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(54) Title: POLYNUCLEOTIDE DECOYS THAT INHIBIT MHC-II EXPRESSION AND USES THEREOF			
(57) Abstract			
<p>The invention is directed to a newly discovered class of polynucleotide decoys that is capable of competitively inhibiting the binding of transcription factors to the X-box sequence. This binding is necessary for the expression of MHC-II genes. The invention is also directed to methods of preparing these polynucleotide decoys, and methods of use thereof. In particular, we have identified a class of polynucleotide decoys that mimic the X-Box of MHC-II and competitively bind the MHC-II transcription factor RF-X, resulting in the modulation of MHC-II antigen expression. Thus, the invention can be used to inhibit the expression of HLA molecules on the surface of donor cells or organs, in order to render them invisible to the host's immune system, or in methods of treating an individual with an autoimmune disease characterized by dysfunctional expression of an MHC class II antigen. Further, because of the role of RF-X in the expression of several viral proteins, the polynucleotide decoys of the invention can be used in methods of treating an individual infected with hepatitis B virus, or cytomegalovirus.</p>			

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POLYNUCLEOTIDE DECOYS THAT INHIBIT MHC-II
EXPRESSION AND USES THEREOF

5 Technical Field

This invention relates to nucleotide therapeutics, transplantation and immunology. More specifically, it relates to a newly discovered class of polynucleotide decoy molecules and methods for making cells and organs
10 that are less likely to be rejected when transplanted into a recipient host using these polynucleotide decoys, which are capable of binding to a specific gene regulatory factor and reducing the expression of MHC class II transplantation antigens.

15

Background of the Invention

Among gene products that relate to transplantation antigens are the products of the Human Leukocyte Antigen (HLA) complex, also known as the
20 major histocompatibility complex (MHC), located on the short arm of chromosome 6. The HLA antigens are divided into two classes depending on their structure. The genetic loci denoted HLA-A -B, and -C code for the HLA Class I antigens, and HLA-DP, -DQ and -DR code for
25 the HLA Class II antigens.

HLA Class II molecules are composed of two non-covalently linked glycoproteins, the α chain and the highly polymorphic β chain. Each chain contains one
30 extracellular domain, a transmembrane segment and a cytoplasmic tail. The structure of the α and β chains and their genes have been elucidated. All known Class II genes are similar in structure and encoded by exons 1 - 4, with exon 5 coding for an untranslated region. The DP, DQ and DR loci all consist of multiple genes.
35 A total of twelve class II genes have been identified.

In some haplotypes, some class II genes do not code for a functional peptide and are classified as pseudogenes.

Regulation of HLA class II antigen expression occurs in part through a series of promoter regions such as the J, W, X (including X₁ and X₂), and Y boxes, and the gamma interferon response element. The X (including X₁ and X₂) and Y boxes are known to be required in the transcriptional regulation of all class II promoters. Ono, S.J. et al., Proc. Natl. Acad. Sci. (USA) (1991) 88: 4304-4308.

Transcription of HLA-DR α Class II can be activated by RF-X (Regulatory Factors-X) which binds to the X-box region (-110 to -95) of the DR α promoter. RF-X and its binding site, the X-box are unique and have a high specificity for each other. The DNA binding domain of RF-X consists of 91 amino acids with a basic stretch and shares no notable homology with other known DNA binding motifs (Reith et al. (1990), Genes Dev., Vol. 4(9), pp. 1528-40). RF-X binds only the X-box; substitutions in the X-box are generally not well tolerated by RF-X. (Hasegawa et al. (1991), Nucleic Acids Res., Vol 19(6), pp. 1243-49). The X-box sequence is an atypical promoter site, being neither palindromic nor dyad symmetric. Additionally no other sequence (using the program Eugene) shows exact homology with the X-box. The X-box is conserved in humans. No other known cloned transcription factors bind to this entire region of the X-box in the same manner.

HLA antigens are implicated in the survival of cell grafts or transplants. Although there is acceptable graft survival in the first year for nearly all types of transplants, by five and ten years after transplantation only 40-50% of all grafts are still functioning. This low rate is due to the relentless attack of the immune system on the graft. In addition, death rates of 1-5% are recorded even at the best

transplant centers. Drugs are commonly used to control immune responses and prevent graft rejection, and death is often an indirect result of this drug administration.

5 The drugs used to control immune responses usually cause a non-specific depression of the immune system. A patient with a depressed immune system is far more susceptible to develop life-threatening infections and a variety of neoplasia. The low rate of long term
10 success, and serious risks of infection and cancer are the two main challenges now facing the entire field of tissue and organ transplantation.

 It has been suggested that graft rejection can be prevented or reduced by reducing the levels of exposed
15 HLA antigens on the surface of transplant cells. Faustman, D. et al., Science (1991) 252:1700-1702, observed that xenograft survival was increased by masking HLA class I surface antigens with F(ab')₂ antibody fragments to HLA class I or tissue specific
20 epitopes.

 One way to reduce the level of cell surface transplantation antigens is to retard (downregulate) the expression of the transplantation antigen genes.

 Generally, eucaryotic gene expression may be
25 regulated at any of the steps from DNA transcription to RNA translation to protein; and it is generally agreed that gene expression is at the level of transcription. In order for transcription to occur, transcription factors must bind distinct regulatory sites or
30 promoters on the gene. Once bound, transcription factors may interact with RNA polymerase or other factors to activate or repress transcription. Some transcription factors are constitutively expressed in specific cells while others may be transiently
35 activated in response to various physiological signals (such as cAMP, IFN- γ , etc.). Thus in a given cell transcription of particular genes depends on which

transcription factors are present in that cell type and/or whether the signals to activate the transcription factors are present.

Agents such as actinomycin (an intercalator) have
5 been used to block transcription in a nonspecific manner. A variety of approaches to sequence-specific gene modulation include use of antisense oligonucleotides and antigene oligonucleotides (triple helix formers). These are limited in general or in
10 particular instances. Antisense oligonucleotides block gene expression by targeting mRNA while triple helix forming oligos target double-stranded DNA. Inaccessibility of the target mRNA due to RNA secondary structure can limit the usefulness of antisense
15 methods; triple helix approaches are limited by poor nuclear access, chromatin structure (bypassing histones), and the need for targeting a homopurine-homopyrimidine stretch. Degradation of oligonucleotides by exonucleases and excessive binding
20 to untargeted cellular factors can limit the effectiveness of both antisense and triple helix methods. Chemical modification of oligos can improve nuclease resistance but can also result in increased toxicity, reduced binding affinity, and lower activity
25 (Cook 1991, Crooke 1991).

J.T. Holt (1991), Antisense Res. Dev., Vol. 1(4), pp. 65-9, has shown that by providing excess DNA binding sites, specific transcription factors can be quenched and are thereby prevented from binding to
30 endogenous DNA.

Chu et al. (1991) Nucl. Acids Res. 19: 6958, describe DNA structures in "hairpin" and "dumbbell" configurations containing CRE and TRE sequences, and reported using them *in vitro* as substitutes for regular
35 double-stranded DNA to bind CREB and JUN, respectively, in gel shift assays.

Chu et al. (1992) Nucl. Acids Res. 21: 5857-5858, demonstrate that in nuclear extracts, dumbbell DNA is much more stable than double-stranded or hairpin DNA. However, the nicking of dumbbell DNA in human serum, caused by endonuclease degradation of the single-stranded loops, is a potential problem, since it converts dumbbell DNA to a double-stranded form that is no more stable in the nucleus than standard double-stranded DNA molecules.

10 International patent publication WO92/19732 describes "closed" oligonucleotides that can be used as "sense" or "antisense" molecules, with the advantage of being resistant to exonucleases. Among the variety of "closed" structures are "dumbbell" configurations in
15 which the ends are closed by virtue of addition links of thymidine nucleotides. Use of the closed "sense" oligonucleotides to bind protein factors having an affinity for RNA or DNA sequences or structures is suggested.

20 Single-strand circular DNA where a portion becomes double-stranded have been used as an experimental system for studying local thermal stability in DNA (Wemmer et al. (1985), Nucl. Acid. Res., Vol. 13(23), pp. 8611-21); as models for hairpins and cruciforms
25 (Erie et al. (1989), Biochemistry, Vol. 28(1), pp. 268-73); and as models for comparison to nicked or gapped DNA (Snowden-Ifft et al. (1990), Biochemistry, Vol. 29(25), pp. 6017-25).

30 Summary of the Invention

We have discovered a class of polynucleotide decoys that competitively bind transcription factors necessary for transcription of MHC-II genes. Particularly, for example, we have identified
35 oligonucleotides that mimic the X-Box of MHC-II and that competitively bind the MHC-II transcription factor RF-X. Exposing the cell to such an oligonucleotide can

result in production of a MHC-II-depleted cell. That is, the polynucleotide decoys of the invention act as modulators of MHC-II expression. The invention can be used, therefore, to inhibit the expression of HLA
5 molecules on the surface of donor organ/cell, in order to reduce its visibility to the host's immune system.

The polynucleotide decoy of the invention is a centrally double-stranded (but potentially lacking as many as two base pairings), "dumbbell" shaped
10 structure, with linking cap oligonucleotides that is covalently closed in some embodiments and in other embodiments is linear. The single-stranded caps linking each of the two ends of the polynucleotide decoy are present to stabilize the structure, prevent
15 degradation by exonucleases, and are not intended to be involved in recognition of the protein of interest. Compared to antisense and antigene approaches, target inaccessibility may be of less concern for the polynucleotide decoy, because its target is a protein
20 which may be found in the cytoplasm as well as the nucleus. Lower doses of the polynucleotide decoy may suffice to effectively block transcription of the gene of interest, as compared for example to dosages of nucleotides required for an antisense approach, because
25 transcription factors are ordinarily present in low copy number.

Preferred polynucleotide decoys are short covalently closed nucleotides, with an entirely double-stranded internal segment, and wherein the
30 polynucleotide decoys have a sequence that mimics the double-stranded sequence of the X-box, with polythymidine segments linking the ends together. Administration of a polynucleotide of the invention results in downregulation of HLA II DR α proteins or
35 mRNA. The polynucleotide decoys of the invention can be useful in regulating gene expression in the HLA system and in other systems as well.

The invention contemplates the development of a "universal donor cell" reduced in one or more HLA antigens by treatment of the cell with a polynucleotide decoy of the invention. The absence of certain HLA
5 antigens on the surface of donor cells, tissues or organs comprising these cells will cause them not to be recognized as foreign and not to elicit a rejection response. By the selective introduction of polynucleotide decoys into a cell it is possible to
10 block the expression of MHC class II genes, particularly constitutive expression, thereby rendering a graft "invisible" to the immune system. Thus, the problem of rejection is eliminated without nonspecific suppression of the immune system, and the immune system
15 remains active to defend against infection and neoplasia. By implementing the methods of the present invention, even xenogeneic donor cells, tissues and organs can be rendered "invisible" to the immune system. Thus, the target cells or organs of the
20 present invention may be derived from non-human mammalian species, made into transplantation antigen-depleted cells or universal donor organs, respectively, by a method of the invention, and subsequently transplanted into a human individual.

25

In particular the present invention embodies a class of polynucleotide decoys which comprise: (a) an internal oligonucleotide (I) having a length of X bases, where X is a number from about 10 to about 40;
30 (b) two cap oligonucleotides (P_1 and P_2), each having a length of from about 3 to about 8 bases, wherein each of said cap oligonucleotides is comprised of bases which are unable to bind to any other base within the same cap oligonucleotide; (c) a first complementary
35 oligonucleotide (C_1) having a length of Q bases, where Q is a number from about 5 to $(X - 5)$, said C_1 having a 3' to 5' nucleic acid sequence capable of Watson-Crick-

type binding to the first Q bases in the 5' to 3' nucleic acid sequence of said I; and (d) a second complementary oligonucleotide (C_2) having a length of Z bases, where Z is a number greater than equal to about 5 and from $((X - (Q + \text{about } X/8)))$ to $(X - Q)$, said C_2 having a 5' to 3' nucleic acid sequence capable of Watson-Crick-type binding to the first Z bases in the 3' to 5' nucleic acid sequence of said I; wherein the 3' end of said C_1 is covalently linked to the 5' end of said P_1 , the 3' end of said P_1 is covalently linked to the 5' end of said I, the 3' end of said I is covalently linked to the 5' end of said P_2 , and the 3' end of said P_2 is covalently linked to the 5' end of said C_2 ; wherein at least one of the oligonucleotides selected from the group consisting of said C_1 , said C_2 , and said I, comprises an RF-X recognition sequence of Type-1 or Type-2; and wherein said polynucleotide decoy is capable of binding to an RF-X transcription factor.

The present invention also embodies methods of making the polynucleotide decoys of the invention comprising covalently linking oligonucleotides such that the resulting product has the structure of a polynucleotide decoy of the invention.

The present invention embodies methods for making an MHC-II-depleted cell from a target cell, the method comprising: (a) obtaining the target cell; and (b) exposing the target cell to a polynucleotide decoy of the invention, said polynucleotide decoy being preset in an amount sufficient to make the target cell a MHC-II-depleted cell. The present invention additionally embodies the MHC-II-depleted cells prepared by the fore-mentioned method.

The present invention further embodies polynucleotide decoys of the invention for use in the preparation of a composition for treating target cells to make them MHC-II-depleted cells.

The present invention further embodies MHC-II-depleted donor organs prepared by the method comprising: (a) obtaining a target organ from an individual; and (b) exposing the target organ to a
5 polynucleotide decoy of the invention, said polynucleotide decoy being present in an amount sufficient to make the target organ a MHC-II-depleted donor organ.

The present invention further embodies methods of
10 treating an individual with an autoimmune disease characterized by dysfunctional expression of an MHC class II antigen, the method comprising administering to that individual a polynucleotide decoy of the invention in an amount sufficient to inhibit expression
15 of the MHC class II antigen.

The present invention also embodies methods of treating an individual infected with hepatitis B virus, the method comprising administering to that individual a polynucleotide decoy of the invention, in an amount
20 sufficient to inhibit expression of hepatitis B surface antigen.

The present invention further embodies methods of treating an individual infected with cytomegalovirus, the method comprising administering to that individual
25 a polynucleotide decoy of the invention in an amount sufficient to inhibit RF-X binding to the enhanced factor C sites of the cytomegalovirus genome.

The present invention further embodies methods of using a polynucleotide decoy to obtain a substantially
30 purified, covalently closed polynucleotide decoy, comprising the steps of: (a) providing a linear polynucleotide decoy of the invention; (b) treating said linear polynucleotide decoy with a kinase enzyme, thereby obtaining 5' phosphorylated polynucleotide
35 decoy; (c) heating said 5' phosphorylated polynucleotide decoy to inactivate said kinase enzyme, thereby obtaining heated, 5' phosphorylated

polynucleotide decoy; (d) cooling said heated, 5' phosphorylated polynucleotide decoy mixture slowly to facilitate intramolecular annealing, thereby obtaining cooled, 5' phosphorylated polynucleotide decoy; (e) 5 treating said cooled, 5' phosphorylated polynucleotide decoy with a ligating enzyme, thereby obtaining a covalently closed polynucleotide decoy; and (f) purifying said covalently closed polynucleotide decoy, thereby obtaining a substantially purified, covalently 10 closed polynucleotide decoy.

The present invention also embodies nucleic acids comprising a linear polynucleotide decoy of the invention.

The present invention further embodies host cells 15 comprising a nucleic acid comprising a linear polynucleotide decoy of the invention.

Brief Description of the Drawings

Figs. 1 - 4 are half-tone reproductions from 20 minigels, visualized with ethidium bromide on an ultraviolet illuminator, showing migration of, and degradation of, the 42-mer oligonucleotide and the polynucleotide decoy, both untreated (Fig. 1) and following various enzymatic treatments, namely (Fig. 2) klenow fragment, (Fig. 3) shrimp alkaline phosphatase, 25 and (Fig. 4) S1 nuclease.

Fig. 5 is a half-tone reproduction from a gel from an electrophoretic mobility shift assay of RF-X 30 incubated with various nucleotides at various concentrations.

Fig. 6 A-C are graphs summarizing the effects of various decoys and control sequences on the CAT activity of Raji cells transfected with the plasmid pDRASCAT which contains the DRA promoter from positions 35 -150 to +31 linked to the CAT gene.

Fig. 7 is a schematic drawing of one embodiment of the polynucleotide decoy structure according to the invention.

5 Detailed Description of the Invention

The present invention provides an improved source of transplantable cells for medical treatment. These transplantation antigen-depleted cells give rise to improved graft survival rates in the recipient or
10 require lower levels of immunosuppressant drug administration in the recipient. These cells may also be useful in treating patients with autoimmune diseases.

The practice of the present invention encompasses
15 conventional techniques of chemistry, molecular biology, biochemistry, protein chemistry, and recombinant DNA technology, which are within the skill of the art. Such techniques are explained fully in the literature. See, e.g., Oligonucleotide Synthesis (M.J. Gait ed. 1984); Nucleic Acid Hybridization (B.D. Hames & S.J. Higgins, eds., 1984); Sambrook, Fritsch & Maniatis, Molecular Cloning: A Laboratory Manual, Second Edition (1989); PCR Technology (H.A. Erlich ed., Stockton Press); R.K. Scope, Protein Purification
20 Principles and Practice (Springer-Verlag); and the series Methods in Enzymology (S. Colowick and N. Kaplan eds., Academic Press, Inc.).

All patents, patent applications and publications mentioned herein, whether supra or infra, are hereby
30 incorporated by reference in their entirety.

Definitions:

As used herein, the term "transplantation antigen" is used to refer to antigenic molecules that are
35 expressed on the cell surface of transplanted cells, either at the time of transplantation, or at some point following transplantation. Generally these antigenic

molecules are proteins and glycoproteins. The primary transplantation antigens are products of the major histocompatibility complex (MHC), located on chromosome 6 in humans. The human MHC complex is also called the human leukocyte antigen (HLA) complex. MHC antigens are divided into MHC class I antigens (in humans, this class includes HLA-A, -B, and -C antigens) and MHC class II antigens (in humans, this class includes HLA-DP, -DQ, and -DR antigens). Thus, the terms "MHC-II antigens", "MHC class II antigens", and "MHC class II transplantation antigens" are used interchangeably herein to refer to the class of proteins, which in humans, includes HLA-DP, -DQ and -DR antigens. While the terms "MHC class II genes" and "MHC-II genes" are used interchangeably herein to refer to the genes which encode the MHC class II transplantation antigens. The term "MHC-II" is used herein to refer to the gene locus which encodes the MHC class II transplantation antigens, as well as the group of proteins encoded by that locus. Transplantation antigens also include cell surface molecules other than MHC class I and II antigens. These antigens include the following: (1) the ABO antigens involved in blood cell recognition; (2) cell adhesion molecules such as ICAM, which is involved in leukocyte cell-cell recognition; and (3) β_2 -microglobulin, a polypeptide associated with the 44 kd heavy chain polypeptide that comprises the HLA-I antigens but is not encoded by the MHC complex.

As used herein, the term "transplantation antigen nucleotide sequence" refers to nucleotide sequences associated with genes encoding transplantation antigens. Nucleotide sequences associated with genes include the region of the gene encoding the structural product, including intron and exon regions, and regions upstream of the structural gene associated with transcription, transcription initiation (including transcription factor binding sites), translation

initiation, operator and promoter regions, ribosome binding regions, as well as regions downstream of the structural gene, including termination sites.

Nucleotide sequences associated with genes also include
5 sequences found on any form of messenger RNA (mRNA) derived from the gene, including the pre-mRNA, spliced mRNA, and polyadenylated mRNA.

As used herein, the term "MHC-II-depleted cell" refers to cells that are in some way depleted in the
10 expression of at least one MHC class II transplantation antigen. This depletion may be manifested by a reduced amount of antigen present on the cell surface.

Preferably, at least about 50%, more preferably about 80%, and even more preferably about 90% of the MHC-II
15 antigen is eliminated at the cell surface. Most preferably, this depletion results in essentially total absence of the antigen at the cell surface. The amount of MHC-II antigens on the cell surface can be determined by a number of techniques known in the art.
20 Usually such techniques make use of an antibody that specifically binds to the MHC-II antigen, wherein the antibody is linked to a compound that is easily quantified, such as a radionuclide, enzyme (e.g., horseradish peroxidase), or fluorescent dye. The
25 techniques for measuring the amount of MHC-II antigens on the cell surface can be performed on living cells or on dead, fixed cell or tissue samples. Preferably the techniques are performed on living cells using a fluorescence-activated cell sorting analysis and
30 fluorescently tagged antibodies that specifically bind to an MHC-II antigen.

The term "MHC-II-depleted donor organ", as used herein, refers to an organ comprising MHC-II-depleted cells. An MHC-II-depleted cell or an MHC-II-depleted
35 donor organ will have at least one of two properties: (1) the cell or organ will survive in the transplant recipient for time periods significantly longer than

normal cells; or (2) the cell or organ will survive in the transplant recipient for time periods commensurate to normal, or untreated, cells or organs, but will require lower doses of immunosuppressive agents to the
5 transplant recipient.

As used interchangeably herein, the terms "oligonucleotides", "polynucleotides", and "nucleic acids" include RNA, DNA, or RNA/DNA hybrid sequences of more than one nucleotide in either single chain or duplex
10 form. The term "nucleotide" as used herein as an adjective to describe molecules comprising RNA, DNA, or RNA/DNA hybrid sequences of any length in single-stranded or duplex form. The term "nucleotide" is also used herein as a noun to refer to individual
15 nucleotides or varieties of nucleotides, meaning a molecule, or individual unit in a larger nucleic acid molecule, comprising a purine or pyrimidine, a ribose or deoxyribose sugar moiety, and a phosphate group, or phosphodiester linkage in the case of nucleotides
20 within an oligonucleotide or polynucleotide. Although the term "nucleotide" is also used herein to encompass "modified nucleotides" which comprise at least one modifications (a) an alternative linking group, (b) an analogous form of purine, (c) an analogous form of
25 pyrimidine, or (d) an analogous sugar, all as described herein.

Generally the terms "oligonucleotides", "polynucleotides", and "nucleic acids" are used herein to refer to "straight-chained" molecules, in which the
30 individual nucleotides are joined by linking groups which bridge the 5' carbon of one nucleotide's sugar moiety to the 3' carbon of a second nucleotide's sugar moiety. Although "branch-chained" nucleic acids, including those with linking groups bridging carbons
35 other than 3' and 5' of two nucleotide may be used in the invention, particularly in the attachment of targeting compounds to polynucleotide decoys. Thus, a

single-stranded nucleic acid molecule can either be "linear" or "covalently closed". "Covalently closed" nucleic acids (or polynucleotide decoys) form a continuous circle of nucleotides without an end. In the case of "linear" nucleic acids or (polynucleotide decoys), there would be at least two ends, exactly two ends in a straight-chained molecule and more in a branch chained molecule. Therefore, a straight-chained "linear" nucleic acid or decoy would have both a "5' end", in which the 5' carbon of the terminal nucleotide is not joined to any other nucleotide, and a "3' end", in which the 3' carbon of the terminal nucleotide is not joined to any other nucleotide. The term "linear" is used herein to refer to any nucleic acid molecules regardless of the degree of folding or degree to which the bases of an individual molecule bind to one another.

The phrase "having a length of N bases" or "having a length of N nucleotides" is used herein to describe lengths along a single nucleotide strand, of a nucleic acid molecule, consisting of N individual nucleotides.

As used herein, the term "bind", in the phrase "bases which are unable to bind to any other base", refers to an interaction between the bases of an oligonucleotide which is mediated through base-base hydrogen bonding. One type of binding is "Watson-Crick-type" binding interactions in which adenine-thymine (or adenine-uracil) and guanine-cytosine base-pairs are formed through hydrogen bonding between the bases. An example of this type of binding is the binding traditionally associated with the DNA double helix.

The term "complementary" and "complementary oligonucleotide" are used herein to refer to oligonucleotides or portions of polynucleotides which are capable of forming Watson-Crick-type binding interactions with another particular oligonucleotide or

particular region of a polynucleotide. Generally, unless otherwise noted, the use of the term "complementary" means that all of the bases of the shorter of the two nucleotides, or portions of the 5 nucleotides being discussed, are capable of Watson-Crick-type binding to a particular region of the other longer, or equal sized, oligonucleotide.

The term "substantially purified" is used herein to describe a polynucleotide or polynucleotide decoy 10 sample which has been separated from other compounds including, but not limited to other nucleic acids, and proteins (such as the enzymes used in the synthesis of the decoy), or the separation of covalently closed polynucleotide decoys from linear decoys. A 15 polynucleotide decoy is substantially pure when at least about 60 to 75% of a sample exhibits a single polynucleotide sequence and conformation (linear versus covalently close). A substantially pure polynucleotide decoy typically comprises about 60 to 90% W/W of a 20 nucleic acid sample, more usually about 95%, and preferably is over about 99% pure. Polynucleotide purity or homogeneity may be indicated by a number of means well known in the art, such as agarose or polyacrylamide gel electrophoresis of a sample, 25 followed by visualizing a single polynucleotide band upon staining the gel, as exemplified in Example 7 below. For certain purposes higher resolution can be provided by using HPLC or other means well known in the art.

30 The "polynucleotide decoys" (also referred to herein as just "decoys") of the invention are derived from the DNA sequence of the X-box, the latter having a sequence as follows:

35 5' -CCCCTAGCAACAGATG-3'
3' -GGGGATCGTTGTCTAC-5'

The polynucleotide decoys of the invention comprise an internal oligonucleotide (I) having a length of X bases, where X is a number from about 10 to about 40, preferably 12 to 25, most preferably 14 to 20. The size of the I segments is bounded on the lower end by their ability to maintain the relative binding affinity of the larger segments to RF-X transcription factors. The size of the I segments is bounded on the upper end by their ability to remain relatively insensitive to endonucleases. Thus, the length limits of the I segment of a decoy can be determined empirically by one of skill in the art.

The polynucleotide decoys of the invention further comprise two cap oligonucleotides (P_1 and P_2), each having a length of from about 3 to about 8 bases, preferably 4 to 6 base most preferably 4 bases. Each of the cap oligonucleotides is comprised of bases which are unable to bind to any other base within the same cap oligonucleotide. Preferably each of the cap oligonucleotides consists of a single variety of nucleotide comprising a base selected from the group consisting of adenine, cytosine, thymidine, and modified nucleotides thereof. Most preferably both of the cap oligonucleotides consist of a single variety of nucleotide comprising a thymidine base.

The polynucleotide decoys of the invention further comprise a first complementary oligonucleotide (C_1) having a length of Q bases, where Q is a number from about 5 to $(X - 5)$, said C_1 having a 3' to 5' nucleic acid sequence capable of Watson-Crick-type binding to the first Q bases in the 5' to 3' nucleic acid sequence of said I. Thus, the entire C_1 segment beginning with its 3' end is able to bind to the first Q bases of the I segment beginning with its 5' end.

The polynucleotide decoys of the invention further comprise a second complementary oligonucleotide (C_2) having a length of Z bases, where Z is a number greater

than equal to about 5 and from $(X - (Q + \text{about } X/8))$ to $(X - Q)$, said C_2 , having a 5' to 3' nucleic acid sequence capable of Watson-Crick-type binding to the first Z bases in the 3' to 5' nucleic acid sequence of said I. Thus, the entire C_2 segment beginning with its 5' end is able to bind to the first Z bases of the I segment beginning with its 3' end, and the C_1 and C_2 segments are at least about five nucleotides in length. The minimal length of the complementary oligonucleotides is bounded on the lower end by their ability to effectively anneal to the I segment and maintain the polynucleotide decoy's relative insensitivity to exonucleases. Thus, a lower limit on the size of the C_1 and C_2 segments can be determined empirically by one of skill in the art. The upper limit on the size of each of the C_1 and C_2 segments is determined by the size of the I segment and the size of the other complementary oligonucleotide, such that the combined length of the C_1 and C_2 segments can not be longer than the I segment. However, there can be a gap between the C_1 and the C_2 segments, such that there are up to about one eighth the nucleotides of the I segment ($X/8$) which do not form Watson-Crick-type base pairing with any nucleotide of the C_1 and C_2 segments. Thus, a decoy of the invention comprising an I segment of 16 nucleotides in length could have a gap between its C_1 and C_2 segments of up to about 2 nucleotides. The maximal length of this gap is bounded by the decoy's relative insensitivity to exonucleases. Thus the maximal length of this gap between the C_1 and the C_2 segments can be determined empirically by one of skill in the art. Preferably the C_1 and C_2 segments are of roughly equal length, and the polynucleotide decoys of the invention contain no gap between the C_1 and the C_2 segments.

The oligonucleotides described above are covalently linked to form a polynucleotide decoy of the

invention, wherein the 3' end of the C₁ segment is covalently linked to the 5' end of the P₁ segment, the 3' end of the P₁ segment is covalently linked to the 5' end of the I segment, the 3' end of the I segment is covalently linked to the 5' end of the P₂ segment, and the 3' end of the P₂ segment is covalently linked to the 5' end of said C₂.

Finally the polynucleotide decoys of the invention are comprised of an RF-X recognition sequence of Type-1 or Type-2, in at least one of the oligonucleotides selected from the group consisting of C₁, C₂, and I.

Reith *et al.* (1994) Mol. and Cell. Biol. 14: 1230-1244, demonstrated that the RF-X proteins comprise a family of DNA-binding proteins of which they have cloned three members, RF-X1, RF-X2, and RF-X3, from humans and mice. Members of the RFX family constitute the nuclear complexes that have been previously referred to as enhancer factor C, EP, methylation-dependent DNA-binding protein, and rpL30 α . RF-X proteins share five strongly conserved regions which include the two domains required for DNA binding and dimerization, resulting in very similar DNA-binding specificities.

Reith *et al.* (1994) Mol. and Cell. Biol. 14: 1230-1244 have proposed a set of rules governing RF-X transcription factor binding to native double-stranded DNA sequences with the following consensus sequence: RTY*RYAY*RGY*RAY, where R is a purine, Y is a pyrimidine, and Y* is either a 5-methyl cytosine or a thymidine residue.

However, within a polynucleotide decoy of the present invention, an RF-X recognition sequence is derived from the following "decoy RF-X consensus sequence":

35

20

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18
 5' -N T Y* N Y N R Y* N N N N A Y R R A N-
 3'

5 where:

N is any nucleotide;

R is a nucleotide comprising a purine, preferably an adenine or guanine deoxyribonucleotide;

10 Y is a nucleotide comprising a pyrimidine, preferably a cytosine or thymidine deoxyribonucleotide;

Y* is a nucleotide comprising a 5-methylated pyrimidine, preferably a 5-methyl cytosine or a thymidine deoxyribonucleotide;

15 A is a nucleotide comprising an adenine, preferably an adenine deoxyribonucleotide;

the nucleotide at position 3 may also be any nucleotide comprising a pyrimidine, preferably a cytosine, a 5-methyl cytosine, or a thymidine deoxyribonucleotide; and

20 the nucleotide at position 10 is preferably a nucleotide comprising guanine, more preferably a guanine deoxyribonucleotide.

25 The term "RF-X recognition sequence" is used herein to refer to both "Type-1 RF-X recognition sequences" and "Type-2 RF-X recognition sequences", each defined hereafter.

30 As used herein, the term "Type-1 RF-X recognition sequence" refers to a nucleotide sequence, which: (1) has a span of the decoy RF-X consensus sequence of at least about 8 nucleotides in length; (2) comprises at least one of the spans selected from the following groups: (a) the nucleotides from positions 3 through 8 (5'-Y*-N-Y-N-R-Y*-3'), and (b) the nucleotides from position 8 through 14 (5'-Y*-N-N-N-N-A-Y-3') of the
 35 decoy RF-X consensus sequence; and (3) may contain up to about 2 non-adjacent changes to the decoy RF-X consensus sequence, said changes being selected from

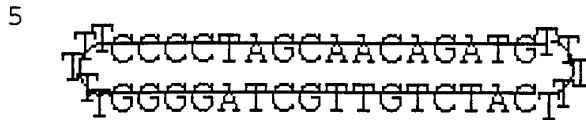
the group consisting of (a) the insertion of a single nucleotide into a span of the decoy RF-X consensus sequence, (b) the deletion of a single nucleotide from the span of the decoy RF-X consensus sequence, and (c) the replacement of a single nucleotide of the span of the decoy RF-X consensus sequence for a spacer compound.

As used herein, the term "Type-2 RF-X recognition sequence" refers to a nucleotide sequence, which: (1) has a span of the decoy RF-X consensus sequence of at least 12 nucleotides in length; (2) comprises at least one of the spans selected from the following groups: (a) the nucleotides from positions 3 through 8 (5'-Y*-N-Y-N-R-Y*-3'), and (b) the nucleotides from position 5 through 14 (5'-Y-N-R-Y*-N-N-N-N-A-Y-3') of the decoy RF-X consensus sequence; and (3) may contain up to 1 non-adjacent change to the decoy RF-X consensus sequence, said changes being selected from the group consisting of (a) the insertion of a single nucleotide into a span of the decoy RF-X consensus sequence, (b) the deletion of a single nucleotide from the span of the decoy RF-X consensus sequence, and (c) the replacement of a single nucleotide of the span of the decoy RF-X consensus sequence for a spacer compound.

Further, the polynucleotide decoys of the invention are capable of binding to an RF-X transcription factor. Such binding can be demonstrated by any method known in the art, including for example methods that directly measure binding like the Electrophoretic Mobility Shift Assay described below in Example 4, and the Gel Shift Assay described below in Example 8, as well as functional assays which indirectly demonstrate polynucleotide decoy/RF-X transcription factor binding like the CAT Assay described below in example 9.

Accordingly, a polynucleotide decoy of the invention mimics the X-box sequence, but is capped by

short DNA oligonucleotides, preferably consisting of thymidine nucleotides, on the ends to improve resistance to exonucleases, for example:



The polynucleotide decoys of the invention are
 10 linear in some embodiments and covalently closed in other embodiments. Initial experiments have indicated that the linear molecules have a marginally high affinity to RF-X transcription factors, while the covalently closed molecules have a marginally greater
 15 stability, being less sensitive to exonucleases. Thus, the choice of linear versus covalently closed decoys can be made on the basis of the requirements of a particular method or use of the invention for affinity and stability.

20 The invention is particularly drawn to the polynucleotide decoys consisting essentially of an oligonucleotide selected from the group consisting of:
 RF-X: 5'-CTAGGGGTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3',
 RF-X-2: 5'-GACTGGGTTTTCCCAGTCCATACGAAGTTTTCTTCGTATG-3',
 25 42 mer: 5'-GCTAGGGGTTTTTCCCCTAGCAACAGATGTTTTTCATCTGTT-3',
 42* mer: 5'-CTAGGGGTTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3',

hlyn
 30 site: 5'-TAACAACTTTTGTTGTTATAGTAACTTTTGTTACTA-3',
 κB: 5'-TCCCTTGGTTTTTCCAAGGACTTTCCGCTTTTGCGGAAAG-3',
 and

38 mer: 5'-TGCTAGGTTTTTCCCTAGCAACAGATGTTTTTCATCTGT-3',
 as well as the polynucleotide decoys formed by
 35 covalently linking the 3' end and the 5' end of an individual decoy such that it is covalently closed.

The polynucleotide decoys and other nucleic acids of the invention usually comprise the naturally-

occurring bases, sugars and phosphodiester linkages. However, any of the hydroxyl groups ordinarily present in the sugars may be replaced by phosphonate groups, phosphate groups, protected by a standard protecting
5 group, or activated to prepare additional linkages to additional nucleotides, or may be conjugated to solid supports. If the 5' and 3' terminal OH groups are free they may be phosphorylated or substituted with amines or organic capping group moieties of from 1 to 20
10 carbon atoms. Other hydroxyl groups may also be derivatized to standard protecting groups.

The individual nucleotides or oligonucleotides which comprise the polynucleotide decoys or other nucleic acids of the invention are usually covalently
15 bonded by phosphodiester linkages, one or more of which may be replaced by alternative linking groups. Thus, the term "covalently linked" is used herein to refer to the joining of two or more nucleotides by phosphodiester linkages or alternative linking groups.
20 These alternative linking groups include, but are not limited to embodiments wherein phosphate is replaced by P(O)S ("thioate"), P(S)S ("dithioate"), P(O)NR₂ ("amidate"), P(O)R, P(O)OR', CO or CH₂ ("formacetal"), wherein each R or R' is independently H or substituted
25 or unsubstituted alkyl (1-20C) optionally containing an ether (-O-) linkage, aryl, alkenyl, cycloalkyl, cycloalkenyl or aralkyl. Not all linkages in an oligomer need to be identical.

Also included within this invention are synthetic
30 procedures in which the resultant oligonucleotides or polynucleotide decoys of the invention include incorporation of analogous forms of purines and pyrimidines. "Analogous" forms of purines and pyrimidines are those generally known in the art, many
35 of which are used as chemotherapeutic agents. An exemplary but not exhaustive list includes aziridinylcytosine, 4-acetylcytosine, 5-fluorouracil,

5-bromouracil, 5-carboxymethylaminomethyl-2-thiouracil, 5-carboxymethylaminomethyluracil, inosine, N6-isopentenyladenine, 1-methyladenine, 1-methylpseudouracil, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-methyladenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid methylester, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid, 5-pentynyluracil and 2,6-diaminopurine. The use of uracil as a substitute base for thymine in deoxyribonucleic acid (hereinafter referred to as "dU") is considered to be an "analogous" form of pyrimidine in this invention.

The oligonucleotides of polynucleotide decoys of the invention may contain analogous forms of ribose or deoxyribose sugars that are generally known in the art. An exemplary, but not exhaustive list includes 2' substituted sugars such as 2'-O-methyl-, 2'-O-allyl, 2'-fluoro- or 2'-azido-ribose, carbocyclic sugar analogs, α -anomeric sugars, epimeric sugars such as arabinose, xyloses or lyxoses, pyranose sugars, furanose sugars, sedoheptuloses, acyclic analogs and abasic nucleoside analogs such as methyl riboside.

Although the conventional sugars and bases may be used in applying the methods of the invention, substitution of analogous forms of sugars, purines and pyrimidines can be advantageous in designing the final product, as can alternative backbone structures like a polyamide backbone.

In addition the polynucleotide decoys of the invention may be comprised of short (the equivalent of up to 3 nucleotides in length) "spacer compounds" which duplicate the length and spacial geometry of a

nucleotide, but do not engage in Watson-Crick-type binding with other nucleotides, and are not subject to cleavage by endonucleases. The various oligonucleotides which comprise the polynucleotide decoys (I, P₁, P₂, C₁, and C₂) of the invention are considered to encompass oligonucleotides in which up to about 3 nucleotides are replaced with one or more of the spacer compounds that are known in the art. Further, such replacements may be used to overcome the sensitivity of a decoy of the invention to particular endonucleases. If, for example, the stability of a particular decoy of the invention were found to be compromised by the cleavage of a particular endonuclease of a target cell, one or more of the nucleotides at the cleavage site could be replaced by a spacer compound.

However, the polynucleotide decoys of the invention are preferably comprised of greater than 50% conventional deoxyribose nucleotides, and most preferably greater than 90% conventional deoxyribose nucleotides.

The polynucleotide decoy sequences of the invention may be prepared by any known method, including synthetic, recombinant, *ex vivo* generation, or a combination thereof, as well as utilizing any purification methods known in the art.

The invention embodies nucleic acids comprising a linear polynucleotide decoy of the invention. The nucleic acids of the invention include expression vectors, amplification vectors, and PCR-suitable nucleic acids. In addition, the invention embodies amplification vectors, which comprise a decoy of the invention, and an origin of replication. Preferably, such amplification vectors further comprise restriction endonuclease sites flanking the decoy, so as to facilitate cleavage and purification of the decoy from the remainder of the amplification vector, and a

selectable marker, so as to facilitate amplification of the amplification vector. Most preferably, the restriction endonuclease sites in the amplification vector are situated such that cleavage at those site
5 would result in no other amplification vector fragments of a similar size.

Thus, such an amplification vector may be transfected into a host cell compatible with the origin of replication of said amplification vector, wherein
10 the host cell is a prokaryotic or eukaryotic cell, preferably a mammalian, insect, yeast, or bacterial cell, most preferably an *Escherichia coli* cell. The resulting transfected host cells may be grown by culture methods known in the art, preferably under
15 selection compatible with the selectable marker (e.g., antibiotics). The amplification vectors can be isolated and purified by methods known in the art (e.g., standard plasmid prep procedures). The decoy of the invention can be cleaved with restriction enzymes
20 that specifically cleave at the restriction endonuclease sites flanking the decoy, and the double-stranded decoy fragment purified by techniques known in the art, including gel electrophoresis. Finally the decoy of the invention can be purified from its
25 complementary fragment by repeatedly heating the double stranded decoy fragments, and allowing them to partially cool while passing them through a column designed to bind regions of the complementary non-decoy strand.

30 Alternatively linear polynucleotides comprising a decoy of the invention may be amplified by PCR. The PCR method is well known in the art and described in, e.g., U.S. Patent Nos. 4,683,195 and 4,683,202 and Saiki, R.K., et al., Science (1988) 239:487-491, and
35 European patent applications 86302298.4, 86302299.2 and 87300203.4, as well as Methods in Enzymology (1987) 155:335-350. The amplified DNA may then be recovered,

in a single-stranded or dumbbell conformation, using conventional techniques, as well as those described above for decoys derived from amplification vectors.

The polynucleotide decoys of the invention can also be derivatized in various ways. The decoys can be derivatized by attaching a nuclear localization signal to it to improve targeted delivery to the nucleus. One well-characterized nuclear localization signal is the heptapeptide PKKKRKV (pro-lys-lys-lys-arg-lys-val). Preferably, in the case of polynucleotide decoys in the form of a closed circle, the nuclear localization signal is attached via a modified loop nucleotide or spacer that forms a branching structure.

Also, if the polynucleotide decoy is to be used for separation of transcription factors, one or more of the oligonucleotides of the decoy may be derivatized to a solid support to permit chromatographic separation. If the decoy is to be used to label the target transcriptional factor or otherwise attach a detectable moiety to this target, the oligonucleotide of the decoy will be derivatized to include a radionuclide, a fluorescent molecule, a chromophore or the like.

If it is to be used *in vivo*, the polynucleotide decoy of the invention may be derivatized to include ligands and/or delivery vehicles which provide dispersion through the blood, targeting to specific cell types, or permit easier transit of cellular barriers. Thus, the polynucleotide decoys of the invention may be linked or combined with any targeting or delivery agent known in the art, including but not limited to, cell penetration enhancers, lipofectin, liposomes, dendrimers, DNA intercalators, and nanoparticles. In particular, nanoparticles for use in the delivery of the polynucleotide decoys of the invention are particles of less than about 50 nanometers diameter, nontoxic, non-antigenic, and comprised of albumin and surfactant, or iron as in the

nanoparticle particle technology of SynGenix. In general the delivery vehicles used to target the polynucleotide decoys of the invention may further comprise any cell specific or general targeting agents known in the art, and will have a specific trapping efficiency to the target cells or organs of from about 5 to about 35%.

The polynucleotide decoys of the invention may be used *ex vivo* in a method of obtaining MHC-II-depleted cells from target cells. The cells are created by incubation of the target cell with one or more of the above-described decoys under standard conditions for uptake of nucleic acids, including electroporation or lipofection. In practicing an *ex vivo* method of treating cells or organs, the concentration of decoys of the invention in a solution prepare to treat target cells or organs is from about 0.1 to about 100 μM , preferably 0.5 to 50 μM , most preferably from 1 to 10 μM .

Alternatively, the oligonucleotides can be modified or co-administered for targeted delivery to the nucleus. Improved oligonucleotide stability is expected in the nucleus due to: (1) lower levels of DNases and RNases; and (2) higher oligonucleotide concentrations due to lower total volume.

Alternatively, the decoys of the invention can be covalently bonded to biotin to form a biotin-decoy prodrug by methods known in the art, and co-administered with a receptor ligand bound to avidin or receptor specific antibody bound to avidin, wherein the receptor is capable of causing uptake of the resulting decoy-biotin-avidin complex into the cells. Receptors that cause uptake are known to those of skill in the art.

Any transplantable cell type is a potential target cell for this invention. Preferably, the target cell is selected from corneal endothelial cells, thyroid

cells, parathyroid cells, brain cells, adrenal gland cells, bone marrow cells, pancreatic islet cells, hepatic cells, lymphoid cells, fibroblasts, epithelial cells, chondrocytes, endocrine cells, renal cells,
5 cardiac muscle cells, keratinocytes, and hair follicle cells. Most preferably, the target cell is selected from corneal endothelial cells, thyroid cells, parathyroid cells, brain cells, adrenal gland cells, bone marrow cells, pancreatic islet cells
10 keratinocytes, and hepatic cells.

In another aspect of this invention, the above-described decoys may be incorporated into a vector through methods well known in the art, and then inserted into the target cell via standard techniques
15 such as electroporation, lipofection, or calcium phosphate or calcium salt mediation. In this fashion, the desired oligonucleotides can be produced *in situ* by the vector. When using an expression vector, the target cell will continue to express an RNA version of
20 a decoy of the invention for at least a period of time following transplant.

Also, this invention is applicable to the field of solid organ transplants. Organs are normally perfused *ex vivo* prior to transplantation. By adding an amount
25 of the above-described oligonucleotides to the perfusion medium, MHC-II-depleted cells can be created from perfusion-accessible cells in the organ to create an MHC-II-depleted organ useful in solid organ transplants. Any transplantable organ is a potential
30 target organ for this invention. Preferably the target organ is selected from the group consisting of heart, liver, kidney, adrenal cortex, lung, intestine, pancreas, cornea and skin. Most preferably, the target organ is selected from the group consisting of
35 heart, kidney, liver, cornea, and skin.

Finally, local administration of the polynucleotide decoys directly into the transplanted

organ within several days after the transplant is within the scope of this invention. Also, sustained releases of the of these drugs are also contemplated.

As described above the polynucleotide decoys of this invention are useful in creating the MHC-II-depleted cells of this invention. These cells are then directly transplanted to an individual. For the purposes of the invention, an individual, treated with the cells or decoys of the invention, can be any animal, preferably any mammal, most preferably a human.

The polynucleotide decoys of this invention are also useful in treating autoimmune diseases characterized by dysfunctional or aberrant expression of an MHC class II antigen. The autoimmune diseases characterized by dysfunctional or aberrant expression of an MHC class II antigen include, but are not limited to, glomerulonephritis, thyroiditis, pancreatitis, rheumatoid arthritis, and primary biliary cirrhosis. In such a case, the oligonucleotides described herein may be administered systemically in an amount sufficient to inhibit expression of an MHC class II antigen, as measured by the techniques described herein.

As Reith *et al.* (1994) Mol. and Cell. Biol. 14: 1230-1244 note, RF-X1 is a transcription factor that was initially cloned by virtue of its affinity for the X-box motif, a conserved *cis*-acting regulatory element present in the promoters of all MHC class II genes from all species examined. RF-X was subsequently also found to bind with high affinity to the inverted repeats known as enhanced factor C (EF-C) sites, which are *cis*-acting regulatory elements present in the enhancers of several unrelated viruses, including hepatitis B virus (HBV), polyomavirus, and cytomegalovirus (CMV). Among these EF-C sites, the functional importance of the site in the HBV enhancer (EnhI) has been clearly demonstrated in several studies, including Dikstein *et*

al. (1990) Mol. Cell. Biol. 10: 3683-3689, Garcia et al. (1993) J. Virol. 67: 3940-3950, Guo et al. (1991) J. Virol. 65: 6686-6692, Ostapchuk et al. (1989) Mol. Cell. Biol. 9:2787-2797, Siegrist et al. (1993) Mol. Cell. Biol. 13: 6375-6384, and Trujillo et al. (1991) Proc. Natl. Acad. Sci. - U.S.A. 88: 3797-3801. Of particular interest, RF-X1 specific antisense oligonucleotides have been shown to inhibit expression of the hepatitis B surface antigen gene, which is under the control of EnhI. In addition in a transient-transfection assays, RF-X1 was shown to be a potent transactivator of EnhI *in vivo* (Siegrist et al. (1993) Mol. Cell. Biol. 13: 6375-6384).

Thus, the invention embodies methods of treating an individual infected with viruses, which contain EF-C sites in their gene regulator elements (e.g., in enhancers or near transcription initiation sites), said methods of treatment comprising the systemic administration of a polynucleotide decoy of the invention to that individual. The viruses, which contain EF-C sites in their gene regulator elements may be determined by the methods described by Reith et al. (1994) Mol. and Cell. Biol. 14: 1230-1244. Such viruses include, for example, hepatitis B and cytomegalovirus.

In the case of treating an individual infected with hepatitis B virus, the decoy of the invention is administered in an amount sufficient to inhibit expression of hepatitis B surface antigen, antibody-based assays for measuring the levels of hepatitis B surface antigen are known in the art. In the case of treating an individual infected with cytomegalovirus, the method comprises administering to that individual a polynucleotide decoy of the invention in an amount sufficient to inhibit RF-X binding to the enhanced factor C sites of the cytomegalovirus genome. Appropriate assays for measuring the RF-X binding to

cytomegalovirus EF-C sites can be developed by replacing the X-box sequences in the Gel shift assay and Cat assays of Examples 8 and 9, with cytomegalovirus EF-C sites, and cytomegalovirus enhancers respectively. The use of the decoys of the invention for the treatment of cytomegalovirus is particularly directed to the prevention of the onset of cytomegalovirus retinitis in HIV infected patients. Thus, the decoys of the invention can be used as a prophylaxis in any HIV infected patient. The HIV-induced suppression of the immune system is thought to allow the normally benign cytomegalovirus to cross the blood-retina barrier, infecting the choroid and retinal tissues, and cause blindness.

Under physiological conditions, the polynucleotide decoys of the invention, whether linear or covalently closed, are considered to exist in a dumbbell conformation. Thus, while all of the polynucleotide decoys of the present invention maintain some sequence homology with the X box of MHC-II, the dumbbell conformation should prevent these polynucleotide decoys from being capable of binding to any transplantation antigen nucleotide sequence.

The mechanism by which the polynucleotide decoys of the invention interfere with or inhibit the production of one or more MHC-II antigens is not always established, and is not a necessary part of the invention. The polynucleotide decoys of the invention are characterized structurally and by their ability to bind to RF-X transcription factors, regardless of the specific mechanisms of RF-X binding, or the mechanism of the effect thereof.

Described below are examples of the present invention which are provided for illustrative purposes, and not to limit the scope of the present invention. In light of the disclosure, numerous embodiments within

the scope of the claims will be apparent to those of ordinary skill in the art.

Examples

5

Example 1: Synthesis of a polynucleotide decoy

A linear, HPLC purified 42-mer oligonucleotide was purchased from Keystone Labs (Menlo Park, CA). The sequence of the linear 42-mer oligonucleotide is as follows:

10 5' -GCTAGGGGTTTTTCCCCTAGCAACAGATGTTTTTCATCTGTT-3'.

The 42-mer oligonucleotide was phosphorylated with T4 polynucleotide kinase, then heated to inactivate the kinase, and subsequently allowed to anneal by cooling slowly to room temperature. It was then ligated with T4 DNA ligase.

Example 2: Purification of a polynucleotide decoy

For large scale purification, the polynucleotide decoy (about 20 μ g per lane) was run on a polyacrylamide gel. The DNA was visualized on a silica gel plate using UV light. The band of interest was excised, "crushed and soaked" in detergent buffer, and purified through a Sep-pak C₁₈ reverse-phase column.

25 A polynucleotide decoy synthesized as described above in Example 1 was run on a 15% polyacrylamide gel. Fig. 1 shows a print from a minigel, visualized with ethidium bromide on an ultraviolet illuminator. The polynucleotide decoy (Lane 5) migrated faster than the unligated linear 42-mer (Lane 8).

Example 3: Proof of Synthesis & Resistance to Nucleases

To demonstrate that the product resulting from the synthesis was indeed the polynucleotide decoy, it was subjected to the following enzymes, each of which is expected to degrade only non-ligated material:

(a) Klenow fragment from *E. coli* DNA polymerase, a 3' to 5' exonuclease;

(b) Shrimp alkaline phosphatase, a 5'-dephosphorylating agent; and

5 (c) Nuclease S1, a nuclease that is relatively single-strand specific at high salt and low enzyme conditions.

Figs. 2 - 4 are prints from minigels showing results from the enzymatic treatments, visualized with
10 ethidium bromide on an ultraviolet illuminator.

Klenow fragment acts as a 3' to 5' exonuclease, degrading from free 3'-hydroxyl termini. The polynucleotide decoy should be resistant to this activity compared to linear sequence. Fig. 3 shows
15 migration of the untreated polynucleotide decoy (Lane 2) and the untreated linear oligonucleotide (Lane 3); after 8 hours of enzyme treatment, the linear oligonucleotide (Lane 4) is completely degraded, while the polynucleotide decoy (Lane 5) migrates as the
20 untreated control, showing that it is resistant to exonuclease; after 12 hours of enzyme treatment, the linear oligonucleotide (Lane 8) has completely degraded, while polynucleotide decoy (Lane 7) has begun to degrade.

25 Shrimp alkaline phosphatase cleaves off phosphate from the 5' end. The polynucleotide decoy should be resistant to this treatment, as it has no free 5' phosphate. Fig. 3 shows identical migration of the treated polynucleotide decoy (Lane 6) and the untreated
30 polynucleotide decoy (Lane 5); treated linear oligonucleotide (Lane 2), linear phosphorylated oligonucleotide (Lane 3), and treated linear phosphorylated oligonucleotide (Lane 4) are all shortened.

35 S1 nuclease degrades single strand DNA more rapidly than double stranded DNA under reaction conditions of high salt and low enzyme concentration.

The polynucleotide decoy should be comparatively resistant to the treatment, and the middle portion is base paired. Fig. 4 shows migration of untreated polynucleotide decoy (Lane 8) and untreated oligonucleotide (Lane 7); after 5 minutes of treatment linear oligonucleotide (Lane 4) has been degraded to a great extent, while the polynucleotide decoy is still present (Lane 5); after 20 minutes of treatment the oligonucleotide has been completely degraded (Lane 1), while the polynucleotide decoy is still present (Lane 2).

Example 4: Electrophoretic Mobility Shift Assays
("EMSA")

Preliminary *in vitro* binding experiments (EMSA) demonstrated specific binding between the polynucleotide decoy prepared in Example 1 and RF-X (of which a sample was obtained from Dr. B.M. Peterlin, Medicine; Microbiology and Immunology, University of California, San Francisco). In such a gel shift assay, the RF-X-polynucleotide decoy complex should show reduced mobility as compared to RF-X alone. Initial studies involved competition for RF-X by the polynucleotide decoy of the invention and by a radiolabeled, double-stranded X-box probe.

The binding assays were performed generally as described in Hasegawa *et al.* (1991), *Nucleic Acids Res.*, Vol. 19(6), pp. 1243-9, under the following conditions. The samples were preincubated for 10 minutes on ice with: binding buffer, polydIdC, RF-X (from rabbit reticulocyte lysate), and a competitor (one of: polynucleotide decoy, X-box probe, non-specific double-stranded 10-mer or 20-mer) or no competitor. Following preincubation, ³²P, end-labeled, double-stranded DRA X-box probe (~30-mer) was added, and allowed to incubate 20 minutes on ice.

The resulting reactants were run on a 5 % native polyacrylamide gel (TBE). Fig. 5 is an autoradiograph of a gel showing the results for RF-X preincubated with: (Lane 1) double strand X box probe (100 excess compared to ³²P-labelled probe); (Lane 2) control (no DNA added); (Lane 3) polynucleotide decoy (100 fold excess); (Lane 4) polynucleotide decoy (200 fold excess); (Lane 5) non-specific double-stranded 10-mer (500 fold excess); (Lane 6) non-specific double-stranded 10-mer (500 fold excess); (Lane 7) control (no DNA added). The ³²P-labeled DRA X-box probe was unable to compete off the polynucleotide decoy from RF-X at selected concentrations.

15 Example 5: Activity of the Polynucleotide Decoys in Cells is Measured by Flow Cytometry:

The effect of the polynucleotide decoy on HLA II DR α gene expression are determined using fluorescence flow cytometry. Fluorescently tagged antibodies are used for fluorescence-activated cell sorting ("FACS") analysis of HLA II DR α cell surface expression. An RNase protection assay is used to determine the levels of HLA II DR α mRNA. The polynucleotide decoy may inhibit constitutively expressed HLA or inducible HLA (for example, γ -IFN inducible), or both inducible and constitutively expressed HLA. Many different human cell lines are used to assay for specific inhibition; as noted above, for example, the B-lymphoma cell line Raji constitutively expresses HLA class II; the HeLa S3 cell line is inducible with γ -IFN.

Example 6: Method for Synthesis of Polynucleotide Decoys:

Oligonucleotides. Oligonucleotides (oligos) were purchased from either Keystone Laboratories (Menlo Park, CA) or Oligos Etc. (Wilsonville, OR). Synthesis of polynucleotide decoys from linear oligos involves

phosphorylation followed by ligation (Wemmer *et al.* (1985), Nucl. Acid. Res., Vol. 13(23), pp. 8611-21; Erie *et al.* (1987), Biochemistry, Vol. 26(22), pp. 7150-9; Erie *et al.* (1989), Biochemistry, Vol. 28(1), pp. 268-73; and Chu *et al.* (1991) Nucl. Acids Res. 19: 6958.

Oligonucleotides used to make polynucleotide decoys.
Internal double stranded sequences underlined.

10 5 T oligos

42 mer: 5'-CTAGGGGTTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3'

38 mer: 5'- TGCTAGGTTTTTCCCTAGCAACAGATGTTTTTCATCTGT-3'

15 4 T oligos, all 40 mers

RF-X-2: 5'-GACTGGGTTTTCCCAGTCCATACGAAGTTTTCTTCGTATG-3'

RF-X: 5'-CTAGGGGTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3'

NF-κB: 5'-TCCCTTGGTTTTCCAAGGGACTTTCCGCTTTTGCGGAAAG-3'

M6

20 mutant: 5'-TATACGGGTTTTCCCGTATACTACTCTGTTTTTCAGAGTGG-3'

κB

mutant: 5'-TGAGTTGGTTTTCCAACTCACTTTCCGCTTTTGCGGAAAG-3'

κB: 5'-TCCCTTGGTTTTCCAAGGGACTTTCCGCTTTTGCGGAAAG-3'

hlyn

25 site: 5'-TAACAACCTTTTGTTGTTATAGTAACTTTTGTTACTA-3'

Annealed oligos:

Top Strand 5'-CCCCTAGCAACAGATG

Bottom Strand 3'-GGGGATCGTTGTCTAC

30

Phosphorylation. 100 nmol of oligo was phosphorylated with T4 polynucleotide kinase (USB), 5 units/nmol, 2 mM ATP (Mg salt), 500 mM Tris HCl pH 7.6, 10 mM MgCl₂, 10 mM 2-mercaptoethanol at 37°C. After 1 hour

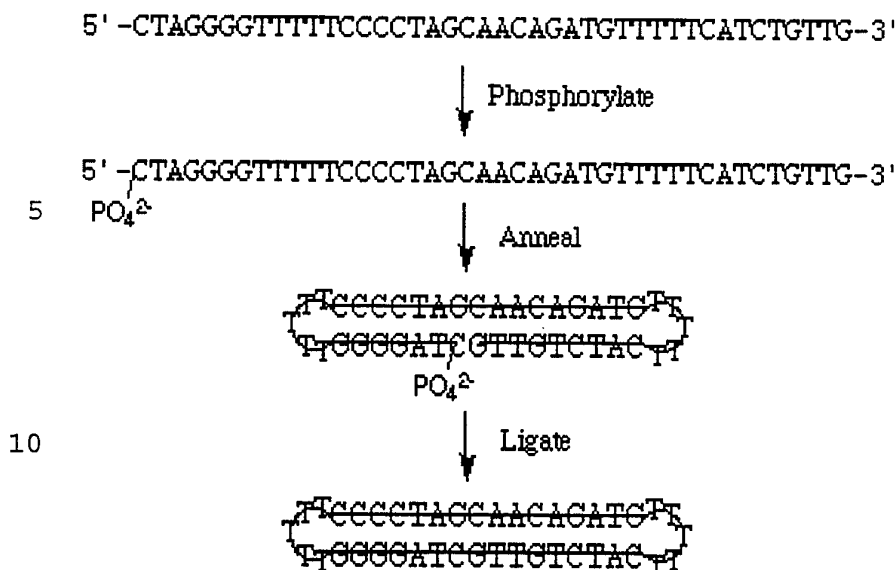
35 additional ATP and 1 unit/nmol T4 polynucleotide kinase were added; the reaction was incubated for 1 hour at 37°C. A final aliquot of ATP and T4 polynucleotide

kinase was added, and the reaction was incubated for 24 hours at 37°C.

Ligation. The phosphorylated oligo reaction mixture
5 was heated to 80-90°C for 10 minutes to inactivate the
kinase and to remove secondary structure. It was
cooled slowly to room temperature (3 hours) to
facilitate intramolecular annealing. Ligation was
performed with a 3-fold molar excess of T4 DNA ligase
10 (USB), 5% polyethylene glycol, 66 mM Tris HCl pH 7.6,
6.6 mM MgCl₂, 10 mM DTT, and 66 mM ATP at 16°C for 48
hours. The sample was frozen at -20°C.

Nuclease Reactions. The reaction product (at various
15 stages) was tested for resistance to various nucleases:
shrimp alkaline phosphatase (USB), Klenow fragment of
e. coli (Worthington), and S1 nuclease (Worthington),
and Mae I (Boehringer Mannheim).

20 *Purification.* Ligated material was separated from
unreacted material on a 15% denaturing polyacrylamide
gel. For the gel shift assay, ligated products were
sliced from the gel and transferred to a microfuge
tube, then crushed and soaked (in TE, pH 7.8) overnight
25 in a 37°C shaking water bath. It was then Speed Vac'd
(Savant) until dry. The decoy was then reconstituted
to a concentration of 10 pmol/ml for the gel shift
assay. For cell studies, oligos were eluted from gel
slices using BioRad's Model 422 electro-eluter, then
30 ethanol precipitated.



Results of Synthesis. Successful synthesis was
 15 determined by analyzing the products on a
 polyacrylamide gel and subjecting the products to
 various nucleases.

Nuclease reactions. The RF-X polynucleotide decoy was
 found to be more resistant to the exonucleitic action of
 20 the Klenow fragment compared to the non-ligated
 sequence. The polynucleotide decoy was degraded more
 slowly by S1 nuclease (a relatively single-strand
 specific nuclease) compared to the non-ligated
 sequence. The polynucleotide decoy was completely
 25 resistant to shrimp alkaline phosphatase. In addition,
 the polynucleotide decoy was cleaved more readily into
 2 segments by the restriction enzyme Mae I, which
 recognizes the double-strand sequence 5'-CTAG found in
 the polynucleotide decoy. All nuclease reactions were
 30 analyzed using gel electrophoresis as described in
 Example 2.

Example 7: Staining to Distinguish Between Covalently
 Closed Polynucleotide Decoys & Single-stranded
 35 Oligonucleotides

Often these decoys are analyzed on polyacrylamide
 gels to determine purity (i.e., shorter failed

sequences) or to check for secondary structure (such as hairpins or dumbbells). Two non-radioactive stains are used to visualize such decoys on polyacrylamide gels: ethidium bromide and methylene blue. Methylene blue
5 has been touted as an inexpensive, non-toxic alternative to ethidium bromide (Young-Sharp, D. and R. Kumar. (1989) Technique 1(3): 183-187), staining both single- and double-stranded oligos with equal intensity. For most purposes methylene blue is
10 adequate for staining short oligos. Ethidium bromide has an advantage when looking for oligo secondary structure because it preferentially stains double-stranded DNA. However, ethidium bromide is inefficient for visualizing short single-stranded
15 oligos at low concentrations. By using these dyes sequentially we can both detect single- and double-stranded oligos at low concentrations and differentiate between them.

Table 1 illustrates the minimum amounts of single-
20 and double-stranded oligos of variable sequences/lengths detectable using the 2 different dyes. Similar results were obtained using phosphorothioate and 3'-amino-modified oligos. Single-stranded oligos were analyzed on denaturing 12%
25 polyacrylamide gels (Sambrook, J., E. F. Fritsch and T. Maniatis. (1989) Molecular Cloning: A Laboratory Manual. Cold Spring Harbor Laboratory Press. Plainview, NY.) (5 g urea, 4.69 ml 30% acrylamide solution, 1.88 ml 5x TBE, 56 ml 10% ammonium
30 persulfate, 3.5 ml TEMED, and 0.5 ml of 10 mg/ml ethidium bromide solution) and loaded with formamide-containing sample buffer. Double-stranded oligos were analyzed on non-denaturing 20% gels (Sambrook, J., E. F. Fritsch and T. Maniatis. (1989)
35 Molecular Cloning: A Laboratory Manual. Cold Spring Harbor Laboratory Press. Plainview, NY.) (6.66 ml 30% acrylamide solution, 2 ml 5x TBE, 1.27 ml water, 56 ml

10% ammonium persulfate, 3.5 ml TEMED, and 0.5 ml of 10 mg/ml ethidium bromide solution) and loaded with glycerol-containing sample buffer. Cassettes (1 mm), 10-well combs, and the Xcell Mini-cell system from
5 Novex (San Diego, CA) were used. Running buffer (1.25x TBE) included 0.5 mg/ml ethidium bromide. After 1.5 hours of electrophoresis at 150 V (~15 mAmp), the gel was removed from the cassette and photographed in the dark using a Polaroid camera and ultraviolet light
10 source. Immediately after photography, the gel was stained for approximately 5 minutes using 0.02% methylene blue. The gel was destained with warm water (2-5 washes) until background was clear, then photographed on top of a fluorescent light box.

15 We have used this sequential staining technique to distinguish (on a 19% denaturing acrylamide gel) products from reactants in a ligation reaction (Chu, B. C. and L. E. Orgel. (1991) Nucl. Acid. Res. 19(24): 6958) where the reactant is a 42 mer single-stranded,
20 intramolecularly annealing oligo and the product is a closed, double-stranded, dumbbell-shaped oligo of the same length. Although the molecular weights of the product and reactant are similar, they migrate at significantly distinct rates due to their structural
25 difference (Chu, B. C. and L. E. Orgel. (1991) Nucl. Acid. Res. 19(24): 6958). This technique has also been successfully used to analyze ligation reactions involving oligos with different sequences and lengths. Ethidium bromide helps to distinguish between our
30 reactants (single-stranded) and products (double-stranded) while methylene blue may allow quantification of percent yield for the reaction. We are thus able to obtain more information from gels stained with two consecutive agents rather than just
35 one.

The polynucleotide decoys used in the gels:

Lane 1. 42mer product:

5 $\begin{array}{c} \text{T}^{\text{T}}\text{T}^{\text{T}}\text{CCCCTAGCAACAGATG}^{\text{T}}\text{T}^{\text{T}} \\ \text{T}^{\text{T}}\text{GGGGATCGTTGTCTAC}^{\text{T}}\text{T}^{\text{T}} \end{array}$

2. 42mer ~~kan~~etant:

10 5'-CTAGGGGTTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3'

Table 1: Approximate amounts (mg) of oligonucleotide detectable with methylene blue and ethidium bromide.

		<u>Approximate Amounts (mg)</u>	
<u>Detectable:</u>			
	<u>Oligonucleotide</u>	<u>Methylene Blue</u>	<u>Ethidium Bromide</u>
5			
10	single-stranded 16mer 5' -CCCCTAGCAACAGATG-3'	0.05	>10
15	single-stranded 20mer 5' -GCCACGGAGCGAGACATCTC-3'	0.05	>1
20	single-stranded 40mer 5' -CTAGGGGTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3'	0.05	0.01
25	double-stranded 16 bp 5' -CCCCTAGCAACAGATG-3' 3' -GGGGATCGTTGTCTAC-5'	0.05	0.1
30	double-stranded 20 bp 5' -GCCACGGAGCGAGACATCTC-3' 3' -CGGTGCCTCGCTCTGTAGAG-5'	0.05	0.05
35	40mer decoy (double-stranded 16 bp)	0.005-0.01	0.005-0.01
40			
45	T ^T CCCCTAGCAACAGATG ^T T ^T GGGGATCGTTGTCTAC ^T		
50			

Example 8: Gel Shift Assays - Methods and Results

There were 2 types of gel shift assays performed: the direct binding assay and the competition assay. The direct binding assay was able to test whether or not a radiolabeled polynucleotide decoy was able to directly bind to RF-X transcription factors. The competition assay was able to test whether a decoy was able to compete with a radiolabeled control sequence, usually the double-stranded extended X-box, also referred to as "AX", for binding to RF-X. RF-X transcription factors were obtained from a nuclear extracts and incubated with the polynucleotide decoy of interest. In the case of the competition assay the competing oligo was added to see if it could displace the other oligo from RF-X. The mixtures were then run on a gel, and radioactive bands were detectable by autoradiography.

Detailed Methods:20 *Nuclear Extracts*

Nuclear extracts, containing RF-X transcription factors, were obtained from Raji cells using the method of Ohlsson et al. (1986), Cell, Vol. 45(1), pp. 35-44.

25 *Direct Binding Assay Conditions*

The following mixture was made and incubated for 20 minutes on ice:

- a. binding buffer (12 mM HEPES-KOH, 12% glycerol, 60 mM KCl, 5 mM MgCl₂, 0.12 mM EDTA, 0.3 mM DTT)
- b. polydIdC (non-specific DNA competitor, 2 μ g)
- c. RF-X transcription factor from nuclear extract: ~3 μ g
- d. test compound (DNA decoy, X-box probe = AX, or other nucleic acid of interest)

Competition Assay Conditions (similar to the assay performed by Hasegawa et al. (1991), *Nucleic Acids Res.*, Vol. 19(6), pp. 1243-9)

1. The following mixture was made and incubated
5 for 10 minutes on ice:
 - a. binding buffer (12 mM HEPES-KOH, 12%
glycerol, 60 mM KCl, 5 mM MgCl₂, 0.12 mM
EDTA, 0.3 mM DTT)
 - 10 b. polydIdC (non-specific DNA competitor, 2
μg)
 - c. RF-X transcription factors from nuclear
extract: ~3 μg
 - 15 d. competitor (polynucleotide decoy; or
controls of X-box probe, non-specific
double-stranded 10-mer, or nothing)
2. After the 10 minute incubation, ³²P
end-labeled, double-stranded DRA X-box probe
(~50-mer) (the sequence of which is described
by Hasegawa et al. (1991), *Nucleic Acids*
20 *Res.*, Vol. 19(6), pp. 1243-9) was added and
incubated for 30 minutes.

Gel: Run reaction mixtures on 5% native polyacrylamide
gel; re-circulate 0.25 x TBE buffer. Run time:
25 ~2 hours (150-175 V, 200 Watts, 10 mAmp)

Autoradiograph: expose X-ray film O/N at -70°C.

Detailed Results:

30 When the Direct Binding Assay was performed, Lane
1 demonstrated RF-X transcription factor binding to the
hot AX probe. Lane 2 demonstrated that the cold AX
probe was able to compete off the hot AX probe (cold AX
in a 30-fold excess). These 2 lanes served as controls
35 for comparison with all lanes with test compounds. The
16 bp X-box, M6 decoy, unligated M6 decoy, RF-X-2
decoy, unligated RF-X-2 decoy, unligated RF-X decoy T5,

and unligated κ B mutant (lanes 3, 6, 7, 8, 9, and 12, respectively) demonstrated no measurable binding to the RF-X transcription factors. The RF-X decoy (lane 4) bound to RF-X, as did the unligated RF-X decoy (lane 5) and the RF-X decoy T5 (lane 10), but the latter 2 to a lesser degree.

When the Competition Assay was performed, Lane 1 exhibited RF-X transcription factor binding to the hot AX probe. For competition to occur, the competing oligo should "displace" the hot AX probe from the RF-X transcription factors. The oligos containing the core X-box sequence could compete with hot AX for binding to RF-X; cold AX (lane 2), unligated RF-X decoy (lanes 3, 4, 9), RF-X decoy (lanes 5, 6), RF-X decoy T5 (lane 7), and 16 bp X-box (lane 9). Lane 11, the κ B mutant, did not compete, nor did a non-specific 20-mer oligo (lane 12).

Example 9: CAT Assays - Methods and Results

When Raji cells were transfected with pDRASCAT plasmid, basal CAT activity could be measured. When Raji cells are co-transfected with pDRASCAT and a decoy, we were able to measure the specific effect of the various decoys on the basal CAT activity. Decoys that diminished CAT activity were considered "active." "Inactive" oligos demonstrated little or no effect on CAT activity.

Detailed Methods

Cell culture. MHC Class II expressing Raji cells (ATCC COWL 86, a human Burkitt's lymphoma B-cell line) were grown in RPMI 1640 medium with fetal bovine serum (20%), penicillin (100 U/ml), streptomycin (100 μ g/ml), and L-glutamine (292 μ g/ml).

Plasmids. The plasmid pDRASCAT has been described previously (Tsang et al. (1990), Mol. Cell. Biol., Vol.

10(2), pp. 711-719; and Voliva et al. (1992), Mol. Cell. Biol., Vol. 12(5), pp. 2383-90). Briefly, it contains the DRA promoter from positions -150 to +31 linked to the CAT gene.

5

Transfections. 1×10^6 to 4×10^6 Raji cells (in 1/2 ml) were electroporated with 20 to 40 μ g pDRASCAT plasmid, along with 10 μ M of a test decoy (or no nucleic acid) using the Bio-Rad Gene Pulsar[®] set at 300
10 V and 960 μ F. After 10 minutes, the cells were transferred to 4.5 ml complete media.

CAT assays. Cells were harvested for CAT assays approximately 48 hours after electroporation. The
15 polynucleotide decoys and other oligonucleotides were synthesized as described in Example 6. CAT assays were performed using the method of Neumann et al. (1987), Biotechniques, Vol. 5(5), pp. 444-447, which involves measurement of CAT by scintillation counting. Counts
20 were directly proportional to CAT activity.

Detailed Results

For all figures, basal CAT activity (pDRASCAT plasmid transfected alone) shown by white bar, CAT activity in
25 the presence of oligos shown by shaded bars. The shaded bars should be compared to the white bar in order to quantify the decreased CAT activity as a result of the addition of a decoy.

30 *Figure 6A (plasmid, ds DNA, unlig RF-X decoy, RF-X-2 decoy, RF-X decoy):*

Doubled stranded X-box DNA demonstrated no effect on CAT activity. While unligated RF-X decoy, RF-X-2 decoy, and RF-X decoy all decrease CAT
35 activity significantly.

Figure 6B (plasmid, unlig RF-X decoy, cells only):

Unligated RF-X decoy, again decreased CAT activity. When no plasmid is transfected (cells alone), Raji cells displayed minimal basal CAT activity, as expected. This "cells-alone" data point was generated as an internal control.

Figure 6C (plasmid, unlig M6 mutant, unlig κB mutant, unlig RF-X decoy T4, unlig RF-X decoy T5):

Only the unligated RF-X decoy T4 (4 T's in the loop) decreased CAT activity in this experiment. All other decoys were inactive.

Summary of Data for CAT Assays of Decoys

All oligos co-transfected (electroporated) with pDRASCAT plasmid unless otherwise indicated

Active Decoy: Down-regulated pDRASCAT

RF-X decoy
unligated RF-X decoy
RF-X-2 decoy
unligated RF-X-2 decoy
unligated κB decoy

Inactive Decoys: No Significant Down-regulation of pDRASCAT

double-stranded (preannealed) X-box, 16 bp
unligated 5T loop RF-X decoy
unligated M6 mutant decoy
unligated κB mutant decoy
passively administered unligated RF-X decoy

Claims

1. A polynucleotide decoy comprising:
 - (a) an internal oligonucleotide (I) having a length of X bases, where X is a number from about 10 to about 40;
 - (b) two cap oligonucleotides (P_1 and P_2), each having a length of from about 3 to about 8 bases, wherein each of said cap oligonucleotides is comprised of bases which are unable to bind to any other base within the same cap oligonucleotide;
 - (c) a first complementary oligonucleotide (C_1) having a length of Q bases, where Q is a number from about 5 to $(X - 5)$, said C_1 having a 3' to 5' nucleic acid sequence capable of Watson-Crick-type binding to the first Q bases in the 5' to 3' nucleic acid sequence of said I; and
 - (d) a second complementary oligonucleotide (C_2) having a length of Z bases, where Z is a number greater than equal to about 5 and from $((X - (Q + X/8))$ to $(X - Q)$, said C_2 having a 5' to 3' nucleic acid sequence capable of Watson-Crick-type binding to the first Z bases in the 3' to 5' nucleic acid sequence of said I;wherein the 3' end of said C_1 is covalently linked to the 5' end of said P_1 , the 3' end of said P_1 is covalently linked to the 5' end of said I, the 3' end of said I is covalently linked to the 5' end of said P_2 , and the 3' end of said P_2 is covalently linked to the 5' end of said C_2 ;
 - wherein at least one of the oligonucleotides selected from the group consisting of said C_1 , said C_2 , and said I, comprises an RF-X recognition sequence; and
 - wherein said polynucleotide decoy is capable of binding to an RF-X transcription factor.
2. The polynucleotide decoy of claim 1, wherein said RF-X recognition sequence is Type-1.

3. The polynucleotide decoy of claim 1, wherein said RF-X recognition sequence is Type-2.

4. The polynucleotide decoy of claim 1, wherein $Z = X - Q$.

5. The polynucleotide decoy of claim 4, wherein the 3' end of C_2 is covalently linked to the 5' end of C_1 .

10

6. The polynucleotide decoy of claim 1, wherein P_1 and P_2 each consists of a single variety of nucleotide comprising a base selected from the group consisting of adenine, cytosine, thymidine, and modified nucleotides thereof.

15

7. The polynucleotide decoy of claim 6, wherein P_1 and P_2 consist of a single variety of nucleotide comprising a thymidine base.

20

8. The polynucleotide decoy of claim 7, wherein the cap oligonucleotides have a length of 4 nucleotides.

9. A polynucleotide decoy consisting essentially of an oligonucleotide selected from the group consisting of:

5' -CTAGGGGTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3',
 5' -GACTGGGTTTTCCAGTCCATACGAAGTTTTCTTCGTATG-3',
 5' -GCTAGGGGTTTTCCCCTAGCAACAGATGTTTTTCATCTGTT-3',
 5' -CTAGGGGTTTTCCCCTAGCAACAGATGTTTTTCATCTGTTG-3',
 5' -TAACAACTTTTGTTGTTATAGTAACTTTTGTACTA-3',
 5' -TCCCTTGGTTTTCCAAGGACTTTCCGCTTTTGCGGAAAG-3',

and

5' -TGCTAGGTTTTTCCTAGCAACAGATGTTTTTCATCTGT-3'.

35

10. A polynucleotide decoy according to claim 9, wherein the 3' end and the 5' end of said oligonucleotide are covalently linked.

5 11. A method for making a polynucleotide decoy, comprising covalently linking nucleotides such that the resulting product has the structure of a polynucleotide decoy of claim 1, 5, 9, or 10.

10 12. A method for making an MHC-II-depleted cell from a target cell, the method comprising:
(a) obtaining the target cell; and
(b) exposing the target cell to a polynucleotide
15 decoy of claim 1, 5, 9, or 10, said polynucleotide decoy being preset in an amount sufficient to make the target cell a MHC-II-depleted cell.

13. The method of claim 12 wherein the target cell is selected from the group consisting of corneal
20 endothelial cells, thyroid cells, parathyroid cells, brain cells, adrenal gland cells, bone marrow cells, pancreatic islet cells and hepatic cells.

14. A MHC-II-depleted cell prepared by the method
25 of claim 12.

15. A polynucleotide decoy of claim 1, 5, 9, or
10 for use in the preparation of a composition for
treating target cells to make them MHC-II-depleted
30 cells.

16. A MHC-II-depleted donor organ prepared by a method comprising the steps of:

(a) obtaining a target organ from an individual;
and

5 (b) exposing the target organ to a polynucleotide decoy of claim 1, 5, 9, or 10, said polynucleotide decoy being preset in an amount sufficient to make the target organ a MHC-II-depleted donor organ.

10 17. A method of treating an individual with an autoimmune disease characterized by dysfunctional expression of a MHC class II antigen, the method comprising administering to that individual a
15 amount sufficient to inhibit expression of the MHC class II antigen.

18. A method of treating an individual infected with hepatitis B virus, the method comprising
20 administering to that individual a polynucleotide decoy of claim 1, 5, 9, or 10 in an amount sufficient to inhibit expression of hepatitis B surface antigen.

19. A method of treating an individual infected
25 with cytomegalovirus, the method comprising administering to that individual a polynucleotide decoy of claim 1, 5, 9, or 10 in an amount sufficient to inhibit RF-X binding to the enhanced factor C sites of the cytomegalovirus genome.

30

35

20. A method of using a polynucleotide decoy to obtain a substantially purified, covalently closed polynucleotide decoy, comprising the steps of:

- 5 of claim 1 or 9;
- (a) providing a linear polynucleotide decoy
- (b) treating said linear polynucleotide decoy with a kinase enzyme, thereby obtaining 5' phosphorylated polynucleotide decoy;
- 10 (c) heating said 5' phosphorylated polynucleotide decoy to inactivate said kinase enzyme, thereby obtaining heated, 5' phosphorylated polynucleotide decoy;
- (d) cooling said heated, 5' phosphorylated polynucleotide decoy mixture slowly to facilitate
- 15 intramolecular annealing, thereby obtaining cooled, 5' phosphorylated polynucleotide decoy;
- (e) treating said cooled, 5' phosphorylated polynucleotide decoy with a ligating enzyme, thereby obtaining a covalently closed polynucleotide decoy; and
- 20 (f) purifying said covalently closed polynucleotide decoy, thereby obtaining a substantially purified, covalently closed polynucleotide decoy.

21. A nucleic acid comprising a linear

25 polynucleotide decoy of claim 1 or 9.

22. A host cell comprising a nucleic acid of claim 21.

FIG. 1

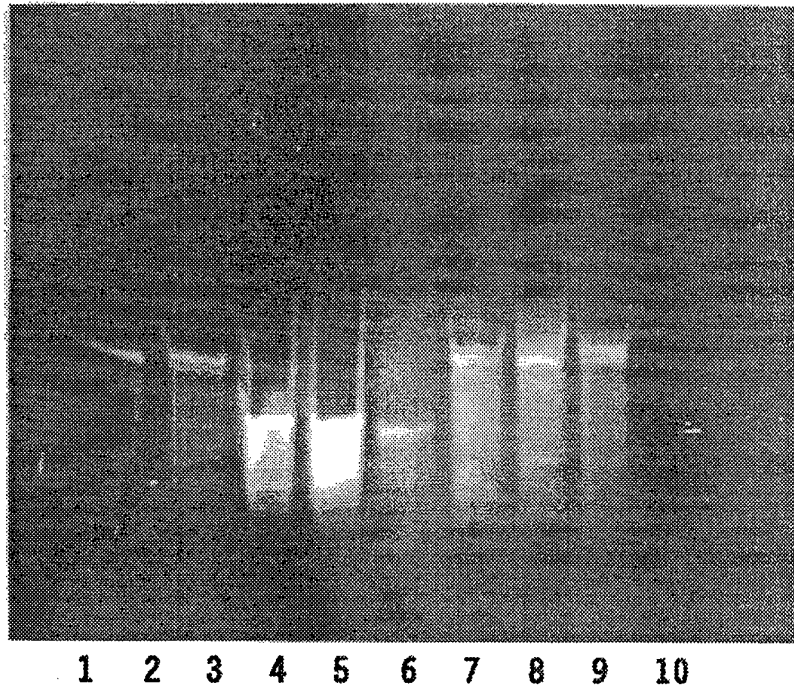


FIG. 2

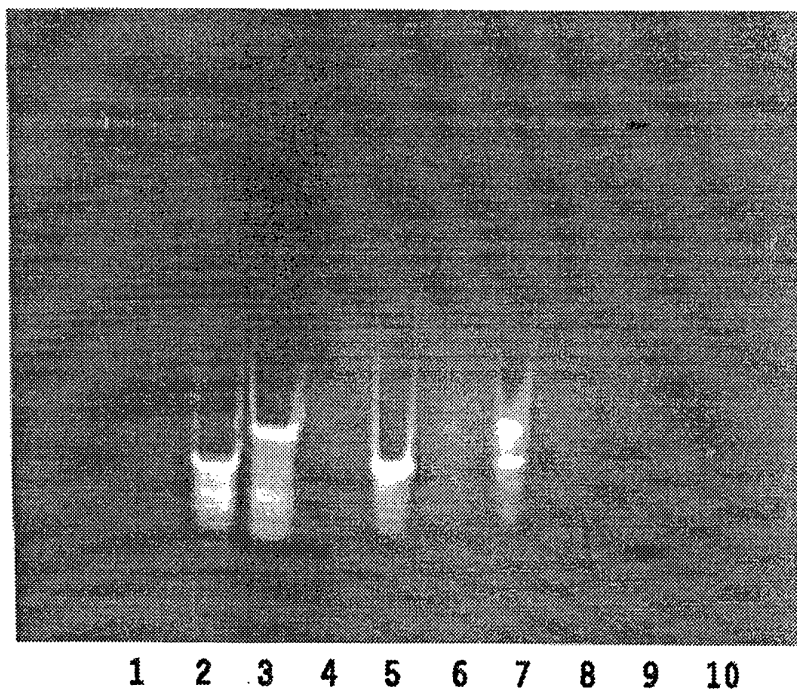


FIG. 3

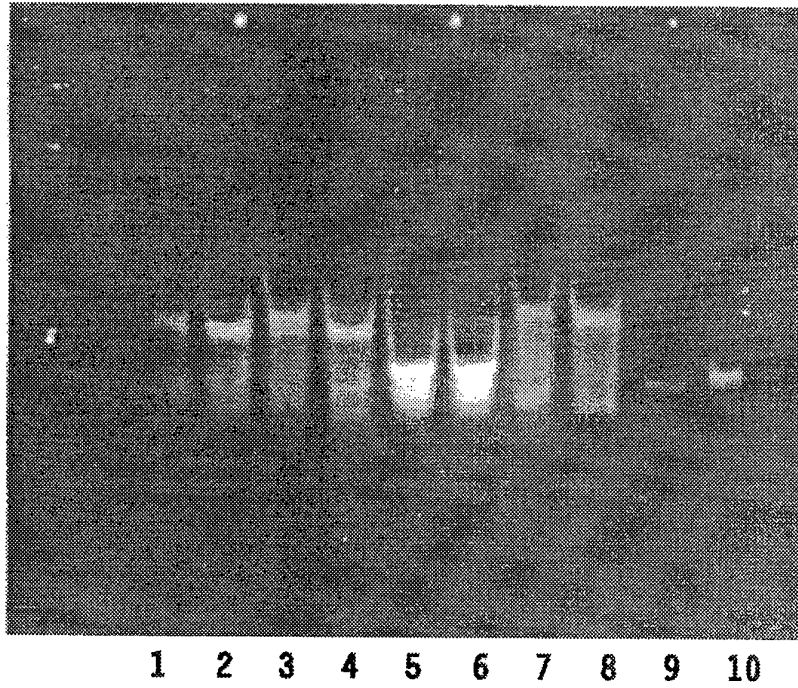
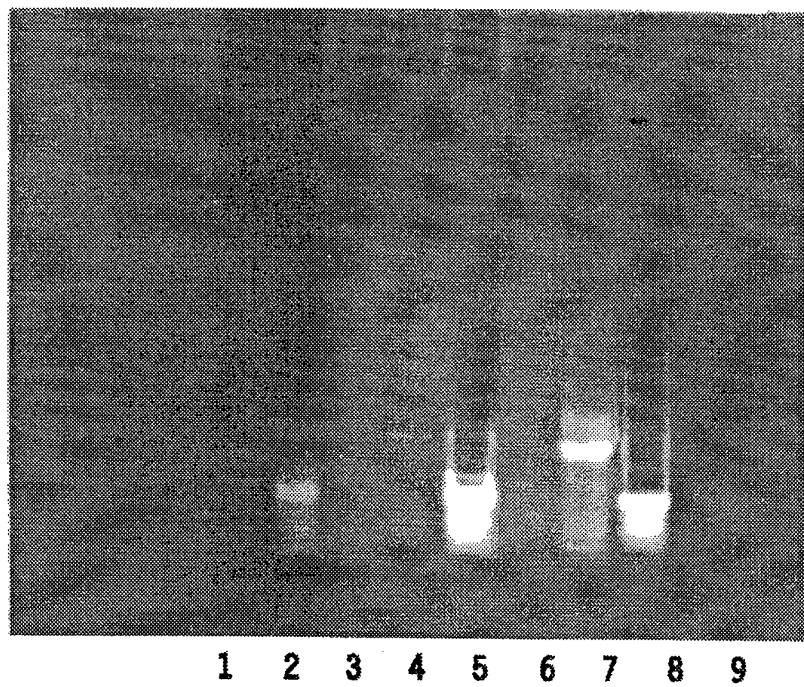


FIG. 4

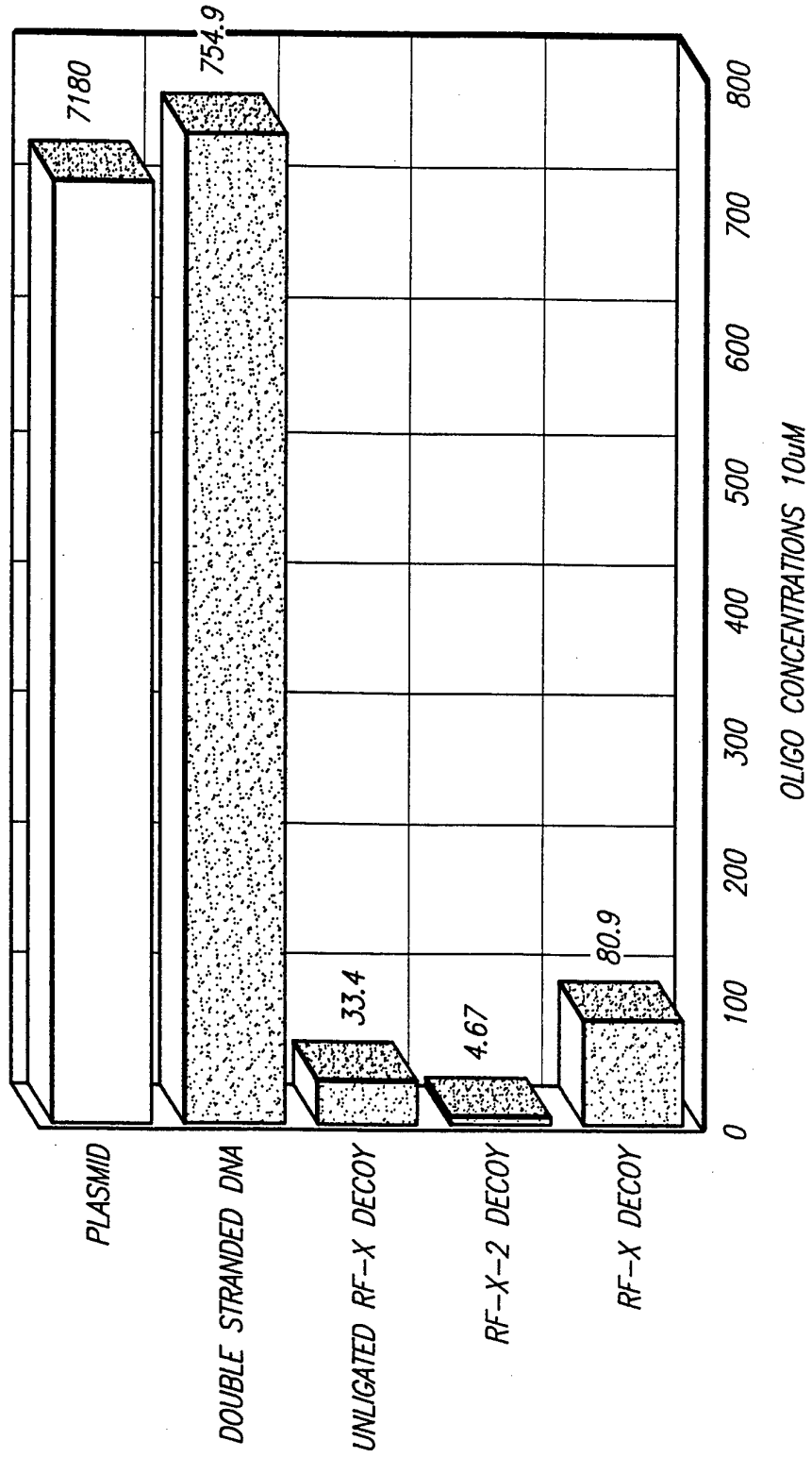


1 2 3 4 5 6 7



FIG. 5

FIG. 6A



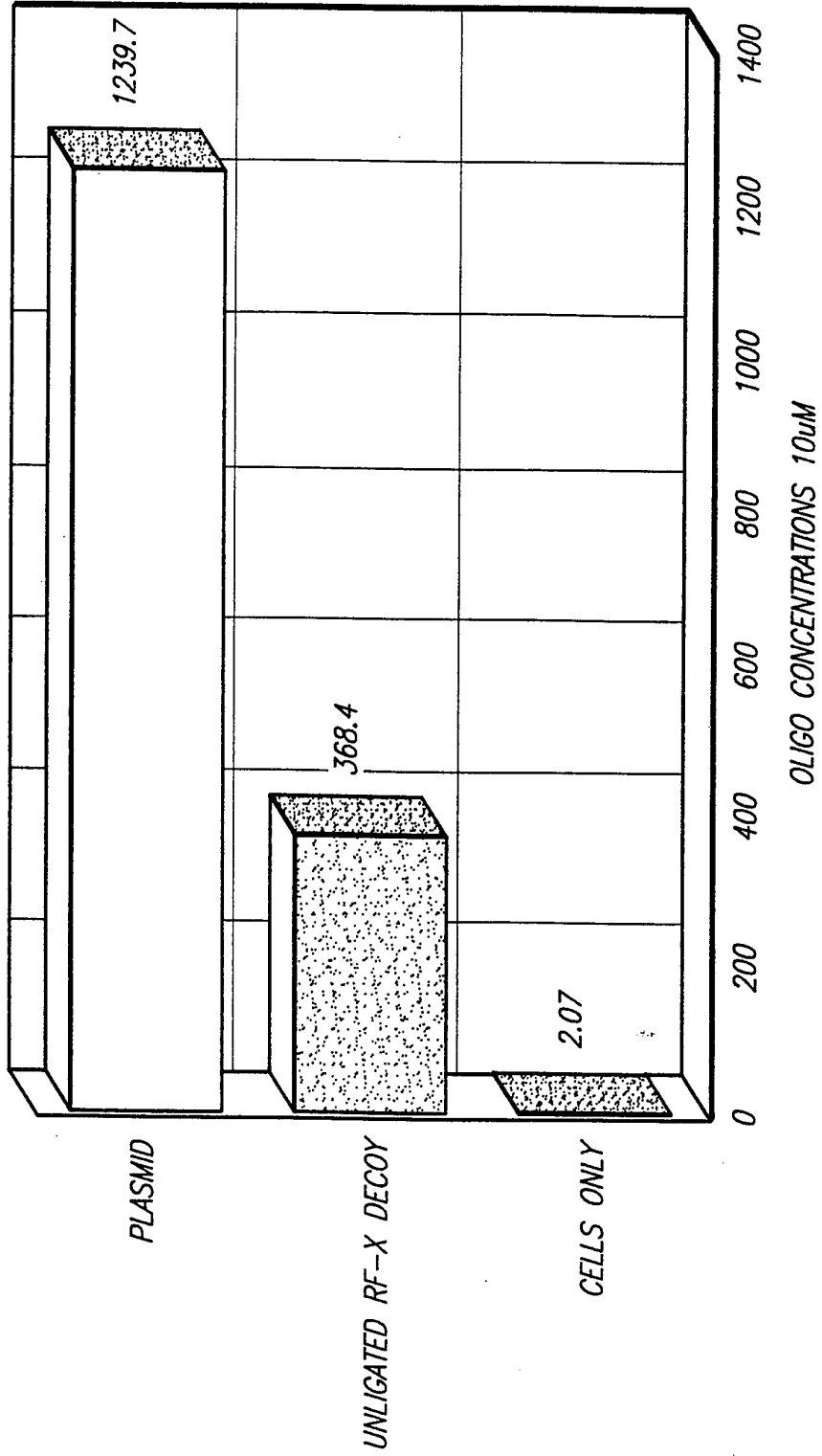


FIG. 6B

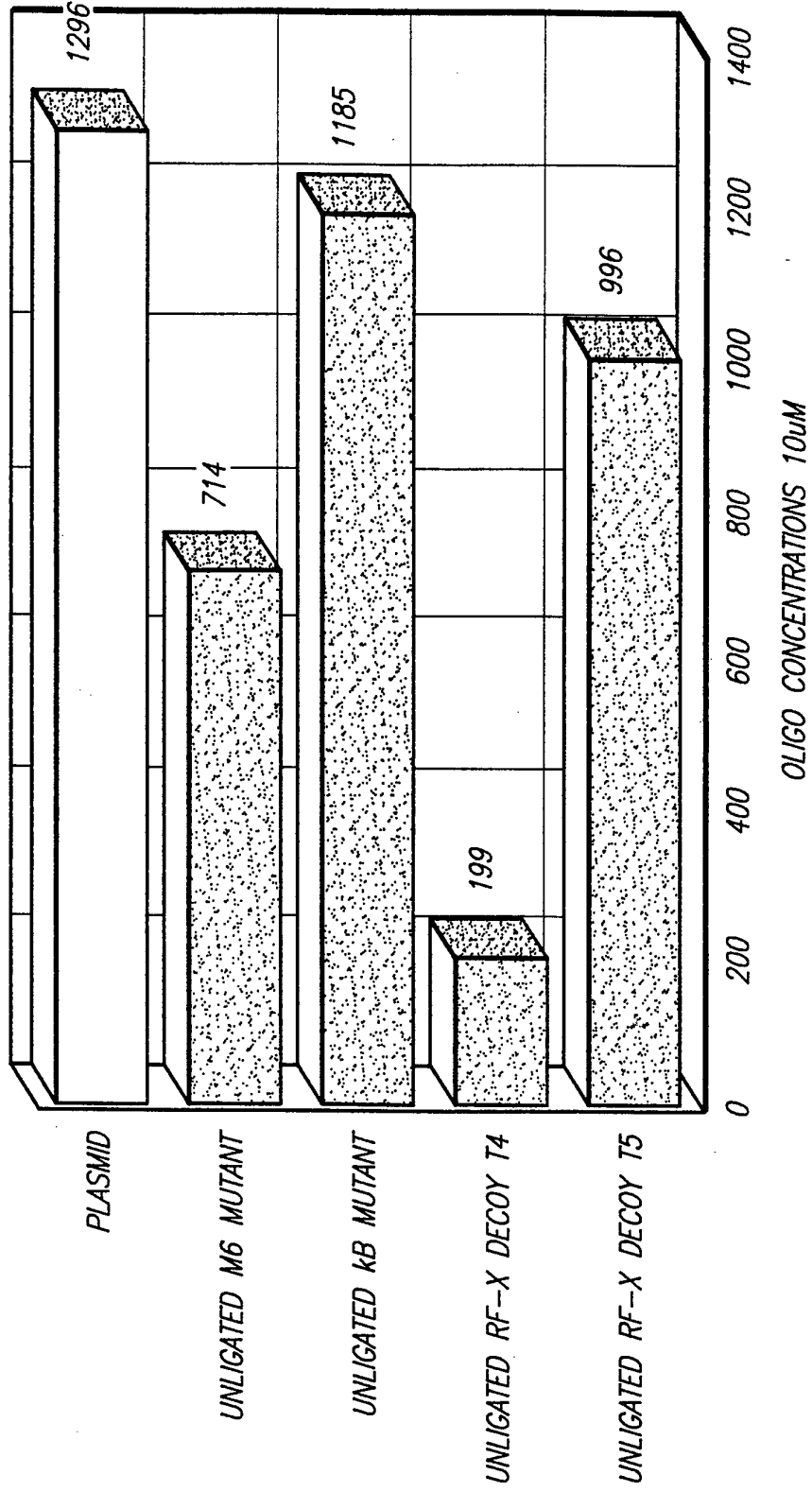


FIG. 6C

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/08667

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C07H 17/00; C12N 5/00; C12Q 1/68

US CL :435/6, 240.2; 514/44; 536/24.1

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/6, 240.2; 514/44; 536/24.1

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

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

APS, DIALOG

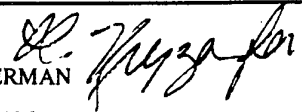
C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Nucleic Acids Research, Volume 19, Number 24, issued 25 December 1991, Chu et al, "Binding of Hairpin and Dumbbell DNA to Transcription Factors", page 6958, see entire document.	1-22
Y	Proceedings of the National Academy of Sciences USA, Volume 88, issued May 1991, Ono et al. "Transcription of a Subset of Human Class II Major Histocompatibility Complex Genes is Regulated by a Nucleoprotein Complex that Contains c-fos or an Antigenically Related Protein", pages 4304-4308, see entire document.	1-22

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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Date of the actual completion of the international search 31 OCTOBER 1994	Date of mailing of the international search report NOV 07 1994
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Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized officer DOUGLAS GURIAN-SHERMAN  Telephone No. (703) 308-0196
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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/08667

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Science, Volume 252, issued 21 June 1991, Faustman et al, "Prevention of Xenograft Rejection by Masking Donor HLA Class I Antigens", pages 1700-1702, see entire document.	1-17, 20-22
Y,P	Molecular and Cellular Biology, Volume 13, Number 10, issued October 1993, Siegrist et al., "RFX1 is Identical to Enhancer Factor C and Functions as a Transactivator of the Hepatitis B Virus Enhancer", pages 6375-6384, see entire document.	17-19