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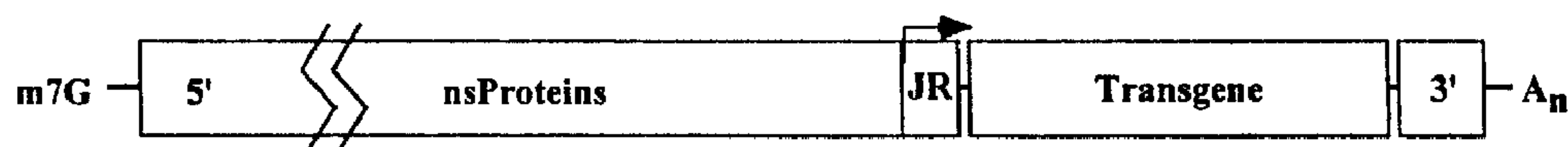
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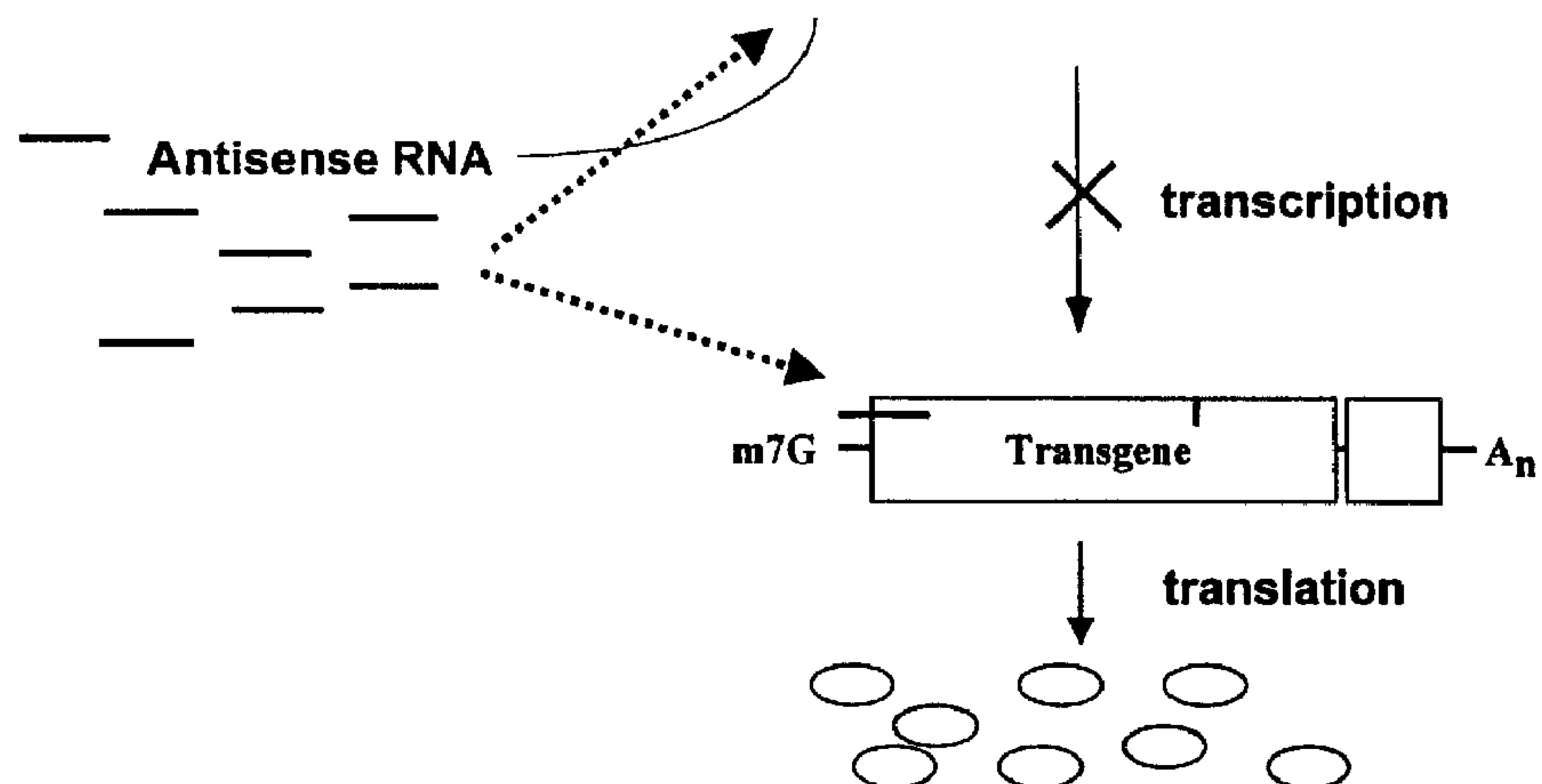
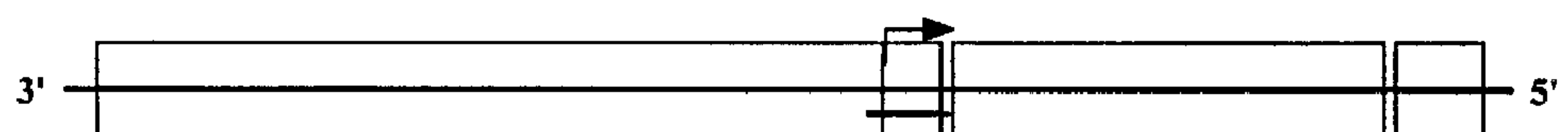
(54) Titre : COMPOSITIONS ET METHODES PERMETTANT L'ENCAPSIDATION DE VECTEURS D'ALPHAVIRUS
(54) Title: COMPOSITIONS AND METHODS FOR PACKAGING OF ALPHAVIRUS VECTORS

Reduction of Transgene Translation Using an Antisense RNA

Vector Replicon RNA



Vector Replicon minus strand RNA



(57) Abrégé/Abstract:

Nucleic acid molecules are provided comprising a nucleic acid sequence which encodes, in order, an alphavirus capsid, a signal peptide, and an alphavirus E1 or E2 glycoprotein. Also provided are vectors encoding such nucleic acid molecules, and use of

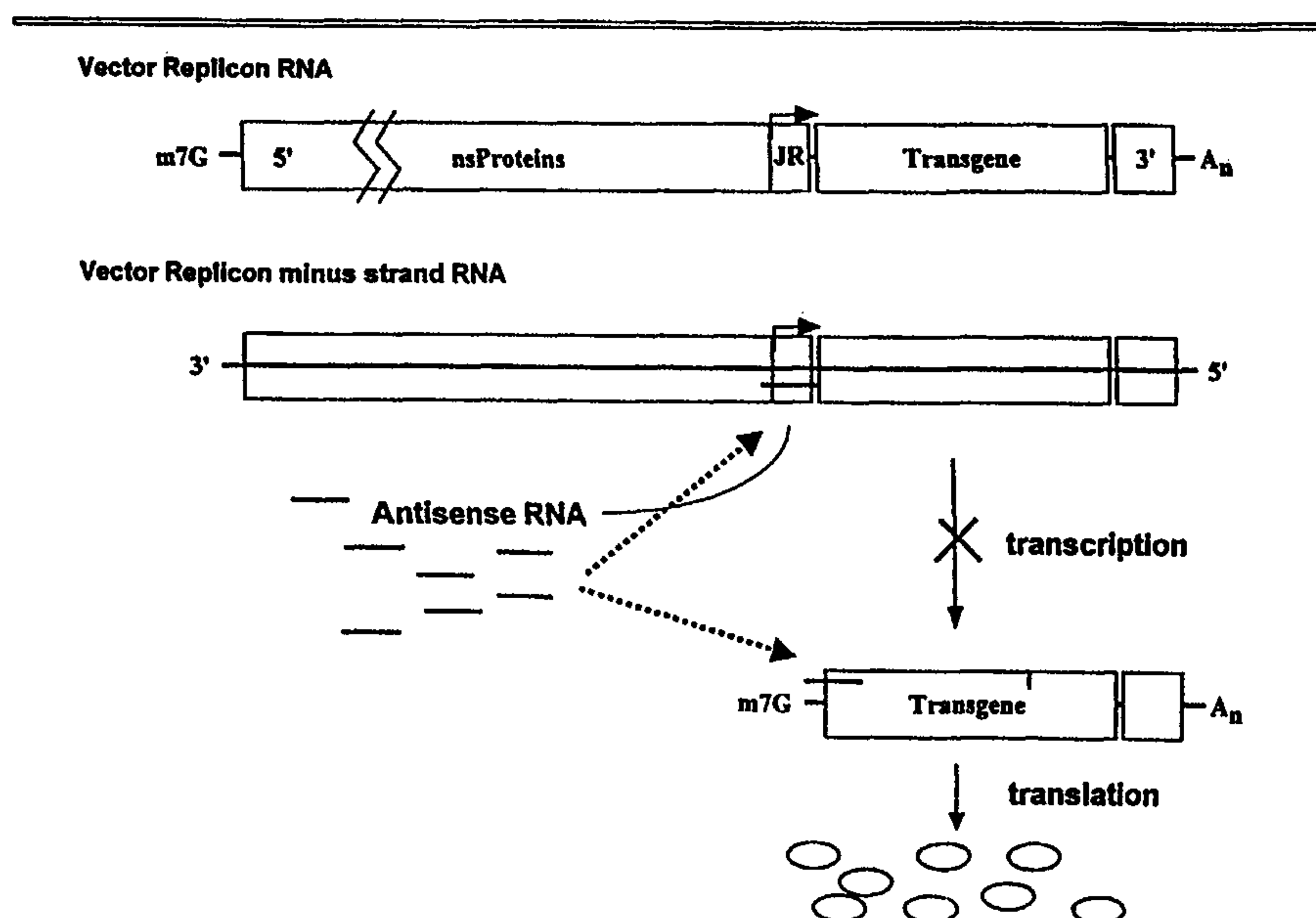
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such vectors or expression cassettes to generate recombinant alphavirus particles and alphavirus packaging cell lines. In addition, modified alphavirus vector constructs are provided that permit reduced transgene expression during vector packaging, as well as methods of using such vector constructs for the production of alphavirus vector particles.

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(54) Title: COMPOSITIONS AND METHODS FOR PACKAGING OF ALPHAVIRUS VECTORS**Reduction of Transgene Translation Using an Antisense RNA****(57) Abstract**

Nucleic acid molecules are provided comprising a nucleic acid sequence which encodes, in order, an alphavirus capsid, a signal peptide, and an alphavirus E1 or E2 glycoprotein. Also provided are vectors encoding such nucleic acid molecules, and use of such vectors or expression cassettes to generate recombinant alphavirus particles and alphavirus packaging cell lines. In addition, modified alphavirus vector constructs are provided that permit reduced transgene expression during vector packaging, as well as methods of using such vector constructs for the production of alphavirus vector particles.

COMPOSITIONS AND METHODS FOR PACKAGING OF ALPHAVIRUS VECTORS

TECHNICAL FIELD

The present invention relates generally to recombinant DNA technology;
5 and more specifically, to the development of packaging systems for the high level
production of recombinant alphavirus vector particles useful for directing expression of
one or more heterologous gene products.

BACKGROUND OF THE INVENTION

Alphaviruses comprise a group of genetically, structurally, and
10 serologically related arthropod-borne viruses of the *Togaviridae* family. These viruses
are distributed worldwide, and persist in nature through a mosquito to vertebrate cycle.
Birds, rodents, horses, primates, and humans are among the defined alphavirus
vertebrate reservoir/hosts.

Twenty-six known viruses and virus subtypes have been classified
15 within the alphavirus genus utilizing the hemagglutination inhibition (HI) assay. This
assay segregates the 26 alphaviruses into three major complexes: the Venezuelan equine
encephalitis (VEE) complex, the Semliki Forest (SF) complex, and the western equine
encephalitis (WEE) complex. In addition, four other viruses, eastern equine
encephalitis (EEE), Barmah Forest, Middelburg, and Ndumu, receive individual
20 classification based on the HI serological assay.

Members of the alphavirus genus also are classified based on their
relative clinical features in humans: alphaviruses associated primarily with
encephalitis, and alphaviruses associated primarily with fever, rash, and polyarthrititis.
Included in the former group are the VEE and WEE complexes, and EEE. In general,
25 infection with this group can result in permanent sequelae, including death. In the latter
group is the SF complex, comprised of the individual alphaviruses Semliki Forest,
Sindbis, Ross River, Chikungunya, O'nyong-nyong, and Mayaro. Although serious

epidemics have been reported, infection by viruses of this group is generally self-limiting, without permanent sequelae.

Sindbis virus is the prototype member of the *Alphavirus* genus of the *Togaviridae* family. Its replication strategy is well characterized and serves as a model
5 for other alphaviruses (Strauss and Strauss, 1994, *Microbio. Rev.*, 58:491-562). The genome of Sindbis virus (like other alphaviruses) is an approximately 12 kb single-stranded, positive-sense RNA molecule that is capped and polyadenylated. Genome RNA is contained within a virus-encoded capsid protein shell which is, in turn, surrounded by a host-derived lipid envelope from which two viral-specific
10 glycoproteins, E1 and E2, protrude as spikes from the virion surface. Certain alphaviruses (*e.g.*, SF) also maintain an additional protein, E3, which is a cleavage product from the E2 precursor protein, PE2.

After virus particle absorption to target cells, penetration, and uncoating of the nucleocapsid to release viral genomic RNA into the cytoplasm, the replication
15 process is initiated by translation of four nonstructural replicase proteins (nsP1-nsP4) from the 5' two-thirds of the viral genome. The four nsPs are translated as one of two polyproteins (nsP123 or nsP1234), and processed post-translationally into mature monomeric proteins by an active protease in the C-terminal domain of nsP2. Both of the nonstructural polyproteins and their derived monomeric units may participate in the
20 RNA replication process, which involves nsP binding to the conserved nucleotide sequence elements (CSEs) present at the 5' and 3' ends, and an internal subgenomic junction region promoter.

The positive strand genome RNA serves as template for the nsP-catalyzed synthesis of a full-length complementary negative strand RNA. Synthesis of
25 the negative strand RNA is catalyzed by binding of an nsP complex to the 3' terminal CSE of the positive strand genome RNA. The negative strand, in turn, serves as template for the synthesis of additional positive strand genome RNA, as well as an abundant subgenomic RNA, initiated internally at the junction region promoter. Synthesis of additional positive strand genome RNA occurs after binding of an nsP
30 complex to the 3' terminal CSE of the complementary negative strand genome-length

RNA template. Synthesis of the subgenomic mRNA from the negative strand RNA template is initiated from the junction region promoter. Thus, the 5' end and junction region CSEs of the positive strand genome RNA are functional only after being transcribed into the negative strand RNA complement (*i.e.*, the 5' end CSE is functional when it is the 3' end of the genomic negative stranded complement).

Alphavirus structural proteins (sPs) are translated from the subgenomic RNA, which represents the 3' one-third of the genome, and like the nsPs, are processed post-translationally into the individual proteins (Figure 1). Translation of this subgenomic mRNA produces a single polyprotein consisting of the structural proteins capsid (C) glycoprotein E2 and glycoprotein E1, plus the corresponding leader/signal sequences (E3, 6k) for glycoprotein insertion into the endoplasmic reticulum. The structural gene polyprotein is processed into the mature protein species by a combination of viral (capsid autoprotease) and cellular proteases (*e.g.*, signal peptidase). Alphavirus structural proteins are produced at very high levels due to the abundance of subgenomic mRNA transcribed, as well as the presence of a translational enhancer element (Frolov and Schlesinger, 1994, *J. Virol.* 68:8111-8117; Sjoberg et al., 1994, *Bio/Technol.* 12:1127-1131) within the mRNA, located in the capsid gene coding sequence. Because all structural proteins are synthesized at equimolar ratios, as part of the polyprotein, the translation enhancer element exerts its effect equally on each of the genes.

The general strategy for construction of alphavirus-based expression vectors has been to substitute the viral structural protein genes with a heterologous gene, maintaining transcriptional control via the highly active subgenomic RNA promoter. Vectors of this configuration are termed RNA "replicons" and may be transcribed *in vitro* from cDNA using a bacteriophage promoter, or, generated *in vivo* directly from DNA when linked to a eukaryotic promoter. Currently, alphavirus replicon RNA is packaged into recombinant vector particles by transient co-transfection with *in vitro* transcribed defective helper RNA, or, using stable packaging cell lines having structural protein expression cassettes. The structural protein expression cassette(s) used for vector packaging encode either the intact "native" alphavirus

structural polyprotein that is post-translationally processed into mature C, E2, and E1; or, alphavirus structural proteins that have been split into separate cassettes encoding either C or E2/E1.

As described in the detailed description and examples below, the present invention provides new compositions and methods for packaging of alphavirus particles, including novel alphavirus structural polyprotein genes, expression cassettes containing the genes, polypeptides expressed from the cassettes, and methods of using the genes, cassettes, and polypeptides for the high level packaging of alphavirus vector RNA into recombinant alphavirus vector particles in the absence of contaminating replication-competent virus. The invention also provides modifications to alphavirus vector systems wherein expression of the heterologous transgene from an alphavirus vector replicon may be reduced (suppressed) in desired cells, including during the vector packaging process. This specific reduction in transgene expression provides a means to increase the level of recombinant alphavirus vector particles produced.

SUMMARY OF THE INVENTION

Briefly stated, the present invention provides methods for generating high titer recombinant alphavirus particle preparations in the absence of contaminating replication-competent virus, as well as the generation of stable alphavirus packaging cells. Within one aspect of the invention nucleic acid molecules are provided wherein the molecule encodes an alphavirus envelope glycoprotein E2 or E1 independent of the other glycoprotein, as an in-frame fusion with a leader/signal peptide sequence of alphavirus (*e.g.*, E3, 6K) or non-alphavirus (*e.g.*, TPA) origin.

In another aspect of the invention nucleic acid molecules are provided wherein the molecule encodes in a single open-reading frame (ORF), a polypeptide that is a polyprotein comprising in order, an alphavirus capsid protein, a leader/signal peptide, and an alphavirus glycoprotein E2, with the proviso that the molecule does not encode an alphavirus glycoprotein E1. In one embodiment, the leader/signal sequence is an alphavirus E3 peptide. In another embodiment, the leader/signal sequence is of non-alphavirus origin.

In another aspect of the invention nucleic acid molecules are provided wherein the molecule encodes in a single open-reading frame (ORF), a polypeptide that is a polyprotein comprising in order, an alphavirus capsid protein, a leader/signal peptide, and an alphavirus glycoprotein E1, with the proviso that the molecule does not
5 encode an alphavirus glycoprotein E2.. In one embodiment, the encoded leader/signal peptide is an alphavirus E3 peptide. In another embodiment, the encoded leader/signal peptide is an alphavirus 6K peptide. In yet another embodiment, the encoded leader/signal peptide is of non-alphavirus origin.

In another aspect of the invention nucleic acid molecules are provided
10 wherein the molecule encodes a single open-reading frame (ORF), a polypeptide that is a polyprotein comprising in order, an alphavirus capsid protein, a first leader/signal peptide, a second leader/signal peptide, and an alphavirus glycoprotein E1. In a preferred embodiment, the first encoded leader/signal peptide is an alphavirus E3 peptide and the second encoded leader/signal peptide is an alphavirus 6K peptide.
15 Within this aspect it is preferred that the nucleic acid molecule not encode an alphavirus E2 protein

Within any of the above-noted aspects, the nucleic acid molecules provided herein may be 'isolated' (*i.e.*, not integrated into the genomic DNA of an organism, or, in the case of a virus, is separated from the complete virus genome). Within
20 preferred embodiments, the nucleic acid molecules provided herein do not contain all elements of the wild-type structural polyprotein gene (*i.e.*, C, E3, E2, 6k and E1) in order.

In other aspects of the invention expression cassettes are provided wherein the cassette directs the expression of an alphavirus structural polypeptide of the present invention. In preferred embodiments, the cassette directs the expression of a
25 polypeptide that is a polyprotein selected from the group consisting of 1) an alphavirus capsid protein, a leader/signal peptide, and an alphavirus glycoprotein E2; 2) an alphavirus capsid protein, a leader/signal peptide, and an alphavirus glycoprotein E1; 3) an alphavirus capsid protein, a first leader/signal peptide, a second leader/signal peptide, and an alphavirus glycoprotein E1; and 4) an alphavirus capsid protein, a first
30 leader/signal peptide, a second leader/signal peptide, and an alphavirus glycoprotein E2.

In particularly preferred embodiments, the cassette directs the expression of a polypeptide that is a polyprotein selected from the group consisting of 1) an alphavirus capsid protein, an alphavirus E3 peptide, and an alphavirus glycoprotein E2; 2) an alphavirus capsid protein, a leader/signal peptide of non-alphavirus origin, and an alphavirus glycoprotein E1; 3) an alphavirus capsid protein, an alphavirus E3 peptide, and an alphavirus glycoprotein E1; 4) an alphavirus capsid protein, an alphavirus 6K peptide, and an alphavirus glycoprotein E1; and 5) an alphavirus capsid protein, an alphavirus E3 peptide, an alphavirus 6K peptide, and an alphavirus glycoprotein E1.

In one embodiment, alphavirus structural protein expression cassettes of the present invention are provided wherein the cassette comprises a DNA promoter of RNA transcription (*e.g.*, RNA polymerase II promoter) operably linked to a structural protein open reading frame (ORF), and a transcription termination/polyadenylation sequence.

In another embodiment, the alphavirus structural protein expression cassette is an alphavirus defective helper RNA transcribed *in vitro* or *in vivo*, comprising the ordered elements: 5' viral or defective-interfering RNA sequence required *in cis* for alphaviral replication (also referred to as 5' CSE, or, the "5' sequence which initiates transcription of alphavirus RNA", in the background), viral subgenomic junction region promoter, alphavirus structural protein sequence of the present invention, 3' alphaviral sequence required *in cis* for replication (also referred to as 3' CSE or the "3' alphavirus RNA polymerase recognition sequence", in the background), and, optionally, a polyadenylate tract. Preferably, such expression cassettes do not encode all functional non-structural proteins. Within further embodiments, the expression cassettes further comprise a 5' promoter upstream of the expression cassette which drives transcription of the expression cassette (*e.g.*, *in vitro*, or alternatively, in a eukaryotic cell).

In yet another embodiment, the alphavirus structural protein expression cassette is a DNA cassette comprising the ordered elements: DNA promoter of RNA transcription that functions within a eukaryotic cell (*e.g.*, RNA polymerase II promoter), 5' viral or defective-interfering RNA sequence required *in cis* for alphaviral replication,

viral subgenomic junction region promoter, alphavirus structural protein sequence of the present invention. 3' alphaviral sequence required in *cis* for replication, polyadenylate tract, and transcription termination/polyadenylation sequence.

In another aspect of the invention, alphavirus structural polypeptides are provided wherein the polypeptides are polyproteins selected from the group consisting of: 1) an alphavirus capsid protein, an alphavirus E3 peptide, and an alphavirus glycoprotein E2; 2) an alphavirus capsid protein, a leader/signal peptide of non-alphavirus origin, and an alphavirus glycoprotein E1; 3) an alphavirus capsid protein, an alphavirus E3 peptide, and an alphavirus glycoprotein E1; 4) an alphavirus capsid protein, an alphavirus 6K peptide, and an alphavirus glycoprotein E1; and 5) an alphavirus capsid protein, an alphavirus E3 peptide, an alphavirus 6K peptide, and an alphavirus glycoprotein E1.

In another aspect, expression cassettes are provided comprising a promoter which is operably linked to a nucleic acid molecule, which when transcribed produces an RNA sequence complementary to an alphavirus junction region promoter, or, alphavirus subgenomic RNA, wherein said nucleic acid molecule is less than 500, 250, 100, or 50 nucleotides in length. Preferably the RNA sequence complementary to an alphavirus subgenomic RNA is complementary to at least a portion of the 5' end nontranslated region and translation initiation codon of the subgenomic RNA.

In another aspect of the invention, host cells are provided which contain the alphavirus expression cassettes described herein. In preferred embodiments, the cells are stably transformed with the structural protein expression cassettes. In particularly preferred embodiments, the cells are of mammalian origin and are alphavirus packaging cells.

Within certain embodiments of the invention, the elements of the alphavirus expression cassettes may be selected from different alphaviruses (*e.g.*, a packaging cell may be comprised of a first alphavirus expression cassette: C, E3, E2 and a second alphavirus expression cassette: C, 6k, E1, wherein E3 and 6k are from different alphaviruses). Within certain embodiments the two expression cassettes may

be placed onto a single vector (*e.g.*, with opposite and divergent transcriptional orientation).

Within other aspects of the present invention alphavirus packaging cells are provided, comprising (a) a first expression cassette which directs the expression of a first nucleic acid molecule, comprising a nucleic acid sequence which encodes, in order, an alphavirus capsid, a signal peptide, and an alphavirus E1 glycoprotein, with the proviso that the first nucleic acid molecule does not encode an alphavirus E2 glycoprotein; and (b) a second expression cassette which directs the expression of a second nucleic acid molecule, comprising a nucleic acid sequence which encodes, in order, an alphavirus capsid, a signal peptide, and an alphavirus E2 glycoprotein, with the proviso that the second nucleic acid molecule does not encode an alphavirus E1 glycoprotein. Within various embodiments the signal peptide may be an alphavirus signal peptide (*e.g.* an alphavirus E3 peptide or an alphavirus 6k peptide), or, a non-alphavirus signal peptide (*e.g.*, a tissue plasminogen activator signal peptide). Within further embodiments, the signal peptide of the first or second expression cassette is a first signal peptide, and that cassette further comprises an additional, second signal peptide (which may be of alphavirus or non-alphavirus origin, as discussed above).

Within further aspects of the invention, methods of producing alphavirus particles are provided herein, comprising the step of introducing into a packaging cell line as described herein a vector selected from the group consisting of alphavirus vector constructs, RNA vector replicons, eukaryotic layered vector initiation systems (*e.g.*, U.S. Patent No. 5,814,482), and alphavirus vector particles, such that recombinant alphavirus vector particles are produced.

Within another aspect of the invention methods of producing alphavirus vector particles are provided, comprising introducing into a host cell (a) a first expression cassette which directs the expression of a first nucleic acid molecule, comprising a nucleic acid sequence which encodes, in order, an alphavirus capsid, a signal peptide, and an alphavirus E1 glycoprotein, with the proviso that the first nucleic acid molecule does not encode an alphavirus E2 glycoprotein; and (b) a second expression cassette which directs the expression of a second nucleic acid molecule,

comprising a nucleic acid sequence which encodes, in order, an alphavirus capsid, a signal peptide, and an alphavirus E2 glycoprotein, with the proviso that the second nucleic acid molecule does not encode an alphavirus E1 glycoprotein; and (c) a vector selected from the group consisting of alphavirus vector constructs, RNA vector
5 replicons, eukaryotic layered vector initiation systems, and alphavirus vector particles; such that alphavirus vector particles are produced.

Within various embodiments, methods of producing recombinant alphavirus particles are provided comprising introducing into a population of cells (a) an alphavirus vector construct, RNA vector replicon, eukaryotic layered vector
10 initiation system, or, recombinant alphavirus particle, (b) one or more alphavirus structural protein expression cassettes, and (c) an RNA sequence complementary to an alphavirus junction region promoter, or alphavirus subgenomic RNA. In preferred embodiments, the complementary RNA sequence is introduced as an expression cassette that transcribes within the population of cells the complementary RNA
15 sequence.

Within other aspects of the invention, expression cassettes are provided comprising a promoter which is operably linked to a nucleic acid molecule, which when transcribed produces an RNA sequence complementary to an alphavirus junction region promoter, or, alphavirus subgenomic RNA, wherein said nucleic acid molecule is less
20 than 500, 250, or, 100 nucleotides in length. Also provided in other aspects are host cells which contain such expression cassettes, and alphavirus packaging cell lines which contain such expression cassettes.

In another aspect of the invention, methods of packaging recombinant alphavirus vector particles are provided, comprising introducing into a population of
25 cells (a) an alphavirus vector construct or vector RNA replicon, (b) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E2, and (c) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E1, such that recombinant alphavirus particles are produced. In preferred embodiments, the alphavirus structural protein expression
30 cassettes encoding capsid and glycoprotein E2 or E1 are defective helper RNA

transcribed in vitro or in vivo. In particularly preferred embodiments, the cells are alphavirus packaging cells.

In yet another aspect of the invention, preparations of alphavirus vector particles free from contaminating replication-competent virus are provided, wherein the
5 vector particles are obtained by the method comprising introducing into a population of cells (a) an alphavirus vector construct or vector RNA replicon, (b) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E2, and (c) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E1, such that recombinant alphavirus particles are
10 produced. In preferred embodiments, the alphavirus vector particle preparations are obtained by said method wherein the alphavirus structural protein expression cassettes encoding capsid and glycoprotein E2 and E1 are defective helper RNA transcribed in vitro or in vivo. In particularly preferred embodiments, the alphavirus vector particle preparations are obtained by said method wherein the cells are alphavirus packaging
15 cells.

Within other aspects of the invention RNA vector replicons are provided, comprising: a 5' sequence which initiates transcription of alphavirus RNA; a nucleic acid sequence that codes for biologically active alphavirus nonstructural proteins; an alphavirus subgenomic junction region promoter; a non-alphavirus nucleotide sequence
20 which, when bound by a ligand reduces transcription of subgenomic RNA or translation of a heterologous gene of interest (*i.e.*, "transgene") encoded by the subgenomic RNA; a heterologous gene of interest; and a 3' alphavirus RNA polymerase recognition sequence.

Within other aspects, RNA vector replicons are provided comprising a 5'
25 sequence which initiates transcription of alphavirus RNA; a nucleic acid sequence that codes for biologically active alphavirus nonstructural proteins from a first alphavirus; an alphavirus subgenomic junction region promoter; an alphavirus nucleotide sequence which, when bound by a ligand reduces transcription of subgenomic RNA or translation of a heterologous gene of interest encoded by the subgenomic RNA, wherein said
30 alphavirus nucleotide sequence is from a second alphavirus different from said first

alphavirus; a heterologous gene of interest; and a 3' alphavirus RNA polymerase recognition sequence.

Within another aspect of the invention RNA vectors are provided comprising a 5' sequence which initiates transcription of alphavirus RNA; an alphavirus
5 subgenomic junction region promoter, a non-alphavirus nucleotide sequence which, when bound by a ligand reduces transcription of subgenomic RNA or translation of a heterologous gene of interest encoded by the subgenomic RNA, a heterologous gene of interest; and a 3' alphavirus RNA polymerase recognition sequence, wherein said RNA vector does not encode all biologically active alphavirus nonstructural proteins.

10 Within yet another aspect of the present invention RNA vectors are provided comprising a 5' sequence which initiates transcription of alphavirus RNA; an alphavirus subgenomic junction region promoter, an alphavirus nucleotide sequence which, when bound by a ligand reduces transcription of subgenomic RNA or translation
15 of a heterologous gene of interest encoded by the subgenomic RNA. a heterologous gene of interest and a 3' alphavirus RNA polymerase recognition sequence, wherein said 5' sequence which initiates transcription and said 3' alphavirus RNA polymerase recognition sequence are from a first alphavirus, and the alphavirus nucleotide sequence is from a second alphavirus different from said first alphavirus, and wherein said RNA vector does not encode all biologically active alphavirus nonstructural proteins.

20 Within preferred embodiments of the invention, the above-noted RNA vector replicons and RNA vectors are comprised of elements that are ordered in the manner set forth above.

A wide variety of non-alphavirus (*i.e.*, not obtained or derived from an alphavirus) nucleotide sequences may be specifically selected and utilized within the
25 context of the present invention. For example, within certain embodiments of the above aspects, the non-alphavirus nucleotide sequence is a binding site for an RNA binding protein, such as, for example an R17 coat binding protein, or comprises a sequence as shown in Figure 7 (*i.e.*, 5TOP or TOP).. Within other embodiments the non-alphavirus nucleotide sequence is a binding site for an antibiotic (*e.g.*, tobramycin), or a binding
30 site for Hoechst dyes (*e.g.*, H33258 or H33342).

Within further embodiments the alphavirus nucleotide sequence is a sequence from a subgenomic 5' end nontranslated region of VEE.

Within related embodiments of the invention, the ligand may be an RNA binding protein (*e.g.*, R17 coat protein), an antisense sequence, a dye (*e.g.*, Hoechst
5 dyes H33258 or H3342), or an antibiotic.

As utilized within the context of the present invention, a reduction of either transcription of subgenomic RNA, or, a reduction of translation of a heterologous gene of interest (or "transgene") encoded by the subgenomic RNA, should be understood to refer to a statistically significant decrease of either transcription, or,
10 translation respectively, in the presence of the selected ligand. . In particularly preferred embodiments, the level either transcription of subgenomic RNA, or, level of heterologous transgene expression in cells is reduced at least 25%, 50%, 75%, or 90%, or 3-fold, 5-fold, or 10-fold as compared to the level of expression without the presence of the binding ligand. A wide variety of assays may be utilized to assess a reduced level
15 of transcription or translation, including for example, enzymatic assays of a reporter gene, northern blots, metabolic RNA labeling, as well as the assays provided in Example 8.

Within other embodiments of the invention, the alphavirus nucleotide sequence from the second alphavirus comprises a sequence from a subgenomic 5' end
20 non-translated region of an alphavirus. Within various embodiments, the alphavirus nucleotide sequence is positioned downstream from the subgenomic junction region promoter of the RNA vector, or, upstream from the heterologous gene of interest. Within further embodiments, the first alphavirus may be Sindbis virus or Semliki Forest virus, and the second alphavirus can be VEE.

25 Within further aspects of the invention, RNA vector replicons are provided comprising a 5' sequence which initiates transcription of alphavirus RNA, nucleic acid sequences that code for biologically active alphavirus nonstructural proteins, an alphavirus subgenomic junction region promoter, a sequence from a subgenomic 5' end nontranslated region from VEE, a heterologous gene of interest and
30 a 3' alphavirus RNA polymerase recognition sequence. with the proviso that said

nonstructural proteins are not from VEE.

Within a related aspect, RNA vectors are provided comprising a 5' sequence which initiates transcription of alphavirus RNA, an alphavirus subgenomic junction region promoter, a sequence from a subgenomic 5' end nontranslated region
5 from VEE, a heterologous gene of interest and a 3' alphavirus RNA polymerase recognition sequence, wherein the vector does not encode all biologically active alphavirus nonstructural proteins, and wherein said 5' sequence which initiates transcription and said 3' alphavirus RNA polymerase recognition sequence are not from VEE.

10 Within various embodiments of the above, the RNA vectors provided herein may further comprise a polyadenylation tract. Within other embodiments of the invention, the non-alphavirus sequence in the RNA vector or RNA vector replicons provided herein are not internal ribosome entry sites "IRES" or ribosomal readthrough sequences).

15 Within other aspects of the invention, alphavirus vector constructs are provided, comprising a 5' promoter operably linked to a nucleic acid molecule, wherein said nucleic acid molecule is complementary DNA to the RNA vectors described herein. As utilized within this context, it should be understood that an RNA vector refers to an RNA molecule which may be either an RNA vector replicon as described
20 herein, or, an RNA molecule which, unlike the RNA vector replicon lacks all sequences necessary for biologically active alphavirus nonstructural proteins (*e.g.*, is a defective helper construct). Within certain embodiments, the promoter can be a eukaryotic promoter, or, a bacteriophage promoter.

Other aspects of the present invention provide methods of producing
25 packaged alphavirus vector particles wherein the level of expression of a heterologous transgene from an alphavirus vector construct or vector RNA replicon is reduced in cells used for packaging as compared to the level of expression in a target cell. In preferred embodiments, the method of reducing expression of a heterologous transgene from the alphavirus vector is by reducing transcription from the subgenomic junction
30 region promoter or reducing translation from the subgenomic mRNA. In particularly

preferred embodiments, the level of heterologous transgene expression in cells used for packaging is reduced at least 25%, 50%, 75%, or 90%, or 3-fold, 5-fold, or 10-fold as compared to the level of expression in a target cell.

In one embodiment, a method of producing alphavirus vector particles is provided, comprising introducing into a population of cells (a) an alphavirus vector construct, RNA vector replicon, eukaryotic layered vector initiation system, or, recombinant alphavirus particle, (b) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E2, and (c) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E1, such that recombinant alphavirus particles are produced, wherein the cells contain a complementary RNA sequence (*e.g.*, antisense) of the present invention, the sequence being complementary to the junction region promoter or subgenomic RNA of the alphavirus vector RNA. In preferred embodiments, the complementary or antisense RNA sequence is expressed from a cassette within the cells and the cells are stably transformed with said cassette that expresses an antisense RNA. In another embodiment, cells are provided wherein the cells contain a cassette that expresses an RNA sequence complementary to (*e.g.*, antisense) the junction region promoter or subgenomic RNA of an alphavirus vector, with the proviso that the cells are permissive for alphavirus vector RNA replication.

In another embodiment, a method of producing alphavirus vector particles is provided, comprising introducing into a population of cells (a) an alphavirus vector construct, vector RNA replicon, eukaryotic layered vector initiation system or recombinant alphavirus particle (b) an alphavirus structural protein expression cassette encoding capsid but not glycoproteins E2 or E1, and (c) an alphavirus structural protein expression cassette encoding glycoproteins E2 and E1 but not capsid, such that recombinant alphavirus particles are produced, wherein the cells contain a complementary RNA sequence (*e.g.*, antisense) of the present invention, said sequence being complementary to the junction region promoter or subgenomic RNA of the alphavirus vector RNA. In preferred embodiments, the complementary or antisense RNA sequence is expressed from a cassette within the cells and the cells are stably

transformed with said cassette that expresses an antisense RNA. In another embodiment, cells are provided wherein the cells contain a cassette that expresses an RNA sequence complementary to (*e.g.*, antisense) the junction region promoter or subgenomic RNA of an alphavirus vector, with the proviso that the cells are permissive
5 for alphavirus vector RNA replication.

In a further embodiment, a method of producing alphavirus vector particles is provided, comprising introducing into a population of cells (a) an alphavirus vector construct, RNA vector replicon, eukaryotic layered vector initiation system, or recombinant alphavirus particle, (b) an alphavirus structural protein expression cassette
10 described herein encoding capsid and glycoprotein E2, and (c) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E1, such that recombinant alphavirus particles are produced, wherein the alphavirus vector construct, RNA vector replicon, eukaryotic layered vector initiation system, or alphavirus particle further comprises an RNA aptamer sequence that binds to a ligand
15 provided, therein reducing translation from the subgenomic RNA, said RNA aptamer being inserted within the vector sequence corresponding to said subgenomic RNA. As utilized herein, an RNA aptamer should be understood to refer to a non-alphavirus nucleotide sequence that is one member of a ligand binding pair that reduces or prevents translation of the alphavirus vector subgenomic RNA, as compared to wild-type
20 alphavirus, in the presence of a ligand.

In another embodiment, a method of producing alphavirus vector particles is provided, comprising introducing into a population of cells (a) an alphavirus vector construct, RNA vector replicon, eukaryotic layered vector initiation system, or alphavirus particle, (b) an alphavirus structural protein expression cassette encoding
25 capsid but not glycoproteins E2 and E1, and (c) an alphavirus structural protein expression cassette encoding glycoproteins E2 and E1 but not capsid, such that recombinant alphavirus particles are produced, wherein the alphavirus vector construct , RNA vector replicon, eukaryotic layered vector initiation system, or alphavirus particle further comprises an RNA aptamer sequence that binds to a ligand provided, therein
30 reducing translation from the subgenomic RNA, said RNA aptamer being inserted

within the vector sequence corresponding to said subgenomic RNA. As utilized herein, an RNA aptamer should be understood to refer to a non-alphavirus nucleotide sequence that is one member of a ligand binding pair that reduces or prevents translation of the alphavirus vector subgenomic RNA, as compared to wild-type alphavirus, in the
5 presence of a ligand.

In preferred embodiments, the RNA aptamer within the alphavirus vector construct or vector RNA replicon binds to a ligand that is an antibiotic. In particularly preferred embodiments, the RNA aptamer binds to Tobramycin or Kanamycin.

In another preferred embodiment, the RNA aptamer within the
10 alphavirus vector construct or vector RNA replicon binds to Hoechst dyes H33258 or H33342.

Also provided within the present invention are alphavirus vector constructs, RNA vector replicons, eukaryotic layered vector initiation systems, and alphavirus particles which further comprise an RNA aptamer sequence that binds to a
15 ligand, therein reducing translation from the vector subgenomic RNA, said RNA aptamer being inserted within the vector sequence corresponding to said subgenomic RNA. In preferred embodiments the RNA aptamer binds to an antibiotic or Hoechst dyes H33258 or H33342.

Within further embodiments of the invention, methods are provided for
20 suppressing expression of a transgene (or desired heterologous sequence) during the generation of recombinant alphavirus particles from a packaging or producer cell line, by culturing the packaging or producer cell line with the second member of the ligand binding pair (the ligand) which binds the RNA aptamer, thus preventing or reducing expression of the transgene (or desired heterologous sequence). In one embodiment, the
25 ligand may be an antisense RNA molecule.

Within yet other aspects of the invention, methods are for reducing transcription of subgenomic RNA or translation of a heterologous gene of interest encoded by subgenomic RNA of an alphavirus RNA vector replicon or alphavirus vector construct, comprising the steps of (a) introducing an RNA vector or alphavirus
30 vector construct as described herein; and (b) introducing into a cell an expression

cassette which directs the expression of a ligand that reduces transcription of subgenomic RNA or translation of a heterologous gene of interest encoded by the subgenomic RNA according to the present invention, such that transcription of subgenomic RNA or translation of a heterologous gene of interest encoded by
5 subgenomic RNA is reduced. Within various embodiments, the ligand is an R17 coat protein, or, an antisense sequence.

These and other aspects of the present invention will become evident upon reference to the following detailed description and attached drawings. In addition, various references are identified below which describe in more detail various
10 compositions (*e.g.*, plasmids or cell lines) and methods, and therefore are herein expressly incorporated by reference in their entirety.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic illustration of the expression and processing of a wild-type alphavirus structural polyprotein.

15 Figure 2 is a schematic illustration of selected representative alphavirus structural polyproteins suitable for use in vector replicon RNA packaging.

Figure 3 is a schematic illustration of representative expression cassettes used for expression of alphavirus structural polyproteins of the present invention and packaging of vector replicon RNA.

20 Figure 4 is a graph demonstrating alphavirus vector packaging using representative structural polyprotein cassettes of the present invention.

Figure 5 is a schematic illustration of two representative methods for reducing heterologous transgene expression from alphavirus vectors using antisense RNA.

25 Figure 6 is a schematic illustration of a method of reducing heterologous transgene expression from alphavirus vectors using a specific inserted nucleotide sequence and supplied corresponding ligand.

Figure 7 is an illustration of sequence modifications (SEQ ID NOs: 41, 50 and 51) to an alphavirus vector that allow for binding of a bacteriophage protein.

Figure 8 is a graph demonstrating reduced expression of a heterologous transgene by an RNA binding protein specific for the sequence modifications in Figure 7.

DETAILED DESCRIPTION OF THE INVENTION

5

DEFINITIONS

The following terms are used throughout the specification. Unless otherwise indicated, these terms are defined as follows:

10 An “isolated nucleic acid molecule” is a nucleic acid molecule that is not integrated into the genomic DNA of an organism, or, in the case of a virus, is separated from the complete virus genome. One example of an isolated nucleic acid molecule is a chemically-synthesized nucleic acid molecule, or, a nucleic acid molecule that is produced by recombinant (*e.g.*, PCR) techniques.

15 “Genomic RNA” refers to an RNA that contains all of the genetic information required to direct its own amplification or self-replication *in vivo*, within a target cell. An alphavirus-derived genomic RNA molecule should contain the following ordered elements: 5' viral or defective-interfering RNA sequence(s) required in *cis* for replication, sequences which, when expressed, code for biologically active alphavirus nonstructural proteins (*e.g.*, nsP1, nsP2, nsP3, nsP4), 3' viral sequences required in *cis* for replication, and a polyadenylate tract. The alphavirus-derived genomic RNA vector replicon may also contain a viral subgenomic “junction region” promoter and sequences which, when expressed, code for biologically active alphavirus structural proteins (*e.g.*, C, E3, E2, 6K, E1). Generally, the term genomic RNA refers to a molecule of positive
20 polarity, or “message” sense, and the genomic RNA may be of length different from
25 that of any known, naturally-occurring alphavirus.

“Subgenomic RNA” refers to an RNA molecule of a length or size which is smaller than the genomic RNA from which it was derived. Subgenomic RNA is transcribed from an internal promoter whose sequences reside within the genomic RNA

or its complement. In preferred embodiments, the subgenomic RNA is produced from an alphavirus vector construct, RNA vector replicon, or defective helper construct and encodes one or more alphavirus structural proteins or other heterologous sequences of interest. Generally, the subgenomic RNA resembles a typical mRNA with 5' and 3' end non-translated regions and a protein encoding open reading frame.

"Alphavirus vector construct" refers to an assembly which is capable of directing the expression of a sequence or gene of interest. Such vector constructs are comprised of a 5' sequence which is capable of initiating transcription of an alphavirus RNA (also referred to as 5' CSE, or, 5' sequence which is capable of initiating transcription of an alphavirus RNA, in the background), as well as sequences which, when expressed, code for biologically active alphavirus nonstructural proteins (*e.g.*, nsP1, nsP2, nsP3, nsP4), an alphavirus RNA polymerase recognition sequence (also referred to as 3' CSE, or, alphavirus RNA polymerase recognition sequence, in the background), and, optionally a polyadenylate tract. In addition, the vector construct may include a viral subgenomic "junction region" promoter, sequences from one or more structural protein genes or portions thereof, extraneous nucleic acid molecule(s) which are of a size sufficient to allow production of viable virus, a 5' promoter which is capable of initiating the synthesis of viral RNA from cDNA *in vitro* or *in vivo*, a heterologous sequence to be expressed, and one or more restriction sites for insertion of heterologous sequences.

"Alphavirus RNA vector replicon", "RNA vector replicon" and "replicon" refers to an RNA molecule which is capable of directing its own amplification or self-replication *in vivo*, within a target cell. An alphavirus-derived RNA vector replicon should contain the following ordered elements: 5' viral sequences required in *cis* for replication (also referred to as 5' CSE, in background), sequences which, when expressed, code for biologically active alphavirus nonstructural proteins (*e.g.*, nsP1, nsP2, nsP3, nsP4), 3' viral sequences required in *cis* for replication (also referred to as 3' CSE, in background), and a polyadenylate tract. The alphavirus-derived RNA vector replicon may also contain a viral subgenomic "junction region" promoter, sequences from one or more structural protein genes or portions thereof,

extraneous nucleic acid molecule(s) which are of a size sufficient to allow production of recombinant alphavirus particles, as well as heterologous sequence(s) to be expressed.

“Recombinant Alphavirus Particle” refers to a virion-like structural unit containing an alphavirus RNA vector replicon. Generally, a recombinant alphavirus particle comprises one or more alphavirus structural proteins, a lipid envelope and an RNA vector replicon. Preferably, the recombinant alphavirus particle contains a nucleocapsid structure that is contained within a host cell-derived lipid bilayer, such as a plasma membrane, in which alphaviral-encoded envelope glycoproteins are embedded. The particle may also contain other components (*e.g.*, targeting elements, other viral structural proteins, or other receptor binding ligands) which direct the tropism of the particle from which the alphavirus was derived.

“Structural protein expression cassette” refers to a nucleic acid molecule that is capable of directing the synthesis of one or more alphavirus structural proteins. The expression cassette should include a 5' promoter which is capable of initiating the synthesis of RNA from cDNA, as well as sequences which, when expressed, code for one or more biologically active alphavirus structural proteins (*e.g.*, C, E3, E2, 6K, E1), and a 3' sequence which controls transcription termination. The expression cassette also may include a 5' sequence which is capable of initiating transcription of an alphavirus RNA (also referred to as 5' CSE, in background), a viral subgenomic “junction region” promoter, an alphavirus RNA polymerase recognition sequence (also referred to as 3' CSE, in background), and a polyadenylate tract. In certain embodiments, the expression cassette also may include splice recognition sequences, a catalytic ribozyme processing sequence, a sequence encoding a selectable marker, or a nuclear export signal. In addition, expression of the alphavirus structural protein(s) may, in certain embodiments, be regulated by the use of an inducible promoter.

“Stable Transformation” refers to the introduction of a nucleic acid molecule into a living cell, and long-term or permanent maintenance of that nucleic acid molecule in progeny cells through successive cycles of cell division. The nucleic acid molecule may be maintained in any cellular compartment, including, but not limited to, the nucleus, mitochondria, or cytoplasm. In preferred embodiments, the nucleic acid

molecule is maintained in the nucleus. Maintenance may be intrachromosomal (integrated) or extrachromosomal, as an episomal event.

“Alphavirus packaging cell line” refers to a cell which contains an alphavirus structural protein expression cassette and which produces recombinant
5 alphavirus particles after introduction of an alphavirus vector construct, RNA vector replicon, eukaryotic layered vector initiation system (*e.g.*, U.S. Patent No. 5,814,482), or recombinant alphavirus particle. The parental cell may be of mammalian or non-mammalian origin. Within preferred embodiments, the packaging cell line is stably transformed with the structural protein expression cassette.

10 “Defective helper construct” refers to an assembly which is capable of RNA amplification or replication, and expression of one or more alphavirus structural proteins in response to biologically active alphavirus nonstructural proteins supplied in *trans*. The defective helper construct should contain the following ordered elements: 5' viral or defective-interfering RNA sequences required in *cis* for replication (also
15 referred to as 5' CSE, in background), a viral subgenomic junction region promoter, sequences which, when expressed, code for one or more biologically active alphavirus structural proteins (*e.g.*, C, E3, E2, 6K, E1), 3' viral sequences required in *cis* for replication (also referred to as 3' CSE, in background), and a polyadenylate tract. The defective helper construct may also contain a 5' promoter which is capable of initiating
20 the synthesis of viral RNA from cDNA in vitro or in vivo, a 3' sequence which controls transcription termination, splice recognition sequences, a catalytic ribozyme processing sequence, a sequence encoding a selectable marker, and a nuclear export signal. A defective helper construct should not encode all four functional alphavirus nonstructural proteins.

25 “Eukaryotic Layered Vector Initiation System” refers to an assembly which is capable of directing the expression of a sequence(s) or gene(s) of interest. The eukaryotic layered vector initiation system should contain a 5' promoter which is capable of initiating *in vivo* (*i.e.*, within a cell) the synthesis of RNA from cDNA, and a viral vector sequence which is capable of directing its own replication in a eukaryotic
30 cell and also expressing a heterologous sequence. In preferred embodiments, the

nucleic acid vector sequence is an alphavirus-derived sequence and is comprised of a 5' sequence which is capable of initiating transcription of an alphavirus RNA (also referred to as 5' CSE, in background), as well as sequences which, when expressed, code for biologically active alphavirus nonstructural proteins (*e.g.*, nsP1, nsP2, nsP3, nsP4), and an alphavirus RNA polymerase recognition sequence (also referred to as 3' CSE, in background). In addition, the vector sequence may include a viral subgenomic “junction region” promoter, sequences from one or more structural protein genes or portions thereof, extraneous nucleic acid molecule(s) which are of a size sufficient to allow optimal amplification, a heterologous sequence to be expressed, one or more restriction sites for insertion of heterologous sequences, as well as a polyadenylation sequence. The eukaryotic layered vector initiation system may also contain splice recognition sequences, a catalytic ribozyme processing sequence, a nuclear export signal, and a transcription termination sequence. (See U.S. Patent No. 5,814,482 or PCT Publication No. WO 97/38087.)

15 “Polyprotein” refers to a single polypeptide comprising two or more proteins, wherein the polypeptide is processed by proteases into the individual proteins during or after translation of the polypeptide. The polyprotein should be encoded by one open-reading frame (ORF) within an mRNA molecule, and may be naturally occurring or generated using recombinant DNA techniques by combining separate genes (*e.g.*, removing translation initiator or terminator codons as necessary) and inserting appropriate signals for processing of the polyprotein into individual protein species.

25 “Leader/signal peptide” refers to a peptide sequence contained within a polypeptide, which is necessary for transport of the polypeptide into / across the endoplasmic reticulum (ER) membrane. In preferred embodiments, the leader/signal peptide is located at the amino-terminus of the polypeptide and is cleaved from the remainder of the polypeptide (*e.g.*, by signal peptidase) after its function for ER transport has been performed.

As discussed in more detail below, the present invention provides: (A) sources of wild-type alphaviruses suitable for constructing the nucleic acid sequences, expression cassettes, polypeptides and vectors; (B) alphavirus vector constructs and alphavirus RNA vector replicons; (C) alphavirus structural polyprotein genes; (D) expression cassettes containing alphavirus structural polyprotein genes; (E) alphavirus packaging cell lines; (F) methods of packaging recombinant alphavirus particles; (G) methods of suppressing heterologous gene expression during vector packaging; and (H) heterologous sequences (also referred to as "transgenes") which may be expressed within the context of the present invention.

10 A. Sources of Alphaviruses

As noted above, the present invention provides a wide variety of alphavirus-derived nucleic acid sequences, expression cassettes, polypeptides, and vectors, as well as methods for utilizing such sequences, cassettes, and polypeptides for high level packaging of the vectors into recombinant alphavirus vector particles in the absence of contaminating replication-competent virus. Briefly, sequences encoding alphaviruses suitable for use in preparing the above-described compositions can be readily obtained given the disclosure provided herein from naturally occurring sources, or from depositories (*e.g.*, the American Type Culture Collection, Rockville, Maryland). Representative examples of suitable alphaviruses are described in more detail in U.S. Patent No. 5,843,723 and PCT Publication No. WO 97/38087.

B. Alphavirus Vector Constructs and Alphavirus RNA Vector Replicons

As noted above, the present invention provides both DNA and RNA vector constructs which are derived from alphaviruses. Briefly, within one aspect of the present invention alphavirus vector constructs are provided, comprising a 5' promoter which initiates synthesis of viral RNA from cDNA, a 5' sequence which initiates transcription of alphavirus RNA, a nucleic acid sequence encoding all four alphaviral nonstructural proteins, an alphavirus RNA polymerase recognition sequence and a 3' polyadenylate tract. Within other aspects, RNA vector replicons are provided,

comprising a 5' sequence which initiates transcription of alphavirus RNA, a nucleic acid sequence which encodes all four alphaviral nonstructural proteins, an alphavirus RNA polymerase recognition sequence and a 3' polyadenylate tract. Within preferred embodiments of the above, the above constructs further comprise a viral junction
5 region. Each of these aspects is discussed in more detail below (see also Strauss and Strauss, *Microbiol. Rev.* 58(3):491-562, 1994).

1. 5' Promoters which initiate synthesis of viral RNA

As noted above, within certain embodiments of the invention, alphavirus vector constructs and defective helper constructs are provided which contain 5'
10 promoters (*e.g.*, DNA dependent RNA polymerase promoters) that initiate synthesis of viral RNA from cDNA by a process of *in vitro* transcription. Within preferred embodiments such promoters include, for example, the bacteriophage T7, T3, and SP6 RNA polymerase promoters. Similarly, alphavirus vector constructs and defective helper constructs are provided which contain 5' promoters (*e.g.*, DNA dependent RNA
15 polymerase promoters) that initiate synthesis of viral RNA from cDNA *in vivo* (*i.e.*, within a cell). Within certain embodiments, such RNA polymerase promoters may be derived from both prokaryotic and eukaryotic organisms, and include, for example, the bacterial β -galactosidase and *trpE* promoters, and the eukaryotic viral simian virus 40 (SV40) (*e.g.*, early or late), cytomegalovirus (CMV) (*e.g.*, immediate early), Moloney
20 murine leukemia virus (MoMLV) or Rous sarcoma virus (RSV) LTR, and herpes simplex virus (HSV) (thymidine kinase) promoters.

2. Sequences Which Initiate Transcription

As noted above, within preferred embodiments, the alphavirus vector constructs, RNA vector replicons and defective helper constructs of the present
25 invention contain a 5' sequence which is capable of initiating transcription of an alphavirus RNA (also referred to as 5'-end CSE, or 5' *cis* replication sequence). Representative examples of such sequences include nucleotides 1-60, and to a lesser extent nucleotides through bases 150-210, of the Sindbis virus, nucleotides 10-75 for tRNA^{Asp} (aspartic acid, U.S. Patent 5,091,309), and 5' sequences from other
30 alphaviruses which initiate transcription. It is the complement of these sequences,

3. Alphavirus Nonstructural Proteins

Alphaviruses express four nonstructural proteins, designated nsP1, nsP2, nsP3, and nsP4. Vectors of the present invention derived from alphaviruses should contain sequences encoding the four nonstructural proteins. In wild-type Sindbis virus, nonstructural proteins 1-3 are encoded by nucleotides 60 to 5747, while nsP4 is encoded by nucleotides 5769 to 7598. The nonstructural proteins are translated from the genomic positive strand RNA as one of two large polyproteins, known as P123 or P1234, respectively, depending upon (i) whether there is an opal termination codon between the coding regions of nsP3 and nsP4 and (ii) if there is such an opal codon present, whether there is translation termination of the nascent polypeptide at that point

or readthrough and hence production of P1234. The opal termination codon is present at the nsP3/nsP4 junction of the alphaviruses SIN, AURA, WEE, EEE, VEE, and RR, and thus the P123 and P1234 species are expressed in cells infected with these viruses. In contrast, no termination codon is present at the nsP3/nsP4 junction of the
5 alphaviruses SF and ONN, and thus only the P1234 species is expressed in cells infected with these viruses. Both the polyprotein and processed monomeric forms of the nonstructural proteins function in the replication of the alphavirus RNA genome. Translational readthrough generally occurs about 10%-20% of the time in cells infected with wild type Sindbis virus containing the opal termination codon at the nsP3/nsP4
10 junction. Processing of P123 and P1234 is by a proteinase activity encoded by the one of the nonstructural proteins, and is discussed further below. Each nonstructural protein has several functions, some of which are discussed in more detail within U.S. Patent No. 5,843,723 and PCT Publication No. WO 97/38087.

4. Viral Junction Regions

15 The alphavirus viral junction region promoter normally controls transcription initiation of the subgenomic mRNA. Thus, this element also is referred to as the subgenomic mRNA promoter. In the case of Sindbis virus, the normal viral junction region typically begins at approximately nucleotide number 7579 and continues through at least nucleotide number 7612 (and possibly beyond). At a
20 minimum, nucleotides 7579 to 7602 (5'- ATC TCT ACG GTG GTC CTA AAT AGT - SEQ. ID NO. 1) are believed necessary for transcription of the subgenomic fragment. This region (nucleotides 7579 to 7602) is hereinafter referred to as the "minimal junction region core." It is the complement of these junction region sequences, present in alphavirus minus strand RNA, that has functional promoter activity and allows for
25 nsP-mediated transcription of subgenomic mRNA.

5. Alphavirus RNA Polymerase Recognition Sequence, Poly(A) Tract

As noted above, alphavirus vector constructs or RNA vector replicons of the present invention also should include an alphavirus RNA polymerase recognition
30 sequence (also termed "alphavirus replicase recognition sequence", "3' terminal CSE",

or "3' *cis* replication sequence"). Briefly, the alphavirus RNA polymerase recognition sequence, which is located at the 3' end region of positive stranded genomic RNA, provides a recognition site at which the virus begins replication by synthesis of the negative strand. A wide variety of sequences may be utilized as an alphavirus RNA
5 polymerase recognition sequence. For example, within one embodiment, Sindbis virus vector constructs in which the polymerase recognition is truncated to the smallest region that can still function as a recognition sequence (*e.g.*, nucleotides 11,684 to 11,703) can be utilized. Within another embodiment of the invention, Sindbis virus vector constructs in which the entire nontranslated region downstream from the glycoprotein
10 E1 gene to the 3' end of the viral genome including the polymerase recognition site (*e.g.*, nucleotides 11,382 to 11,703), can be utilized.

Within preferred embodiments of the invention, the alphavirus vector construct or RNA vector replicon may additionally contain a poly(A) tract. Briefly, the poly(A) tract may be of any size which is sufficient to promote recognition of the
15 alphavirus 3' *cis* replication sequence by nonstructural proteins and stability in the cytoplasm, thereby increasing the efficiency of initiating the viral life cycle. Within various embodiments of the invention, the poly(A) tract comprises at least 25 adenosine nucleotides, and most preferably, at least 40 adenosine nucleotides. Within one embodiment, the poly(A) sequence is attached directly to Sindbis virus nucleotide
20 11,703.

C. Alphavirus Structural Polyprotein Genes

Wild-type alphavirus virions consist of four elements: a single-stranded genome RNA, repeating units of a capsid (C) protein monomer which interacts with the RNA to form an icosahedral nucleocapsid, a host-derived lipid bilayer (envelope), and
25 two viral-encoded envelope glycoproteins, E1 and E2, which protrude as heterodimeric spikes. In addition, some alphavirus virions also contain another protein, E3. The alphavirus structural proteins are synthesized during virus replication as a single polyprotein from subgenomic RNA. The polyprotein (shown in Figure 1), comprises the ordered elements C, E3, E2, 6K, E1, and proteolytic processing gives rise to the

final protein products. E3 and 6K serve as signal peptides for insertion of glycoproteins E2 and E1, respectively, into the endoplasmic reticulum.

Recombinant alphavirus vector particles are packaged using these same structural proteins as wild-type virions. In general, alphavirus vector RNA packaging
5 has been performed by expressing in cells an intact, wild-type structural polyprotein which maintains its ability to be processed into the individual protein species by normal mechanisms (Figure 2). More recently, it was shown that the capsid protein and envelope glycoproteins could be expressed separately from separate cassettes and still result in functional packaging of vector (U.S. Patent 5,789,245). In this configuration, a
10 polyprotein comprising E3-E2-6K-E1, which maintains the wild-type virus gene order and processing, was utilized. The present invention provides alphavirus structural polyproteins significantly different from those previously described, and makes use of altered gene order and both alphavirus and non-alphavirus derived leader/signal peptides for novel polyproteins (*e.g.*, C-L-E1, C-6K-E1, C-E3-6K-E1, C-E3-E1, C-E3-
15 E2, *see* Figure 2). As discussed above, as well as in the examples below, the novel alphavirus structural polyproteins provide certain advantages when used for packaging vectors.

D. Expression Cassettes Containing Alphavirus Structural Polyprotein Genes

Within one aspect of the invention, a variety of expression cassettes are
20 provided, which contain the sequences coding for and operably express one or more alphavirus structural polypeptides provided herein. Generally, the expression cassettes fall within one of three categories: 1) a DNA promoter of RNA transcription (*e.g.*, RNA polymerase II promoter) directly and operably linked to the structural protein open reading frame (ORF), and a transcription termination/polyadenylation sequence; 2) an
25 alphavirus defective helper RNA transcribed *in vitro* or *in vivo*, comprising the ordered elements 5' viral or defective-interfering RNA sequence required *in cis* for alphaviral replication (also referred to as 5' CSE, in background), viral subgenomic junction region promoter, alphavirus structural protein sequence of the present invention, 3' alphaviral sequence required *in cis* for replication (also referred to as 3' CSE, in background), and

polyadenylate tract; and 3) DNA cassettes comprising the ordered elements of a DNA promoter of RNA transcription that functions within a eukaryotic cell (*e.g.*, RNA polymerase II promoter) and is operably linked to a 5' viral or defective-interfering RNA sequence required in *cis* for alphaviral replication, viral subgenomic junction region promoter, alphavirus structural protein sequence of the present invention. 3' alphaviral sequence required in *cis* for replication, polyadenylate tract, and transcription termination/polyadenylation sequence. In preferred embodiments, the structural proteins of the present invention are synthesized at high levels by the cassettes only after induction by the RNA vector replicon itself or some other provided stimulus.

10 E. Alphavirus Packaging Cell Lines

Within further aspects of the invention, alphavirus packaging cell lines are provided. In particular, within one aspect of the present invention, alphavirus packaging cell lines are provided wherein the viral structural proteins are supplied in *trans* from one or more stably transformed expression cassettes, and are able to encapsidate transfected, transduced, or intracellularly produced alphavirus vector RNA transcripts in the cytoplasm and release infectious packaged vector particles through the plasma membrane. In preferred embodiments, the structural proteins necessary for packaging are synthesized at high levels only after induction by the RNA vector replicon itself or some other provided stimulus, and the transcripts encoding these structural proteins are capable of cytoplasmic amplification in a manner that will allow expression levels sufficient to mimic that of a natural viral infection (WO 97/38087 and U.S. Patent 5,789,245). Furthermore, in other embodiments, expression of a selectable marker is operably linked to the structural protein expression cassette. Such a linked selectable marker allows efficient generation of functional, stably transformed packaging cell lines.

F. Methods of Packaging Recombinant Alphavirus Particles

As provided by the invention, generation (packaging) of recombinant alphavirus vector particles may be readily accomplished by, for example, co-

transfection of complementing vector and defective helper (DH) molecules derived from *in vitro* transcribed RNA, or plasmid DNA, or by co-infection with virus (*see* Bredenbeek et al., *J. Virol.* 67:6439-6446, 1993, Dubensky et al., *J. Virol* 70:508-519, 1996 and U.S. Patents 5,814,482, 5,739,026, 5,766,602, 5,789,245 and 5,792,462.

5 Alternatively, vector particles may be generated by introduction of vector RNA into stable alphavirus packaging cell lines (PCL, U.S. Patent 5,789,245). Briefly, such PCL and their stably transformed structural protein expression cassettes can be derived using methods described within U.S. Patent 5,789,245, or using novel methods described within this invention. For example, the production of recombinant alphavirus vector

10 particles by PCL may be accomplished following introduction of alphavirus-based vector molecules into the PCL, the vectors being derived from *in vitro* transcribed RNA, plasmid DNA, or previously obtained recombinant alphavirus particles, incubating the PCL for a under conditions and for a time necessary for vector particle packaging, and harvesting of the packaged vector particles. As shown in the detailed

15 examples provided herein, utilization of the novel alphavirus structural polyproteins of the present invention for efficient vector packaging in such approaches is readily accomplished.

G. Methods of Reducing Heterologous Gene Expression During Packaging

Also provided in the invention are methods to reduce expression of a

20 heterologous sequence (also referred to as a "transgene") from an alphavirus vector during the production of recombinant alphavirus vector particles (*e.g.*, packaging of vector RNA). Such methods are advantageous given the very high level expression of transgene in cells during vector packaging and the often deleterious effects of transgene expression on final vector particle titer due to interference with glycoprotein transport

25 and processing, competition with alphavirus gene product function, and general toxicity for the host or packaging cell. Reduction of transgene expression during vector packaging provides one mechanism to addresses these issues. In preferred embodiments, the level of heterologous transgene expression in cells used for packaging

is reduced at least 25%, 50%, 75%, 95%, or, 3-fold, 5-fold, 10-fold, or even about 100-fold as compared to the level of expression in a target cell.

The approaches provided herein to reduce expression of a transgene in alphavirus vectors are directed at the reduction of transcription or translation of the
5 vector subgenomic RNA. Such approaches are deemed necessary because reduction in the overall ability of the vector itself to replicate would have an adverse effect of the production of alphavirus vector particles.

In one embodiment, a method of producing alphavirus vector particles is provided, comprising introducing into a population of cells (a) an alphavirus vector
10 construct or vector RNA replicon, (b) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E2, and (c) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E1, such that recombinant alphavirus particles are produced, wherein the cells contain a cassette which expresses an RNA sequence complementary to (*e.g.*, antisense) the
15 junction region promoter or subgenomic RNA of the alphavirus vector RNA. The antisense sequence may be operably linked to and expressed directly from a eukaryotic promoter (*e.g.*, RNA polymerase I, II, or III promoter), or it may be expressed from an alphavirus junction region promoter, such as in the various cassettes (*e.g.*, defective helper RNA) provided herein. In those instances where it is expressed from an
20 alphavirus junction region promoter, sufficient sequence differences should be included to prevent binding of the antisense RNA to its own junction region promoter or 5'-end. In addition, antisense RNA molecules of the present invention may be complementary to any sequence of the subgenomic RNA, but preferably the antisense RNA is complementary to a sequence at or near the 5'-nontranslated region or AUG initiation
25 codon of the subgenomic RNA. In preferred embodiments, the cells that contain a cassette that expresses an antisense RNA of the present invention are stably transformed with said cassette.

In another embodiment, a method of producing alphavirus vector particles is provided, comprising introducing into a population of cells (a) an alphavirus
30 vector construct or vector RNA replicon, (b) an alphavirus structural protein expression

cassette described herein encoding capsid and glycoprotein E2, and (c) an alphavirus structural protein expression cassette described herein encoding capsid and glycoprotein E1, such that recombinant alphavirus particles are produced, wherein the alphavirus vector construct or vector RNA replicon further comprises an RNA aptamer sequence
5 that binds to a ligand provided, therein reducing translation from the subgenomic RNA, said RNA aptamer being inserted within the vector sequence corresponding to said subgenomic RNA. The RNA aptamer sequence used in methods of the present invention is not limited to those explicitly described herein. Rather, any RNA aptamer with a known binding ligand may be readily substituted by one of skill in the art, using
10 this disclosure. In utilizing an RNA aptamer and ligand binding pair, it is preferable to choose an aptamer and site for insertion into the vector that does not interfere with replication of the alphavirus vector or with translation of the subgenomic mRNA in the absence of its ligand. In preferred embodiments, the RNA aptamer within the alphavirus vector construct or vector RNA replicon binds to a ligand that is an
15 antibiotic, such as Tobramycin or Kanamycin. In another preferred embodiment, the RNA aptamer within the alphavirus vector construct or vector RNA replicon binds to Hoechst dyes H33258 or H33342 (Werstuck and Green, 1998, *Science* 282:296-298). In particularly preferred embodiments, the RNA aptamer is inserted within the 5'-nontranslated region of the alphavirus subgenomic RNA.

20 H. Heterologous Sequences

As briefly noted above, a wide variety of heterologous sequences may be included within the alphavirus vectors described herein including, for example, sequences which encode palliatives such as lymphokines or cytokines, toxins, and prodrug converting enzyme, sequences which encode antigens that stimulate an immune
25 response, ribozymes or antisense sequences, sequences which encode proteins for therapeutic application such as growth or regulatory factors, and sequences which encode proteins that assist or inhibit an immune response.

Preferably, the nucleotide sequences should be of a size sufficient to allow efficient production of viable vector particles. Within the context of the present

invention, the production of any measurable titer of recombinant alphavirus particles, for example, by transfer of expression assay, titering cell line assay, reporter assay, or plaque assay on appropriate susceptible monolayers, is considered to be "production of viable vector particles". This may be, at a minimum, an alphavirus vector construct
5 which does not contain any additional heterologous sequence. However, within other embodiments, the vector construct may contain additional heterologous or foreign sequences. Within preferred embodiments, the heterologous sequence can comprise a heterologous sequence of at least about 100 bases, 2 kb, 3.5 kb, 5 kb, 7 kb, or even a heterologous sequence of at least about 8 kb. The above-described heterologous
10 sequences may be readily obtained from a variety of sources, including for example, depositories such as the American Type Culture Collection (ATCC, Rockville, MD), or from commercial sources such as British Bio-Technology Limited (Cowley, Oxford, England). Alternatively, cDNA sequences which encode the above-described heterologous sequences may be obtained from cells which express or contain the
15 sequences, utilizing standard procedures known in the art. In addition, heterologous sequences also may be synthesized, for example, on an Applied Biosystems, Inc. DNA synthesizer.

Representative examples of suitable heterologous sequences are discussed in more detail within U.S. Patent No. 5,843,723 and PCT Publication No.
20 WO 97/38087.

The following examples are included to more fully illustrate the present invention. Additionally, these examples provide preferred embodiments of the invention and are not meant to limit the scope thereof. Standard methods for many of the procedures described in the following examples, or suitable alternative procedures,
25 are provided in widely reorganized manuals of molecular biology, such as, for example, "Molecular Cloning," Second Edition (Sambrook et al., Cold Spring Harbor Laboratory Press, 1987) and "Current Protocols in Molecular Biology" (Ausubel et al., eds. Greene Associates/Wiley Interscience, NY, 1990).

EXAMPLE 1

CONSTRUCTION OF THE C-E3-E2 STRUCTURAL POLYPROTEIN AND CASSETTE

The C-E3-E2 polyprotein and expression cassette comprises the coding regions for an alphavirus capsid protein and glycoprotein E2. As described above, a leader/signal sequence also is included to facilitate insertion of the envelope glycoprotein into the endoplasmic reticulum. As will be evident from the examples provided herein, a variety of alphavirus (*e.g.*, E3, 6k) or non-alphavirus (*e.g.*, TPA) leader/signal sequences may be incorporated as part of the polyprotein open reading frame. For example, one version of the cassette comprises C-E3-E2 and was assembled by PCR amplification to include the insertion of a stop codon after the final amino acid (aa806) of E2. Specifically, PCR was performed with the following primers:

XCAPSIDF:

5'CTC GAG ACC ATG AAT AGA GGA TTC TTT AAC (SEQ. ID NO. 2)

NGLYCOE2R:

15 5'GCG GCC GCT CAA GCA TTG GCC GAC CTA ACG CAG CAC (SEQ. ID NO. 3)

The capsid forward primer (XCAPSIDF) contained a *Xho*I site to facilitate subcloning, a Kozak consensus sequence, and the first twenty one bases of the capsid open reading frame (ORF). The E2 reverse primer (NGLYCOE2R) contained a *Not*I site to facilitate subcloning, a UGA stop codon, and the last 25 bases of the glycoprotein E2 ORF.

PCR amplification was performed using PFU polymerase (Stratagene Inc., San Diego, CA) as per the manufacturer's instructions. The PCR template was the CMV Promoter driven Sindbis virus genomic clone pDCMVSINg (Dubensky et al., 1996, *J. Virol.* 70:508-519). The PCR reaction was cycled 20x (95°C 30 sec, 60°C 30 sec, 72°C 4 min), and the PCR product and corresponding polyprotein referred to as C-E3-E2 (Figure 2).

The 2.3kbp C-E3-E2 PCR product was cloned into the pCR-Blunt vector using the Zero Blunt PCR Cloning Kit (Invitrogen, Inc. Carlsbad, CA) as per the

manufacturer's instructions. Plasmid DNA was isolated and screened for the presence of the PCR insert by digestion with *XhoI* and *NotI* restriction endonucleases. The C-E3-E2 fragment was isolated from pCR-Blunt by digestion with *XhoI* and *NotI*, purification from a 1% agarose gel using the GeneCleanII kit (GENECLEAN, Bio 101 Inc., Vista, CA) as per the manufacturer's instructions, and inserted into appropriate expression cassettes (Figure 3) as described below.

Construction of a C-E3-E2 expression cassette was performed by insertion of the isolated C-E3-E2 fragment into a Sindbis virus derived defective helper configuration, comprising the following ordered elements: a 5' sequence which initiates transcription of alphavirus RNA from viral RNA, a viral subgenomic junction region promoter, an alphavirus polyprotein gene described herein, an alphavirus RNA polymerase recognition sequence, and a polyadenylate tract. In this example, the defective helper is a cDNA contained within a plasmid, wherein the defective helper is operably linked to a bacteriophage (*e.g.*, SP6) promoter for in vitro transcription, and also contains a unique restriction site downstream of the polyadenylate tract for linearization of the plasmid. Specifically, the defective helper plasmid was constructed by modification of the previously described Sindbis virus vector pRSIN-luc (Dubensky et al., 1996, *ibid*). Plasmid pRSIN-luc was digested with *BspEI* to remove most Sindbis virus nonstructural protein gene sequences and the vector was then re-ligated to itself, resulting in a construct referred to as pRSINdl-luc. This plasmid was next digested with *NotI* and *SacI* to remove the minimal Sindbis 3'-end sequence and A₂₅ tract, which were replaced with an approximately 0.4kbp fragment from pKSSIN1-BV (WO 97/38087) containing the complete Sindbis virus 3'-end, an A₄₀ tract and a *PmeI* site for linearization prior to transcription, obtained from pKSSIN1-BV after digestion with *NotI* and *SacI*, and purification from an agarose gel. This new construct was designated SINdl-luc. Plasmid SINdl-luc was digested with *XhoI* and *NotI* to remove the luciferase gene insert and the remaining vector portion was purified from an agarose gel using GENECLEAN. The isolated C-E3-E2 fragment (from above) was then ligated into the vector, resulting in the expression construct SINdl-CE3E2.

EXAMPLE 2

CONSTRUCTION OF THE C-E3-6K-E1 STRUCTURAL POLYPROTEIN AND CASSETTE

This polyprotein and expression cassette comprises the coding regions for an alphavirus capsid protein and glycoprotein E1. As described above, a leader/signal sequence also is included to facilitate insertion of the envelope glycoprotein into the endoplasmic reticulum. For example, one version of the cassette comprises C-E3-6K-E1 (Figure 2) and was assembled by using two separate PCR amplification reactions for C-E3 and 6K-E1. Specifically, a C-E3 PCR product was generated with the following primers:

10 XCAPSIDF:

5'CTC GAG ACC ATG AAT AGA GGA TTC TTT AAC (SEQ. ID NO. 2)

BE3R:

5'GGA TCC GTC GTC AAC GAC GCT TCT TTT GC (SEQ. ID NO. 4)

15 The capsid forward primer (XCAPSIDF) was described above. The E3 reverse primer (BE3R) contains a *Bam*HI site to facilitate subcloning. The PCR reaction and subsequent subcloning into pCR-Blunt was performed as described above, using the Sindbis virus genomic clone pDCMVSING (Dubensky et al., *ibid*) as template. Clones were screened with the restriction endonucleases *Xho*I and *Bam*HI, the plasmid
20 then digested with *Xho*I and *Bam*HI, and the 1kbp C-E3 fragment purified from an agarose gel using GeneCleanII. This fragment was subcloned into *Xho*I/*Bam*HI digested pBluescript-KS plasmid (Stratagene Inc.), resulting in a plasmid called pKS-C-E3.

The 6K-E1 PCR product was generated with the following primers:

25 B6KF:5'GGA TCC AGG TCG GCC AAT GCT GAA ACG TTC ACC GAG ACC ATG AGT
TAC (SEQ. ID NO. 5)

NEIR:

5'GCG GCC GCT CAT CTT CGT GTG CTA GTC AGC ATC (SEQ. ID NO. 6)

The B6KF forward primer contains a *Bam*HI site and the NEIR reverse
5 primer contains a *Not*I site to facilitate subcloning. The PCR reaction and subsequent
subcloning into pCR-Blunt was performed as described above, using the Sindbis virus
genomic clone pDCMVSING (Dubensky et al., *ibid*) as template. Clones were screened
with the restriction endonucleases *Bam*HI and *Not*I, the plasmid then was digested with
*Bam*HI and *Not*I, and the 1.5kbp C-E3 fragment purified from an agarose gel using
10 GeneCleanII. This fragment was subcloned into *Bam*HI/*Not*I digested pKS-C-E3
plasmid, resulting in the new plasmid called pKS-C-E3-6K-E1.

The C-E3-6K-E1 fragment was isolated by digestion with *Xho*I and *Not*I,
purified from a 1% agarose gel using GENE CLEAN, and ligated into SINdl-luc that
also was digested with *Xho*I and *Not*I to remove the luciferase gene insert (as described
15 above). The resulting expression construct was designated SINdl-CE36KE1.

EXAMPLE 3

CONSTRUCTION OF THE C-6K-E1 STRUCTURAL POLYPROTEIN AND CASSETTE

This polyprotein and expression cassette comprises the coding regions
for an alphavirus capsid protein and glycoprotein E1. As described above, a
20 leader/signal sequence also is included to facilitate insertion of the envelope
glycoprotein into the endoplasmic reticulum. For example, one version of the cassette
comprises C-6K-E1 (Figure 2) and was assembled by using two separate PCR
amplification reactions for C-6K and 6K-E1, followed by an overlapping PCR reaction.
The last two codons of E3 are retained to preserve the capsid protease cleavage site.
25 Specifically, a C-6K PCR product was generated with the following primers:

XCAPSIDF:

5'CTC GAG ACC ATG AAT AGA GGA TTC TTT AAC (SEQ. ID NO. 2)

6KE3CR:

5'CTC GGT GAA CGT TTC TGC GGA CCA CTC TTC TGT CCC TTC (SEQ. ID NO. 7)

- 5 The capsid forward primer (XCAPSIDF) was described above. The 6K reverse primer (BE3R) contains the last six codons of capsid, the first two codons of E3, and the first six codons of 6K. The resulting 0.8kbp PCR product contains: 5'XhoI site/capsid ORF/first two codons of E3/first five codons of 6K.

In the other reaction, a 6K-E1 PCR product was generated with the following primers:

CE36KF:

5'GAA GAG TGG TCC GCA GAA ACG TTC ACC GAG ACC ATG AG (SEQ. ID NO. 8)

NE1R:

- 15 5'GCG GCC GCT CAT CTT CGT GTG CTA GTC AGC ATC (SEQ. ID NO. 6)

The 6K forward primer (CE36KF) contains the last three codons of capsid, the first two codons of E3, and the first twenty-three nucleotides of the 6K ORF. The NE1R primer was described previously. The resulting 1.5 kbp PCR product contains: 5' last three codons of capsid/first two codons of E3/6K ORF/E1 ORF/NotI site.

Both PCR reactions were performed as above using pDCMVSING as template. The separate PCR products were purified from an agarose gel using GeneCleanII, and then combined as template for an "overlapping" PCR reaction containing the primers XCAPSIDF and NE1R. The resulting 2.3 kbp C-6K-E1 fragment was then gel purified, subcloned into pCR-Blunt, and screened with *XhoI* and *NotI* to identify the new construct pCR-Blunt-C-6K-E1.

The C-6K-E1 fragment was isolated from pCR-Blunt-C-6K-E1 by digestion with *XhoI* and *NotI*, purified from a 1% agarose gel using GENECLEAN, and ligated into SINDI-luc that also was digested with *XhoI* and *NotI* to remove the

luciferase gene insert (as described above). The resulting expression construct was designated SINdl-C6KE1.

EXAMPLE 4

CONSTRUCTION OF THE C-E3-E1 STRUCTURAL POLYPROTEIN AND CASSETTE

5 This polyprotein and expression cassette comprises the coding regions for an alphavirus capsid protein and glycoprotein E1. As described above, a leader/signal sequence also is included to facilitate insertion of the envelope glycoprotein into the endoplasmic reticulum. For example, one version of the cassette comprises C-E3-E1 (Figure 2) and was assembled by using two separate PCR
10 amplification reactions for C-E3 and E3-E1, followed by an overlapping PCR reaction. Specifically, a C-E3 PCR product was generated with the following primers:

SINCAPF:

5'ATATATCTCGAGCACCATGAATAGAGGATTCTTTAAC (SEQ. ID NO. 9)

CE3E1R:

15 5'GTGGTCGCATGTTCGCTTCTTTTGCTTCTGCCAGACG (SEQ. ID NO. 10)

The capsid forward primer (SINCAPF) contains a *Xho*I site to facilitate subcloning and the structural polyprotein Kozak consensus sequence. The E3 reverse primer (CE3E1R) contains the last six codons of E3, a serine codon in place of the first
20 E1 codon, and the next four codons of E1. The resulting 1kbp PCR product contains: 5'*Xho*I site/capsid ORF/E3 OFR/SER codon/first four codons of E1.

In the other reaction, an E3-E1 PCR product was generated with the following primers:

E3E1F:

25 5'GAAGCAAAAGAAGCGAACATGCGACCACTGTTCCAAATG (SEQ. ID NO. 11)

SINE1R:

5'TATATAGCGGCCGCTCATCTTCGTGTGCTAGTCAGCATC (SEQ. ID NO. 12)

The E3 forward primer (E3E1F) contains the last four codons of E3, a serine codon in place of the first E1 codon, and the first 8 codons of the E1 ORF. The E1 reverse primer (SINE1R) contains a *NotI* site to facilitate subcloning. The resulting 1.3 kbp PCR product contains: 5' last four codons of E3/SER codon/E1 ORF/*NotI* site.

Both PCR reactions were performed as above using the Sindbis virus genomic pDCMVSING as template. The separate PCR products were purified using QIAquick PCR purification kit (QIAGEN, Santa Clarita, CA), and then combined as template for an "overlapping" PCR reaction containing the primers SINCAPF and SINE1R. The resulting 2.3 kbp C-E3-E1 PCR product was purified using the QIAquick kit, digested with *XhoI* and *NotI*, and isolated from an agarose gel using GENECLAN. This isolated C-E3-E1 PCR fragment was then cloned directly into SINdl-luc that also was digested with *XhoI* and *NotI* to remove the luciferase gene insert (as described above). The resulting expression construct was designated SINdl-CE3E1.

EXAMPLE 5

CONSTRUCTION OF THE C-TPA-E1 STRUCTURAL POLYPROTEIN AND CASSETTE

This polyprotein and expression cassette comprises the coding regions for an alphavirus capsid protein and glycoprotein E1. As described above, a leader/signal sequence also is included to facilitate insertion of the envelope glycoprotein into the endoplasmic reticulum. For example, one version of the cassette utilizes a non-alphavirus derived leader/signal peptide sequence, such as that of tissue plasminogen activator (TPA), and comprises the polyprotein C-leader-E1 (Figure 2). For use of the TPA leader, this construct is assembled by using two separate PCR amplification reactions for C-TPA and TPA-E1, followed by an overlapping PCR reaction. Specifically, a C-TPA PCR product is generated with the following primers:

SINCAPF:

5'ATATATCTCGAGCACCATGAATAGAGGATTCTTTAAC (SEQ. ID NO. 9)

TPACAPR:

5'TGCTCCACACAGCAGCAGCACACAGCAGAGCCCTCTCTT
 CATTGCATCTGCGGACCACTCTTCTGTCCCTTCCGG
 (SEQ. ID NO. 13)

5

The capsid forward primer (SINCAPF) was described above. The TPA reverse primer (TPACAPR) contains the last seven codons of capsid, the first two codons of E3, and the first 16 codons of TPA leader. The resulting 0.9 kbp PCR product contains: 5'*Xho*I site/capsid ORF/TPA ORF.

10

In the other reaction, a TPA-E1 PCR product was generated with the following primers:

TPAE1F:

5'TGTGTGCTGCTGCTGTGTGGAGCAGTCTTCGTTTCGCCCAGCGC-
 TAGCTACGAACATGCGACCACTGTTCCAAAT (SEQ. ID NO. 14)

15

SINE1R:

5'TATATAGCGGCCGCTCATCTTCGTGTGCTAGTCAGCATC (SEQ. ID NO. 12)

20

The TPA forward primer (TPAE1F) contains the last 16 codons of TPA, including the cleavage signal, and the first nine codons of the E1 ORF. The E1 reverse primer (SINE1R) was described above. The resulting 1.4 kbp PCR product contains: 5' last 16 codons of TPA/E1 ORF/*Not*I site.

25

Both PCR reactions were performed as above using the Sindbis virus genomic pDCMVSiNg as template. The separate PCR products were purified using the GENECLAN II Kit and then combined as template for an "overlapping" PCR reaction containing the primers SINCAPF and SINE1R. The resulting 2.3 kbp C-TPA-E1 PCR product was purified using the QIAquick kit, digested with *Xho*I and *Not*I, and isolated from an agarose gel using GENECLAN. This isolated C-TPA-E1 PCR fragment was then cloned directly into SINDl-luc that also was digested with *Xho*I and *Not*I to remove the luciferase gene insert (as described above). The resulting expression construct was designated SINDl-CTPAE1.

30

Similarly to that described above, a non-alphavirus leader (*e.g.*, TPA) also may be used to generate polyproteins and expression cassettes with an alphavirus E2 glycoprotein.

EXAMPLE 6

5 VECTOR PACKAGING WITH STRUCTURAL POLYPROTEIN EXPRESSION CASSETTES

Packaging of alphavirus vectors with various combinations of the above structural protein expression cassettes was demonstrated using, as an example, Sindbis vector replicons expressing a β -galactosidase or GFP reporter (Dubensky et al., 1996, *ibid*; Polo et al., 1999, *PNAS* 96:4598-4603). The SIN vector was linearized as
10 described previously, and the various structural protein expression cassettes also were linearized using *PmeI*. Vector and defective helper RNAs then were transcribed in vitro from the linearized plasmids, with bacteriophage SP6 polymerase as described in Dubensky et al. (*ibid*). To package vector particles, the SIN vector RNA was co-transfected into BHK-21 cells (Dubensky, *ibid*) together with the C-E3-E2 defective
15 helper RNA and one of the E1 expressing defective helper RNAs, selected from the group C-6K-E1, C-E3-6K-E1, C-E3-E1, C-tPA-E1. Transfected cells were incubated at 37°C for 24 hr, at which time the culture supernatants were harvested and clarified by centrifugation. Clarified supernatants were then serially diluted and used to infect in duplicate, naïve BHK-21 cells for approximately 14 hr. The infected cells were
20 analyzed for reporter expression and the number of cells exhibiting such expression counted to determine vector particle titer in the original supernatants. Results demonstrating functionality of different E1 expressing constructs for vector packaging are shown in Figure 4.

EXAMPLE 7

25 DERIVATION OF STABLE ALPHAVIRUS VECTOR PACKAGING CELL LINES

Stable alphavirus vector packaging cell lines that express the structural polyproteins of the present invention are generated using expression cassettes that function directly in eukaryotic cells, to stably transform an appropriate cell type (*e.g.*, BHK-21, CHO cells), rather than requiring a prior step of in vitro transcription. The

expression cassettes may comprise a DNA promoter of RNA synthesis (*e.g.*, RNA polymerase II promoter) operably linked directly to the structural polyprotein genes, or a configuration wherein the structural polyprotein genes are operably linked to their native subgenomic promoter and require induction by vector supplied replicase (*e.g.*,
5 alphavirus nonstructural proteins, see US 5,789,245). For example, cassettes expressing either Sindbis virus-derived polyproteins C-E3-E2 and C-E3-6K-E1 are constructed in a configuration comprising the following ordered elements: RNA polymerase II promoter, 5' viral or defective-interfering RNA sequence required in *cis* for alphaviral replication, viral subgenomic junction region promoter, alphavirus structural
10 polyprotein sequence, 3' alphaviral sequence required in *cis* for replication, polyadenylate tract, and transcription termination sequence.

As a first step, an expression cassette backbone for use in packaging cells was constructed. This backbone can serve as the initial starting material for any of the structural polyprotein genes of the present invention. All components used in this
15 construction have been described in detail previously (WO 97/38087). Specifically, the expression cassette backbone was generated by step-wise insertions into plasmid pBGSV3'. A Sindbis virus junction region promoter plus *Xho*I and *Not*I cloning sites were obtained as a luciferase reporter-containing fragment from plasmid pDCMVSIN-luc, by digestion with *Bam*HI and *Fsp*I, and purification of the luciferase reporter-containing fragment from a 0.7% agarose gel using GENECLAN. The fragment was
20 ligated into plasmid pBGSV3' that also had been digested with *Bam*HI and *Fsp*I, and treated with alkaline phosphatase to produce a plasmid designated pBGSV3'BaFLuc. Next, an RSV (polII) promoter/5'-end tRNA sequence was obtained from 987DHBB by digestion with *Bgl*II and *Bam*HI and purification from a 1% agarose gel using
25 GENECLAN. This fragment was ligated into pBGSV3'BaFLuc that was similarly digested with *Bgl*II and *Bam*HI, to produce the expression cassette backbone construct pBRSV987dl-Luc. Alphavirus structural polyprotein genes and selectable markers are next inserted into this cassette in appropriate combinations.

The C-E3-E2 polyprotein gene is obtained from plasmid SINdl-CE3E2
30 (Example 1) by digestion with *Xho*I and *Not*I, and purification from an agarose gel

using GENECLAN. The C-E3-E2 fragment is then ligated into the packaging cell line expression plasmid, pBRSV987dl-Luc, which is also digested with *XhoI* and *NotI* to remove the luciferase gene insert, and purified from an agarose gel using GENECLAN. The resulting construct is designated pBRSV987dl-CE3E2. Insertion
5 of a neomycin phosphotransferase gene (*neo^r*) into the region of nonstructural protein gene deletion as a selectable marker, is next accomplished by digestion with *BspEI* and *BamHI*, purification from an agarose gel using GENECLAN, and ligation with a PCR-amplified *neo^r* gene (see WO 97/38087) that is also digested with *BspEI* and *BamHI*, and purified from an agarose gel. The resulting construct is designated
10 pBRSV987dlneo-CE3E2.

The C-E3-6K-E1 polyprotein gene is obtained from plasmid SINdl-CE36KE1 (Example 2) by digestion with *XhoI* and *NotI*, and purification from an agarose gel using GENECLAN. The C-E3-6K-E1 fragment is then ligated into the packaging cell line expression plasmid pBRSV987dl-Luc, which is also digested with
15 *XhoI* and *NotI* to remove the luciferase gene insert, and purified from an agarose gel using GENECLAN. The resulting construct is designated pBRSV987dl-CE36KE1. Insertion of a hygromycin phosphotransferase gene (*hyg^r*) into the region of nonstructural protein gene deletion as a selectable marker is next accomplished by digestion with *BspEI*, blunt-ending with Klenow, and further digesting with *BamHI*.
20 The *hyg^r* insert is obtained as a PCR-amplified product (see WO 97/38087) digested with *EcoRV* and *BamHI*, and purified from an agarose gel. This fragment is ligated into the prepared pBRSV987dl-CE36KE1 vector, resulting in the construct pBRSV987dlhyg-CE36KE1.

To generate stable alphavirus packaging cell lines, cells (*e.g.*, BHK-21)
25 are transfected initially with plasmid pBRSV98dlneo-CE3E2 using Lipofectamine, as described by the manufacturer. Approximately 24 hr post-transfection, the cells are trypsinized and re-plated in media containing 600 ug/ml of the drug G418 (neomycin). The media is exchanged periodically with fresh G418-containing media and foci of resistant cells are allowed to grow. Cells are trypsinized and cloned by limiting dilution
30 in 96 well tissue culture dishes, and individual cell clones are grown and expanded for

screening. Cells which inducibly expressed capsid protein and glycoprotein E2 in response to input vector are identified by transfecting with Sindbis luciferase vector RNA or Sindbis β -galactosidase DNA vectors (Dubensky et al., 1996, *ibid*), making cell lysates approximately 24 or 48 hr post-transfection, and performing western blot
5 analysis with a rabbit anti-Sindbis polyclonal antibody. A positive CE3E2 cell line demonstrating expression of the Sindbis virus capsid and E2 is then used for subsequent steps.

Next, the positive CE3E2 cell line is transfected with pBRSV987dlhyg-CE36KE1 using Lipofectamine, as described by the manufacturer. Approximately 24
10 hr post-transfection, the cells are trypsinized and re-plated in media containing 500 ug/ml of hygromycin (Boehringer Mannheim). The media is exchanged periodically with fresh media containing hygromycin and foci of resistant cells are allowed to grow. Cells are trypsinized and cloned by limiting dilution in 96 well tissue culture dishes, and individual cell clones grown and expanded for screening. Split
15 structural gene PCL derived in this manner are designated CE2/CE1 PCL. Positive cells which inducibly express all Sindbis virus structural proteins in a biologically active form in response to input vector are identified in transfer of expression (TOE) assays, which demonstrate that transfected vector molecules could induce structural protein expression, resulting in packaging and secretion of vector particles that could in
20 turn be used to infect naive cells. Packaging of Sindbis virus RNA vectors expressing β -galactosidase (Dubensky et al, 1996, *ibid*) is accomplished by RNA transfection into PCL clones and harvesting of supernatants at 24-48 hr post-transfection. The harvested supernatants may be used to infect naive BHK-21 cells for an additional 18 hr, wherein infected cell lysates are harvested and enzymatic β -galactosidase activity determined to
25 demonstrate packaging. Alternatively, alphavirus vectors also may be packaged by infecting the packaging cells with recombinant alphavirus vector particles or transfecting the packaging cells with an alphavirus-derived Eukaryotic Layered Vector Initiation System (US 5,814,482), incubating the cells for a period of time sufficient for vector packaging and production (*e.g.*, 48 hr), and harvesting the culture supernatants
30 containing the packaged vector particles.

As should be readily apparent, substitution of the other alphavirus structural polyproteins of the present invention into these or other expression cassettes is easily accomplished using the teachings provided herein.

EXAMPLE 8

5 VECTOR PACKAGING WITH REDUCED HETEROLOGOUS GENE EXPRESSION

During the packaging of alphavirus vector replicons into recombinant alphavirus particles, either by RNA co-transfection with one or more defective helper RNAs or by introduction into stable alphavirus packaging cell lines, high level expression of the encoded heterologous gene (transgene) occurs. In those instances
10 where expression of the transgene results in toxicity for the host cells, or interference with alphaviral vector replication and packaging, the level of recombinant alphavirus vector particles produced may be substantially decreased. To overcome this issue, the present invention provides methods for the reduction of transgene expression in specific cells that are used for alphavirus vector replicon packaging and recombinant vector
15 particle production. The reduction of transgene expression during the packaging and particle production process provides a method to overcome the issue of toxic or interfering transgene expression, as well as to increase the overall titer of vector particles produced.

A. Use of Complementary (Antisense) RNA to Reduce Transgene Expression

20 For example, the cells used for alphavirus vector packaging may further comprise a cassette that expresses an RNA sequence complementary (*e.g.*, antisense) to the junction region promoter or subgenomic mRNA of the alphavirus vector replicon. In preferred embodiments, the cells are stably transformed with said cassette that expresses intracellularly an antisense RNA. In other embodiments, the antisense
25 molecules may be supplied exogenously. A variety of promoters may be used for intracellular expression of the antisense RNA sequence, including for example RNA polymerase I, II, and III promoters and an alphavirus junction region promoter. Stably transformed cassettes which produce antisense RNA complementary to the minus strand

subgenomic junction region promoter region of Sindbis virus (see Figure 4), for example, may be constructed as follows:

A plasmid known as pBGS131dL*Xho*I-BGHTT (WO 97/38087) is used as initial material for constructions. This plasmid contains a bovine growth hormone (BGH) transcription termination/polyadenylation signal with a *Sac*I site immediately upstream. First, a fragment consisting of the 40 terminal nucleotides from the Sindbis virus 3'-end plus a 25 nucleotide synthetic A-tract, and flanked by *Not*I and *Sac*I sites, is generated using two oligonucleotides (ID No. 20 and 21) from U.S. Patent 5,814,482 as described therein. The fragment is digested with *Not*I and *Sac*I, and ligated into plasmid pBGS131dL*Xho*I-BGHTT that has also been digested with *Not*I and *Sac*I, and purified from an agarose gel using GENECLAN. The resulting construct is designated pBGSdIX-3'BGH. Next, a Sindbis virus junction region promoter plus *Xho*I and *Not*I cloning sites is inserted into the plasmid similarly to that described above in Example 7. The junction region promoter fragment is obtained as a luciferase reporter-containing fragment from plasmid pDCMVSIN-luc, by digestion with *Bam*HI and *Not*I, and purification of the luciferase reporter-containing fragment from a 0.7% agarose gel using GENECLAN. The fragment is ligated into plasmid pBGSdIX-3'BGH that also has been digested with *Bam*HI and *Not*I, to generate a plasmid designated pBGSBNluc-3'BGH. Next, an RSV (polII) promoter/5'-end tRNA sequence is obtained from 987DHBB (WO 97/38087) by digestion with *Bgl*II and *Bam*HI and purification from an agarose gel using GENECLAN. This fragment is ligated into pBGSBNluc-3'BGH that is similarly digested with *Bgl*II and *Bam*HI, to produce the construct pBRSV987dl-Luc3'sh. Insertion of a neomycin phosphotransferase gene (*neo*^r) into the region of nonstructural protein gene deletion as a selectable marker, is next accomplished by digestion with *Bsp*EI and *Bam*HI, purification from an agarose gel using GENECLAN, and ligation with a PCR-amplified *neo*^r gene (see WO 97/38087) that is also digested with *Bsp*EI and *Bam*HI, and purified from an agarose gel. The resulting expression cassette backbone is designated p987dlneo-Luc3'sh.

Finally, a cDNA corresponding to a desired antisense sequence is generated using two overlapping synthetic oligonucleotides, which will produce a fragment with *Xho*I and *Not*I sites.

AntiJRFwd

5 5'ATATATCTCGAGGCCATCAGAGGGGAAATAAAGCATCTCTACGGTGG
(SEQ. ID NO. 15)

AntiJRRev

10 5'TATATATGCGGCCGCTGACTATTTAGGACCACCGTAGAGATGCTTTAT
(SEQ. ID NO. 16)

For example, the oligonucleotides listed above are mixed together at equimolar concentrations in the presence of 10 mM MgCl₂, heated to 100°C for 5 minutes, and cooled slowly to room temperature. The partially double-stranded molecule is then filled in using Klenow enzyme and 50 uM dNTPs. The resultant
15 molecule is digested with *Xho*I and *Not*I, purified from a 2% NuSieve/1% agarose gel, and ligated to p987dlneo-Luc3'sh that has also been digested with *Xho*I and *Not*I, and purified from an agarose gel to remove the luciferase insert. The resulting construct is designated p987dlneo-antiJR.

In addition to this construct, variations in the junction region promoter
20 antisense sequence may be made by altering the overall sequence length or by directing hybridizing to sequences 5' or 3' to those provided, or by utilizing chimeric vector replicons that have incorporated sequences (*e.g.*, subgenomic promoter or subgenomic 5'NTR sequence) from other alphaviruses. Furthermore, and as described below, similar antisense sequences also may be designed to hybridize directly to the alphavirus
25 vector subgenomic mRNA.

To generate stable cell lines containing p987dlneo-antiJR or other similar cassettes, cells (*e.g.*, BHK-21) are transfected with the plasmid using Lipofectamine, as described by the manufacturer. Approximately 24 hr post-transfection, the cells are trypsinized and re-plated in media containing 600 ug/ml of the
30 drug G418 (neomycin). The media is exchanged periodically with fresh G418-

containing media and foci of resistant cells are allowed to grow. Cells are trypsinized and cloned by limiting dilution in 96 well tissue culture dishes, and individual cell clones are grown and expanded prior to screening.

Alternatively, an antisense sequence is expressed from a cassette, such as
 5 an RNA polymerase III (pol III) cassette. For example, an antisense sequence which binds to the region of minus strand vector RNA, inclusive of the subgenomic junction region promoter, is expressed from a cassette comprising the following ordered elements: Adenovirus 2 VA1 RNA promoter (nucleotides -70/+30), nucleotides 7562-7606 of Sindbis virus, and the RNA polymerase III consensus transcription termination
 10 sequence. These elements are cloned into a pUC-derived kanamycin-resistant plasmid DNA vector, pBGSVG (WO 97/38087), with a <*PmeI* - *BglIII* - *XhoI* - *NotI* - *EcoRI* - *PmeI*> polylinker sequence and unique *PacI* site on the opposite side of the vector backbone.

Assembly of the desired components in the pol III-based expression
 15 cassette is performed by PCR to juxtapose the following ordered sequences:

Ad2 Stuffer (nts. 1051-10584):

5'CCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGACGCTC
 TGGA-3' (SEQ. ID NO. 17)

Ad2 VA1RNA promoter [(-70/+30) (nts. 10585-10682)]:

20 5'CCGTGCAAAAGGAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGA
 TAAATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGA-3'
 (SEQ. ID NO. 18)

Sindbis virus nucleotides 7562-7606

CAGAGGGGAAATAAAGCATCTCTACGGTGGTCCTAAATAGTCAGC
 25 (SEQ. ID NO. 19)

containing media and foci of resistant cells are allowed to grow. Cells are trypsinized and cloned by limiting dilution in 96 well tissue culture dishes, and individual cell clones are grown and expanded prior to screening.

Alternatively, an antisense sequence is expressed from a cassette, such as
 5 an RNA polymerase III (pol III) cassette. For example, an antisense sequence which binds to the region of minus strand vector RNA, inclusive of the subgenomic junction region promoter, is expressed from a cassette comprising the following ordered elements: Adenovirus 2 VA1 RNA promoter (nucleotides -70/+30), nucleotides 7562-7606 of Sindbis virus, and the RNA polymerase III consensus transcription termination
 10 sequence. These elements are cloned into a pUC-derived kanamycin-resistant plasmid DNA vector, pBGSVG (WO 97/38087), with a <PmeI - BglII - XhoI - NotI - EcoRI - PmeI> polylinker sequence and unique PaeI site on the opposite side of the vector backbone.

Assembly of the desired components in the pol III-based expression
 15 cassette is performed by PCR to juxtapose the following ordered sequences:

Ad2 Stuffer (nts. 10531-10584):

5'CCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGACGCTC
 TGGA-3' (SEQ. ID NO. 17)

Ad2 VA1RNA promoter [(-70/+30) (nts. 10585-10682)]:

20 5'CCGTGCAAAAGGAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGA-3'
 (SEQ. ID NO. 18)

Sindbis virus nucleotides 7562-7606

25 CAGAGGGGAAATAAAGCATCTCTACGGTGGTCCTAAATAGTCAGC
 (SEQ. ID NO. 19)

AMENDED SHEET

51

Temperature (°C)	Time (Min.)	No. Cycles
95	2	1
95	0.5	
55	0.5	10
72	0.5	

The PCR product is purified using the QIAquick PCR kit, the purified product then used in a second PCR amplification, with the primer pair shown below:

Ad2 PCR 1F:

5 5'-ATCTTCATGCGGCCGCCATGGTCGGGAC-3' (SEQ. ID NO. 25)

Ad2 PCR 1R:

5'-ATCTTCATGCGGCCGCTCCGGAGCGCAA-3' (SEQ. ID NO. 26)

The second PCR amplification of the pol III-based cassette is performed with the Vent polymerase and similar reaction conditions, with the PCR amplification
10 protocol, as shown below.

Temperature (°C)	Time (Min.)	No. Cycles
95	2	1
95	0.5	
55	0.5	20
72	0.5	
72		1

The products of this second PCR reaction are purified with the QIAquick kit, digested with *NotI*, re-purified and then inserted into the vector pBGSVG that has
15 also been digested with *NotI* and treated with alkaline phosphatase. This plasmid is designated pAntiSINjr.

Insertion of a hygromycin drug selectable marker into pAntiSINjr is accomplished using the unique *PacI* site in the vector backbone. A hygromycin marker cassette, under the control of an HSV TK promoter, is isolated from plasmid
20 pBGSVhygro-G (WO 97/38087) by digestion with *PacI* and purification from an agarose gel using GENECLAN. The isolated fragment then is ligated into pAntiSINjr

that has also been digested with *PacI*, resulting in a construct designated pAntiSINjrHyg.

Plasmid pAntiSINjrHyg is transfected into a desired cell type (*e.g.*, BHK-21) using Lipofectamine, as described by the manufacturer. Approximately 24 hr
5 post-transfection, the cells are trypsinized and re-plated in media containing 500 ug/ml of hygromycin (Boehringer Mannheim). The media is exchanged periodically with fresh drug-containing media and foci of resistant cells are allowed to grow. Cells are trypsinized and cloned by limiting dilution in 96 well plates.

Initially, demonstration of reduced transgene expression from Sindbis or
10 other similar replicon vectors in such cell lines is accomplished by linearization and in vitro transcription of a reporter vector, as described previously. The in vitro transcribed vector RNA is then transfected or electroporated into the modified cells, as well as control parental cells, followed by harvest of cell lysates at 12 and 24 hr post-transfection. Reporter gene expression is quantitated as previously described
15 (Dubensky et al, 1996 *ibid*, U.S. Patent 5,789,245, and WO 97/38087), and compared between the cell lines. Decreased expression in the modified cells, as compared to the parental cells, is indicative of specific transgene suppression. These systems may be similarly used in a variety of cell lines for the packaging of alphavirus vector particles, and established methods for packaging are provided elsewhere (US 5,789,245,
20 Dubensky et al, 1996, *ibid*; Polo et al., 1999, *ibid*).

In another embodiment of the present invention, chimeric alphavirus replicons, as well as chimeric alphavirus glycoprotein defective helper cassettes are disclosed. In such chimeras, the wild-type subgenomic 5'-nontranslated (NTR) region of one alphavirus is substituted with the corresponding region from another alphavirus.
25 Such a chimera provides a number of advantages for use in the present invention. For example, by using different subgenomic 5'-NTR sequences, transgene expression from one vector or defective helper RNA can be preferentially targeted with antisense molecules, while not affecting other vector or defective helper RNAs that contain the wild-type sequence. Furthermore, the incorporation of a specific 5'-NTR from an
30 alphavirus whose subgenomic translational enhancer element does not extend into the

capsid gene coding sequence (*e.g.*, Venezuelan equine encephalitis virus, VEE), may overcome potential expression issues for vector and defective helper RNAs derived from alphaviruses that do require such a capsid-inclusive element (*e.g.*, Sindbis and Semliki Forest virus).

5 Specifically, a Sindbis virus derived vector replicon and glycoprotein defective helper (US 5,789,245) may be modified to contain the subgenomic 5'NTR sequence from VEE, as follows. A cDNA fragment comprising both the Sindbis virus subgenomic promoter and the VEE subgenomic 5'-NTR is synthesized by PCR using overlapping oligonucleotides. These oligonucleotides further contain *Bam*HI and *Xho*I
10 restriction sites for convenient substitution into the final constructs.

VEE5'NTR1:

5'ATATAGGATCCCCTGAAAAGGCTGTTTAAGTTGGGTAAACCGCTCCCAGC
CGACGACGAGCAAGACGAAGACAGAAGACGC (SEQ ID NO:35)

VEE5'NTR2:

15 5'CCACTGCTAAAGTGCCTGTTATACCTACTCTAAACCACGCCTTTGTTTCAT
CTAGCAGAGCGCGTCTTCTGTCTTCGTCTT (SEQ ID NO:36)

VEE5'NTR3:

5'AACAGGCACTTTAGCAGTGGCCGTGACGACCCGGTATGAGGTAGACAATA
TTACACCTGTCCTACTGGCATTGAGAACTTTTGCCCAG
20 AGCAAAAGAGCATTCCAAGC (SEQ ID NO:37)

VEE5'NTR4:

5'ATATACTCGAGCTTGGCGGACTAGACTATGTCGTAGTCTATTTAGGACCA
CCGTAGAGATGCTTTATTTCCCCTCTGATGGCTTGGAATGCTCTTTTGCT
(SEQ ID NO:38)

PCR amplification of the oligonucleotides shown above is performed in a single reaction, using Vent polymerase and reaction conditions as suggested by the supplier, containing in addition 2 mM MgSO₄, 5% DMSO, and Hot Start Wax beads, with the PCR amplification protocol, as shown below.

5

Temperature (°C)	Time (Min.)	No. Cycles
95	2	1
95	0.5	
55	0.5	10
72	0.5	

The PCR product is purified using the QIAquick PCR kit, and then the purified product is used in a second PCR amplification, with the primer pair shown below:

10 VEE5'NTR5:

5'ATATAGGATCCCCTGA (SEQ ID NO:39)

VEE5'NTR6:

5'ATATACTCGAGCTTGG (SEQ ID NO:40)

15

Following an additional purification step, the synthesized PCR fragment is digested with *Bam*HI and *Xho*I, and ligated into a Sindbis replicon vector and glycoprotein defective helper that were similarly digested with *Bam*HI and *Xho*I. An additional final cloning step is required for the replicon construct, due to the unwanted deletion of a *Bam*HI – *Bam*HI nonstructural gene fragment. This fragment is obtained from another preparation of replicon DNA by *Bam*HI digestion and gel purification, and then it is ligated into the new VEE 5'NTR containing replicon that also has been digested with *Bam*HI and treated with alkaline phosphatase.

20

B. Use of Aptamer-Ligand Combinations to Reduce Transgene Expression

Alternatively, aptamer RNA as described herein may be using to reduce transgene expression by inserting the aptamer sequence into vector RNA to be

25

packaged, and providing the appropriate ligand to the population of cells during vector packaging (see Figure 6). For example, H10 and H19 aptamers (Werstuck and Green, *Science* 282:296-298), are generated as a tandem pair (single copy each) by PCR using overlapping oligonucleotides which produce flanking and internal restriction sites for
5 manipulation. The aptamer sequences are shown below:

H10 Aptamer sequence

GGTGATCAGATTCTGATCCAATGTTATGCTTCTCTGCCTGGGAACAGCTGCC
TGAAGCTTTGGATCCGTCGC (SEQ. ID NO. 27)

H19 Aptamer sequence

10 GGTGATCAGATTCTGATCCAACAGGTTATGTAGTCTCCTACCTCTGCGCCTG
AAGCTTGGATCCGTCGC (SEQ. ID NO. 28)

To demonstrate the ability of these aptamers to reduce transgene expression from alphavirus vectors in the presence of their ligands, Hoechst dye
15 H33258 or H33342, a reporter vector is first constructed with the aptamer sequence inserted at a site which corresponds to the 5'-nontranslated region of the subgenomic RNA, immediately upstream of the AUG initiator codon for the reporter gene. Vectors containing other transgenes may be constructed in a similar manner, and alternatively, may also contain aptamer insertion at other sites.

20 In order to construct an alphavirus-derived vector containing the above aptamer insert, a previously described Sindbis vector is modified to position a unique *Xho*I site just downstream and adjacent to the subgenomic transcription start site. This is accomplished by PCR amplification using the following two oligonucleotides:

SIN5122F:

25 5'ATATATGGCCGAAGAGGCCCCCGAAGTTGTAG
(SEQ. ID NO. 29)

SIN7600R:

5'ATATATCTCGAGTATTTAGGACCACCGTAGAGATGC
(SEQ. ID NO. 30)

The forward primer (SIN5122F) contains a unique *Sfi*I site within the Sindbis virus nsP3 sequence, while the reverse primer (SIN7600R) contains the junction region promoter and a *Xho*I site. PCR is performed using VENT polymerase and plasmid pRSIN- β -gal (Dubensky et al., 1996, *ibid*) as template. The approximately 2.5 kbp fragment is purified using the QIAquick kit, digested with *Sfi*I and *Xho*I, and ligated to plasmid pRSIN- β -gal that has also been digested with *Sfi*I and *Xho*I to remove the corresponding fragment, and purified from an agarose gel using GENECLAN. The new Sindbis virus vector construct is designated pRSINjr- β gal.

Generation of the aptamer sequence for insertion into this vector is accomplished using a series of overlapping oligonucleotides listed below to produce a fragment comprising the following ordered elements: 5'-*Sal*I site/*Aat*II site/H10 Aptamer/*Pac*I site/H19 Aptamer/*Xho*I site/*Not*I site-3'.

AptH10Fwd1:

15 5'ATATATGTCGACGTC
GGTGATCAGATTCTGATCCAATGTTATGCTTCTCTGCCTGGGAACAGC
(SEQ. ID NO. 31)

AptH10Rev1:

20 5'TCTGATCACCTTAATTAAGCGACGGATCCAAAGCTTCAGGCAGCTGTTCCC
AGGCAGAGAAGCATA (SEQ. ID NO. 32)

AptH19Fwd1:

5'GCTTAATTAA
GGTGATCAGATTCTGATCCAACAGGTTATGTAGTCTCCTACCTCTGCG
(SEQ. ID NO. 33)

AptH19Rev1:

25 5'ATATATGCGGCCGCATATCTCGAG
GCGACGGATCCAAGCTTCAGGCGCAGAGGTAGGAGACTACATAAC
(SEQ. ID NO. 34)

The partially complementary oligonucleotides listed above are first used at 1 μ M concentration in a short (10 cycle) single PCR amplification reaction, with

VENT polymerase as suggested by the supplier, containing in addition 2 mM MgSO₄, 5% DMSO, and Hot Start Wax beads (Perkin-Elmer), with a PCR amplification protocol, as shown below.

Temperature (°C)	Time (Min.)	No. Cycles
95	2	1
95	0.5	
55	0.5	10
72	0.5	

5

The PCR product is purified with the QIAquick kit and then used in a second PCR amplification, with the primer pair AptH10Fwd1 and AptH19Rev1 and additional cycles, under similar conditions. The products of this second PCR reaction again are purified with QIAquick, digested with *SalI* and *NotI*, and then inserted into
 10 plasmid pRSINjr- β gal that has been digested with *XhoI* and *NotI*, and gel-purified to remove the β gal insert. This vector plasmid is designated pRSINapt-empty. The β -gal insert is then re-inserted into the vector by digestion of pRSINjr- β gal with *XhoI* and *NotI*, purification of the fragment from an agarose gel, and ligation into similarly digested and gel-purified plasmid pRSINapt-empty. The resulting reporter vector
 15 construct is pRSINapt- β gal.

Demonstration of the reduction in transgene translation as compared to the unmodified parental vector is accomplished by linearization and in vitro transcription of both vectors as described previously. The in vitro transcribed vector RNAs are then transfected or electroporated into cells that are growing in Hoechst dyes
 20 H33258 or H33342, ranging in concentration from 5 to 50 mM. At 12 and 24 hr post-transfection, cell lysates are obtained and reporter gene expression is quantitated as previously described (Dubensky et al, 1996 *ibid*, U.S. Patent 5,789,245, and WO 97/38087).

Of particular interest, insertion of artificial stem-loop structures (*e.g.*,
 25 aptamers) into the 5' non-translated region of alphavirus subgenomic RNA as disclosed herein, also may provide a mechanism to enhance translation (*e.g.*, in the absence of

ligand), by substituting for the native alphavirus stem-loop translational enhancer structure in this region.

In another embodiment of this technology, the bacteriophage R17 coat protein and corresponding specific RNA ligand may be used to reduce transgene
5 expression. The RNA binding site for bacteriophage R17 coat protein (Krug et al. 1982, *Biochemistry* 21:4713-4720) can be introduced in the 5' end of the subgenomic RNA. The minimal R17 binding site comprises a sequence AAACAUGAGGAUUACCCAUGU (SEQ ID NO:41) and forms a stem loop structure absolutely required for binding. Analysis of a population of high affinity RNA ligands
10 to the coat protein revealed the components of a hairpin that promote favorable interactions with the coat protein (Schneider et al. 1992 *J. Mol. Biol.* 228:862-869).

To demonstrate the ability of this binding site to reduce the transgene expression from alphavirus vectors in the presence of the R17 coat protein, variants of the binding site were introduced into the 5'-nontranslated (NTR) region of a Sindbis
15 vector subgenomic RNA, immediately upstream of the AUG initiator codon for a reporter gene. Additional vectors containing other transgenes may be readily constructed in a similar manner by one of skill in the art using the teachings provided herein.

Two binding sites were designed to include components essential for
20 both high affinity binding to the coat protein and high efficiency of replication and subgenomic transcription of the Sindbis virus derived vectors (Figure 7). In both binding sites, specific elements were introduced comprising: 1) in the loop, a C replaced U₁₃ to maximize functionality of this structure in Eukaryotic cells, and 2) the two pairs G₉-C₁₆, G₁₀-C₁₅ were inverted. Furthermore, one binding site also included 1) the
25 substitution of the pair U₆-A₁₈ into G-C in the stem loop to eliminate the AUG codon, and 2) the substitution of A₁A₂ with first 5 nucleotides (AUAGU) of Sindbis 5'-nontranslated region which are essential for subgenomic transcription. This binding site was named 5TOP. The other binding site was designed to include the first 5 nucleotides of Sindbis 5'-nontranslated region in the stem, resulting in the substitution of the pairs

A₃-U₂₁, C₄-G₂₀, A₅-U₁₉, and G₇C₁₇ with the pairs U-A, A-U, G-C, and C-G respectively. This binding site was named TOP.

Introduction of the 5TOP binding site was accomplished by PCR amplification of two overlapping fragments also generated by PCR amplification, as follows:

5TOPF:

5'-CAGCACCATCAGGGCTGGCAGCATAGTACATTTTCATCTGAC (SEQ ID NO:42)

SCG8120R:

10 5'-CGTTGTGGCTGTTGTAGTTGTAC (SEQ ID NO:43)

In the forward primer (5TOPF), bases 1-19 are the R17 coat protein-binding site, while bases 20-42 are complementary to nt. 7603-7625 of the Sindbis replicon vector expressing a GFP reporter (SINCR-GFP). The reverse primer is complementary to nt. 8105-8083 of the same vector.

The second fragment was amplified using the following two oligonucleotides:

SCR728F:

5'-TGCGGCGGATTATCTTGCAAG (SEQ ID NO:44)

20 5TOPR:

5' -CAGCCCTGATGGTGCTGGACTATTTAGGACCACCGTAGAG (SEQ ID NO:45)

The forward primer is complementary to the 7281-7302 nt. of the Sindbis GFP reporter vector, while the reverse primer contains the R17 coat protein binding site in nts. 1-19, and nts. 20-41 are complementary to 7581-7602 of the Sindbis vector. The oligonucleotides were used at 2 µM concentration with 0.1µg of plasmid template in a single, 10-cycle PCR reaction with Pfu Polymerase, as suggested by the supplier. The amplification protocol is shown below.

60

Temperature (°C)	Time (Min.)	No. Cycles
94	2	1
94	0.5	
60	0.5	10
72	2	

The two amplified fragments were purified from an agarose gel using a QIAquick gel extraction kit, and an aliquot of each fragment was used as template for a second PCR amplification. The two fragments were mixed with Vent Polymerase as suggested by supplier, and one PCR amplification cycle was performed:

Temperature (°C)	Time (Min.)	No. Cycles
94	2	1
94	0.5	
42	1	1
72	3	

The following primers, overlapping respectively with the *Bam*HI₇₃₃₅ and *Xho*I sites of the Sindbis replicon, then were added at a 2 μ M concentration:

SCR7320F:

5' ATATATGCGTGCCGCGTGGCGGATCCCC (SEQ ID NO:46)

SCRG7672R:

5' ATATATCATGGTGGCTCGAGGGTGGTGTT (SEQ ID NO:47)

PCR amplification was performed as follows:

Temperature (°C)	Time (Min.)	No. Cycles
94	2	1
94	0.5	
60	0.5	30
72	2	

The PCR product (300bp) was purified using the QIAquick kit, digested with *Bam*HI and *Xho*I, gel purified from agarose gel as described above, and ligated

into plasmid SINCR-GFP that had also been digested with *Bam*HI and *Xho*I and purified from an agarose gel. Clones containing the inserts were verified by sequencing and, finally, the *Bam*HI₄₆₃₄-*Bam*HI₇₃₅₅ was reinserted into this construct. The new Sindbis virus vector construct was designated SINCR-5TOP-GFP.

5 The second Sindbis derived vector was constructed in a similar manner, with the only differences being the forward oligonucleotide primer for the PCR amplification of the first fragment, as follows:

TOPF:

10 5' CACCATCAGGGACTACAGCATAGTACATTTTCATCTGAC (SEQ ID NO:48)

and, the reverse primer for the PCR amplification of the second fragment, as follows:

TOPR:

5' TAGTCCCTGATGGTGACTATTAGGACCACCGTAGAG (SEQ ID NO:49)

15 Following insertion of the overlapping PCR fragment into the SINCR-GFP vector, this new Sindbis replicon construct was designated SINCR-TOP-GFP.

Next, the ability of these new vectors to express the reporter gene GFP and to be packaged into vector particles was demonstrated. Plasmid DNA from these two new constructs and from the parental construct were linearized and used for in vitro
20 transcription as described previously. Each transcript was co-transfected into BHK cells with together with helper RNAs expressing capsid and glycoproteins (Polo et al., 1999, *ibid*). Transfected cells were incubated for 24 hr, at which time the culture supernatants were collected and the cells harvested. The harvested cells were analyzed by flow cytometry. The supernatants were clarified by centrifugation, serially diluted
25 and used to infect in duplicate, naïve BHK-21 cells for approximately 14 hr. The infected cells were counted based on GFP fluorescence to determine vector particle titer in the original supernatants. Both constructs efficiently expressed GFP reporter and were packaged into recombinant alphavirus vector particles.

To demonstrate that transgene expression can be down-regulated in the

presence of R17 coat protein, vector particles derived from both modified constructs, as well as the parental construct, were used to infect at an MOI=3, 293 cells and 293 derivative cells that express the R17 coat protein. Following incubation for 24 hrs, GFP expression was analyzed by flow cytometry. As shown in Figure 8, expression of the GFP transgene was reduced significantly in R17-expressing cells (as compared to 293 cells) for each of the modified vector replicons, but not for the wild-type SINCR replicon. The R17 coat protein similarly may be expressed in a variety of cell lines for use in the packaging of alphavirus vector particles, and established methods for packaging are provided elsewhere (US 5,789,245, Dubensky et al, 1996, *ibid*; Polo et al., 1999, *ibid*) These data demonstrate the ability to specifically down-regulate transgene expression from alphavirus vectors, using methods of the present invention.

While the present invention has been described above both generally and in terms of preferred embodiments, it is understood that variations and modifications will occur to those skilled in the art in light of the description, *supra*. Therefore, it is intended that the appended claims cover all such variations coming within the scope of the invention as claimed.

SEQUENCE LISTING

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<120> COMPOSITIONS AND METHODS FOR PACKAGING OF ALPHAVIRUS
VECTORS

<130> PAT 49626W-1

<140> PCT/US99/31193

<141> 1999-12-30

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<151> 1998-12-31

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<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 6

gcggccgctc atcttcgtgt gctagtcagc atc

33

<210> 7

<211> 39

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 7

ctcgggtgaac gtttctgcgg accactcttc tgcccttc 39

<210> 8

<211> 38

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 8

gaagagtggc ccgcagaaac gttcaccgag accatgag 38

<210> 9

<211> 37

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 9

atatatctcg agcaccatga atagaggatt ctttaac 37

<210> 10

<211> 37

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 10

gtggtcgcat gttcgcttct tttgcttctg ccagacg 37

<210> 11

<211> 39

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 11

66

gaagcaaaaag aagcgaacat gcgaccactg ttccaaatg

39

<210> 12

<211> 39

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 12

tatatagcgg ccgctcatct tcgtgtgcta gtcagcatc

39

<210> 13

<211> 75

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 13

tgctccacac agcagcagca cacagcagag ccctctcttc attgcatctg cggaccactc
ttctgtccct tccgg

60

75

<210> 14

<211> 75

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 14

tgtgtgctgc tgctgtgtgg agcagtcttc gtttcgcccc gcgctagcta cgaacatgcg
accactgttc caaat

60

75

<210> 15

<211> 47

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

67

<400> 15

atatatctcg aggccatcag aggggaaata aagcatctct acggtgg

47

<210> 16

<211> 48

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 16

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48

<210> 17

<211> 54

<212> DNA

<213> adeno-associated virus 2

<400> 17

ccatggtcgg gacgctctgg ccggtgaggc gtgcgcagtc gttgacgctc tgga

54

<210> 18

<211> 98

<212> DNA

<213> adeno-associated virus 2

<400> 18

ccgtgcaaaa ggagagcctg taagcgggca ctcttccgtg gtctggtgga taaattcgca
agggtatcat ggcggacgac cggggttcga accccgga

60

98

<210> 19

<211> 45

<212> DNA

<213> Sindbis virus

<400> 19

cagaggggaa ataaagcatc tctacggtgg tcctaaatag tcagc

45

<210> 20

<211> 13

<212> DNA

<213> Unknown

68

<220>

<223> Description of Unknown Organism: RNA polymerase
III consensus transcription termination sequence

<400> 20

gcgcttttttg cgc

13

<210> 21

<211> 79

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 21

gcggccgcca tggtcgggac gctctggccg gtgaggcgtg cgcagtcgtt gacgctctgg
accgtgcaaa aggagagcc

60

79

<210> 22

<211> 58

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 22

cccttgcgaa tttatccacc agaccacgga agagtgcccg cttacaggct ctcctttt

58

<210> 23

<211> 80

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 23

aattcgcaag ggtatcatgg cggacgaccg gggttcgaac cccggatcta gacctaaaac
caaagtacag aggggaaata

60

80

<210> 24

<211> 72

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 24

gcggccgctc cggagcgcaa aaagcgcgct gactatttag gaccaccgta gagatgcttt 60
atttcccctc tg 72

<210> 25

<211> 28

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 25

atcttcatgc ggccgccatg gtcgggac 28

<210> 26

<211> 28

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence: Primer for
PCR amplification

<400> 26

atcttcatgc ggccgctccg gagcgcaa 28

<210> 27

<211> 72

<212> DNA

<213> Unknown

<220>

<223> Description of Unknown Organism: See Werstuck &
Green, Science 282: 296-298, 1998. Isolated
aptamer that binds specifically to Hoechst dye
H33258.

<400> 27

70

ggtgatcaga ttctgatcca atgttatgct tctctgcctg ggaacagctg cctgaagctt 60
 tggatccgtc gc 72

<210> 28
 <211> 69
 <212> DNA
 <213> Unknown

<220>
 <223> Description of Unknown Organism: See Werstuck &
 Green, Science 282: 296-298, 1998. Isolated
 aptamer that binds specifically to Hoechst dye
 H33258.

<400> 28

ggtgatcaga ttctgatcca acagggttatg tagtctccta cctctgcgcc tgaagcttgg 60
 atccgtcgc 69

<210> 29
 <211> 32
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence:
 Oligonucleotide for PCR amplification

<400> 29

atatatggcc gaagaggccc ccgaagttgt ag 32

<210> 30
 <211> 36
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence:
 Oligonucleotide for PCR amplification

<400> 30

atatatctcg agtathtag accaccgtag agatgc 36

<210> 31
 <211> 63
 <212> DNA
 <213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 31

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agc 63

<210> 32

<211> 66

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 32

tctgatcacc ttaattaagc gacggatcca aagcttcagg cagctgttcc caggcagaga 60
agcata 66

<210> 33

<211> 58

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 33

gcttaattaa ggtgatcaga ttctgatcca acaggttatg tagtctccta cctctgcg 58

<210> 34

<211> 69

<212> DNA

<213> Artificial Sequence

<220>

<223> Description of Artificial Sequence:
Oligonucleotide for PCR amplification

<400> 34

atatatgcgg ccgcatatct cgaggcgacg gatccaagct tcaggcgag aggtaggaga 60
ctacataac 69

72

<210> 35
 <211> 81
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> PCR amplified oligonucleotide

<400> 35

atataggatc ccctgaaaag gctgtttaag ttgggtaaac cgctcccagc cgacgacgag 60
 caagacgaag acagaagacg c 81

<210> 36
 <211> 81
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> PCR amplified oligonucleotide

<400> 36

ccactgctaa agtgccctgtt atacctactc taaaccacgc ctttgtttca tctagcagag 60
 cgcgtcttct gtcttcgtct t 81

<210> 37
 <211> 108
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> PCR amplified oligonucleotide

<400> 37

aacaggcact ttagcagtgg ccgtgacgac ccggtatgag gtagacaata ttacacctgt 60
 cctactggca ttgagaactt ttgcccagag caaaagagca ttccaagc 108

<210> 38
 <211> 100
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> PCR amplified oligonucleotide

<400> 38

atatactcga gcttggcgga ctagactatg tcgtagtcta tttaggacca ccgtagagat 60
 gctttatttc ccctctgatg gcttggaatg ctcttttget 100

73

<210> 39
 <211> 16
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Primer for PCR amplification

<400> 39

atataggatc ccctga

16

<210> 40
 <211> 16
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Primer for PCR amplification

<400> 40

atatactcga gcttgg

16

<210> 41
 <211> 21
 <212> RNA
 <213> Artificial Sequence

<220>
 <223> Synthesized oligoribonucleotide identical with the
 bacteriophage R17 replicase initiator region

<400> 41

aaacaugagg auuacccaug u

21

<210> 42
 <211> 41
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Primer for PCR amplification

<400> 42

cagcaccatc agggctggca gcatagtaca tttcatctga c

41

<210> 43
 <211> 23

<212> DNA
<213> Artificial Sequence

<220>
<223> Primer for PCR amplification

<400> 43

cgttgtggct gttgtagttg tac

23

<210> 44
<211> 22
<212> DNA
<213> Artificial Sequence

<220>
<223> primer for PCR amplification

<400> 44

tgcggcggat ttatcttgca ag

22

<210> 45
<211> 40
<212> DNA
<213> Artificial Sequence

<220>
<223> Primer for PCR amplification

<400> 45

cagccctgat ggtgctggac tatttaggac caccgtagag

40

<210> 46
<211> 28
<212> DNA
<213> Artificial Sequence

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<210> 47
<211> 29
<212> DNA
<213> Artificial Sequence

75

<220>

<223> Primer for PCR amplification

<400> 47

atatatcatg gtggctcgag ggtggtggt

29

<210> 48

<211> 38

<212> DNA

<213> Artificial Sequence

<220>

<223> Primer for PCR amplification

<400> 48

caccatcagg gactacagca tagtacattt catctgac

38

<210> 49

<211> 37

<212> DNA

<213> Artificial Sequence

<220>

<223> Primer for PCR amplification

<400> 49

tagtccctga tggtagctat ttaggaccac cgtagag

37

<210> 50

<211> 24

<212> RNA

<213> Artificial Sequence

<220>

<223> Modified bacteriophage R17 replicase initiator
oligoribonucleotide

<400> 50

auaguccagc accaucaggg cugg

24

<210> 51

<211> 20

<212> RNA

<213> Artificial Sequence

76

<220>

<223> Modified bacteriophage R17 replicase initiator
oligoribonucleotide

<400> 51

auagucacca ucagggacua

20

CLAIMS

1. An RNA vector replicon, comprising:

- (a) a 5' sequence which initiates transcription of alphavirus RNA;
- (b) a nucleic acid sequence that codes for biologically active alphavirus nonstructural proteins;
- (c) an alphavirus subgenomic junction region promoter;
- (d) a non-alphavirus nucleotide sequence which, when bound by a ligand reduces transcription or translation of a heterologous gene of interest within the subgenomic RNA;
- (e) a heterologous gene of interest; and
- (f) a 3' alphavirus RNA polymerase recognition sequence.

2. An RNA vector replicon, comprising:

- (a) a 5' sequence which initiates transcription of alphavirus RNA;
- (b) a nucleic acid sequence that codes for biologically active alphavirus nonstructural proteins from a first alphavirus;
- (c) an alphavirus subgenomic junction region promoter;
- (d) an alphavirus nucleotide sequence which, when bound by a ligand reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA, wherein said alphavirus nucleotide sequence is from a second alphavirus different from said first alphavirus;
- (e) a heterologous gene of interest; and
- (f) a 3' alphavirus RNA polymerase recognition sequence.

3. An RNA vector, comprising:

- (a) a 5' sequence which initiates transcription of alphavirus RNA;
- (b) an alphavirus subgenomic junction region promoter,
- (c) a non-alphavirus nucleotide sequence which, when bound by a ligand reduces transcription or translation of a heterologous gene of interest within the subgenomic RNA,
- (d) a heterologous gene of interest; and
- (e) a 3' alphavirus RNA polymerase recognition sequence, wherein said RNA vector does not encode all biologically active alphavirus nonstructural proteins.

4. An RNA vector, comprising:

- (a) a 5' sequence which initiates transcription of alphavirus RNA;
- (b) an alphavirus subgenomic junction region promoter,
- (c) an alphavirus nucleotide sequence which, when bound by a ligand reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA,
- (d) a heterologous gene of interest and
- (e) a 3' alphavirus RNA polymerase recognition sequence, wherein said 5' sequence which initiates transcription and said 3' alphavirus RNA polymerase recognition sequence are from a first alphavirus, and said alphavirus nucleotide sequence is from a second alphavirus different from said first alphavirus, and wherein said RNA vector does not encode all biologically active alphavirus nonstructural proteins.

5. The RNA vector according to claim 1 or 3 wherein said non-alphavirus nucleotide sequence is a binding site for a R17 coat binding protein.

6. The RNA vector according to claim 5 wherein said non-alphavirus nucleotide sequence comprises a sequence 5TOP consisting of Sequence ID No. 50 or TOP consisting of Sequence ID No. 51.

7. The RNA vector according to claims 1 or 3 wherein said non-alphavirus nucleotide sequence is a binding site for an antibiotic.

8. The RNA vector according to claim 7 wherein said antibiotic is Tobramycin.

9. The RNA vector according to claims 1 or 3 wherein said non-alphavirus nucleotide sequence is a binding site for Hoechst dyes H33258 or H33342.

10. The RNA vector according to claim 2 or 4, wherein said alphavirus nucleotide sequence is a sequence from a subgenomic 5' end nontranslated region of Venezuelan equine encephalitis (VEE).

11. The RNA vector according to any one of claims 1 to 4 wherein said non-alphavirus nucleotide sequence or said alphavirus nucleotide sequence from a second alphavirus is positioned downstream from said alphavirus subgenomic junction region promoter.

12. The RNA vector according to any one of claims 1 to 4 wherein said non-alphavirus nucleotide sequence or said alphavirus nucleotide sequence from a second alphavirus is positioned upstream from the heterologous gene of interest.

13. The RNA vector according to claim 2 or 4 wherein said first alphavirus is Sindbis virus.

14. The RNA vector according to claim 2 or 4 wherein said first alphavirus is Semliki Forest virus.

15. The RNA vector according to claim 2 or 4 wherein said second alphavirus is VEE.

16. The RNA vector according to any one of claims 1, 2, 3 or 4, further comprising a polyadenylation tract.

17. An alphavirus vector construct, comprising a 5' promoter operably linked to a nucleic acid molecule, wherein said nucleic acid molecule is complementary DNA to the RNA vector according to any one of claims 1 to 4.

18. The alphavirus vector construct according to claim 17 wherein said promoter is a eukaryotic promoter.

19. The alphavirus vector construct according to claim 17 wherein said promoter is a bacteriophage promoter.

20. A method for reducing transcription or translation of a heterologous gene of interest encoded by subgenomic RNA of an alphavirus RNA vector replicon or alphavirus vector construct, comprising:

- (a) introducing an RNA vector according to any one of claims 1 to 4 into a cell; and
- (b) introducing into a cell (i) a ligand that reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA, or, (ii) an expression cassette which directs the expression of a ligand that reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA, such that transcription or translation of a heterologous gene of interest encoded by subgenomic RNA is reduced.

21. A method for reducing transcription or translation of a heterologous gene of interest encoded by subgenomic RNA of an alphavirus RNA vector replicon or alphavirus vector construct, comprising:

- (a) introducing an alphavirus vector construct according to claim 17 into a cell; and
- (b) introducing into a cell (i) a ligand that reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA, or, (ii) an expression cassette which directs the expression of a ligand that reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA, such that transcription or translation of a heterologous gene of interest encoded by subgenomic RNA is reduced.

22. The method according to claim 20 or claim 21 wherein said ligand is R17 coat binding protein.

23. The method according to claim 20 or claim 21 wherein said ligand is an antibiotic.

24. The method according to claim 23 wherein said antibiotic is Tobramycin.

25. The method according to claim 20 or claim 21 wherein said ligand is Hoechst dye H33258 or H33342.

26. The method according to claim 20 or claim 21 wherein said ligand is an antisense sequence.

27. A method for reducing transcription or translation of a heterologous gene of interest encoded by subgenomic RNA of an alphavirus RNA vector replicon or alphavirus vector construct, comprising:

- a) introducing an alphavirus RNA vector or alphavirus vector construct into a cell; and
- b) introducing into the cell an expression cassette which directs the expression of a ligand that reduces transcription or translation of a heterologous gene of interest encoded by the subgenomic RNA of the alphavirus RNA vector or alphavirus vector construct, such that the transcription or translation of the heterologous gene of interest encoded by the subgenomic RNA is reduced.

28. The method according to claim 27, wherein the ligand is an R17 coat protein or an antisense sequence.

Expression and Processing of the Alphavirus Structural Polypeptide

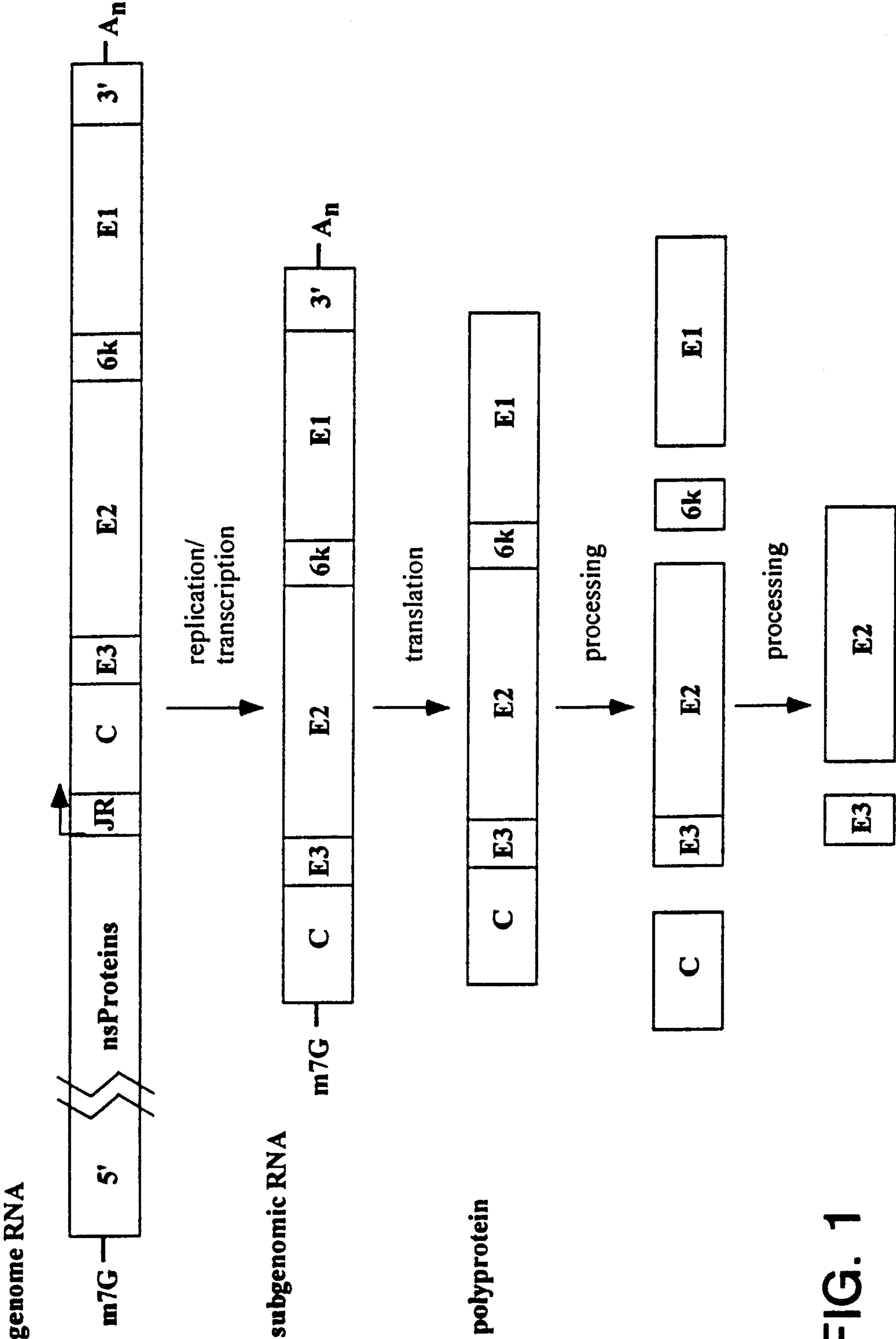


FIG. 1

Alphavirus Structural Protein Gene Open Reading Frames (ORFs)
for Vector Replicon Packaging

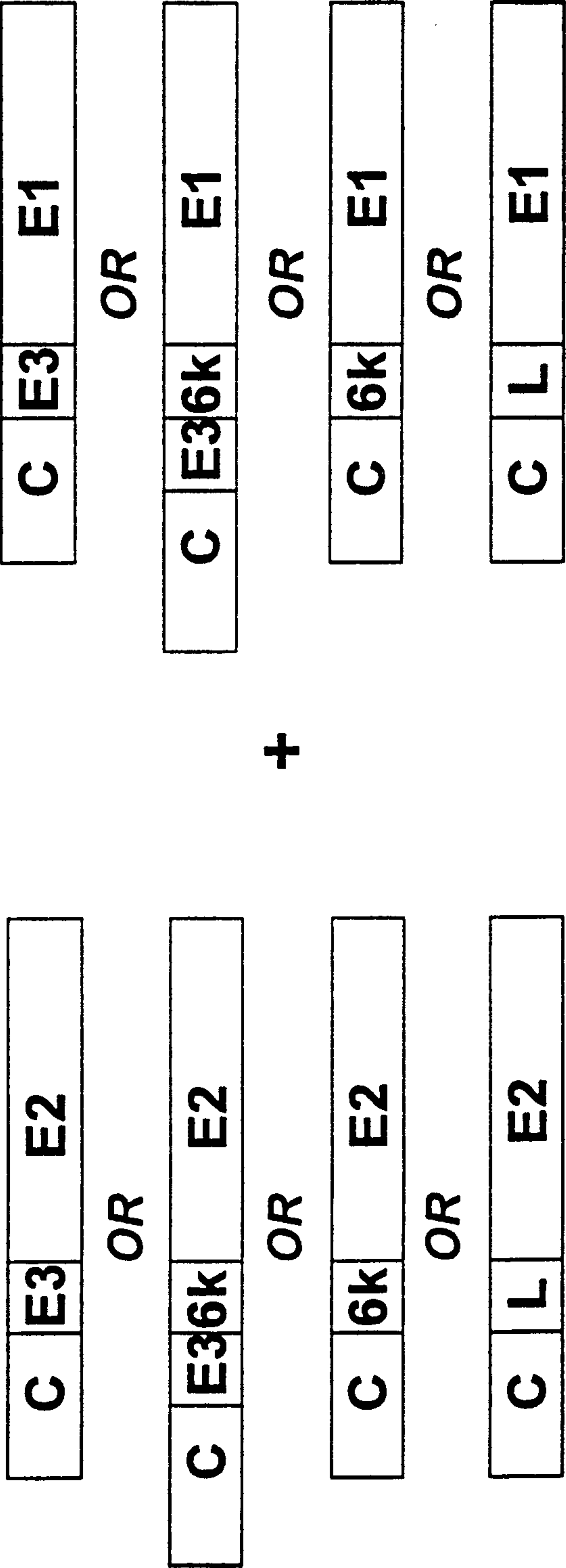


FIG. 2

Expression Cassettes Containing Alphavirus Structural Polyproteins
(representative examples)

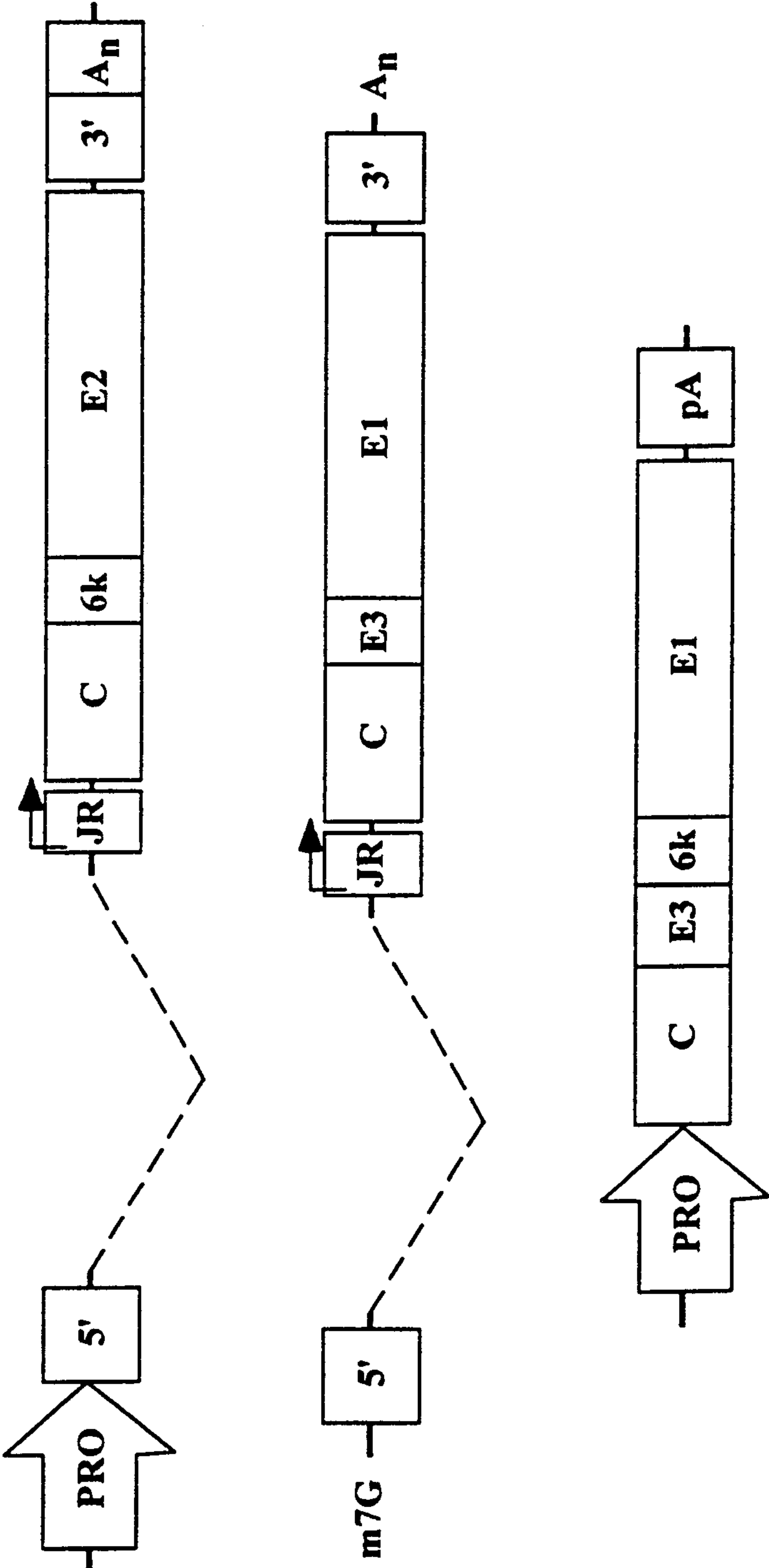
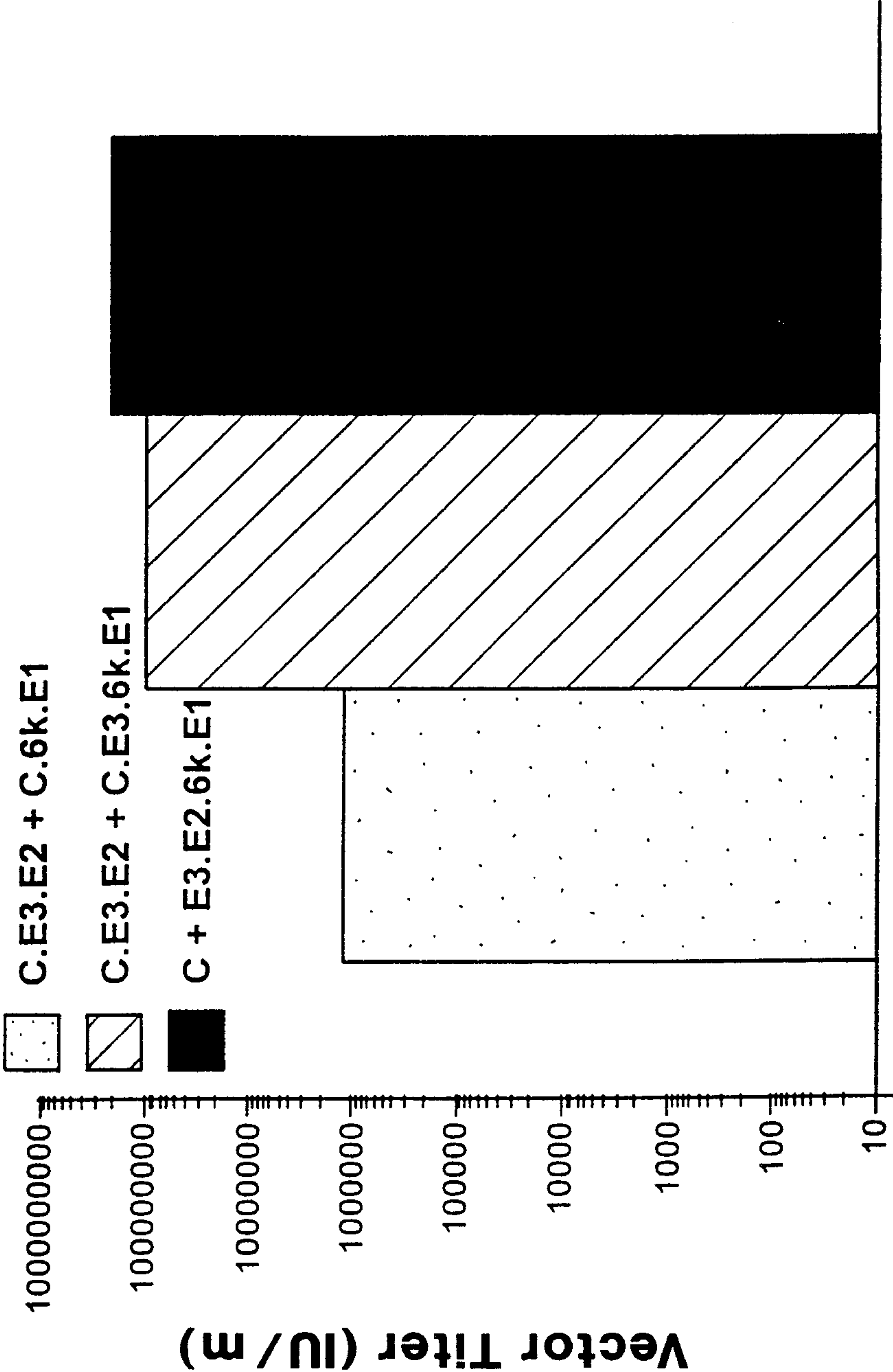


FIG. 3

Alphavirus replicon vector packaging using
various structural protein expression cassettes



Structural Protein Cassettes

FIG. 4

Reduction of Transgene Translation Using an Antisense RNA

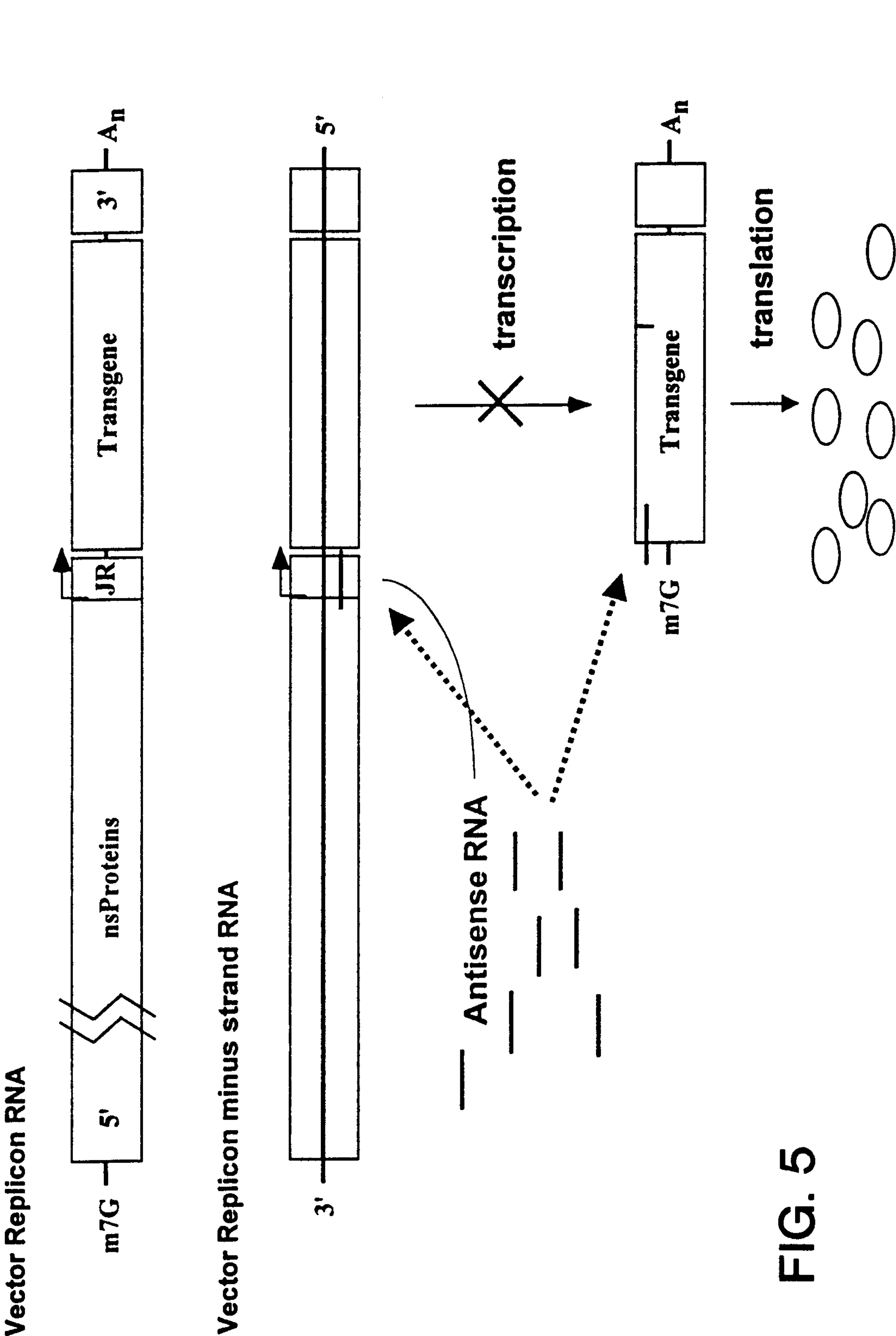


FIG. 5

Reduction of Transgene Translation Using a Specific Inserted Nucleotide Sequence and Supplied Corresponding Ligand

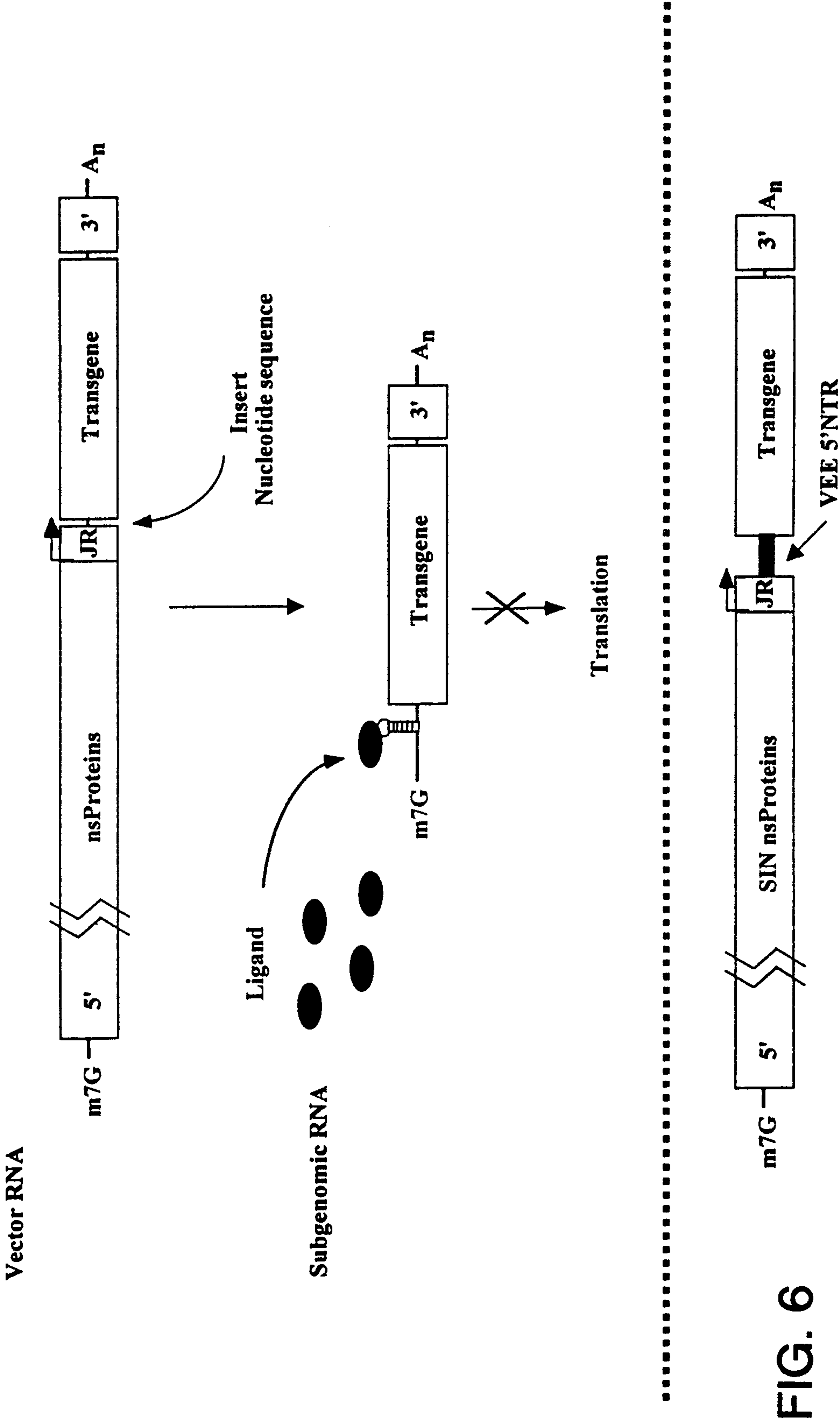
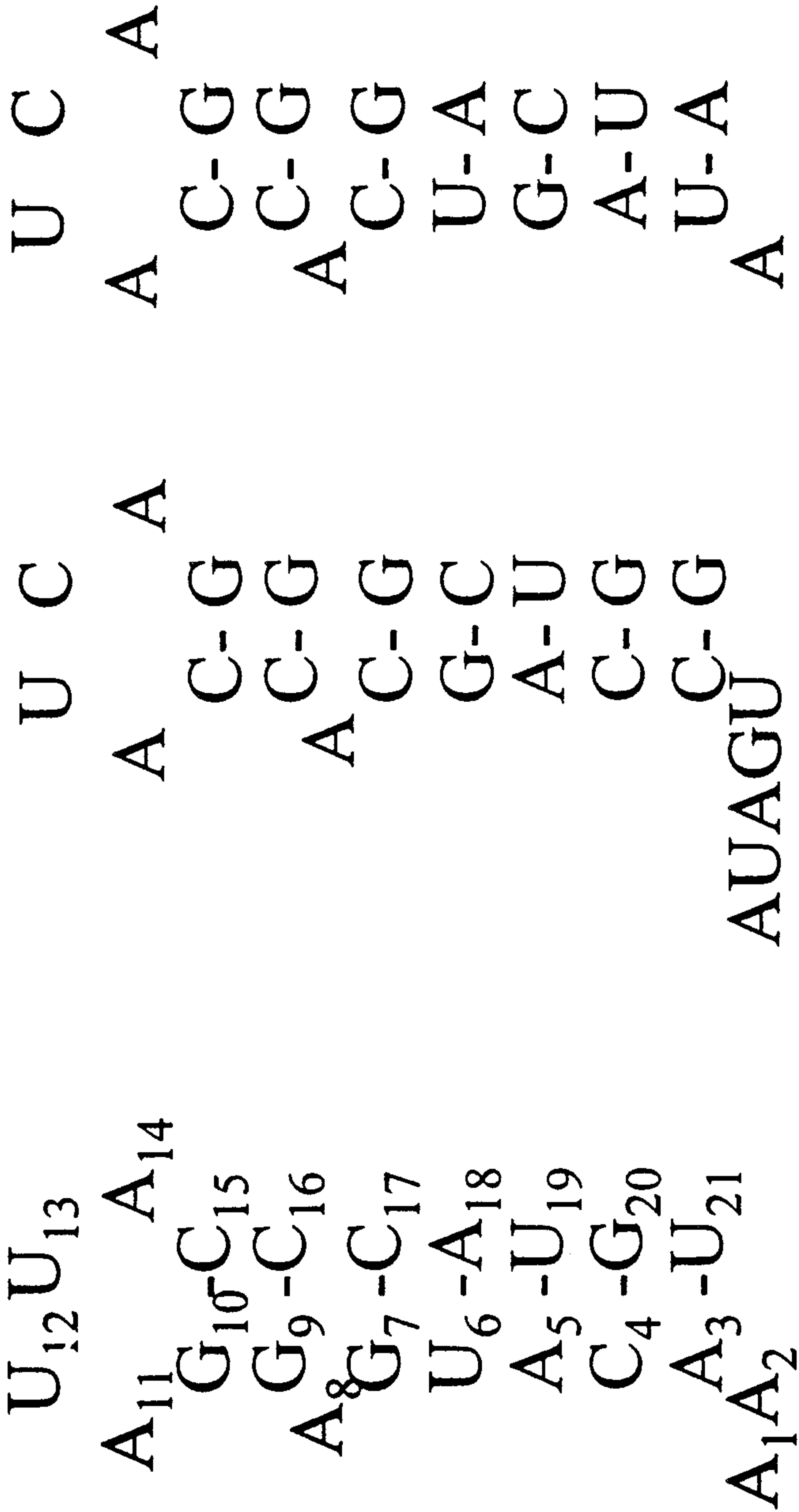


FIG. 6

Sequences for Binding of R17 Ligand



MS2 operator 5TOP TOP

FIG. 7

Reduced heterologous gene expression
from a modified alphavirus replicon

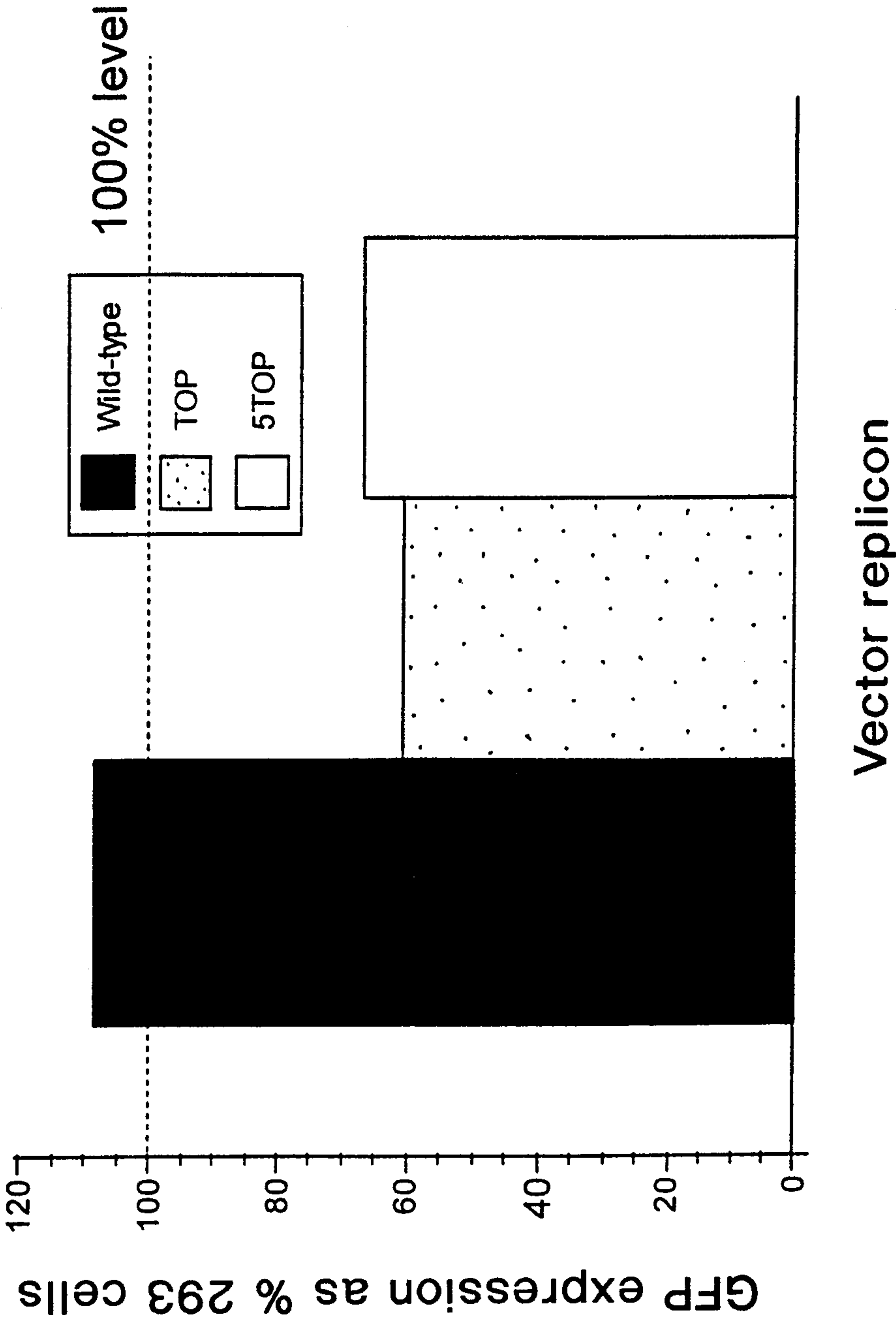
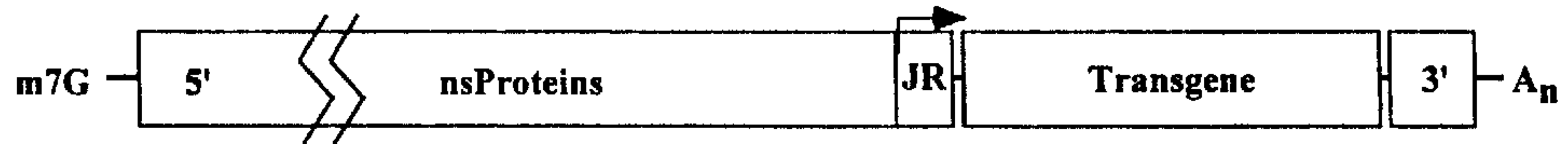


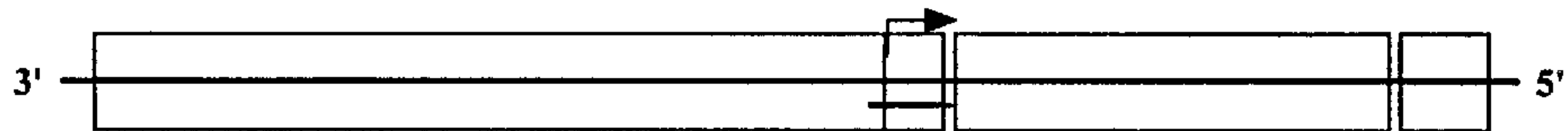
FIG. 8

Reduction of Transgene Translation Using an Antisense RNA

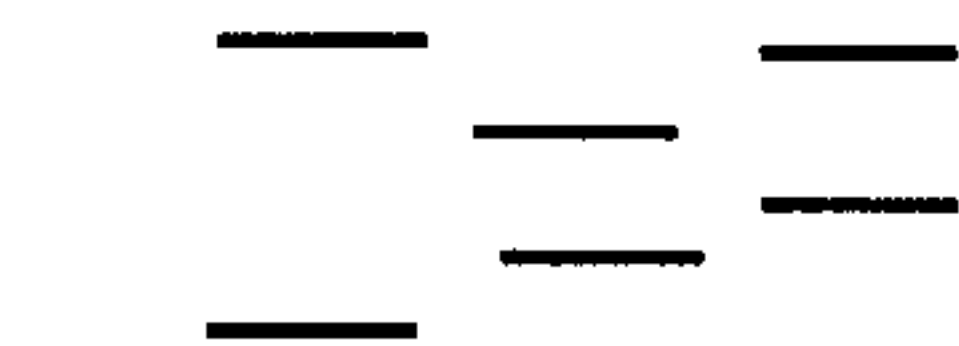
Vector Replicon RNA



Vector Replicon minus strand RNA



Antisense RNA



transcription

m7G

Transgene

A_n

translation

