cells, where a cell comprises (a) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and (b)

a second polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site; and culturing the cells in a culture medium to allow the cells to produce AAV Cap protein, wherein the polynucleotide allows for production of AAV Cap proteins VP1, VP2 and VP3. The (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (a) first polynucleotide can be operably linked, and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein of (b) second polynucleotide can be operably linked. The polynucleotide can be integrated into a cell genome. The cell can be a CHO cell, a HEK 293 cell, a BHK cell, a human amniotic cell or other eukaryotic cell. The cell can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. The cell can further comprise: a polynucleotide encoding AAV Rep, a polynucleotide encoding Ad E1A, a polynucleotide encoding Ad E1B, a polynucleotide encoding Ad E2A or E2A orf, a polynucleotide encoding Ad E4 or E4 orf 6, a polynucleotide encoding VA RNA, and a polynucleotide encoding AAV ITRs and a protein of interest, wherein the cell can produce recombinant AAV.

[0019] There are also described methods of producing adeno-associated virus (AAV) Cap protein in cell culture, wherein a method comprises the steps of: providing eukaryotic cells, where a cell comprises (a) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and (b) a second polynucleotide comprising (i) a promoter, (ii) an internal ribosome entry site, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site; and culturing the cells in a culture medium to allow the cells to produce AAV Cap

protein, wherein the polynucleotide allows for production of AAV Cap proteins VP1, VP2 and VP3. The (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (a) first polynucleotide can be operably linked, and the (i) promoter, (ii) internal ribosome entry site, and (iii) polynucleotide encoding AAV Cap protein of (b) second polynucleotide can be operably linked. The polynucleotide can be integrated into a cell genome. The cell can be a CHO cell, a HEK 293 cell, a BHK cell, a human amniotic cell or other eukaryotic cell. The cell can further comprise an operator. The promoter can be a CMV promoter and the operator can be a Tet operator. The cell can further comprise: a polynucleotide encoding AAV Rep, a polynucleotide encoding Ad E1A, a polynucleotide encoding Ad E1B, a polynucleotide encoding Ad E2A or E2A orf, a polynucleotide encoding Ad E4 or E4 orf 6, a polynucleotide encoding VA RNA, and a polynucleotide encoding AAV ITRs and a protein of interest, wherein the cell can produce recombinant AAV.

BRIEF DESCRIPTION OF THE FIGURES

[0020] **FIGURE 1** is a schematic diagram of a polynucleotide comprising a promoter, an intron, an internal ribosome entry site (IRES), a polynucleotide encoding AAV Cap protein and a polyadenylation site (poly A).

[0021] **FIGURE 2** is a schematic diagram of a polynucleotide comprising a promoter, an intron, two internal ribosome entry sites, two polynucleotides encoding AAV Cap protein and a polyadenylation site.

[0022] **FIGURE 3** is a schematic diagram of a polynucleotide comprising a promoter, an intron, a polynucleotide encoding AAV Cap protein and a polyadenylation site.

[0023] **FIGURE 4** is a schematic diagram of a polynucleotide comprising a promoter, an internal ribosome entry site, a polynucleotide encoding AAV Cap protein and a polyadenylation site.

[0024] **FIGURE 5** is a schematic diagram of a polynucleotide comprising a promoter with operator (Op), an intron, an internal ribosome entry site, a polynucleotide encoding AAV Cap protein and a polyadenylation site.

[0025] **FIGURE 6** is a schematic diagram of a polynucleotide comprising a promoter with operator (Op), an intron, two internal ribosome entry sites, two polynucleotides encoding AAV Cap protein and a polyadenylation site.

[0026] **FIGURE 7** is a schematic diagram of a polynucleotide comprising a promoter with operator (Op), an intron, a polynucleotide encoding AAV Cap protein and a polyadenylation site.

[0027] **FIGURE 8** is a schematic diagram of a polynucleotide comprising a promoter with operator (Op), an internal ribosome entry site, a polynucleotide encoding AAV Cap protein and a polyadenylation site.

DETAILED DESCRIPTION OF THE INVENTIONS

[0028] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which these inventions belong.

Definitions

[0029] The term “about” in the context of numerical values and ranges refers to values or ranges that approximate or are close to the recited values or ranges such that the inventions can perform, such as having a sought rate, amount,

degree, increase, decrease, or extent of expression, concentration, or time, as is apparent from the teachings contained herein. Thus, this term encompasses values beyond those simply resulting from systematic error. For example, “about” can signify values either above or below the stated value in a range of approx. +/- 10% or more or less depending on the ability to perform.

[0030] “Intron” is a section of DNA located between exons. An intron is removed to form a mature messenger RNA. Preferred introns are those that can affect the starting point of translation, and exemplars are the hCMV-IE intron (Human cytomegalovirus immediate early protein) and FMDV intron (Foot and Mouth Disease Virus). The globin gene intron also has been reportedly used for expression.

[0031] A “nucleic acid moiety” includes any arrangement of single stranded or double stranded nucleotide sequences. Nucleic acid moieties can include, but are not limited to, polynucleotides, promoters, enhancers, operators, repressors, transcription termination signals, ribosomal entry sites and polyadenylation signals.

[0032] A “DNA cassette” or “cassette” is a type of nucleic acid moiety that comprises at least a promoter, at least one open reading frame and optionally a polyadenylation signal, for example an SV40 polyadenylation signal. Other nucleic acid moieties, such as operators, also are optional. A DNA cassette thus is a polynucleotide that comprises two or more shorter polynucleotides.

[0033] "Operably linked" refers to one or more nucleotide sequences in functional relationships with one or more other nucleotide sequences. Such functional relationships can directly or indirectly control, cause, regulate, enhance,

facilitate, permit, attenuate, repress or block an action or activity in accordance with the selected design. Exemplars include single-stranded or double-stranded nucleic acid moieties, and can comprise two or more nucleotide sequences arranged within a given moiety in such a way that sequence(s) can exert at least one functional effect on other(s). For example, a promoter operably linked to the coding region of a DNA polynucleotide sequence can facilitate transcription of the coding region. Other elements, such as enhancers, operators, repressors, transcription termination signals, ribosomal entry sites and polyadenylation signals also can be operably linked with a polynucleotide of interest to control its expression. Arrangements and spacing to achieve operable linkages can be ascertained by approaches available to the person skilled in the art, such as screening using western blots and RT-PCR.

[0034] "Operator" indicates a DNA sequence that is introduced in or near a polynucleotide sequence in such a way that the polynucleotide sequence may be regulated by the interaction of a molecule capable of binding to the operator and, as a result, prevent or allow transcription of the polynucleotide sequence, as the case may be. One skilled in the art will recognize that the operator must be located sufficiently in proximity to the promoter such that it is capable of controlling or influencing transcription by the promoter, which can be considered a type of operable linkage. The operator may be placed either downstream or upstream of the promoter. These include, but are not limited to, the operator region of the Lex A gene of *E. coli*, which binds the Lex A peptide and the lactose and 45 tryptophan operators, which bind the repressor proteins encoded by the Lad and trpR genes of *E. coli*. The bacteriophage operators from the lambda Pi and the phage P22 Mnt and Arc. Preferred operators are the Tet (tetracycline) operator and the Arc

operator. Operators can have a native sequence or a mutant sequence. For example, mutant sequences of the Tet operator are disclosed in Wissmann *et al., Nucleic Acids Res.* 14: 4253-66 (1986).

[0035] The phrases "percent identity" or "% identical," in their various grammatical forms, when describing a sequence is meant to include homologous sequences that display the recited identity along regions of contiguous homology, but the presence of gaps, deletions, or insertions that have no homolog in the compared sequence are not taken into account in calculating percent identity. As used herein, a "percent identity" or "% identical" determination between homologs would not include a comparison of sequences where the homolog has no homologous sequence to compare in an alignment. Thus, "percent identity" and "% identical" do not include penalties for gaps, deletions, and insertions.

[0036] A "homologous sequence" in the context of nucleic acid sequences refers to a sequence that is substantially homologous to a reference nucleic acid sequence. In some embodiments, two sequences are considered to be substantially homologous if at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more of their corresponding nucleotides are identical over a relevant stretch of residues. In some embodiments, the relevant stretch is a complete (i.e., full) sequence.

[0037] "Polynucleotide" includes a sequence of nucleotides covalently joined, and includes RNA and DNA. Oligonucleotides are considered shorter polynucleotides. Genes are DNA polynucleotides (polydeoxyribonucleic acid) that ultimately encode polypeptides, which are translated from RNA (polyribonucleic acid) that was typically transcribed from DNA. DNA polynucleotides also can

encode RNA polynucleotides that is not translated, but rather function as RNA "products". The type of polynucleotide (that is, DNA or RNA) is apparent from the context of the usage of the term. A polynucleotide referred to or identified by the polypeptide it encodes sets forth and covers all suitable sequences in accordance with codon degeneracy. Polynucleotides, including those disclosed herein, include percent identity sequences and homologous sequences when indicated.

[0038] "Polypeptide" or "peptide" refers to sequence(s) of amino acids covalently joined. Polypeptides include natural, semi-synthetic and synthetic proteins and protein fragments. "Polypeptide" and "protein" can be used interchangeably. Oligopeptides are considered shorter polypeptides.

[0039] "Protein of interest" or "polypeptide of interest" can have any amino acid sequence, and includes any protein, polypeptide, or peptide, and derivatives, components, domains, chains and fragments thereof. Included are, but not limited to, viral proteins, bacterial proteins, fungal proteins, plant proteins and animal (including human) proteins. Protein types can include, but are not limited to, antibodies, bi-specific antibodies, multi-specific antibodies, antibody chains (including heavy and light), antibody fragments, Fv fragments, Fc fragments, Fc-containing proteins, Fc-fusion proteins, receptor Fc-fusion proteins, receptors, receptor domains, trap and mini-trap proteins, enzymes, factors, repressors, activators, ligands, reporter proteins, selection proteins, protein hormones, protein toxins, structural proteins, storage proteins, transport proteins, neurotransmitters and contractile proteins. Derivatives, components, chains and fragments of the above also are included. The sequences can be natural, semi-synthetic or synthetic. Proteins of interest and polypeptides of interest are encoded by "genes

of interest," which also can be referred to as "polynucleotides of interest." Where multiple genes (same or different) are integrated, they can be referred to as "first," "second", "third," "fourth," "fifth," "sixth," "seventh," "eighth," "ninth," "tenth," etc. as is apparent from the context of use.

[0040] "Promoter" indicates a DNA sequence that cause transcription of a DNA sequence to which it is operably linked, i.e., linked in such a way as to permit transcription of the nucleotide sequence of interest when the appropriate signals are present and repressors are absent. The expression of a polynucleotide of interest may be placed under control of any promoter or enhancer element known in the art. A eukaryotic promoter can be operably linked to a TATA Box. The TATA Box is typically located upstream of the transcription start site.

[0041] Useful promoters that may be used include, but are not limited to, the SV40 early promoter region, SV40 E/L (early late) promoter, the promoter contained in the 3' long terminal repeat of Rous sarcoma virus, the regulatory sequences of the metallothionein gene, mouse or human cytomegalovirus major immediate early (CMV-MIE) promoter and other CMV promoters, including CMVmin promoters. Plant expression vectors comprising the nopaline synthetase promoter region, the cauliflower mosaic virus 35S RNA promoter, and the promoter of the photosynthetic enzyme ribulose biphosphate carboxylase; promoter elements from yeast or other fungi such as the Gal 4 promoter, the ADC (alcohol dehydrogenase) promoter, PGK (phosphoglycerol kinase) promoter, alkaline phosphatase promoter, and the following animal transcriptional control regions, which exhibit tissue specificity and have been utilized in transgenic animals: elastase I; insulin; immunoglobulin; mouse mammary tumor virus; albumin; C.-feto protein; C.1-

antitrypsin; 3-globin, and myosin light chain-2. Various forms of the CMV promoter can be used according to the inventions.

[0042] Minimal promoters, such as CMVmin promoters, can be truncated promoters or core promoters and are preferred for use in controlled expression systems. Minimal promoters and development approaches are widely known and disclosed in, for example, Saxena *et al., Methods Molec. Biol.* 1651:263-73 (2017); Ede *et al., ACS Synth Biol.* 5:395-404 (2016); Brown *et al., Biotech Bioeng.* 111:1638-47 (2014); Morita *et al., Biotechniques* 0:1-5 (2012); Lagrange *et al., Genes Dev.* 12:34-44 (1998). There are many CMVmin promoters described in the field.

[0043] "Reporter proteins" as used herein, refers to any protein capable of generating a detectable signal. Reporter proteins typically fluoresce, or catalyze a colorimetric or fluorescent reaction, and often are referred to as "fluorescent proteins" or "color proteins." However, a reporter protein also can be non-enzymatic and non-fluorescent as long as it can be detected by another protein or moiety, such as a cell surface protein detected with a fluorescent ligand. A reporter protein also can be an inactive protein that is made functional through interaction with another protein that is fluorescent or catalyzes a reaction. Accordingly, any suitable reporter protein, as understood by one of skill in the art, could be used. In some aspects, the reporter protein may be selected from fluorescent protein, luciferase, alkaline phosphatase, β-galactosidase, β-lactamase, dihydrofolate reductase, ubiquitin, and variants thereof. Fluorescent proteins are useful for the recognition of gene cassettes that have or have not been successfully inserted and/or replaced, as the case may be. Fluid cytometry and fluorescence–activated cell sorting are

suitable for detection. Examples of fluorescent proteins are well-known in the art, including, but not limited to Discosoma coral (DsRed), green fluorescent protein (GFP), enhanced green fluorescent protein (eGFP), cyano fluorescent protein (CFP), enhanced cyano fluorescent protein (eCFP), yellow fluorescent protein (YFP), enhanced yellow fluorescent protein (eYFP) and far-red fluorescent protein (e.g. mKate, mKate2, mPlum, mRaspberry or E2-crimson. See, for example, U.S. Patent Nos. 9,816,110. Reporter proteins are encoded by polynucleotides, and are referred to herein as "reporter genes" or "reporter protein genes." Reporters can be considered a type of marker. "Color" or "fluorescent," in their various grammatical forms, also can be used the more specifically refer to a reporter protein or gene..

[0044] A " repressor protein", also referred to as a "repressor," is a protein that can bind to DNA in order to repressor transcription. Repressors are of eukaryotic and prokaryotic origin. Prokaryotic repressors are preferred. Examples of repressor families include: TetR, LysR, LacI, ArsR, IcIR, MerR, AsnC, MarR, DeoR, GntR and Crp families. Repressor proteins in the TetR family include: ArcR, ActII, AmeR, AmrR, ArpR, BpeR, EnvR, EthR, HemR, HydR, IfeR, LanK, LfrR, LmrA, MtrR, Pip, PqrA, QacR, RifQ, RmrR, SimReg2, SmeT, SrpR, TcmR, TetR, TtgR, TrgW, UrdK, VarR YdeS, ArpA., BarA, Aur1B, CalR1, CprB, FarA, JadR*, JadR2, MphB, NonG, PhlF, TylQ, VanT, TarA, TylP, BM1P1, Bm3R1, ButR, CampR, CamR, DhaR, KstR, LexA-like, AcnR, PaaRR, PsbI, Th1R, UidR, YDH1, BetI, McbR, MphR, PhaD, Q9ZF45, TtK, Yhgd, YixD, CasR, IcaR, LitR, LuxR, LuxT, OpaR, Orf2, SmcR, HapR, Ef0113, HlyIIR, BarB, ScbR, MmfR, AmtR, PsrA andYjdC proteins See Ramos *et al., Microbiol. Mol. Biol. Rev.,* 69: 326-56 (2005). Still other repressors

include PurR, LacR, MetJ and PadR, Repressor proteins are encoded by genes referred to as “repressor genes” or “repressor protein genes.”

[0045] “Selectable” or “selection” marker proteins include proteins conferring certain traits, including but not limited to drug resistance or other selective advantages. Selection markers can give the cell receiving the selectable marker gene resistance towards a certain toxin, drug, antibiotic or other compound and permit the cell to produce protein and propagate in the presence of the toxin, drug, antibiotic or other compound, and are often referred to as “positive selectable markers.” Suitable examples of antibiotic resistance markers include, but are not limited to, proteins that impart resistance to various antibiotics, such as kanamycin, spectinomycin, neomycin, gentamycin (G418), ampicillin, tetracycline, chloramphenicol, puromycin, hygromycin, zeocin, and/or blasticidin. There are other selectable markers, often referred to as ”negative selectable markers,” which cause a cell to stop propagating, stop protein production and/or are lethal to the cell in the presence of the negative selectable marker proteins. Thymidine kinase and certain fusion proteins can serve as negative selectable markers, including but not limited to GyrB-PKR. See White *et al., Biotechniques*, 50: 303-309 (May 2011). Selectable marker proteins and corresponding genes can be referred to generically as first (1), second (2), third (3), fourth (4), fifth (5), sixth (6), seventh (7), eighth (8), ninth (9), tenth (10), etc., as is apparent from the context of usage..

[0046] All numerical limits and ranges set forth herein include all numbers or values thereabout or there between of the numbers of the range or limit. The ranges and limits described herein expressly denominate and set forth all integers, decimals and fractional values defined and encompassed by the range or limit.

Description

[0047] The inventions provide cells comprising AAV and optionally Ad polynucleotide sequences to allow production of recombinant AAV comprising a polynucleotide of interest, such as a gene or other sequence encoding a polypeptide of interest. The AAV and Ad polynucleotides provide the requisite structural and helper products required for AAV production.

[0048] AAV polynucleotides, and optionally Ad polynucleotides, can be integrated using a recombinase-mediated cassette exchange (RMCE), for example. “Stable” in the context of cell integration refers to a polynucleotide of interest, such as a gene, introduced into the genome of a cell and can be passed to subsequent generations of cells, and thereby can provide cell lines that are genetically homogeneous for a period of time.

[0049] Cells that are suitable for use with the inventions can be readily selected by those of skill in the art. In some embodiments the cell line is a eukaryotic cell line such as a yeast cell line, insect cell line (for example, Sf9 and Sf21 cells) or a mammalian cell line. Preferred mammalian cells include primate cells (including human), canine cells and rodent cells. Cells can be primary cells or immortalized cells. Suitable cells can be selected from Vero cells, COS cells, HEK 293 cells, HeLa cells, CHO cells, BHK cells, MDCK cells, amniotic cells (human),

embryonic cells, cell lines transfected with viral genes, for example, AD5 E1, including but not limited to an immortalized human retinal cell transfected with an adenovirus gene, for example, a PER.C6 cell, or an NSO cell. In some embodiments, the cell is a Chinese hamster ovary (CHO) cell line. Some examples of CHO cells include, but are not limited to, CHO-ori, CHO-K1, CHO-s, CHO-DHB11, CHO-DXB11, CHO-K1SV, and mutants and variants thereof. In other embodiments, the cell is a HEK293 cell. Some examples of HEK293 cells include, but are not limited, to HEK293, HEK293A, HEK293E, HEK293F, HEK293FT, HEK293FTM, HEK293H, HEK293MSR, HEK293S, HEK293SG, HEK293SGGD, HEK293T and mutants and variants thereof.

[0050] For hamster cells such as CHO and BHK, integration can be accomplished by inventions disclosed in U.S. Patent Nos. 7,771,997 ("Stable Site 1") and 9,816,110 ("Stable Site 2"), which are hereby incorporated by reference, including sequence information. Regeneron provides a suite of goods and services referred to as EESYR®. CHO cells with integrated sequences in Stable Site 1 and Stable Site 2 are disclosed in US 2019/0233544 A1, which is hereby incorporated by reference, including sequence information. Sequences set forth in these patents and Examples 14 and 15 can be used according to the inventions described and depicted herein. Additionally, an AAVS1-like region and the COSMC locus in hamster cells can be used according to the inventions.

[0051] Where human cells are employed, integration into adeno-associated virus integration site 1 (AAVS1) can be undertaken. See Lou *et al., Human Gene Therapy Methods*, 28: 124-38 (2017); Liu *et al., BMC Research Note*,

7:626 (2014). AAVS1 is reported to be located on chromosome 19. Other integration sites in human cells can be used as well, such as CCR5 and hROSA26.

[0052] Modification of cellular genomes can be undertaken with known approaches, such as Cre/Lox, Flp/Frt, transcription activator-like effector nuclease (TALEN), a TAL effector domain fusion protein, zinc finger nuclease (ZFN), a ZFN dimer, or a RNA-guided DNA endonuclease system, such as CRISPR/Cas9. See U.S. Patent No. 9,816,110 at cols. 17-18. Integration using Bxb1 integrase in human, mouse and rat cells also can be undertaken. Russell *et al., Biotechniques* 40: 460-64 (2006).

[0053] To maximize stability and efficiency and facilitate integration and control of the inventions, Stable Integration Sites (SIS) can be created using Genomic Safe Harbors and the like in a wide variety of cell types and lines according to the teachings of U.S. Serial No. 63/256,675. The descriptions (including examples) and figures providing methods and cells resulting from the methods of U.S. Serial No. 63/256,675 are hereby incorporated by reference.

[0054] For production of recombinant AAV, the inventions provide for integration of AAV Cap to produce size variants VP1 (about 90kDa), VP2 (about 72kDa) and VP3 (about 60kDa). The variants differ at their N-terminus.

[0055] Usually, recombinant AAV will contain a gene-of interest (GOI) flanked by AAV ITRs (inverted terminal repeats), For production of recombinant AAV, seven additional polynucleotides, namely adenovirus E1A, E1B, E4, E2A, VA RNA and AAV Rep and Cap, are typically employed for production.

[0056] Figures 1-4 depict constructs that permit constitutive expression of Cap protein. Figures 5-8 depict constructs that permit controlled expression of

Cap protein by inclusion of an operator downstream of the promoter. A preferred operator is the tetracycline operator (TetO), which binds the tetracycline repressor (TetR). Tetracycline, doxycycline and derivatives thereof can bind TetR so that TetR no longer binds TetO, and thus is permissive for transcription. An example of a CMV promoter and TetO is set forth in Example 13.

[0057] The inventions is further described by the following examples, which are illustrative of the many aspects of the invention, but do not limit the inventions in any manner.

EXAMPLE 1 – CHO Cells

[0058] One or more Cap-containing polynucleotides according to Figures 1-8 are stably inserted into the CHO genome. A preferred promoter is the hCMV-IE promoter, and optionally a tet operator can be operably linked to the promoter for expression control. Optionally, an intron can be located 3' of the promoter. A preferred intron is an hCMV-IE intron. AAV Cap, Rep and ITRs can be obtained from any AAV serotype. Preferred AAV serotypes are AAV2 and AAV5. AAV polynucleotide sequences are set forth in Example 11. Promoter, operator, IRES and intron sequences are set forth in Example 13.

[0059] AAV ITRs flanking a gene of interest, AAV Rep and Ad E1A, E1B, E2A (or E2A partial sequence (E2A orf)), E4 (or E4 partial sequence (E4 orf 6)) and VA RNA can be randomly integrated, site-specifically integrated or remain on a plasmid. Adenovirus polynucleotide sequences are available and are exemplified in Example 12. Adenovirus (Ad) proteins and polynucleotides can be selected from any serotype.

EXAMPLE 2 – HEK 293 Cells

[0060] One or more Cap-containing polynucleotides according to Figures 1-8 are stably inserted into the HEK 293 genome. A preferred promoter is the hCMV-IE promoter, and optionally a tet operator can be operably linked to the promoter for expression control. Optionally, an intron can be located 3' of the promoter. A preferred intron is an hCMV-IE intron. AAV Cap, Rep and ITRs can be obtained from any AAV serotype. Preferred AAV serotypes of AAV2 and AAV5. AAV polynucleotide sequences are set forth in Example 11.

[0061] AAV ITRs and Rep and Ad E1A, E1B, E2A (or E2A partial sequence (E2A orf)), E4 (or E4 partial sequence (E4 orf 6)) and VA RNA can be randomly integrated, site-specifically integrated or remain on a plasmid. Adenovirus polynucleotide sequences are available and are exemplified in Example 12.

EXAMPLE 3 – BHK Cells

[0062] BHK cells are fibroblast cells from baby hamster kidneys. There are adherent BHK lines and BHK lines that can propagate in suspension. Wentz and Schügerl, *Enzyme Microbial Tech.* 14: 68-75 (1992).

[0063] One or more Cap-containing polynucleotides according to Figures 1-8 are stably inserted into the BHK genome. A preferred promoter is the hCMV-IE promoter, and optionally a tet operator can be operably linked to the promoter for expression control. Optionally, an intron can be located 3' of the promoter. A preferred intron is an hCMV-IE intron. AAV Cap, Rep and ITRs can be obtained from any AAV serotype. Preferred AAV serotypes of AAV2 and AAV5. AAV polynucleotide sequences are set forth in Example 11.

[0064] AAV ITRs and Rep and Ad E1A, E1B, E2A (or E2A partial sequence (E2A orf)), E4 (or E4 partial sequence (E4 orf 6)) and VA RNA can be randomly integrated, site-specifically integrated or remain on a plasmid. Adenovirus polynucleotide sequences are available and are exemplified in Example 12.

EXAMPLE 4 – Intron IRES CAP

[0065] Embodiments of this construct are depicted in Figures 1 (constitutive) and 5 (controllable). In a cell, both embodiments can primarily produce VP2 and VP3, with more VP3 being produced than VP2. In an experiment in CHO cells using Cap5, an average ratio of about 1:5.7 of VP2 to VP3 was observed by conducting densitometry analyses on western blots. Some VP1 also can be produced, but the level observed was typically less than 1% the level of VP2 that is produced. Accordingly, this construct can be used with a construct that will produce primarily VP1. See Examples 6 and 7. Production ratios can vary based upon experimental conditions and analytical techniques. A preferred IRES is the encephalomyocarditis virus (referred to as “EMCV” or “ECMV”) IRES.

EXAMPLE 5 – Intron IRES CAP IRES CAP

[0066] Embodiments of this construct are depicted in Figures 2 (constitutive) and 6 (controllable). This construct contains two IRES polynucleotides and two Cap polynucleotides. In a cell, both embodiments can produce a VP1, VP2 and VP3. The amount of VP3 production can be greater than the amount of VP1 production and the amount of VP2 production. The amount of VP2 production can be greater than the amount of VP1 production. In an experiment using CHO cells containing Cap5, an average ratio of about 1:2:9.3 of VP1 to VP2 to VP3 was observed by conducting densitometry analyses on western blots. Production ratios

can vary based upon experimental conditions and analytical techniques. A preferred IRES is the encephalomyocarditis virus (referred to as “EMCV” or “ECMV”) IRES.

EXAMPLE 6 – Intron CAP

[0067] Embodiments of this construct are depicted in Figures 3 (constitutive) and 7 (controllable). This construct contains an Intron and Cap polynucleotide, and produces predominantly VP1, and can be used with the constructs of Example 4 to produce VP1, VP2 and VP3.

Example 7 – IRES CAP

[0068] Embodiments of this construct are depicted in Figures 4 (constitutive) and 8 (controllable). This construct contains an Intron and Cap polynucleotide, and produces predominantly VP1, and can be used with the constructs of Example 4 to produce VP1, VP2 and VP3. A preferred IRES is the encephalomyocarditis virus (referred to as “EMCV” or “ECMV”) IRES.

Example 8 – CHO Cells Comprising Cap constructs

[0069] CHO cells of Example 1 can comprise an Intron IRES CAP IRES CAP polynucleotide of Example 5 (Figures 2 or 6). In an experiment, CHO cells containing Cap5 expressed VP1:VP2:VP3 in an observed ratio of about 1:2:9.3 by conducting densitometry analyses on western blots.

[0070] As an alternative, CHO cells of Example 1 can comprise CAP polynucleotides of Example 4 (Figs. 1 or 5) and Examples 6 (Figs. 3 or 7) or 6 (Figs. 4 and 8). Such cells would express VP1, VP2 and VP3, although possibly at different ratios than according to Example 5.

Example 9 – HEK 293 Cells Comprising Cap constructs

[0071] HEK 293 cells of Example 2 can comprise an Intron IRES CAP IRES CAP polynucleotide of Example 5 (Figures 2 or 6).

[0072] As an alternative, HEK 293 cells of Example 2 can comprise CAP polynucleotides of Example 4 (Figs. 1 or 5) and Examples 6 (Figs. 3 or 7) or 7 (Figs. 4 and 8). Such cells would express VP1, VP2 and VP3, although possibly at different ratios than according to Example 5.

EXAMPLE 10 – BHK Cells Comprising Cap constructs

[0073] BHK cells of Example 3 can comprise an Intron IRES CAP IRES CAP polynucleotide of Example 5 (Figures 2 or 6).

[0074] As an alternative, BHK cells of Example 3 can comprise CAP polynucleotides of Example 4 (Figs. 1 or 5) and Examples 6 (Figs. 3 or 7) or 7 (Figs. 4 and 8). Such cells would express VP1, VP2 and VP3, although possibly at different ratios than according to Example 5.

EXAMPLE 11 – AAV Polynucleotide Sequences

[0075] AAV Rep, Cap and ITR sequences are known in the art. The present inventions are amenable to all AAV serotypes. AAV sequences from various AAV serotypes are set forth below. Many of these sequences are available from the National Center for Biotechnology Information (NCBI).

AAV-1

Full Genome: NC_002077

CapVP1: (SEQ ID NO: 1)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAGGGCATTCGCGAGTGGTGGGACTTG
AAACCTGGAGCCCCGAAGCCCAAAGCCAACCAGCAAAAGCAGGACGACGGCCGGGGTCTGGTGCTTCCTGGCTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGGGAGCCCGTCAACGCGGCGGACGCAGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTCAAAGCGGGTGACAATCCGTACCTGCGGTATAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTGCAAGAAGATACGTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAGAAGCGGGTT

CTCGAACCTCTCGGTCTGGTTGAGGAAGGCGCTAAGACGGCTCCTGGAAAGAAACGTCCGGTAGAGCAGTCGCCA
CAAGAGCCAGACTCCTCCTCGGGCATCGGCAAGACAGGCCAGCAGCCCGCTAAAAAGAGACTCAATTTTGGTCAG
ACTGGCGACTCAGAGTCAGTCCCCGATCCACAACCTCTCGGAGAACCTCCAGCAACCCCCGCTGCTGTGGGACCT
ACTACAATGGCTTCAGGCGGTGGCGCACCAATGGCAGACAATAACGAAGGCGCCGACGGAGTGGGTAATGCCTCA
GGAAATTGGCATTGCGATTCCACATGGCTGGGCGACAGAGTCATCACCACCAGCACCCGCACCTGGGCCTTGCCC
ACCTACAATAACCACCTCTACAAGCAAATCTCCAGTGCTTCAACGGGGGCCAGCAACGACAACCACTACTTCGGC
TACAGCACCCCCTGGGGGTATTTTGATTTCAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAGCGACTC
ATCAACAACAATTGGGGATTCCGGCCCAAGAGACTCAACTTCAAACTCTTCAACATCCAAGTCAAGGAGGTCACG
ACGAATGATGGCGTCACAACCATCGCTAATAACCTTACCAGCACGGTTCAAGTCTTCTCGGACTCGGAGTACCAG
CTTCCGTACGTCCTCGGCTCTGCGCACCAGGGCTGCCTCCCTCCGTTCCCGGCGGACGTGTTCATGATTCCGCAA
TACGGCTACCTGACGCTCAACAATGGCAGCCAAGCCGTGGGACGTTCATCCTTTTACTGCCTGGAATATTTCCCT
TCTCAGATGCTGAGAACGGGCAACAACTTTACCTTCAGCTACACCTTTGAGGAAGTGCCTTTCCACAGCAGCTAC
GCGCACAGCCAGAGCCTGGACCGGCTGATGAATCCTCTCATCGACCAATACCTGTATTACCTGAACAGAACTCAA
AATCAGTCCGGAAGTGCCCAAAACAAGGACTTGCTGTTTAGCCGTGGGTCTCCAGCTGGCATGTCTGTTCAGCCC
AAAAACTGGCTACCTGGACCCTGTTATCGGCAGCAGCGCGTTTCTAAAACAAAAACAGACAACAACAACAGCAAT
TTTACCTGGACTGGTGCTTCAAAATATAACCTCAATGGGCGTGAATCCATCATCAACCCTGGCACTGCTATGGCC
TCACACAAAGACGACGAAGACAAGTTCTTTCCCATGAGCGGTGTCATGATTTTTGGAAAAGAGAGCGCCGGAGCT
TCAAACACTGCATTGGACAATGTCATGATTACAGACGAAGAGGAAATTAAAGCCACTAACCCTGTGGCCACCGAA
AGATTTGGGACCGTGGCAGTCAATTTCCAGAGCAGCAGCACAGACCCTGCGACCGGAGATGTGCATGCTATGGGA
GCATTACCTGGCATGGTGTGGCAAGATAGAGACGTGTACCTGCAGGGTCCCATTTGGGCCAAAATTCCTCACACA
GATGGACACTTTCACCCGTCTCCTCTTATGGGCGGCTTTGGACTCAAGAACCCGCCTCCTCAGATCCTCATCAAA
AACACGCCTGTTCCTGCGAATCCTCCGGCGGAGTTTTCAGCTACAAAGTTTGCTTCATTCATCACCCAATACTCC
ACAGGACAAGTGAGTGTGGAAATTGAATGGGAGCTGCAGAAAGAAAACAGCAAGCGCTGGAATCCCGAAGTGCAG
TACACATCCAATTATGCAAAATCTGCCAACGTTGATTTTACTGTGGACAACAATGGACTTTATACTGAGCCTCGC
CCCATTGGCACCCGTTACCTTACCCGTCCCCTGTAA

Rep78: (SEQ ID NO: 2)

ATGCCGGGCTTCTACGAGATCGTGATCAAGGTGCCGAGCGACCTGGACGAGCACCTGCCGGGCATTTCTGACTCG
TTTGTGAGCTGGGTGGCCGAGAAGGAATGGGAGCTGCCCCCGGATTCTGACATGGATCTGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTCCTGGTCCAATGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTTCAGTTCGAGAAGGGCGAGTCCTACTTCCACCTCCATATTCTGGTGGAGACCACGGGGGTC
AAATCCATGGTGCTGGGCCGCTTCCTGAGTCAGATTAGGGACAAGCTGGTGCAGACCATCTACCGCGGGATCGAG
CCGACCCTGCCCAACTGGTTCGCGGTGACCAAGACGCGTAATGGCGCCGGAGGGGGGAACAAGGTGGTGGACGAG
TGCTACATCCCCAACTACCTCCTGCCCAAGACTCAGCCCGAGCTGCAGTGGGCGTGGACTAACATGGAGGAGTAT
ATAAGCGCCTGTTTGAACCTGGCCGAGCGCAAACGGCTCGTGGCGCAGCACCTGACCCACGTCAGCCAGACCCAG
GAGCAGAACAAGGAGAATCTGAACCCCAATTCTGACGCGCCTGTCATCCGGTCAAAAACCTCCGCGCGCTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGGGGCATCACCTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCGTAC
ATCTCCTTCAACGCCGCTTCCAACTCGCGGTCCCAGATCAAGGCCGCTCTGGACAATGCCGGCAAGATCATGGCG
CTGACCAAATCCGCGCCCGACTACCTGGTAGGCCCCGCTCCGCCCGCGGACATTAAAACCAACCGCATCTACCGC
ATCCTGGAGCTGAACGGCTACGAACCTGCCTACGCCGGCTCCGTCTTTCTCGGCTGGGCCCAGAAAAGGTTCGGG
AAGCGCAACACCATCTGGCTGTTTGGGCCGGCCACCACGGGCAAGACCAACATCGCGGAAGCCATCGCCCACGCC
GTGCCCTTCTACGGCTGCGTCAACTGGACCAATGAGAACTTTCCCTTCAATGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTCGGCGGCAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCGTCCGCCCAGATCGACCCCACCCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCGTTGCAGGACCGGATGTTCAAATTTGAACTC
ACCCGCCGTCTGGAGCATGACTTTGGCAAGGTGACAAAGCAGGAAGTCAAAGAGTTCTTCCGCTGGGCGCAGGAT
CACGTGACCGAGGTGGCGCATGAGTTCTACGTCAGAAAGGGTGGAGCCAACAAAAGACCCGCCCCCGATGACGCG
GATAAAAGCGAGCCCAAGCGGGCCTGCCCCTCAGTCGCGGATCCATCGACGTCAGACGCGGAAGGAGCTCCGGTG
GACTTTGCCGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGCATGCTTCAGATGCTGTTTCCCTGCAAGACA
TGCGAGAGAATGAATCAGAATTTCAACATTTGCTTCACGCACGGGACGAGAGACTGTTCAGAGTGCTTCCCCGGC
GTGTCAGAATCTCAACCGGTCGTCAGAAAGAGGACGTATCGGAAACTCTGTGCCATTCATCATCTGCTGGGGCGG
GCTCCCGAGATTGCTTGCTCGGCCTGCGATCTGGTCAACGTGGACCTGGATGACTGTGTTTCTGAGCAATAA

AAV-2
Full Genome: NC_001401

Rep78: (SEQ ID NO: 3)

ATGCCGGGGTTTTACGAGATTGTGATTAAGGTCCCCAGCGACCTTGACGAGCATCTGCCCGGCATTTCTGACAGC
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGTTGCCGCCAGATTCTGACATGGATCTGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTTCTGACGGAATGGCGCCGTGTGAGTAAGGCCCCGGAG
GCCCTTTTCTTTGTGCAATTTGAGAAGGGAGAGAGCTACTTCCACATGCACGTGCTCGTGGAAACCACCGGGGTG
AAATCCATGGTTTTGGGACGTTTCCTGAGTCAGATTCGCGAAAAACTGATTCAGAGAATTTACCGCGGGATCGAG
CCGACTTTGCCAAACTGGTTCGCGGTCACAAAGACCAGAAATGGCGCCGGAGGCGGGAACAAGGTGGTGGATGAG
TGCTACATCCCCAATTACTTGCTCCCCAAAACCCAGCCTGAGCTCCAGTGGGCGTGGACTAATATGGAACAGTAT
TTAAGCGCCTGTTTGAATCTCACGGAGCGTAAACGGTTGGTGGCGCAGCATCTGACGCACGTGTCGCAGACGCAG
GAGCAGAACAAAGAGAATCAGAATCCCAATTCTGATGCGCCGGTGATCAGATCAAAAACTTCAGCCAGGTACATG
GAGCTGGTCGGGTGGCTCGTGGACAAGGGGATTACCTCGGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCATAC
ATCTCCTTCAATGCGGCCTCCAACTCGCGGTCCCAAATCAAGGCTGCCTTGGACAATGCGGGAAAGATTATGAGC
CTGACTAAAACCGCCCCCGACTACCTGGTGGGCCAGCAGCCCGTGGAGGACATTTCCAGCAATCGGATTTATAAA
ATTTTGGAACTAAACGGGTACGATCCCCAATATGCGGCTTCCGTCTTTCTGGGATGGGCCACGAAAAAGTTCGGC
AAGAGGAACACCATCTGGCTGTTTGGGCCTGCAACTACCGGGAAGACCAACATCGCGGAGGCCATAGCCCACACT
GTGCCCTTCTACGGGTGCGTAAACTGGACCAATGAGAACTTTCCCTTCAACGACTGTGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGGAAGATGACCGCCAAGGTCGTGGAGTCGGCCAAAGCCATTCTCGGAGGAAGCAAGGTGCGC
GTGGACCAGAAATGCAAGTCCTCGGCCCAGATAGACCCGACTCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACTCAACGACCTTCGAACACCAGCAGCCGTTGCAAGACCGGATGTTCAAATTTGAACTC
ACCCGCCGTCTGGATCATGACTTTGGGAAGGTCACCAAGCAGGAAGTCAAAGACTTTTTCCGGTGGGCAAAGGAT
CACGTGGTTGAGGTGGAGCATGAATTCTACGTCAAAAAGGGTGGAGCCAAGAAAAGACCCGCCCCCAGTGACGCA
GATATAAGTGAGCCCAAACGGGTGCGCGAGTCAGTTGCGCAGCCATCGACGTCAGACGCGGAAGCTTCGATCAAC
TACGCAGACAGGTACCAAAACAAATGTTCTCGTCACGTGGGCATGAATCTGATGCTGTTTCCCTGCAGACAATGC
GAGAGAATGAATCAGAATTCAAATATCTGCTTCACTCACGGACAGAAAGACTGTTTAGAGTGCTTTCCCGTGTCA
GAATCTCAACCCGTTTCTGTCGTCAAAAAGGCGTATCAGAAACTGTGCTACATTCATCATATCATGGGAAAGGTG
CCAGACGCTTGCACTGCCTGCGATCTGGTCAATGTGGATTTGGATGACTGCATCTTTGAACAATAA

Rep52: (SEQ ID NO: 4)

ATGGAGCTGGTCGGGTGGCTCGTGGACAAGGGGATTACCTCGGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCA
TACATCTCCTTCAATGCGGCCTCCAACTCGCGGTCCCAAATCAAGGCTGCCTTGGACAATGCGGGAAAGATTATG
AGCCTGACTAAAACCGCCCCCGACTACCTGGTGGGCCAGCAGCCCGTGGAGGACATTTCCAGCAATCGGATTTAT
AAAATTTTGGAACTAAACGGGTACGATCCCCAATATGCGGCTTCCGTCTTTCTGGGATGGGCCACGAAAAAGTTC
GGCAAGAGGAACACCATCTGGCTGTTTGGGCCTGCAACTACCGGGAAGACCAACATCGCGGAGGCCATAGCCCAC
ACTGTGCCCTTCTACGGGTGCGTAAACTGGACCAATGAGAACTTTCCCTTCAACGACTGTGTCGACAAGATGGTG
ATCTGGTGGGAGGAGGGGAAGATGACCGCCAAGGTCGTGGAGTCGGCCAAAGCCATTCTCGGAGGAAGCAAGGTG
CGCGTGGACCAGAAATGCAAGTCCTCGGCCCAGATAGACCCGACTCCCGTGATCGTCACCTCCAACACCAACATG
TGCGCCGTGATTGACGGGAACTCAACGACCTTCGAACACCAGCAGCCGTTGCAAGACCGGATGTTCAAATTTGAA
CTCACCCGCCGTCTGGATCATGACTTTGGGAAGGTCACCAAGCAGGAAGTCAAAGACTTTTTCCGGTGGGCAAAG
GATCACGTGGTTGAGGTGGAGCATGAATTCTACGTCAAAAAGGGTGGAGCCAAGAAAAGACCCGCCCCCAGTGAC
GCAGATATAAGTGAGCCCAAACGGGTGCGCGAGTCAGTTGCGCAGCCATCGACGTCAGACGCGGAAGCTTCGATC
AACTACGCAGACAGGTACCAAAACAAATGTTCTCGTCACGTGGGCATGAATCTGATGCTGTTTCCCTGCAGACAA
TGCGAGAGAATGAATCAGAATTCAAATATCTGCTTCACTCACGGACAGAAAGACTGTTTAGAGTGCTTTCCCGTG
TCAGAATCTCAACCCGTTTCTGTCGTCAAAAAGGCGTATCAGAAACTGTGCTACATTCATCATATCATGGGAAAG
GTGCCAGACGCTTGCACTGCCTGCGATCTGGTCAATGTGGATTTGGATGACTGCATCTTTGAACAATAA

CapVP1: (SEQ ID NO: 5)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACACTCTCTCTGAAGGAATAAGACAGTGGTGGAAGCTC
AAACCTGGCCCACCACCACCAAAGCCCGCAGAGCGGCATAAGGACGACAGCAGGGGTCTTGTGCTTCCTGGGTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGAGAGCCGGTCAACGAGGCAGACGCCGCGGCCCTCGAGCAC
GACAAAGCCTACGACCGGCAGCTCGACAGCGGAGACAACCCGTACCTCAAGTACAACCACGCCGACGCGGAGTTT
CAGGAGCGCCTTAAAGAAGATACGTCTTTTGGGGGCAACCTCGGACGAGCAGTCTTCCAGGCGAAAAAGAGGGTT
CTTGAACCTCTGGGCCTGGTTGAGGAACCTGTTAAGACGGCTCCGGGAAAAAAGAGGCCGGTAGAGCACTCTCCT
GTGGAGCCAGACTCCTCCTCGGGAACCGGAAAGGCGGGCCAGCAGCCTGCAAGAAAAAGATTGAATTTTGGTCAG
ACTGGAGACGCAGACTCAGTACCTGACCCCCAGCCTCTCGGACAGCCACCAGCAGCCCCCTCTGGTCTGGGAACT
AATACGATGGCTACAGGCAGTGGCGCACCAATGGCAGACAATAACGAGGGCGCCGACGGAGTGGGTAATTCCTCG

GGAAATTGGCATTGCGATTCCACATGGATGGGCGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTGCCC
ACCTACAACAACCACCTCTACAAACAAATTTCCAGCCAATCAGGAGCCTCGAACGACAATCACTACTTTGGCTAC
AGCACCCCTTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAAAGACTCATC
AACAACAACTGGGGATTCCGACCCAAGAGACTCAACTTCAAGCTCTTTAACATTCAAGTCAAAGAGGTCACGCAG
AATGACGGTACGACGACGATTGCCAATAACCTTACCAGCACGGTTCAGGTGTTTACTGACTCGGAGTACCAGCTC
CCGTACGTCCTCGGCTCGGCGCATCAAGGATGCCTCCCGCCGTTCCCAGCAGACGTCTTCATGGTGCCACAGTAT
GGATACCTCACCCTGAACAACGGGAGTCAGGCAGTAGGACGCTCTTCATTTTACTGCCTGGAGTACTTTCCTTCT
CAGATGCTGCGTACCGGAAACAACTTTACCTTCAGCTACACTTTTGAGGACGTTCCTTTCCACAGCAGCTACGCT
CACAGCCAGAGTCTGGACCGTCTCATGAATCCTCTCATCGACCAGTACCTGTATTACTTGAGCAGAACAAACACT
CCAAGTGGAACCACCACGCAGTCAAGGCTTCAGTTTTCTCAGGCCGGAGCGAGTGACATTCGGGACCAGTCTAGG
AACTGGCTTCCTGGACCCTGTTACCGCCAGCAGCGAGTATCAAAGACATCTGCGGATAACAACAACAGTGAATAC
TCGTGGACTGGAGCTACCAAGTACCACCTCAATGGCAGAGACTCTCTGGTGAATCCGGGCCCGGCCATGGCAAGC
CACAAGGACGATGAAGAAAAGTTTTTTCCTCAGAGCGGGGTTCTCATCTTTGGGAAGCAAGGCTCAGAGAAAACA
AATGTGGACATTGAAAAGGTCATGATTACAGACGAAGAGGAAATCAGGACAACCAATCCCGTGGCTACGGAGCAG
TATGGTTCTGTATCTACCAACCTCCAGAGAGGCAACAGACAAGCAGCTACCGCAGATGTCAACACACAAGGCGTT
CTTCCAGGCATGGTCTGGCAGGACAGAGATGTGTACCTTCAGGGGCCCATCTGGGCAAAGATTCCACACACGGAC
GGACATTTTCACCCCTCTCCCCTCATGGGTGGATTCGGACTTAAACACCCTCCTCCACAGATTCTCATCAAGAAC
ACCCCGGTACCTGCGAATCCTTCGACCACCTTCAGTGCGGCAAAGTTTGCTTCCTTCATCACACAGTACTCCACG
GGACAGGTCAGCGTGGAGATCGAGTGGGAGCTGCAGAAGGAAAACAGCAAACGCTGGAATCCCGAAATTCAGTAC
ACTTCCAACTACAACAAGTCTGTTAATGTGGACTTTACTGTGGACACTAATGGCGTGTATTCAGAGCCTCGCCCC
ATTGGCACCAGATACCTGACTCGTAATCTGTAA

CapVP2: (SEQ ID NO: 6)

ACGGCTCCGGGAAAAAAGAGGCCGGTAGAGCACTCTCCTGTGGAGCCAGACTCCTCCTCGGGAACCGGAAAGGCG
GGCCAGCAGCCTGCAAGAAAAAGATTGAATTTTGGTCAGACTGGAGACGCAGACTCAGTACCTGACCCCCAGCCT
CTCGGACAGCCACCAGCAGCCCCCTCTGGTCTGGGAACTAATACGATGGCTACAGGCAGTGGCGCACCAATGGCA
GACAATAACGAGGGCGCCGACGGAGTGGGTAATTCCTCGGGAAATTGGCATTGCGATTCCACATGGATGGGCGAC
AGAGTCATCACCACCAGCACCCGAACCTGGGCCCTGCCCACCTACAACAACCACCTCTACAAACAAATTTCCAGC
CAATCAGGAGCCTCGAACGACAATCACTACTTTGGCTACAGCACCCCTTGGGGGTATTTTGACTTCAACAGATTC
CACTGCCACTTTTCACCACGTGACTGGCAAAGACTCATCAACAACAACTGGGGATTCCGACCCAAGAGACTCAAC
TTCAAGCTCTTTAACATTCAAGTCAAAGAGGTCACGCAGAATGACGGTACGACGACGATTGCCAATAACCTTACC
AGCACGGTTCAGGTGTTTACTGACTCGGAGTACCAGCTCCCGTACGTCCTCGGCTCGGCGCATCAAGGATGCCTC
CCGCCGTTCCCAGCAGACGTCTTCATGGTGCCACAGTATGGATACCTCACCCTGAACAACGGGAGTCAGGCAGTA
GGACGCTCTTCATTTTACTGCCTGGAGTACTTTCCTTCTCAGATGCTGCGTACCGGAAACAACTTTACCTTCAGC
TACACTTTTGAGGACGTTCCTTTCCACAGCAGCTACGCTCACAGCCAGAGTCTGGACCGTCTCATGAATCCTCTC
ATCGACCAGTACCTGTATTACTTGAGCAGAACAAACACTCCAAGTGGAACCACCACGCAGTCAAGGCTTCAGTTT
TCTCAGGCCGGAGCGAGTGACATTCGGGACCAGTCTAGGAACTGGCTTCCTGGACCCTGTTACCGCCAGCAGCGA
GTATCAAAGACATCTGCGGATAACAACAACAGTGAATACTCGTGGACTGGAGCTACCAAGTACCACCTCAATGGC
AGAGACTCTCTGGTGAATCCGGGCCCGGCCATGGCAAGCCACAAGGACGATGAAGAAAAGTTTTTTCCTCAGAGC
GGGGTTCTCATCTTTGGGAAGCAAGGCTCAGAGAAAACAAATGTGGACATTGAAAAGGTCATGATTACAGACGAA
GAGGAAATCAGGACAACCAATCCCGTGGCTACGGAGCAGTATGGTTCTGTATCTACCAACCTCCAGAGAGGCAAC
AGACAAGCAGCTACCGCAGATGTCAACACACAAGGCGTTCTTCCAGGCATGGTCTGGCAGGACAGAGATGTGTAC
CTTCAGGGGCCCATCTGGGCAAAGATTCCACACACGGACGGACATTTTCACCCCTCTCCCCTCATGGGTGGATTC
GGACTTAAACACCCTCCTCCACAGATTCTCATCAAGAACACCCCGGTACCTGCGAATCCTTCGACCACCTTCAGT
GCGGCAAAGTTTGCTTCCTTCATCACACAGTACTCCACGGGACAGGTCAGCGTGGAGATCGAGTGGGAGCTGCAG
AAGGAAAACAGCAAACGCTGGAATCCCGAAATTCAGTACACTTCCAACTACAACAAGTCTGTTAATGTGGACTTT
ACTGTGGACACTAATGGCGTGTATTCAGAGCCTCGCCCCATTGGCACCAGATACCTGACTCGTAATCTGTAA

CapVP3: (SEQ ID NO:7)

ATGGCTACAGGCAGTGGCGCACCAATGGCAGACAATAACGAGGGCGCCGACGGAGTGGGTAATTCCTCGGGAAAT
TGGCATTGCGATTCCACATGGATGGGCGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTGCCCACCTAC
AACAACCACCTCTACAAACAAATTTCCAGCCAATCAGGAGCCTCGAACGACAATCACTACTTTGGCTACAGCACC
CCTTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAAAGACTCATCAACAAC
AACTGGGGATTCCGACCCAAGAGACTCAACTTCAAGCTCTTTAACATTCAAGTCAAAGAGGTCACGCAGAATGAC
GGTACGACGACGATTGCCAATAACCTTACCAGCACGGTTCAGGTGTTTACTGACTCGGAGTACCAGCTCCCGTAC
GTCCTCGGCTCGGCGCATCAAGGATGCCTCCCGCCGTTCCCAGCAGACGTCTTCATGGTGCCACAGTATGGATAC

CTCACCCTGAACAACGGGAGTCAGGCAGTAGGACGCTCTTCATTTTACTGCCTGGAGTACTTTCCTTCTCAGATG
CTGCGTACCGGAAACAACTTTACCTTCAGCTACACTTTTGAGGACGTTCCTTTCCACAGCAGCTACGCTCACAGC
CAGAGTCTGGACCGTCTCATGAATCCTCTCATCGACCAGTACCTGTATTACTTGAGCAGAACAAACACTCCAAGT
GGAACCACCACGCAGTCAAGGCTTCAGTTTTCTCAGGCCGGAGCGAGTGACATTCGGGACCAGTCTAGGAACTGG
CTTCCTGGACCCTGTTACCGCCAGCAGCGAGTATCAAAGACATCTGCGGATAACAACAACAGTGAATACTCGTGG
ACTGGAGCTACCAAGTACCACCTCAATGGCAGAGACTCTCTGGTGAATCCGGGCCCGGCCATGGCAAGCCACAAG
GACGATGAAGAAAAGTTTTTTCCTCAGAGCGGGGTTCTCATCTTTGGGAAGCAAGGCTCAGAGAAAACAAATGTG
GACATTGAAAAGGTCATGATTACAGACGAAGAGGAAATCAGGACAACCAATCCCGTGGCTACGGAGCAGTATGGT
TCTGTATCTACCAACCTCCAGAGAGGCAACAGACAAGCAGCTACCGCAGATGTCAACACACAAGGCGTTCTTCCA
GGCATGGTCTGGCAGGACAGAGATGTGTACCTTCAGGGGCCCATCTGGGCAAAGATTCCACACACGGACGGACAT
TTTCACCCCTCTCCCCTCATGGGTGGATTCGGACTTAAACACCCTCCTCCACAGATTCTCATCAAGAACACCCCG
GTACCTGCGAATCCTTCGACCACCTTCAGTGCGGCAAAGTTTGCTTCCTTCATCACACAGTACTCCACGGGACAG
GTCAGCGTGGAGATCGAGTGGGAGCTGCAGAAGGAAAACAGCAAACGCTGGAATCCCGAAATTCAGTACACTTCC
AACTACAACAAGTCTGTTAATGTGGACTTTACTGTGGACACTAATGGCGTGTATTCAGAGCCTCGCCCCATTGGC
ACCAGATACCTGACTCGTAATCTGTAA

CapAAP: (SEQ ID NO:8)

CTGGAGACGCAGACTCAGTACCTGACCCCCAGCCTCTCGGACAGCCACCAGCAGCCCCCTCTGGTCTGGGAACTA
ATACGATGGCTACAGGCAGTGGCGCACCAATGGCAGACAATAACGAGGGCGCCGACGGAGTGGGTAATTCCTCGG
GAAATTGGCATTGCGATTCCACATGGATGGGCGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTGCCCA
CCTACAACAACCACCTCTACAAACAAATTTCCAGCCAATCAGGAGCCTCGAACGACAATCACTACTTTGGCTACA
GCACCCCTTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAAAGACTCATCA
ACAACAACTGGGGATTCCGACCCAAGAGACTCAACTTCAAGCTCTTTAACATTCAAGTCAAAGAGGTCACGCAGA
ATGACGGTACGACGACGATTGCCAATAACCTTACCAGCACGGTTCAGGTGTTTACTGACTCGGAGTACCAGCTCC
CGTACGTCCTCGGCTCGGCGCATCAAGGATGCCTCCCGCCGTTCCCAGCAGACGTCTTCATGGTGCCACAGTATG
GATACCTCACCCTGA

AAV-3
Full Genome: NC_001729

Rep78: (SEQ ID NO:9)

ATGCCGGGGTTCTACGAGATTGTCCTGAAGGTCCCGAGTGACCTGGACGAGCGCCTGCCGGGCATTTCTAACTCG
TTTGTTAACTGGGTGGCCGAGAAGGAATGGGACGTGCCGCCGGATTCTGACATGGATCCGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAAAAGCTTCAGCGCGAGTTCCTGGTGGAGTGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTTTTTGTCCAGTTCGAAAAGGGGGAGACCTACTTCCACCTGCACGTGCTGATTGAGACCATCGGGGTC
AAATCCATGGTGGTCGGCCGCTACGTGAGCCAGATTAAAGAGAAGCTGGTGACCCGCATCTACCGCGGGGTCGAG
CCGCAGCTTCCGAACTGGTTCGCGGTGACCAAAACGCGAAATGGCGCCGGGGGCGGGAACAAGGTGGTGGACGAC
TGCTACATCCCCAACTACCTGCTCCCCAAGACCCAGCCCGAGCTCCAGTGGGCGTGGACTAACATGGACCAGTAT
TTAAGCGCCTGTTTGAATCTCGCGGAGCGTAAACGGCTGGTGGCGCAGCATCTGACGCACGTGTCGCAGACGCAG
GAGCAGAACAAAGAGAATCAGAACCCCAATTCTGACGCGCCGGTCATCAGGTCAAAAACCTCAGCCAGGTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGCGGGATCACGTCAGAAAAGCAATGGATTCAGGAGGACCAGGCCTCGTAC
ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCCCAGATCAAGGCCGCGCTGGACAATGCCTCCAAGATCATGAGC
CTGACAAAGACGGCTCCGGACTACCTGGTGGGCAGCAACCCGCCGGAGGACATTACCAAAAATCGGATCTACCAA
ATCCTGGAGCTGAACGGGTACGATCCGCAGTACGCGGCCTCCGTCTTCCTGGGCTGGGCGCAAAAGAAGTTCGGG
AAGAGGAACACCATCTGGCTCTTTGGGCCGGCCACGACGGGTAAAACCAACATCGCGGAAGCCATCGCCCACGCC
GTGCCCTTCTACGGCTGCGTAAACTGGACCAATGAGAACTTTCCCTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTGGAGAGCGCCAAGGCCATTCTGGGCGGAAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCATCGGCCCAGATCGAACCCACTCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACAGCACCACCTTCGAGCATCAGCAGCCGCTGCAGGACCGGATGTTTGAATTTGAACTT
ACCCGCCGTTTGGACCATGACTTTGGGAAGGTCACCAAACAGGAAGTAAAGGACTTTTTCCGGTGGGCTTCCGAT
CACGTGACTGACGTGGCTCATGAGTTCTACGTCAGAAAGGGTGGAGCTAAGAAACGCCCCGCCTCCAATGACGCG
GATGTAAGCGAGCCAAAACGGGAGTGCACGTCACTTGCGCAGCCGACAACGTCAGACGCGGAAGCACCGGCGGAC
TACGCGGACAGGTACCAAAACAAATGTTCTCGTCACGTGGGCATGAATCTGATGCTTTTTCCCTGTAAAACATGC
GAGAGAATGAATCAAATTTCCAATGTCTGTTTTACGCATGGTCAAAGAGACTGTGGGGAATGCTTCCCTGGAATG
TCAGAATCTCAACCCGTTTCTGTCGTCAAAAAGAAGACTTATCAGAAACTGTGTCCAATTCATCATATCCTGGGA
AGGGCACCCGAGATTGCCTGTTCGGCCTGCGATTTGGCCAATGTGGACTTGGATGACTGTGTTTCTGAGCAATAA

CapVP1: (SEQ ID NO:10)

ATGGCTGCTGACGGTTATCTTCCAGATTGGCTCGAGGACAACCTTTCTGAAGGCATTCGTGAGTGGTGGGCTCTG
AAACCTGGAGTCCCTCAACCCAAAGCGAACCAACAACACCAGGACAACCGTCGGGGTCTTGTGCTTCCGGGTTAC
AAATACCTCGGACCCGGTAACGGACTCGACAAAGGAGAGCCGGTCAACGAGGCGGACGCGGCAGCCCTCGAACAC
GACAAAGCTTACGACCAGCAGCTCAAGGCCGGTGACAACCCGTACCTCAAGTACAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTTCAAGAAGATACGTCTTTTGGGGGCAACCTTGGCAGAGCAGTCTTCCAGGCCAAAAAGAGGATC
CTTGAGCCTCTTGGTCTGGTTGAGGAAGCAGCTAAAACGGCTCCTGGAAAGAAGGGGGCTGTAGATCAGTCTCCT
CAGGAACCGGACTCATCATCTGGTGTTGGCAAATCGGGCAAACAGCCTGCCAGAAAAAGACTAAATTTCGGTCAG
ACTGGAGACTCAGAGTCAGTCCCAGACCCTCAACCTCTCGGAGAACCACCAGCAGCCCCCACAAGTTTGGGATCT
AATACAATGGCTTCAGGCGGTGGCGCACCAATGGCAGACAATAACGAGGGTGCCGATGGAGTGGGTAATTCCTCA
GGAAATTGGCATTGCGATTCCCAATGGCTGGGCGACAGAGTCATCACCACCAGCACCAGAACCTGGGCCCTGCCC
ACTTACAACAACCATCTCTACAAGCAAATCTCCAGCCAATCAGGAGCTTCAAACGACAACCACTACTTTGGCTAC
AGCACCCCTTGGGGGTATTTTGACTTTAACAGATTCCACTGCCACTTCTCACCACGTGACTGGCAGCGACTCATT
AACAACAACTGGGGATTCCGGCCCAAGAAACTCAGCTTCAAGCTCTTCAACATCCAAGTTAGAGGGGTCACGCAG
AACGATGGCACGACGACTATTGCCAATAACCTTACCAGCACGGTTCAAGTGTTTACGGACTCGGAGTATCAGCTC
CCGTACGTGCTCGGGTCGGCGCACCAAGGCTGTCTCCCGCCGTTTCCAGCGGACGTCTTCATGGTCCCTCAGTAT
GGATACCTCACCCTGAACAACGGAAGTCAAGCGGTGGGACGCTCATCCTTTTACTGCCTGGAGTACTTCCCTTCG
CAGATGCTAAGGACTGGAAATAACTTCCAATTCAGCTATACCTTCGAGGATGTACCTTTTCACAGCAGCTACGCT
CACAGCCAGAGTTTGGATCGCTTGATGAATCCTCTTATTGATCAGTATCTGTACTACCTGAACAGAACGCAAGGA
ACAACCTCTGGAACAACCAACCAATCACGGCTGCTTTTTAGCCAGGCTGGGCCTCAGTCTATGTCTTTGCAGGCC
AGAAATTGGCTACCTGGGCCCTGCTACCGGCAACAGAGACTTTCAAAGACTGCTAACGACAACAACAACAGTAAC
TTTCCTTGGACAGCGGCCAGCAAATATCATCTCAATGGCCGCGACTCGCTGGTGAATCCAGGACCAGCTATGGCC
AGTCACAAGGACGATGAAGAAAAATTTTTCCCTATGCACGGCAATCTAATATTTGGCAAAGAAGGGACAACGGCA
AGTAACGCAGAATTAGATAATGTAATGATTACGGATGAAGAAGAGATTCGTACCACCAATCCTGTGGCAACAGAG
CAGTATGGAACTGTGGCAAATAACTTGCAGAGCTCAAATACAGCTCCCACGACTGGAACTGTCAATCATCAGGGG
GCCTTACCTGGCATGGTGTGGCAAGATCGTGACGTGTACCTTCAAGGACCTATCTGGGCAAAGATTCCTCACACG
GATGGACACTTTCATCCTTCTCCTCTGATGGGAGGCTTTGGACTGAAACATCCGCCTCCTCAAATCATGATCAAA
AATACTCCGGTACCGGCAAATCCTCCGACGACTTTCAGCCCGGCCAAGTTTGCTTCATTTATCACTCAGTACTCC
ACTGGACAGGTCAGCGTGGAAATTGAGTGGGAGCTACAGAAAGAAAACAGCAAACGTTGGAATCCAGAGATTCAG
TACACTTCCAACTACAACAAGTCTGTTAATGTGGACTTTACTGTAGACACTAATGGTGTTTATAGTGAACCTCGC
CCTATTGGAACCCGGTATCTCACACGAAACTTGTGA

AAV-4

Full Genome: NC_001829

Rep78: (SEQ ID NO:11)

ATGCCGGGGTTCTACGAGATCGTGCTGAAGGTGCCCAGCGACCTGGACGAGCACCTGCCCGGCATTTCTGACTCT
TTTGTGAGCTGGGTGGCCGAGAAGGAATGGGAGCTGCCGCCGGATTCTGACATGGACTTGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAAAAGCTGCAACGCGAGTTCCTGGTCGAGTGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTCCAGTTCGAGAAGGGGGACAGCTACTTCCACCTGCACATCCTGGTGGAGACCGTGGGCGTC
AAATCCATGGTGGTGGGCCGCTACGTGAGCCAGATTAAAGAGAAGCTGGTGACCCGCATCTACCGCGGGGTCGAG
CCGCAGCTTCCGAACTGGTTCGCGGTGACCAAGACGCGTAATGGCGCCGGAGGCGGGAACAAGGTGGTGGACGAC
TGCTACATCCCCAACTACCTGCTCCCCAAGACCCAGCCCGAGCTCCAGTGGGCGTGGACTAACATGGACCAGTAT
ATAAGCGCCTGTTTGAATCTCGCGGAGCGTAAACGGCTGGTGGCGCAGCATCTGACGCACGTGTCGCAGACGCAG
GAGCAGAACAAGGAAAACCAGAACCCCAATTCTGACGCGCCGGTCATCAGGTCAAAAACCTCCGCCAGGTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGCGGGATCACGTCAGAAAAGCAATGGATCCAGGAGGACCAGGCGTCCTAC
ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCACAAATCAAGGCCGCGCTGGACAATGCCTCCAAAATCATGAGC
CTGACAAAGACGGCTCCGGACTACCTGGTGGGCCAGAACCCGCCGGAGGACATTTCCAGCAACCGCATCTACCGA
ATCCTCGAGATGAACGGGTACGATCCGCAGTACGCGGCCTCCGTCTTCCTGGGCTGGGCGCAAAAGAAGTTCGGG
AAGAGGAACACCATCTGGCTCTTTGGGCCGGCCACGACGGGTAAAACCAACATCGCGGAAGCCATCGCCCACGCC
GTGCCCTTCTACGGCTGCGTGAACTGGACCAATGAGAACTTTCCGTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTAGAGAGCGCCAAGGCCATCCTGGGCGGAAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCATCGGCCCAGATCGACCCAACTCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCGGTCATCGACGGAAACTCGACCACCTTCGAGCACCAACAACCACTCCAGGACCGGATGTTCAAGTTCGAGCTC

ACCAAGCGCCTGGAGCACGACTTTGGCAAGGTCACCAAGCAGGAAGTCAAAGACTTTTTCCGGTGGGCGTCAGAT
CACGTGACCGAGGTGACTCACGAGTTTTACGTCAGAAAGGGTGGAGCTAGAAAGAGGCCCGCCCCCAATGACGCA
GATATAAGTGAGCCCAAGCGGGCCTGTCCGTCAGTTGCGCAGCCATCGACGTCAGACGCGGAAGCTCCGGTGGAC
TACGCGGACAGGTACCAAAACAAATGTTCTCGTCACGTGGGTATGAATCTGATGCTTTTTCCCTGCCGGCAATGC
GAGAGAATGAATCAGAATGTGGACATTTGCTTCACGCACGGGGTCATGGACTGTGCCGAGTGCTTCCCCGTGTCA
GAATCTCAACCCGTGTCTGTCGTCAGAAAGCGGACGTATCAGAAACTGTGTCCGATTCATCACATCATGGGGAGG
GCGCCCGAGGTGGCCTGCTCGGCCTGCGAACTGGCCAATGTGGACTTGGATGACTGTGACATGGAACAATAA

CapVP1: (SEQ ID NO:12)

ATGACTGACGGTTACCTTCCAGATTGGCTAGAGGACAACCTCTCTGAAGGCGTTCGAGAGTGGTGGGCGCTGCAA
CCTGGAGCCCCTAAACCCAAGGCAAATCAACAACATCAGGACAACGCTCGGGGTCTTGTGCTTCCGGGTTACAAA
TACCTCGGACCCGGCAACGGACTCGACAAGGGGGAACCCGTCAACGCAGCGGACGCGGCAGCCCTCGAGCACGAC
AAGGCCTACGACCAGCAGCTCAAGGCCGGTGACAACCCCTACCTCAAGTACAACCACGCCGACGCGGAGTTCCAG
CAGCGGCTTCAGGGCGACACATCGTTTGGGGGCAACCTCGGCAGAGCAGTCTTCCAGGCCAAAAAGAGGGTTCTT
GAACCTCTTGGTCTGGTTGAGCAAGCGGGTGAGACGGCTCCTGGAAAGAAGAGACCGTTGATTGAATCCCCCCAG
CAGCCCGACTCCTCCACGGGTATCGGCAAAAAAGGCAAGCAGCCGGCTAAAAAGAAGCTCGTTTTCGAAGACGAA
ACTGGAGCAGGCGACGGACCCCCTGAGGGATCAACTTCCGGAGCCATGTCTGATGACAGTGAGATGCGTGCAGCA
GCTGGCGGAGCTGCAGTCGAGGGCGGACAAGGTGCCGATGGAGTGGGTAATGCCTCGGGTGATTGGCATTGCGAT
TCCACCTGGTCTGAGGGCCACGTCACGACCACCAGCACCAGAACCTGGGTCTTGCCCACCTACAACAACCACCTC
TACAAGCGACTCGGAGAGAGCCTGCAGTCCAACACCTACAACGGATTCTCCACCCCCTGGGGATACTTTGACTTC
AACCGCTTCCACTGCCACTTCTCACCACGTGACTGGCAGCGACTCATCAACAACAACTGGGGCATGCGACCCAAA
GCCATGCGGGTCAAAATCTTCAACATCCAGGTCAAGGAGGTCACGACGTCGAACGGCGAGACAACGGTGGCTAAT
AACCTTACCAGCACGGTTCAGATCTTTGCGGACTCGTCGTACGAACTGCCGTACGTGATGGATGCGGGTCAAGAG
GGCAGCCTGCCTCCTTTTCCCAACGACGTCTTTATGGTGCCCCAGTACGGCTACTGTGGACTGGTGACCGGCAAC
ACTTCGCAGCAACAGACTGACAGAAATGCCTTCTACTGCCTGGAGTACTTTCCTTCGCAGATGCTGCGGACTGGC
AACAACTTTGAAATTACGTACAGTTTTGAGAAGGTGCCTTTCCACTCGATGTACGCGCACAGCCAGAGCCTGGAC
CGGCTGATGAACCCTCTCATCGACCAGTACCTGTGGGGACTGCAATCGACCACCACCGGAACCACCCTGAATGCC
GGGACTGCCACCACCAACTTTACCAAGCTGCGGCCTACCAACTTTTCCAACTTTAAAAAGAACTGGCTGCCCGGG
CCTTCAATCAAGCAGCAGGGCTTCTCAAAGACTGCCAATCAAAACTACAAGATCCCTGCCACCGGGTCAGACAGT
CTCATCAAATACGAGACGCACAGCACTCTGGACGGAAGATGGAGTGCCCTGACCCCCGGACCTCCAATGGCCACG
GCTGGACCTGCGGACAGCAAGTTCAGCAACAGCCAGCTCATCTTTGCGGGGCCTAAACAGAACGGCAACACGGCC
ACCGTACCCGGGACTCTGATCTTCACCTCTGAGGAGGAGCTGGCAGCCACCAACGCCACCGATACGGACATGTGG
GGCAACCTACCTGGCGGTGACCAGAGCAACAGCAACCTGCCGACCGTGGACAGACTGACAGCCTTGGGAGCCGTG
CCTGGAATGGTCTGGCAAAACAGAGACATTTACTACCAGGGTCCCATTTGGGCCAAGATTCCTCATACCGATGGA
CACTTTCACCCCTCACCGCTGATTGGTGGGTTTGGGCTGAAACACCCGCCTCCTCAAATTTTTATCAAGAACACC
CCGGTACCTGCGAATCCTGCAACGACCTTCAGCTCTACTCCGGTAAACTCCTTCATTACTCAGTACAGCACTGGC
CAGGTGTCGGTGCAGATTGACTGGGAGATCCAGAAGGAGCGGTCCAAACGCTGGAACCCCGAGGTCCAGTTTACC
TCCAACTACGGACAGCAAAACTCTCTGTTGTGGGCTCCCGATGCGGCTGGGAAATACACTGAGCCTAGGGCTATC
GGTACCCGCTACCTCACCCACCACCTGTAA

AAV-5
Full Genome: NC_006152

Rep78: (SEQ ID NO:13)

ATGGCTACCTTCTATGAAGTCATTGTTCGCGTCCCATTTGACGTGGAGGAACATCTGCCTGGAATTTCTGACAGC
TTTGTGGACTGGGTAACTGGTCAAATTTGGGAGCTGCCTCCAGAGTCAGATTTAAATTTGACTCTGGTTGAACAG
CCTCAGTTGACGGTGGCTGATAGAATTCGCCGCGTGTTCCTGTACGAGTGGAACAAATTTTCCAAGCAGGAGTCC
AAATTCTTTGTGCAGTTTGAAAAGGGATCTGAATATTTTCATCTGCACACGCTTGTGGAGACCTCCGGCATCTCT
TCCATGGTCCTCGGCCGCTACGTGAGTCAGATTCGCGCCCAGCTGGTGAAAGTGGTCTTCCAGGGAATTGAACCC
CAGATCAACGACTGGGTCGCCATCACCAAGGTAAAGAAGGGCGGAGCCAATAAGGTGGTGGATTCTGGGTATATT
CCCGCCTACCTGCTGCCGAAGGTCCAACCGGAGCTTCAGTGGGCGTGGACAAACCTGGACGAGTATAAATTGGCC
GCCCTGAATCTGGAGGAGCGCAAACGGCTCGTCGCGCAGTTTCTGGCAGAATCCTCGCAGCGCTCGCAGGAGGCG
GCTTCGCAGCGTGAGTTCTCGGCTGACCCGGTCATCAAAAGCAAGACTTCCCAGAAATACATGGCGCTCGTCAAC
TGGCTCGTGGAGCACGGCATCACTTCCGAGAAGCAGTGGATCCAGGAAAATCAGGAGAGCTACCTCTCCTTCAAC
TCCACCGGCAACTCTCGGAGCCAGATCAAGGCCGCGCTCGACAACGCGACCAAAATTATGAGTCTGACAAAAAGC

GCGGTGGACTACCTCGTGGGGAGCTCCGTTCCCGAGGACATTTCAAAAAACAGAATCTGGCAAATTTTTGAGATG
AATGGCTACGACCCGGCCTACGCGGGATCCATCCTCTACGGCTGGTGTCAGCGCTCCTTCAACAAGAGGAACACC
GTCTGGCTCTACGGACCCGCCACGACCGGCAAGACCAACATCGCGGAGGCCATCGCCCACACTGTGCCCTTTTAC
GGCTGCGTGAACTGGACCAATGAAAACTTTCCCTTTAATGACTGTGTGGACAAAATGCTCATTTGGTGGGAGGAG
GGAAAGATGACCAACAAGGTGGTTGAATCCGCCAAGGCCATCCTGGGGGGCTCAAAGGTGCGGGTCGATCAGAAA
TGTAAATCCTCTGTTCAAATTGATTCTACCCCTGTCATTGTAACTTCCAATACAAACATGTGTGTGGTGGTGGAT
GGGAATTCCACGACCTTTGAACACCAGCAGCCGCTGGAGGACCGCATGTTCAAATTTGAACTGACTAAGCGGCTC
CCGCCAGATTTTGGCAAGATTACTAAGCAGGAAGTCAAGGACTTTTTTGCTTGGGCAAAGGTCAATCAGGTGCCG
GTGACTCACGAGTTTAAAGTTCCCAGGGAATTGGCGGGAACTAAAGGGGCGGAGAAATCTCTAAAACGCCCACTG
GGTGACGTCACCAATACTAGCTATAAAAGTCTGGAGAAGCGGGCCAGGCTCTCATTTGTTCCCGAGACGCCTCGC
AGTTCAGACGTGACTGTTGATCCCGCTCCTCTGCGACCGCTCAATTGGAATTCAAGGTATGATTGCAAATGTGAC
TATCATGCTCAATTTGACAACATTTCTAACAAATGTGATGAATGTGAATATTTGAATCGGGGCAAAAATGGATGT
ATCTGTCACAATGTAACTCACTGTCAAATTTGTCATGGGATTCCCCCCTGGGAAAAGGAAAACTTGTCAGATTTT
GGGGATTTTGACGATGCCAATAAAGAACAGTAA

CapVP1: (SEQ ID NO:14)

ATGTCTTTTGTTGATCACCCTCCAGATTGGTTGGAAGAAGTTGGTGAAGGTCTTCGCGAGTTTTTGGGCCTTGAA
GCGGGCCCACCGAAACCAAAACCCAATCAGCAGCATCAAGATCAAGCCCGTGGTCTTGTGCTGCCTGGTTATAAC
TATCTCGGACCCGGAAACGGTCTCGATCGAGGAGAGCCTGTCAACAGGGCAGACGAGGTCGCGCGAGAGCACGAC
ATCTCGTACAACGAGCAGCTTGAGGCGGGAGACAACCCCTACCTCAAGTACAACCACGCGGACGCCGAGTTTCAG
GAGAAGCTCGCCGACGACACATCCTTCGGGGGAAACCTCGGAAAGGCAGTCTTTCAGGCCAAGAAAAGGGTTCTC
GAACCTTTTGGCCTGGTTGAAGAGGGTGCTAAGACGGCCCCTACCGGAAAGCGGATAGACGACCACTTTCCAAAA
AGAAAGAAGGCTCGGACCGAAGAGGACTCCAAGCCTTCCACCTCGTCAGACGCCGAAGCTGGACCCAGCGGATCC
CAGCAGCTGCAAATCCCAGCCCAACCAGCCTCAAGTTTGGGAGCTGATACAATGTCTGCGGGAGGTGGCGGCCCA
TTGGGCGACAATAACCAAGGTGCCGATGGAGTGGGCAATGCCTCGGGAGATTGGCATTGCGATTCCACGTGGATG
GGGGACAGAGTCGTCACCAAGTCCACCCGAACCTGGGTGCTGCCCAGCTACAACAACCACCAGTACCGAGAGATC
AAAAGCGGCTCCGTCGACGGAAGCAACGCCAACGCCTACTTTGGATACAGCACCCCCTGGGGGTACTTTGACTTT
AACCGCTTCCACAGCCACTGGAGCCCCCGAGACTGGCAAAGACTCATCAACAACTACTGGGGCTTCAGACCCCGG
TCCCTCAGAGTCAAAATCTTCAACATTCAAGTCAAAGAGGTCACGGTGCAGGACTCCACCACCACCATCGCCAAC
AACCTCACCTCCACCGTCCAAGTGTTTACGGACGACGACTACCAGCTGCCCTACGTCGTCGGCAACGGGACCGAG
GGATGCCTGCCGGCCTTCCCTCCGCAGGTCTTTACGCTGCCGCAGTACGGTTACGCGACGCTGAACCGCGACAAC
ACAGAAAATCCCACCGAGAGGAGCAGCTTCTTCTGCCTAGAGTACTTTCCCAGCAAGATGCTGAGAACGGGCAAC
AACTTTGAGTTTACCTACAACTTTGAGGAGGTGCCCTTCCACTCCAGCTTCGCTCCCAGTCAGAACCTGTTCAAG
CTGGCCAACCCGCTGGTGGACCAGTACTTGTACCGCTTCGTGAGCACAAATAACACTGGCGGAGTCCAGTTCAAC
AAGAACCTGGCCGGGAGATACGCCAACACCTACAAAAACTGGTTCCCGGGGCCCATGGGCCGAACCCAGGGCTGG
AACCTGGGCTCCGGGGTCAACCGCGCCAGTGTCAGCGCCTTCGCCACGACCAATAGGATGGAGCTCGAGGGCGCG
AGTTACCAGGTGCCCCCGCAGCCGAACGGCATGACCAACAACCTCCAGGGCAGCAACACCTATGCCCTGGAGAAC
ACTATGATCTTCAACAGCCAGCCGGCGAACCCGGGCACCACCGCCACGTACCTCGAGGGCAACATGCTCATCACC
AGCGAGAGCGAGACGCAGCCGGTGAACCGCGTGGCGTACAACGTCGGCGGGCAGATGGCCACCAACAACCAGAGC
TCCACCACTGCCCCCGCGACCGGCACGTACAACCTCCAGGAAATCGTGCCCGGCAGCGTGTGGATGGAGAGGGAC
GTGTACCTCCAAGGACCCATCTGGGCCAAGATCCCAGAGACGGGGGCGCACTTTCACCCCTCTCCGGCCATGGGC
GGATTCGGACTCAAACACCCACCGCCCATGATGCTCATCAAGAACACGCCTGTGCCCGGAAATATCACCAGCTTC
TCGGACGTGCCCGTCAGCAGCTTCATCACCCAGTACAGCACCGGGCAGGTCACCGTGGAGATGGAGTGGGAGCTC
AAGAAGGAAAACTCCAAGAGGTGGAACCCAGAGATCCAGTACACAAACAACTACAACGACCCCCAGTTTGTGGAC
TTTGCCCCGGACAGCACCGGGGAATACAGAACCACCAGACCTATCGGAACCCGATACCTTACCCGACCCCTTTAA

AAV-6
Full Genome: AF028704

Rep78: (SEQ ID NO:15)

ATGCCGGGGTTTTACGAGATTGTGATTAAGGTCCCCAGCGACCTTGACGAGCATCTGCCCGGCATTTCTGACAGC
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGTTGCCGCCAGATTCTGACATGGATCTGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTCCTGGTCCAGTGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTTCAGTTCGAGAAGGGCGAGTCCTACTTCCACCTCCATATTCTGGTGGAGACCACGGGGGTC
AAATCCATGGTGCTGGGCCGCTTCCTGAGTCAGATTAGGGACAAGCTGGTGCAGACCATCTACCGCGGGATCGAG
CCGACCCTGCCCAACTGGTTCGCGGTGACCAAGACGCGTAATGGCGCCGGAGGGGGGAACAAGGTGGTGGACGAG

TGCTACATCCCCAACTACCTCCTGCCCAAGACTCAGCCCGAGCTGCAGTGGGCGTGGACTAACATGGAGGAGTAT
ATAAGCGCGTGTTTAAACCTGGCCGAGCGCAAACGGCTCGTGGCGCACGACCTGACCCACGTCAGCCAGACCCAG
GAGCAGAACAAGGAGAATCTGAACCCCAATTCTGACGCGCCTGTCATCCGGTCAAAAACCTCCGCACGCTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGGGGCATCACCTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCGTAC
ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCCCAGATCAAGGCCGCTCTGGACAATGCCGGCAAGATCATGGCG
CTGACCAAATCCGCGCCCGACTACCTGGTAGGCCCCGCTCCGCCCGCCGACATTAAAACCAACCGCATTTACCGC
ATCCTGGAGCTGAACGGCTACGACCCTGCCTACGCCGGCTCCGTCTTTCTCGGCTGGGCCCAGAAAAGGTTCGGA
AAACGCAACACCATCTGGCTGTTTGGGCCGGCCACCACGGGCAAGACCAACATCGCGGAAGCCATCGCCCACGCC
GTGCCCTTCTACGGCTGCGTCAACTGGACCAATGAGAACTTTCCCTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTCGGCGGCAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCGTCCGCCCAGATCGATCCCACCCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCGTTGCAGGACCGGATGTTCAAATTTGAACTC
ACCCGCCGTCTGGAGCATGACTTTGGCAAGGTGACAAAGCAGGAAGTCAAAGAGTTCTTCCGCTGGGCGCAGGAT
CACGTGACCGAGGTGGCGCATGAGTTCTACGTCAGAAAGGGTGGAGCCAACAAGAGACCCGCCCCCGATGACGCG
GATAAAAGCGAGCCCAAGCGGGCCTGCCCCTCAGTCGCGGATCCATCGACGTCAGACGCGGAAGGAGCTCCGGTG
GACTTTGCCGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGCATGCTTCAGATGCTGTTTCCCTGCAAAACA
TGCGAGAGAATGAATCAGAATTTCAACATTTGCTTCACGCACGGGACCAGAGACTGTTCAGAATGTTTCCCCGGC
GTGTCAGAATCTCAACCGGTCGTCAGAAAGAGGACGTATCGGAAACTCTGTGCCATTCATCATCTGCTGGGGCGG
GCTCCCGAGATTGCTTGCTCGGCCTGCGATCTGGTCAACGTGGATCTGGATGACTGTGTTTCTGAGCAATAA

CapVP1: (SEQ ID NO:16)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAGGGCATTCGCGAGTGGTGGGACTTG
AAACCTGGAGCCCCGAAACCCAAAGCCAACCAGCAAAAGCAGGACGACGGCCGGGGTCTGGTGCTTCCTGGCTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGGGAGCCCGTCAACGCGGCGGATGCAGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTCAAAGCGGGTGACAATCCGTACCTGCGGTATAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTGCAAGAAGATACGTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAGAAGAGGGTT
CTCGAACCTTTTGGTCTGGTTGAGGAAGGTGCTAAGACGGCTCCTGGAAAGAAACGTCCGGTAGAGCAGTCGCCA
CAAGAGCCAGACTCCTCCTCGGGCATTGGCAAGACAGGCCAGCAGCCCGCTAAAAAGAGACTCAATTTTGGTCAG
ACTGGCGACTCAGAGTCAGTCCCCGACCCACAACCTCTCGGAGAACCTCCAGCAACCCCCGCTGCTGTGGGACCT
ACTACAATGGCTTCAGGCGGTGGCGCACCAATGGCAGACAATAACGAAGGCGCCGACGGAGTGGGTAATGCCTCA
GGAAATTGGCATTGCGATTCCACATGGCTGGGCGACAGAGTCATCACCACCAGCACCCGAACATGGGCCTTGCCC
ACCTATAACAACCACCTCTACAAGCAAATCTCCAGTGCTTCAACGGGGGCCAGCAACGACAACCACTACTTCGGC
TACAGCACCCCCTGGGGGTATTTTGATTTCAACAGATTCCACTGCCATTTCTCACCACGTGACTGGCAGCGACTC
ATCAACAACAATTGGGGATTCCGGCCCAAGAGACTCAACTTCAAGCTCTTCAACATCCAAGTCAAGGAGGTCACG
ACGAATGATGGCGTCACGACCATCGCTAATAACCTTACCAGCACGGTTCAAGTCTTCTCGGACTCGGAGTACCAG
TTGCCGTACGTCCTCGGCTCTGCGCACCAGGGCTGCCTCCCTCCGTTCCCGGCGGACGTGTTCATGATTCCGCAG
TACGGCTACCTAACGCTCAACAATGGCAGCCAGGCAGTGGGACGGTCATCCTTTTACTGCCTGGAATATTTCCCA
TCGCAGATGCTGAGAACGGGCAATAACTTTACCTTCAGCTACACCTTCGAGGACGTGCCTTTCCACAGCAGCTAC
GCGCACAGCCAGAGCCTGGACCGGCTGATGAATCCTCTCATCGACCAGTACCTGTATTACCTGAACAGAACTCAG
AATCAGTCCGGAAGTGCCCAAAACAAGGACTTGCTGTTTAGCCGGGGGTCTCCAGCTGGCATGTCTGTTCAGCCC
AAAAACTGGCTACCTGGACCCTGTTACCGGCAGCAGCGCGTTTCTAAAACAAAAACAGACAACAACAACAGCAAC
TTTACCTGGACTGGTGCTTCAAAATATAACCTTAATGGGCGTGAATCTATAATCAACCCTGGCACTGCTATGGCC
TCACACAAAGACGACAAAGACAAGTTCTTTCCCATGAGCGGTGTCATGATTTTTGGAAAGGAGAGCGCCGGAGCT
TCAAACACTGCATTGGACAATGTCATGATCACAGACGAAGAGGAAATCAAAGCCACTAACCCCGTGGCCACCGAA
AGATTTGGGACTGTGGCAGTCAATCTCCAGAGCAGCAGCACAGACCCTGCGACCGGAGATGTGCATGTTATGGGA
GCCTTACCTGGAATGGTGTGGCAAGACAGAGACGTATACCTGCAGGGTCCTATTTGGGCCAAAATTCCTCACACG
GATGGACACTTTCACCCGTCTCCTCTCATGGGCGGCTTTGGACTTAAGCACCCGCCTCCTCAGATCCTCATCAAA
AACACGCCTGTTCCTGCGAATCCTCCGGCAGAGTTTTCGGCTACAAAGTTTGCTTCATTCATCACCCAGTATTCC
ACAGGACAAGTGAGCGTGGAGATTGAATGGGAGCTGCAGAAAGAAAACAGCAAACGCTGGAATCCCGAAGTGCAG
TATACATCTAACTATGCAAAATCTGCCAACGTTGATTTCACTGTGGACAACAATGGACTTTATACTGAGCCTCGC
CCCATTGGCACCCGTTACCTCACCCGTCCCCTGTAA

AAV-7
Full Genome: NC_006260

Rep78: (SEQ ID NO:17)

ATGCCGGGTTTCTACGAGATCGTGATCAAGGTGCCGAGCGACCTGGACGAGCACCTGCCGGGCATTTCTGACTCG
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGCTGCCCCCGGATTCTGACATGGATCTGAATCTGATCGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTCCTGGTCCAATGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTGTTCTTTGTTCAGTTCGAGAAGGGCGAGAGCTACTTCCACCTTCACGTTCTGGTGGAGACCACGGGGGTC
AAGTCCATGGTGCTAGGCCGCTTCCTGAGTCAGATTCGGGAGAAGCTGGTCCAGACCATCTACCGCGGGGTCGAG
CCCACGCTGCCCAACTGGTTCGCGGTGACCAAGACGCGTAATGGCGCCGGCGGGGGGAACAAGGTGGTGGACGAG
TGCTACATCCCCAACTACCTCCTGCCCAAGACCCAGCCCGAGCTGCAGTGGGCGTGGACTAACATGGAGGAGTAT
ATAAGCGCGTGTTTGAACCTGGCCGAACGCAAACGGCTCGTGGCGCAGCACCTGACCCACGTCAGCCAGACGCAG
GAGCAGAACAAGGAGAATCTGAACCCCAATTCTGACGCGCCCGTGATCAGGTCAAAAACCTCCGCGCGCTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGGGGCATCACCTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCGTAC
ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCCCAGATCAAGGCCGCGCTGGACAATGCCGGCAAGATCATGGCG
CTGACCAAATCCGCGCCCGACTACCTGGTGGGGCCCTCGCTGCCCGCGGACATTAAAACCAACCGCATCTACCGC
ATCCTGGAGCTGAACGGGTACGATCCTGCCTACGCCGGCTCCGTCTTTCTCGGCTGGGCCCAGAAAAAGTTCGGG
AAGCGCAACACCATCTGGCTGTTTGGGCCCGCCACCACCGGCAAGACCAACATTGCGGAAGCCATCGCCCACGCC
GTGCCCTTCTACGGCTGCGTCAACTGGACCAATGAGAACTTTCCCTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTCGGCGGCAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCGTCCGCCCAGATCGACCCCACCCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCGTTGCAGGACCGGATGTTCAAATTTGAACTC
ACCCGCCGTCTGGAGCACGACTTTGGCAAGGTGACGAAGCAGGAAGTCAAAGAGTTCTTCCGCTGGGCCAGTGAT
CACGTGACCGAGGTGGCGCATGAGTTCTACGTCAGAAAGGGCGGAGCCAGCAAAAGACCCGCCCCCGATGACGCG
GATATAAGCGAGCCCAAGCGGGCCTGCCCCTCAGTCGCGGATCCATCGACGTCAGACGCGGAAGGAGCTCCGGTG
GACTTTGCCGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGCATGATTCAGATGCTGTTTCCCTGCAAAACG
TGCGAGAGAATGAATCAGAATTTCAACATTTGCTTCACACACGGGGTCAGAGACTGTTTAGAGTGTTTCCCCGGC
GTGTCAGAATCTCAACCGGTCGTCAGAAAAAAGACGTATCGGAAACTCTGCGCGATTCATCATCTGCTGGGGCGG
GCGCCCGAGATTGCTTGCTCGGCCTGCGACCTGGTCAACGTGGACCTGGACGACTGCGTTTCTGAGCAATAA

CapVP1: (SEQ ID NO:18)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAGGGCATTCGCGAGTGGTGGGACCTG
AAACCTGGAGCCCCGAAACCCAAAGCCAACCAGCAAAAGCAGGACAACGGCCGGGGTCTGGTGCTTCCTGGCTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGGGAGCCCGTCAACGCGGCGGACGCAGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTCAAAGCGGGTGACAATCCGTACCTGCGGTATAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTGCAAGAAGATACGTCATTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAGAAGCGGGTT
CTCGAACCTCTCGGTCTGGTTGAGGAAGGCGCTAAGACGGCTCCTGCAAAGAAGAGACCGGTAGAGCCGTCACCT
CAGCGTTCCCCCGACTCCTCCACGGGCATCGGCAAGAAAGGCCAGCAGCCCGCCAGAAAGAGACTCAATTTCGGT
CAGACTGGCGACTCAGAGTCAGTCCCCGACCCTCAACCTCTCGGAGAACCTCCAGCAGCGCCCTCTAGTGTGGGA
TCTGGTACAGTGGCTGCAGGCGGTGGCGCACCAATGGCAGACAATAACGAAGGTGCCGACGGAGTGGGTAATGCC
TCAGGAAATTGGCATTGCGATTCCACATGGCTGGGCGACAGAGTCATTACCACCAGCACCCGAACCTGGGCCCTG
CCCACCTACAACAACCACCTCTACAAGCAAATCTCCAGTGAAACTGCAGGTAGTACCAACGACAACACCTACTTC
GGCTACAGCACCCCCTGGGGGTATTTTGACTTTAACAGATTCCACTGCCACTTCTCACCACGTGACTGGCAGCGA
CTCATCAACAACAACTGGGGATTCCGGCCCAAGAAGCTGCGGTTCAAGCTCTTCAACATCCAGGTCAAGGAGGTC
ACGACGAATGACGGCGTTACGACCATCGCTAATAACCTTACCAGCACGATTCAGGTATTCTCGGACTCGGAATAC
CAGCTGCCGTACGTCCTCGGCTCTGCGCACCAGGGCTGCCTGCCTCCGTTCCCGGCGGACGTCTTCATGATTCCT
CAGTACGGCTACCTGACTCTCAACAATGGCAGTCAGTCTGTGGGACGTTCCTCCTTCTACTGCCTGGAGTACTTC
CCCTCTCAGATGCTGAGAACGGGCAACAACTTTGAGTTCAGCTACAGCTTCGAGGACGTGCCTTTCCACAGCAGC
TACGCACACAGCCAGAGCCTGGACCGGCTGATGAATCCCCTCATCGACCAGTACTTGTACTACCTGGCCAGAACA
CAGAGTAACCCAGGAGGCACAGCTGGCAATCGGGAACTGCAGTTTTACCAGGGCGGGCCTTCAACTATGGCCGAA
CAAGCCAAGAATTGGTTACCTGGACCTTGCTTCCGGCAACAAAGAGTCTCCAAAACGCTGGATCAAAACAACAAC
AGCAACTTTGCTTGGACTGGTGCCACCAAATATCACCTGAACGGCAGAAACTCGTTGGTTAATCCCGGCGTCGCC
ATGGCAACTCACAAGGACGACGAGGACCGCTTTTTCCCATCCAGCGGAGTCCTGATTTTTGGAAAAACTGGAGCA
ACTAACAAAACTACATTGGAAAATGTGTTAATGACAAATGAAGAAGAAATTCGTCCTACTAATCCTGTAGCCACG
GAAGAATACGGGATAGTCAGCAGCAACTTACAAGCGGCTAATACTGCAGCCCAGACACAAGTTGTCAACAACCAG
GGAGCCTTACCTGGCATGGTCTGGCAGAACCGGGACGTGTACCTGCAGGGTCCCATCTGGGCCAAGATTCCTCAC
ACGGATGGCAACTTTCACCCGTCTCCTTTGATGGGCGGCTTTGGACTTAAACATCCGCCTCCTCAGATCCTGATC
AAGAACACTCCCGTTCCCGCTAATCCTCCGGAGGTGTTTACTCCTGCCAAGTTTGCTTCGTTCATCACACAGTAC
AGCACCGGACAAGTCAGCGTGGAAATCGAGTGGGAGCTGCAGAAGGAAAACAGCAAGCGCTGGAACCCGGAGATT
CAGTACACCTCCAACTTTGAAAAGCAGACTGGTGTGGACTTTGCCGTTGACAGCCAGGGTGTTTACTCTGAGCCT
CGCCCTATTGGCACTCGTTACCTCACCCGTAATCTGTAA

AAV-8

Full Genome: NC_006261

Rep78: (SEQ ID NO:19)

ATGCCGGGCTTCTACGAGATCGTGATCAAGGTGCCGAGCGACCTGGACGAGCACCTGCCGGGCATTTCTGACTCG
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGCTGCCCCCGGATTCTGACATGGATCGGAATCTGATCGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTCCTGGTCCAATGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTTCAGTTCGAGAAGGGCGAGAGCTACTTTCACCTGCACGTTCTGGTCGAGACCACGGGGGTC
AAGTCCATGGTGCTAGGCCGCTTCCTGAGTCAGATTCGGGAAAAGCTTGGTCCAGACCATCTACCCGCGGGGTCG
AGCCCCACCTTGCCCAACTGGTTCGCGGTGACCAAAGACGCGGTAATGGCGCCGGCGGGGGGGAACAAGGTGGTG
GACGAGTGCTACATCCCCAACTACCTCCTGCCCAAGACTCAGCCCGAGCTGCAGTGGGCGTGGACTAACATGGAG
GAGTATATAAGCGCGTGCTTGAACCTGGCCGAGCGCAAACGGCTCGTGGCGCAGCACCTGACCCACGTCAGCCAG
ACGCAGGAGCAGAACAAGGAGAATCTGAACCCCAATTCTGACGCGCCCGTGATCAGGTCAAAAACCTCCGCGCGC
TATATGGAGCTGGTCGGGTGGCTGGTGGACCGGGGCATCACCTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCC
TCGTACATCTCCTTCAACGCCGCCTCCAACTCGCGGTCCCAGATCAAGGCCGCGCTGGACAATGCCGGCAAGATC
ATGGCGCTGACCAAATCCGCGCCCGACTACCTGGTGGGGCCCTCGCTGCCCGCGGACATTACCCAGAACCGCATC
TACCGCATCCTCGCTCTCAACGGCTACGACCCTGCCTACGCCGGCTCCGTCTTTCTCGGCTGGGCTCAGAAAAAG
TTCGGGAAACGCAACACCATCTGGCTGTTTGGACCCGCCACCACCGGCAAGACCAACATTGCGGAAGCCATCGCC
CACGCCGTGCCCTTCTACGGCTGCGTCAACTGGACCAATGAGAACTTTCCCTTCAATGATTGCGTCGACAAGATG
GTGATCTGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTCGGCGGCAGCAAG
GTGCGCGTGGACCAAAAGTGCAAGTCGTCCGCCCAGATCGACCCCACCCCCGTGATCGTCACCTCCAACACCAAC
ATGTGCGCCGTGATTGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCTCTCCAGGACCGGATGTTTAAGTTC
GAACTCACCCGCCGTCTGGAGCACGACTTTGGCAAGGTGACAAAGCAGGAAGTCAAAGAGTTCTTCCGCTGGGCC
AGTGATCACGTGACCGAGGTGGCGCATGAGTTTTACGTCAGAAAGGGCGGAGCCAGCAAAAGACCCGCCCCCGAT
GACGCGGATAAAAGCGAGCCCAAGCGGGCCTGCCCCTCAGTCGCGGATCCATCGACGTCAGACGCGGAAGGAGCT
CCGGTGGACTTTGCCGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGCATGCTTCAGATGCTGTTTCCCTGC
AAAACGTGCGAGAGAATGAATCAGAATTTCAACATTTGCTTCACACACGGGGTCAGAGACTGCTCAGAGTGTTTC
CCCGGCGTGTCAGAATCTCAACCGGTCGTCAGAAAGAGGACGTATCGGAAACTCTGTGCGATTCATCATCTGCTG
GGGCGGGCTCCCGAGATTGCTTGCTCGGCCTGCGATCTGGTCAACGTGGACCTGGATGACTGTGTTTCTGAGCAA
TAA

CapVP1: (SEQ ID NO:20)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAGGGCATTCGCGAGTGGTGGGCGCTG
AAACCTGGAGCCCCGAAGCCCAAAGCCAACCAGCAAAAGCAGGACGACGGCCGGGGTCTGGTGCTTCCTGGCTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGGGAGCCCGTCAACGCGGCGGACGCAGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTGCAGGCGGGTGACAATCCGTACCTGCGGTATAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTGCAAGAAGATACGTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAGAAGCGGGTT
CTCGAACCTCTCGGTCTGGTTGAGGAAGGCGCTAAGACGGCTCCTGGAAAGAAGAGACCGGTAGAGCCATCACCC
CAGCGTTCTCCAGACTCCTCTACGGGCATCGGCAAGAAAGGCCAACAGCCCGCCAGAAAAAGACTCAATTTTGGT
CAGACTGGCGACTCAGAGTCAGTTCCAGACCCTCAACCTCTCGGAGAACCTCCAGCAGCGCCCTCTGGTGTGGGA
CCTAATACAATGGCTGCAGGCGGTGGCGCACCAATGGCAGACAATAACGAAGGCGCCGACGGAGTGGGTAGTTCC
TCGGGAAATTGGCATTGCGATTCCACATGGCTGGGCGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTG
CCCACCTACAACAACCACCTCTACAAGCAAATCTCCAACGGGACATCGGGAGGAGCCACCAACGACAACACCTAC
TTCGGCTACAGCACCCCCTGGGGGTATTTTGACTTTAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAG
CGACTCATCAACAACAACTGGGGATTCCGGCCCAAGAGACTCAGCTTCAAGCTCTTCAACATCCAGGTCAAGGAG
GTCACGCAGAATGAAGGCACCAAGACCATCGCCAATAACCTCACCAGCACCATCCAGGTGTTTACGGACTCGGAG
TACCAGCTGCCGTACGTTCTCGGCTCTGCCCACCAGGGCTGCCTGCCTCCGTTCCCGGCGGACGTGTTCATGATT
CCCCAGTACGGCTACCTAACACTCAACAACGGTAGTCAGGCCGTGGGACGCTCCTCCTTCTACTGCCTGGAATAC
TTTCCTTCGCAGATGCTGAGAACCGGCAACAACTTCCAGTTTACTTACACCTTCGAGGACGTGCCTTTCCACAGC
AGCTACGCCCACAGCCAGAGCTTGGACCGGCTGATGAATCCTCTGATTGACCAGTACCTGTACTACTTGTCTCGG
ACTCAAACAACAGGAGGCACGGCAAATACGCAGACTCTGGGCTTCAGCCAAGGTGGGCCTAATACAATGGCCAAT
CAGGCAAAGAACTGGCTGCCAGGACCCTGTTACCGCCAACAACGCGTCTCAACGACAACCGGGCAAAACAACAAT
AGCAACTTTGCCTGGACTGCTGGGACCAAATACCATCTGAATGGAAGAAATTCATTGGCTAATCCTGGCATCGCT
ATGGCAACACACAAAGACGACGAGGAGCGTTTTTTTCCCAGTAACGGGATCCTGATTTTTGGCAAACAAAATGCT
GCCAGAGACAATGCGGATTACAGCGATGTCATGCTCACCAGCGAGGAAGAAATCAAAACCACTAACCCTGTGGCT
ACAGAGGAATACGGTATCGTGGCAGATAACTTGCAGCAGCAAAACACGGCTCCTCAAATTGGAACTGTCAACAGC

CAGGGGGCCTTACCCGGTATGGTCTGGCAGAACCGGGACGTGTACCTGCAGGGTCCCATCTGGGCCAAGATTCCT
CACACGGACGGCAACTTCCACCCGTCTCCGCTGATGGGCGGCTTTGGCCTGAAACATCCTCCGCCTCAGATCCTG
ATCAAGAACACGCCTGTACCTGCGGATCCTCCGACCACCTTCAACCAGTCAAAGCTGAACTCTTTCATCACGCAA
TACAGCACCGGACAGGTCAGCGTGGAAATTGAATGGGAGCTGCAGAAGGAAAACAGCAAGCGCTGGAACCCCGAG
ATCCAGTACACCTCCAACTACTACAAATCTACAAGTGTGGACTTTGCTGTTAATACAGAAGGCGTGTACTCTGAA
CCCCGCCCCATTGGCACCCGTTACCTCACCCGTAATCTGTAA

AAV-9
Cap only: AY530579

CapVP1: (SEQ ID NO:21)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACAACCTTAGTGAAGGAATTCGCGAGTGGTGGGCTTTG
AAACCTGGAGCCCCTCAACCCAAGGCAAATCAACAACATCAAGACAACGCTCGAGGTCTTGTGCTTCCGGGTTAC
AAATACCTTGGACCCGGCAACGGACTCGACAAGGGGGAGCCGGTCAACGCAGCAGACGCGGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTCAAGGCCGGAGACAACCCGTACCTCAAGTACAACCACGCCGACGCCGAGTTC
CAGGAGCGGCTCAAAGAAGATACGTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAAAAGAGGCTT
CTTGAACCTCTTGGTCTGGTTGAGGAAGCGGCTAAGACGGCTCCTGGAAAGAAGAGGCCTGTAGAGCAGTCTCCT
CAGGAACCGGACTCCTCCGCGGGTATTGGCAAATCGGGTGCACAGCCCGCTAAAAAGAGACTCAATTTCGGTCAG
ACTGGCGACACAGAGTCAGTCCCAGACCCTCAACCAATCGGAGAACCTCCCGCAGCCCCCTCAGGTGTGGGATCT
CTTACAATGGCTTCAGGTGGTGGCGCACCAGTGGCAGACAATAACGAAGGTGCCGATGGAGTGGGTAGTTCCTCG
GGAAATTGGCATTGCGATTCCCAATGGCTGGGGGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTGCCC
ACCTACAACAATCACCTCTACAAGCAAATCTCCAACAGCACATCTGGAGGATCTTCAAATGACAACGCCTACTTC
GGCTACAGCACCCCCTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTCTCACCACGTGACTGGCAGCGA
CTCATCAACAACAACTGGGGATTCCGGCCTAAGCGACTCAACTTCAAGCTCTTCAACATTCAGGTCAAAGAGGTT
ACGGACAACAATGGAGTCAAGACCATCGCCAATAACCTTACCAGCACGGTCCAGGTCTTCACGGACTCAGACTAT
CAGCTCCCGTACGTGCTCGGGTCGGCTCACGAGGGCTGCCTCCCGCCGTTCCCAGCGGACGTTTTCATGATTCCT
CAGTACGGGTATCTGACGCTTAATGATGGAAGCCAGGCCGTGGGTCGTTCGTCCTTTTACTGCCTGGAATATTTC
CCGTCGCAAATGCTAAGAACGGGTAACAACTTCCAGTTCAGCTACGAGTTTGAGAACGTACCTTTCCATAGCAGC
TACGCTCACAGCCAAAGCCTGGACCGACTAATGAATCCACTCATCGACCAATACTTGTACTATCTCTCAAAGACT
ATTAACGGTTCTGGACAGAATCAACAAACGCTAAAATTCAGTGTGGCCGGACCCAGCAACATGGCTGTCCAGGGA
AGAAACTACATACCTGGACCCAGCTACCGACAACAACGTGTCTCAACCACTGTGACTCAAAACAACAACAGCGAA
TTTGCTTGGCCTGGAGCTTCTTCTTGGGCTCTCAATGGACGTAATAGCTTGATGAATCCTGGACCTGCTATGGCC
AGCCACAAAGAAGGAGAGGACCGTTTCTTTCCTTTGTCTGGATCTTTAATTTTTGGCAAACAAGGAACTGGAAGA
GACAACGTGGATGCGGACAAAGTCATGATAACCAACGAAGAAGAAATTAAAACTACTAACCCGGTAGCAACGGAG
TCCTATGGACAAGTGGCCACAAACCACCAGAGTGCCCAAGCACAGGCGCAGACCGGCTGGGTTCAAAACCAAGGA
ATACTTCCGGGTATGGTTTGGCAGGACAGAGATGTGTACCTGCAAGGACCCATTTGGGCCAAAATTCCTCACACG
GACGGCAACTTTCACCCTTCTCCGCTGATGGGAGGGTTTGGAATGAAGCACCCGCCTCCTCAGATCCTCATCAAA
AACACACCTGTACCTGCGGATCCTCCAACGGCCTTCAACAAGGACAAGCTGAACTCTTTCATCACCCAGTATTCT
ACTGGCCAAGTCAGCGTGGAGATCGAGTGGGAGCTGCAGAAGGAAAACAGCAAGCGCTGGAACCCGGAGATCCAG
TACACTTCCAACTATTACAAGTCTAATAATGTTGAATTTGCTGTTAATACTGAAGGTGTATATAGTGAACCCCGC
CCCATTGGCACCAGATACCTGACTCGTAATCTGTAA

AAV-10
Partial Genome: AY631965

Rep78: (SEQ ID NO:22)

ATGCCGGGCTTCTACGAGATCGTGATCAAGGTGCCGAGCGACCTGGACGAGCACCTGCCGGGCATTTCTGACTCG
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGCTGCCCCCGGATTCTGACATGGATCGGAATCTGATCGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTCCTGGTCCACTGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTTCAGTTCGAGAAGGGCGAGTCCTACTTTCACCTGCACGTTCTGGTCGAGACCACGGGGGTC
AAGTCCATGGTCCTGGGCCGCTTCCTGAGTCAGATCAGAGACAGGCTGGTGCAGACCATCTACCGCGGGGTAGAG
CCCACGCTGCCCAACTGGTTCGCGGTGACCAAGACGCGAAATGGCGCCGGCGGGGGGAACAAGGTGGTGGACGAG
TGCTACATCCCCAACTACCTCCTGCCCAAGACGCAGCCCGAGCTGCAGTGGGCGTGGACTAACATGGAGGAGTAT
ATAAGCGCGTGTCTGAACCTCGCGGAGCGTAAACGGCTCGTGGCGCAGCACCTGACCCACGTCAGCCAGACGCAG
GAGCAGAACAAGGAGAATCTGAACCCGAATTCTGACGCGCCCGTGATCAGGTCAAAAACCTCCGCGCGCTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGGGGCATCACCTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCGTAC

ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCCCAGATCAAGGCCGCGCTGGACAATGCCGGAAAGATCATGGCG
CTGACCAAATCCGCGCCCGACTACCTGGTAGGCCCGTCCTTACCCGCGGACATTAAGGCCAACCGCATCTACCGC
ATCCTGGAGCTCAACGGCTACGACCCCGCCTACGCCGGCTCCGTCTTCCTGGGCTGGGCGCAGAAAAAGTTCGGT
AAAAGGAATACAATTTGGCTGTTCGGGCCCGCCACCACCGGCAAGACCAACATCGCGGAAGCCATCGCCCACGCC
GTGCCCTTCTACGGCTGCGTCAACTGGACCAATGAGAACTTTCCCTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACCGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTGGGCGGAAGCAAGGTGCGC
GTCGACCAAAAGTGCAAGTCCTCGGCCCAGATCGACCCCACGCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATCGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCCCTGCAGGACCGCATGTTCAAGTTCGAGCTC
ACCCGCCGTCTGGAGCACGACTTTGGCAAGGTGACCAAGCAGGAAGTCAAAGAGTTCTTCCGCTGGGCTCAGGAT
CACGTGACTGAGGTGACGCATGAGTTCTACGTCAGAAAGGGCGGAGCCACCAAAAGACCCGCCCCCAGTGACGCG
GATATAAGCGAGCCCAAGCGGGCCTGCCCCTCAGTTGCGGAGCCATCGACGTCAGACGCGGAAGCACCGGTGGAC
TTTGCGGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGCATGCTTCAGATGCTGTTTCCCTGCAAGACATGC
GAGAGAATGAATCAGAATTTCAACGTCTGCTTCACGCACGGGGTCAGAGACTGCTCAGAGTGCTTCCCCGGCGCG
TCAGAATCTCAACCTGTCGTCAGAAAAAAGACGTATCAGAAACTGTGCGCGATTCATCATCTGCTGGGGCGGGCA
CCCGAGATTGCGTGTTCGGCCTGCGATCTCGTCAACGTGGACTTGGATGACTGTGTTTCTGAGCAATAA

CapVP1: (SEQ ID NO:23)

ATGGCTGCTGACGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAGGGCATTCGCGAGTGGTGGGACCTG
AAACCTGGAGCCCCCAAGCCCAAGGCCAACCAGCAGAAGCAGGACGACGGCCGGGGTCTGGTGCTTCCTGGCTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGGGAGCCCGTCAACGCGGCGGACGCAGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTCAAAGCGGGTGACAATCCGTACCTGCGGTATAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTGCAAGAAGATACGTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAGAAGCGGGTT
CTCGAACCTCTCGGTCTGGTTGAGGAAGCTGCTAAGACGGCTCCTGGAAAGAAGAGACCGGTAGAACCGTCACCT
CAGCGTTCCCCCGACTCCTCCACGGGCATCGGCAAGAAAGGCCAGCAGCCCGCTAAAAAGAGACTGAACTTTGGG
CAGACTGGCGAGTCAGAGTCAGTCCCCGACCCTCAACCAATCGGAGAACCACCAGCAGGCCCCTCTGGTCTGGGA
TCTGGTACAATGGCTGCAGGCGGTGGCGCTCCAATGGCAGACAATAACGAAGGCGCCGACGGAGTGGGTAGTTCC
TCAGGAAATTGGCATTGCGATTCCACATGGCTGGGCGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTG
CCCACCTACAACAACCACCTCTACAAGCAAATCTCCAACGGGACATCGGGAGGAAGCACCAACGACAACACCTAC
TTCGGCTACAGCACCCCCTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTCTCACCACGTGACTGGCAG
CGACTCATCAACAACAACTGGGGATTCCGGCCAAAAAGACTCAGCTTCAAGCTCTTCAACATCCAGGTCAAGGAG
GTCACGCAGAATGAAGGCACCAAGACCATCGCCAATAACCTTACCAGCACGATTCAGGTATTTACGGACTCGGAA
TACCAGCTGCCGTACGTCCTCGGCTCCGCGCACCAGGGCTGCCTGCCTCCGTTCCCGGCGGATGTCTTCATGATT
CCCCAGTACGGCTACCTGACACTGAACAATGGAAGTCAAGCCGTAGGCCGTTCCTCCTTCTACTGCCTGGAATAT
TTTCCATCTCAAATGCTGCGAACTGGAAACAATTTTGAATTCAGCTACACCTTCGAGGACGTGCCTTTCCACAGC
AGCTACGCACACAGCCAGAGCTTGGACCGACTGATGAATCCTCTCATTGACCAGTACCTGTACTACTTATCCAGA
ACTCAGTCCACAGGAGGAACTCAAGGTACCCAGCAATTGTTATTTTCTCAAGCTGGGCCTGCAAACATGTCGGCT
CAGGCCAAGAACTGGCTGCCTGGACCTTGCTACCGGCAGCAGCGAGTCTCCACGACACTGTCGCAAAACAACAAC
AGCAACTTTGCTTGGACTGGTGCCACCAAATATCACCTGAACGGAAGAGACTCTCTGGTGAATCCCGGTGTCGCC
ATGGCAACCCACAAGGACGACGAGGAACGCTTCTTCCCGTCGAGCGGAGTCCTGATGTTTGGAAAACAGGGTGCT
GGAAGAGACAATGTGGACTACAGCAGCGTTATGCTAACAAGCGAAGAAGAAATTAAAACCACTAACCCTGTAGCC
ACAGAACAATACGGCGTGGTGGCTGACAACTTGCAGCAAGCCAATACAGGGCCTATTGTGGGAAATGTCAACAGC
CAAGGAGCCTTACCTGGCATGGTCTGGCAGAACCGAGACGTGTACCTGCAGGGTCCCATCTGGGCCAAGATTCCT
CACACGGACGGCAACTTTCACCCGTCTCCTCTGATGGGCGGCTTTGGACTTAAACACCCGCCTCCACAGATCCTG
ATCAAGAACACGCCGGTACCTGCGGATCCTCCAACAACGTTCAGCCAGGCGAAATTGGCTTCCTTCATCACGCAG
TACAGCACCGGACAGGTCAGCGTGGAAATCGAGTGGGAGCTGCAGAAGGAGAACAGCAAACGCTGGAACCCAGAG
ATTCAGTACACTTCAAACTACTACAAATCTACAAATGTGGACTTTGCTGTCAATACAGAGGGAACTTATTCTGAG
CCTCGCCCCATTGGTACTCGTTATCTGACACGTAATCTGTAA

AAV-11
Partial Genome: AY631966

Rep78: (SEQ ID NO:24)

ATGCCGGGCTTCTACGAGATCGTGATCAAGGTGCCGAGCGACCTGGACGAGCACCTGCCGGGCATTTCTGACTCG
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGCTGCCCCCGGATTCTGACATGGATCGGAATCTGATCGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTCCTGGTCCACTGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTTCAGTTCGAGAAGGGCGAGTCCTACTTCCACCTCCACGTTCTCGTCGAGACCACGGGGGTC

AAGTCCATGGTCCTGGGCCGCTTCCTGAGTCAGATCAGAGACAGGCTGGTGCAGACCATCTACCGCGGGGTCGAG
CCCACGCTGCCCAACTGGTTCGCGGTGACCAAGACGCGAAATGGCGCCGGCGGGGGGAACAAGGTGGTGGACGAG
TGCTACATCCCCAACTACCTCCTGCCCAAGACCCAGCCCGAGCTGCAGTGGGCGTGGACTAACATGGAGGAGTAT
ATAAGCGCGTGTCTAAACCTCGCGGAGCGTAAACGGCTCGTGGCGCAGCACCTGACCCACGTCAGCCAGACGCAG
GAGCAGAACAAGGAGAATCTGAACCCGAATTCTGACGCGCCCGTGATCAGGTCAAAAACCTCCGCGCGCTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGGGGCATCACCTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCGTAC
ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCCCAGATCAAGGCCGCGCTGGACAATGCCGGAAAGATCATGGCG
CTGACCAAATCCGCGCCCGACTACCTGGTAGGCCCGTCCTTACCCGCGGACATTAAGGCCAACCGCATCTACCGC
ATCCTGGAGCTCAACGGCTACGACCCCGCCTACGCCGGCTCCGTCTTCCTGGGCTGGGCGCAGAAAAAGTTCGGT
AAACGCAACACCATCTGGCTGTTTGGGCCCGCCACCACCGGCAAGACCAACATCGCGGAAGCCATAGCCCACGCC
GTGCCCTTCTACGGCTGCGTGAACTGGACCAATGAGAACTTTCCCTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACCGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTGGGCGGAAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCCTCGGCCCAGATCGACCCCACGCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATCGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCGCTGCAGGACCGCATGTTCAAGTTCGAGCTC
ACCCGCCGTCTGGAGCACGACTTTGGCAAGGTGACCAAGCAGGAAGTCAAAGAGTTCTTCCGCTGGGCTCAGGAT
CACGTGACTGAGGTGGCGCATGAGTTCTACGTCAGAAAGGGCGGAGCCACCAAAAGACCCGCCCCCAGTGACGCG
GATATAAGCGAGCCCAAGCGGGCCTGCCCCTCAGTTCCGGAGCCATCGACGTCAGACGCGGAAGCACCGGTGGAC
TTTGCGGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGCATGCTTCAGATGCTGTTTCCCTGCAAGACATGC
GAGAGAATGAATCAGAATTTCAACGTCTGCTTCACGCACGGGGTCAGAGACTGCTCAGAGTGCTTCCCCGGCGCG
TCAGAATCTCAACCCGTCGTCAGAAAAAAGACGTATCAGAAACTGTGCGCGATTCATCATCTGCTGGGGCGGGCA
CCCGAGATTGCGTGTTCGGCCTGCGATCTCGTCAACGTGGACTTGGATGACTGTGTTTCTGAGCAATAA

CapVP1: (SEQ ID NO:25)

ATGGCTGCTGACGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAGGGCATTCGCGAGTGGTGGGACCTG
AAACCTGGAGCCCCGAAGCCCAAGGCCAACCAGCAGAAGCAGGACGACGGCCGGGGTCTGGTGCTTCCTGGCTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGGGAGCCCGTCAACGCGGCGGACGCAGCGGCCCTCGAGCAC
GACAAGGCCTACGACCAGCAGCTCAAAGCGGGTGACAATCCGTACCTGCGGTATAACCACGCCGACGCCGAGTTT
CAGGAGCGTCTGCAAGAAGATACGTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAGAAGAGGGTA
CTCGAACCTCTGGGCCTGGTTGAAGAAGGTGCTAAAACGGCTCCTGGAAAGAAGAGACCGTTAGAGTCACCACAA
GAGCCCGACTCCTCCTCGGGCATCGGCAAAAAAGGCAAACAACCAGCCAGAAAGAGGCTCAACTTTGAAGAGGAC
ACTGGAGCCGGAGACGGACCCCCTGAAGGATCAGATACCAGCGCCATGTCTTCAGACATTGAAATGCGTGCAGCA
CCGGGCGGAAATGCTGTCGATGCGGGACAAGGTTCCGATGGAGTGGGTAATGCCTCGGGTGATTGGCATTGCGAT
TCCACCTGGTCTGAGGGCAAGGTCACAACAACCTCGACCAGAACCTGGGTCTTGCCCACCTACAACAACCACTTG
TACCTGCGTCTCGGAACAACATCAAGCAGCAACACCTACAACGGATTCTCCACCCCCTGGGGATATTTTGACTTC
AACAGATTCCACTGTCACTTCTCACCACGTGACTGGCAAAGACTCATCAACAACAACTGGGGACTACGACCAAAA
GCCATGCGCGTTAAAATCTTCAATATCCAAGTTAAGGAGGTCACAACGTCGAACGGCGAGACTACGGTCGCTAAT
AACCTTACCAGCACGGTTCAGATATTTGCGGACTCGTCGTATGAGCTCCCGTACGTGATGGACGCTGGACAAGAG
GGGAGCCTGCCTCCTTTCCCCAATGACGTGTTCATGGTGCCTCAATATGGCTACTGTGGCATCGTGACTGGCGAG
AATCAGAACCAAACGGACAGAAACGCTTTCTACTGCCTGGAGTATTTTCCTTCGCAAATGTTGAGAACTGGCAAC
AACTTTGAAATGGCTTACAACTTTGAGAAGGTGCCGTTCCACTCAATGTATGCTCACAGCCAGAGCCTGGACAGA
CTGATGAATCCCCTCCTGGACCAGTACCTGTGGCACTTACAGTCGACTACCTCTGGAGAGACTCTGAATCAAGGC
AATGCAGCAACCACATTTGGAAAAATCAGGAGTGGAGACTTTGCCTTTTACAGAAAGAACTGGCTGCCTGGGCCT
TGTGTTAAACAGCAGAGATTCTCAAAAACTGCCAGTCAAAATTACAAGATTCCTGCCAGCGGGGGCAACGCTCTG
TTAAAGTATGACACCCACTATACCTTAAACAACCGCTGGAGCAACATCGCGCCCGGACCTCCAATGGCCACAGCC
GGACCTTCGGATGGGGACTTCAGTAACGCCCAGCTTATATTCCCTGGACCATCTGTTACCGGAAATACAACAACT
TCAGCCAACAATCTGTTGTTTACATCAGAAGAAGAAATTGCTGCCACCAACCCAAGAGACACGGACATGTTTGGC
CAGATTGCTGACAATAATCAGAATGCTACAACTGCTCCCATAACCGGCAACGTGACTGCTATGGGAGTGCTGCCT
GGCATGGTGTGGCAAAACAGAGACATTTACTACCAAGGGCCAATTTGGGCCAAGATCCCACACGCGGACGGACAT
TTTCATCCTTCACCGCTGATTGGTGGGTTTGGACTGAAACACCCGCCTCCCCAGATATTCATCAAGAACACTCCC
GTACCTGCCAATCCTGCGACAACCTTCACTGCAGCCAGAGTGGACTCTTTCATCACACAATACAGCACCGGCCAG
GTCGCTGTTCAGATTGAATGGGAAATTGAAAAGGAACGCTCCAAACGCTGGAATCCTGAAGTGCAGTTTACTTCA
AACTATGGGAACCAGTCTTCTATGTTGTGGGCTCCTGATACAACTGGGAAGTATACAGAGCCGCGGGTTATTGGC
TCTCGTTATTTGACTAATCATTTGTAA

AAV-12
Partial Genome: DQ813647

Rep78: (SEQ ID NO:26)

ATGCCGGGGTTCTACGAGGTGGTGATCAAGGTGCCCAGCGACCTGGACGAGCACCTGCCCGGCATTTCTGACTCC
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGTTGCCCCCGGATTCTGACATGGATCAGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGAGTTCCTGGTGGAATGGCGCCGAGTGAGTAAATTTCTGGAG
GCCAAGTTTTTTGTGCAGTTTGAAAAGGGGGACTCGTACTTTCATTTGCATATTCTGATTGAAATTACCGGCGTG
AAATCCATGGTGGTGGGCCGCTACGTGAGTCAGATTAGGGATAAACTGATCCAGCGCATCTACCGCGGGGTCGAG
CCCCAGCTGCCCAACTGGTTCGCGGTCACAAAGACCCGAAATGGCGCCGGAGGCGGGAACAAGGTGGTGGACGAG
TGCTACATCCCCAACTACCTGCTCCCCAAGGTCCAGCCCGAGCTTCAGTGGGCGTGGACTAACATGGAGGAGTAT
ATAAGCGCCTGTTTGAACCTCGCGGAGCGTAAACGGCTCGTGGCGCAGCACCTGACGCACGTCTCCCAGACCCAG
GAGGGCGACAAGGAGAATCTGAACCCGAATTCTGACGCGCCGGTGATCCGGTCAAAAACCTCCGCCAGGTACATG
GAGCTGGTCGGGTGGCTGGTGGACAAGGGCATCACGTCCGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCGTAC
ATCTCCTTCAACGCGGCCTCCAACTCCCGGTCGCAGATCAAGGCGGCCCTGGACAATGCCTCCAAAATCATGAGC
CTCACCAAAACGGCTCCGGACTATCTCATCGGGCAGCAGCCCGTGGGGGACATTACCACCAACCGGATCTACAAA
ATCCTGGAACTGAACGGGTACGACCCCCAGTACGCCGCCTCCGTCTTTCTCGGCTGGGCCCAGAAAAAGTTTGGA
AAGCGCAACACCATCTGGCTGTTTGGGCCCGCCACCACCGGCAAGACCAACATCGCGGAAGCCATCGCCCACGCG
GTCCCCTTCTACGGCTGCGTCAACTGGACCAATGAGAACTTTCCCTTCAACGACTGCGTCGACAAAATGGTGATT
TGGTGGGAGGAGGGCAAGATGACCGCCAAGGTCGTAGAGTCCGCCAAGGCCATTCTGGGCGGCAGCAAGGTGCGC
GTGGACCAAAAATGCAAGGCCTCTGCGCAGATCGACCCCACCCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACAGCACCACCTTCGAGCACCAGCAGCCCCTGCAGGACCGGATGTTCAAGTTTGAACTC
ACCCGCCGCCTCGACCACGACTTTGGCAAGGTCACCAAGCAGGAAGTCAAGGACTTTTTCCGGTGGGCGGCTGAT
CACGTGACTGACGTGGCTCATGAGTTTTACGTCACAAAGGGTGGAGCTAAGAAAAGGCCCGCCCCCTCTGACGAG
GATATAAGCGAGCCCAAGCGGCCGCGCGTGTCATTTGCGCAGCCGGAGACGTCAGACGCGGAAGCTCCCGGAGAC
TTCGCCGACAGGTACCAAAACAAATGTTCTCGTCACGCGGGTATGCTGCAGATGCTCTTTCCCTGCAAGACGTGC
GAGAGAATGAATCAGAATTCCAACGTCTGCTTCACGCACGGTCAGAAAGATTGCGGGGAGTGCTTTCCCGGGTCA
GAATCTCAACCGGTTTCTGTCGTCAGAAAAACGTATCAGAAACTGTGCATCCTTCATCAGCTCCGGGGGGCACCC
GAGATCGCCTGCTCTGCTTGCGACCAACTCAACCCCGATTTGGACGATTGCCAATTTGAGCAATAA

CapVP1: (SEQ ID NO:27)

ATGGCTGCTGACGGTTATCTTCCAGATTGGCTCGAGGACAACCTCTCTGAAGGCATTCGCGAGTGGTGGGCGCTG
AAACCTGGAGCTCCACAACCCAAGGCCAACCAACAGCATCAGGACAACGGCAGGGGTCTTGTGCTTCCTGGGTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGAGAGCCGGTCAACGAGGCAGACGCCGCGGCCCTCGAGCAC
GACAAGGCCTACGACAAGCAGCTCGAGCAGGGGGACAACCCGTATCTCAAGTACAACCACGCCGACGCCGAGTTC
CAGCAGCGCTTGGCGACCGACACCTCTTTTGGGGGCAACCTCGGGCGAGCAGTCTTCCAGGCCAAAAAGAGGATT
CTCGAGCCTCTGGGTCTGGTTGAAGAGGGCGTTAAAACGGCTCCTGGAAAGAAACGCCCATTAGAAAAGACTCCA
AATCGGCCGACCAACCCGGACTCTGGGAAGGCCCCGGCCAAGAAAAAGCAAAAAGACGGCGAACCAGCCGACTCT
GCTAGAAGGACACTCGACTTTGAAGACTCTGGAGCAGGAGACGGACCCCCTGAGGGATCATCTTCCGGAGAAATG
TCTCATGATGCTGAGATGCGTGCGGCGCCAGGCGGAAATGCTGTCGAGGCGGGACAAGGTGCCGATGGAGTGGGT
AATGCCTCCGGTGATTGGCATTGCGATTCCACCTGGTCAGAGGGCCGAGTCACCACCACCAGCACCCGAACCTGG
GTCCTACCCACGTACAACAACCACCTGTACCTGCGAATCGGAACAACGGCCAACAGCAACACCTACAACGGATTC
TCCACCCCCTGGGGATACTTTGACTTTAACCGCTTCCACTGCCACTTTTCCCCACGCGACTGGCAGCGACTCATC
AACAACAACTGGGGACTCAGGCCGAAATCGATGCGTGTTAAAATCTTCAACATACAGGTCAAGGAGGTCACGACG
TCAAACGGCGAGACTACGGTCGCTAATAACCTTACCAGCACGGTTCAGATCTTTGCGGATTCGACGTATGAACTC
CCATACGTGATGGACGCCGGTCAGGAGGGGAGCTTTCCTCCGTTTCCCAACGACGTCTTTATGGTTCCCCAATAC
GGATACTGCGGAGTTGTCACTGGAAAAAACCAGAACCAGACAGACAGAAATGCCTTTTACTGCCTGGAATACTTT
CCATCCCAAATGCTAAGAACTGGCAACAATTTTGAAGTCAGTTACCAATTTGAAAAAGTTCCTTTCCATTCAATG
TACGCGCACAGCCAGAGCCTGGACAGAATGATGAATCCTTTACTGGATCAGTACCTGTGGCATCTGCAATCGACC
ACTACCGGAAATTCCCTTAATCAAGGAACAGCTACCACCACGTACGGGAAAATTACCACTGGAGACTTTGCCTAC
TACAGGAAAAACTGGTTGCCTGGAGCCTGCATTAAACAACAAAAATTTTCAAAGAATGCCAATCAAAACTACAAG
ATTCCCGCCAGCGGGGGAGACGCCCTTTTAAAGTATGACACGCATACCACTCTAAATGGGCGATGGAGTAACATG
GCTCCTGGACCTCCAATGGCAACCGCAGGTGCCGGGGACTCGGATTTTAGCAACAGCCAGCTGATCTTTGCCGGA
CCCAATCCGAGCGGTAACACGACCACATCTTCAAACAATTTGTTGTTTACCTCAGAAGAGGAGATTGCCACAACA
AACCCACGAGACACGGACATGTTTGGACAGATTGCAGATAATAATCAAAATGCCACCACCGCCCCTCACATCGCT
AACCTGGACGCTATGGGAATTGTTCCCGGAATGGTCTGGCAAAACAGAGACATCTACTACCAGGGCCCTATTTGG
GCCAAGGTCCCTCACACGGACGGACACTTTCACCCTTCGCCGCTGATGGGAGGATTTGGACTGAAACACCCGCCT
CCACAGATTTTCATCAAAAACACCCCCGTACCCGCCAATCCCAATACTACCTTTAGCGCTGCAAGGATTAATTCT
TTTCTGACGCAGTACAGCACCGGACAAGTTGCCGTTCAGATCGACTGGGAAATTCAGAAGGAGCATTCCAAACGC

TGGAATCCCGAAGTTCAATTTACTTCAAACTACGGCACTCAAAATTCTATGCTGTGGGCTCCCGACAATGCTGGC
AACTACCACGAACTCCGGGCTATTGGGTCCCGTTTCCTCACCCACCACTTGTAA

AAV-13
Partial Genome: EU285562

Rep78: (SEQ ID NO:28)

ATGCCGGGATTCTACGAGATTGTCCTGAAGGTGCCCAGCGACCTGGACGAGCACCTGCCTGGCATTTCTGACTCT
TTTGTAAACTGGGTGGCGGAGAAGGAATGGGAGCTGCCGCCGGATTCTGACATGGATCTGAATCTGATTGAGCAG
GCACCCCTAACCGTGGCCGAAAAGCTGCAACGCGAATTCCTGGTCGAGTGGCGCCGCGTGAGTAAGGCCCCGGAG
GCCCTCTTCTTTGTTCAGTTCGAGAAGGGGGACAGCTACTTCCACCTACACATTCTGGTGGAGACCGTGGGCGTG
AAATCCATGGTGGTGGGCCGCTACGTGAGCCAGATTAAAGAGAAGCTGGTGACCCGCATCTACCGCGGGGTCGAG
CCGCAGCTTCCGAACTGGTTCGCGGTGACCAAGACGCGTAATGGCGCCGGAGGCGGGAACAAGGTGGTGGACGAC
TGCTACATCCCCAACTACCTGCTCCCCAAGACCCAGCCCGAGCTCCAGTGGGCGTGGACTAATATGGACCAGTAT
TTAAGCGCCTGTTTGAATCTCGCGGAGCGTAAACGGCTGGTGGCGCAGCATCTGACGCACGTGTCGCAGACGCAG
GAGCAGAACAAAGAGAACCAGAATCCCAATTCTGACGCGCCGGTGATCAGATCAAAAACCTCCGCGAGGTACATG
GAGCTGGTCGGGTGGCTGGTGGACCGCGGGATCACGTCAGAAAAGCAATGGATCCAGGAGGACCAGGCCTCTTAC
ATCTCCTTCAACGCCGCCTCCAACTCGCGGTCACAAATCAAGGCCGCACTGGACAATGCCTCCAAATTTATGAGC
CTGACAAAAACGGCTCCGGACTACCTGGTGGGAAACAACCCGCCGGAGGACATTACCAGCAACCGGATCTACAAA
ATCCTCGAGATGAACGGGTACGATCCGCAGTACGCGGCCTCCGTCTTCCTGGGCTGGGCGCAAAAGAAGTTCGGG
AAGAGGAACACCATCTGGCTCTTTGGGCCGGCCACGACGGGTAAAACCAACATCGCTGAAGCTATCGCCCACGCC
GTGCCCTTTTACGGCTGCGTGAACTGGACCAATGAGAACTTTCCGTTCAACGATTGCGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGCAAGATGACGGCCAAGGTCGTGGAGTCCGCCAAGGCCATTCTGGGCGGAAGCAAGGTGCGC
GTGGACCAAAAGTGCAAGTCATCGGCCCAGATCGACCCAACTCCCGTCATCGTCACCTCCAACACCAACATGTGC
GCGGTCATCGACGGAAATTCCACCACCTTCGAGCACCAACAACCACTCCAAGACCGGATGTTCAAGTTCGAGCTC
ACCAAGCGCCTGGAGCACGACTTTGGCAAGGTCACCAAGCAGGAAGTCAAGGACTTTTTCCGGTGGGCGTCAGAT
CACGTGACTGAGGTGTCTCACGAGTTTTACGTCAGAAAGGGTGGAGCTAGAAAGAGGCCCGCCCCCAATGACGCA
GATATAAGTGAGCCCAAGCGGGCCTGTCCGTCAGTTGCGCAGCCATCGACGTCAGACGCGGAAGCTCCGGTGGAC
TACGCGGACAGGTACCAAAACAAATGTTCTCGTCACGTGGGCATGAATCTGATGCTTTTTCCCTGCCGGCAATGC
GAGAGAATGAATCAGAATGTGGACATTTGCTTCACGCACGGGGTCATGGACTGTGCCGAGTGCTTCCCCGTGTCA
GAATCTCAACCCGTGTCTGTCGTCAGAAAGCGGACATATCAGAAACTGTGTCCGATTCATCACATCATGGGGAGG
GCGCCCGAGGTGGCTTGTTCGGCCTGCGATCTGGCCAATGTGGACTTGGATGACTGTGACATGGAGCAATAA

CapVP1: (SEQ ID NO:29)

ATGACTGACGGTTACCTTCCAGATTGGCTAGAGGACAACCTCTCTGAAGGCGTTCGAGAGTGGTGGGCGCTGCAA
CCTGGAGCCCCTAAACCCAAGGCAAATCAACAACATCAGGACAACGCTCGGGGTCTTGTGCTTCCGGGTTACAAA
TACCTCGGACCCGGCAACGGACTTGACAAGGGGGAACCCGTCAACGCAGCGGACGCGGCAGCCCTCGAACACGAC
AAGGCCTACGACCAGCAGCTCAAGGCCGGTGACAACCCCTACCTCAAGTACAACCACGCCGACGCCGAGTTTCAG
GAGCGTCTTCAAGAAGATACGTCTTTTGGGGGCAACCTCGGACGAGCAGTCTTCCAGGCCAAAAAGAGGATCCTT
GAGCCTCTGGGTCTGGTTGAGGAAGCGGCTAAGACGGCTCCTGGAAAAAAGAGACCTGTAGAGCAATCTCCAGCA
GAACCGGACTCCTCTTCGGGCATCGGCAAATCAGGCCAGCAGCCCGCTAGAAAAAGACTGAATTTTGGTCAGACT
GGCGACACAGAGTCAGTCCCAGACCCTCAACCACTCGGACAACCTCCCGCAGCCCCCTCTGGTGTGGGATCTACT
ACAATGGCTTCAGGCGGTGGCGCACCAATGGCAGACAATAACGAGGGTGCCGATGGAGTGGGTAATTCCTCAGGA
AATTGGCATTGCGATTCCCAATGGCTGGGCGACAGAGTCATCACCACCAGCACCCGCACCTGGGCCCTGCCCACC
TACAACAATCACCTCTACAAGCAAATCTCCAGCCAATCAGGAGCCACCAACGACAACCACTACTTTGGCTACAGC
ACCCCCTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAAAGACTCATCAAC
AACAACTGGGGATTCCGACCCAAGAGACTCAACTTCAAGCTCTTTAACATTCAAGTCAAAGAGGTCACGCAGAAT
GACGGTACGACGACGATTGCCAATAACCTTACCAGCACGGTTCAGGTGTTTACTGACTCCGAGTACCAGCTCCCG
TACGTCCTCGGCTCGGCGCATCAGGGATGCCTCCCGCCGTTCCCAGCAGACGTCTTCATGGTCCCACAGTATGGA
TACCTCACCCTGAACAACGGGAGTCAGGCGGTAGGACGCTCTTCCTTTTACTGCCTGGAGTACTTTCCTTCTCAG
ATGCTGCGTACTGGAAACAACTTTCAGTTTAGCTACACTTTTGAAGACGTGCCTTTCCACAGCAGCTACGCTCAC
AGCCAAAGTCTGGACCGTCTCATGAATCCTCTGATCGACCAGTACCTGTACTATCTGAACAGGACACAAACAGCC
AGTGGAACTCAGCAGTCTCGGCTACTGTTTAGCCAAGCTGGACCCACCAGTATGTCTCTTCAAGCTAAAAACTGG
CTGCCTGGACCTTGCTACAGACAGCAGCGTCTGTCAAAGCAGGCAAACGACAACAACAACAGCAACTTTCCCTGG
ACTGGTGCCACCAAATATCATCTGAATGGCCGGGACTCATTGGTGAACCCGGGCCCTGCTATGGCCAGTCACAAG
GATGACAAAGAAAAGTTTTTCCCCATGCATGGAACCCTGATATTTGGTAAAGAAGGAACAAATGCCAACAACGCG

GATTTGGAAAATGTCATGATTACAGATGAAGAAGAAATCCGCACCACCAATCCCGTGGCTACGGAGCAGTACGGG
ACTGTGTCAAATAATTTGCAAAACTCAAACGCTGGTCCAACTACTGGAACTGTCAATCACCAAGGAGCGTTACCT
GGTATGGTGTGGCAGGATCGAGACGTGTACCTGCAGGGACCCATTTGGGCCAAGATTCCTCACACCGATGGACAC
TTTCATCCTTCTCCACTGATGGGAGGTTTTGGGCTCAAACACCCGCCTCCTCAGATCATGATCAAAAACACTCCC
GTTCCAGCCAATCCTCCCACAAACTTTAGTGCGGCAAAGTTTGCTTCCTTCATCACACAGTACTCCACGGGGCAG
GTCAGCGTGGAGATCGAGTGGGAGCTGCAGAAGGAGAACAGCAAACGCTGGAATCCCGAAATTCAGTACACTTCC
AACTACAACAAATCTGTTAATGTGGACTTTACTGTGGACACTAATGGTGTGTATTCAGAGCCTCGCCCCATTGGC
ACCAGATACCTGACTCGTAATCTGTAA

ITR Sequence(SEQ ID NO:30)

CCTGCAGGCAGCTGCGCGCTCGCTCGCTCACTGAGGCCGCCCGGGCAAAGCCCGGGCGTCGGGCGACCTTTGGTC
GCCCGGCCTCAGTGAGCGAGCGAGCGCGCAGAGAGGGAGTGGCCAACTCCATCACTAGGGGTTCCT

Rep2 Sequence – Contains Rep78 and Rep52 (start codon underlined) (SEQ ID NO:31)

ATGCCGGGGTTTTACGAGATTGTGATTAAGGTCCCCAGCGACCTTGACGAGCATCTGCCCGGCATTTCTGACAGC
TTTGTGAACTGGGTGGCCGAGAAGGAATGGGAGTTGCCGCCAGATTCTGACATGGATCTGAATCTGATTGAGCAG
GCACCCCTGACCGTGGCCGAGAAGCTGCAGCGCGACTTTCTGACGGAATGGCGCCGTGTGAGTAAGGCCCCGGAG
GCCCTTTTCTTTGTGCAATTTGAGAAGGGAGAGAGCTACTTCCACATGCACGTGCTCGTGGAAACCACCGGGGTG
AAATCCATGGTTTTGGGACGTTTCCTGAGTCAGATTCGCGAAAAACTGATTCAGAGAATTTACCGCGGGATCGAG
CCGACTTTGCCAAACTGGTTCGCGGTCACAAAGACCAGAAATGGCGCCGGAGGCGGGAACAAGGTGGTGGATGAG
TGCTACATCCCCAATTACTTGCTCCCCAAAACCCAGCCTGAGCTCCAGTGGGCGTGGACTAATATGGAACAGTAT
TTAAGCGCCTGTTTGAATCTCACGGAGCGTAAACGGTTGGTGGCGCAGCATCTGACGCACGTGTCGCAGACGCAG
GAGCAGAACAAAGAGAATCAGAATCCCAATTCTGATGCGCCGGTGATCAGATCAAAAACTTCAGCCAGGTACATG
GAGCTGGTCGGGTGGCTCGTGGACAAGGGGATTACCTCGGAGAAGCAGTGGATCCAGGAGGACCAGGCCTCATAC
ATCTCCTTCAATGCGGCCTCCAACTCGCGGTCCCAAATCAAGGCTGCCTTGGACAATGCGGGAAAGATTATGAGC
CTGACTAAAACCGCCCCCGACTACCTGGTGGGCCAGCAGCCCGTGGAGGACATTTCCAGCAATCGGATTTATAAA
ATTTTGGAACTAAACGGGTACGATCCCCAATATGCGGCTTCCGTCTTTCTGGGATGGGCCACGAAAAAGTTCGGC
AAGAGGAACACCATCTGGCTGTTTGGGCCTGCAACTACCGGGAAGACCAACATCGCGGAGGCCATAGCCCACACT
GTGCCCTTCTACGGGTGCGTAAACTGGACCAATGAGAACTTTCCCTTCAACGACTGTGTCGACAAGATGGTGATC
TGGTGGGAGGAGGGGAAGATGACCGCCAAGGTCGTGGAGTCGGCCAAAGCCATTCTCGGAGGAAGCAAGGTGCGC
GTGGACCAGAAATGCAAGTCCTCGGCCCAGATAGACCCGACTCCCGTGATCGTCACCTCCAACACCAACATGTGC
GCCGTGATTGACGGGAACTCAACGACCTTCGAACACCAGCAGCCGTTGCAAGACCGGATGTTCAAATTTGAACTC
ACCCGCCGTCTGGATCATGACTTTGGGAAGGTCACCAAGCAGGAAGTCAAAGACTTTTTCCGGTGGGCAAAGGAT
CACGTGGTTGAGGTGGAGCATGAATTCTACGTCAAAAAGGGTGGAGCCAAGAAAAGACCCGCCCCCAGTGACGCA
GATATAAGTGAGCCCAAACGGGTGCGCGAGTCAGTTGCGCAGCCATCGACGTCAGACGCGGAAGCTTCGATCAAC
TACGCAGACAGGTACCAAAACAAATGTTCTCGTCACGTGGGCATGAATCTGATGCTGTTTCCCTGCAGACAATGC
GAGAGAATGAATCAGAATTCAAATATCTGCTTCACTCACGGACAGAAAGACTGTTTAGAGTGCTTTCCCGTGTCA
GAATCTCAACCCGTTTCTGTCGTCAAAAAGGCGTATCAGAAACTGTGCTACATTCATCATATCATGGGAAAGGTG
CCAGACGCTTGCACTGCCTGCGATCTGGTCAATGTGGATTTGGATGACTGCATCTTTGAACAATAA

Cap2 Sequence – contains sequentially VP1, VP2, AAP, VP3 (start codons underlined) (SEQ ID NO:32)

ATGGCTGCCGATGGTTATCTTCCAGATTGGCTCGAGGACACTCTCTCTGAAGGAATAAGACAGTGGTGGAAGCTC
AAACCTGGCCCACCACCACCAAAGCCCGCAGAGCGGCATAAGGACGACAGCAGGGGTCTTGTGCTTCCTGGGTAC
AAGTACCTCGGACCCTTCAACGGACTCGACAAGGGAGAGCCGGTCAACGAGGCAGACGCCGCGGCCCTCGAGCAC
GACAAAGCCTACGACCGGCAGCTCGACAGCGGAGACAACCCGTACCTCAAGTACAACCACGCCGACGCGGAGTTT
CAGGAGCGCCTTAAAGAAGATACGTCTTTTGGGGGCAACCTCGGACGAGCAGTCTTCCAGGCGAAAAAGAGGGTT
CTTGAACCTCTGGGCCTGGTTGAGGAACCTGTTAAGACGGCTCCGGGAAAAAAGAGGCCGGTAGAGCACTCTCCT
GTGGAGCCAGACTCCTCCTCGGGAACCGGAAAGGCGGGCCAGCAGCCTGCAAGAAAAAGATTGAATTTTGGTCAG
ACTGGAGACGCAGACTCAGTACCTGACCCCCAGCCTCTCGGACAGCCACCAGCAGCCCCCTCTGGTCTGGGAACT
AATACGATGGCTACAGGCAGTGGCGCACCAATGGCAGACAATAACGAGGGCGCCGACGGAGTGGGTAATTCCTCG
GGAAATTGGCATTGCGATTCCACATGGATGGGCGACAGAGTCATCACCACCAGCACCCGAACCTGGGCCCTGCCC
ACCTACAACAACCACCTCTACAAACAAATTTCCAGCCAATCAGGAGCCTCGAACGACAATCACTACTTTGGCTAC

AGCACCCCTTGGGGGTATTTTGACTTCAACAGATTCCACTGCCACTTTTCACCACGTGACTGGCAAAGACTCATC
AACAACAACTGGGGATTCCGACCCAAGAGACTCAACTTCAAGCTCTTTAACATTCAAGTCAAAGAGGTCACGCAG
AATGACGGTACGACGACGATTGCCAATAACCTTACCAGCACGGTTCAGGTGTTTACTGACTCGGAGTACCAGCTC
CCGTACGTCCTCGGCTCGGCGCATCAAGGATGCCTCCCGCCGTTCCCAGCAGACGTCTTCATGGTGCCACAGTAT
GGATACCTCACCCTGAACAACGGGAGTCAGGCAGTAGGACGCTCTTCATTTTACTGCCTGGAGTACTTTCCTTCT
CAGATGCTGCGTACCGGAAACAACTTTACCTTCAGCTACACTTTTGAGGACGTTCCTTTCCACAGCAGCTACGCT
CACAGCCAGAGTCTGGACCGTCTCATGAATCCTCTCATCGACCAGTACCTGTATTACTTGAGCAGAACAAACACT
CCAAGTGGAACCACCACGCAGTCAAGGCTTCAGTTTTCTCAGGCCGGAGCGAGTGACATTCGGGACCAGTCTAGG
AACTGGCTTCCTGGACCCTGTTACCGCCAGCAGCGAGTATCAAAGACATCTGCGGATAACAACAACAGTGAATAC
TCGTGGACTGGAGCTACCAAGTACCACCTCAATGGCAGAGACTCTCTGGTGAATCCGGGCCCGGCCATGGCAAGC
CACAAGGACGATGAAGAAAAGTTTTTTCCTCAGAGCGGGGTTCTCATCTTTGGGAAGCAAGGCTCAGAGAAAACA
AATGTGGACATTGAAAAGGTCATGATTACAGACGAAGAGGAAATCAGGACAACCAATCCCGTGGCTACGGAGCAG
TATGGTTCTGTATCTACCAACCTCCAGAGAGGCAACAGACAAGCAGCTACCGCAGATGTCAACACACAAGGCGTT
CTTCCAGGCATGGTCTGGCAGGACAGAGATGTGTACCTTCAGGGGCCCATCTGGGCAAAGATTCCACACACGGAC
GGACATTTTCACCCCTCTCCCCTCATGGGTGGATTCGGACTTAAACACCCTCCTCCACAGATTCTCATCAAGAAC
ACCCCGGTACCTGCGAATCCTTCGACCACCTTCAGTGCGGCAAAGTTTGCTTCCTTCATCACACAGTACTCCACG
GGACAGGTCAGCGTGGAGATCGAGTGGGAGCTGCAGAAGGAAAACAGCAAACGCTGGAATCCCGAAATTCAGTAC
ACTTCCAACTACAACAAGTCTGTTAATGTGGACTTTACTGTGGACACTAATGGCGTGTATTCAGAGCCTCGCCCC
ATTGGCACCAGATACCTGACTCGTAATCTGTAA

Cap5 Sequence – contains sequentially VP1, VP2, AAP, VP3 (start codons underlined) (SEQ ID NO:33)

ATGGCTTTTGTTGATCACCCTCCAGATTGGTTGGAAGAAGTTGGTGAAGGTCTTCGCGAGTTTTTGGGCCTTGAA
GCGGGCCCACCGAAACCAAAACCCAATCAGCAGCATCAAGATCAAGCCCGTGGTCTTGTGCTGCCTGGTTATAAC
TATCTCGGACCCGGAAACGGTCTCGATCGAGGAGAGCCTGTCAACAGGGCAGACGAGGTCGCGCGAGAGCACGAC
ATCTCGTACAACGAGCAGCTTGAGGCGGGAGACAACCCCTACCTCAAGTACAACCACGCGGACGCCGAGTTTCAG
GAGAAGCTCGCCGACGACACATCCTTCGGGGGAAACCTCGGAAAGGCAGTCTTTCAGGCCAAGAAAAGGGTTCTC
GAACCTTTTGGCCTGGTTGAAGAGGGTGCTAAGACGGCCCCTACCGGAAAGCGGATAGACGACCACTTTCCAAAA
AGAAAGAAGGCTCGGACCGAAGAGGACTCCAAGCCTTCCACCTCGTCAGACGCCGAAGCTGGACCCAGCGGATCC
CAGCAGCTGCAAATCCCAGCCCAACCAGCCTCAAGTTTGGGAGCTGATACAATGTCTGCGGGAGGTGGCGGCCCA
TTGGGCGACAATAACCAAGGTGCCGATGGAGTGGGCAATGCCTCGGGAGATTGGCATTGCGATTCCACGTGGATG
GGGGACAGAGTCGTCACCAAGTCCACCCGAACCTGGGTGCTGCCCAGCTACAACAACCACCAGTACCGAGAGATC
AAAAGCGGCTCCGTCGACGGAAGCAACGCCAACGCCTACTTTGGATACAGCACCCCCTGGGGGTACTTTGACTTT
AACCGCTTCCACAGCCACTGGAGCCCCCGAGACTGGCAAAGACTCATCAACAACTACTGGGGCTTCAGACCCCGG
TCCCTCAGAGTCAAAATCTTCAACATTCAAGTCAAAGAGGTCACGGTGCAGGACTCCACCACCACCATCGCCAAC
AACCTCACCTCCACCGTCCAAGTGTTTACGGACGACGACTACCAGCTGCCCTACGTCGTCGGCAACGGGACCGAG
GGATGCCTGCCGGCCTTCCCTCCGCAGGTCTTTACGCTGCCGCAGTACGGTTACGCGACGCTGAACCGCGACAAC
ACAGAAAATCCCACCGAGAGGAGCAGCTTCTTCTGCCTAGAGTACTTTCCCAGCAAGATGCTGAGAACGGGCAAC
AACTTTGAGTTTACCTACAACTTTGAGGAGGTGCCCTTCCACTCCAGCTTCGCTCCCAGTCAGAACCTCTTCAAG
CTGGCCAACCCGCTGGTGGACCAGTACTTGTACCGCTTCGTGAGCACAAATAACACTGGCGGAGTCCAGTTCAAC
AAGAACCTGGCCGGGAGATACGCCAACACCTACAAAAACTGGTTCCCGGGGCCCATGGGCCGAACCCAGGGCTGG
AACCTGGGCTCCGGGGTCAACCGCGCCAGTGTCAGCGCCTTCGCCACGACCAATAGGATGGAGCTCGAGGGCGCG
AGTTACCAGGTGCCCCCGCAGCCGAACGGCATGACCAACAACCTCCAGGGCAGCAACACCTATGCCCTGGAGAAC
ACTATGATCTTCAACAGCCAGCCGGCGAACCCGGGCACCACCGCCACGTACCTCGAGGGCAACATGCTCATCACC
AGCGAGAGCGAGACGCAGCCGGTGAACCGCGTGGCGTACAACGTCGGCGGGCAGATGGCCACCAACAACCAGAGC
TCCACCACTGCCCCCGCGACCGGCACGTACAACCTCCAGGAAATCGTGCCCGGCAGCGTGTGGATGGAGAGGGAC
GTGTACCTCCAAGGACCCATCTGGGCCAAGATCCCAGAGACGGGGGCGCACTTTCACCCCTCTCCGGCCATGGGC
GGATTCGGACTCAAACACCCACCGCCCATGATGCTCATCAAGAACACGCCTGTGCCCGGAAATATCACCAGCTTC
TCGGACGTGCCCGTCAGCAGCTTCATCACCCAGTACAGCACCGGGCAGGTCACCGTGGAGATGGAGTGGGAGCTC
AAGAAGGAAAACTCCAAGAGGTGGAACCCAGAGATCCAGTACACAAACAACTACAACGACCCCCAGTTTGTGGAC
TTTGCCCCGGACAGCACCGGGGAATACAGAAGCACCAGACCTATCGGAACCCGATACCTTACCCGACCCCTTTAA

EXAMPLE 12 – Adenovirus Polynucleotide Sequences

Adenovirus (Ad) polynucleotides can be selected from any serotype, and representative polynucleotides are exemplified below.

E2A Full Sequence (SEQ ID NO:34)

CGACCGCACCCTGTGACGAAAGCCGCCCGCAAGCTGCGCCCCTGAGTTAGTCATCTGAACTTCGGCCTGGGCGT
CTCTGGGAAGTACCACAGTGGTGGGAGCGGGACTTTCCTGGTACACCAGGGCAGCGGGCCAACTACGGGGATTAA
GGTTATTACGAGGTGTGGTGGTAATAGCCGCCTGTTCGAGGAGAATTCGGTTTCGGTGGGCGCGGATTCCGTTGA
CCCGGGATATCATGTGGGGTCCCGCGCTCATGTAGTTTATTCGGGTTGAGTAGTCTTGGGCAGCTCCAGCCGCAA
GTCCCATTTGTGGCTGGTAACTCCACATGTAGGGCGTGGGAATTTCCTTGCTCATAATGGCGCTGACGACAGGTG
CTGGCGCCGGGTGTGGCCGCTGGAGATGACGTAGTTTTCGCGCTTAAATTTGAGAAAGGGCGCGAAACTAGTCCT
TAAGAGTCAGCGCGCAGTATTTGCTGAAGAGAGCCTCCGCGTCTTCCAGCGTGCGCCGAAGCTGATCTTCGCTTT
TGTGATACAGGCAGCTGCGGGTGAGGGAGCGCAGAGACCTGTTTTTTATTTTCAGCTCTTGTTCTTGGCCCCTGC
TTTGTTGAAATATAGCATACAGAGTGGGAAAAATCCTATTTCTAAGCTCGCGGGTCGATACGGGTTCGTTGGGCG
CCAGACGCAGCGCTCCTCCTCCTGCTGCTGCCGCCGCTGTGGATTTCTTGGGCTTTGTCAGAGTCTTGCTATCCG
GTCGCCTTTGCTTCTGTGTGACCGCTGCTGTTGCTGCCGCTGCCGCTGCCGCCGGTGCAGTAGGGGCTGTAGAGA
TGACGGTAGTAATGCAGGATGTTACGGGGGAAGGCCACGCCGTGATGGTAGAGAAGAAAGCGGCGGGCGAAGGAG
ATGTTGCCCCCACAGTCTTGCAAGCAAGCAACTATGGCGTTCTTGTGCCCGCGCCACGAGCGGTAGCCTTGGCGC
TGTTGTTGCTCTTGGGCTAACGGCGGCGGCTGCTTAGACTTACCGGCCCTGGTTCCAGTGGTGTCCCATCTACGG
TTGGGTCGGCGAACAGGCAGTGCCGGCGGCGCCTGAGGAGCGGAGGTTGTAGCGATGCTGGGAACGGTTGCCAAT
TTCTGGGGCGCCGGCGAGGGGAATGCGACCGAGGGTGACGGTGTTTCGTCTGACACCTCTTCGGCCTCGGAAGCT
TCGTCTAGGCTGTCCCAGTCTTCCATCATCTCCTCCTCCTCGTCCAAAACCTCCTCTGCCTGACTGTCCCAGTAT
TCCTCCTCGTCCGTGGGTGGCGGCGGCGGCAGCTGCAGCTTCTTTTTGGGTGCCATCCTGGGAAGCAAGGGCCCG
CGGCTGCTGATAGGGCTGCGGCGGCGGGGGGATTGGGTTGAGCTCCTCGCCGGACTGGGGGTCCAGGTAAACCCC
CCGTCCCTTTCGTAGCAGAAACTCTTGGCGGGCTTTGTTGATGGCTTGCAATTGGCCAAGGATGTGGCCCTGGGT
AATGACGCAGGCGGTAAGCTCCGCATTTGGCGGGCGGGATTGGTCTTCGTAGAACCTAATCTCGTGGGCGTGGTA
GTCCTCAGGTACAAATTTGCGAAGGTAAGCCGACGTCCACAGCCCCGGAGTGAGTTTCAACCCCGGAGCCGCGGA
CTTTTCGTCAGGCGAGGGACCCTGCAGCTCAAAGGTACCGATAATTTGACTTTCGCTAAGCAGTTGCGAATTGCA
GACCAGGGAGCGGTGCGGGGTGCATAGGTTGCAGCGACAGTGACACTCCAGTAGGCCGTCACCGCTCACGTCTTC
CATGATGTCGGAGTGGTAGGCAAGGTAGTTGGCTAGCTGCAGAAGGTAGCAGTGACCCCAAAGCGGCGGAGGGCA
TTCACGGTACTTAATGGGCACAAAGTCGCTAGGAAGCGCACAGCAGGTGGCGGGCAGAATTCCTGAACGCTCTAG
GATAAAGTTCCTAAAGTTTTGCAACATGCTTTGACTGGTGAAGTCTGGCAGACCCTGTTGCAGGGTTTTAAGCAG
GCGTTCGGGGAAGATAATGTCCGCCAGGTGCGCGGCCACGGAGCGCTCGTTGAAGGCCGTCCATAGGTCCTTCAA
GTTTTGCTTTAGCAGCTTCTGCAGCTCCTTTAGGTTGCGCTCCTCCAGGCATTGCTGCCACACGCCCATGGCCGT
TTGCCAGGTGTAGCACAGAAATAAGTAAACGCAGTCGCGGACGTAGTCGCGGCGCGCCTCGCCCTTGAGCGTGGA
ATGAAGCACGTTTTGCCCGAGGCGGTTTTCGTGCAAAATTCCAAGGTAGGAGACCAGGTTGCAGAGCTCCACGTT
GGAAATTTTGCAGGCCTGGCGCACGTAGCCCTGGCGAAAGGTGTAGTGCAACGTTTCCTCTAGCTTGCGCTGCAT
CTCCGGGTCAGCAAAGAACCGCTGCATGCACTCAAGCTCCACGGTAACAAGCACTGCGGCCATCATTAGCTTGCG
TCGCTCCTCCAAGTCGGCAGGCTCGCGCGTCTCAAGCCAGCGCGCCAGCTGCTCATCGCCAACTGCGGGTAGGCC
CTCCTCGGTTTGTTCTTGCAAGTTTGCATCCCTCTCCAGGGGTCGTGCACGGCGCACGATCAGCTCGCTCATGAC
TGTGCTCATAACCTTGGGGGGTAGGTTAAGTGCCGGGTAGGCAAAGTGGGTGACCTCGATGCTGCGTTTCAGCAC
GGCTAGGCGCGCGTTGTCACCCTCAAGTTCCACCAGCACTCCACAGTGACTTTCATTTTCGCTGTTTTCTTGTTG
CAGAGCGTTTGCCGCGCGTTTCTCGTCGCGTCCAAGACCCTCAAAGATTTTTGGCACTTCGTCGAGCGAGGCGAT
ATCAGGTATGACAGCGCCCTGCCGCAAGGCCAGCTGCTTGTCCGCTCGGCTGCGGTTGGCACGGCAGGATAGGGG
TATCTTGCAGTTTTGGAAAAAGATGTGATAGGTGGCAAGCACCTCTGGCACGGCAAATACGGGGTAGAAGTTGAG
GCGCGGGTTGGGCTCGCATGTGCCGTTTTCTTGGCGTTTGGGGGGTACGCGCGGTGAGAACAGGTGGCGTTCGTA
GGCAAGGCTGACATCCGCTATGGCGAGGGGCACATCGCTGCGCTCTTGCAACGCGTCGCAGATAATGGCGCACTG
GCGCTGCAGATGCTTCAACAGCACGTCGTCTCCCACATCTAGGTAGTCGCCATGCCTTTGGTCCCCCCGCCCGAC
TTGTTCCTCGTTTGCCTCTGCGTCGTCCTGGTCTTGCTTTTTATCCTCTGTTGGTACTGAGCGATCCTCGTCGTC
TTCGCTTACAAAACCTGGGTCCTGCTCGATAATCACTTCCTCCTCCTCAAGCGGGGGTGCCTCGACGGGGAAGGT
GGTAGGCGCGTTGGCGGCATCGGTGGAGGCGGTGGTGGCGAACTCAAAGGGGGCGGTTAGGCTGTCCTCCTTCTC
GACTGACTCCATGATCTTTTTCTGCCTATAGGAGAAGGAAATGGCCAGTCGGGAAGAGGAGCAGCGCGAAACCAC
CCCCGAGCGCGGACGCGGTGCGGCGCGACGTCCACCAACCATGGAGGACGTGTCGTCCCCGTCGCCGTCGCCGCC
GCCTCCCCGCGCGCCCCCAAAAAAGCGGCTGAGGCGGCGTCTCGAGTCCGAGGACGAAGAAGACTCGTCACAAGA
TGCGCTGGTGCCGCGCACACCCAGCCCGCGGCCATCGACCTCGACGGCGGATTTGGCCATTGCGTCCAAAAAGAA
AAAGAAGCGCCCCTCTCCCAAGCCCGAGCGCCCGCCATCCCCAGAGGTGATCGTGGACAGCGAGGAAGAAAGAGA
AGATGTGGCGCTACAAATGGTGGGTTTCAGCAACCCACCGGTGCTAATCAAGCACGGCAAGGGAGGTAAGCGCAC
GGTGCGGCGGCTGAATGAAGACGACCCAGTGGCGCGGGGTATGCGGACGCAAGAGGAAAAGGAAGAGTCCAGTGA
AGCGGAAAGTGAAAGCACGGTGATAAACCCGCTGAGCCTGCCGATCGTGTCTGCGTGGGAGAAGGGCATGGAGGC
TGCGCGCGCGTTGATGGACAAGTACCACGTGGATAACGATCTAAAGGCAAACTTCAAGCTACTGCCTGACCAAGT
GGAAGCTCTGGCGGCCGTATGCAAGACCTGGCTAAACGAGGAGCACCGCGGGTTGCAGCTGACCTTCACCAGCAA

CAAGACCTTTGTGACGATGATGGGGCGATTCCTGCAGGCGTACCTGCAGTCGTTTGCAGAGGTAACCTACAAGCA
CCACGAGCCCACGGGCTGCGCGTTGTGGCTGCACCGCTGCGCTGAGATCGAAGGCGAGCTTAAGTGTCTACACGG
GAGCATTATGATAAATAAGGAGCACGTGATTGAAATGGATGTGACGAGCGAAAACGGGCAGCGCGCGCTGAAGGA
GCAGTCTAGCAAGGCCAAGATCGTGAAGAACCGGTGGGGCCGAAATGTGGTGCAGATCTCCAACACCGACGCAAG
GTGCTGCGTGCATGACGCGGCCTGTCCGGCCAATCAGTTTTCCGGCAAGTCTTGCGGCATGTTCTTCTCTGAAGG
CGCAAAGGCTCAGGTGGCTTTTAAGCAGATCAAGGCTTTCATGCAGGCGCTGTATCCTAACGCCCAGACCGGGCA
CGGTCACCTTCTGATGCCACTACGGTGCGAGTGCAACTCAAAGCCTGGGCATGCACCCTTTTTGGGAAGGCAGCT
ACCAAAGTTGACTCCGTTCGCCCTGAGCAACGCGGAGGACCTGGACGCGGATCTGATCTCCGACAAGAGCGTGCT
GGCCAGCGTGCACCACCCGGCGCTGATAGTGTTCCAGTGCTGCAACCCTGTGTATCGCAACTCGCGCGCGCAGGG
CGGAGGCCCCAACTGCGACTTCAAGATATCGGCGCCCGACCTGCTAAACGCGTTGGTGATGGTGCGCAGCCTGTG
GAGTGAAAACTTCACCGAGCTGCCGCGGATGGTTGTGCCTGAGTTTAAGTGGAGCACTAAACACCAGTATCGCAA
CGTGTCCCTGCCAGTGGCGCATAGCGATGCGCGGCAGAACCCCTTTGATTTTTAAACGGCGCAGACGGCAAGGGT
GGGGGGTAAATAATCACCCGAGAGTGTACAAATAAAAACATTTGCCTTTATTGAAAGTGTCTCCTAGTACATTAT
TTTTACATGTTTTTCAAGTGACAAAAAGAAGTGGCGCTCCTAATCTGCGCACTGTGGCTGCGGAAGTAGGGCGAG
TGGCGCTCCAGGAAGCTGTAGAGCTGTTCCTGGTTGCGACGCAGGGTGGGCTGTACCTGGGGACTGTTAAGCATG
GAGTTGGGTACC

E2A ORF Sequence (SEQ ID NO:35)

ATGGCCAGTCGGGAAGAGGAGCAGCGCGAAACCACCCCCGAGCGCGGACGCGGTGCGGCGCGACGTCCACCAACC
ATGGAGGACGTGTCGTCCCCGTCGCCGTCGCCGCCGCCTCCCCGCGCGCCCCCAAAAAAGCGGCTGAGGCGGCGT
CTCGAGTCCGAGGACGAAGAAGACTCGTCACAAGATGCGCTGGTGCCGCGCACACCCAGCCCGCGGCCATCGACC
TCGACGGCGGATTTGGCCATTGCGTCCAAAAAGAAAAAGAAGCGCCCCTCTCCCAAGCCCGAGCGCCCGCCATCC
CCAGAGGTGATCGTGGACAGCGAGGAAGAAAGAGAAGATGTGGCGCTACAAATGGTGGGTTTCAGCAACCCACCG
GTGCTAATCAAGCACGGCAAGGGAGGTAAGCGCACGGTGCGGCGGCTGAATGAAGACGACCCAGTGGCGCGGGGT
ATGCGGACGCAAGAGGAAAAGGAAGAGTCCAGTGAAGCGGAAAGTGAAAGCACGGTGATAAACCCGCTGAGCCTG
CCGATCGTGTCTGCGTGGGAGAAGGGCATGGAGGCTGCGCGCGCGTTGATGGACAAGTACCACGTGGATAACGAT
CTAAAGGCAAACTTCAAGCTACTGCCTGACCAAGTGGAAGCTCTGGCGGCCGTATGCAAGACCTGGCTAAACGAG
GAGCACCGCGGGTTGCAGCTGACCTTCACCAGCAACAAGACCTTTGTGACGATGATGGGGCGATTCCTGCAGGCG
TACCTGCAGTCGTTTGCAGAGGTAACCTACAAGCACCACGAGCCCACGGGCTGCGCGTTGTGGCTGCACCGCTGC
GCTGAGATCGAAGGCGAGCTTAAGTGTCTACACGGGAGCATTATGATAAATAAGGAGCACGTGATTGAAATGGAT
GTGACGAGCGAAAACGGGCAGCGCGCGCTGAAGGAGCAGTCTAGCAAGGCCAAGATCGTGAAGAACCGGTGGGGC
CGAAATGTGGTGCAGATCTCCAACACCGACGCAAGGTGCTGCGTGCATGACGCGGCCTGTCCGGCCAATCAGTTT
TCCGGCAAGTCTTGCGGCATGTTCTTCTCTGAAGGCGCAAAGGCTCAGGTGGCTTTTAAGCAGATCAAGGCTTTC
ATGCAGGCGCTGTATCCTAACGCCCAGACCGGGCACGGTCACCTTCTGATGCCACTACGGTGCGAGTGCAACTCA
AAGCCTGGGCATGCACCCTTTTTGGGAAGGCAGCTACCAAAGTTGACTCCGTTCGCCCTGAGCAACGCGGAGGAC
CTGGACGCGGATCTGATCTCCGACAAGAGCGTGCTGGCCAGCGTGCACCACCCGGCGCTGATAGTGTTCCAGTGC
TGCAACCCTGTGTATCGCAACTCGCGCGCGCAGGGCGGAGGCCCCAACTGCGACTTCAAGATATCGGCGCCCGAC
CTGCTAAACGCGTTGGTGATGGTGCGCAGCCTGTGGAGTGAAAACTTCACCGAGCTGCCGCGGATGGTTGTGCCT
GAGTTTAAGTGGAGCACTAAACACCAGTATCGCAACGTGTCCCTGCCAGTGGCGCATAGCGATGCGCGGCAGAAC
CCCTTTGATTTTTAA

E4 Full Sequence (SEQ ID NO:36)

CCCGGGCGTTTTAGGGCGGAGTAACTTGCATGTATTGGGAATTGTAGTTTTTTTAAAATGGGAAGTGACGTATCG
TGGGAAAACGGAAGTGAAGATTTGAGGAAGTTGTGGGTTTTTTGGCTTTCGTTTCTGGGCGTAGGTTCGCGTGCG
GTTTTCTGGGTGTTTTTTGTGGACTTTAACCGTTACGTCATTTTTTAGTCCTATATATACTCGCTCTGTACTTGG
CCCTTTTTACACTGTGACTGATTGAGCTGGTGCCGTGTCGAGTGGTGTTTTTTAATAGGTTTTTTTACTGGTAAG
GCTGACTGTTATGGCTGCCGCTGTGGAAGCGCTGTATGTTGTTCTGGAGCGGGAGGGTGCTATTTTGCCTAGGCA
GGAGGGTTTTTCAGGTGTTTATGTGTTTTTCTCTCCTATTAATTTTGTTATACCTCCTATGGGGGCTGTAATGTT
GTCTCTACGCCTGCGGGTATGTATTCCCCCGGGCTATTTCGGTCGCTTTTTAGCACTGACCGATGTTAACCAACC
TGATGTGTTTACCGAGTCTTACATTATGACTCCGGACATGACCGAGGAACTGTCGGTGGTGCTTTTTAATCACGG
TGACCAGTTTTTTTACGGTCACGCCGGCATGGCCGTAGTCCGTCTTATGCTTATAAGGGTTGTTTTTCCTGTTGT
AAGACAGGCTTCTAATGTTTAAATGTTTTTTTTTTTGTTATTTTATTTTGTGTTTAATGCAGGAACCCGCAGACA
TGTTTGAGAGAAAAATGGTGTCTTTTTCTGTGGTGGTTCCGGAACTTACCTGCCTTTATCTGCATGAGCATGACT
ACGATGTGCTTGCTTTTTTGCGCGAGGCTTTGCCTGATTTTTTGAGCAGCACCTTGCATTTTATATCGCCGCCCA
TGCAACAAGCTTACATAGGGGCTACGCTGGTTAGCATAGCTCCGAGTATGCGTGTCATAATCAGTGTGGGTTCTT
TTGTCATGGTTCCTGGCGGGGAAGTGGCCGCGCTGGTCCGTGCAGACCTGCACGATTATGTTCAGCTGGCCCTGC

GAAGGGACCTACGGGATCGCGGTATTTTTGTTAATGTTCCGCTTTTGAATCTTATACAGGTCTGTGAGGAACCTG
AATTTTTGCAATCATGATTCGCTGCTTGAGGCTGAAGGTGGAGGGCGCTCTGGAGCAGATTTTTACAATGGCCGG
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TGCTGGAATGTTTATAGAGGAGATTCACCCTGAAGGGTTTAGCCTTTACGTCCACTTGGACGTGAGGGCAGTTTG
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CGGAGGGGAGCGCGTTCACTTAATAGATCTTCATTTTGAGGTTTTGGATAATCTTTTGGAATAAAAAAAAAAAAA
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TGCGAGGTCTTCCCTGCAGTGTGGGATTTACGCTGATTCAGGAATGGGTTGTTCCCTGGGATATGGTTCTGACGC
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TGATGATCCATGGTTACGAGTCCTGGGCTCTCCACTGTCATTGTTCCAGTCCCGGTTCCCTGCAGTGCATAGCCG
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GCAATGTAGGAGAAGGTGTAAGAGTTGGTAGCAAAAGTTTCAGTGGTGTATTTTCCACTTTCCCAGGACCATGTA
AAAGACATAGAGTAAGTGCTTACCTCGCTAGTTTCTGTGGATTCACTAGAA

E4 Orf6 Sequence(SEQ ID NO:37)

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TGTAACACTTTGACAATGCACAACGTGAGTTACGTGCGAGGTCTTCCCTGCAGTGTGGGATTTACGCTGATTCAG
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GTCCCCGCCATGAGCTTTGGATACAGCGCCTTGCACTGTGGGATTTTGAACAATATTGTGGTGCTGTGCTGCAGT
TACTGTGCTGATTTAAGTGAGATCAGGGTGCGCTGCTGTGCCCGGAGGACAAGGCGTCTCATGCTGCGGGCGGTG
CGAATCATCGCTGAGGAGACCACTGCCATGTTGTATTCCTGCAGGACGGAGCGGCGGCGGCAGCAGTTTATTCGC
GCGCTGCTGCAGCACCACCGCCCTATCCTGATGCACGATTATGACTCTACCCCCATGTAG

VA Sequence (VA transcripts I and II are underlined) (SEQ ID NO:38)

CGTAATCCGTAGATGTACCTGGACATCCAGGTGATGCCGGCGGCGGTGGTGGAGGCGCGCGGAAAGTCGCGGAC
GCGGTTCCAGATGTTGCGCAGCGGCAAAAAGTGCTCCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTC
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GGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGATCCGGCCGTCCGCCGTGATCCATGCGGTTACCGCCCGC
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TGCGCTAGCTTTTTTGGCCACTGGCCGCGCGCGGCGTAAGCGGTTAGGCTGGAAAGCGAAAGCATTAAGTGGCTC
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GCAGCGGCAGACATGCAGGGCACCCTCCCCTTCTCCTACCGCGTCAGGAGGGGCAACATCCTACATCGA

Sequences for E1A and E1B are both contained within Accession AY339865.1

Ad5 E1A

Two proteins can be transcribed, a 32 kDa protein (first accession number) and a 27 kDa protein (second accession number). These are both splice variants from the transcript:
Accession 1: AAQ19284.1
Accession 2: AAQ19285.1
(SEQ ID NO:39)

ATGAGACATATTATCTGCCACGGAGGTGTTATTACCGAAGAAATGGCCGCCAGTCTTTTGGACCAGCTGATCGAA
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GTGACGGCCCCCGAAGATCCCAACGAGGAGGCGGTTTCGCAGATTTTTCCCGACTCTGTAATGTTGGCGGTGCAG
GAAGGGATTGACTTACTCACTTTTCCGCCGGCGCCCGGTTCTCCGGAGCCGCCTCACCTTTCCCGGCAGCCCGAG
CAGCCGGAGCAGAGAGCCTTGGGTCCGGTTTCTATGCCAAACCTTGTACCGGAGGTGATCGATCTTACCTGCCAC
GAGGCTGGCTTTCCACCCAGTGACGACGAGGATGAAGAGGGTGAGGAGTTTGTGTTAGATTATGTGGAGCACCCC
GGGCACGGTTGCAGGTCTTGTCATTATCACCGGAGGAATACGGGGGACCCAGATATTATGTGTTCGCTTTGCTAT
ATGAGGACCTGTGGCATGTTTGTCTACAGTCCTGTGTCTGAACCTGAGCCTGAGCCCGAGCCAGAACCGGAGCCT
GCAAGACCTACCCGCCGTCCTAAAATGGCGCCTGCTATCCTGAGACGCCCGACATCACCTGTGTCTAGAGAATGC
AATAGTAGTACGGATAGCTGTGACTCCGGTCCTTCTAACACACCTCCTGAGATACACCCGGTGGTCCCGCTGTGC
CCCATTAAACCAGTTGCCGTGAGAGTTGGTGGGCGTCGCCAGGCTGTGGAATGTATCGAGGACTTGCTTAACGAG
CCTGGGCAACCTTTGGACTTGAGCTGTAAACGCCCCAGGCCATAA

(SEQ ID NO:40)
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GTGACGGCCCCCGAAGATCCCAACGAGGAGGCGGTTTCGCAGATTTTTCCCGACTCTGTAATGTTGGCGGTGCAG
GAAGGGATTGACTTACTCACTTTTCCGCCGGCGCCCGGTTCTCCGGAGCCGCCTCACCTTTCCCGGCAGCCCGAG
CAGCCGGAGCAGAGAGCCTTGGGTCCGGTTTCTATGCCAAACCTTGTACCGGAGGTGATCGATCTTACCTGCCAC
GAGGCTGGCTTTCCACCCAGTGACGACGAGGATGAAGAGGGTCCTGTGTCTGAACCTGAGCCTGAGCCCGAGCCA
GAACCGGAGCCTGCAAGACCTACCCGCCGTCCTAAAATGGCGCCTGCTATCCTGAGACGCCCGACATCACCTGTG
TCTAGAGAATGCAATAGTAGTACGGATAGCTGTGACTCCGGTCCTTCTAACACACCTCCTGAGATACACCCGGTG
GTCCCGCTGTGCCCCATTAAACCAGTTGCCGTGAGAGTTGGTGGGCGTCGCCAGGCTGTGGAATGTATCGAGGAC
TTGCTTAACGAGCCTGGGCAACCTTTGGACTTGAGCTGTAAACGCCCCAGGCCATAA

Ad5 E1B_19K

Accession: AAQ19286.1
(SEQ ID NO:41)

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TGGTTTTGGAGGTTTCTGTGGGGCTCATCCCAGGCAAAGTTAGTCTGCAGAATTAAGGAGGATTACAAGTGGGAA
TTTGAAGAGCTTTTGAAATCCTGTGGTGAGCTGTTTGATTCTTTGAATCTGGGTCACCAGGCGCTTTTCCAAGAG
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GTTGTGAGACACAAGAATCGCCTGCTACTGTTGTCTTCCGTCCGCCCGGCGATAATACCGACGGAGGAGCAGCAG
CAGCAGCAGGAGGAAGCCAGGCGGCGGCGGCAGGAGCAGAGCCCATGGAACCCGAGAGCCGGCCTGGACCCTCGG
GAATGA

Ad5 E1B_55K

Accession: AAQ19287.1
(SEQ ID NO:42)

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GCAGGAGGAAGCCAGGCGGCGGCGGCAGGAGCAGAGCCCATGGAACCCGAGAGCCGGCCTGGACCCTCGGGAATG
AATGTTGTACAGGTGGCTGAACTGTATCCAGAACTGAGACGCATTTTGACAATTACAGAGGATGGGCAGGGGCTA
AAGGGGGTAAAGAGGGAGCGGGGGGCTTGTGAGGCTACAGAGGAGGCTAGGAATCTAGCTTTTAGCTTAATGACC
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AAGTATTCCATAGAGCAGCTGACCACTTACTGGCTGCAGCCAGGGGATGATTTTGAGGAGGCTATTAGGGTATAT
GCAAAGGTGGCACTTAGGCCAGATTGCAAGTACAAGATCAGCAAACTTGTAAATATCAGGAATTGTTGCTACATT
TCTGGGAACGGGGCCGAGGTGGAGATAGATACGGAGGATAGGGTGGCCTTTAGATGTAGCATGATAAATATGTGG
CCGGGGGTGCTTGGCATGGACGGGGTGGTTATTATGAATGTAAGGTTTACTGGCCCCAATTTTAGCGGTACGGTT
TTCCTGGCCAATACCAACCTTATCCTACACGGTGTAAGCTTCTATGGGTTTAACAATACCTGTGTGGAAGCCTGG
ACCGATGTAAGGGTTCGGGGCTGTGCCTTTTACTGCTGCTGGAAGGGGGTGGTGTGTCGCCCCAAAAGCAGGGCT
TCAATTAAGAAATGCCTCTTTGAAAGGTGTACCTTGGGTATCCTGTCTGAGGGTAACTCCAGGGTGCGCCACAAT
GTGGCCTCCGACTGTGGTTGCTTCATGCTAGTGAAAAGCGTGGCTGTGATTAAGCATAACATGGTATGTGGCAAC
TGCGAGGACAGGGCCTCTCAGATGCTGACCTGCTCGGACGGCAACTGTCACCTGCTGAAGACCATTCACGTAGCC
AGCCACTCTCGCAAGGCCTGGCCAGTGTTTGAGCATAACATACTGACCCGCTGTTCCTTGCATTTGGGTAACAGG
AGGGGGGTGTTCCTACCTTACCAATGCAATTTGAGTCACACTAAGATATTGCTTGAGCCCGAGAGCATGTCCAAG
GTGAACCTGAACGGGGTGTTTGACATGACCATGAAGATCTGGAAGGTGCTGAGGTACGATGAGACCCGCACCAGG
TGCAGACCCTGCGAGTGTGGCGGTAAACATATTAGGAACCAGCCTGTGATGCTGGATGTGACCGAGGAGCTGAGG
CCCGATCACTTGGTGCTGGCCTGCACCCGCGCTGAGTTTGGCTCTAGCGATGAAGATACAGATTGA

Sequences for E2A and E4A are both contained within Accession MN088492

Ad5 E2A orf:
Accession: QHX41645.1
(SEQ ID NO:43)
ATGGCCAGTCGGGAAGAGGAGCAGCGCGAAACCACCCCCGAGCGCGGACGCGGTGCGGCGCGACGTCCACCAACC
ATGGAGGACGTGTCGTCCCCGTCGCCGTCGCCGCCGCCTCCCCGCGCGCCCCCAAAAAAGCGGCTGAGGCGGCGT
CTCGAGTCCGAGGACGAAGAAGACTCGTCACAAGATGCGCTGGTGCCGCGCACACCCAGCCCGCGGCCATCGACC
TCGACGGCGGATTTGGCCATTGCGTCCAAAAAGAAAAAGAAGCGCCCCTCTCCCAAGCCCGAGCGCCCGCCATCC
CCAGAGGTGATCGTGGACAGCGAGGAAGAAAGAGAAGATGTGGCGCTACAAATGGTGGGTTTCAGCAACCCACCG
GTGCTAATCAAGCACGGCAAGGGAGGTAAGCGCACGGTGCGGCGGCTGAATGAAGACGACCCAGTGGCGCGGGGT
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GTGACGAGCGAAAACGGGCAGCGCGCGCTGAAGGAGCAGTCTAGCAAGGCCAAGATCGTGAAGAACCGGTGGGGC
CGAAATGTGGTGCAGATCTCCAACACCGACGCAAGGTGCTGCGTGCATGACGCGGCCTGTCCGGCCAATCAGTTT
TCCGGCAAGTCTTGCGGCATGTTCTTCTCTGAAGGCGCAAAGGCTCAGGTGGCTTTTAAGCAGATCAAGGCTTTC
ATGCAGGCGCTGTATCCTAACGCCCAGACCGGGCACGGTCACCTTCTGATGCCACTACGGTGCGAGTGCAACTCA
AAGCCTGGGCATGCACCCTTTTTGGGAAGGCAGCTACCAAAGTTGACTCCGTTCGCCCTGAGCAACGCGGAGGAC
CTGGACGCGGATCTGATCTCCGACAAGAGCGTGCTGGCCAGCGTGCACCACCCGGCGCTGATAGTGTTCCAGTGC
TGCAACCCTGTGTATCGCAACTCGCGCGCGCAGGGCGGAGGCCCCAACTGCGACTTCAAGATATCGGCGCCCGAC
CTGCTAAACGCGTTGGTGATGGTGCGCAGCCTGTGGAGTGAAAACTTCACCGAGCTGCCGCGGATGGTTGTGCCT
GAGTTTAAGTGGAGCACTAAACACCAGTATCGCAACGTGTCCCTGCCAGTGGCGCATAGCGATGCGCGGCAGAAC
CCCTTTGATTTTTAA

Ad5 E4A:
Two proteins are present in this ORF. The first is a splice variant contained within the ORF. The second is a non-spliced transcript present in the ORF. Accession 1: QHX41659.1
Accession 2: QHX41660.1

(SEQ ID NO:44)

ATGACTACGTCCGGCGTTCCATTTGGCATGACACTACGACCAACACGATCTCGGTTGTCTCGGCGCACTCCGTAC
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TGTAACACTTTGACAATGCACAACGCGTGGACTTCCCCTTCGCCGCCCGTTGAGCAACCGCAAGTTGGACAGCAG
CCTGTGGCTCAGCAGCTGGACAGCGACATGAACTTAAGCGAGCTGCCCGGGGAGTTTATTAATATCACTGATGAG
CGTTTGGCTCGACAGGAAACCGTGTGGAATATAACACCTAAGAATATGTCTGTTACCCATGATATGATGCTTTTT
AAGGCCAGCCGGGGAGAAAGGACTGTGTACTCTGTGTGTTGGGAGGGAGGTGGCAGGTTGAATACTAGGGTTCTG
TGA

(SEQ ID NO:45)

ATGACTACGTCCGGCGTTCCATTTGGCATGACACTACGACCAACACGATCTCGGTTGTCTCGGCGCACTCCGTAC
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GAATGGGTTGTTCCCTGGGATATGGTTCTGACGCGGGAGGAGCTTGTAATCCTGAGGAAGTGTATGCACGTGTGC
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Ad5 VA:
Accession: AF369965.1
(SEQ ID NO:46)
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GCACGCCTCACCGGCCAGAGCGTCCCGACCATGGAGCACTTTTTGCCGCTGCGCAACATCTGGAACCGCGTCCGC
GACTTTCCGCGCGCCTCCACCACCGCCGCCGGCATCACCTGGATGTCCAGGTACATCTACGGATTACG

EXAMPLE 13 – Promoter, Operator, IRES and Intron Sequences

CMV Promoter (SEQ ID NO:47)

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AACGCCAATAGGGACTTTCCATTGACGTCAATGGGTGGAGTATTTACGGTAAACTGCCCACTTGGCAGTACATCA
AGTGTATCATATGCCAAGTACGCCCCCTATTGACGTCAATGACGGTAAATGGCCCGCCTGGCATTATGCCCAGTA
CATGACCTTATGGGACTTTCCTACTTGGCAGTACATCTACGTATTAGTCATCGCTATTACCATGGTGATGCGGTT
TTGGCAGTACATCAATGGGCGTGGATAGCGGTTTGACTCACGGGGATTTCCAAGTCTCCACCCCATTGACGTCAA
TGGGAGTTTGTTTTGGCACCAAAATCAACGGGACTTTCCAAAATGTCGTAACAACTCCGCCCCATTGACGCAAAT
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2xTet Operator Sequence (SEQ ID NO: 48)

TCCCTATCAGTGATAGAGATCTCCCTATCAGTGATAGAGA

hCMV Intron Sequence (SEQ ID NO: 49)

GTAAGTACCGCCTATAGAGTCTATAGGCCCACCCCCTTGGCTTCTTATGCATGCTATACTGTTTTTGGCTTGGGG
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TGACCACTCCCCTATTGGTGACGATACTTTCCATTACTAATCCATAACATGGCTCTTTGCCACAACTCTCTTTAT
TGGCTATATGCCAATACACTGTCCTTCAGAGACTGACACGGACTCTGTATTTTTACAGGATGGGGTCTCATTTAT
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ECMV IRES Sequence (SEQ ID NO: 50)

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FMDV IRES (SEQ ID NO: 51)
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TTAATGACCCT

EXAMPLE 14 – CHO and Mouse Stable Site 1 Sequences – U.S. Patent No. 7,771,997

211> 6473
<212> DNA
<213> Cricetulus griseus

<400> 1
(SEQ ID NO:52)

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gatggtagaa taaaaatctg aaacataggt gatgtatctg ccatactgca tgggtgtgta 180
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cacagagagg gccagaagca ctcagaactc caggggggtca ggagtggttc tctggaggct 420
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EXAMPLE 15 – CHO Stable Site 2 Sequences – U.S. Patent No. 9,816,110

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It is to be understood that the description, specific examples and data, while indicating exemplary embodiments, are given by way of illustration and are not intended to limit the present inventions. Various changes and modifications within the present invention, including combining embodiments in whole and in part, will become apparent to the skilled artisan from the discussion, disclosure and data contained herein, and thus are considered part of the inventions.

What is Claimed Is:

1. A polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding Adeno-associated virus (AAV) Cap protein, and (v) a polyadenylation site, wherein the polynucleotide allows for production of AAV Cap VP2 and VP3.

2. The polynucleotide according to claim 1, wherein the polynucleotide is in a CHO cell and the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein are operably linked.

3. The polynucleotide according to claim 2, wherein the polynucleotide is integrated into a CHO cell genome or a BHK cell genome.

4. The polynucleotide according to claim 1, wherein the polynucleotide is in a HEK 293 cell and the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein are operably linked.

5. The polynucleotide according to claim 2, wherein the polynucleotide is integrated into a HEK 293 cell genome or a human amniotic cell genome.

6. The polynucleotide according to claim 1, further comprising an operator.

7. The polynucleotide according to claim 6, wherein the promoter is a CMV promoter and the operator is a Tet operator.

8. A polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a first internal ribosome entry site, (iv) a first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, (vi) a second polynucleotide encoding AAV Cap protein, and (vii) a polyadenylation site.

9. The polynucleotide according to claim 8, wherein the polynucleotide is in a CHO cell and the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked.

10. The polynucleotide according to claim 9, wherein the polynucleotide is integrated into a CHO cell genome or a BHK cell genome.

11. The polynucleotide according to claim 8, wherein the polynucleotide is in a HEK 293 cell and the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked.

12. The polynucleotide according to claim 11 wherein the polynucleotide is integrated into a HEK 293 cell genome or a human amniotic cell genome.

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13. The polynucleotide according to claim 8, further comprising an operator.

14. The polynucleotide according to claim 13, wherein the promoter is a CMV promoter and the operator is a Tet operator.

15. A polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site, wherein the polynucleotide allows for production of AAV Cap VP1 protein when expressed.

16. The polynucleotide according to claim 15, wherein the polynucleotide is in a CHO cell and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein are operably linked.

17. The polynucleotide according to claim 16, wherein the polynucleotide is integrated into a CHO cell genome or a BHK cell genome .

18. The polynucleotide according to claim 15, wherein the polynucleotide is in a HEK 293 cell and the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein are operably linked.

19. The polynucleotide according to claim 16, wherein the polynucleotide is integrated into a HEK 293 cell genome or a human amniotic cell genome .

20. The polynucleotide according to claim 15, further comprising an operator.

21. The polynucleotide according to claim 20, wherein the promoter is a CMV promoter and the operator is a Tet operator.

22. A polynucleotide comprising (i) a promoter, (ii) an internal ribosome binding site, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site, wherein the polynucleotide allows for production of AAV VP1 Cap protein when expressed.

23. The polynucleotide according to claim 22 wherein the polynucleotide is in a CHO cell and the (i) promoter, (ii) internal ribosome binding site, and (iii) polynucleotide encoding AAV Cap protein are operably linked.

24. The polynucleotide according to claim 23, wherein the polynucleotide is integrated into a CHO genome or a BHK cell genome.

25. The polynucleotide according to claim 22 wherein the polynucleotide is in a HEK 293 cell and the (i) promoter, (ii) internal ribosome binding site, and (iii) polynucleotide encoding AAV Cap protein are operably linked.

26. The polynucleotide according to claim 25, wherein the polynucleotide is integrated into a HEK 293 cell genome or a human amniotic cell genome.

27. The polynucleotide according to claim 22, further comprising an operator.

28. The polynucleotide according to claim 27, wherein the promoter is a CMV promoter and the operator is a Tet operator.

29. A eukaryotic cell comprising a polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a first internal ribosome entry site, (iv) a first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, (vi) a second polynucleotide encoding AAV Cap protein, and (vii) a polyadenylation site.

30. The cell according to claim 29, wherein the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked.

31. The cell according to claim 30, wherein the polynucleotide is integrated into a cell genome.

32. The cell according to claim 29, wherein the cell is a CHO cell or a BHK cell.

33. The cell according to claim 29, wherein the cell is a HEK 293 cell or a human amniotic cell.

34. The cell according to claim 29, further comprising an operator.

35. The cell according to claim 34, wherein the promoter is a CMV promoter and the operator is a Tet operator.

36. The cell according to claim 29, further comprising:

a polynucleotide encoding AAV Rep;

a polynucleotide encoding Ad E1A;

a polynucleotide encoding Ad E1B;

a polynucleotide encoding Ad E2A or E2A orf;

a polynucleotide encoding Ad E4 or E4 orf 6;

a polynucleotide encoding VA RNA; and

a polynucleotide encoding AAV ITRs and a protein of interest.

37. A eukaryotic cell comprising

(A) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and

(B) a second polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site.

38. The cell according to claim 37, wherein

the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (A) first polynucleotide are operably linked, and wherein

the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein of (B) second polynucleotide are operably linked.

39. The cell according to claim 38, wherein at least one polynucleotide is integrated into a cell genome.

40. The cell according to claim 37, wherein the cell is a CHO cell or a BHK cell.

41. The cell according to claim 37, wherein the cell is a HEK 293 cell or a human amniotic cell.

42. The cell according to claim 37, further comprising an operator.

43. The cell according to claim 42, wherein the promoter is a CMV promoter and the operator is a Tet operator.

44. The cell according to claim 37, further comprising:

a polynucleotide encoding AAV Rep;

a polynucleotide encoding Ad E1A;

a polynucleotide encoding Ad E1B;

a polynucleotide encoding Ad E2A or E2A orf;

a polynucleotide encoding Ad E4 or E4 orf 6;

a polynucleotide encoding VA RNA; and

a polynucleotide encoding AAV ITRs and a protein of interest.

45. A eukaryotic cell comprising

(A) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and

(B) a second polynucleotide comprising (i) a promoter, (ii) an internal ribosome entry site, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site.

46. The cell according to claim 45, wherein

the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (A) first polynucleotide are operably linked, and wherein

the (i) promoter, (ii) internal ribosome entry site, and (iii) polynucleotide encoding AAV Cap protein of (B) second polynucleotide are operably linked.

47. The cell according to claim 46, wherein at least one polynucleotide is integrated into a cell genome.

48. The cell according to claim 45, wherein the cell is a CHO cell or a BHK cell.

49. The cell according to claim 45, wherein the cell is a HEK 293 cell or a human amniotic cell.

50. The cell according to claim 45, further comprising an operator.

51. The cell according to claim 50, wherein the promoter is a CMV promoter and the operator is a Tet operator.

52. The cell according to claim 45, further comprising:

a polynucleotide encoding AAV Rep;

a polynucleotide encoding Ad E1A;

a polynucleotide encoding Ad E1B;

a polynucleotide encoding Ad E2A or E2A orf;

a polynucleotide encoding E4 or E4 orf 6;

a polynucleotide encoding VA RNA; and

a polynucleotide encoding AAV ITRs and a protein of interest.

53. A cell culture comprising more than one cell according to claims 29-36 in a culture medium.

54. A cell culture comprising more than one cell according to claims 37-44 in a culture medium.

55. A cell culture comprising more than one cell according to claims 45-52 in a culture medium.

56. A method of producing adeno-associated virus (AAV) Cap protein in cell culture, wherein the method comprises the steps of:

providing eukaryotic cells, wherein a cell comprises a polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a first internal ribosome entry site, (iv) a first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, (vi) a second polynucleotide encoding AAV Cap protein, and (vii) a polyadenylation site; and

culturing the cells in a culture medium to allow the cells to produce AAV Cap protein.

57. The method according to claim 56, wherein the (i) promoter, (ii) intron, (iii) first internal ribosome entry site, (iv) first polynucleotide encoding AAV Cap protein, (v) a second internal ribosome entry site, and (vi) second polynucleotide encoding AAV Cap protein are operably linked.

58. The method according to claim 57, wherein the polynucleotide is integrated into a cell genome.

59. The method according to claim 56, wherein the cell is a CHO cell or a BHK cell.

60. The method according to claim 56, wherein the cell is a HEK 293 cell or a human amniotic cell.

61. The method according to claim 56, wherein the cell further comprises an operator.

62. The method according to claim 61, wherein the promoter is a CMV promoter and the operator is a Tet operator.

63. The method according to claim 56, wherein the cell further comprises:
a polynucleotide encoding AAV Rep;
a polynucleotide encoding Ad E1A;
a polynucleotide encoding Ad E1B;
a polynucleotide encoding Ad E2A or E2A orf;
a polynucleotide encoding E4 or E4 orf 6;
a polynucleotide encoding VA RNA; and
a polynucleotide encoding AAV ITRs and a protein of interest, wherein the cell can produce recombinant AAV.

64. A method of producing adeno-associated virus (AAV) Cap protein in cell culture, wherein the method comprises the steps of

providing eukaryotic cells, where a cell comprises

(a) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and

(b) a second polynucleotide comprising (i) a promoter, (ii) an intron, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site; and

culturing the cells in a culture medium to allow the cells to produce AAV Cap protein, wherein the polynucleotide allows for production of AAV Cap proteins VP1, VP2 and VP3.

65. The method according to claim 64, wherein

the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (a) first polynucleotide are operably linked, and wherein

the (i) promoter, (ii) intron, and (iii) polynucleotide encoding AAV Cap protein of (b) second polynucleotide are operably linked.

66. The method according to claim 64, wherein the polynucleotide is integrated into a cell genome.

67. The method according to claim 64, wherein the cell is a CHO cell or a BHK cell.

68. The method according to claim 64, wherein the cell is a HEK 293 cell or a human amniotic cell.

69. The method according to claim 64, wherein the cell further comprises an operator.

70. The method according to claim 69, wherein the promoter is a CMV promoter and the operator is a Tet operator.

71. The method according to claim 64, wherein the cell further comprises:
a polynucleotide encoding AAV Rep;
a polynucleotide encoding Ad E1A;
a polynucleotide encoding Ad E1B;
a polynucleotide encoding Ad E2A or E2A orf;
a polynucleotide encoding Ad E4 or E4 orf 6;
a polynucleotide encoding VA RNA,; and
a polynucleotide encoding AAV ITRs and a protein of interest, wherein the cell can produce recombinant AAV.

72. A method of producing adeno-associated virus (AAV) Cap protein in cell culture, wherein the method comprises the steps of
providing eukaryotic cells, where a cell comprises

(a) a first polynucleotide comprising (i) a promoter, (ii) an intron, (iii) an internal ribosome entry site, (iv) a polynucleotide encoding AAV Cap protein, and (v) a polyadenylation site; and

(b) a second polynucleotide comprising (i) a promoter, (ii) an internal ribosome entry site, (iii) a polynucleotide encoding AAV Cap protein, and (iv) a polyadenylation site; and

culturing the cells in a culture medium to allow the cells to produce AAV Cap protein, wherein the polynucleotide allows for production of AAV Cap proteins VP1, VP2 and VP3.

73. The method according to claim 72, wherein

the (i) promoter, (ii) intron, (iii) internal ribosome entry site and (iv) polynucleotide encoding AAV Cap protein of (a) first polynucleotide are operably linked, and wherein

the (i) promoter, (ii) internal ribosome entry site, and (iii) polynucleotide encoding AAV Cap protein of (b) second polynucleotide are operably linked.

74. The method according to claim 72, wherein the polynucleotide is integrated into a cell genome.

75. The method according to claim 72, wherein the cell is a CHO cell or a BHK cell.

76. The method according to claim 72, wherein the cell is a HEK 293 cell or a human amniotic cell.

77. The method according to claim 72, wherein the cell further comprises an operator.

78. The method according to claim 77, wherein the promoter is a CMV promoter and the operator is a Tet operator.

79. The method according to claim 72, wherein the cell further comprises:
a polynucleotide encoding AAV Rep;
a polynucleotide encoding Ad E1A;
a polynucleotide encoding Ad E1B;
a polynucleotide encoding Ad E2A or E2A orf;
a polynucleotide encoding Ad E4 or E4 orf 6;
a polynucleotide encoding VA RNA; and
a polynucleotide encoding AAV ITRs and a protein of interest, wherein the cell can produce recombinant AAV.

80. A cell according to any of the above methods.

81. A method using any of the above cells.

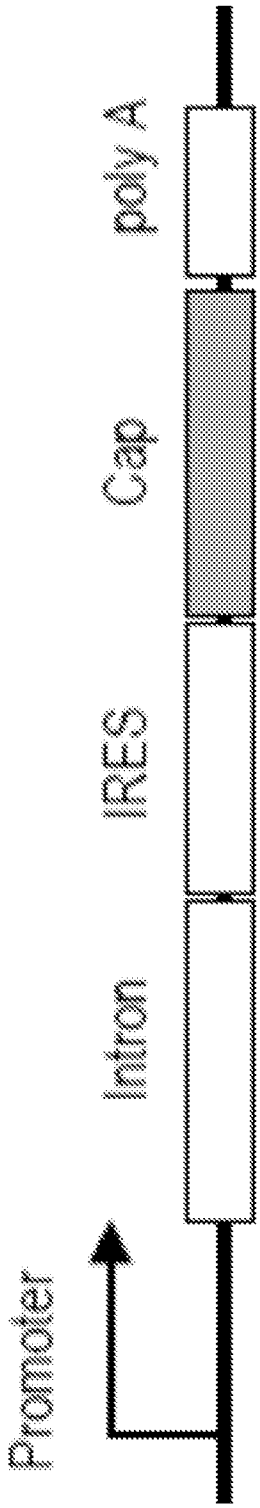

Figure 1

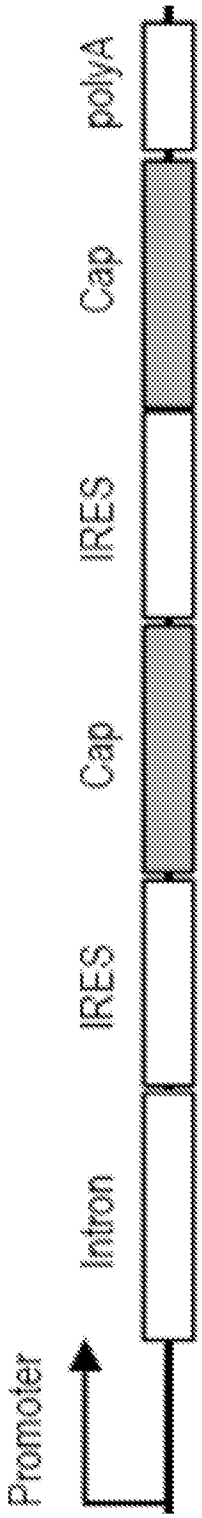

Figure 2

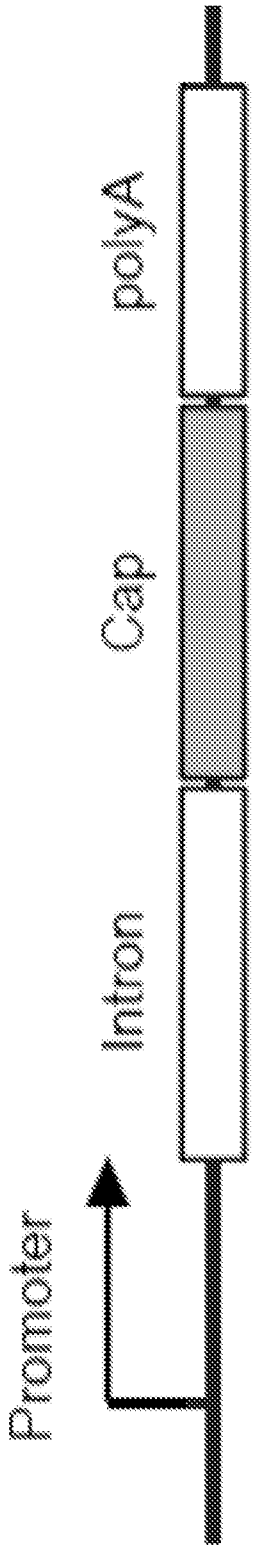

Figure 3

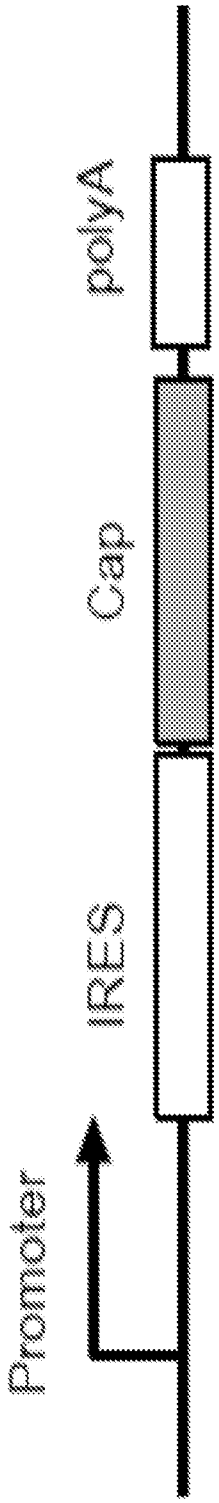

Figure 4

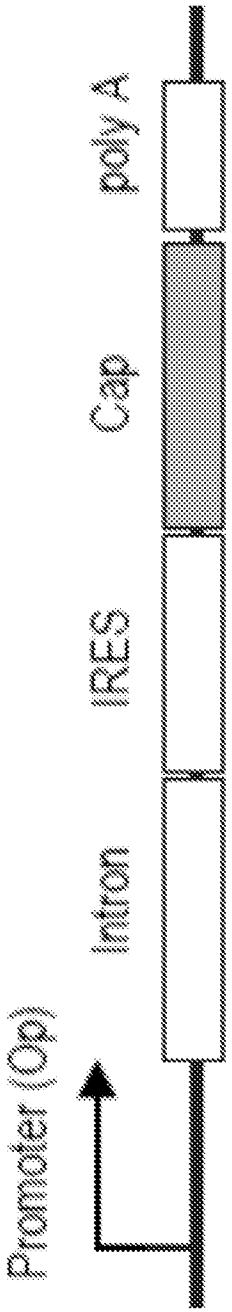

Figure 5

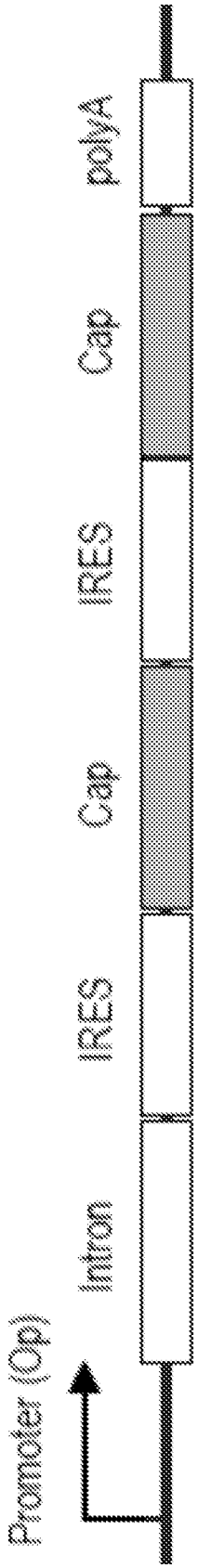

Figure 6

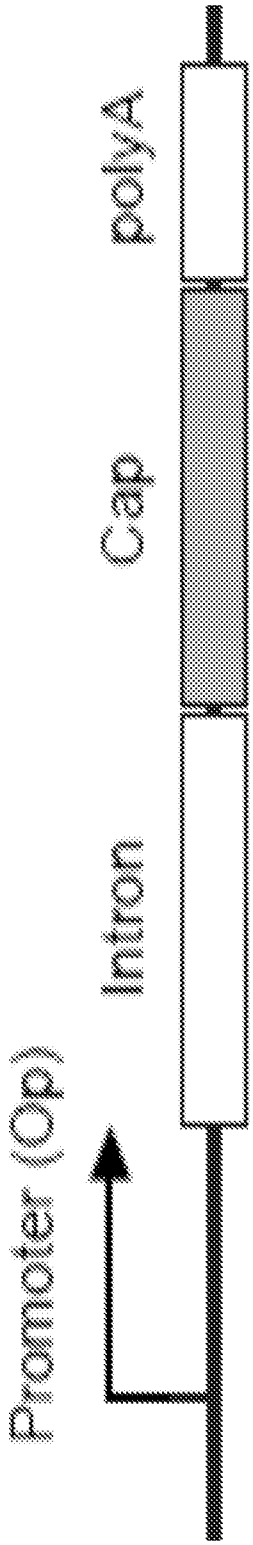

Figure 7

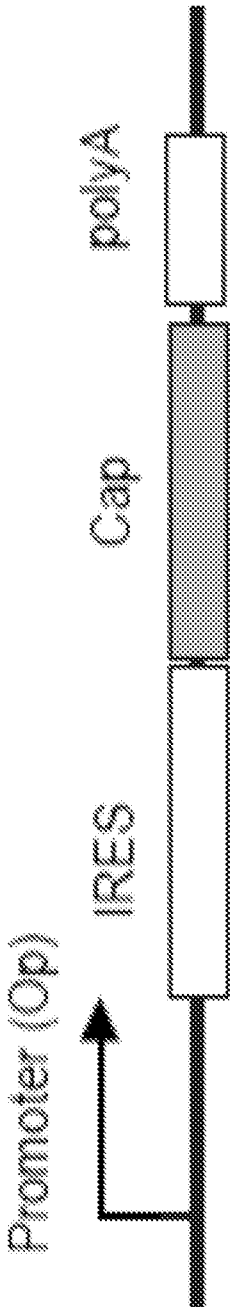

Figure 8

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2022/078266

A. CLASSIFICATION OF SUBJECT MATTER
INV. C12N15/86
ADD.

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C12N C07K

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

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

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	AYUSO EDUARD ET AL: "Production, Purification and Characterization of Adeno-Associated Vectors", CURRENT GENE THERAPY, BENTHAM SCIENCE PUBLISHERS LTD, NL, vol. 10, no. 6, 30 November 2010 (2010-11-30), pages 423-436, XP009516742, ISSN: 1566-5232, DOI: 10.2174/156652310793797685	15,22
Y	figure 1	1-14, 16-21, 23-81

X	US 10 711 274 B2 (UNIV MASSACHUSETTS [US]) 14 July 2020 (2020-07-14) page 35 - page 37	1-55

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[X] Further documents are listed in the continuation of Box C.

[X] See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

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"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance;; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
6 February 2023	17/02/2023

Name and mailing address of the ISA/	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Merkl, Philipp

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Form PCT/ISA/210 (second sheet) (April 2005)

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2022/078266

Box No. I Nucleotide and/or amino acid sequence(s) (Continuation of item 1.c of the first sheet)

1. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international search was carried out on the basis of a sequence listing:

   a. [X] forming part of the international application as filed.

   b. [ ] furnished subsequent to the international filing date for the purposes of international search (Rule 13*ter*.1(a)).

      [ ] accompanied by a statement to the effect that the sequence listing does not go beyond the disclosure in the international application as filed.

2. [ ] With regard to any nucleotide and/or amino acid sequence disclosed in the international application, this report has been established to the extent that a meaningful search could be carried out without a WIPO Standard ST.26 compliant sequence listing.

3. Additional comments:

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2022/078266

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	DOUIN VICTORINE ET AL: "Use and comparison of different internal ribosomal entry sites (IRES) in tricistronic retroviral vectors", BMC BIOTECHNOLOGY, BIOMED CENTRAL LTD, vol. 4, no. 1, 27 July 2004 (2004-07-27), page 16, XP021005932, ISSN: 1472-6750, DOI: 10.1186/1472-6750-4-16 the whole document	1-14, 16-21, 23-81
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