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[Continued on next page]

(54) Title: CD154 VARIANTS

TNFH(-) CD154 variant is absent from cell surface

Transfection

Wild-type
TNFH(-)

p33
p31

p33
p31

p33
p31

TNFH(-)-----

i.p. with:

CD40-Fc

Rb 779

Rb 784

9E10

1 2 3 4 5 6

(57) Abstract: Methods of decreasing (e.g., inhibiting) the expression of wildtype CD154 on the surface of a target cell and methods of treating a patient suffering from or predisposed to a CD154-mediated disease. In these methods, a nucleic acid construct that directs expression of a mutant CD154 lacking at least a portion of the tumor necrosis factor homologous domain ("TNFH") is introduced into a target cell (such as a T helper cell or a cytotoxic T cell). The expressed mutant CD154 binds to wildtype CD154 inside the cell, rendering the wildtype protein unable to reach the cell surface.

WO 01/96397 A2



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CD154 VARIANTS

TECHNICAL FIELD OF THE INVENTION

This invention relates to methods of using
5 CD154 variants lacking at least a portion of a tumor
necrosis factor homologous domain to inhibit the
expression of wildtype CD154 on the surface of target
cells. The methods of the invention are useful in
treating or inhibiting CD154-dependent immune
10 disorders.

BACKGROUND OF THE INVENTION

CD154 (i.e., CD40 ligand or CD40L) is a
type II membrane protein expressed primarily on
activated T cells. The interaction of CD154 with its
15 receptor, CD40, is critical for the functions of T
helper cells to induce differentiation, proliferation,
and immunoglobulin isotype switching in B cells (for
review see Foy et al., *Annu Rev Immunol* 14:591-617
(1996); van Kooten et al., *J Leukoc Biol* 67:2-17
20 (2000)). The CD154 gene, located at chromosomal region
Xq2.6-2.7, spans more than 12 kilobase pairs and
contains five exons (Villa et al., *Proc Natl Acad Sci
USA* 91:2110-4 (1994)).

The first exon encodes the cytoplasmic
25 region, the transmembrane domain, and six amino acids
of the extracellular domain. The second and third

- 2 -

exons encode the extracellular stalk region. The fourth and fifth exons encode the C-terminal 147 amino acids (Villa et al., *supra*), a region that shares limited homology with other members of the tumor
5 necrosis factor ("TNF") family and is therefore called the TNF homologous ("TNFH") domain. The X-ray structure of the CD154 TNFH domain reveals that it contains a sandwich-like fold of two β sheets with jellyroll or Greek key topology. Although members of
10 the TNF family share the configuration of the Type II membrane protein, the limited homology between these members is located in the C-terminal TNFH domain of approximately 150 amino acid residues.

Like TNF and lymphotoxin- α , wildtype CD154
15 exists as trimers (Karpusas et al., *Structure* 3:1031-9 (1995)). The formation of CD154 trimers is mediated by the TNFH domain. It has been shown that the TNFH domain alone is capable of forming trimers (PCT patent application WO 97/00895; Karpusas et al., *supra*; Mazzei
20 et al., *J Biol Chem* 270:7025-8 (1995)). Deletion mutants missing a major portion of this domain do not seem to exist as trimers (Garber et al., *J Biol Chem* 274:33545-50 (1999)). Thus, it appears that the TNFH domain is necessary and sufficient for the assembly of
25 trimeric CD154 proteins.

Mutations of the CD154 gene preventing the expression of functional CD154 protein can lead to an immunodeficiency characterized by elevated IgM levels and low IgG and IgA levels in serum. Since the CD154
30 gene is located on the X chromosome in humans, this immunodeficiency is called X-linked hyper-IgM syndrome ("XHIM") (Allen et al., *Science* 259:990-3 (1993); Korthauer et al., *Nature* 361:539-41 (1993); DiSanto et

- 3 -

al., *Nature* 361:541-3 (1993); Aruffo et al., *Cell* 72:291-300 (1993); Fuleihan et al., *Proc Natl Acad Sci USA* 90:2170-3 (1993)). Over 70 unique mutations in the CD154 gene have been identified in more than one
5 hundred XHIM patients (Notarangelo et al., *Immunol Today* 17:511-6 (1996)). These mutations are very heterogeneous. They include insertions, deletions, and point mutations. Thus, it is conceivable that the underlying mechanisms for the functional defects of
10 CD154 in XHIM patients are different (Garber et al., *supra*; Seyama et al., *Blood* 92:2421-34 (1998)).

Because the CD154 gene is X-linked, each cell from normal individuals or XHIM patients makes a single species of CD154-encoding transcript. However, in some
15 XHIM patients, mutations in the donor splicing sites (Seyama et al., *supra*; and references cited therein) or the acceptor splicing sites (Ameratunga et al., *Clin Diagn Lab Immunol* 3:722-6 (1996)) lead to generation of multiple species of mRNA transcripts in a single cell.
20 These transcripts include the normally spliced transcripts encoding wildtype CD154, as well as the misspliced transcripts encoding variant CD154 proteins lacking either a major portion of the TNFH domain, or the entire TNFH domain (Seyama et al., *supra*;
25 Ameratunga et al. *supra*).

Because the TNFH domain alone appears to be responsible for the assembly of trimeric CD154 protein, variants lacking the TNFH domain were predicted not to affect the trimerization of the wildtype protein.
30 Thus, it is unclear why patients with mutations at the splicing sites of their CD154 genes exhibit the XHIM syndrome, which presumably results from a lack of functional CD154 protein.

- 4 -

SUMMARY OF THE INVENTION

The present invention is based on our discovery that CD154 variants lacking at least a portion of the TNFH domain can nonetheless interact with a wildtype CD154 protein via the stalk region. This interaction retains the wildtype CD154 protein intracellularly, thereby preventing the wildtype protein from being expressed on the cell surface and participating in any CD154 functional activity. This discovery reveals the stalk region of the CD154 protein as a previously unrecognized structural element that contributes to CD154 trimer assembly.

Accordingly, this invention provides a method of decreasing (including inhibiting) the expression of wildtype CD154 protein on the surface of target cells. In this method, a recombinant nucleic acid construct is introduced into a CD154-expressing target cell, where the construct is capable of directing the expression of a CD154 variant protein lacking a functional TNFH domain of the wild type protein, such that the variant protein is incapable of trimerization and yet can bind to the wildtype protein, rendering the wildtype protein unable to reach the cell surface. For instance, the variant lacks at least a portion of the TNFH domain. By way of example, a CD154 variant useful in the invention may lack at least 5 (e.g., at least 10; at least 15; or at least 20) amino acids of the TNFH domain. In one embodiment, a CD154 variant may lack the entire TNFH domain. In another embodiment, a CD154 variant may comprise a TNFH domain with insertional mutations and/or point mutations.

As used herein, the TNFH domain of CD154 corresponds to a region spanning from amino acid

- 5 -

residues 116 to 261 of SEQ ID NO:1:

```
1      MIETYNQTSP RSAATGLPIS MKIFMYLLTV FLITQMIGSA LFAVYLHRRL
51     DKIEDERNLH EDFVFMKTIQ RCNTGERSLS LLNCEEIKSQ FEGFVKDIML
101    NKEETKKENS FEMQKGDQNP QIAAHVISEA SSKTTSVLQW AEKGYTMSN
5  151    NLVTLENGKQ LTVKRQGLYY IYAQVTFCNS REASSQAPFI ASLCLKSPGR
201    FERILLRAAN THSSAKPCGQ QSIHLGGVFE LQPGASVFN VTDPSQVSHG
251    TGFTSFGLLK L      (SEQ ID NO:1)
```

SEQ ID NO:1 represents a full-length sequence of human CD154, and is available in GenBank under
10 accession number CAA48554 (referencing Graf et al.,
Eur. J. Immunol. 22(12):3191-4 (1992)). Allelic isoforms of SEQ ID NO:1 can also be used as the cognate sequence for the CD154 variants useful in this invention.

15 Examples of CD154 variants useful in this invention include those that lack (1) amino acid residues 116-136, (2) amino acid residues 137-261, (3) amino acid residues 115-261, and (4) amino acid residues 97-261, respectively, of SEQ ID NO:1. Methods
20 of generating CD154 variants are known in the art. See, e.g., Fig. 1 and Table 1 of Garber et al., *J. Biol. Chem.* 274:33545-550 (1999).

This invention further provides a method of treating or inhibiting a CD154-dependent disease (e.g.,
25 an immune disorder mediated by CD154:CD40 interaction) in a subject (i.e., a mammal such as a primate, preferably a human). The method involves administering to target cells of the subject, such as T cells (e.g., CD4⁺ and CD8⁺ T cells) or megakaryocytes, a CD154
30 variant of the present invention, thereby decreasing the expression of wildtype CD154 on the surface of the target cells. In one embodiment, the CD154 variant is

- 6 -

generated inside the target cell, based on uptake by the cell of a gene expression construct comprising a nucleic acid sequence encoding that variant.

The methodology of the present invention can also be used to generate variants of other TNF ligand family proteins (e.g., the FAS ligand), where these proteins' TNFH domains are rendered incapable of trimerization by mutations (e.g., deletions, insertions and/or point mutations), such that the resultant variants bind to and retain the corresponding wildtype protein intracellularly, interfering with the cell surface expression of the wildtype protein. For a general review of the TNF receptor family, see Fahrner et al., *Nature* 409:836-8 (2001) and Locksley et al., *Cell* 104:487-501 (2001).

Within the scope of the present invention are also pharmaceutical compositions comprising a nucleic acid construct that directs expression of a CD154 mutant useful in a method of the invention. The invention also provides the use of such a construct for the manufacture of a medicament for decreasing the expression of wildtype CD154 on the surface of a target cell, and/or for treating a patient suffering from or predisposed to a CD154-mediated disease.

The present invention is well-suited for local delivery and uptake by target cells at a locus in need of therapeutic treatment in the body of the subject. For example, the gene expression construct described herein can be administered locally (e.g., by injection) to spleen or other lymphoid tissues, or into a locus of inflammation, such as a wound site or pathological lesion. Such local treatment circumvents any complications that may be encountered with systemic delivery of immunomodulating agents.

- 7 -

In addition, the present gene expression construct avoids potential drawbacks of extracellular immunomodulating agents, such as antibodies.

Antibodies, when bound to a cell-surface antigen, may
5 trigger cell destruction, for instance, via an antibody-dependent cell cytotoxicity response, or by macrophages or other effector cells that display Fc receptors on their cell surface. Such cell destruction may not be desirable in certain therapeutic settings.

10 Other features and advantages of the present invention will be apparent from the following drawings and detailed description, and also from the appended claims.

Unless otherwise defined, all technical and
15 scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention pertains. Exemplary methods and materials are described below, although methods and materials similar or equivalent to those described
20 herein can also be used in the practice of the present invention. All publications and other references mentioned herein are incorporated by reference in their entirety. In case of conflict, the present specification, including definitions, will control.
25 The materials, methods, and examples are illustrative only and not intended to be limiting.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is an autoradiograph demonstrating the association of TNFH(-) CD154 with wildtype CD154.

30 Fig. 2 is a panel of autoradiographs demonstrating the dose-dependent inhibitory effect of TNFH(-) CD154 on the production of functional CD154 trimers.

- 8 -

Figs. 3A and 3B are a bar graph and a table, respectively, showing that a CD154 mutant reduces the lymphotoxin- α up-regulating activity of wildtype CD154.

Fig. 4 is a panel of autoradiographs
5 demonstrating that TNFH(-) CD154 prevents the cell surface expression of wildtype CD154.

Fig. 5 is a schematic diagram illustrating that TNFH(-) CD154 binds to wildtype CD154 intracellularly and prevents it from reaching the cell
10 surface.

DETAILED DESCRIPTION OF THE INVENTION

It has been established in the past decades that T cell activation requires both T cell antigen receptor ("TCR") mediated signals and simultaneously
15 delivered costimulatory signals. For example, antibody production by B cells in response to protein antigens requires an antigen-specific interaction between B cells and helper T cells as well as non-antigen-specific, costimulatory receptor-ligand interactions
20 between the B and T cells. These non-antigen-specific interactions include the binding of CD40 on B cells to CD154 on T cells.

Human CD40 is a 50 kD cell surface protein expressed on mature B cells, macrophages and activated
25 endothelial cells. CD40 belongs to a class of receptors involved in proliferation and apoptosis, including Fas/CD95, TNF receptors and lymphotoxin receptors. Human CD154 is a 32 kD type II membrane glycoprotein transiently expressed primarily on
30 activated T cells. CD40:CD154 interaction is required for essentially all T cell-dependent immune responses, including antibody responses. In particular,

- 9 -

CD40:CD154 interaction provides anti-apoptotic and/or lymphokine stimulatory signals.

The present invention rests on the surprising discovery that CD154 mutational variants lacking at least a portion of the TNFH domain can bind to wildtype CD154 intracellularly and prevent the wildtype protein from reaching the cell surface. Thus, expression of such a CD154 variant in the T cells of a subject can specifically suppress CD154-dependent immune responses by eliminating the cell surface expression of wildtype CD154.

Treatment of Diseases

CD40:CD154 interaction is known to be pivotal in the induction of T cell dependent immune responses, including antibody-mediated humoral immune responses and T-cell mediated inflammatory responses to protein antigens. Although considerable efforts have been invested by several groups of investigators in developing means for intervening in this interaction to avert or to treat unwanted or pathological immune responses, there remains a need for improved means of interrupting CD40:CD154 interaction in therapeutic settings.

The present invention provides the improved means. According to this invention, a CD154 variant lacking a functional TNFH domain (e.g., a TNFH-minus CD154 mutant) can be introduced into target T cells in a subject to treat (e.g., mitigate, delay or reverse) or inhibit (e.g., prevent the onset or progression of) CD154-dependent diseases that may be characterized by significant inflammatory system or immune system involvement. Such diseases include, but are not limited to, lupus, systemic lupus erythematosus, lupus nephritis, lupus neuritis, asthma, chronic obstructive

- 10 -

pulmonary disease, bronchitis, emphysema, multiple sclerosis, uveitis, Alzheimer's disease, traumatic brain injury, traumatic spinal cord injury, stroke, atherosclerosis, coronary restenosis, ischemic
5 congestive heart failure, cirrhosis, hepatitis C, diabetic nephropathy, glomerulonephritis, autoimmune disease, osteoarthritis, rheumatoid arthritis, psoriasis, atopic dermatitis, systemic sclerosis, radiation-induced fibrosis, Crohn's disease, ulcerative
10 colitis, multiple myeloma, ocular inflammatory disease, graft versus host disease, graft rejection (e.g., corneal and retinal graft rejection) or cachexia.

The CD154 mutational variants can also be administered prophylactically to a patient who has not
15 yet shown symptoms of the diseases. For instance, the CD154 variants can be used to treat patients who will undergo transplantation so as to prevent or mitigate possible graft rejection.

In one embodiment, the CD154 variant protein
20 can be introduced to a target cell by local injection of liposomes or other suitable carriers (e.g., microspheres) that contain the variant protein. For enhanced targeting, the liposomes or other suitable carriers may be coated with molecules which function as
25 ligands of tissue-specific receptors.

Gene Therapy

According to this invention, a TNFH-minus CD154 mutant may also be introduced into a target cell by expressing within the cell a nucleic acid construct
30 comprising a promoter sequence operably linked to a sequence encoding the mutant CD154 protein.

(1) VECTORS

A nucleic acid construct according to this invention may be derived from a non-replicating linear

- 11 -

or circular DNA or RNA vector, or from an autonomously replicating plasmid or viral vector. Alternatively, the construct may be integrated into the host genome. Any vector that can transfect or transduce a T cell may
5 be used. Preferred vectors are viral vectors, including those derived from replication-defective retroviruses (see, e.g., WO 89/07136; Rosenberg et al., *N. Eng. J. Med.* 323(9): 570-578 (1990)), adenovirus (see, e.g., Morsey et al., *J. Cell. Biochem., Supp.* 17E
10 (1993)), adeno-associated virus (Kotin et al., *Proc. Natl. Acad. Sci. USA* 87:2211-2215 (1990)), replication-defective herpes simplex viruses (HSV; Lu et al., Abstract, page 66, Abstracts of the Meeting on Gene Therapy, Sept. 22-26, 1992, Cold Spring Harbor
15 Laboratory, Cold Spring Harbor, New York), vaccinia virus (Mukherjee et al., *Cancer Gene Ther.* 7:663-70 (2000)), and any modified versions of these vectors. Methods for constructing expression vectors are well known in the art. See, e.g., Sambrook et al.,
20 *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory, 2nd Edition, Cold Spring Harbor, New York, 1989).

The vectors of this invention may target T cells specifically. T-cell-specific viral vectors
25 useful in gene therapy are known in the art. For instance, one can use (1) the retroviral vectors of Annenkov et al., *Gene Therapy* 7:714-22 (2000); Cavazzana-Calvo et al., *Science* 288:669-72 (2000); and Farson et al., *J. Gene Med.* 1:195-209 (1999); (2) the
30 herpesvirus saimiri vector of Hiller et al., *Gene Therapy* 7:664-74 (2000); (3) the HIV-based hybrid vectors of Kung et al., *J. Virol.* 74:3668-81 (2000); (4) the HIV-derived lentiviral vectors of Costello et

- 12 -

al., *Gene Therapy* 7:596-604 (2000); or (5) any modified versions of the above-mentioned vectors.

(2) EXPRESSION CONTROL SEQUENCES

In these vectors, expression control
5 sequences are operably linked to the nucleic acid sequence encoding the mutant protein of the invention. Any expression control sequences than can direct a desired level of transcription in T cells may be used. For eukaryotic cells, expression control sequences may
10 include a promoter, an enhancer, such as one derived from an immunoglobulin gene, SV40, cytomegalovirus, etc., and a polyadenylation sequence. A nucleic acid construct of this invention may also contain an internal ribosome entry site ("IRES"), and an intron
15 that may be desirably located between the promoter/enhancer sequence and the mutant CD154-coding sequence. Selection of these and other common vector elements are conventional. See, e.g., Sambrook et al, *supra*; Ausubel et al., *Current Protocols in Molecular Biology*, John Wiley & Sons, New York, (1989); and
20 references cited therein.

In one embodiment of the present invention, the native promoter for CD154 is used. The native promoter may be preferred when it is desired that
25 expression of mutant CD154 should mimic the native expression. The native promoter may be used when expression of mutant CD154 must be regulated temporally or developmentally, or in a tissue-specific manner, or in response to specific native transcriptional stimuli.
30 In a further embodiment, other native expression control elements, such as enhancer elements, polyadenylation sites or Kozak consensus sequences may also be used to mimic the native expression.

- 13 -

In another embodiment of the present invention, T-cell-specific promoters are desired. Such promoters include two classes: cell type specific promoters and activation specific promoters. Examples
5 of such promoters include, without limitation, promoters derived from the genes of CD2, CD4, CD3, T cell receptor α and β chains and IL-2.

To prevent prolonged immunosuppression, it may be desirable to use inducible promoters to regulate
10 the expression of mutant CD154. Such promoters are known in the art. They include, without limitation, (1) tetracycline-inducible promoters (Gossen et al., *Science* 268:1766-1769 (1995); Harvey et al., *Curr. Opin. Chem. Biol.* 2:512-518 (1998)); (2) tetracycline-
15 suppressible promoters (Alvarez-Vallina et al., *Cancer Gene Therapy* 7:526-9 (2000); Gossen et al., *Proc. Natl. Acad. Sci. USA* 89:5547-5551 (1992)); (3) the rapamycin-inducible promoter systems of Ye et al., *Science* 283:88-91 (1999); Magari et al., *J. Clin. Invest.* 100:2865-2872 (1997); and Rivera et al., *Nat. Medicine* 2:1028-1032 (1996); (4) the zinc-inducible metallothionine promoter; (5) the dexamethasone (Dex)-inducible mouse mammary tumor virus (MMTV) promoter; (6) the T7 polymerase promoter system (WO
25 98/10088); (7) the ecdysone insect promoter (No et al., *Proc. Natl. Acad. Sci. USA* 93:3346-3351 (1996)); (8) the RU486-inducible promoter systems (Wang et al., *Nat. Biotech.* 15:239-243 (1997); Wang et al., *Gene Ther.* 4:432-441 (1997)); and (9) the modified versions of the
30 the above promoter systems. Other types of inducible promoters useful in this invention are those regulated by a specific physiological state, e.g., temperature, acute phase, or in replicating cells only.

- 14 -

In yet another embodiment of the present invention, high-level constitutive expression is desired. Exemplary promoters for this purpose include, without limitation, the retroviral Rous sarcoma virus (RSV) LTR promoter/enhancer, the cytomegalovirus (CMV) immediate early promoter/enhancer (see, e.g., Boshart et al, *Cell* 41:521-530 (1985)), the SV40 promoter, the dihydrofolate reductase promoter, the cytoplasmic β -actin promoter and the phosphoglycerol kinase (PGK) promoter.

Using the guidance provided by this application, one of skill in the art may make a selection among the above expression control sequences and modified versions thereof without departing from the scope of this invention.

(3) ADMINISTRATION OF NUCLEIC ACID CONSTRUCTS

The nucleic acid constructs of this invention may be formulated as a pharmaceutical composition for use in any form of transient and/or stable gene transfer *in vivo* and *in vitro*. The composition comprises at least the nucleic acid construct and a pharmaceutically acceptable carrier such as saline. Other aqueous and non-aqueous sterile suspensions known to be pharmaceutically acceptable carriers and well known to those of skill in the art may be employed also. The construct may be used for *in vivo* and *ex vivo* gene therapy, *in vitro* protein production and diagnostic assays.

The nucleic acid construct can be introduced into target cells as naked DNA, or by, e.g., liposome fusion (see, e.g., Nabel et al., *Science* 249:1285-8 (1990); Ledley, *J Pediatrics* 110:1-8 and 167-74 (1987); Nicolau et al., *Proc Natl Acad Sci USA* 80:1068-72

- 15 -

(1983)), erythrocyte ghosts, or microsphere methods (microparticles; see, e.g., United States patents 4,789,734, 4,925,673, and 3,625,214; Gregoriadis, *Drug Carriers in Biology and Medicine*, pp. 287-341, Academic Press, 1979). Alternatively, the nucleic acid construct can be coupled to ligands of T-cell-specific receptors, and thereby enter T cells via receptor-mediated endocytosis.

If the nucleic acid construct is viral-based, it can also be packaged as a virion which then is used to transduce a cell (e.g., an autologous T cell isolated from a patient) ex vivo. The infected cell is then returned to the body of the patient. Alternatively, the recombinant virus may be administered to a patient directly, e.g., intravenously, intraperitoneally, intranasally, intramuscularly, subcutaneously, and/or intradermally, as determined by one skilled in the gene therapy art. A slow-release device, such as an implantable pump, may be used to facilitate delivery of the recombinant virus to a cell. Where the virus is administered to a subject, the specific cells to be infected may be targeted by controlling the method of delivery. For example, intravenous injection of the virus may be used to facilitate targeting the virus to a circulating T cell. Target areas of the body for local delivery sites include, for example, the lungs, skin, lymph nodes, thymus, spleen and bone marrow. Such delivery may be, for example, by topical, inhalation, aerosol or local injection routes, including, for example, portal vein catheters. The treatments of the invention may be repeated as needed, as determined by one skilled in the art.

- 16 -

Dosages of the nucleic acid construct of this invention in gene therapy will depend primarily on factors such as the condition being treated. The dosage may also vary depending upon the age, weight and health of the patient. For example, an effective human dosage of a mutant CD154-coding virus is generally in the range of from about 0.5 ml to 50 ml of saline solution containing the virus at concentrations of about 1×10^7 , 1×10^8 , 1×10^9 , 1×10^{10} , 1×10^{11} , 1×10^{12} , 1×10^{13} , 1×10^{14} , 1×10^{15} , or 1×10^{16} viral particles per dose administered. The dosage will be adjusted to balance the corrective benefits against any adverse side effects. The levels of expression of mutant CD154 may be monitored to determine the type and frequency of dosage administration.

The pharmaceutical compositions of the invention may be used alone or in a mixture, or in chemical combination, with one or more materials, including other proteins or recombinant vectors that increase the biological stability of the proteins or the recombinant vectors, or with materials that increase the compositions' ability to target T cells selectively.

Combined Therapy

In addition, the pharmaceutical compositions of the invention can be used in combination with another immunomodulating regimen to achieve desired immunosuppression, e.g., long-term, rejection-free integration of heterologous donor tissue into a primate recipient. For instance, an agent that blocks the CD154:CD40 interaction, or blocks costimulation via CD28, CD80 or CD86 can be used.

Exemplary CD154:CD40 interaction inhibitors are antibodies against CD154, such as monoclonal

- 17 -

antibodies ("mAbs") 5c8 (produced by the hybridoma having ATCC Accession Number HB 10916; disclosed in United States patent 5,474,771); ImxM90, ImxM91 and ImxM92 (described in United States patent 5,961,974);
5 and those commercially available from Ancell (clone 24-31, catalog # 353-020, Bayport, MN), Genzyme (Cambridge, MA, catalog # 80-3703-01), and PharMingen (San Diego, catalog #33580D). Numerous additional anti-CD154 antibodies have been produced and
10 characterized (see, e.g., PCT patent application WO 96/23071 of Bristol-Myers Squibb).

Other known immunomodulators that block CD154:CD40 interaction include anti-CD154 molecules of other types, such as complete Fab fragments, F(ab')₂,
15 compounds, V_H regions, F_V regions, single chain antibodies (see, e.g., PCT patent application WO 96/23071), polypeptides, fusion proteins (such as CD40Ig, as in Hollenbaugh et al., *J. Immunol. Meth.* 188:1-7 (1995)), and small molecule compounds such as
20 small semi-peptidic compounds or non-peptidic compounds. All of these immunomodulators are capable of blocking or interrupting CD40:CD154 interaction. Procedures for designing, screening and optimizing small molecules are provided in PCT patent publication
25 WO 97/00895, the specification of which is hereby incorporated by reference.

To avoid potential immune responses to the recombinant virus during gene therapy, it may also be desired to adopt the treatment regimen identical or
30 similar to that described in Chirmule et al., *J. Virol.* 74:3345-52 (2000). In this regimen, a patient is first treated concurrently with (1) the recombinant virus without the mutant CD154 insert and (2) an immunomodulator such as a humanized anti-CD154 antibody

- 18 -

or another costimulation inhibitor. Then the patient is treated with the mutant CD154-coding virus. It has been demonstrated that such a two-step regimen results in significant and prolonged inhibition of the
5 recombinant virus-specific humoral response that often causes side effects in gene therapy patients.

See also United States patent 5,872,174 and PCT patent application WO 96/26285.

10 **Pre-Clinical Model Systems for
Evaluating Mutant CD154 Gene Therapy Regimens**

An exemplary model system for testing efficacy of mutant CD154 gene therapy regimens is the primate renal allograft model disclosed in Kirk et al., *Proc. Natl. Acad. Sci. USA* 94:8789-94 (1997), the
15 teachings of which are incorporated by reference herein. This rhesus monkey model can serve as a rigorous test of immune manipulation: one that is exquisitely sensitive to even minor changes in allograft function or adverse effects on recipient
20 wound healing and immune system function. In addition, it has biological similarity to human renal transplantation: specifically, genes that encode MHC proteins are well conserved between rhesus monkeys and humans, and rhesus monkeys' rejection of vascularized
25 organs closely parallels that seen clinically.

It will be readily appreciated that this model system is suitable for evaluating grafts comprising renal (kidney) tissue. Other art-recognized preclinical model systems, preferably primate model
30 systems, are suitable for assessing efficacy of mutant CD154 gene therapy in suppressing rejection of other graft tissue types such as liver, heart, lung, pancreas, pancreatic islet, skin, peripheral and central nerve. In addition, efficacy of mutant CD154

- 19 -

gene therapy according to the present invention may be assessed in animal models of lupus nephritis, such as those described in PCT patent applications WO 98/30240 and WO 98/30241. Such efficacy may also be assessed in
5 animal corneal allograft models, such as murine corneal allograft models.

EXAMPLES

The following examples are meant to illustrate the methods and materials of the present
10 invention. Suitable modifications and adaptations of the described conditions and parameters normally encountered in the art which are obvious to those skilled in the art are within the spirit and scope of the present invention.

15 Example 1: Procedures

(1) Cell lines, Antibodies, and Ig Fusion Protein

BJAB, a human B cell line, was a gift from Dr. George Mosialos at Harvard Medical School. The cell line was maintained in a RPMI medium supplemented
20 with penicillin, streptomycin, 10% heat inactivated fetal bovine serum, and 4 mM glutamine. The mAb 9E10 was produced and purified from the culture of the 9E10 hybridoma, available from the American Type Culture Collection. See also G.I. Evan et al., "Isolation of
25 Monoclonal Antibodies Specific for Human c-myc Proto-Oncogene Product", *Mol Cell Biol* 5:3610-3616 (1985) regarding mAb 9E10 and the EQKLISEEDL myc tag. Procedures for engineering, producing and purifying
CD40-Fc fusion protein, humanized 5c8, and rabbit
30 polyclonal antibodies Rb779 and Rb784 were carried out as described in Hsu et al., *J Biol Chem* 272:911-5 (1997).

- 20 -

**(2) Transient Expression of Wildtype
and Mutant CD154 Proteins**

A wildtype CD154 cDNA was isolated from a cDNA library made from activated human peripheral blood
5 cells. The coding sequence of this cDNA encodes a protein having an amino acid sequence of SEQ ID NO:1. Procedures for constructing mutants of CD154 were carried out as described in Garber et al., *supra*. The wildtype and mutant cDNAs were subcloned or
10 reconstructed from restriction fragments with confirmed sequence into a unique NotI site in a CMV, immediate early promoter-driven, expression vector containing an SV40 origin for amplification in COS7 cells. COS7 cells were transfected with supercoiled plasmid DNA
15 using lipofectamine (GibcoBRL, Grand Island, New York) following the manufacture's instructions. Plasmid DNA lacking the CD154 coding sequence was used as a negative control, and a vector containing the wild-type CD154 coding sequence was used as a positive control in
20 all examples. Expression of CD154 and its variants was analyzed on transfected cells harvested 72 hours after transfection. Metabolic labeling and biotinylation of cell surface proteins, immunoprecipitation, SDS-polyacrylamide gel electrophoresis ("SDS-PAGE"),
25 and western blotting analysis were performed using procedures described in Hsu et al., *supra*.

Preparation of membrane fractions from transfected COS7 cells and the procedures for analysis of lymphotoxin- α upregulation were performed as
30 previously described (Garber et al., *supra*).

- 21 -

**Example 2: TNFH(-) mutant is
associated with wildtype CD154.**

To investigate the effect of TNFH(-) mutant on the expression of wildtype CD154, lysates of
5 metabolically labeled COS7 cells co-transfected with cDNAs encoding these proteins were analyzed by immunoprecipitation using CD154-specific antibodies. To distinguish the mutant protein from the p18 component of the wildtype heterotrimeric complexes (Hsu
10 et al., *supra*), a myc tag was engineered at the C-terminal of the first 96 amino acids of CD154 to replace the non-CD154 amino acids (11 or 21 amino acids) predicted in the aberrantly spliced transcripts (patients 19, 20 and 21 in Seyama et al., *supra*.

15 To determine the association of CD154 mutants with the wildtype protein, cell lysates prepared from ³⁵S-metabolically labeled COS7 cells transfected with cDNA encoding either the full length wildtype or the myc-tagged mutant CD154 (amino acid residues 1-96 of
20 SEQ ID NO:1) or both were immunoprecipitated with anti-myc mAb 9E10, anti-human CD154 mAb 5c8, or anti-CD154-C-terminal peptide antiserum Rb784 (or "784"). Immunoprecipitates were analyzed by electrophoresis on 10-20% gradient SDS-PAGE gel, followed by
25 autoradiography (Fig. 1).

Fig. 1 shows that when wildtype CD154 was expressed alone, immunoprecipitates of anti-CD154-N-terminal peptide antiserum Rb784 (lane 3) and anti-CD154 mAb 5c8 (lane 4) contained primarily the
30 full length p33 protein, some p31 protein, and a small amount of p18 (p33, p31 and p18 are components of wildtype CD154). This is consistent with our previous observation (Hsu et al., *supra*). The p33, p31 and p18 components were not observed in the immunoprecipitates

- 22 -

of either anti-myc mAb 9E10 (lane 1) or the control rabbit antiserum (lane 2). When a mutant CD154 protein (containing amino acid residues 1-96 OF SEQ ID NO:1 and linked to a myc tag) ("TNFH(-)") was expressed alone, a p17 component, corresponding to the mutant protein itself, was immunoprecipitated by both 9E10 and Rb784 but not by mAb 5c8 or the control rabbit antiserum (lanes 5 to 8). When mutant and wildtype proteins were co-expressed, immunoprecipitates of 9E10 contained not only the p17 myc-tagged mutant protein but also the p33, p31, and p18 wildtype proteins (lane 9). Similar protein patterns were found in immunoprecipitates of Rb784 and mAb 5c8, but not the control serum (lanes 10 to 12). Notably, mAb 5c8 immunoprecipitates exhibited a less amount of these proteins, suggesting that not all of the wildtype proteins expressed were recognized by 5c8.

The above data reveal three important findings. First, mutant CD154 proteins can be stably produced and readily detected. Second, mutant CD154 proteins missing the TNFH domain can associate with wildtype CD154. Third, at least some wildtype proteins, while associated with the mutant proteins, can interact with mAb 5c8 whose binding epitope has been shown to be conformational and similar to the CD154 binding site for CD40.

Example 3: Association of TNFH(-) mutant protein diminishes the receptor-binding activity of the wildtype protein in a dose dependent fashion.

The data shown in Fig. 1 suggest that the association of TNFH(-) CD154 with wildtype CD154 may compromise the ability of the wildtype protein to interact with its receptor. To further examine this, we transfected COS7 cells with (1) a constant amount of

- 23 -

a plasmid containing cDNA encoding full length CD154 and a varied amount of a plasmid containing cDNA encoding TNFH(-) CD154. The transfectants were then metabolically labeled with ³⁵S and lysed. Immuno-
5 precipitates of the lysates were analyzed by a 10-20% gradient SDS-polyacrylamide gel followed by autoradiography (Fig. 2).

Fig. 2 shows that the amount of wildtype CD154 immunoprecipitated by Rb784 was the same
10 regardless of the amount of TNFH(-) DNA used for transfection (Fig. 2, bottom panel). In lane 1, the cells were transfected with only the plasmid encoding wildtype CD154. In lanes 2-4, the cells were transfected with the same amount of the wildtype CD154
15 plasmid and a varied amount of the TNFH(-) plasmid; and the ratios of the former plasmid to the latter plasmid were 3:1, 1:1, and 1:3, respectively. In lane 6, the cells were transfected with only a control plasmid.

The data shown in Fig. 2 indicates that the
20 expression of the wildtype protein was not affected by the introduction of the TNFH(-) mutant. However, the amount of wildtype protein recognized by a CD40-Fc fusion protein or by mAb 5c8 was inversely proportional to the amount of the TNFH(-) plasmid used for
25 transfection (Fig. 2, top and middle panels, respectively). Interestingly, only a small portion of the total TNFH(-) mutant expressed (Fig. 2, bottom panel) was immunoprecipitated by either CD40-Fc or mAb 5c8. These results show that expression of the TNFH(-)
30 mutant affected the receptor-interacting activity, but not the expression level, of wildtype CD154 in a dose dependent fashion.

- 24 -

**Example 4: Effect of TNFH(-) mutant
protein on the function of wildtype CD154.**

To test if the co-expression of mutant CD154 affects the function of the wildtype protein,
5 UV-irradiated membranes from transfected COS7 cells transfected with cDNA(s) encoding wildtype and/or mutant CD154 were incubated with BJAB cells for 24 h. Ratios of wildtype to mutant cDNAs used for COS7 transfections are indicated in Fig. 3B. The BJAB cells
10 were stained with biotin-labeled anti-lymphotoxin- α mAb, NC2, then with phycoerythrin-labeled streptavidin, and finally fixed with 1% paraformaldehyde. FACS analysis of the BJAB cells was subsequently used to determine the level of up-regulated lymphotoxin- α on
15 the surface of the cells (Fig. 3A, in which the bars represent the mean fluorescence intensities).

Figs. 3A and 3B show that co-expression of the mutant CD154 protein with the wildtype protein reduced the lymphotoxin- α up-regulating ability of the
20 wildtype protein present on the plasma membranes. This inhibitory effect was dose-dependent. When the amount of wildtype cDNA used for transfection was one third of the mutant cDNA, the functional activity of CD154 was down to the background level.

**25 Example 5: TNFH(-) mutant
does not express on the cell surface.**

There are at least two possible mechanisms by which mutant CD154 inhibits wildtype CD154 functions in a dose-dependent manner. First, a wildtype CD154
30 associated with a mutant CD154 may be less active in interacting with CD40. Second, a wildtype CD154 associated with a mutant CD154 may not be functional at all and the activity observed in the up-regulation of

- 25 -

lymphotoxin- α may be due exclusively to the wildtype proteins that are free from association with mutant proteins. To distinguish these two possibilities, we next examined the biochemical properties of CD154 proteins expressed on the cell surface.

To do this, COS7 cells were transfected on day one with cDNA encoding wildtype CD154, or cDNA encoding the TNFH(-) mutant, or both. The transfectants were rinsed on day four with PBS to remove cell debris and unattached cells, and then labeled with biotin *in situ*. Labelling was carried out using biotin-sulfo-NHS at 0.5 mg/ml in water left on the cells for 3 minutes at room temperature. The reaction was stopped using glycine, with the transfectant cells remaining in a vast molar excess of glycine for 15 minutes, after which they were rinsed with PBS.

The transfectant cells were then lysed, centrifuged to remove cell debris, and immunoprecipitated with CD40-Fc, Rb779, Rb784, or 9E10. Immunoprecipitates were subjected to electrophoresis in 10-20% SDS-polyacrylamide gel, transferred to nitrocellulose membrane and probed with horseradish peroxidase-conjugated streptavidin (Fig. 4).

Fig. 4 shows that CD40-Fc fusion protein, Rb784, and Rb779, but not 9E10, immunoprecipitated the biotin-labeled wildtype proteins. In addition, the amount of wildtype CD154 detected at the cell surface was inversely proportional to the ratio of mutant cDNA to wildtype cDNA used for co-transfection. Thus, the expression of the TNFH(-) mutant protein prevented the surface expression of the wildtype protein in a dose-dependent fashion. Importantly, we did not detect biotin-labeled mutant protein (p17) in

- 26 -

immunoprecipitates with CD40-Fc fusion protein, Rb784, and Rb779, suggesting that the biotin-labeled, i.e. cell surface expressed, wildtype proteins were not associated with the TNFH(-) mutant protein. Since
5 there is a total of eight lysine residues in the extracellular domain of the TNFH(-) mutant, it was unlikely that none of these lysine residues was accessible for biotinylation under the experimental conditions such that mutant proteins present on the
10 surface were undetected. Moreover, the fact that 9E10 did not immunoprecipitate any wildtype proteins indicates that none of the cell surface wildtype protein was associated with the TNFH(-) mutant.

Together with the results shown in Fig. 1,
15 these data indicate that wildtype CD154 proteins free of association with the TNFH(-) mutant were expressed on the cell surface. However, wildtype proteins complexed with the TNFH(-) mutant were retained inside the cell. Thus, association with the TNFH(-) mutant
20 prevents wildtype CD154 from being expressed on the cell surface.

Fig. 5 illustrates this inhibitory process. When the wildtype and mutant proteins are co-expressed, wildtype proteins can form trimers with or without the
25 TNFH(-) mutant associated. However, those bound by the mutant protein cannot mature onto the cell surface. The association with TNFH(-) per se does not compromise the ability of the wildtype protein to interact with the receptor, CD40.

30 **Example 6: Discussion of experimental results.**

To study the potential dominant negative effect of the TNFH(-) variant on wildtype CD154, we co-expressed these two proteins in COS cells and examined their biochemical and functional properties.

- 27 -

Our results indicate that by complexing with the wildtype protein, the mutant protein prevents the cell surface expression of wildtype CD154. This observation is significant in at least two aspects. First, it
5 implies that while a patient's cells are capable of making both wildtype and TNFH(-) mutant CD154 proteins, they fail to produce functional CD154 on the cell surface and lead to a hyper-IgM phenotype. Second, our observation reveals that in addition to the TNFH
10 domain, other parts of CD154 also participate in the assembly of CD154 trimers.

As CD154 is transiently expressed on the cell surface of T lymphocytes upon activation, even a partial reduction of surface expression of this protein
15 will be beneficial in suppressing undesired T cell mediated immunological responses. Thus, the introduction of a CD154 variant lacking a functional TNFH domain into target T cells will provide a therapeutic opportunity by effectively blocking the
20 expression of functional CD154 on the surface of the target cells.

OTHER EMBODIMENTS

It is to be understood that while the invention has been described in conjunction with the
25 detailed description thereof, the foregoing description is intended to illustrate and not to limit the scope of the invention, which is defined by the scope of the appended claims.

Other aspects, advantages, and modifications
30 are within the scope of the following claims.

- 28 -

CLAIMS

What is claimed is:

1. A method of decreasing the expression of wildtype CD154 on the surface of a cell, the method comprising
5 the step of introducing into the cell a nucleic acid construct that directs expression of a mutant CD154 lacking at least five amino acid residues of the tumor necrosis factor homologous domain of wildtype CD154, whereby the expressed mutant CD154 binds to the
10 wildtype CD154 inside the cell, rendering the wildtype CD154 unable to reach the cell surface.
2. A method of treating a patient suffering from or predisposed to a CD154-mediated disease, the method comprising the step of delivering to said patient a
15 nucleic acid construct that directs expression of a mutant CD154 lacking at least five amino acid residues of the tumor necrosis factor homologous domain of wildtype CD154 in a cell of the patient, whereby the expressed mutant CD154 binds to the wildtype CD154
20 inside the cell, rendering the wildtype CD154 unable to reach the cell surface.
3. The method according to claim 1 or 2, wherein the nucleic acid construct comprises a virus-derived vector.
- 25 4. The method according to claim 3, wherein the virus-derived vector is a retroviral vector.
5. The method according to claim 3, wherein the virus-derived vector is a lentiviral vector.

- 29 -

6. The method according to claim 3, wherein the virus-derived vector is an adenoviral vector.
7. The method according to claim 3, wherein the virus-derived vector is an adeno-associated viral
5 vector.
8. The method according to claim 3, wherein the nucleic acid construct is introduced into the cell via viral transduction.
9. The method according to claim 1 or 2, wherein the
10 cell is a T cell or a megakaryocyte.
10. The method according to claim 1 or 2, wherein the cell is a mammalian cell.
11. The method according to claim 10, wherein the cell is a human cell.
- 15 12. The method according to claim 10, wherein the cell is a human T cell.
13. The method according to claim 1 or 2, wherein the nucleic acid construct is introduced into the cell *in vivo*.
- 20 14. The method according to claim 1 or 2, wherein the nucleic acid construct is introduced into the cell *ex vivo*.
15. The method according to claim 2, wherein the CD154-mediated disease is graft rejection.

- 30 -

16. The method according to claim 2, wherein the CD154-mediated disease is an autoimmune disease.

17. The method according to claim 2, wherein the CD154-mediated disease is an inflammatory disease.

5 18. The method according to claim 2, wherein the CD154-mediated disease is selected from the group consisting of lupus, systemic lupus erythematosus, lupus nephritis, lupus neuritis, asthma, chronic obstructive pulmonary disease, bronchitis, emphysema,
10 multiple sclerosis, uveitis, Alzheimer's disease, traumatic brain injury, traumatic spinal cord injury, stroke, atherosclerosis, coronary restenosis, ischemic congestive heart failure, cirrhosis, hepatitis C virus, diabetic nephropathy, glomerulonephritis, autoimmune
15 disease, osteoarthritis, rheumatoid arthritis, psoriasis, atopic dermatitis, systemic sclerosis, radiation-induced fibrosis, Crohn's disease, ulcerative colitis, multiple myeloma, ocular inflammatory disease, graft versus host disease, graft rejection and
20 cachexia.

19. The method according to claim 1 or 2, wherein the mutant CD154 lacks a portion of wildtype CD154, said portion corresponds to (1) amino acid residues 116-136, (2) amino acid residues 137-261, (3) amino acid
25 residues 115-261, or (4) amino acid residues 97-261 of SEQ ID NO:1.

20. The method according to claim 19, wherein the mutant CD154 lacks a portion of wildtype CD154, said portion corresponds to amino acid residues 116-261 of
30 SEQ ID NO:1.

- 31 -

21. A pharmaceutical composition comprising a nucleic acid construct that directs expression of a mutant CD154 lacking at least five amino acid residues of the tumor necrosis factor homologous domain of wildtype CD154, whereby the expressed mutant CD154 binds to the wildtype CD154 inside a cell, rendering the wildtype CD154 unable to reach the cell surface.
22. The pharmaceutical composition according to claim 21, wherein the mutant CD154 lacks a portion of wildtype CD154, said portion corresponds to (1) amino acid residues 116-136, (2) amino acid residues 137-261, (3) amino acid residues 115-261, or (4) amino acid residues 97-261 of SEQ ID NO:1.
23. Use of a nucleic acid that directs expression of a mutant CD154 lacking at least five amino acid residues of the tumor necrosis factor homologous domain of wildtype CD154 for the manufacture of a medicament for decreasing the expression of wildtype CD154 on the surface of a cell.
24. Use of a nucleic acid that directs expression of a mutant CD154 lacking at least five amino acid residues of the tumor necrosis factor homologous domain of wildtype CD154 for the manufacture of a medicament for treating a CD154-mediated disease.
25. The use according to claim 23 or 24, wherein the mutant CD154 lacks a portion of wildtype CD154, said portion corresponds to (1) amino acid residues 116-136, (2) amino acid residues 137-261, (3) amino acid

- 32 -

residues 115-261, or (4) amino acid residues 97-261 of
SEQ ID NO:1.

1 / 5

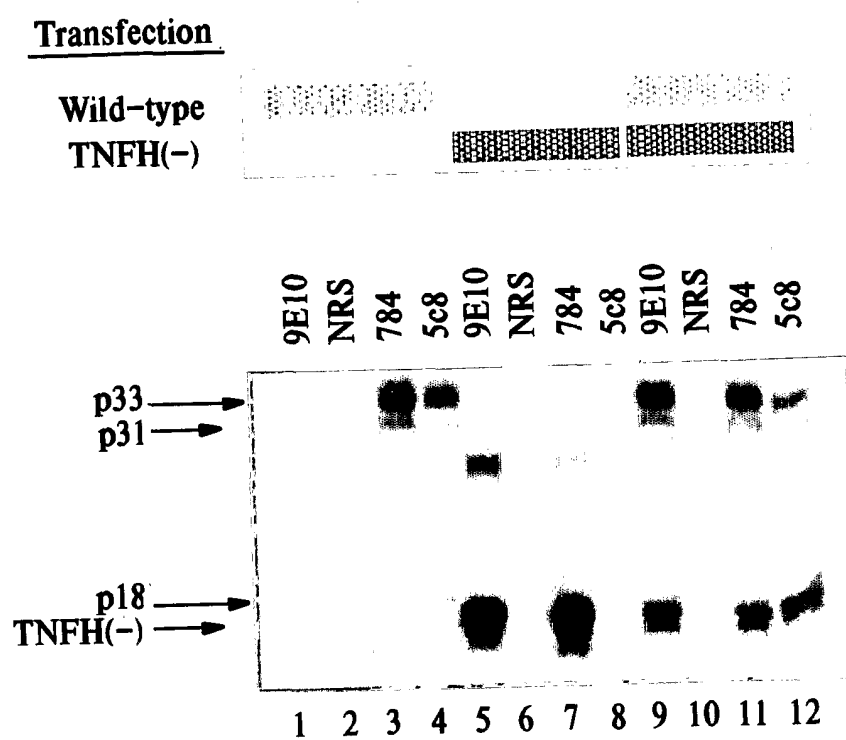


FIG. 1

2 / 5

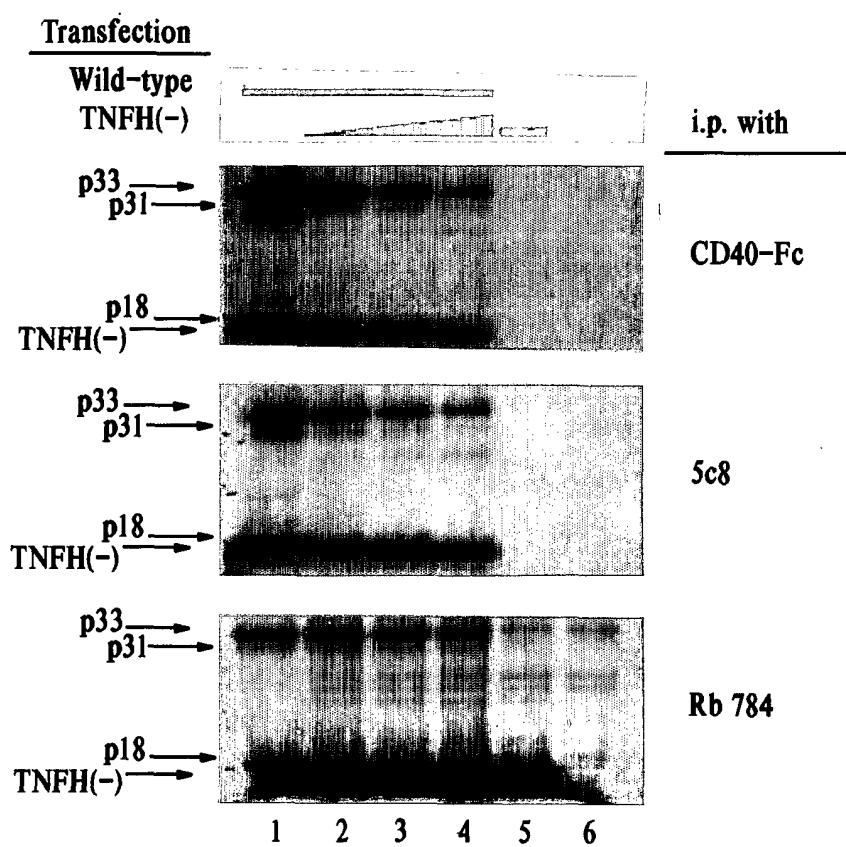


FIG. 2

3/5

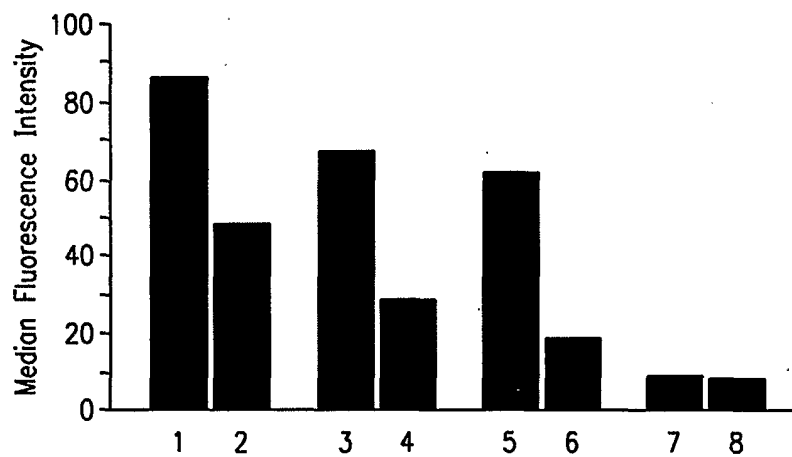


FIG. 3A

	CD154 Transfection		CD154 (ng /mg protein)	MFI	% Activity Reduction
	Wild-type	TNFH(-)			
1	3	0	41.50	86.60	
2	3	1	31.96	48.26	48.47
3	1	0	26.85	66.71	
4	1	1	17.37	27.63	66.00
5	1/3	0	24.82	60.98	
6	1/3	1	14.29	18.11	80.16
7	0	1	0.52	8.35	
8	0	0	0.07	7.50	N/A

FIG. 3B

4 / 5

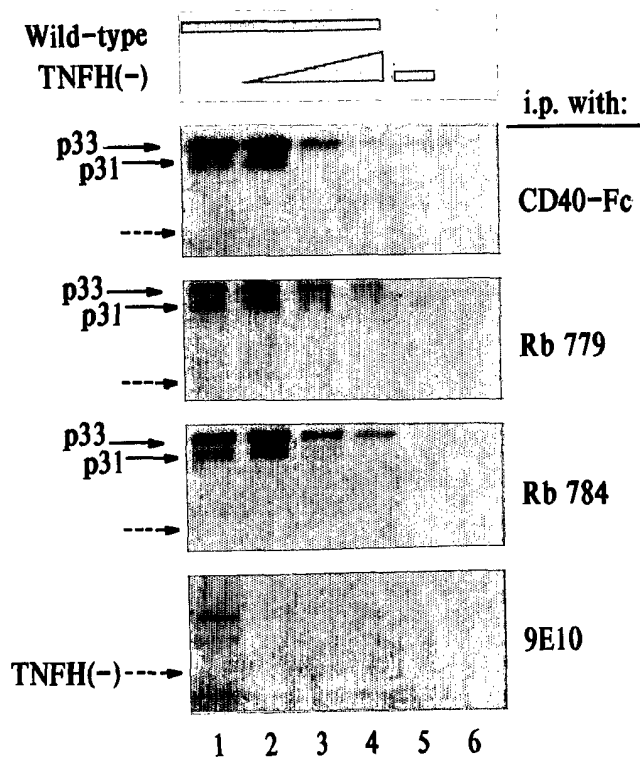
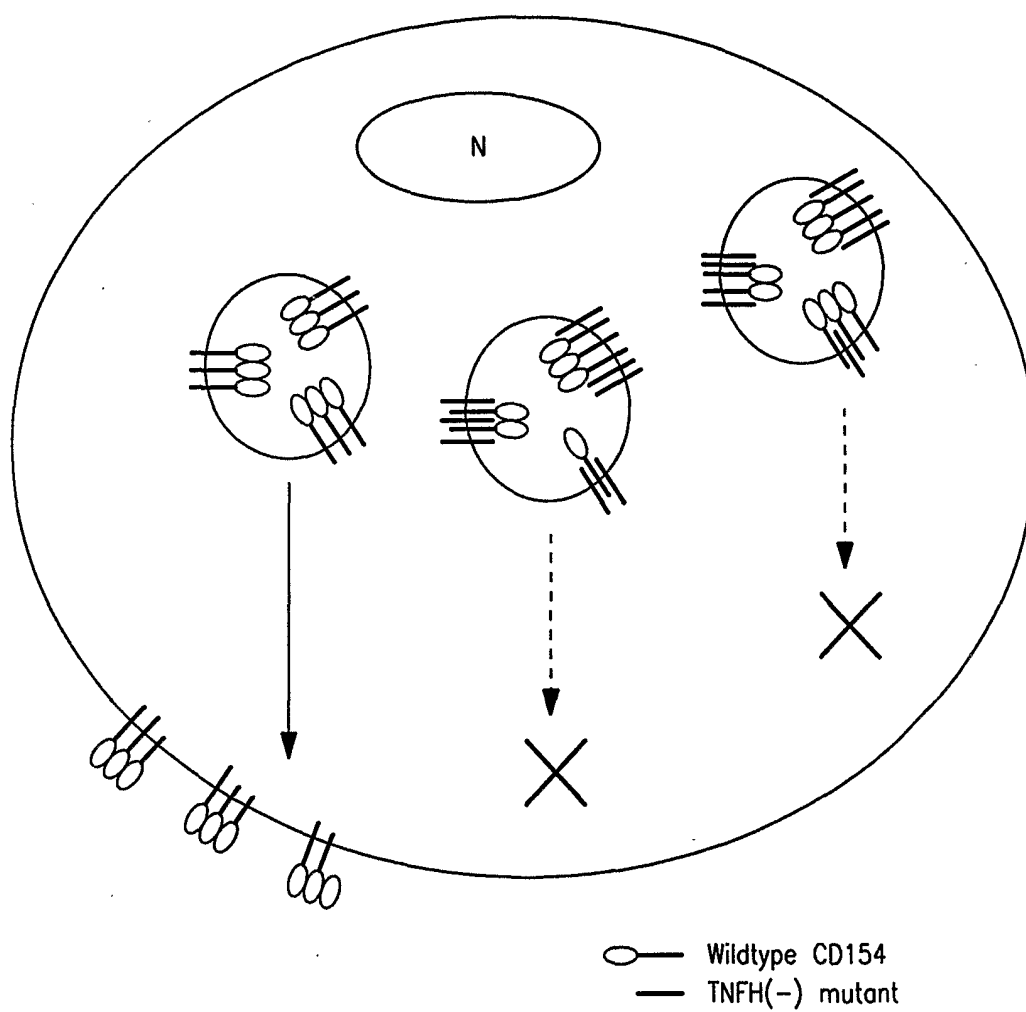
TNFH(-) CD154 variant is absent from cell surfaceTransfection

FIG. 4

5/5

**FIG. 5**

SEQUENCE LISTING

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<120> CD154 VARIANT

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<140>

<141>

<160> 1

<170> PatentIn Ver. 2.1

<210> 1

<211> 261

<212> PRT

<213> Homo sapiens

<400> 1

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		20						25					30		

Ile	Thr	Gln	Met	Ile	Gly	Ser	Ala	Leu	Phe	Ala	Val	Tyr	Leu	His	Arg
		35					40					45			

Arg	Leu	Asp	Lys	Ile	Glu	Asp	Glu	Arg	Asn	Leu	His	Glu	Asp	Phe	Val
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Phe	Met	Lys	Thr	Ile	Gln	Arg	Cys	Asn	Thr	Gly	Glu	Arg	Ser	Leu	Ser
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Leu	Leu	Asn	Cys	Glu	Glu	Ile	Lys	Ser	Gln	Phe	Glu	Gly	Phe	Val	Lys
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Asp	Ile	Met	Leu	Asn	Lys	Glu	Glu	Thr	Lys	Lys	Glu	Asn	Ser	Phe	Glu
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Glu	Ala	Ser	Ser	Lys	Thr	Thr	Ser	Val	Leu	Gln	Trp	Ala	Glu	Lys	Gly
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Tyr	Tyr	Thr	Met	Ser	Asn	Asn	Leu	Val	Thr	Leu	Glu	Asn	Gly	Lys	Gln
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Leu	Thr	Val	Lys	Arg	Gln	Gly	Leu	Tyr	Tyr	Ile	Tyr	Ala	Gln	Val	Thr
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