

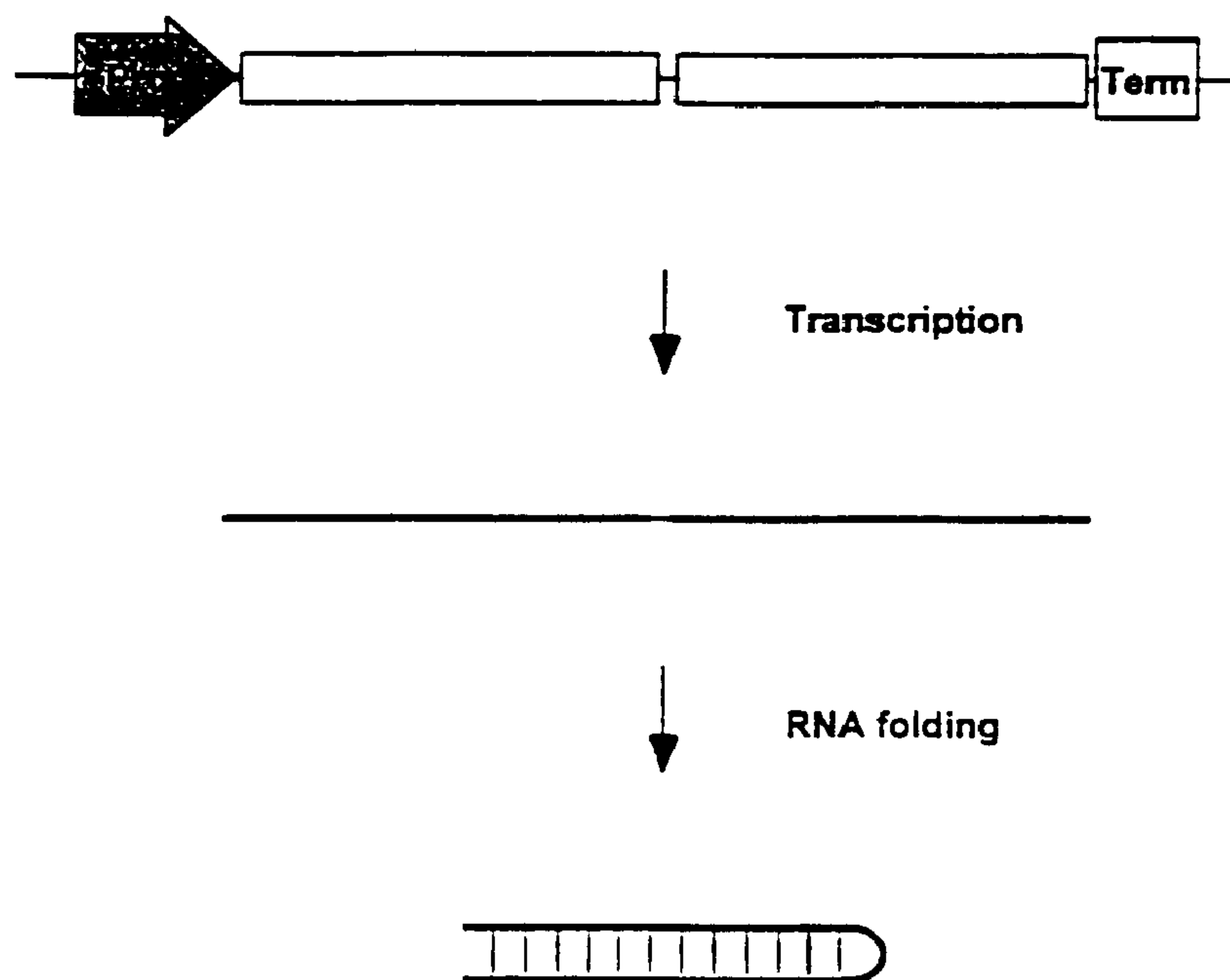


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(72) Inventeurs/Inventors:
POLO, JOHN M., US;
DUBENSKY, THOMAS W., JR., US;
BELLI, BARBARA A., US;
PERRI, SILVIA, US;
FONG, TIMOTHY, US
(73) Propriétaire/Owner:
NOVARTIS VACCINES AND DIAGNOSTICS, INC., US
(74) Agent: BORDEN LADNER GERVAIS LLP

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dsRNA Expression Cassette I: Hairpin RNA



(57) Abrégé/Abstract:

Expression cassettes are provided comprising a promoter operably linked to a nucleic acid molecule which, when transcribed in vivo, forms double-stranded RNA that induces the production of interferon. Expression cassettes also are provided comprising a

(57) Abrégé(suite)/Abstract(continued):

promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed in vivo, forms a ribozyme or antisense RNA molecule that stimulates an immune response. In addition, expression cassettes are provided comprising a promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed in vivo, stimulates apoptosis. Finally, gene delivery vectors are provided which contain such expression cassettes, host cells containing the gene delivery vectors, and methods of utilizing the expression cassettes, gene delivery vectors, and host cells.

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PERRI, Silvia [US/US]; 18426 Ogilvie Drive, Castro Valley, CA 94546 (US). **FONG, Timothy** [US/US]; 170 Sandingham South, Moraga, CA 94556 (US).

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(74) Agents: **BLACKBURN, Robert, P.**; Chiron Corporation, 4560 Horton Street, Emeryville, CA 94608-2916 et al. (US).

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(71) Applicant (*for all designated States except US*): **CHIRON CORPORATION** [US/US]; 4560 Horton Street, Emeryville, CA 94608 (US).

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(72) Inventors; and

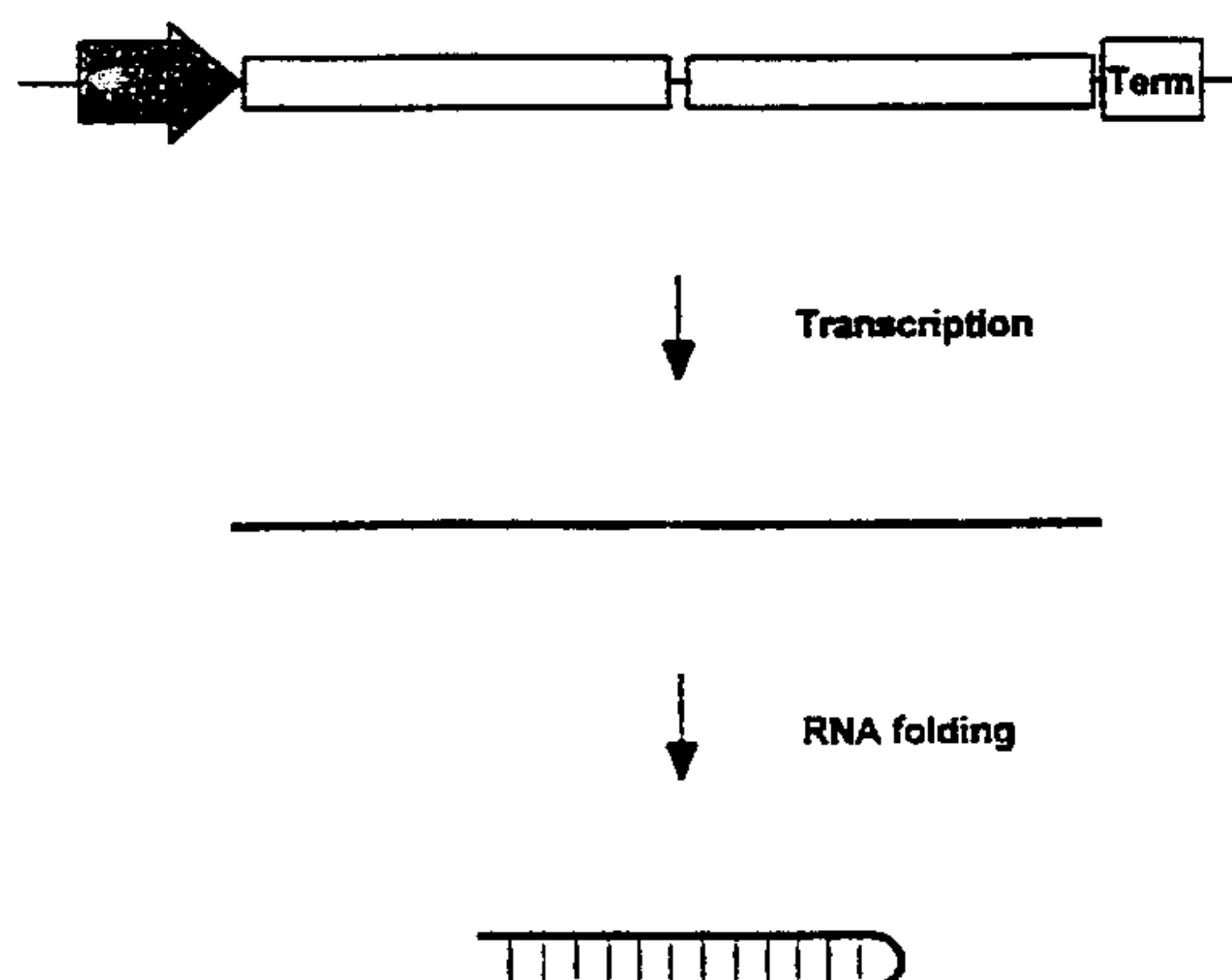
(75) Inventors/Applicants (*for US only*): **POLO, John, M.** [US/US]; 221 Witham Road, Encinitas, CA 92024 (US). **DUBENSKY, Thomas, W., Jr.** [US/US]; 6 Pacific Avenue, Piedmont, CA 94611 (US). **BELLI, Barbara, A.** [US/US]; 5850 Despejo Place, San Diego, CA 92124 (US).

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(54) Title: ENHANCEMENT OF THE IMMUNE RESPONSE FOR VACCINE AND GENE THERAPY APPLICATIONS

dsRNA Expression Cassette I: Hairpin RNA



(57) Abstract: Expression cassettes are provided comprising a promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double-stranded RNA that induces the production of interferon. Expression cassettes also are provided comprising a promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed *in vivo*, forms a ribozyme or antisense RNA molecule that stimulates an immune response. In addition, expression cassettes are provided comprising a promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed *in vivo*, stimulates apoptosis. Finally, gene delivery vectors are provided which contain such expression cassettes, host cells containing the gene delivery vectors, and methods of utilizing the expression cassettes, gene delivery vectors, and host cells.

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ENHANCEMENT OF THE IMMUNE RESPONSE FOR VACCINE AND GENE THERAPY APPLICATIONS

TECHNICAL FIELD

The present invention relates generally to pharmaceutical compositions
5 and methods, and more specifically, to a variety of compositions and methods suitable
for increasing, enhancing, or stimulating an immune response, particularly for
prophylactic and therapeutic vaccine applications in the areas of infectious disease and
cancer.

BACKGROUND OF THE INVENTION

10 Substantial research has been undertaken in order to realize the
possibility that the course of disease may be affected through the introduction of
nucleic acids into living organisms. When utilized for medical purposes, such therapy,
termed "gene therapy", allows the alteration of the genetic repertoire of cells for a
therapeutic benefit.

15 A variety of methods have been described for delivering nucleic acids to
cells, including for example, the use of viral vectors derived from retrovirus,
adenovirus, poxvirus, herpes virus, and adeno-associated virus (see Jolly, *Cancer Gene
Therapy* 1:51-64, 1994), as well as direct nucleic acid transfer techniques, including
direct DNA injection (Donnelly, J.J. et al., *Ann. Rev. Immunol.* 15:617-648, 1997),
20 micro-projectile bombardment (Williams et al., *PNAS* 88:2726-2730, 1991), and
nucleic acid complexes with lipids of several types (Felgner et al., *Proc. Natl. Acad.
Sci. USA* 84:7413-7417, 1989). Many of these strategies are being applied widely, with
clinical trials now ongoing for a wide range of hereditary (*e.g.*, ADA deficiency, cystic
fibrosis, hemophilia) and acquired (*e.g.*, cancer, viral infection) diseases (Crystal,
25 *Science* 270:404-410, 1995).

One particularly important application of gene therapy is in the vaccine
field. Briefly, live virus vaccines (*e.g.*, vaccinia, polio, measles) have been enormously
successful and have made a dramatic and historic impact on public health. However,

for certain pathogens, such as HIV, safety concerns about either reversion of attenuated vaccine strains to virulent phenotypes or inducing fulminate infection in immune compromised individuals have forced the development of subunit or inactivated virus vaccines. Unfortunately, in many cases, these vaccines have not elicited the potent
5 broad-based humoral, cellular, and mucosal immune responses or long-term memory necessary to confer life-long protection in immunized individuals. Induction of such robust immune responses will be particularly important for vaccines against HIV and HCV infection, both of which have reached worldwide epidemic proportions and have, to date, proved elusive to candidate vaccines.

10 Still relatively new, plasmid DNA-based vaccines combine the latest developments of molecular biology with an ever-increasing understanding of immunology, and show promise for immunization against infection with pathogens, such as HIV, as well as other diseases for which improved vaccines are needed (*e.g.*, influenza virus). Importantly, DNA immunization elicits humoral and cellular immune
15 responses in rodents and non-human primates. DNA is an attractive mode for vaccination, as it provides the safety advantages of subunit or inactivated virus vaccines, and induces the MHC class I-restricted cytotoxic T cell responses typical of live virus, or "replicating antigen" vaccines, which may be critical for efficacy (Donnelly, J.J. et al., *ibid*).

20 Unfortunately however, recent DNA vaccine clinical trials have demonstrated that, although antigen-specific cytotoxic T lymphocyte responses were observed in vaccinated individuals, these responses were insufficient to afford protection against challenge with the infectious agent (Wang, R. et al., *Science* 282:476-480, 1998; Calarota, S. et al., *Lancet* 351:1320-1325, 1998). Thus, for DNA-
25 based vaccination to become a widely used method for protection against infectious disease and cancer, the technology must be further improved.

The present invention provides compositions and methods for enhancing the efficacy of DNA vectors, as well as a variety of other gene delivery vectors, used for genetic immunization. Such improvements can include immunostimulatory

modifications to the antigen-expressing vector itself, or the *in vivo* administration, along with the antigen-expressing vector, of one or more additional gene delivery vectors that are immunostimulatory. The improvements described herein address previous difficulties associated with the use of gene delivery vectors by enhancing the
5 extent of the antigen-specific immune response, thus providing the broad and robust responses critical for successful prophylactic or therapeutic vaccination against infectious agents and cancer.

SUMMARY OF THE INVENTION

Briefly stated, the present invention provides compositions and methods
10 suitable for increasing, enhancing, or stimulating an immune response. For example, within one aspect of the invention expression cassettes are provided comprising a promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double stranded RNA that induces the production of interferon (alpha, beta, or gamma) either directly or indirectly, which in turn may upregulate MHC Class I and/or
15 Class II molecules, thereby stimulating an immune response.

Within one aspect of the present invention, expression cassettes are provided comprising a promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double stranded RNA that induces the production of interferon, wherein the expression cassette is selected from the group consisting of: (a)
20 an expression cassette which, when transcribed *in vivo*, forms self-complementary RNA; and (b) an expression cassette comprising a first promoter operably linked to a first nucleic acid molecule, and a second promoter operably linked to a second nucleic acid molecule, wherein the first and second nucleic acid molecules, when transcribed *in vivo*, form double stranded RNA that induces the production of interferon. Within
25 various embodiments of the invention, the RNA may be either coding or non-coding.

Within other aspects of the invention, expression cassettes are provided comprising a promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed *in vivo*, forms a ribozyme or antisense RNA molecule that stimulates an immune response. Within one embodiment, the ribozyme

or antisense molecule inhibits expression of a protein that can inhibit at least one aspect of an immune response. Representative examples of suitable proteins that may be inhibited according to the present invention include IRF2, YY1, IL-10, TGF- β and cyclooxygenase. It should be noted that inhibition of protein expression may in certain instances occur by binding and/or cleavage of mRNA by the ribozyme or antisense molecule.

Within another aspect, expression cassettes are provided comprising a promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed *in vivo*, stimulates apoptosis. Representative examples of suitable ribozymes or antisense molecules include those that cleave or inhibit targets within an apoptosis cascade, including for example, Bcl-2 and Bcl-xL.

Within other aspects, expression cassettes are provided comprising a first promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double stranded RNA that induces the production of interferon, and a second promoter operably linked to a nucleic acid molecule that encodes an antigen from a pathogenic agent. Within yet other aspects, expression cassettes are provided comprising a first promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed *in vivo*, stimulates apoptosis, and a second promoter operably linked to a nucleic acid molecule that encodes an antigen from a pathogenic agent. Within certain embodiments, the pathogenic agent is a virus, bacteria, fungus, parasite or cancerous cell. In a further aspect, expression cassettes are provided which direct the expression of both a polypeptide that promotes apoptosis and an antigen from a pathogenic agent. Within certain embodiments, the antigen from the pathogenic agent and the polypeptide that promotes apoptosis are expressed from separate promoters. Representative examples of pro-apoptotic polypeptides include Bax, Bik, Bcl-xL. Alternatively, the polypeptide and antigen can be transcribed from the same promoter and expressed by an Internal Ribosome Entry Site or ribosome readthrough.

Within various embodiments, any of the above expression cassettes may contain an RNA polymerase I, RNA polymerase II (*e.g.*, CMV, SV40, MoMLV LTR and RSV LTR), or RNA polymerase III promoter (*e.g.*, an Adenovirus VA1 promoter).

Within certain embodiments of the invention, the above noted
5 expression cassettes may further comprise an additional promoter operably linked to a nucleic acid molecule that encodes a polypeptide of interest. Within certain embodiments, the promoter is a RNA polymerase II promoter (*e.g.*, CMV, SV40, MoMLV LTR and RSV LTR). A wide variety of polypeptides may be expressed, including for example polypeptides that promote apoptosis, antigens from a pathogenic
10 agent, prodrug converting enzymes, and cytokines (*e.g.*, gamma interferon, IL-2, IL-12, and IL-15). Pathogenic agents can include viruses (*e.g.*, HIV, HSV, HBV, HCV, HPV, and FIV), bacteria, parasites, fungus and tumors.

Within related aspects, the present invention further provides gene delivery vectors that contain one or more of the expression cassettes described herein.
15 Representative examples of gene delivery vectors include plasmids and recombinant viral vectors derived from retrovirus, herpesvirus, poxvirus, adenovirus, AAV, parvovirus, alphavirus, polyoma virus, and vesiculovirus.

Expression cassettes and gene delivery vehicles as described herein can be transformed, transfected or otherwise introduced into a wide variety of host cells,
20 including for example, a wide variety of vertebrate cells (*e.g.*, mammalian cells such as human, non-human primate, horse, dog, mouse).

Expression cassettes and gene delivery vehicles as provided herein can be utilized to stimulate an immune response within a desired host, comprising the general step of administering to an animal an expression cassette or gene delivery
25 vector as described herein, such that an immune response is generated.

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 set forth herein which describe in more detail certain procedures or compositions (*e.g.*, DNA constructs), and are therefore incorporated by reference in
30 their entirety.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic illustration of one class of dsRNA expression cassette that produces RNA with a hairpin.

Figure 2 is a schematic illustration of one class of dsRNA expression cassette that produces true double-stranded RNA.

Figure 3 is a schematic illustration of one class of dsRNA expression cassette that produces circular RNA.

Figure 4 is a schematic illustration of DNA vectors with double-stranded RNA and antigen cassettes.

Figures 5A and 5B are schematic illustrations of plasmid vectors with pro-apoptotic and antigen cassettes.

Figure 6 depicts several target cleavage sites in IRF2, YY1, IL-10 and Cox2 mRNA.

DETAILED DESCRIPTION OF THE INVENTION

DEFINITIONS

Prior to setting forth the invention, it may be helpful to an understanding thereof to first set forth definitions of certain terms that will be used hereinafter.

"Expression cassette" refers to a transcription unit comprising a promoter operably linked to a nucleic acid molecule, such that transcription from the promoter results in synthesis of an RNA corresponding to the nucleic acid molecule. The expression cassette may further comprise a transcription termination and/or polyadenylation site.

"Isolated nucleic acid molecule" is a nucleic acid molecule that is not integrated in the genomic DNA of an organism or cell(s) derived from an organism. For example, a DNA molecule encoding a gene that has been separated from the genomic DNA of a eukaryotic cell is an isolated DNA molecule. Other examples of

isolated nucleic acid molecules include chemically synthesized nucleic acid molecules, and nucleic acid molecules that have been made by recombinant means (*e.g.*, PCR).

“Recombinant host” refers to any prokaryotic or eukaryotic cell that contains either a gene delivery vector or expression vector. This term also includes
5 those prokaryotic or eukaryotic cells that have been genetically engineered to contain the cloned gene(s) in the chromosome or genome of the host cell.

An “antisense oligonucleotide” or “antisense molecule” refers to an oligonucleotide having a sequence (a) capable of forming a stable triplex with a portion of the gene, or (b) capable of forming a stable duplex with a portion of a mRNA
10 transcript.

“Ribozyme” refers to a nucleic acid molecule that is capable of cleaving a specific ribonucleic acid sequence. Ribozymes may be composed of RNA, DNA, nucleic acid analogues (*e.g.*, phosphorothioates), or any combination of these (*e.g.*, DNA/RNA chimerics). Within particularly preferred embodiments, a ribozyme should
15 be understood to refer to RNA molecules that contain antisense sequences for specific recognition, and an RNA-cleaving enzymatic activity.

“Ribozyme gene” refers to a nucleic acid molecule (*e.g.*, DNA) consisting of the ribozyme sequence which, when transcribed into RNA, will yield the ribozyme.

20 “Promoter” is a nucleotide sequence that directs the transcription of a gene or non-protein encoding RNA. Typically, a promoter is located in the 5’ region adjacent to the coding sequence of a gene or RNA to be expressed, and proximal to the transcription start site. If a promoter is an inducible promoter, then the level of transcription increases in response to an inducing agent.

25 “Gene delivery vector” or “GDV” is a vehicle for delivering nucleic acid sequences to cells. GDVs include viral vectors, nucleic acid vectors such as plasmids, naked nucleic acid molecules such as DNA or RNA, nucleic acid molecules complexed to one or more polycationic molecules capable of neutralizing part or all of the negative charge on a nucleic acid molecule and condensing the nucleic acid molecule into a
30 compact structure, nucleic acid associated with liposomes, bacteria, and certain

eukaryotic cells such as producer cells, capable of delivering a nucleic acid molecule having one or more desirable properties to host cells in an organism. As discussed below, the desirable properties include the ability to express a desired substance, such as a protein (*e.g.*, an enzyme, antibody, antibody fragment, regulatory factor, ligand) or
5 nucleic acid molecule (*e.g.*, an antisense RNA, sense RNA, ribozyme), wherein the nucleic acid molecule carried by the GDV may itself be the active agent without requiring the expression of a desired substance. One example of a nucleic acid molecule is an antisense molecule that binds to mRNA and inhibits translation. Another example, is where the nucleic acid molecule encodes a ribozyme that binds and
10 cleaves mRNA thereby inhibiting translation.

"Recombinant retroviral vector" refers to a gene delivery vector which is capable of directing the expression of a sequence(s) or gene(s) of interest. Preferably, the recombinant retroviral vector should include a 5' LTR, a tRNA binding site, a packaging signal, one or more heterologous sequences, an origin of second strand DNA
15 synthesis and a 3' LTR. The recombinant retroviral vector also may include a transcriptional promoter/enhancer, or other elements which control gene expression by means such as alternate splicing, nuclear RNA export, post-translational modification of messenger, or post-transcriptional modification of protein. Optionally, the recombinant retroviral vector may also include selectable markers that confer, to
20 transduced or transfected cells, resistance or sensitivity to one or more compounds.

The present invention relates to the introduction of nucleic acid sequences into a vertebrate to achieve expression of molecules (*e.g.*, protein, RNA) that directly or indirectly stimulate one or more components of the immune response. The molecules may be expressed from any number of different delivery systems, either
25 alone or co-expressed with an antigenic or therapeutic polypeptide.

Thus, the present invention can be utilized for a variety of therapeutic purposes, including for example, for the purpose of prophylactic and therapeutic vaccination, and for other gene therapy applications. In order to further an understanding of the inventions provided herein, a more detailed discussion is provided

of (A) expression cassettes; (B) generation of gene delivery vectors; and (c) methods for utilizing expression cassettes and gene delivery vectors for a variety of applications (*e.g.*, for therapeutic purpose, or, for use in laboratory setting).

A. GENERATION OF EXPRESSION CASSETTES

5 As noted above, the present invention provides a variety of expression cassettes that direct the generation of RNA, as well as optionally, a selected gene of interest, which increases, enhances, or stimulates an immune response. As described in more detail below, expression cassettes (comprising a promoter operably linked to a nucleic acid molecule of interest) can be generated to express double-stranded RNA,
10 ribozymes and antisense RNA, as well as antigens or therapeutic polypeptides of interest. Each of these is discussed in more detail below.

Briefly, through use of the expression cassettes and/or gene delivery vectors described herein, double-stranded RNA (dsRNA) can be produced intracellularly for a variety of applications. The intracellular presence of dsRNA
15 induces the synthesis of interferons α , β , and γ (IFN- α , IFN- β , and IFN- γ ; Type I interferon: IFN- α and IFN- β ; Type II interferon: IFN- γ). Production of Type I and II interferons in turn induces both an antigen-specific and non-specific immune response, by directly stimulating CD8+ cytotoxic T lymphocytes (CTL), natural killer (NK) cells, and monocytes/macrophages. Furthermore these interferons enhance antigen
20 presentation through increased expression of MHC class I and class II on the cell surface. Compositions and methods are provided herein for the expression of non-coding dsRNA in the context of expression of a desired antigen. Expression of dsRNA according to the invention enhances the overall robustness of antigen-specific immune responses.

25 Induction of Type I and II interferons, as a result of the intracellular presence of dsRNA in turn induces the synthesis of protein kinase R (PKR), and 2', 5' oligoadenylate synthetase (2',5' OAS) (Jacobs and Langland, *Virology* 219:339-349, 1996). Activation of PKR and 2', 5' OAS can in turn lead to apoptosis in a given cell, when the ratio of death agonists, for example Bax and Bak, to death antagonists, for

example Bcl-2, exceeds unity (for reviews, see Kroemer, *Nat. Med.* 6:614-620, 1997; and Adams and Cory, *Science* 281:1322-1326, 1998). Activated PKR can directly effect this ratio through phosphorylation and inactivation of the translation factor eif-2 α and I κ B. Phosphorylation of I κ B by PKR can lead to I κ B degradation and subsequent
5 activation of NF- κ B (Maran, *Virology* 164:106-113, 1994). NF- κ B is a transcription factor that activates transcription of genes for inflammatory cytokines, including TNF α (Gilmore, *Cancer Biol.* 8:61-62, 1997). Induction of TNF α can induce apoptosis through the CrmA/P35-sensitive caspase-dependent pathway (Thornberry and Lazebnik, *Science* 281:1312-1316, 1998).

10 Induction of Type I and II interferons and apoptosis in the context of antigen expression enhances the specific immune response, through efficient priming of, for example, dendritic cells (DCs). As antigen presenting cells, DCs phagocytize apoptotic cells, and process antigens into peptides in the cytosol, which in turn bind to MHC class I or class II peptides, and are presented to T lymphocytes (Banchereau and
15 Steinman, *Nature* 392:245-252, 1998). Induction of apoptosis, or the death pathway, in the context of antigen expression in order to enhance the immune response can be accomplished by a variety of different methods including, for example, expression of either caspase-dependent or caspase-independent genes. expression of death receptors (e.g., CD95 or TNFR1).

20 Representative examples of suitable expression cassettes are described in more detail below, as well as in the figures (see Figures 1 – 3 for schematic illustrations of expression cassettes which create dsRNA through the use of hairpins (Figure 1), true dsRNA (Figure 2), and circular RNA (Figure 3).

A wide variety of nucleic acid molecules may be expressed from the
25 expression cassettes and/or gene delivery vectors of the present invention. Representative examples include (1) antisense RNA; (2) ribozymes; (3) polypeptides, including antigens and therapeutic proteins, and apoptotic polypeptides.

1. Antisense RNA

Within one embodiment of the invention delivery systems are provided which produce an antisense RNA sequence upon introduction into a host cell. Briefly, antisense RNA sequences are designed to bind to target RNA transcripts, and thereby
5 prevent cellular synthesis of a particular protein or prevent use of that RNA sequence by the cell. In preferred embodiments, target RNA transcripts encode gene products that control or regulate various aspects of the immune response or apoptotic pathway.

Antisense sequences of the present invention may be of a variety of sizes, including for example 40, 60, 100, 250, 500, or even 1000 nucleotides. For
10 preferred embodiments, the antisense sequence is less than 100 nucleotides. The antisense RNA sequence also may be 100% complementary to the target sequence, or may contain one or more "mismatches" that are not complementary provided that the antisense RNA maintains its ability to bind the target sequence under physiological conditions of the host.

Representative examples of such antisense sequences suitable for use
15 within the present invention include antisense IRF2, YYK1, Cox2, IL-10, Bcl-2, Bcl-xL, and TGF- β . The antisense sequence may also be an antisense RNA complementary to RNA sequences necessary for pathogenicity. Alternatively, the biologically active nucleic acid molecule may be a sense RNA (or DNA) complementary to RNA
20 sequences necessary for pathogenicity

2. Ribozymes

Within other aspects of the present invention, delivery systems are provided which produce ribozymes upon introduction into a host cell. Briefly, ribozymes are RNA molecules used to cleave specific RNAs and are designed such that
25 it can only affect one specific RNA sequence. Generally, the substrate binding sequence of a ribozyme typically is between 10 and 20 nucleotides long, although longer sequences also may be utilized. The length of this sequence is sufficient to allow hybridization with the target RNA and disassociation of the ribozyme from the cleaved RNA. Several different types of ribozymes may be constructed for use within
30 the present invention, including for example, hammerhead ribozymes (Rossi, J.J. et al.,

Pharmac. Ther. 50:245-254, 1991) (Forster and Symons, *Cell* 48:211-220, 1987; Haseloff and Gerlach, *Nature* 328:596-600, 1988; Walbot and Bruening, *Nature* 334:196, 1988; Haseloff and Gerlach, *Nature* 334:585, 1988; Haseloff et al., U.S. Patent No. 5,254,678), hairpin ribozymes (Hampel et al., *Nucl. Acids Res.* 18:299-304, 5 1990, and U.S. Patent No. 5,254,678), hepatitis delta virus ribozymes (Perrotta and Been, *Biochem.* 31:16, 1992), Group I intron ribozymes (Cech et al., U.S. Patent No. 4,987,071) and RNase P ribozymes (Takada et al., *Cell* 35:849, 1983); (see also, WO 95/29241, entitled "Ribozymes with Product Ejection by Strand Displacement"; and WO 95/31551, entitled "Novel Enzymatic RNA Molecules." (see also U.S. Patent Nos. 10 5,116,742; 5,225,337 and 5,246,921). Preferred ribozymes for use within the present invention include those which target RNA transcripts encoding gene products that control or regulate various aspects of the immune response or apoptotic pathway. Particularly preferred ribozymes are disclosed in more detail below.

15 3. Co-Expression of Antigens, Therapeutic Polypeptides, and Apoptotic Genes

In addition to the immunostimulatory RNAs of the present invention, a wide variety of genes encoding antigenic or therapeutic polypeptides also may be included in such delivery systems, either by co-expression from one vector or co-administration of separate vectors, including for example, sequences which encode 20 palliatives such as lymphokines, toxins, or prodrugs, antigens which stimulate an immune response, and proteins which modulate the immune response or apoptotic pathway. Within various embodiments of the invention, the delivery systems provided herein may contain genes for two or more antigenic or therapeutic polypeptides.

a. Antigens

25 Within the present invention, constructs are provided which direct the expression of immunogenic portions of antigens from foreign organisms or other pathogenic agents (e.g., cancerous cells). Preferred vaccine antigens from foreign organisms are those that protect warm-blooded animals from diseases caused by infectious agents such as bacteria, fungi, parasites, and viruses.

Representative examples of such antigens particularly relevant for human vaccines include: viral antigens from arenaviruses such as lassa fever virus; rotavirus; adenovirus; human papillomavirus; retroviruses such as HIV, HTLV-I and -II; herpes viruses such as herpes simplex virus, Epstein-Barr virus, cytomegalovirus, varicella-zoster virus, and human herpes virus 6; picornaviruses such as poliovirus, human rhinovirus, and hepatitis A virus; togaviruses such as Venezuelan equine encephalitis, and rubella virus; hepatitis B virus; flaviviruses such as Dengue virus, tick-borne encephalitis virus, and hepatitis C virus; human coronavirus; rhabdoviruses such as rabies virus, and vesicular stomatitis virus; filoviruses such as ebola virus and marburg virus; paramyxoviruses such as human parainfluenza virus, mumps virus, respiratory syncytial virus, and measles virus; influenza virus; bunyaviruses such as hantaan virus, and Rift Valley fever virus; bacterial antigens (e.g., from *E. coli*, streptococcus, staphylococcus, mycobacteria; meningococcus or helicobacter) fungal antigens and parasitic antigens (e.g. from *Leishmania* or malaria).

Representative examples of antigens particularly relevant for veterinary applications are derived from *Toxoplasma*, *Dirofilaria*, *Acanthocheilonema*, *Babesia*, *Brugia*, *Candida*, *Cryptococcus*, *Cryptosporidium*, *Dipetalonema*, *Eimeria*, *Encephalitozoon*, *Hepatozoon*, *Histoplasma*, *Isospora*, *Loa*, *Microsporidia*, *Neospora*, *Nosema*, *Onchocerca*, *Parafilaria*, *Plasmodium*, *Pneumocystis*, *Rochalimaea*, *Setaria*, *Stephanofilaria*, *Theileria*, *Toxoplasma*, and *Wuchereria*. viruses such as equine herpes virus, feline rhinotracheitis virus, porcine rotavirus, blue tongue virus, FIV, FeLV, BIV, EIAV, bovine papillomavirus, canine parvovirus, feline panleukopenia virus, foot-and-mouth disease virus, porcine enterovirus, TGEV, feline infectious peritonitis virus, bovine coronavirus, rabies virus, canine distemper virus.

As utilized within the context of the present invention, "immunogenic portion" refers to a portion of the respective antigen which is capable, under the appropriate conditions, of inducing an immune response (*i.e.*, cell-mediated or humoral). "Portions" may be of variable size, but are preferably at least 9 amino acids long, and may include the entire antigen.

b. Therapeutic Polypeptides

Within one embodiment of the invention, expression cassettes and gene delivery vectors are provided which direct the expression of a gene product(s) that activates a compound with little or no cytotoxicity into a toxic product. Briefly, a wide
5 variety of gene products which either directly or indirectly activate a compound with little or no cytotoxicity into a toxic product may be utilized within the context of the present invention. Representative examples of such gene products include HSVTK and VZVTk which selectively monophosphorylate certain purine arabinosides and substituted pyrimidine compounds, converting them to cytotoxic or cytostatic
10 metabolites. More specifically, exposure of the drugs ganciclovir, acyclovir, or any of their analogues (*e.g.*, FIAC, DHPG) to HSVTK, phosphorylates the drug into its corresponding active nucleotide triphosphate form.

Other therapeutic polypeptides that can be expressed from the cassettes and gene delivery vectors of the present invention include cytokines, such as IL-1, IL-2
15 (Karupiah et al., *J. Immunology* 144:290-298, 1990; Weber et al., *J. Exp. Med.* 166:1716-1733, 1987; Gansbacher et al., *J. Exp. Med.* 172:1217-1224, 1990; U.S. Patent No. 4,738,927), IL-3, IL-4 (Tepper et al., *Cell* 57:503-512, 1989; Golumbek et al., *Science* 254:713-716, 1991; U.S. Patent No. 5,017,691), IL-5, IL-6 (Brakenhof et al., *J. Immunol.* 139:4116-4121, 1987; WO 90/06370), IL-7 (U.S. Patent
20 No. 4,965,195), IL-8, IL-9, IL-10, IL-11, IL-12, IL-13 (*Cytokine Bulletin*, Summer 1994), IL-14 and IL-15, particularly IL-2, IL-4, IL-6, IL-12, and IL-13, alpha interferon (Finter et al., *Drugs* 42(5):749-765, 1991; U.S. Patent No. 4,892,743; U.S. Patent No. 4,966,843; WO 85/02862; Nagata et al., *Nature* 284:316-320, 1980; Familletti et al., *Methods in Enz.* 78:387-394, 1981; Twu et al., *PNAS* 86:2046-2050, 1989; Faktor
25 et al., *Oncogene* 5:867-872, 1990), beta interferon (Seif et al., *J. Virol.* 65:664-671, 1991), gamma interferons (Radford et al., *The American Society of Hepatology* 2008-2015, 1991; Watanabe et al., *PNAS* 86:9456-9460, 1989; Gansbacher et al., *Cancer Research* 50:7820-7825, 1990; Maio et al., *Can. Immunol. Immunother.* 30:34-42, 1989; U.S. Patent No. 4,762,791; U.S. Patent No. 4,727,138), G-CSF (U.S. Patent

Nos. 4,999,291 and 4,810,643), GM-CSF (WO 85/04188), tumor necrosis factors (TNFs) (Jayaraman et al., *J. Immunology* 144:942-951, 1990), CD3 (Krissanen et al., *Immunogenetics* 26:258-266, 1987), ICAM-1 (Altman et al., *Nature* 338:512-514, 1989; Simmons et al., *Nature* 331:624-627, 1988), ICAM-2, LFA-1, LFA-3 (Wallner et al., *J. Exp. Med.* 166(4):923-932, 1987), MHC class I molecules, MHC class II molecules. B7.1-3, b₂-microglobulin (Parnes et al., *PNAS* 78:2253-2257, 1981), chaperones such as calnexin, MHC linked transporter proteins or analogs thereof (Powis et al., *Nature* 354:528-531, 1991).

c. Apoptotic genes

10 Apoptosis is the normal process of programmed cell death, which maintains tissue homeostasis (Kerr, J. F. et al., *Br. J. Cancer* 26:239-257, 1972). The process of apoptosis is regulated by signals generated when cytokines bind to their receptors, which in turn may either produce inductive or inhibitory signals that regulate the apoptotic process. Upon induction of apoptosis a cascade of zymogen activation
15 occurs which eventually leads to the activation of endogenous nucleases that cleave chromatin between nucleosomes and reduce the content of intact DNA in apoptotic cells, thereby leading to cell death. While apoptosis is mediated by a variety of signals and complex interactions of cellular gene products, the results of these interactions ultimately feed into a cell death pathway that is, itself, a cascade of zymogen activation
20 (*i.e.*, proteolytic events) analogous to that of the blood coagulation cascade.

Several gene families and products that modulate the apoptotic process have now been identified. One such family is the Bcl-2 family, which includes cell death inhibitory proteins Bcl-2, Bcl-xL, and CED-9 as well as apoptotic promoting proteins, bax, bak, bcl-xS, and bad. Bcl-2 was the first recognized component of
25 programmed cell death (Tsujimoto et al., *Science* 228:1440-1443, 1985; Nunez et al., *J. Immunol.* 144:3602-3610, 1990; Reed, *Nature* 387:773, 1997).

Briefly, Bcl-2 and its homologue Bcl-xL are death antagonists that associate mainly with the outer mitochondrial membrane, the endoplasmic reticulum, and nuclear envelope and moreover, have documented ion channel activity (Reed,
30 *Nature* 387:773, 1997). These proteins may prevent apoptosis by regulating the

electrical and osmotic homeostasis of the mitochondria, a process that is required to prevent mitochondrial swelling, outer membrane rupture, and subsequent cytochrome c release (Heiden et al., *Cell* 91:627-637, 1997). Release of cytochrome c from the mitochondria is believed to be the triggering event of the cell death cysteine protease cascade, through formation of the Apaf-1/caspase-9/cytochrome c complex (Li et al.,
5 *Cell* 91:479-489, 1997; Reed, *Cell* 91:559-562, 1997). This complex in combination with dATP in turn leads to the proteolytic activation of the caspase-9 zymogen thereby initiating the proteolytic cascade.

The cysteine proteases responsible for this cascade are aspartate-specific
10 cysteine proteases and are collectively referred to as "caspase" proteins. The first member of the caspase family was identified as Interleukin-1 β Converting Enzyme (ICE) which was found to be structurally and functionally related to the CED-3 protease that functions as a cell death effector in the roundworm *C. elegans* (Yuan et al., *Cell* 75:641, 1993). Following this discovery many other members of the
15 ICE/CED-3 family were identified and include, for example, human ICE (interleukin-1- β converting enzyme) (caspase-1), ICH-1 (caspase-2), CPP32 (caspase-3), ICE_{rel}II (caspase-4), ICE_{rel}III (caspase-5), Mch2 (caspase-6), ICE-LAP3 (caspase-7), Mch5 (caspase-8), ICE-LAP6 (caspase-9), Mch4 (caspase-10), and others.

Given that the apoptotic process functions to maintain tissue
20 homeostasis, a variety of pathological states can be associated with abnormal apoptotic regulation. Further, the loss or inhibition of apoptosis can lead to accumulation of virally infected cells or hyperproliferative cells such as tumor cells. Similarly, the inappropriate activation of apoptosis can also contribute to a variety of pathological disease states including, for example, ischemic injury. Accordingly, treatments which
25 are specifically designed to modulate the apoptotic pathways in these and other pathological conditions can alter the natural progression of many of these diseases.

4. *Obtaining a desired nucleic acid molecule of interest*

Sequences that encode the above-described antisense or ribozyme sequences, antigens, therapeutic polypeptides and apoptotic genes may be generated synthetically or recombinantly, or, obtained from a variety of sources, including for example, depositories such as the American Type Culture Collection (ATCC, Rockville, MD). Isolated nucleic acid molecules encoding such antigens are readily generated by one of skill in the art using standard PCR methodologies, provided that at least some sequence data are available. Alternatively, cDNA sequences that encode the above-described antigens may be obtained from cells or cultures that express or contain the sequences. Briefly, within one embodiment, mRNA from a cell which expresses the gene of interest is reverse transcribed with reverse transcriptase using oligonucleotide dT or random primers. The single stranded cDNA may then be amplified by PCR (*see* U.S. Patent Nos. 4,683,202; 4,683,195 and 4,800,159. *See also PCR Technology: Principles and Applications for DNA Amplification*, Erlich (ed.), Stockton Press, 1989) utilizing oligonucleotide primers complementary to sequences on either side of desired sequences. In particular, a double-stranded DNA is denatured by heating in the presence of heat stable Taq polymerase, sequence-specific DNA primers, dATP, dCTP, dGTP and dTTP. Double-stranded DNA is produced when synthesis is complete. This cycle may be repeated many times, resulting in a factorial amplification of the desired DNA.

Sequences encoding the above-described antigens also may be synthesized, for example, on an Applied Biosystems Inc. DNA synthesizer (Applied Biosystems, Foster City, CA).

B. PREPARATION OF GENE DELIVERY VECTORS

A wide variety of gene delivery vectors may be utilized to contain and express the expression cassettes described herein. The gene delivery vector may be of either viral or non-viral origin (*see* generally, Jolly, *Cancer Gene Therapy* 1:51-64, 1994; Kimura, *Human Gene Therapy* 5:845-852, 1994; Connelly, *Human Gene Therapy* 6:185-193, 1995; and Kaplitt, *Nature Genetics* 6:148-153, 1994; Donnelly et

al., *ibid*). Gene therapy vehicles for delivery of constructs including a coding sequence of a therapeutic of the invention can be administered either locally or systemically (e.g., intravenously, intramuscularly, intradermally, subcutaneously, intratumorally, intranasally, orally, intrahepatically, intratracheally). These constructs can utilize viral or non-viral vector approaches as either an *in vivo* or *ex vivo* modality. Expression of such coding sequence can be induced using endogenous mammalian or heterologous promoters. Expression of the coding sequence *in vivo* can be either constitutive or regulated as is described in detail below.

Within one aspect of the present invention, the gene delivery vector is a plasmid-based vector. Representative examples of suitable vectors in this regard include those disclosed within U.S. Patent No. 5,580,859 entitled "Delivery of Exogenous DNA Sequences in a Mammal," U.S. Patent No. 5,589,466 entitled "Induction of a Protective Immune Response in a Mammal by Injecting a DNA Sequence," U.S. Patent Nos. 5,688,688, 5,814,482 and 5,580,859, and Donnelly et al., *ibid*)

Within other aspects of the present invention, gene delivery vectors can be generated based upon a retrovirus. Briefly, retroviral gene delivery vectors of the present invention may be readily constructed from a wide variety of retroviruses, including for example, B, C, and D type retroviruses as well as spumaviruses and lentiviruses (see RNA Tumor Viruses, Second Edition, Cold Spring Harbor Laboratory, 1985). Such retroviruses may be readily obtained from depositories or collections such as the American Type Culture Collection ("ATCC"; Rockville, Maryland), or isolated from known sources using commonly available techniques. Representative examples of retroviral gene delivery vectors are described in more detail in EP 0,415,731; PCT Publication Nos. WO 90/07936; WO 91/0285, WO 93/11230; WO 93/10218, WO 94/03622; WO 93/25698; WO 93/25234; and U.S. Patent Nos. 5,219,740, 5,716,613, 5,851,529, 5,591,624, 5,716,826, 5,716,832, and 5,817,491.

Other suitable gene delivery vectors can be generated from alphaviruses (see, e.g., U.S. Patent Nos. 5,091,309 and 5,217,879, 5,843,723, and 5,789,245),

recombinant adenoviral vectors (see, *e.g.*, U.S. Patent No. 5,872,005), and numerous other viruses such as pox viruses, such as canary pox virus or vaccinia virus (Fisher-Hoch et al., *PNAS* 86:317-321, 1989; Flexner et al., *Ann. N.Y. Acad. Sci.* 569:86-103, 1989; Flexner et al., *Vaccine* 8:17-21, 1990; U.S. Patent Nos. 4,603,112, 4,769,330 and 5,017,487; WO 89/01973); adeno-associated vectors (see, *e.g.*, U.S. Patent No. 5,872,005); SV40 (Mulligan et al., *Nature* 277:108-114, 1979); herpes (Kit, *Adv. Exp. Med. Biol.* 215:219-236, 1989; U.S. Patent No. 5,288,641); and lentivirus such as HIV (Poznansky, *J. Virol.* 65:532-536, 1991) or FIV.

C. METHODS OF STIMULATING AN IMMUNE RESPONSE

10 The ability to recognize and defend against foreign pathogens is central to the function of the immune system. This system, through immune recognition, must be capable of distinguishing "self" from "non-self" (foreign), which is essential to ensure that defensive mechanisms are directed toward invading entities rather than against host tissues. The fundamental features of the immune system are the presence
15 of highly polymorphic cell surface recognition structures (receptors) and effector mechanisms (antibodies and cytolytic cells) for the destruction of invading pathogens.

Cytolytic or cytotoxic T lymphocytes (CTL) are normally induced by the display of processed pathogen-specific peptides in conjunction with the MHC class I or class II cell surface proteins. Also stimulated by this type of antigen presentation
20 are the generation and production antibodies, helper cells and memory cells. Within one embodiment of the present invention, presentation of immunogenic viral determinants in the context of appropriate MHC molecules efficiently induces optimal CTL responses without exposing the patient to the pathogen. This vector approach to immune stimulation provides a more effective means of inducing potent class I -
25 restricted protective and therapeutic CTL responses, because the type of immunity induced by the vector more closely resembles that induced by exposure to natural infection. Based on current knowledge of several viral systems, it is unlikely that exogenously supplied, non-replicating viral antigens, such as peptides and purified recombinant proteins, will provide sufficient stimulus to induce optimal class I-

restricted CTL responses. Alternatively, vector-delivered expression of selected viral proteins or other antigens corresponding to a pathogenic condition, such as cancer, within target cells as described within the present invention provides such a stimulus.

By way of example, in the case of HIV-1 infections, patients develop
5 antibodies specific for a variety of viral envelope-region determinants, some of which are capable of in vitro virus neutralization. Nevertheless, disease progression continues and the patients eventually succumb to the disease. Low-level CTL responses against infected patients' cells (Plata et al., *Nature* 328:348-351, 1987) and against target cells
10 infected with recombinant vaccinia vectors expressing HIV gag, pol, or env (Walker et al., *Nature* 328:345-348, 1987; Walker et al., *Science* 240:64-66, 1988) have been detected in some HIV-1 seropositive patients. In addition, it has recently been shown that murine as well as human CTL can be induced by autologous stimulator cells expressing HIV gp 120 via transfection (Langlade-Demoyan et al., *J. Immunol.* 141:1949, 1988). Improved CTL induction could be therapeutically advantageous to
15 infected patients and provide effective preventive therapy to individuals under noninfectious conditions. HIV infection itself may not be producing an adequate CTL response because other elements associated with HIV infection may prevent proper immune stimulation. In addition, it may be that stimulation of T-cells by infected cells is an interaction that leads to infection of the stimulated T-cells.

20 HIV is only one example. This approach can be effective against many virally linked diseases or cancers where a characteristic antigen (which does not need to be a membrane protein) is expressed, such as in HPV and cervical carcinoma, HTLV-I-induced leukemia's, prostate-specific antigen (PSA) and prostate cancer, mutated p53 and colon carcinoma, GD2 antigen and melanoma.

25 1. *In Vivo Administration*

Within certain aspects of the present invention, methods are provided for delivering one or more nucleotide sequences to a warm-blooded animal, comprising the step of administering to the warm-blooded animal a gene delivery vector, such as

plasmid DNA, retrovirus vector, adenovirus vector, AAV vector, poxvirus vector, herpesvirus vector, alphavirus vector, and eukaryotic layered vector initiation system, expressing the nucleotide sequence as described above.

Gene delivery vectors may be administered to warm-blooded animals
5 either directly (*e.g.*, intravenously, intramuscularly, intraperitoneally, subcutaneously, orally, rectally, intraocularly, intranasally, intradermally, intratumorally), or by various physical methods such as lipofection (Felgner et al., *Proc. Natl. Acad. Sci. USA* 84:7413-7417, 1989), direct DNA injection (Acsadi et al., *Nature* 352:815-818, 1991); microprojectile bombardment (Williams et al., *PNAS* 88:2726-2730, 1991); liposomes
10 of several types (*see, e.g.*, Wang et al., *PNAS* 84:7851-7855, 1987); CaPO₄ (Dubensky et al., *PNAS* 81:7529-7533, 1984); DNA ligand (Wu et al., *J. of Biol. Chem.* 264:16985-16987, 1989); administration of nucleic acids alone (WO 90/11092); or administration of DNA linked to killed adenovirus (Curiel et al., *Hum. Gene Ther.* 3:147-154, 1992); via polycation compounds such as polylysine, utilizing receptor specific ligands; as
15 well as with psoralen inactivated viruses such as Sendai or Adenovirus. In addition, the eukaryotic layered vector initiation systems may either be administered directly (*i.e.*, *in vivo*), or to cells which have been removed (*ex vivo*), and subsequently returned.

Within one embodiment of the invention, the expression cassette or gene delivery vector is delivered (either *in vivo* or *ex vivo*) to dendritic cells (DC). Briefly,
20 dendritic cells (DC) are professional antigen presenting cells, which initiate and modulate the immune response. In particular, DCs are potent stimulators of B and T lymphocytes (for review, see Banchereau and Steinman, *Nature* 392:245-252, 1998). Stimulation of the T and B lymphocyte antigen-specific effectors occurs by presentation by the dendritic antigen presentation cells, in which processed antigen is
25 displayed in conjunction with major histocompatibility complex (MHC) molecules of two alternative types (MHC class I, or MHC class II), to T lymphocytes, via the T-cell antigen receptor. Antigen presentation via MHC class I results in a cellular CD8⁺ cell (cytotoxic T cell, CTL) response, whereas antigen presentation via MHC class II results in stimulation of CD4⁺ cells, and subsequently B lymphocytes, resulting in a humoral
30 (antibody, Ab) response.

The present invention relates broadly to compositions and methods for stimulating the immune response through the utilization of various RNA and other molecules synthesized *in vivo* from an expression cassette contained within an appropriate gene delivery vehicle. As will become evident from the detailed examples
5 below, such immunostimulatory RNA molecules may comprise sequences which form double-stranded (ds) RNA structures within a desired cell, either by base-pairing with RNAs already present in the cell or via self-complementing sequences within the expressed RNA molecule. The presence of such dsRNA molecules within a cell typically will lead to the induction of one or more elements of the immune system,
10 including interferon responses. Further, the immunostimulatory RNA molecules of the present invention also may comprise antisense or ribozyme sequences that specifically target one or more cellular gene products, such that suppression or down-regulation of the targeted gene product results in the up-regulation and increased expression of a second gene product associated with an immune response. The RNA molecules of the
15 present invention may be expressed alone or co-expressed with one or more antigens derived from a pathogenic agent (*e.g.*, infectious agent, cancer), and are suitable for use in a wide variety of gene delivery systems, such as plasmid DNA, retrovirus vectors, adenovirus vectors, AAV vectors, poxvirus vectors, herpesvirus vectors, and alphavirus vectors.

20 The following examples are offered by way of illustration, and not by way of limitation.

EXAMPLES

25

EXAMPLE 1

CONSTRUCTION OF GENE DELIVERY VECTORS THAT SYNTHESIZE DOUBLE-STRANDED RNA
IN TRANSFECTED CELLS

The construction of gene delivery vectors that provide an enhanced immune response to a desired antigen when inoculated into a warm-blooded animal is described below. As an example, plasmids that express only non-protein encoding double-stranded RNA, or in addition, which also express a desired antigen from a pathogenic agent, are illustrated.

A. EXPRESSION CASSETTES THAT TRANSCRIBE NONCODING dsRNA IN TRANSFECTED CELLS

In the present invention, constructions of expression cassettes that transcribe non-coding dsRNA in desired transfected target cells *in vivo* are provided (see Figures 1-3). As a preferred example, synthesis of dsRNA is from an RNA polymerase III promoter-based expression cassette. However, other promoters (*e.g.*, RNA polymerase I and II) may be readily substituted. Thus, in cells transfected *in vivo* with the plasmid DNA constructs described herein, significant levels of non protein-encoding dsRNA are synthesized either alone, or in addition to a specific desired antigen from a pathogenic agent. Expression of the dsRNA results in an overall enhanced immune response to the antigen.

Specifically, an RNA polymerase III (pol III) promoter-based non-coding dsRNA expression cassette (Figure 1) was constructed to contain the following ordered elements: Adenovirus 2 VA1 RNA promoter (nucleotides -70/+30), nucleotides 447-547 of the Rhinovirus 1A 3C protease gene, in which the A nucleotide present in an ATG codon present in any alternative reading frame was changed, a 10 nucleotide-long stretch of C residues, nucleotides 447-547 of the Rhinovirus 1A 3C protease gene, in an antisense orientation, and the RNA polymerase III consensus transcription termination sequence. Additionally, plasmids were constructed which additionally contained an RNA polymerase II (pol II) expression cassette expressing an antigen (Figure 4), consisting of the following ordered elements: CMV promoter, intron, heterologous gene (lac Z or the glycoprotein B gene from herpes simplex virus type 1, and the SV40 polyadenylation/transcription termination sequence. Plasmids were constructed in which the RNA products transcribed from the pol II and pol III expression cassettes were in the same or in opposite directions, relative to each other.

These ordered elements in the pol II and pol III expression cassettes described above were cloned into a pUC-derived kanamycin-resistant plasmid DNA vector for propagation in *E. coli*, as outlined below, using standard recombinant DNA techniques (Maniatis and Sambrook). The plasmids were constructed on the pBGS131 (Spratt et. al., *Gene* 41:337-342, 1986; ATCC #37443) vector backbone that was modified to remove extraneous sequences, to insert an alternate multiple-cloning site (*PmeI* - *BglII* - *XhoI* - *NotI* - *EcoRI* - *PmeI*), and to render an existing *XhoI* site within the kanamycin resistance gene non-functional. This modified vector known as pBGSVG and has been described previously (WO 97/38087).

For expression of genes encoding desired antigens, the RNA polymerase II cassette from the pCI vector (Promega, Madison, WI) was inserted into the pBGSVG plasmid vector, described above. The pCI vector DNA was digested first with *BamHI* and the 5'-overhang ends were filled in by incubation with the Klenow enzyme and dNTPs. Following purification with GeneCleanII (Bio 101, Vista, CA), the treated plasmid DNA was then digested with *BglII* and the 1348 bp fragment, consisting, in order, of the human cytomegalovirus (CMV) immediate early enhancer/promoter, a chimeric intron composed of the 5'-splice site for the β -globin intron and the 3'-splice site from an IgG intron, a multiple cloning site, and the SV40 late region poly(A) sequence, was inserted into the pBGSVG plasmid digested with *EcoRI*, fill-in of 5'-overhang ends, and digested with *BglII*. Gel purification of the pol II expression cassette-containing fragment was not necessary, as clones were selected with kanamycin-containing media on which bacteria containing the pBGSVG (kan^r gene) vector, but not the pCI (amp^r gene) vector, can grow. This plasmid was designated pBGSVG-pol II.

The lac Z gene or the glycoprotein B (gB) gene from HSV-1 was then inserted into the polylinker of pBGSVG-pol II. The source of the lac Z gene was the pSV- β -Galactosidase Control Vector (Promega, Madison, WI), from which it was isolated as a *HindIII*-*BamHI* 3737 bp fragment, ligated into the pKS+ plasmid (Stratagene, La Jolla, CA), then re-isolated by digestion with *XhoI* and *NotI*, and ligated

into the pBGSVG -pol II plasmid that was digested with *Xho*I and *Not*I. This plasmid was designated pBGSVG -pol II/ β -gal.

The HSV-1 KOS strain glycoprotein B (gB) gene was obtained from Dr. Martin Muggeridge (LSU Medical Center, Shreveport, LA) and subcloned into the
5 pBGSVG -pol II plasmid between the *Xho*I and *Xba*I sites within the multiple cloning sequence. This plasmid was designated pBGSVG -pol II/gB-1.

The pol III-based expression cassette was then inserted into the pBGSVG, pBGSVG -pol II/ β -gal, or pBGSVG -pol II/gB-1 plasmids, at the unique *Bgl*II site. Assembly of the desired components in the pol III-based expression cassette
10 was performed by PCR to juxtapose the following ordered sequences:

Ad2 Stuffer (nts. 1051-10584): (SEQ. ID. NO.)

5'CCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGACGCTC
TGGA-3'

15

Ad2 VA1RNA promoter [(-70/+30) (nts. 10585-10682)]: (SEQ. ID. NO.)

5'CCGTGCAAAAGGAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGA-3'

20 Rhinovirus 1A 3C Protease (nts. 447-457): (SEQ. ID. NO.)

5'CCTAAAACCAAAGTACCTGAAAGAAGAGTAGTTGCTCAAGGTCCAGAAGAAGAATTTGGAAGGTCAATTCTCAAAAACAATACTTGTGTGATTACTACAGG-3'

25 Stem: (SEQ. ID. NO.)

5'-CCCCCCCCCCC-3'

Rhinovirus 1A 3C Protease (nts. 447-457) ANTISENSE: (SEQ. ID. NO.)

5'CCTGTAGTAATCACACAAGTATTGTTTTTGAGAATTGACCTTCCAAATTCT
30 TCTTCTGGACCTTGAGCAACTACTCTTCTTTCAGGTACTTTGGTTTTAGG-3'

RNA polymerase III termination sequence: (SEQ. ID. NO.)

5'-GCGCTTTTGGCGC-3'

5 Additionally, the following restriction sites were interspersed within the pol III expression cassette in order to facilitate interchanging the ordered elements present within the construct:

	<u>Enzyme:</u>	<u>Recognition Sequence:</u>
10	<i>Bgl</i> II	A'GATCT
	<i>Bsp</i> EI	T'CCGGA
	<i>Eco</i> RI	G'AATTC
	<i>Xba</i> I	T'CTAGA

15 The configuration of the ordered sequence elements present in the pol III expression cassette is 5'-*Bgl*II-Ad2 Stuffer-Ad2 VA1 promoter-*Xba*I-Rhinovirus Sense Stem-Loop- Rhinovirus Antisense Stem-*Eco*RI-pol III termination-*Bsp*EI-*Bgl*II-3'

 Juxtaposition of the sequence elements shown above to construct the pol III-based expression cassette was accomplished in two steps. The first step used the
20 partially complementary oligonucleotides shown below, each at 1 μ M concentration in a short (10 cycle) PCR amplification:

Ad2VA 1F: (SEQ. ID. NO.)

5'AGATCTCCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTG
25 ACGCTCTGGACCGTGCAAAAGGAGAGCC-3'

Ad2VA 1R: (SEQ. ID. NO.)

5'CCCTTGCGAATTTATCCACCAGACCACGGAAGAGTGCCCGCTTACAGGCT
CTCCTTTT-3'

Ad2VA 2F: (SEQ. ID. NO.)

5'AATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCTGAACCCCGGATCTA
GACCTAAAACCAAAGTACCTGAAAGAAGA-3'

5

Ad2VA 2R: (SEQ. ID. NO.)

5'TGAGAATTGACCTTCCAAATTCTTCTTCTGGACCTTGAGCAACTACTCTTC
TTTCAGG-3'

10 Ad2VA 3F: (SEQ. ID. NO.)

5'GGTCAATTCTCAAAAACAATACTTGTGTGATTACTACAGGCCCCCCCCCCC
CCTGTAGTAATCACACAAGTATTGTTTTT-3'

Ad2VA 3R: (SEQ. ID. NO.)

15 5'AGTAGTTGCTCAAGGTCCAGAAGAAGAATTTGGAAGGTCAATTCTCAAAA
ACAATACT-3'

Ad2VA 4F: (SEQ. ID. NO.)

5'TGAGCAACTACTCTTCTTTCAGGTACTTTGGTTTTAGGGAATTCGCGCTTT
20 TTGCGCTCCGGAAGATCT-3'

Ad2VA 4R: (SEQ. ID. NO.)

5'-AGATCTTCCGGAGCGCAAAAAG-3'

25 PCR amplification of the oligonucleotides comprising the pol III-based
expression cassette shown above was performed in a single reaction, using the Vent_R
DNA polymerase (New England Biolabs, Beverly, MA), and reaction conditions as
suggested by the supplier, containing in addition 2 mM MgSO₄, 5% DMSO, and Hot
Start Wax beads (Perkin-Elmer), with the PCR amplification protocol, as shown below.

30

28		
Temperature (°C)	Time (Min.)	No. Cycles
95	2	1
95	0.5	
55	0.5	10
72	0.5	

The PCR products were purified (QIAquick PCR purification kit, QIAGEN, Chatsworth, California) and the purified reaction products were used in a second PCR amplification, with the primer pair shown below:

5

Ad2 PCR 1F: (SEQ. ID. NO.)

5'-CCCTTCCCAGATCTCCATGGTCGGGACG-3'

Ad2 PCR 1R: (SEQ. ID. NO.)

10 5'-TTTCCTTTAGATCTTCCGGAGCGCAAAAAG-3'

PCR amplification of the pol III-based expression cassette from the first PCR amplification shown above was performed with the Vent_r DNA polymerase (New England Biolabs, Beverly, MA), and reaction conditions as suggested by the supplier, containing in addition 2 mM MgSO₄, 5% DMSO, and Hot Start Wax beads (Perkin-Elmer), with the PCR amplification protocol 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 were purified with a QIAquick kit, and then inserted directly into the pCR Blunt plasmid DNA vector (Invitrogen, Carlsbad, CA). This plasmid was designated pCR-pol III.

Alternatively, the purified products of this second PCR reaction were
 5 digested with *Bgl*II, purified, and ligated into the pBGSVG, pBGSVG -pol II/ β -gal, or pBGSVG -pol II/gB-1 plasmids, prepared by digestion with *Bgl*II, treatment with calf intestinal alkaline phosphatase, and purification. With all plasmid constructs, clones containing the pol III expression cassette in both orientations (F or R, below) were isolated. These plasmids were designated:

10

pBGSVG -pol III(F)

pBGSVG -pol III(R)

pBGSVG -pol II/ β -gal-pol III(F)pBGSVG -pol II/ β -gal-pol III(R)

15 pBGSVG -pol II/gB-1-pol III(F)

pBGSVG -pol II/gB-1-pol III(R)

To demonstrate functionality of the pol III expression cassette, cultured cells were transfected with pBGSVG-pol II/ β -gal-pol III(F), pBGSVG-pol II/ β -gal-pol
 20 III(R), or pBGSVG-pol II/ β -gal plasmid, and expression of the predicted double-stranded RNA species was determined by RNA dot blot analysis of total RNA purified at 48 hr post-transfection, and treated subsequently with DNase. The predicted sequence of double-stranded (*i.e.*, complementary) RNA, by virtue of its structure, should also be resistant to treatment with RNase. Baby hamster kidney-21 (BHK-21)
 25 cells were transfected with 1.0 μ g of plasmid DNAs complexed with 4.0 μ l of a commercially available lipid (Lipofectamine, GIBCO-BRL). Eagle minimal essential medium supplemented with 10% fetal bovine sera was added to the cells at 4 hours post transfection (hpt), unless otherwise indicated. Transfected cells were incubated at 37°C. At 48 hrs post transfection, β -galactosidase reporter gene expression was
 30 determined by direct *in situ* staining of fixed cells with X-gal as described previously (MacGregor, *Cell Mol. Genet.* 13:253-265, 1987). The efficiency of transfection with

each of the three test plasmids was equivalent and approximately 10%. Alternatively in other samples, total RNA was isolated using a commercially available reagent (Tel-Test, TX), according to the instructions of the supplier. All samples were treated with DNase. Half of the samples also were treated with RNase, to lower the viscosity of the sample and to demonstrate the presence of a double-stranded RNA species, resistant to
5 nuclease digestion. The purified samples were loaded onto Zeta™Probe (Bio-Rad, Richmond CA), as follows, and hybridized with ³²P-labeled Ad2VA 3R probe. The dsRNA was found to be expressed in cells transfected with the pBGSVG-pol II/β-gal-pol III(F) and pBGSVG-pol II/β-gal-pol III(R) plasmids, but not pBGSVG-pol II/β-gal
10 plasmid, demonstrating that as expected, the pol III-based expression cassettes synthesize a double-stranded RNA species that is resistant to RNase.

In addition to the dsRNA configuration described above, a number of dsRNA cassette variations or modifications are readily made by one of skill in the art. These include alternative sequences which form a similar dsRNA hairpin, but with the
15 same or other sequences that form a longer or shorter region of dsRNA, nucleotide substitution(s) resulting in less than 100% base-pairing within the dsRNA structure, insertion of a ribozyme sequence within the non-base paired loop region for cleavage into two separate, complementary RNAs which form dsRNA (Figure 2), sequences which form circular, self-complementary "viroid-like" dsRNA (Figure 3), and the use
20 of two separate expression cassettes for transcription of complementary RNAs which form dsRNA (Figure 2).

EXAMPLE 2

IMMUNE STIMULATION BY PROMOTION OF APOPTOSIS

25

The present invention also provides compositions and methods for stimulation of an immune response specific for a pathogenic agent by expressing, in addition to an antigen from the pathogenic agent, one or more nucleic acid sequences that promote apoptosis. The expressed nucleic acid sequence that promotes apoptosis

may be coding (*e.g.*, encode a polypeptide) or non-coding (*e.g.*, ribozyme, antisense RNA); and, the promotion of apoptosis may be direct (*e.g.*, via an inductive signal) or indirect (*e.g.*, suppression of an anti-apoptotic gene). Furthermore, expression cassettes containing the nucleic acid sequences that promote apoptosis may be part of the same
5 gene delivery vehicle that expresses the antigen (Figure 5), as distinct cassettes or a single cassette (*i.e.*, using an IRES), or may be contained within a different gene delivery vehicle from that which expresses the antigen. In those instances where the pro-apoptotic sequence is expressed from a different gene delivery vehicle than the antigen expressing gene delivery vehicle, administration to a warm-blooded animal
10 may be simultaneous, or separately in any desired order.

Numerous elements of the apoptotic regulatory cascade may be selected for use in accordance with the teachings of the present invention, including for example, the expression of pro-apoptotic gene products such as Bax, Bak, Bok, Bik, Blk, Hrk, BNIP3, Bim_L, Bad, Bid, Fas ligand and EGL-1, or the suppression of anti-
15 apoptotic gene products such as Bcl-2, Bcl-xL and CED-9. In addition, it should be appreciated by those of skill in the art that although vector constructions are illustrated with only one antigen, this antigen is provided as an example and based on the teachings provided herein, alternative antigens may be chosen from a variety of pathogenic agents, including for example, viruses, bacteria, fungi, parasites, and cancer
20 cells, provided that sufficient sequence data is available to facilitate the cloning of the antigens.

A. CO-EXPRESSION OF A PROAPOPTOTIC GENE PRODUCT

To demonstrate the immunostimulatory effects of expressing both a pro-apoptotic sequence and vaccine antigen from a single gene delivery vector, herpes
25 simplex virus type 1 (HSV-1) glycoprotein B, for example, was chosen as the antigen and murine BAX α , for example, was chosen as the pro-apoptotic gene product. Glycoprotein B provides a well-characterized measure of both humoral and cellular immune induction, as well as protection from virulent HSV challenge in a mouse model (Hariharan et al., *J. Virol.* 72:950-958, 1998). Murine BAX α is preferred over human

BAX α for vaccine studies in the mouse model; however, details also are provided for similar constructs containing human BAX α . Construction of a plasmid DNA vector expressing both HSV-1 glycoprotein B and BAX α is accomplished by first isolating a fragment comprising the ordered elements CMV immediate early promoter, HSV-1 glycoprotein B, SV40 polyadenylation signal from plasmid pCI-gB (Hariharan et al., *ibid*). Plasmid pCI-gB is digested with *Bgl*III and *Bam*HI, the DNA is blunt-ended using Klenow enzyme and dNTPs, the glycoprotein B containing fragment is purified from an agarose gel using GeneCleanII (BIO101, Vista, CA), and then ligated into a previously described plasmid vector pBGSVG (WO 97/38087) digested with *Pac*I and also blunt-ended, using T4 DNA polymerase and dNTPs. The resulting construct is designated pBGSV-gB.

Insertion of an expression cassette for BAX α into pBGSV-gB is accomplished by first isolating from plasmid pCI (Promega, Madison, WI), a fragment comprising the ordered elements CMV immediate early promoter, multiple cloning site, SV40 polyadenylation signal. The fragment is isolated by digestion with *Bgl*III and *Bam*HI, blunt-ending with Klenow and dNTPs, and purification from an agarose gel using GeneCleanII. The fragment then is ligated into plasmid pBGSV-gB that has been digested with *Pme*I, treated with alkaline phosphatase, and purified from an agarose gel, to produce the construct pBGSV-gB-cass2. BAX α is obtained by PCR amplification using two oligonucleotide primers:

BAX α fwd

5'ATATATGAATTCGGTGATGGACGGGTCCGGGGAGCAG

25 BAX α rev

5'TATATAGAATTCTCAGCCCATCTTCTTCCAGATGGTG

that contain flanking *Eco*RI restriction sites, PFU Turbo polymerase (Stratagene, San Diego, CA) according to the manufacturer's instructions, and template DNA from

mouse or human B cell cDNA libraries (CLONTECH, Palo Alto, CA) or from commercially available plasmids (INVIVOGEN, San Diego, CA). Alternatively, polyA purified RNA from mouse and human lymphoid cells (*e.g.*, cell lines RL7 and FL5.12; Oltvai et al., *Cell* 74:609-619, 1993) may be used as template for reverse transcriptase coupled PCR reactions. Following amplification, the PCR product is purified with a QIAquick kit (QIAGEN, Chatsworth, CA), digested with EcoRI, re-purified with QIAquick, and ligated with plasmid pBGSV-gB-cass2 that also has been digested with EcoRI and treated with alkaline phosphatase. Constructs are obtained with the BAX α insert in both the correct and opposite (negative control) orientation, and these plasmids are referred to as pBGSV-gB-BAX and pBGSV-gB-BAXrev.

Similar constructions that express other pro-apoptotic genes or antigens, are readily constructed from these teachings by one of skill in the art.

B. CO-EXPRESSION OF ANTISENSE RNA SPECIFIC FOR ANTI-APOPTOTIC GENES

In alternative constructs, an antisense or ribozyme sequence, which specifically hybridizes to the mRNA of an anti-apoptotic gene, is expressed from a cassette, such as an RNA polymerase III (pol III) cassette. As described above, this cassette preferably is constructed as part of the same gene delivery vector (*e.g.*, plasmid DNA) which expresses the desired antigen. However, it may also be contained within a separate vector that is administered to the same vaccinee as the antigen expressing gene delivery vector.

For example, an cassette is constructed wherein an RNA polymerase III promoter is operably linked with a sequence which, when transcribed, gives rise to an antisense RNA that specifically binds to the 5' region of the anti-apoptotic Bcl-2 gene mRNA (Tsujimoto and Croce, *Proc. Natl. Acad. Sci. USA* 83:5214-5218, 1986). Specifically, a cassette comprising the following ordered elements: *EcoRI* restriction site for cloning, Adenovirus 2 VA1 RNA promoter (nucleotides -70/+30), nucleotides 4-44 of the human Bcl-2 gene coding sequence, RNA polymerase III consensus transcription termination sequence, and *EcoRI* restriction site for cloning. Additional or alternative Bcl-2 sequences also may be substituted for nucleotides 4-44. In order to

34

provide a mechanism for demonstrating the immunostimulatory effect of this approach, the elements are inserted into the previously described pBGSV-gB construct, which encodes glycoprotein B of HSV.

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

Ad2 Stuffer (nts. 1051-10584):

5'CCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGACGCTC
TGGA-3'

10

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

5'CCGTGCAAAAGGAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGGA
TAAATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGA-3'

15 Human Bcl-2 coding sequence comprising nucleotides 4-44

5'GCGCACGCTGGGAGAACGGGGTACGACAACCGGGAGATAGT

RNA polymerase III termination sequence:

5'-GCGCTTTTTGCGC-3'

20

Juxtaposition of the sequence elements shown above to construct the pol III-based expression cassette is accomplished in two steps. The first step uses the partially complementary oligonucleotides shown below, each at 1 μ M concentration in a short (10 cycle) PCR amplification:

25

Ad2VAbcl 1F:

5'GAATTCCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGA
CGCTCTGGACCGTGCAAAAGGAGAGCC-3'

Ad2VAbcl 1R:

5'CCCTTGCGAATTTATCCACCAGACCACGGAAGAGTGCCCGCTTACAGGCT
CTCCTTTT-3'

5 Ad2VAbcl 2F:

5'AATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGAACT
ATCTCCCGGTTGTCGTACCCCGTTC-3'

Ad2VAbcl 2R:

10 5'GAATTCGCGCAAAAAGCGCGCGCACGCTGGGAGAACGGGGTACGA-3'

PCR amplification of the oligonucleotides comprising the pol III-based expression cassette shown above is performed in a single reaction, using Vent polymerase and reaction conditions as suggested by the supplier, containing in addition
15 2 mM MgSO₄, 5% DMSO, and Hot Start Wax beads, with the 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	

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

AdVA PCR 1F:

5'-ATATATAGAATTCCATGGTCGGGACGC

AdVA PCR 1R:

5'-ATATATGAATTCGCGCAAAAAGCGC

The second PCR amplification of the pol III-based cassette is performed with the Vent polymerase and similar reaction conditions, with the PCR amplification 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 *EcoRI*, re-purified and then inserted into the vector pBGSV-gB that also has been digested with *EcoRI* and treated with alkaline phosphatase. This plasmid is designated pBGSV-gB α Bcl.

For a separate set of PCR amplifications, an additional pair of oligonucleotide primers (shown below) is substituted for Ad2Vabcl2F and Ad2Vbcl2R, in order to generate a negative control cassette which allows for construction of the Bcl-2 sequence in the sense orientation.

Ad2VAbcl 3F

5'AATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGAGCG
CACGCTGGGAGAACGGGGTACGACA-3'

Ad2VAbcl 3R

5'GAATTCGCGCAAAAAGCGCACTATCTCCCGGTTGTCGTACCCCGT-3'

These primers are used as described above to generate a second fragment with EcoRI ends, and that when inserted into pBGSV-gB, results in a plasmid construct designated pBGSV-gBBcl.

5 Similar constructions that utilize ribozyme sequences (see below) specific for Bcl-2, as well as ribozyme or antisense sequences specific for other anti-apoptotic genes, are readily constructed from these teachings by one of skill in the art.

C. DETERMINATION OF IMMUNOSTIMULATORY EFFECTS FOR PROAPOPTOTIC SEQUENCES

10 To demonstrate immunostimulatory effects of these vector modifications, multiple BALB/c mice are immunized intramuscularly with plasmids pBGSV-gB-BAX, pBGSV-gB-BAXrev, pBGSV-gB α Bcl, and pBGSV-gBBcl in saline, at doses ranging from 10 ug to 10 ng. Following these immunizations, the level of HSV-gB specific antibody and CTL induced by each construct is determined
15 according to Hariharan et al (*ibid*). In addition, representative animals also are challenged with virulent HSV to determine the level of protection (Hariharan et al., *ibid*). Similar levels of immune response or protection at lower DNA doses by the modified vectors compared to the conventional or control vectors is indicative of immune stimulation by the elements of the present invention.

20

EXAMPLE 3

DISRUPTION OF IRF2 EXPRESSION TO ENHANCE IMMUNE EFFICACY

25 In addition to the examples above, the present invention also provides compositions and methods of immune stimulation utilizing specific antisense or ribozyme RNA sequences transcribed *in vivo* that target one or more mRNAs encoding immune regulatory components. The antisense or ribozyme sequences may co-expressed with a desired antigen from a pathogenic agent, or expressed alone and

administered to a warm-blooded animal with or without the desired antigen or antigen-expressing gene delivery vector.

IFN γ plays a major role in the immune response by inducing antiviral and bactericidal activity, stimulating antigen presentation through MHC class I and class II molecules, and by inducing the expression of IFN α and IFN β genes, as well as a variety of genes whose functions have not yet been determined. A primary response gene to IFN γ is the interferon regulatory factor 1 (IRF1). IRF1 is a transcriptional factor that is expressed at the level of few mRNA copies per cell prior to induction by IFN γ (Fujita et al., *Proc. Natl. Acad. Sci. USA* 86:9936-9940, 1989; Harada et al., *Cell* 58:729-739, 1989). Constitutive expression of IRF1 in transgenic mice results in higher resistance to RNA viruses (Pine, *J. Virol.* 66:4470-4478, 1992) and overexpression in COS cells results in the induction of IFN α and IFN β without viral stimulation (Fujita et al., *Nature* 337:270-272, 1989). Interferon regulatory factor 2 (IRF2) is a transcriptional inhibitor that is expressed constitutively in many cells and antagonizes IRF1 activity, by binding to the same promoter element in the IFNs and IFN-inducible genes (Harada et al., *ibid*). Thus, IRF2 is a negative regulator of the IFN γ response.

The present invention provides a mechanism for interfering with IRF2 expression in cells producing an antigen of interest, in order to boost the immune response to that antigen. In particular, a ribozyme or an antisense RNA targeted to IRF2 mRNA is transcribed *in vivo* from an expression cassette contained within a gene delivery vector, to inhibit IRF2 expression and prolong the IFN γ effects induced by the expression of an antigen of interest. Preferably, the ribozyme or antisense RNA is expressed from the same gene delivery vector encoding the antigen(s) of interest, and is expressed utilizing an RNA polymerase III cassette. In preferred embodiments, the antisense or ribozyme RNA is directed (complementary) to mRNA of the first exon of the IRF2 gene. Since the N-terminal 125 amino acids of IRF2, encoding the transcriptional regulatory activity, exhibit a high degree of homology to the same region in IRF1, exons II, III, and IV are less suitable for targeting.

For example, an RNA polymerase III expression cassette first is constructed to express a hammerhead ribozyme specific for IRF2. A variety of target cleavage sites in the IRF2 mRNA may be chosen and several are depicted in Figure 6, shown in boldface type. For purposes of this illustration, the chosen target site is boxed and complementary "antisense" sequences used by the ribozyme for target recognition are underlined.

Specifically, the pol III/ribozyme cassette is constructed to contain the following ordered elements: *PacI* restriction site, Adenovirus 2 VA1 RNA promoter (nucleotides -70/+30), IRF2-specific ribozyme, RNA pol III consensus transcription termination sequence, *PacI* restriction site. These elements are synthesized using overlapping PCR and 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 *PacI* site on the opposite side of the vector backbone.

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

Ad2 Stuffer (nts. 1051-10584):

5'CCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGACGCTC
TGGA-3'

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

5'CCGTGCAAAAGGAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGGA
TAAATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGA-3'

IRF2 specific ribozyme

5'CTCAGTGTGCTTTTTTCACGCCTGATGAGGCCGTGAGGCCGAAACCAAT
ACAATTCCGCAAGG-3'

RNA polymerase III termination sequence:

5'-GCGCTTTTGTGCGC-3'

Juxtaposition of the sequence elements shown above to construct the pol
III-based expression cassette is accomplished in two steps. The first step uses the
partially complementary oligonucleotides shown below, each at 1 μ M concentration in
5 a short (10 cycle) PCR amplification:

VAIRF2-1F:

5' CACACACTTAATTAACCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCG
CAGTCG-3'

10

VAIRF2-1R:

5' CCGCTTACAGGCTCTCCTTTTGCACGGTCCAGAGCGTCAACGACTGCGCA
CGCCT-3'

15 VAIRF2-2F:

5' AGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAAG
GGTAT -3'

VAIRF2-2R:

20 5' CACACTGAGTCCGGGGTTCGAACCCCGGTCGTCCGCCATGATACCCTTGC
GAATT-3'

VAIRF2-3F:

5' CCCGGACTCAGTGTGCTTTTTTTCACGCCTGATGAGGCCGTGAGGCCGAA
25 ACCAAT-3'

VAIRF2-3R:

5' GTGTGTTTAATTAAGCGCAAAAAGCGCCCTTGCGGAATTGTATTGGTTTC
GGCCTC-3'

PCR amplification of the oligonucleotides comprising the pol III-based expression cassette shown above is performed in a single reaction, using Vent polymerase and reaction conditions as suggested by the supplier, containing in addition
 5 2 mM MgSO₄, 5% DMSO, and Hot Start Wax beads, with the 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	

The PCR product is purified using the QIAquick PCR kit, then used in a
 10 second PCR amplification, with the VAIRF2-1F and VAIRF2-3R primers. The second PCR amplification of the pol III-based cassette is performed with Vent polymerase and similar reaction conditions, with the PCR amplification protocol shown below.

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

15 The product of this second PCR reaction is purified with the QIAquick kit, digested with *PacI*, re-purified and then inserted into the vector pBGSVG-polIII (Example 1) that also has been digested with *PacI* and treated with alkaline phosphatase. This plasmid-based gene delivery vector is designated pBGSrIRF2.

To demonstrate disruption of IRF2 expression by directed antisense or
 20 ribozyme RNAs, a reporter gene system similar to that of Harada et al. (*ibid*) is utilized. In this system, an IRF1/IRF2 binding site is inserted into a reporter construct interposed

42

between an SV40 enhancer element and downstream promoter driving expression of a reporter gene. Transfection of this reporter plasmid into cells expressing IRF2 results in suppression of reporter expression. Co-transfection of the IRF2 expressing cells with the reporter plasmid and a gene delivery vector of the present invention (*e.g.*,
5 pBGSrIRF2 above) allows for a direct measurement of inhibition of IRF2 expression.

EXAMPLE 4

DISRUPTION OF CYCLOOXYGENASE-2 EXPRESSION TO INHIBIT CANCER CELL GROWTH AND
10 FACILITATE HOST-MEDIATED CYTOTOXICITY AGAINST TUMORS

Prostaglandin E2 (PGE2) is one of several immunosuppressive molecules secreted by tumors and its secretion correlates with tumor progression. The presence of high levels of immunosuppressive molecules is at least partially
15 accountable for a tumor's ability to escape host immune surveillance mechanisms. Cyclooxygenase-2 (COX2) is involved directly in PGE2 synthesis. Several tumors have been found to express high levels of COX2 (Liu et al., *Cancer Res.* 56:5125-5127, 1996; Ristimaki et al., *Cancer Res.* 57:1276-1280, 1997) and its expression correlated with the metastatic potentials (Tsuji et al., *Proc. Natl. Acad. Sci. USA* 94:3336-3340,
20 1997). The present invention provides gene delivery vectors and methods for targeted disruption (*e.g.*, ribozyme or antisense) of COX2 expression as a mechanism to facilitate anti-tumor therapy. Since the COX2 cDNA sequence has been established (Hla et al., *Proc. Natl. Acad. Sci. USA* 89:7384-7388, 1992; O'Banion et al., *Proc. Natl. Acad. Sci. USA* 89:4888-4892, 1992), antisense or ribozyme RNAs directed against
25 COX2 mRNA may be designed for insertion into the desired expression cassette, preferably an RNA polymerase III cassette. Other gene products of the PGE2 biosynthetic pathway also may be chosen for targeting according to the present invention. Furthermore, the above described gene delivery vectors also may encode one or more cytokines or tumor antigens.

For example, an RNA polymerase III expression cassette first is constructed to express a hairpin ribozyme specific for COX2. A variety of COX2 mRNA target cleavage sites for hammerhead and hairpin ribozymes may be chosen and are depicted in Figure 6, shown in boldface type. For purposes of this illustration, the
 5 chosen target site is boxed and complementary "antisense" sequences used by the ribozyme for target recognition are underlined.

Specifically, the pol III/ribozyme cassette is constructed to contain the following ordered elements: *PacI* restriction site, Adenovirus 2 VA1 RNA promoter (nucleotides -70/+30), COX2-specific ribozyme, RNA pol III consensus transcription
 10 termination sequence, *PacI* restriction site. These elements are synthesized using overlapping PCR and cloned into pBGSVG as described in the previous example. Assembly of the desired components in the pol III-based expression cassette is performed by PCR to juxtapose the following ordered sequences:

15 Ad2 Stuffer (nts. 1051-10584):

5'CCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGCAGTCGTTGACGCTC
 TGGA-3'

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

20 5'CCGTGCAAAAGGAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAAGGGTATCATGGCGGACGACCGGGGTTCGAACCCCGGA-3'

COX2 specific ribozyme

5'GCAAGGATTTGCTGTATGGCTGAGCGCCAGAGAAGCGCACCAAGAGAAAC
 25 ACACGTTGTGGTATATTACCTGGTA-3'

RNA polymerase III termination sequence:

5'-GCGCTTTTTGCGC-3'

Juxtaposition of the sequence elements shown above to construct the pol III-based expression cassette is accomplished in two steps, as described in the previous example, using the following oligonucleotide primers.

5 VACOX2-1F:

5'CACACATTAATTAACCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGC
AGTCGTT

VACOX2-1R:

10 5'GTGCCCCGCTTACAGGCTCTCCTTTTGCTCGGTCCAGAGCGTCAACGACTGC
GCACGC

VACOX2-2F:

5'CCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAAGGGTA
15 TCATGGC

VACOX2-2R:

5'ATACAGCAAATCCTTGCTCCGGGGTTCGAACCCCGGTCGTCCGCCATGAT
ACCCTTG
20

VACOX2-3F:

5'AAGGATTTGCTGTATGGCTGAGCGCCAGAGAAGCGCACCCAGAGAAACAC
ACGTTGTG

25 VACOX2-3R:

5'TGTGTGTTAATTAAGCGCAAAAAGCGCTACCAGGTAATATACCACAACGT
GTGTTTC

The PCR product from the first reaction is purified using a QIAquick PCR kit, and then is used in a second PCR amplification with the VACOX2-1F and VACOX2-3R primers. The product of this second PCR reaction is purified with the QIAquick kit, digested with *PacI*, re-purified and then inserted into the vector
5 pBGSVG-polIII that also has been digested with *PacI* and treated with alkaline phosphatase. This plasmid is designated pBGSrCOX2.

The ability of gene delivery vectors of the present invention to inhibit COX2 expression may be monitored in cell culture by testing for changes in PGE2 secretion into the culture medium after transfection of the cells with pBGSrCOX2 or
10 the control plasmid pBGSVG. Since colon cancer has been found to produce high levels of PGE2, a cell line derived from this cancer may be particularly suitable for the testing. PGE2 may be detected with commercially available ELISA kits.

15

EXAMPLE 5

SUPPRESSION OF YIN-YANG-1 TO ENHANCE THE IMMUNE RESPONSE

Several transcription factors are involved in the regulation of IFN γ expression (Young, *J. Interferon & Cyt. Res.* 16:563-568, 1996). Yin-Yang 1 (YY1) is
20 a ubiquitous DNA binding protein that regulates the transcription of several genes, including the oncogenes c-myc and c-fos. According to the context, YY1 can act either as a positive regulator or as a negative regulator. YY1 appears to be involved in the transcriptional repression of IFN γ expression by interacting with multiple sites in the promoter. Mutations in the YY1 binding sites result in an increase of the IFN γ
25 promoter activity (Ye et al., *Mol. Cell. Biol.* 16:4744-4753, 1996). It appears that YY1 represses the IFN γ promoter by binding to a site overlapping with the AP1 binding site, which is required for IFN γ expression (Ye et al., *ibid*).

Immunization via gene delivery vectors, in particular plasmid DNA, appears to be efficacious in animal models, for inducing antibodies and CD8+
30 associated CTL responses to the expressed antigen. A desirable feature would be to

include some mechanism for the co-induction of CD4⁺ Th1 associated responses. Since IFN γ plays a major role in these responses, the induction of its expression is required. The present invention utilizes a ribozyme or antisense RNA expressed *in vivo*, to target YY1 mRNA and down-regulate expression, consequently facilitating the up-regulation of IFN γ expression. The antisense or ribozyme RNA may be expressed utilizing an RNA polymerase III cassette, as well as other RNA polymerase promoters, thus generating a unit that can be cloned into the same gene delivery vector as the antigen of interest. This unit also may be used in conjunction with other mechanisms to boost the immune response, and in particular gene therapy applications for which IFN γ expression is a desirable element.

For example, an RNA polymerase III expression cassette first is constructed to express a hammerhead ribozyme specific for YY1. A variety of YY1 mRNA target cleavage sites for hammerhead and hairpin ribozymes may be chosen and several are depicted in Figure 6, shown in boldface type. For purposes of this illustration, the chosen target site is boxed and complementary "antisense" sequences used by the ribozyme for target recognition are underlined. Specifically, a pol III-based ribozyme cassette is constructed in pBGSVG, as described in the previous examples. Assembly of the components again is accomplished by overlapping PCR, with the following oligonucleotide primers:

20

VAYY1-1F:

5'CACACACTTAATTAACCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCG
CAGTC-3'

25 VAIYY1-1R:

5'CGCTTACAGGCTCTCCTTTTGCACGGTCCAGAGCGTCAACGACTGCGCAC
GCCTC-3'

47

VAIYY1-2F:

5'GAGAGCCTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAA
GGGTA-3'

5 VAIYY1-2R:

5'CTACATCGTCCGGGGTTCGAACCCCGGTCGTCCGCCATGATACCCTTGCG
AATTT-3'

VAIYY1-3F:

10 5'CCCCGGACGATGTAGAGGGTGTCGCCCCTGATGAGGCCGTGAGGCCGA
AAGGCCA-3'

VAIRF2-3R:

5'GTGTGTTTAATTAAGCGCAAAAAGCGCGAGCCCTCAGCCATGGCCTTTTCG
15 GCCTC

The PCR product is purified using the QIAquick PCR kit, then used in a second PCR amplification, with the VAIYY1-1F and VAIYY1-3R primers. Next, the product of this second PCR reaction is purified with the QIAquick kit, digested with *PacI*, re-
20 purified and inserted into pBGSVG-polIII that also has been digested with *PacI* and treated with alkaline phosphatase. This plasmid-based gene delivery vector is designated pBGSrYY1.

25 EXAMPLE 6

DISRUPTION OF IL10 EXPRESSION IN TUMOR CELLS FOR ANTI-TUMOR RESPONSE

It has been shown that some tumors express and secrete one or more immunosuppressive cytokines. The presence of high levels of these cytokines is at
30 least partially accountable for escape of tumors from host immunosurveillance. IL10 is

capable of blocking the activation of several cytokines and several accessory functions of macrophages, T cells, and NK cells (Fortis et al., *Cancer Lett.* 104:1-5, 1996; Gotlieb et al., *Cytokine* 4:385-90, 1992; Nakagomi et al., *Int. J. Cancer* 63:366-71, 1995; Wojciechowska-Lacka et al., *Neoplasma* 43:155-158, 1996). The present
5 invention describes mechanisms whereby the repression of IL10 expression is used in tumors to facilitate host immune clearance of the tumor. This approach may be particularly efficacious when used in combination with immunization to defined tumor antigens.

Specifically, expression cassettes are constructed for the delivery of
10 antisense or ribozyme RNAs that specifically target and prevent expression of IL10. The IL10 cDNA sequence has been determined (Vieira et al., *Proc. Natl. Acad. Sci. USA* 88:1172-1176, 1991), and such RNAs may be delivered and expressed *in vivo* using any number of different gene delivery vectors, including plasmid DNA. Such vectors may be used alone or in combination with similar cassettes targeted to other
15 immunosuppressive cytokines, and expressed from the same gene delivery vector or from a separate co-administered vector. Furthermore, the vectors of the present invention may be used in conjunction with other genes encoding either cytokines or tumor antigens.

For example, an RNA polymerase III expression cassette first is
20 constructed to express a hairpin ribozyme specific for IL-10. A variety of IL-10 mRNA target cleavage sites for hammerhead and hairpin ribozymes may be chosen and several are depicted in Figure 6, shown in boldface type. For purposes of this illustration, the chosen target site is boxed and complementary "antisense" sequences used by the ribozyme for target recognition are underlined. Specifically, a pol III-based ribozyme
25 cassette is constructed in pBGSVG, as described in the previous examples. Assembly of the components again is accomplished by overlapping PCR, with the following oligonucleotide primers:

VAIL10-1F:

5' CACACATTAATTAACCATGGTCGGGACGCTCTGGCCGGTGAGGCGTGCGC
AGTCGTTG-3'

5 VAIL10-1R:

5' AGTGCCCGCTTACAGGCTCTCCTTTTGCACGGTCCAGAGCGTCAACGACT
GCGCACG-3'

VAIL10-2F:

10 5' CTGTAAGCGGGCACTCTTCCGTGGTCTGGTGGATAAATTCGCAAGGGTAT
CATGGCG-3'

VAIL10-2R:

5' CCCACTTCCCAGGCAACCTCCGGGGTTCGAACCCCGGTCGTCCGCCATGA
15 TACCCTT-3'

VAIL10-3F:

5' TGCCTGGGAAGTGGGTGCAGCTGTTCTCAAGAAGGGTACCAGAGAAACA
CACGTTGT-3'
20

VAIL10-3R:

5' TGTGTGTTAATTAAGCGCAAAAAGCGCTACCAGGTAATATACCACAACG
TGTGTTTCT-3'

25 The PCR product is purified using the QIAquick PCR kit, then used in a second PCR
amplification, with the VAIL10-1F and VAIL10-3R primers. Next, the product of this
second PCR reaction is purified with the QIAquick kit, digested with *PacI*, re-purified
and inserted into pBGSVG-polII that also has been digested with *PacI* and treated with
alkaline phosphatase. This plasmid-based gene delivery vector is designated
30 pBGSrIL10.

50

For each of the previous examples, similar cassettes are readily made by one of skill in the art, wherein the ribozyme sequence is substituted by an alternative ribozyme or an antisense sequence with non-catalytic activity. In such cases, the choice of sequences for targeting may be the same as those described above, or may be
5 specific for other regions of the mRNA molecule.

EXAMPLE 7

ANTIGEN CO-EXPRESSION WITH RIBOZYME/ANTISENSE CASSETTES

10

Antigens derived from a variety of pathogenic agents (*e.g.*, tumor) may be inserted into the RNA polymerase II cassette contained within the above ribozyme or antisense expression plasmids. Alternatively, the antigens may be co-expressed along with the ribozyme or antisense elements in other gene delivery vehicles, such as
15 retroviruses (see for example U.S. Patents 5,693,522, 5,691,177, 6,652,130, 5,591,624, and 5,830,458, referenced in their entirety), adenovirus, or AAV.

For example a plasmid DNA vector is constructed that expresses the Her2neu antigen and a ribozyme specific for COX-2. Specifically, the Her2neu antigen is cloned by PCR amplification, using the following oligonucleotide primers
20 incorporate flanking *Xba*I and *Not*I restrictions sites.

Her2neuFwd

5'ATATATATCTAGACACCATGGAGCTGGCGGCCTTGT

25 Her2neuRev

5'TATATAGCGGCCGCTCACACTGGCACGTCCAGACCCA

Following PCR amplification, the product is purified using the QIAquick kit, digested with *Xba*I and *Not*I, re-purified with the QIAquick kit, and ligated into plasmid

pBGSrCOX2 that also has been digested with *Xba*I and *Not*I. The new construct is designated pBGSrCOX2/Her2. Additional constructs that contain modified Her2neu sequences may be generated in a similar manner.

In another example, a plasmid gene delivery vector is constructed that expresses the gp100 antigen and a ribozyme specific for COX-2. Specifically, the gp100 antigen is cloned by PCR amplification, using the following oligonucleotide primers incorporate flanking *Xba*I and *Not*I restrictions sites.

gp100Fwd

10 5'ATATATATCTAGACACAATGGATCTGGTGCTAAAAAGATG

gp100Rev

5'TATATAGCGGCCGCTCAGACCTGCTGCCCCACTGAGGAG

15 Following PCR amplification, the product is purified using the QIAquick kit, digested with *Xba*I and *Not*I, re-purified with the QIAquick kit, and ligated into plasmid pBGSrCOX2 that also has been digested with *Xba*I and *Not*I. The new construct is designated pBGSrCOX2/gp100.

Alternatively, the pol III/ribozyme cassettes and antigen cassettes may be inserted into a variety of other gene delivery vectors (*e.g.*, retrovirus, adenovirus) by one of skill in the art using the teachings of this invention and those references incorporated in their entirety.

25 EXAMPLE 8

ANIMAL MODELS FOR *IN VIVO* EVALUATION OF THERAPEUTIC EFFECTIVENESS

Several *in vivo* tumor models may be used for evaluating therapeutic efficacy of these recombinant vectors. For example, using a syngeneic mouse model of melanoma, unmodified B16F10 cells are implanted in C57BL/6 mice at doses of

30

1x10⁵ cells either subcutaneously to develop a solid tumor nodule, or intravenously to development pulmonary metastases, or both. B16F10 mouse melanoma cells have been demonstrated to express gp100, a melanoma associated tumor antigen. Human melanoma cells have been demonstrated to express gp100 and an antibody response
5 against gp100 is associated with clinical responses. The human and mouse genes for gp100 have been cloned and characterized. Novel gene delivery vectors that express a combination of tumor antigen (*e.g.*, gp100) and antisense/ribozyme RNA (*e.g.*, antisense specific for COX2, TGF- β) may be evaluated for *in vivo* efficacy as generally described by Karavodin et al. (*Human Gene Therapy* 9:2231-2241, 1998). Specifically,
10 such vectors are evaluated for therapeutic activity *in vivo* by implanting subcutaneously (s.c.) 1x10⁵ B16F10 melanoma cells into the lower ventral abdomen of 6-8 week old female mice. A palpable tumor nodule is allowed to form at the site of injection for 7-10 days or until the nodule is 2-5 mm in diameter. Measurement and recording of tumor diameter or volume of the s.c. tumor before, during and after treatment is used to
15 indicate therapeutic effectiveness. Sacrifice of the animal and removal of the lungs to visually enumerate tumor cell foci is used as a measurement of systemic anti-tumor response. Alternatively, serum anti-tumor antibodies or *in vitro* bulk cytolytic T cell (CTL) activity may be assayed using standard immunological approaches to determine systemic therapeutic responses. The therapeutic vector may be injected intratumorally,
20 peritumorally, or a combination of both in volumes such as 50 - 100 μ L. Treatment schedules may be one or two injections per day given over 5 to 7 consecutive days of vector that is 1 x 10⁸ cfu/ml or 1-100 ug or greater.

Alternatively, evaluation of a gene delivery vector expressing the Her2/neu tumor antigen and an antisense/ribozyme inhibitor (*e.g.*, specific for COX-2)
25 may be performed in a Her2/neu transgenic mouse model as described by Boggio et al., (*J. Exp. Med.* 188:589-596, 1998). Briefly, transgenic mice develop spontaneous mammary tumors that will involve all mammary glands by about 33 months of age. In a vaccination application, mice are given injections of the therapeutic vector intramuscularly or subcutaneously prior to the development of palpable tumors.

Various vaccination schedules are used (*e.g.*, once a week for 4-6 weeks, or once a month for 3-4 months), with 50-100 μL of viral vector at titers of $1-100 \times 10^6$ cfu/ml or 1-100 μg of a DNA vector. These vectors also may be used in a therapeutic application wherein mice with existing tumors are injected. The therapeutic vector is injected
5 intratumorally, peritumorally, or a combination of both, in volumes of 50-100 μL . Various treatment schedules are utilized, for example, including one or two injections per day given over 5 to 7 consecutive days of vector that is 1×10^8 cfu/ml or greater.

From the foregoing, it will be appreciated that, although specific embodiments of the invention have been described herein for purposes of illustration,
10 various modifications may be made without deviating from the spirit and scope of the invention. Accordingly, the invention is not limited except as by the appended claims.

54

SEQUENCE LISTING

<110> Chiron Corporation

<120> ENHANCEMENT OF THE IMMUNE RESPONSE FOR
VACCINE AND GENE THERAPY APPLICATIONS

<130> PAT 50301W-1

<140> PCT/US00/09186

<141> 2000-04-07

<150> US 60/128,409

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<211> 37

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37

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<211> 41

<212> DNA

<213> Homo sapien

<400> 19

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63

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<211> 57

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<223> Oligonucleotide primer

63

<400> 37

gtgcccgcctt acaggctctc cttttgctcg gtccagagcg tcaacgactg cgcacgc 57

<210> 38

<211> 57

<212> DNA

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<400> 38

cctgtaagcg ggcactcttc cgtgggtctgg tggataaatt cgcaagggtg tcatggc 57

<210> 39

<211> 57

<212> DNA

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<210> 48

<211> 58

<212> DNA

<213> Artificial Sequence

<220>

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<400> 48

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<210> 49

<211> 57

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36

<210> 55

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37

<210> 56

<211> 40

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<223> Oligonucleotide primer

<400> 56

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40

<210> 57

<211> 38

<212> DNA

<213> Artificial Sequence

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<223> Oligonucleotide primer

<400> 57

tatatagcgg ccgctcagac ctgctgccca ctgaggag

38

<210> 58

<211> 288

<212> DNA

<213> Unknown

<220>

<223> Target cleavage site in IL10

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60

68

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gtgaagactt	tctttcaaata	gaaggatcag	ctggacaact	tgttggttaa	ggagtccttg	240
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 <212> DNA
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tctaccgaat	gctg					434

CLAIMS:

1. An expression cassette, comprising a promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double stranded RNA that induces the production of interferon, wherein said expression cassette is:
 - (a) an expression cassette which, when transcribed *in vivo*, forms self-complementary RNA; or
 - (b) an expression cassette comprising a first promoter operably linked to a first nucleic acid molecule, and a second promoter operably linked to a second nucleic acid molecule, wherein said first and second nucleic acid molecules, when transcribed *in vivo*, form double stranded RNA that induces the production of interferon.
2. The expression cassette according to claim 1, wherein said cassette is limited to expression cassette according to claim 1(a).
3. The expression cassette according to claim 1, wherein said cassette is limited to an expression cassette according to claim 1(b).
4. The expression cassette according to claim 1 with the proviso that said RNA is not translated into protein *in vivo*.
5. The expression cassette according to claim 1, wherein said promoter is a pol I or a pol III promoter.
6. The expression cassette according to claim 5 wherein said pol III promoter is an Adenovirus VA1 promoter.
7. The expression cassette, comprising a promoter operably linked to a ribozyme or antisense nucleic acid molecule which, when transcribed *in vivo*, promotes apoptosis,

further comprising a promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double stranded RNA that induces the production of interferon.

8. The expression cassette according to claim 1, further comprising a promoter operably linked to a nucleic acid molecule that encodes a polypeptide of interest.

9. The expression cassette according to claim 8, wherein said promoter which is operably linked to a nucleic acid molecule that encodes a polypeptide of interest is a pol II promoter.

10. The expression cassette according to claim 8, wherein said polypeptide promotes apoptosis.

11. The expression cassette according to claim 8, wherein polypeptide encodes an antigen from a pathogenic agent.

12. An expression cassette, comprising a first promoter operably linked to a nucleic acid molecule which, when transcribed *in vivo*, forms double stranded RNA that induces the production of interferon, and a second promoter operably linked to a nucleic acid molecule that encodes an antigen from a pathogenic agent.

13. The expression cassette according to claim 11 or claim 12, wherein said pathogenic agent is a virus.

14. The expression cassette according to claim 13, wherein said virus is selected from the group consisting of HIV, HSV, HBV, HCV, HPV and FIV.

15. The expression cassette according to claim 11 or claim 12, wherein said pathogenic agent is a bacteria, parasite or fungus.

16. The expression cassette according to claim 11 or claim 12, wherein said pathogenic agent is a tumor.
17. The expression cassette according to claim 8 wherein said polypeptide is a cytokine.
18. The expression cassette according to claim 17, wherein said cytokine is selected from the group consisting of IL-2, IL-12, IL-15 and gamma-interferon.
19. The expression cassette according to claim 11 or claim 12, wherein said additional promoter is a pol II promoter.
20. The expression cassette according to claim 9 or 19 wherein said pol II promoter is selected from the group consisting of CMV, SV40, MoMLV LTR and RSV LTR.
21. A gene delivery vector, comprising an expression cassette according to claim 1.
22. The gene delivery vector according to claim 21, wherein said vector is a plasmid, a recombinant retrovirus, a recombinant herpesvirus, a recombinant poxvirus, a recombinant adenovirus, a recombinant parvovirus, a recombinant alphavirus, a recombinant polyoma virus, or a eukaryotic layered vector initiation system.
23. A cell which contains an expression cassette according to claim 1 or a gene delivery vector according to claim 21.
24. The gene delivery vector according to claim 21 for stimulating an immune response to a selected antigen within a desired host.

25. Use of a gene delivery vector according to claim 21 in the manufacture of a medicament for stimulating an immune response to a selected antigen within a desired host.

dsRNA Expression Cassette I: Hairpin RNA

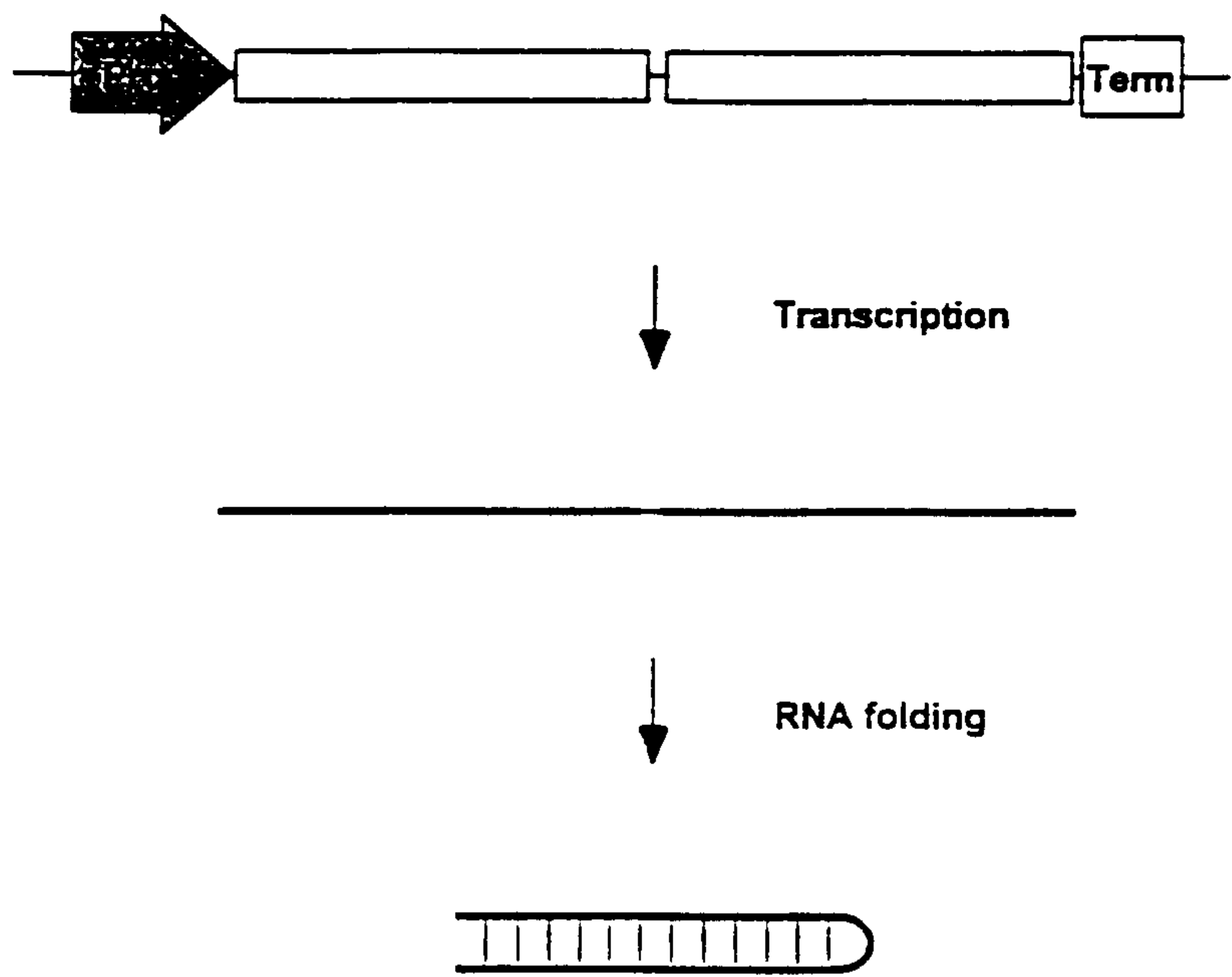
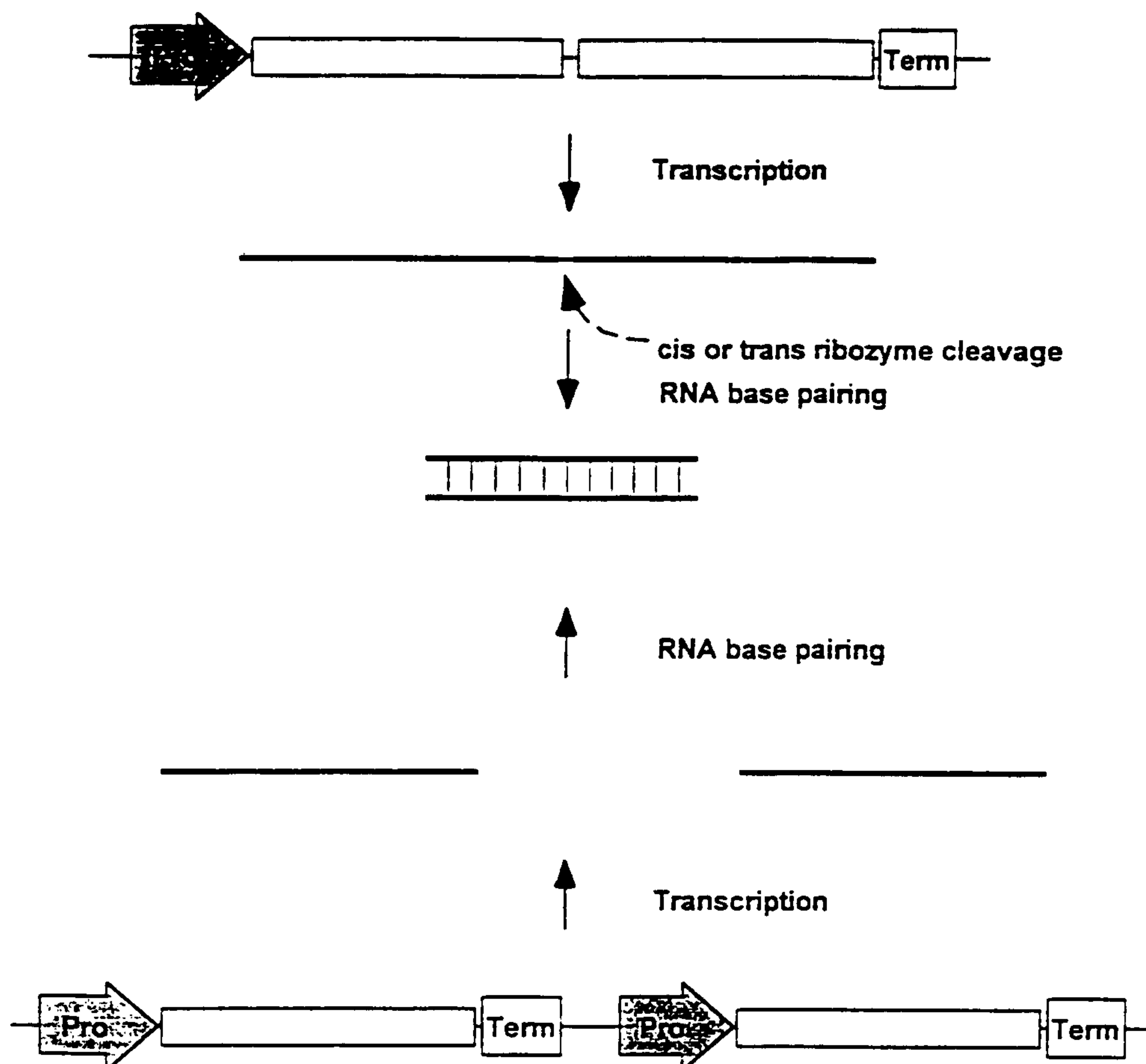
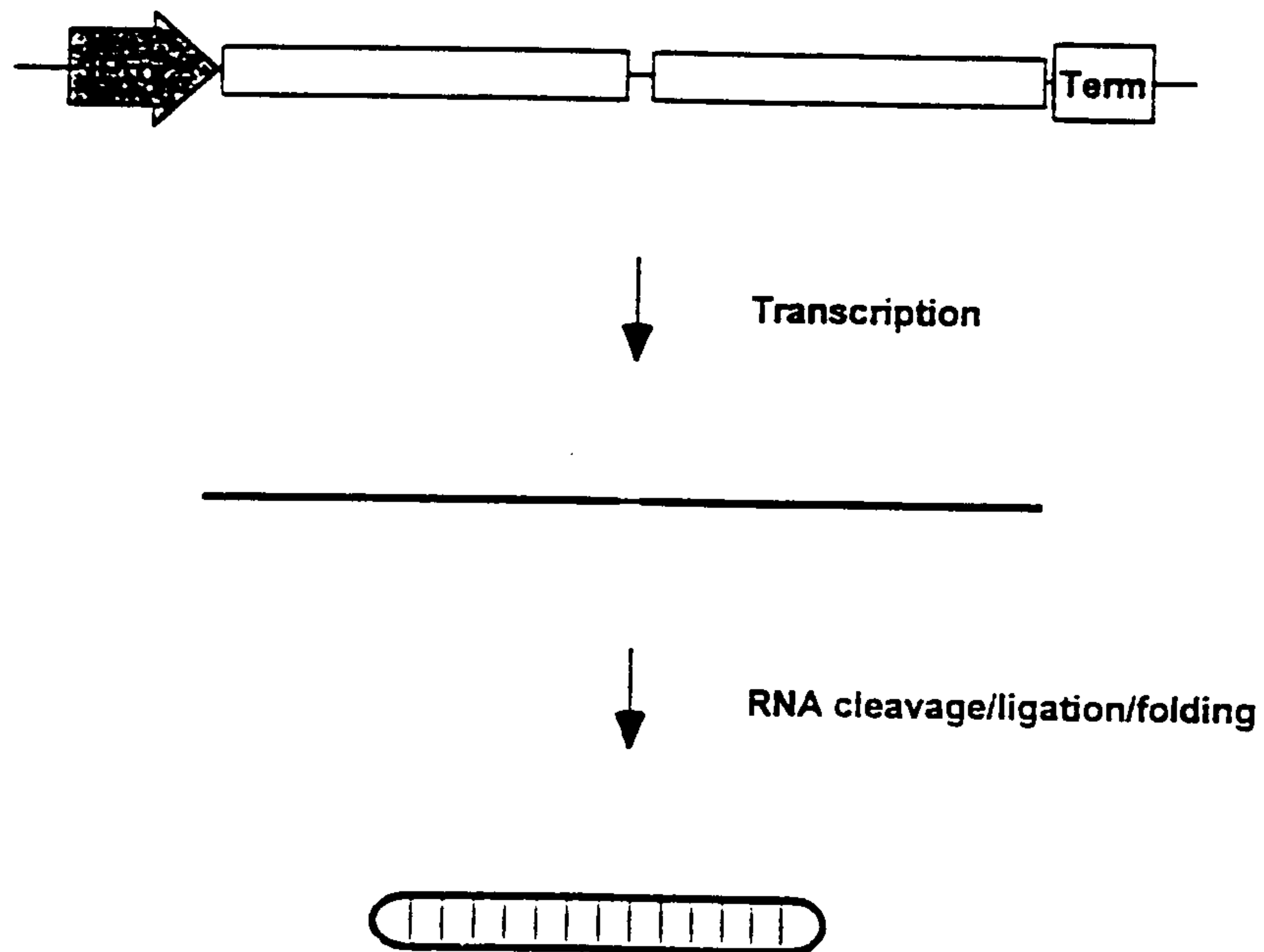


Figure 1

dsRNA Expression Cassette II: "True" dsRNA**Figure 2**

dsRNA Expression Cassette III: Circular RNA**Figure 3**

Plasmid vector with dsRNA and antigen cassettes

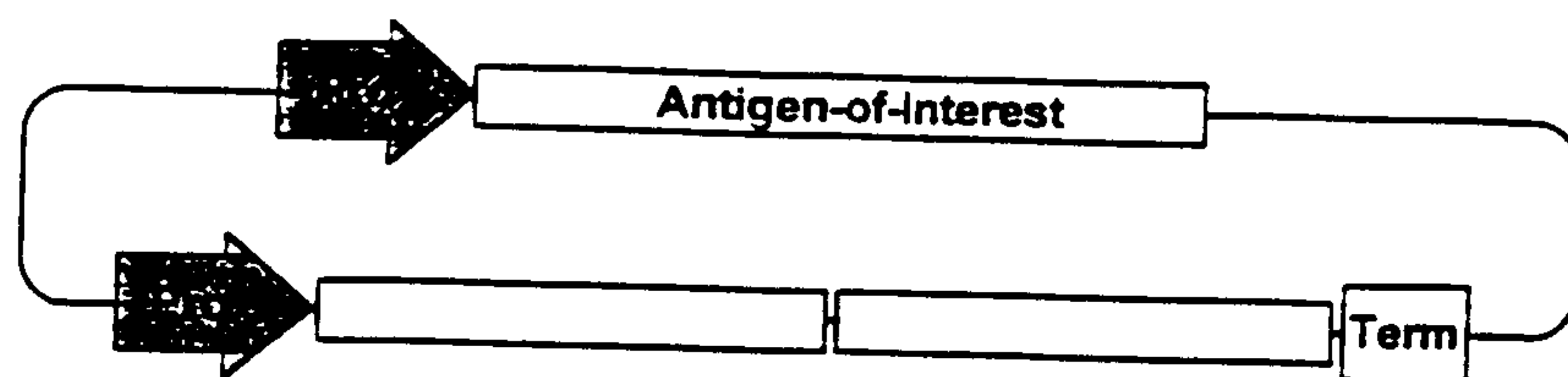
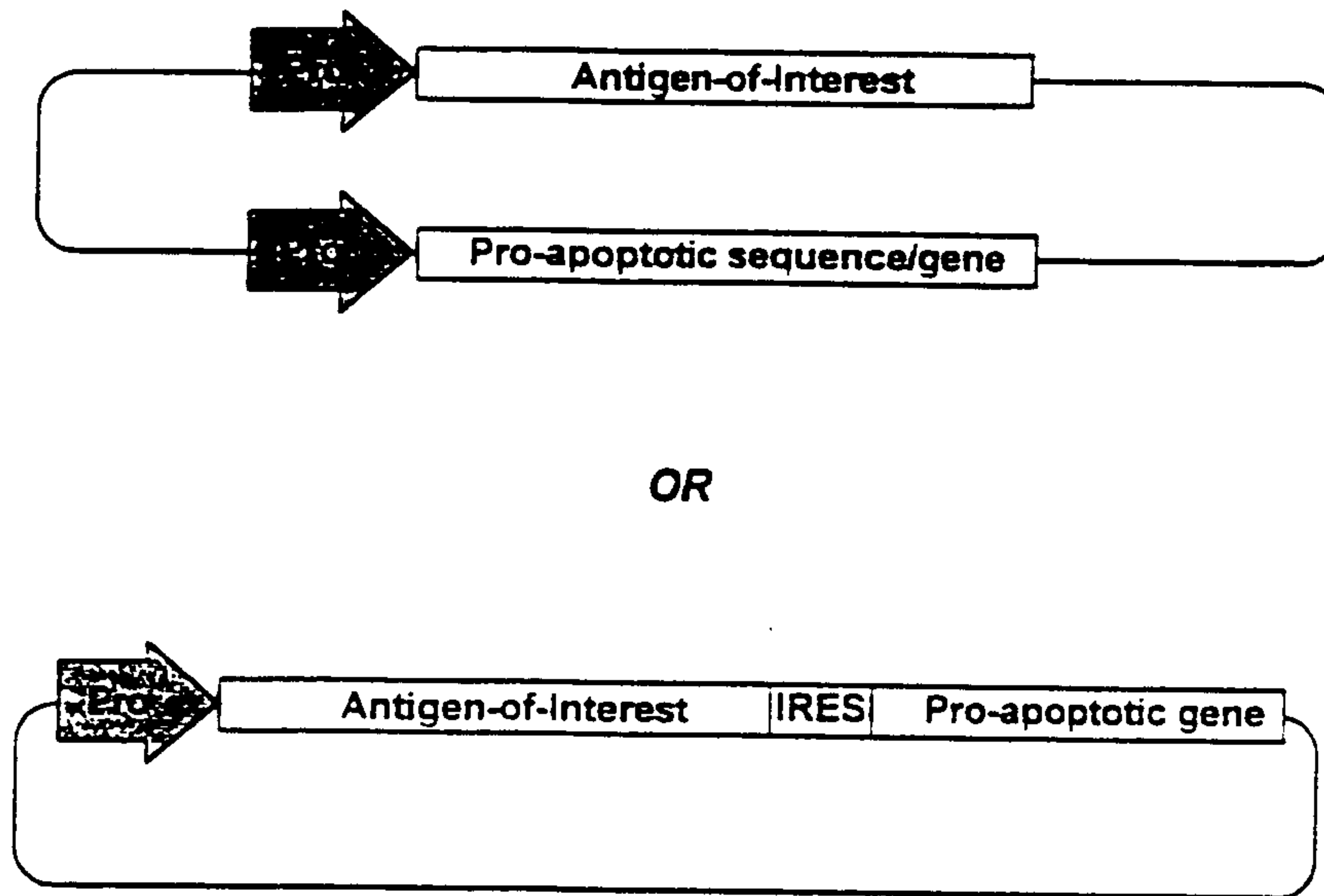


Figure 4

Pro-apoptotic vectors with antigen cassettes**Figure 5**

IL10

1 CTT CTC GGT GTC
GAA GAG CAA CAG

73 C GTC
G CAG

146

220 G GTC
C CAG

COX2

[illegible]

YY1

[illegible]

IRF2

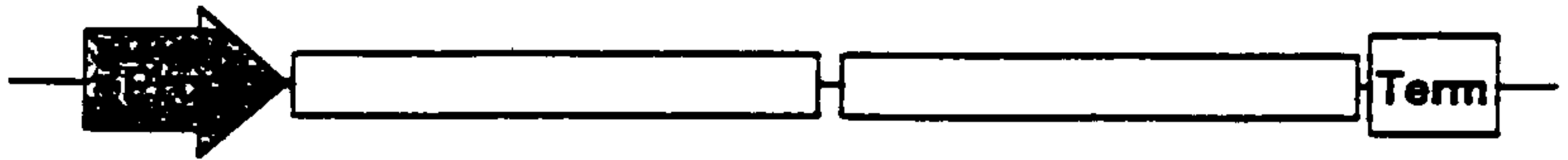
[illegible]

Figure 6

LEGEND

 :starting methionine
 :complementary sequence for the hammerhead ribozyme
 —  :complementary sequence for the hairpin ribozyme
 Boxed sequences:target sequence for the chosen ribozyme
 Bold characters: target sequences for hammerhead ribozymes
 Bold and underlined characters: target sequences for both hammerhead and hairpin ribozymes

dsRNA Expression Cassette I: Hairpin RNA



Transcription



RNA folding

