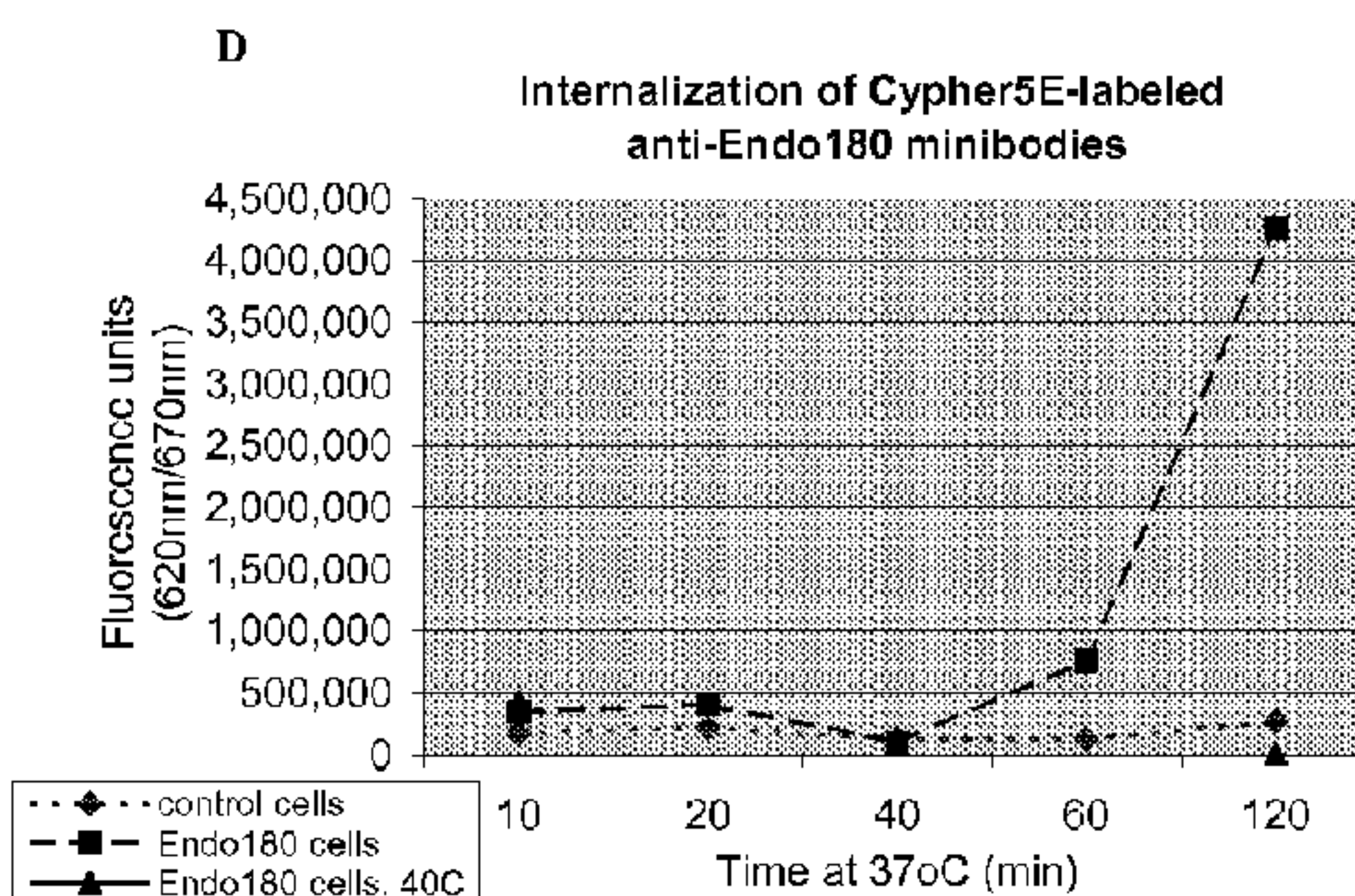
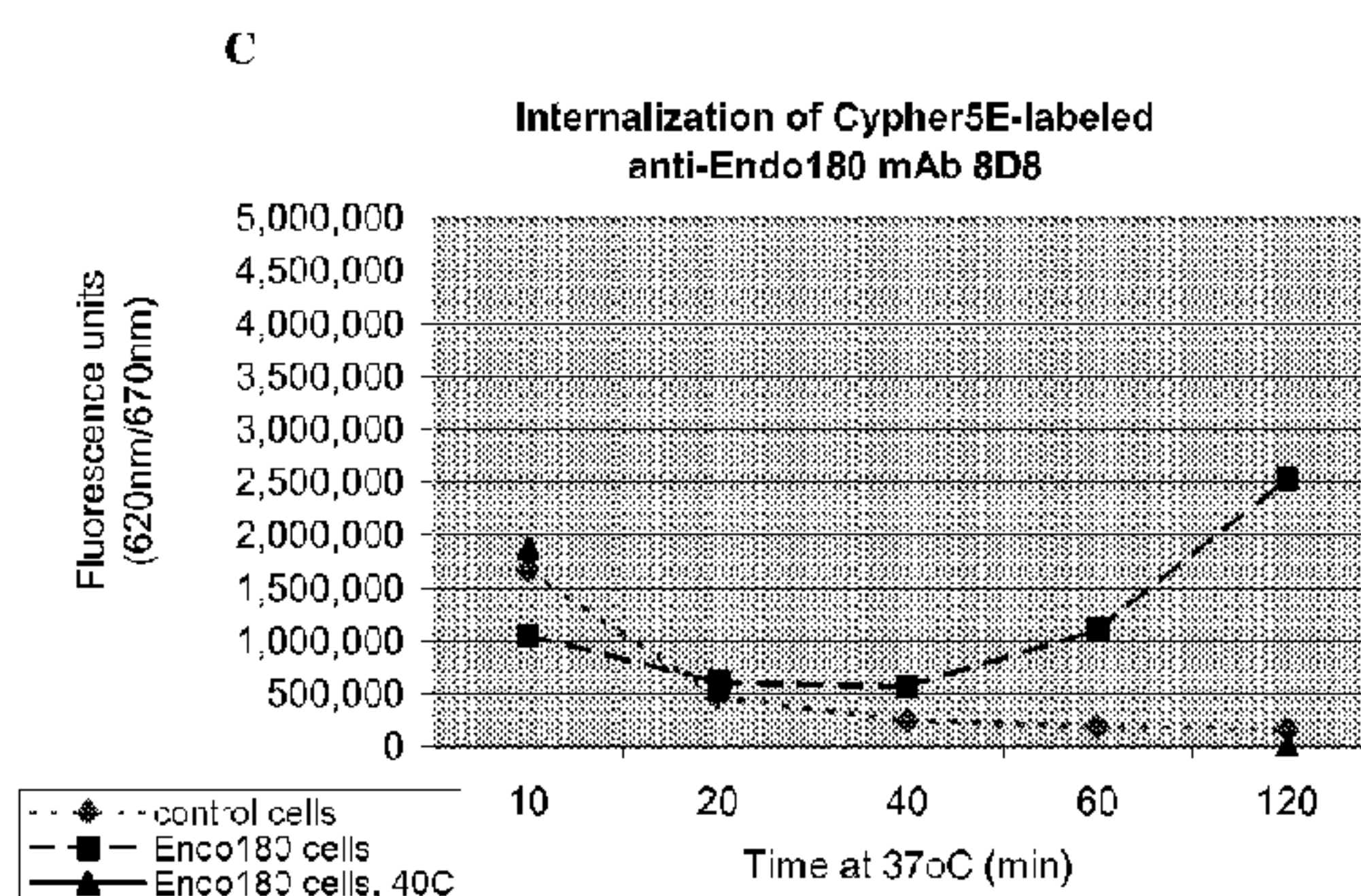




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 (72) **Inventeur/Inventor:**
 FEINSTEIN, ELENA, IL
 (73) **Propriétaire/Owner:**
 QUARK PHARMACEUTICALS, INC., US
 (74) **Agent:** NELLIGAN O'BRIEN PAYNE LLP

(54) **Titre : ANTICORPS ENDO180 SERVANT A TRAITER LE CANCER ET UNE MALADIE FRIBREUSE**
 (54) **Title: ENDO180 ANTIBODY TO TREAT CANCER AND FIBROTIC DISEASE**



(57) **Abrégé/Abstract:**

ENDO180 is a recycling endocytic receptor that directs bound ligands to degradation in the endosomes. The present invention provides, inter-alia, isolation of an anti- ENDO180 antibody (and fragments thereof) that is internalized into a cell upon binding ENDO180 on the cell surface, and is further capable of internalizing moieties conjugated to the antibody. According to some embodiments, the present invention further provides identification of the antibody (i.e., E3-8D8 deposited with the BCCM with Accession Number LMBP 7203CB), conjugates of the antibody with a moiety, compositions of the antibody with a nanoparticle comprising a moiety and use of the antibody, conjugate or composition for delivery of the moiety into an ENDO180 expressing cell. According to some embodiments, the invention provides the ENDO180 antibody for use in treatment of cancer, fibrosis and macrophage- associated disease.

Abstract

ENDO180 is a recycling endocytic receptor that directs bound ligands to degradation in the endosomes. The present invention provides, *inter-alia*, isolation of an anti-ENDO180 antibody (and fragments thereof) that is internalized into a cell upon binding ENDO180 on the cell surface, and is further capable of internalizing moieties conjugated to the antibody. According to some embodiments, the present invention further provides identification of the antibody (i.e., E3-8D8 deposited with the BCCM with Accession Number LMBP 7203CB), conjugates of the antibody with a moiety, compositions of the antibody with a nanoparticle comprising a moiety and use of the antibody, conjugate or composition for delivery of the moiety into an ENDO180 expressing cell. According to some embodiments, the invention provides the ENDO180 antibody for use in treatment of cancer, fibrosis and macrophage-associated disease.

ENDO180 ANTIBODY TO TREAT CANCER AND FIBROTIC DISEASE

5 **FIELD OF THE INVENTION**

The present invention relates to molecules that target the ENDO180 polypeptide and are internalized thereby, to conjugates comprising the molecules, to compositions comprising the molecules and conjugates and to methods of using the same for delivery of therapeutic agents to cells that express an ENDO180 polypeptide on the surface of the cell for treating
10 cell proliferative diseases or disorders and fibrosis, and for controlling (modulating) tumor progression.

BACKGROUND OF THE INVENTIONENDO180 Receptor

ENDO180, also known as CD280, uPARAP (urokinase plasminogen activator receptor associated protein) and mannose receptor C type 2 (MRC2), is a recycling endocytic
15 receptor that directs bound ligands to degradation in the endosomes. It is part of a triple complex with urokinase type plasmin activator (uPA) and urokinase-type plasmin activator receptor (uPAR), thus being involved in the production of plasmin from plasminogen. Plasmin, in turn, is known to play a role in both extracellular matrix (ECM)
20 turnover and proteolytic conversion of latent TGF-beta into its active form.

In addition to its role in the production of plasmin, the triple complex was shown to be involved in the activation of matrix metalloproteinase (MMP) proenzymes, to act on fibrin to bind several collagens and in general turnover of extracellular matrix. This complex also takes part in cell adhesion and signal transduction (Bherendt et al, 2000. JBC
25 275:1993-2002).

ENDO180 is a recycling endocytic receptor that functions in cell motility and remodeling of the extracellular matrix by promoting cell migration and uptake of collagens for 5 aa 2CRD,

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intracellular degradation (Niels. 2004 Biol Chem. 385(2): 103-36; Kjoller et al, 2004 Exp Cell Res. 293(1): 106-16; Wienke et al., 2007 Cancer Res. 67(21): 10230-40.). ENDO180 shares homology with the macrophage mannose receptor family: mannose receptor, phospholipase A2 and DEC-205/MR6 (Isacke et al., 1990 Mol. Cell. Biol. 10:2606-2618; 5 Sheikh et al., 2000, J. Cell. Sci. 113: 1021-1032; Behrendt et al., 2000, J. Biol. Chem. 275: 1993-2002). This family grouping is based on an overall structural conservation: a large extracellular domain comprising an N-terminal signal sequence followed by a cysteine-rich domain, a fibronectin type II domain (FNII), and 8 or 10 C-type lectin-like domains (CTLDs) and small transmembrane and intracellular domains (~66 amino acids together). 10 As a family, these receptors have two striking features: First, although they belong to the large C-type lectin superfamily, they uniquely contain multiple CTLDs within a single polypeptide backbone (Taylor M. E., 1997 Glycobiology 7: v-vii; McKay et al, 1998, Eur. J. Immunol. 28: 4071-4083; Howard and Isacke, 2002, supra). Second, they share the ability to be recycled between the plasma membrane and intercellular compartments of the 15 cell (Isacke et al, 1990, supra; Zvaritch et al., 1996, J. Biol. Chem. 271: 250-257). ENDO180 is unusual in the family of mannose receptors in that it is targeted from the plasma membrane to the recycling endosomes rather than to a late endosome/lysosome compartment (Howard and Isacke, 2002 supra).

ENDO180 is localized on the cell surface, in clathrin coated pits (Isacke et al., 1990 Mol. 20 Cell. Biol. 10: 2606-2618; Sheikh et al., 2000, J. Cell. Sci. 113: 1021-1032) and in endosomes. It is mainly expressed in fibroblasts, endothelial cells and macrophages. In situ hybridization showed its expression in highly vascularized organs. ENDO180 has also been found in bone-forming regions in mouse embryos (Wu et al., 1996, J. Biol. Chem. 271:21323-21330), and in osteoblasts and osteocytes at sites of endochondral and 25 intramembranous ossification during development (Engelholm et al., 2001, Trends Cardiovasc. Med. 11:7-13).

The following patent publications also relate to the ENDO180 receptor: US 6,117,977; US 7,399,468; WO 97/40154 and WO 00/58473. PCT Patent Publication No. WO 2004/100759 and US Patent Publication Nos. 2007/0072244 and 2009/0202566 to the 30 assignee of the present invention and relate to methods of identifying compounds capable of modulating human ENDO180 receptor activity.

Antibody therapy

The search for new therapies to treat cancer and other diseases has resulted in the development of human and humanized antibodies capable of inhibiting receptor function. International patent publication WO 2006/023491 provides a method of RNA interference, which comprises contacting the cell with a fusion protein-double stranded RNA complex, the complex comprising the double stranded RNA segment containing a double stranded RNA of interest and a fusion protein which is an antibody Fab fragment- protamine fusion protein.

SUMMARY OF THE INVENTION

10 The present invention is based in part on the identification of isolated molecules that specifically bind the ENDO180 polypeptide on a cell surface. In some embodiments the molecules bind the extracellular domain of the ENDO180 polypeptide and are internalized into the cell by the polypeptide, thereby providing a vehicle useful for delivery of therapeutic and diagnostic cargo to a cell expressing the ENDO180 polypeptide. 15 Accordingly, in some embodiments the present invention provides a conjugate comprising a molecule that specifically binds the ENDO180 polypeptide and a therapeutic agent useful for the delivery of the therapeutic agent into the cell. In some embodiments the ENDO180 polypeptide is substantially identical to an amino acid sequence set forth in SEQ ID NO:2, encoded by a polynucleotide substantially identical to a nucleic acid 20 sequence set forth in SEQ ID NO:1.

In one aspect the present invention provides an anti-ENDO180 antibody which is produced by hybridoma cell line designated E3-8D8 (BCCM Accession Number LMBP 7203CB), or a fragment of the antibody, which binds to ENDO180 receptor on the surface of a cell. In some embodiments binding of the antibody to the receptor results in 25 internalization of the antibody into the cell. Also provided is the E3-8D8 hybridoma cell line.

In some embodiments the antibody or fragment thereof is humanized or a chimeric antibody or fragment thereof.

The invention provides a composition comprising at least one anti-ENDO180 antibody or 30 fragment thereof, the antibody produced by the E3-8D8 hybridoma or a humanized molecule thereof a chimeric antibody or fragment thereof., together with a carrier.

In some embodiments the isolated antibody is selected from the group consisting of a full IgG, a Fab fragment, a Fab' fragment, an F(ab')₂ fragment, the variable portion of the heavy and/or light chains thereof, Fab miniantibodies, and a scFv. In some embodiments the antibody is a recombinant polypeptide comprising a heavy chain CDR3 domain having an amino acid sequence set forth in SEQ ID NO:7 or a variant thereof which retains the ability to specifically bind ENDO180. In some embodiments the antibody further comprises a light chain CDR3 domain having an amino acid sequence set forth in SEQ ID NO:8 or a variant thereof which retains the ability to specifically bind ENDO180.

In some embodiments the antibody is a scFv recombinant polypeptide comprising an amino acid sequence set forth in SEQ ID NO:6 or a variant thereof, which retains the ability to specifically bind ENDO180. In specific embodiments the antibody exhibiting binding affinity to ENDO180 receptor and comprising CDR3 domains set forth in SEQ ID NOS 7 and 8 is internalized by the receptor into the cell expressing ENDO180 upon contact of the antibody to the receptor.

The invention further provides a composition comprising at least one anti-ENDO180 antibody or fragment thereof, as described above, and a moiety including a radioisotope, a therapeutic agent, a cytotoxic agent, or a detectable label. In some embodiments the moiety is attached (or linked, or conjugated), either covalently, through a linker or a chemical bond, or noncovalently, through ionic, van der Waals, electrostatic, or hydrogen bonds, to the antibody.

In some embodiments provided is an anti-ENDO180 antibody or antigen-binding fragment thereof selected from

- a) the monoclonal antibody produced by the hybridoma cell line E3-8D8 (BCCM Accession Number LMBP 7203CB);
- b) an antibody or fragment thereof that binds to the same epitope as the antibody in (a);
- c) a humanized antibody of (a) or (b);
- d) a fragment of an antibody comprising a polypeptide substantially similar to SEQ ID NO: 6; and
- e) a recombinant polypeptide comprising CDR3 with an amino acid sequence substantially similar to amino acid sequences set forth in SEQ ID NO:7 and 8.

Further provided is a composition comprising an anti-ENDO180 antibody or antigen-binding fragment thereof selected from

- a) the monoclonal antibody produced by the hybridoma cell line E3-8D8 (BCCM Accession Number LMBP 7203CB);
- 5 b) an antibody or fragment thereof that binds to the same epitope as the antibody in (a);
- c) a humanized antibody of (a) or (b);
- d) a fragment of an antibody comprising a polypeptide substantially similar to SEQ ID NO: 6; and
- 10 e) a recombinant polypeptide comprising CDRs having an amino acid sequence substantially similar to amino acid sequences set forth in SEQ ID NO:7 and 8.

In some embodiments the composition further comprises a moiety including a radioisotope, a therapeutic agent, a cytotoxic agent, or a detectable label.

The present invention also provides a method of treating a subject afflicted with a proliferative disorder comprising administering to the subject a composition comprising an
15 anti-ENDO180 antibody or antigen-binding fragment thereof selected from

- a) the monoclonal antibody produced by the hybridoma cell line E3-8D8 (BCCM Accession Number LMBP 7203CB);
- 20 b) an antibody or fragment thereof that binds to the same epitope as the antibody in (a);
- c) a humanized antibody of (a) or (b);
- d) a fragment of an antibody comprising a polypeptide substantially similar to SEQ ID NO: 6; and
- 25 e) a recombinant polypeptide comprising CDRs having an amino acid sequence substantially similar to amino acid sequences set forth in SEQ ID NO:7 and 8.

In some embodiments the proliferative disorder is selected from a solid tumor, a hematopoietic tumor, metastases, fibrosis and a macrophage associated disorder.

In some embodiments the tumor is an ovarian tumor, a breast tumor, osteoblastic/osteocytic cancer, prostate cancer, head and neck cancer, leukemia, renal cell
30 carcinoma, or transitional cell carcinoma.

In some embodiments the fibrosis is liver fibrosis, myelofibrosis, kidney fibrosis for any reason (CKD including end-stage renal disease, ESRD); lung fibrosis (including interstitial lung fibrosis ILF); abnormal scarring (keloids) associated with all possible types of skin injury accidental and iatrogenic (operations); scleroderma; cardiofibrosis, failure of glaucoma filtering operation; intestinal adhesions.

In some embodiments the macrophage-associated disorder is inflammation or atherosclerosis.

In one aspect the present invention provides a conjugate comprising:

- a) an antibody or an antigen binding portion thereof which specifically binds to the extracellular domain of the ENDO180 polypeptide on the surface of a cell;
- b) a moiety including a radioisotope, a therapeutic agent, a cytotoxic agent, or a detectable label.; and
- c) optionally a linking moiety which links (a) to (b).

In some embodiments the moiety is a therapeutic agent selected from an oligonucleotide agent and a non-oligonucleotide agent. In some embodiments the therapeutic agent is an oligonucleotide therapeutic agent, including an inhibitory oligonucleotide. Accordingly, in various embodiments the therapeutic agent is selected from an antisense compound, a chemically modified siRNA compound, an unmodified siRNA compound, a chemically modified shRNA compound, an unmodified shRNA compound, a chemically modified miRNA compound, and an unmodified miRNA compound. In various preferred embodiments the therapeutic agent is chemically modified siRNA. In some embodiments the chemically modified siRNA compound inhibits expression of a target gene associated with cancer, fibrosis or macrophage associated disease. In some embodiments the target gene is selected from any one of the target genes set forth in Table A, hereinbelow.

In certain embodiments the therapeutic agent is attached to the antibody via a nucleotide or non-nucleotide linking moiety.

In yet another aspect the present invention provides a pharmaceutical composition comprising the conjugate of the present invention.

In yet another aspect the present invention provides a method of treating a subject suffering from a proliferative disease comprising administering to the subject a therapeutically effective amount of an antibody that specifically binds ENDO180

polypeptide and is internalized by the ENDO180 polypeptide, wherein the antibody is covalently or non-covalently bound to a therapeutic agent.

In some embodiments the proliferative disease is selected from malignant and benign proliferative disease. In some embodiments proliferative disease is cancer. In other
5 embodiments proliferative disease is fibrosis. Non-limiting examples of diseases and disorders for use of the present invention include

1. soft tissue sarcomas in which ENDO180 is expressed in the tumor and tumor stroma cells (activated myofibroblasts, neovasculature and infiltrating cells of macrophage-monocyte lineage);
- 10 2. carcinomas in which ENDO180 is expressed in the tumor stroma cells (activated myofibroblasts, neovasculature and infiltrating cells of macrophage-monocyte lineage);
3. carcinoma that express ENDO180 and have undergone epithelial-mesenchymal transition thus acquiring high metastatic potential;
4. leukemia expressing ENDO180 for example, from macrophage-monocyte lineage;
- 15 5. fibrotic diseases, for example of kidney, lung and liver with activated myofibroblasts;
6. diseases and disorders associated with macrophage including atherosclerosis and chronic inflammation.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1A-1J provides polynucleotide and amino acid sequences of various compounds
20 according to the present invention. Fig. 1A: human ENDO180 mRNA (SEQ ID NO:1); Fig 1B: human ENDO180 polypeptide (SEQ ID NO:2); Fig. 1C: SEQ ID NO:3 polynucleotide sequence of extracellular domain of human ENDO180 (amino acids 1-522) with FLAG sequence, FLAG domain underlined (pcDNA3-5'hendo180-FLAG construct, SEQ D NO:3); Fig. 1D polypeptide sequence of SEQ ID NO:3 (SEQ ID NO:4); Fig. 1E:
25 polynucleotide sequence of scFv clone G7V (SEQ ID NO:5); Fig. 1F: polypeptide sequence of scFv clone G7V (SEQ ID NO:6); Fig. 1G. heavy chain CDR3 of G7V (SEQ ID NO:7); Fig. 1H. light chain CDR3 of G7V (SEQ ID NO:8); Fig. 1J: polypeptide 1-522 of the extracellular domain of human ENDO180.

Figures 2A-2H. Internalization of CypHer5E fluorophore anti-ENDO180 mAbs to
30 ENDO180 expressing cells.

Figure 3. Internalization of Biotin anti-ENDO180 mAbs to mice having Unilateral Ureter Obstructed kidney.

Figure 4. Internalization of anti-ENDO180 mAbs conjugated to CypHer5E fluorophore into Myelo-Monocytoid human leukemia MonoMac cell line expressing ENDO180.

5 DETAILED DESCRIPTION OF THE INVENTION

Definitions

For convenience certain terms employed in the specification, examples and claims are described herein.

10 It is to be noted that, as used herein, the singular forms “a”, “an” and “the” include plural forms unless the content clearly dictates otherwise.

Where aspects or embodiments of the invention are described in terms of Markush groups or other grouping of alternatives, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the group.

15 An “inhibitor” is a compound, which is capable of reducing (partially or fully) the expression of a gene or the activity of the product of such gene to an extent sufficient to achieve a desired biological or physiological effect. The term "inhibitor" as used herein includes one or more of an oligonucleotide inhibitor, including siRNA, shRNA, synthetic shRNA; miRNA, antisense RNA and DNA and ribozymes. An “inhibitory
20 oligonucleotide” includes an antisense compound, a chemically modified siRNA compound, an unmodified siRNA compound, a chemically modified shRNA compound, an unmodified shRNA compound, a chemically modified miRNA compound, and an unmodified miRNA compound.

A “siRNA inhibitor” is a compound which is capable of reducing the expression of a gene
25 or the activity of the product of such gene to an extent sufficient to achieve a desired biological or physiological effect. The term "siRNA inhibitor" as used herein refers to one or more of a siRNA, shRNA, synthetic shRNA; miRNA. Inhibition may also be referred to as down-regulation or, for RNAi, silencing.

The term "inhibit" as used herein refers to reducing the expression of a gene or the activity
30 of the product of such gene to an extent sufficient to achieve a desired biological or

physiological effect. Inhibition may be complete or partial. As used herein, the term “ENDO180 gene ” is defined as any homolog of the ENDO180 gene having preferably 90% homology, more preferably 95% homology, and even more preferably 98% homology to the amino acid encoding region of SEQ ID NO:1 or nucleic acid sequences which bind to the ENDO180 gene under conditions of highly stringent hybridization, which are well-known in the art (for example, see Ausubel et al., Current Protocols in Molecular Biology, John Wiley and Sons, Baltimore, Maryland (1988), updated in 1995 and 1998).

As used herein, the term “ENDO180” or “ENDO180 polypeptide” or “ENDO180 receptor” is defined as any homolog of the ENDO180 polypeptide having preferably at least 90% homology, more preferably at least 95% homology, and even more preferably at least 98% homology or 100% identity to SEQ ID NO:2, as either full-length or a fragments or a domain thereof, as a mutant or the polypeptide encoded by a spliced variant nucleic acid sequence, as a chimera with other polypeptides, provided that any of the above has the same or substantially the same biological function as the ENDO180 receptor. ENDO180 polypeptide, or an ENDO180 polypeptide homolog, may be present in different forms, including but not limited to soluble protein, membrane-bound (either in purified membrane preparations or on a cell surface), bead-bound, or any other form presenting ENDO180 protein or fragments and polypeptides derived thereof. The term “inhibit” as used herein refers to reducing the expression of a gene or the activity of the product of such gene to an extent sufficient to achieve a desired biological or physiological effect. Inhibition is either complete or partial.

The terms “mRNA polynucleotide sequence”, “mRNA sequence” and “mRNA” are used interchangeably.

As used herein, the terms “polynucleotide” and “nucleic acid” may be used interchangeably and refer to nucleotide sequences comprising deoxyribonucleic acid (DNA), and ribonucleic acid (RNA). The terms are to be understood to include, as equivalents, analogs of either RNA or DNA made from nucleotide analogs. Throughout this application, mRNA sequences are set forth as representing the corresponding genes.

“Oligonucleotide” or “oligomer” refers to a deoxyribonucleotide or ribonucleotide sequence from about 2 to about 50 nucleotides. Each DNA or RNA nucleotide may be independently natural or synthetic, and or modified or unmodified. Modifications include changes to the sugar moiety, the base moiety and or the linkages between nucleotides in

the oligonucleotide. The compounds of the present invention encompass molecules comprising deoxyribonucleotides, ribonucleotides, modified deoxyribonucleotides, modified ribonucleotides and combinations thereof.

Substantially complementary refers to complementarity of greater than about 84%, to another sequence. For example in a duplex region consisting of 19 base pairs one mismatch results in 94.7% complementarity, two mismatches results in about 89.5% complementarity and 3 mismatches results in about 84.2% complementarity, rendering the duplex region substantially complementary. Accordingly substantially identical refers to identity of greater than about 84%, to another sequence.

10 The conjugate of the present invention comprises a) an antibody or fragment thereof, which specifically binds to an ENDO180 polypeptide on the surface of a cell, b) a nucleotide-based therapeutic agent selected from an antisense compound, a chemically modified siRNA compound, an unmodified siRNA compound, a chemically modified shRNA compound, an unmodified shRNA compound, a chemically modified miRNA
15 compound, and an unmodified miRNA compound; and c) a linking moiety which links (a) to (b); wherein the nucleotide-based therapeutic agent inhibits expression of the target gene in the cell.

The “linker” according to the present invention is a nucleotide or non-nucleotide moiety which links the antibody to the therapeutic molecule. In some embodiments the linker is a
20 cleavable moiety. Preferred cleavable groups include a disulfide bond, amide bond, thioamide, bond, ester bond, thioester bond, vicinal diol bond, or hemiacetal. Other cleavable bonds include enzymatically-cleavable bonds, such as peptide bonds (cleaved by peptidases), phosphate bonds (cleaved by phosphatases), nucleic acid bonds (cleaved by endonucleases), and sugar bonds (cleaved by glycosidases).

25 In some embodiments the linker is a non-nucleotide linker including a peptide linker. The choice of peptide sequence is critical to the success of the conjugate. In some embodiments the linker is stable to serum proteases, yet is cleaved by the lysosomal enzymes in the target cell. In a non-limiting example the linker is a peptide selected from a linker set forth in US 5574142, protamine, a fragment of protamine, (Arg)₉, biotin-avidin,
30 biotin-streptavidin and antennapedia peptide. For example, a peptide linker is used to link the antibody to a nucleotide therapeutic agent. Other non-nucleotide linkers include alkyl or aryl chains of about 5 to about 100 atoms.

In some embodiments the linker is a nucleotide linker. In certain embodiments a nucleic acid linker has a length ranging from 2-100, preferably 2-50 or 2-30 nucleotides.

Oligonucleotide Chemical Modifications

“Nucleotide” is meant to encompass deoxyribonucleotides and ribonucleotides, which
5 may be natural or synthetic, and or modified or unmodified. Modifications include changes to the sugar moiety, the base moiety and or the linkages between ribonucleotides in the oligoribonucleotide. As used herein, the term “ribonucleotide” encompasses natural and synthetic, unmodified and modified ribonucleotides. Modifications include changes to the sugar moiety, to the base moiety and/ or to the linkages between ribonucleotides in the
10 oligonucleotide.

The nucleotides useful in preparing a therapeutic agent include naturally occurring or synthetic modified bases. Naturally occurring bases include adenine, guanine, cytosine, thymine and uracil. Modified bases of nucleotides include inosine, xanthine, hypoxanthine, 2- aminoadenine, 6-methyl, 2-propyl and other alkyl adenines, 5-halo
15 uracil, 5-halo cytosine, 6-aza cytosine and 6-aza thymine, pseudo uracil, 4- thiouracil, 8-halo adenine, 8-aminoadenine, 8-thiol adenine, 8-thiolalkyl adenines, 8-hydroxyl adenine and other 8-substituted adenines, 8-halo guanines, 8-amino guanine, 8-thiol guanine, 8-thioalkyl guanines, 8- hydroxyl guanine and other substituted guanines, other aza and deaza adenines, other aza and deaza guanines, 5-trifluoromethyl uracil and 5- trifluoro
20 cytosine. In some embodiments one or more nucleotides in an oligomer is substituted with inosine.

According to some embodiments the present invention provides inhibitory oligonucleotide compounds comprising unmodified and modified nucleotides and or unconventional moieties. In certain embodiments the therapeutic agent is an oligonucleotide. In various
25 preferred embodiments the therapeutic agent is a double stranded oligonucleotide and preferably siRNA.

The selection and synthesis of siRNA corresponding to known genes has been widely reported; (see for example Ui-Tei et al., 2006. J Biomed Biotechnol.; 2006:65052; Chalk et al., 2004. BBRC. 319(1): 264-74; Sioud & Leirdal, 2004. Met. Mol Biol.; 252:457-69; Levenkova et al., 2004, Bioinform. 20(3):430-2; Ui-Tei et al., 2004. NAR 32(3):936-48).
30

For examples of the use of, and production of, modified siRNA see for example Braasch et al., 2003. Biochem., 42(26):7967-75; Chiu et al., 2003, RNA, 9(9):1034-48; PCT

publications WO 2004/015107 (atugen AG) and WO 02/44321 (Tuschl et al). US Patent Nos. 5,898,031 and 6,107,094 teach chemically modified oligomers. US Patent No. 7,452,987 relates to oligomeric compounds having alternating unmodified and 2' sugar modified ribonucleotides. US patent publication No. 2005/0042647 describes dsRNA compounds having chemically modified internucleoside linkages.

5 Amarzguoui et al., (2003, NAR, 31(2):589-595) showed that siRNA activity depended on the positioning of the 2'-O-methyl modifications. Holen et al (2003, NAR, 31(9):2401-2407) report that an siRNA having small numbers of 2'-O-methyl modified nucleosides showed good activity compared to wild type but that the activity decreased as the numbers

10 of 2'-O-methyl modified nucleosides was increased. Chiu and Rana (2003, RNA, 9:1034-1048) teach that incorporation of 2'-O-methyl modified nucleosides in the sense or antisense strand (fully modified strands) severely reduced siRNA activity relative to unmodified siRNA. The placement of a 2'-O-methyl group at the 5'-terminus on the antisense strand was reported to severely limit activity whereas placement at the 3'-15 terminus

15 of the antisense and at both termini of the sense strand was tolerated (Czauderna et al., 2003, NAR, 31(11), 2705-2716).

PCT Patent Application Nos. PCT/IL2008/000248 and PCT/IL2008/001197, assigned to the assignee of the present invention and disclose motifs useful in the preparation of chemically modified siRNA compounds. PCT

20 Patent Publication No. WO 2008/020435 discloses inhibitors, including some siRNA compounds to the target genes set forth herein.

The compound comprises at least one modified nucleotide selected from the group consisting of a sugar modification, a base modification and an internucleotide linkage modification and may contain DNA, and modified nucleotides such as LNA (locked

25 nucleic acid), ENA (ethylene-bridged nucleic acid), PNA (peptide nucleic acid), arabinoside, phosphonocarboxylate or phosphinocarboxylate nucleotide (PACE nucleotide), mirror nucleotide, or nucleotides with a 6 carbon sugar.

All analogs of, or modifications to, a nucleotide / oligonucleotide are employed with the present invention, provided that said analog or modification does not substantially

30 adversely affect the function of the nucleotide / oligonucleotide. Acceptable modifications include modifications of the sugar moiety, modifications of the base moiety, modifications in the internucleotide linkages and combinations thereof.

A sugar modification includes a modification on the 2' moiety of the sugar residue and encompasses amino, fluoro, alkoxy e.g. methoxy, alkyl, amino, fluoro, chloro, bromo, CN, CF, imidazole, carboxylate, thioate, C₁ to C₁₀ lower alkyl, substituted lower alkyl, alkaryl or aralkyl, OCF₃, OCN, O-, S-, or N- alkyl; O-, S, or N-alkenyl; SOCH₃; SO₂CH₃;
5 ONO₂; NO₂, N₃; heterocycloalkyl; heterocycloalkaryl; aminoalkylamino; polyalkylamino or substituted silyl, as, among others, described in European patents EP 0 586 520 B1 or EP 0 618 925 B1.

In one embodiment the siRNA compound comprises at least one ribonucleotide comprising a 2' modification on the sugar moiety ("2' sugar modification"). In certain
10 embodiments the compound comprises 2'O-alkyl or 2'-fluoro or 2'O-allyl or any other 2' modification, optionally on alternate positions. Other stabilizing modifications are also possible (e.g. terminal modifications). In some embodiments a preferred 2'O-alkyl is 2'O-methyl (methoxy) sugar modification.

In some embodiments the backbone of the oligonucleotides is modified and comprises
15 phosphate-D-ribose entities but may also contain thiophosphate-D-ribose entities, triester, thioate, 2'-5' bridged backbone (also may be referred to as 5'-2'), PACE and the like.

As used herein, the terms "non-pairing nucleotide analog" means a nucleotide analog which comprises a non-base pairing moiety including but not limited to: 6 des amino adenosine (Nebularine), 4-Me-indole, 3-nitropyrrole, 5-nitroindole, Ds, Pa, N3-Me ribo U,
20 N3-Me riboT, N3-Me dC, N3-Me-dT, N1-Me-dG, N1-Me-dA, N3-ethyl-dC, N3-Me dC. In some embodiments the non-base pairing nucleotide analog is a ribonucleotide. In other embodiments it is a deoxyribonucleotide. In addition, analogs of polynucleotides may be prepared wherein the structure of one or more nucleotide is fundamentally altered and better suited as therapeutic or experimental reagents. An example of a nucleotide analog is
25 a peptide nucleic acid (PNA) wherein the deoxyribose (or ribose) phosphate backbone in DNA (or RNA is replaced with a polyamide backbone which is similar to that found in peptides. PNA analogs have been shown to be resistant to enzymatic degradation and to have extended stability *in vivo* and *in vitro*. Other modifications that can be made to oligonucleotides include polymer backbones, cyclic backbones, acyclic backbones,
30 thiophosphate-D-ribose backbones, triester backbones, thioate backbones, 2'-5' bridged backbone, artificial nucleic acids, morpholino nucleic acids, glycol nucleic acid (GNA), threose nucleic acid (TNA), arabinoside, and mirror nucleoside (for example, beta-L-deoxyribonucleoside instead of beta-D-deoxyribonucleoside). Examples of siRNA

compounds comprising LNA nucleotides are disclosed in Elmen et al., (NAR 2005, 33(1):439-447).

The compounds of the present invention can be synthesized using one or more inverted nucleotides, for example inverted thymidine or inverted adenine (see, for example, Takei, 5 et al., 2002, JBC 277(26):23800-06).

Other modifications include terminal modifications on the 5' and/or 3' part of the oligonucleotides and are also known as capping moieties. Such terminal modifications are selected from a nucleotide, a modified nucleotide, a lipid, a peptide, a sugar and inverted abasic moiety.

10 What is sometimes referred to in the present invention as an "abasic nucleotide" or "abasic nucleotide analog" is more properly referred to as a pseudo-nucleotide or an unconventional moiety. A nucleotide is a monomeric unit of nucleic acid, consisting of a ribose or deoxyribose sugar, a phosphate, and a base (adenine, guanine, thymine, or cytosine in DNA; adenine, guanine, uracil, or cytosine in RNA). A modified nucleotide 15 comprises a modification in one or more of the sugar, phosphate and or base. The abasic pseudo-nucleotide lacks a base, and thus is not strictly a nucleotide.

In some embodiments the siRNA therapeutic agent comprises a capping moiety. The term "capping moiety" as used herein includes abasic ribose moiety, abasic deoxyribose moiety, modifications abasic ribose and abasic deoxyribose moieties including 2' O alkyl 20 modifications; inverted abasic ribose and abasic deoxyribose moieties and modifications thereof; C6-imino-Pi; a mirror nucleotide including L-DNA and L-RNA; 5'O-Me nucleotide; and nucleotide analogs including 4',5'-methylene nucleotide; 1-(β-D-erythrofuransyl)nucleotide; 4'-thio nucleotide, carbocyclic nucleotide; 5'-amino-alkyl phosphate; 1,3-diamino-2-propyl phosphate, 3-aminopropyl phosphate; 6-aminohexyl 25 phosphate; 12-aminododecyl phosphate; hydroxypropyl phosphate; 1,5-anhydrohexitol nucleotide; alpha-nucleotide; threo-pentofuransyl nucleotide; acyclic 3',4'-seco nucleotide; 3,4-dihydroxybutyl nucleotide; 3,5-dihydroxypentyl nucleotide, 5'-5'-inverted abasic moiety; 1,4-butanediol phosphate; 5'-amino; and bridging or non bridging methylphosphonate and 5'-mercapto moieties.

30 Certain preferred capping moieties are abasic ribose or abasic deoxyribose moieties; inverted abasic ribose or abasic deoxyribose moieties; C6-amino-Pi; a mirror nucleotide including L-DNA and L-RNA.

In some embodiments the therapeutic siRNA comprises a moiety other than a nucleotide. The term “unconventional moiety” as used herein refers to abasic ribose moiety, an abasic deoxyribose moiety, a deoxyribonucleotide, a modified deoxyribonucleotide, a mirror nucleotide, a non-base pairing nucleotide analog and a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide phosphate bond; bridged nucleic acids including LNA and ethylene bridged nucleic acids.

Abasic deoxyribose moiety includes for example abasic deoxyribose-3'-phosphate; 1,2-dideoxy-D-ribofuranose-3-phosphate; 1,4-anhydro-2-deoxy-D-ribitol-3-phosphate. Inverted abasic deoxyribose moiety includes inverted deoxyriboabasic; 3',5' inverted deoxyabasic 5'-phosphate.

A “mirror” nucleotide is a nucleotide with reversed chirality to the naturally occurring or commonly employed nucleotide, i.e., a mirror image (L-nucleotide) of the naturally occurring (D-nucleotide), also referred to as L-RNA in the case of a mirror ribonucleotide, and “spiegelmer”. The nucleotide can be a ribonucleotide or a deoxyribonucleotide and may further comprise at least one sugar, base and or backbone modification. See US Patent No. 6,586,238. Also, US Patent No. 6,602,858 discloses nucleic acid catalysts comprising at least one L-nucleotide substitution. Mirror nucleotide includes for example L-DNA (L-deoxyriboadenosine-3'-phosphate (mirror dA); L-deoxyribocytidine-3'-phosphate (mirror dC); L-deoxyriboguanosine-3'-phosphate (mirror dG); L-deoxyribothymidine-3'-phosphate (mirror image dT)) and L-RNA (L-riboadenosine-3'-phosphate (mirror rA); L-ribocytidine-3'-phosphate (mirror rC); L-riboguanosine-3'-phosphate (mirror rG); L-ribouracil-3'-phosphate (mirror dU)).

Modified deoxyribonucleotide includes, for example 5'OMe DNA (5-methyl-deoxyriboguanosine-3'-phosphate) which may be useful as a nucleotide in the 5' terminal position (position number 1); PACE (deoxyriboadenine 3' phosphonoacetate, deoxyribocytidine 3' phosphonoacetate, deoxyriboguanosine 3' phosphonoacetate, deoxyribothymidine 3' phosphonoacetate).

Bridged nucleic acids include LNA (2'-O,4'-C-methylene bridged Nucleic Acid adenosine 3' monophosphate, 2'-O,4'-C-methylene bridged Nucleic Acid 5-methyl-cytidine 3' monophosphate, 2'-O,4'-C-methylene bridged Nucleic Acid guanosine 3' monophosphate, 5-methyl-uridine (or thymidine) 3' monophosphate); and ENA (2'-O,4'-C-ethylene bridged Nucleic Acid adenosine 3' monophosphate, 2'-O,4'-C-ethylene bridged Nucleic Acid 5-

methyl-cytidine 3' monophosphate, 2'-O,4'-C-ethylene bridged Nucleic Acid guanosine 3' monophosphate, 5-methyl-uridine (or thymidine) 3' monophosphate).

In some embodiments of the present invention a preferred unconventional moiety is an abasic ribose moiety, an abasic deoxyribose moiety, a deoxyribonucleotide, a mirror
5 nucleotide, and a nucleotide joined to an adjacent nucleotide by a 2'-5' internucleotide phosphate bond.

According to one aspect the present invention provides inhibitory oligonucleotide compounds comprising unmodified and modified nucleotides. The compound comprises at least one modified nucleotide selected from the group consisting of a sugar modification, a
10 base modification and an internucleotide linkage modification and may contain DNA, and modified nucleotides such as LNA (locked nucleic acid) including ENA (ethylene-bridged nucleic acid; PNA (peptide nucleic acid); arabinoside; PACE (phosphonoacetate and derivatives thereof), mirror nucleotide, or nucleotides with a six-carbon sugar. In some embodiments the present invention provides methods and compositions for inhibiting
15 expression of a target gene *in vivo*. In general, the method includes administering a delivery -therapeutic agent conjugate. In particular embodiments small interfering RNAs (i.e. siRNAs), that target an mRNA transcribed from the target gene in an amount sufficient to down-regulate expression (reduce mRNA, reduce protein levels) of a target gene by an RNA interference mechanism. In particular, the subject method can be used to
20 inhibit expression of the target gene for treatment of a disease. In accordance with the present invention, the siRNA molecules or inhibitors of the target gene are used as drugs to treat various pathologies.

The synthesis of the nucleic acids described herein, is within the skills of the one of the art. Such synthesis is, among others, described in Beaucage SL and Iyer RP, 1992
25 Tetrahedron; 48: 2223-2311, Beaucage S. and Iyer RP, 1993 Tetrahedron; 49: 6123-6194 and Caruthers MH et. al., 1987 Methods Enzymol.; 154: 287-313, the synthesis of thioates is, among others, described in Eckstein F., 1985 Annu. Rev. Biochem.; 54: 367-402, the synthesis of RNA molecules is described in Sproat B., in Humana Press 2005 Edited by Herdewijn P.; Kap. 2: 17-31 and respective downstream processes are, among others,
30 described in Pingoud A. et. al., in IRL Press 1989 Edited by Oliver R.W.A.; Kap. 7: 183-208 and Sproat B., in Humana Press 2005 Edited by Herdewijn P.; Kap. 2: 17-31 (supra).

siRNA for any one of the target genes is synthesized using methods known in the art as described above, based on the known sequence of the target gene mRNA and is stabilized to serum and/or cellular nucleases by various modifications as described herein.

Target genes

- 5 The conjugates according to the present invention are useful for inhibiting expression of a gene associated with a disease or disorder selected from a proliferative disease a metastatic disease and fibrosis.

10 Target genes include anti-apoptotic genes, genes associated with basic cell division machinery, genes associated with cell cycle regulation/cell proliferation, genes associated with rate-limiting metabolism (nucleotide/nucleic acid synthesis, protein synthesis, energy metabolism), genes associated with protein trafficking (e.g., secretion); proinflammatory genes, cytokines, chemokines, NFkB, growth factors/receptors (TGFβ1 and 2, CTGF, IGF1, PDGF1, PDGF2, VEGF, EGFR, HER2, etc).

A non-limiting list of target genes is set forth in Table A, hereinbelow.

Abbreviation	full name
AARSD1	alanyl-tRNA synthetase domain containing 1
ABCF1	ATP-binding cassette, sub-family F (GCN20), member 1
AKT1	v-akt murine thymoma viral oncogene homolog 1
AKT2	-akt murine thymoma viral oncogene homolog 2
AKT3	v-akt murine thymoma viral oncogene homolog 3 (protein kinase B,
ANG	angiogenin, ribonuclease, RNase A family, 5
BAD	BCL2-associated agonist of cell death
BAG1	BCL2-associated athanogene
BAK1	BCL2-antagonist/killer 1
BAX	BCL2-associated X protein
BCL2	B-cell CLL/lymphoma 2
BCL2A1	BCL2-related protein A1
BCL2L1	BCL2-like 1
BCL2L11	BCL2-like 11 (apoptosis facilitator)
BID	BH3 interacting domain death agonist
CALR	calreticulin
CASP3	caspase 3, apoptosis-related cysteine peptidase
CASP9	caspase 9, apoptosis-related cysteine peptidase
CASP9	caspase 9, apoptosis-related cysteine peptidase
CCNB1	cyclin B1
CD40	CD40 molecule, TNF receptor superfamily member 5
CDC2	cell division cycle 2, G1 to S and G2 to M
CDC73	cell division cycle 73, Paf1/RNA polymerase II complex component,
CDH1	cadherin 1, type 1, E-cadherin (epithelial)
CEBPB	CCAAT/enhancer binding protein (C/EBP), beta

CFLAR	CASP8 and FADD-like apoptosis regulator
CHEK1	CHK1 checkpoint homolog (S. pombe)
CMPK1	cytidine monophosphate (UMP-CMP) kinase 1, cytosolic
COL4A1	collagen, type IV, alpha 1
CTGF	connective tissue growth factor
DDIT4	DNA-damage-inducible transcript 4
DDIT4L	DNA-damage-inducible transcript 4 like
EEF2K	eukaryotic elongation factor-2 kinase
EGF	epidermal growth factor
EIF2AK4	eukaryotic translation initiation factor 2 alpha kinase 4
EPRS	glutamyl-prolyl-tRNA synthetase
ERBB2	-erb-b2 erythroblastic leukemia viral oncogene homolog 2,
ERBB3	v-erb-b2 erythroblastic leukemia viral oncogene homolog 3
ESR1	estrogen receptor 1
F3	coagulation factor III
FAS	Fas (TNF receptor superfamily, member 6)
FEN1	flap structure-specific endonuclease 1
GAPDH	glyceraldehyde-3-phosphate dehydrogenase
H19	H19, imprinted maternally expressed transcript
HDAC1	histone deacetylase 1
HGF	hepatocyte growth factor (hepapoietin A; scatter factor)
HIF1A	hypoxia inducible factor 1, alpha subunit
HSF1	heat shock transcription factor 1
IER3	immediate early response 3
IGF1	insulin-like growth factor 1 (somatomedin C)
IGF1R	insulin-like growth factor 1 receptor
IGFBP5	insulin-like growth factor binding protein 5
IL15	interleukin 15
IL8	interleukin 8
JUN	jun oncogene
MADD	MAP-kinase activating death domain
MAPK1	mitogen-activated protein kinase 1
MCL1	myeloid cell leukemia
MDM2	Mdm2 p53 binding protein homolog (mouse)
MIF	macrophage migration inhibitory factor (glycosylation-inhibiting
MMP3	matrix metalloproteinase 3 (stromelysin 1, progelatinase)
MYC	v-myc myelocytomatosis viral oncogene homolog (avian)
c-MYC	myelocytomatosis viral oncogene homolog (avian)
NFKB1	nuclear factor of kappa light polypeptide gene enhancer in B-cells 1
NOS2	nitric oxide synthase 2, inducible
NOTCH1	Notch homolog 1, translocation-associated (Drosophila)
NOX1	NADPH oxidase 1
NOX2	cytochrome b-245, beta polypeptide (CYBB)
NOX3	NADPH oxidase 3
NOX4	NADPH oxidase 4
NOX5	NADPH oxidase 5
NOXA1	NADPH oxidase activator 1

NOXO1	NADPH oxidase organizer 1
NRF2	nuclear factor (erythroid-derived 2)-like 2
NR4A1	nuclear receptor subfamily 4, group A, member 1
OAS1	2',5'-oligoadenylate synthetase 1, 40/46kDa
OAS2	2'-5'-oligoadenylate synthetase 2, 69/71kDa
OAS3	2'-5'-oligoadenylate synthetase 3, 100kDa
ODC1	ornithine decarboxylase 1
PARP1	poly (ADP-ribose) polymerase 1
PCNA	proliferating cell nuclear antigen
PDGFA	platelet-derived growth factor alpha polypeptide
PIK3R1	phosphoinositide-3-kinase, regulatory subunit 1
PLAU	plasminogen activator, urokinase
PLK1	polo-like kinase 1
POLA1	polymerase (DNA directed), alpha 1, catalytic subunit
POLD1	polymerase (DNA directed), delta 1, catalytic subunit 125kDa
POLE	polymerase (DNA directed), epsilon
PPARD	peroxisome proliferator-activated receptor delta
PRKAR1A	protein kinase, cAMP-dependent, regulatory, type I, alpha (tissue
PRKDC	protein kinase, DNA-activated, catalytic polypeptide
PROK2	prokineticin 2
PTK2	PTK2 protein tyrosine kinase 2
PTK2B	PTK2B protein tyrosine kinase 2 beta
RAC1	ras-related C3 botulinum toxin substrate 1 (rho family, small GTP
RASSF1	Ras association (RalGDS/AF-6) domain family member 1
REG1A	regenerating islet-derived 1 alpha
RFC3	replication factor C (activator 1) 3, 38kDa
RHOA	ras homolog gene family, member A
RPA1	replication protein A1, 70kDa
SIPA1	signal-induced proliferation-associated 1
SOD1	superoxide dismutase 1, soluble
SRC	v-src sarcoma (Schmidt-Ruppin A-2) viral oncogene homolog (avian)
STAT3	signal transducer and activator of transcription 3
STAT6	signal transducer and activator of transcription 6, interleukin-4
TCF7L2	transcription factor 7-like 2
TEK	TEK tyrosine kinase, endothelial
TFAP2B	transcription factor AP-2 beta
TGFβ1	transforming growth factor, beta 1
TIAF1	TGFB1-induced anti-apoptotic factor 1
TIMP1	TIMP metalloproteinase inhibitor 1
TNF	tumor necrosis factor
TNFRSF1B	tumor necrosis factor receptor superfamily, member 1B
TP53	tumor protein p53
TRAF1	TNF receptor-associated factor 1
TYMS	thymidylate synthetase
VEGFA	vascular endothelial growth factor A
XIAP	X-linked inhibitor of apoptosis

Sense and antisense sequences useful in the synthesis of siRNA are selected according to proprietary and publicly available methods and algorithms.

The chemical modifications provided above are useful in synthesizing nucleotide therapeutics that exhibit inter alia, serum stability, activity, reduced immune response,
5 reduced off target effect.

Antibodies

The term "antibody" refers to IgG, IgM, IgD, IgA, and IgE antibody, inter alia. The definition includes polyclonal antibodies or monoclonal antibodies. This term refers to whole antibodies or fragments of antibodies comprising an antigen-binding domain, e.g.
10 antibodies without the Fc portion, single chain antibodies, miniantibodies, fragments consisting of essentially only the variable, antigen-binding domain of the antibody, etc. The term "antibody" may also refer to antibodies against polynucleotide sequences obtained by cDNA vaccination. The term also encompasses antibody fragments which retain the ability to selectively bind with their antigen or receptor and are exemplified as
15 follows, inter alia:

- (1) Fab, the fragment which contains a monovalent antigen-binding fragment of an antibody molecule which can be produced by digestion of whole antibody with the enzyme papain to yield a light chain and a portion of the heavy chain;
- (2) (Fab')₂, the fragment of the antibody that can be obtained by treating whole antibody
20 with the enzyme pepsin without subsequent reduction; F(ab')₂ is a dimer of two Fab fragments held together by two disulfide bonds;
- (3) Fv, defined as a genetically engineered fragment containing the variable region of the light chain and the variable region of the heavy chain expressed as two chains; and
- (4) Single chain antibody (SCA), defined as a genetically engineered molecule containing
25 the variable region of the light chain and the variable region of the heavy chain linked by a suitable polypeptide linker as a genetically fused single chain molecule, including a scFv.

CDR grafting may be performed to alter certain properties of the antibody molecule including affinity or specificity. A non-limiting example of CDR grafting is disclosed in US Patent No. 5,225,539.

30 Single-domain antibodies are isolated from the unique heavy-chain antibodies of immunized Camelidae, including camels and llamas. The small antibodies are very robust

and bind the antigen with high affinity in a monomeric state. US Patent 6838254 describes the production of antibodies or fragments thereof derived from heavy chain immunoglobulins of Camelidae.

5 A monoclonal antibody (mAb) is a substantially homogeneous population of antibodies to a specific antigen. Monoclonal antibodies (mAbs) are obtained by methods known to those skilled in the art. See, for example Kohler et al (1975); US patent 4,376,110; Ausubel et al (1987-1999); Harlow et al (1988); and Colligan et al (1993). The mAbs of the present invention
10 may be of any immunoglobulin class including IgG, IgM, IgE, IgA, and any subclass thereof. A hybridoma producing a mAb may be cultivated in vitro or in vivo. High titers of mAbs are obtained in vivo for example wherein cells from the individual hybridomas are injected intraperitoneally into pristine-primed Balb/c mice to produce ascites fluid containing high concentrations of the desired mAbs. mAbs of isotype IgM or IgG may be purified from such ascites fluid, or from culture supernatants, using column
15 chromatography methods well known to those of skill in the art.

By "specific binding affinity" is meant that the antibody binds to an ENDO180 polypeptide or fragment thereof with greater affinity than it binds to another polypeptide under similar conditions.

The term "epitope" is meant to refer to that portion of a molecule capable of being bound
20 by an antibody which can also be recognized by that antibody. An "antigen" is a molecule or a portion of a molecule capable of being bound by an antibody which is additionally capable of inducing an animal to produce antibody capable of binding to an epitope of that antigen. An antigen may have one or more than one epitope. The specific reaction referred to above is meant to indicate that the antigen will react, in a highly selective manner, with
25 its corresponding antibody and not with the multitude of other antibodies which may be evoked by other antigens.

Epitopes or antigenic determinants usually consist of chemically active surface groupings of molecules such as amino acids or sugar side chains and have specific three-dimensional structural characteristics as well as specific charge characteristics.

30 In one embodiment, the antibody is a monoclonal antibody. In one embodiment, the monoclonal antibody is an IgG, IgM, IgD, IgA, or IgE monoclonal antibody. IgG subclasses are also well known to those in the art and include but are not limited to human

IgG1, IgG2, IgG3 and IgG4. In one embodiment the monoclonal antibody is and IgG monoclonal antibody. In one embodiment, the monoclonal antibody is a human, humanized, or chimeric, antibody. In one embodiment, the portion of the antibody is a Fab fragment of the antibody. In one embodiment, the portion of the antibody comprises the
5 variable domain of the antibody. In one embodiment, the portion of the antibody comprises a CDR portion of the antibody. In other embodiments the antibody is a scFv molecule. The antibodies of the present invention may be produced recombinantly (see generally Marshak et al., 1996 "Strategies for Protein Purification and Characterization. A laboratory course manual." Plainview, N.Y.: Cold Spring Harbor Laboratory Press, 1996)
10 and analogs may be produced by post-translational processing. Differences in glycosylation can provide polypeptide analogs.

The antibody may be a human or nonhuman antibody. A nonhuman antibody may be humanized by recombinant methods to reduce its immunogenicity in man. Methods for humanizing antibodies are known to those skilled in the art.

15 This application provides humanized forms of the above antibodies. As used herein, "humanized" describes antibodies wherein some, most or all of the amino acids outside the CDR regions are replaced with corresponding amino acids derived from human immunoglobulin molecules, e.g. the human framework regions replace the non-human regions. In one embodiment of the humanized forms of the antibodies, some, most or all of
20 the amino acids outside the CDR regions have been replaced with amino acids from human immunoglobulin molecules but where some, most or all amino acids within one or more CDR regions remain unchanged. Small additions, deletions, insertions, substitutions or modifications of amino acids are permissible as long as they would not abrogate the ability of the antibody to bind the antigen, ENDO180.

25 A "humanized" antibody would retain a similar antigenic specificity as the original antibody, i.e. the ability to bind ENDO180, specifically human ENDO180 receptor and would similarly be internalized by the receptor.

One skilled in the art would know how to produce the humanized antibodies of the subject invention. Various publications describe how to make humanized antibodies.

For example, the methods described in U.S. Patent Nos. 4,816,567 and 6,331,415 comprise the production of chimeric antibodies having a variable region of one antibody and a constant region of another antibody.

U.S. Patent No. 5,225,539; 6,548,640 and 6,982,321 describes the use of recombinant DNA technology to produce a humanized antibody wherein the CDRs of one immunoglobulin are replaced with the CDRs from an immunoglobulin with a different specificity such that the humanized antibody would recognize the target antigen but would not illicit an immune response. Specifically, site directed mutagenesis is used to introduce the CDRs onto the framework region.

Other approaches for humanizing an antibody are described in WO 90/07861 and corresponding patents including U.S. Patent Nos. 5,585,089; 5,693,761; 6,180,370 and 7,022,500. These patents describe a method to increase the affinity of an antibody for the desired antigen by combining the CDRs of a mouse monoclonal antibody with human immunoglobulin framework and constant regions. Human framework regions can be chosen to maximize homology with the mouse sequence. Computer modeling can be used to identify amino acids in the framework region which are likely to interact with the CDRs or the specific antigen and then mouse amino acids can be used at these positions to create the humanized antibody.

The above methods are merely illustrative of some of the methods that one skilled in the art could employ to make humanized antibodies.

The monoclonal antibody E3-8D8 represents a suitable anti-ENDO180 antibody for use in the methods of the present invention. The hybridoma cell E3-8D8 was deposited with the Belgian Co-ordinated Collections of Micro-Organisms (BCCM), under the terms of the Budapest Treaty and given Accession Number LMBP 7203CB.

Epitope Mapping

Epitope mapping studies identify the residues that are important for antibody binding. Various methods are known in the art for epitope mapping and are readily performed by one skilled in the art. Certain methods are described in *Epitope Mapping: A Practical Approach* (O. M. R. Westwood, F. C. Hay; Oxford University Press, 2000).

One example of an epitope mapping techniques is Synthetic Labeled Peptides Epitope Mapping whereby a set of overlapping synthetic peptides is synthesized, each

corresponding to a small segment of the linear sequence of the protein antigen, i.e. extracellular domain of ENDO180, and arrayed on a solid phase. The panel of peptides is then probed with the test antibody, and bound antibody is detected using an enzyme-labeled secondary antibody.

- 5 Other techniques include fragmentation or cleavage and gel separation of the protein antigen, transfer to a membrane, probing by test antibody and bound antibody is detected using an enzyme-labeled secondary antibody.

Antibody drug development

In general monoclonal antibodies need to be designed to preserve binding properties
10 (selectivity, internalization etc) and to reduce an immune response in the recipient. Specifically, the monoclonal antibody secreted from hybridoma 3E-8D8 may be optimized for human therapeutics by one of several methods known to those with skill in the art. In one method the variable heavy chain (V_H) and variable light chain (V_L) of the monoclonal antibody are sequenced. Once the amino acid sequence is known, the complementarity
15 determining regions (CDR), heavy chain and light chain CDR3 are identified and degenerate oligonucleotides are used to clone synthetic CDR3 into a vector to produce a recombinant vector or construct. The construct may be for example a Fab fragment, a F(ab')₂ fragment, a Fv fragment, a single chain fragment or a full IgG molecule. The construct(s) is expressed and a polypeptide is isolated. In some embodiments the
20 monoclonal antibody may be further optimized by mutagenesis optimized by site directed mutagenesis to generate a CDR3 domain having substantial identity to the original CDR3.

Therapeutic Agents

The therapeutic agent or active agents according to the present invention includes nucleotide and non-nucleotide agents, including oligonucleotides such as antisense (AS),
25 miRNA and unmodified and chemically modified siRNA compounds. A preferred therapeutic agent is a siRNA compound.

In some embodiments the siRNA targets and reduces expression of a target gene by RNA interference.

The therapeutic oligonucleotides of the present invention are synthesized by any method
30 known in the art for ribonucleic or deoxyribonucleic nucleotides. For example, a commercial polynucleotide synthesizer (e.g. Applied Biosystems 380B DNA synthesizer) can be used. When fragments are used, two or more such sequences can be synthesized

and linked together for use in the present invention. Although a siRNA is the preferred therapeutic agent according to the present invention, the present invention encompasses a conjugate or mixture wherein the therapeutic agent is selected from alkylating agents such as thiotepa and CYTOXAN® cyclophosphamide; alkyl sulfonates such as busulfan, 5 improsulfan and piposulfan; aziridines such as benzodopa, carboquone, meturedopa, and uredopa; ethylenimines and methylamelamines including altretamine, triethylenemelamine, triethylenephosphoramidate, triethylenethiophosphoramidate and trimethylolmelamine; acetogenins (especially bullatacin and bullatacinone); delta-9-tetrahydrocannabinol (dronabinol, MARINOL®); beta-lapachone; lapachol; colchicines; 10 betulinic acid; a camptothecin (including the synthetic analog topotecan (HYCAMTIN®), CPT-11 (irinotecan, CAMPTOSAR®), acetylcamptothecin, scopolectin, and 9-aminocamptothecin); bryostatin; callistatin; CC-1065 (including its adozelesin, carzelesin and bizelesin synthetic analogs); podophyllotoxin; podophyllinic acid; teniposide; cryptophycins (particularly cryptophycin 1 and cryptophycin 8); dolastatin; duocarmycin 15 (including the synthetic analogs, KW-2189 and CB1-TM1); eleutherobin; pancratistatin; a sarcodictyin; spongistatin; nitrogen mustards such as chlorambucil, chlornaphazine, cholophosphamide, estramustine, ifosfamide, mechlorethamine, mechlorethamine oxide hydrochloride, melphalan, novembichin, phenesterine, prednimustine, trofosfamide, uracil mustard; nitrosureas such as carmustine, chlorozotocin, fotemustine, lomustine, nimustine, 20 and ranimustine; antibiotics such as the enediyne antibiotics (e. g., calicheamicin, especially calicheamicin gammaII and calicheamicin omegaII (see, e.g., Agnew, Chem Intl. Ed. Engl., 1994. 33: 183-186); dynemicin, including dynemicin A; an esperamicin; as well as neocarzinostatin chromophore and related chromoprotein enediyne antiobiotic chromophores), aclacinomysins, actinomycin, authramycin, azaserine, bleomycins, 25 cactinomycin, carabycin, carminomycin, carzinophilin, chromomycinis, dactinomycin, daunorubicin, detorubicin, 6-diazo-5-oxo-L-norleucine, doxorubicin (including ADRIAMYCIN®, morpholino-doxorubicin, cyanomorpholino-doxorubicin, 2-pyrrolino-doxorubicin, doxorubicin HCl liposome injection (DOXIL®) and deoxydoxorubicin), epirubicin, esorubicin, idarubicin, marcellomycin, mitomycins such as mitomycin C, 30 mycophenolic acid, nogalamycin, olivomycins, peplomycin, potfiromycin, puromycin, quelamycin, rodorubicin, streptonigrin, streptozocin, tubercidin, ubenimex, zinostatin, zorubicin; anti-metabolites such as methotrexate, gemcitabine (GEMZAR®), tegafur (UFTORAL®), capecitabine (XELODA®), an epothilone, and 5-fluorouracil (5-FU); folic acid analogs such as denopterin, methotrexate, pteropterin, trimetrexate; purine analogs

such as fludarabine, 6-mercaptopurine, thiamiprine, thioguanine; pyrimidine analogs such as ancitabine, azacitidine, 6-azauridine, carmofur, cytarabine, dideoxyuridine, doxifluridine, enocitabine, floxuridine; androgens such as calusterone, dromostanolone propionate, epitiostanol, mepitiothane, testolactone; anti-adrenals such as aminoglutethimide, mitotane, trilostane; folic acid replenisher such as frolinic acid; aceglatone; aldophosphamide glycoside; aminolevulinic acid; eniluracil; amsacrine; bestrabucil; bisantrene; edatraxate; defofamine; demecolcine; diaziquone; elfornithine; elliptinium acetate; etoglucid; gallium nitrate; hydroxyurea; lentinan; lonidainine; maytansinoids such as maytansine and ansamitocins; mitoguazone; mitoxantrone; mopidanmol; nitraerine; pentostatin; phenamet; pirarubicin; losoxantrone; 2-ethylhydrazide; procarbazine; PSK® polysaccharide complex; razoxane; rhizoxin; sizofiran; spirogermanium; tenuazonic acid; triaziqone; 2,2',2"-trichlorotriethylamine; trichothecenes (especially T-2 toxin, verracurin A, roridin A and anguidine); urethan; vindesine (ELDISINE®, FILDESIN®); dacarbazine; mannomustine; mitobronitol; mitolactol; pipobroman; gacytosine; arabinoside ("Ara-C"); thiotepa; taxoids, e.g., paclitaxel (TAXOL®), albumin-engineered nanoparticle formulation of paclitaxel (ABRAXANE.TM.), and doxetaxel (TAXOTERE®); chloranbucil; 6-thioguanine; mercaptopurine; methotrexate; a platinum analog such as cisplatin and carboplatin; vinblastine (VELBAN®); platinum; etoposide (VP-16); ifosfamide; mitoxantrone; vincristine (ONCOVIN®); oxaliplatin; leucovovin; vinorelbine (NAVELBINE®); novantrone; edatrexate; daunomycin; aminopterin; ibandronate; topoisomerase inhibitor RFS 2000; difluoromethylornithine (DMFO); a retinoid such as retinoic acid; pharmaceutically acceptable salts, acids or derivatives of any of the above; as well as combinations of two or more of the above such as CHOP, an abbreviation for a combined therapy of cyclophosphamide, doxorubicin, vincristine, and prednisolone, and FOLFOX, an abbreviation for a treatment regimen with oxaliplatin (ELOXATIN®) combined with 5-FU and leucovovin.

Also included in this definition are anti-hormonal agents that act to regulate, reduce, block, or inhibit the effects of hormones that can promote the growth of cancer, and are often administered as systemic, or whole-body treatment. They may be hormones themselves. Examples include anti-estrogens and selective estrogen receptor modulators (SERMs), including, for example, tamoxifen (including NOLVADEX® tamoxifen), raloxifene (EVISTA®), droloxifene, 4-hydroxytamoxifen, trioxifene, keoxifene,

LY117018, onapristone, and toremifene (FARESTON®); anti-progesterones; estrogen receptor down-regulators (ERDs); agents that function to suppress or shut down the ovaries, for example, leutinizing hormone-releasing hormone (LHRH) agonists such as leuprolide acetate (LUPRON® and ELIGARD®), goserelin acetate, buserelin acetate and triptorelin; other anti-androgens such as flutamide, nilutamide and bicalutamide; and aromatase inhibitors such as, for example, 4(5)-imidazoles, aminoglutethimide, megestrol acetate (MEGASE®), exemestane (AROMASIN®), formestane, fadrozole, vorozole (RIVISOR®), letrozole (FEMARA®), and anastrozole (ARIMIDEX®). In addition, bisphosphonates such as clodronate (for example, BONEFOS® or OSTAC®), etidronate (DIDROCAL®), NE-58095, zoledronic acid/zoledronate (ZOMETA®), alendronate (FOSAMAX®), pamidronate (AREDIA®), tiludronate (SKELID®), or risedronate (ACTONEL®); as well as troxacitabine (a 1,3-dioxolane nucleoside cytosine analog); siRNA, ribozyme and antisense oligonucleotides, particularly those that inhibit expression of genes in signaling pathways implicated in aberrant cell proliferation; vaccines such as THERATOPE® vaccine and gene therapy vaccines, for example, ALLOVECTIN® vaccine, LEUVECTIN® vaccine, and VAXID® vaccine; topoisomerase 1 inhibitor (e.g., LURTOTECAN®); rmRH (e.g., ABARELIX®); lapatinib ditosylate (an ErbB-2 and EGFR dual tyrosine kinase small-molecule inhibitor also known as GW572016); COX-2 inhibitors such as celecoxib (CELEBREX®; 4-(5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl) benzenesulfonamide; and pharmaceutically acceptable salts, acids or derivatives of any of the above.

As used herein, the term “polypeptide” refers to, in addition to a polypeptide, a peptide and a full protein and includes isolated and recombinant polypeptides. As used herein, “biological function” refers to the biological property of the molecule and in this context means an *in vivo* effector or antigenic function or activity that is directly or indirectly performed by a naturally occurring polypeptide or nucleic acid molecule. Biological functions include but are not limited to receptor binding, any enzymatic activity or enzyme modulatory activity, any carrier binding activity, any hormonal activity, any activity in internalizing molecules or translocation from one compartment to another, any activity in promoting or inhibiting adhesion of cells to extracellular matrix or cell surface molecules, or any structural role, as well as having the nucleic acid sequence encode functional protein and be expressible. The antigenic functions essentially mean the possession of an epitope or an antigenic site that is capable of cross-reacting with antibodies raised against

a naturally occurring protein. Biologically active analogs share an effector function of the native polypeptide that may, but need not, in addition possess an antigenic function.

Measurement of the level of the ENDO180 polypeptide may be determined by a method selected from the group consisting of immunohistochemistry, western blotting, ELISA, antibody microarray hybridization and targeted molecular imaging. Such methods are well-known in the art, for example immunohistochemistry, western blotting, antibody microarray hybridization, and targeted molecular imaging.

Measurement of the level of ENDO180 polynucleotide may be determined by a method selected for example from: RT-PCR analysis, *in-situ* hybridization, polynucleotide microarray and Northern blotting. Such methods are well known in the art.

Antisense molecules

In some embodiments the therapeutic agent is an antisense oligonucleotide. By the term "antisense" (AS) or "antisense fragment" is meant a polynucleotide fragment (comprising either deoxyribonucleotides, ribonucleotides or a mixture of both) having inhibitory antisense activity, said activity causing a decrease in the expression of the endogenous genomic copy of the corresponding gene. An AS polynucleotide is a polynucleotide which comprises consecutive nucleotides having a sequence of sufficient length and homology to a sequence present within the sequence of the target gene to permit hybridization of the AS to the gene. Many reviews have covered the main aspects of antisense (AS) technology and its therapeutic potential (Aboul-Fadl T., *Curr Med Chem.* 2005, 12(19):2193-214; Crooke ST, *Curr Mol Med.* 2004, 4(5):465-87; Crooke ST, *Ann Rev Med.* 2004, 55:61-95; Vacek M et al, *Cell Mol Life Sci.* 2003, 60(5):825-33; Cho-Chung YS, *Arch Pharm Res.* 2003, 26(3): 183-91. There are further reviews on the chemical (Crooke et al., *Hematol Pathol.* 1995, 9(2):59-72), cellular (Wagner, *Nature.* 1994, 372(6504):333-5) and therapeutic (Scanlon, et al, *FASEB J.* 1995, 9(13): 1288-96) aspects of AS technology. Antisense intervention in the expression of specific genes can be achieved by the use of modified AS oligonucleotide sequences (for recent reports see Lefebvre-d'Hellencourt et al, 1995; Agrawal, 1996; LevLehman et al, 1997).

AS oligonucleotide sequences may be short sequences of DNA, typically 15-30 mer but may be as small as 7-mer (Wagner et al, *Nat. Biotech.* 1996, 14(7):840-4), designed to complement a target mRNA of interest and form an RNA:AS duplex. This duplex formation can prevent processing, splicing, transport or translation of the relevant mRNA.

Moreover, certain AS nucleotide sequences can elicit cellular RNase H activity when hybridized with their target mRNA, resulting in mRNA degradation (Calabretta et al, Semin Oncol. 1996, 23(1):78-87). In that case, RNaseH will cleave the RNA component of the duplex and can potentially release the AS to further hybridize with additional
5 molecules of the target RNA. An additional mode of action results from the interaction of AS with genomic DNA to form a triple helix, which can be transcriptionally inactive.

The sequence target segment for the antisense oligonucleotide is selected such that the sequence exhibits suitable energy related characteristics important for oligonucleotide duplex formation with their complementary templates, and shows a low potential for self-
10 dimerization or self- complementation (Anazodo et al, 1996, BBRC. 229:305-309). For example, the computer program OLIGO (Primer Analysis Software, Version 3.4), can be used to determine antisense sequence melting temperature, free energy properties, and to estimate potential self-dimer formation and self-complimentary properties. The program allows the determination of a qualitative estimation of these two parameters (potential
15 self-dimer formation and self- complimentary) and provides an indication of "no potential" or "some potential" or "essentially complete potential". Using this program target segments are generally selected that have estimates of no potential in these parameters. However, segments can be used that have "some potential" in one of the categories. A balance of the parameters is used in the selection as is known in the art.
20 Further, the oligonucleotides are also selected as needed so that analog substitution does not substantially affect function.

Phosphorothioate antisense oligonucleotides do not normally show significant toxicity at concentrations that are effective and exhibit sufficient pharmacodynamic half-lives in animals (Agrawal, et al., PNAS U S A. 1997, 94(6):2620-5) and are nuclease resistant.
25 Antisense oligonucleotide inhibition of basic fibroblast growth factor (bFGF), having mitogenic and angiogenic properties, suppressed 80% of growth in glioma cells (Morrison, J Biol Chem. 1991 266(2):728-34) in a saturable and specific manner. Being hydrophobic, antisense oligonucleotides interact well with phospholipid membranes (Akhter et al., NAR. 1991, 19:5551-5559). Following their interaction with the cellular
30 plasma membrane, they are actively (or passively) transported into living cells (Loke et al., PNAS 1989, 86(10):3474-8), in a saturable mechanism predicted to involve specific receptors (Yakubov et al., PNAS, 1989 86(17):6454-58)

Ribozymes

A "ribozyme" is an RNA molecule that possesses RNA catalytic ability (see Cech for review) and cleaves a specific site in a target RNA. In accordance with the present invention, ribozymes which cleave mRNA may be utilized as a therapeutic agent. This
5 may be necessary in cases where antisense therapy is limited by stoichiometric considerations (Sarver et al., 1990, Gene Regulation and Aids, pp. 305-325). Ribozymes can then be used that will target the a gene associated with a bone marrow disease. The number of RNA molecules that are cleaved by a ribozyme is greater than the number predicted by stoichiometry. (Hampel and Tritz, Biochem. 1989, 28(12):4929-33;
10 Uhlenbeck, Nature. 1987.328(6131):596-600). Ribozymes catalyze the phosphodiester bond cleavage of RNA. Several ribozyme structural families have been identified including Group I nitrons, RNase P, the hepatitis delta virus ribozyme, hammerhead ribozymes and the hairpin ribozyme originally derived from the negative strand of the tobacco ringspot virus satellite RNA (sTRSV) (US Patent No. 5,225,347). The latter two
15 families are derived from viroids and virusoids, in which the ribozyme is believed to separate monomers from oligomers created during rolling circle replication (Symons, 1989 and 1992). Hammerhead and hairpin ribozyme motifs are most commonly adapted for trans-cleavage of mRNAs for gene therapy (Sullivan, 1994). In general the ribozyme has a length of from about 30- 100 nucleotides. Delivery of ribozymes is similar to that of
20 AS fragments and/or siRNA molecules.

siRNA and RNA interference

RNA interference (RNAi) is a phenomenon involving double-stranded (ds) RNA-dependent gene-specific posttranscriptional silencing. Initial attempts to study this
25 phenomenon and to manipulate mammalian cells experimentally were frustrated by an active, non-specific antiviral defense mechanism which was activated in response to long dsRNA molecules (Gil et al., Apoptosis, 2000. 5:107-114). Later, it was discovered that synthetic duplexes of 21 nucleotide RNAs could mediate gene specific RNAi in mammalian cells, without stimulating the generic antiviral defense mechanisms Elbashir et al. Nature 2001, 411:494-498 and Caplen et al. PNAS 2001, 98:9742-9747). As a result,
30 small interfering RNAs (siRNAs), which are short double-stranded RNAs, have been widely used to inhibit gene expression and understand gene function.

RNA interference (RNAi) is mediated by small interfering RNAs (siRNAs) (Fire et al, Nature 1998, 391:806) or microRNAs (miRNAs) (Ambros V. Nature 2004, 431:350-355);

and Bartel DP. Cell. 2004 116(2):281-97). The corresponding process is commonly referred to as specific post-transcriptional gene silencing when observed in plants and as quelling when observed in fungi.

A siRNA is a double-stranded RNA which down-regulates or silences (i.e. fully or partially inhibits) the expression of an endogenous or exogenous gene/ mRNA. RNA interference is based on the ability of certain dsRNA species to enter a specific protein complex, where they are then targeted to complementary cellular RNA (i.e. mRNA), which they specifically degrade or cleave. Thus, the RNA interference response features an endonuclease complex containing siRNA, commonly referred to as an RNA-induced silencing complex (RISC), which mediates cleavage of single-stranded RNA having a sequence complementary to the antisense strand of the siRNA duplex. Cleavage of the target RNA may take place in the middle of the region complementary to the antisense strand of the siRNA duplex (Elbashir, et al., Genes Dev., 2001, 15:188). In more detail, longer dsRNAs are digested into short (17-29 bp) dsRNA fragments (also referred to as short inhibitory RNAs or “siRNAs”) by type III RNAses (DICER, DROSHA, etc., (see Bernstein et al., Nature, 2001, 409:363-6 and Lee et al., Nature, 2003, 425:415-9). The RISC protein complex recognizes these fragments and complementary mRNA. The whole process is culminated by endonuclease cleavage of target mRNA (McManus and Sharp, Nature Rev Genet, 2002, 3:737-47; Paddison and Hannon, Curr Opin Mol Ther. 2003, 5(3): 217-24). (For additional information on these terms and proposed mechanisms, see for example, Bernstein, et al., RNA. 2001, 7(11):1509-21; Nishikura, Cell. 2001, 107(4):415-8 and PCT Publication No. WO 01/36646).

Studies have revealed that siRNA can be effective *in vivo* in mammals including humans. Specifically, Bitko et al., showed that specific siRNAs directed against the respiratory syncytial virus (RSV) nucleocapsid N gene are effective in treating mice when administered intranasally (Nat. Med. 2005, 11(1):50-55). For reviews of therapeutic applications of siRNAs see for example Barik (Mol. Med 2005, 83: 764-773) and Chakraborty (Current Drug Targets 2007 8(3):469-82). In addition, clinical studies with short siRNAs that target the VEGFR1 receptor in order to treat age-related macular degeneration (AMD) have been conducted in human patients (Kaiser, Am J Ophthalmol. 2006 142(4):660-8). Further information on the use of siRNA as therapeutic agents may be found in Durcan, 2008. Mol. Pharma. 5(4):559-566; Kim and Rossi, 2008. BioTechniques 44:613-616; Grimm and Kay, 2007, JCI, 117(12):3633-41.

The siRNA according to the invention is unmodified, recombinant or chemically modified. Examples of chemical modifications useful in synthesizing siRNA are disclosed in PCT Patent Publication No. WO 2009/044392, assigned to the assignee of the present invention.

5 Pharmaceutical Compositions

The present invention provides for a pharmaceutical composition comprising any one of the above compounds and a pharmaceutically acceptable excipient. In some embodiments the pharmaceutical composition according to the present invention comprises one of an anti-ENDO180 antibody or antigen-binding fragment thereof selected from

- 10 a) the monoclonal antibody produced by the hybridoma cell line E3-8D8 (BCCM Accession Number LMBP 7203CB);
- b) an antibody or fragment thereof that binds to the same epitope as the antibody of (a);
- c) a humanized antibody of (a) or (b);
- 15 d) a recombinant polypeptide comprising amino acid sequences set forth in SEQ ID NO: 6 or a variant thereof; and
- e) a recombinant polypeptide comprising CDR3 having amino acid sequences set forth in SEQ ID NO:7 and 8, or variants thereof;
- f) a conjugate of any one of the above (a)-(e) conjugated to a moiety;
- 20 wherein upon contact with a cell expressing ENDO180 the antibody or antigen binding fragment thereof is internalized into the cell; and
- a pharmaceutically acceptable vehicle or carrier.

In some embodiments the carrier comprises a lipid particle or a lipidated glycosaminoglycan.

- 25 In another aspect the invention provides compounds including a) an anti-ENDO180 antibody or antigen binding fragment thereof; b) a moiety selected from a detectable label, a cytotoxic agent or a therapeutic agent, and c) a nanocarrier.

In various embodiments the nanocarrier is a polysaccharide-based nanoparticle. In various embodiments, tripartite compounds of the invention can be represented by one of the

30 formulas:

A-X-Y

X-A-Y or

X Y A

5 wherein A represents a detectable label, a cytotoxic agent or a therapeutic agent;

X represents a nanocarrier; and

Y represents an anti-ENDO180 antibody or antigen-binding fragment thereof.

The disclosed compounds are designed to target particular cells or tissues expressing the ENDO180 polypeptide, so that a detectable label, a cytotoxic agent or a therapeutic agent
10 is delivered to the desired cell more effectively and with high specificity. For example, one embodiment of the disclosure includes compounds that target cancerous cell and/or tissues.

As such, certain examples of these compounds include a an anti-ENDO180 antibody or antigen binding fragment thereof that binds to an ENDO180 receptor that is present in a
15 higher concentration on a cancer cell than on a normal cell. Embodiments of the disclosed compounds exploit the up-regulated expression of ENDO180 receptors in diseased cells and tissue to selectively deliver a therapeutic agent to such a cell.

In various embodiments the nanocarrier is a polysaccharide-based nanoparticle. In certain embodiments the polysaccharide is a glycosaminoglycan or a mucopolysaccharide. In
20 various embodiments the glycosaminoglycan is selected from the group comprising, without being limited to, hyaluronic acid, chondroitin sulfate, dermatan sulfate, keratan sulfate, heparin, heparan sulfate and combinations thereof.

In certain embodiments the nanocarrier includes organic polymers, including, without being limited to, organic polymers that self assemble to form a self-assembled
25 nanoparticle, which provides an effectively multivalent species. In such embodiments the self-assembled nanoparticles can include the same or different compounds. For example, the self-assembled nanoparticles include compounds having an anti-ENDO180 antibody or antigen-binding fragment thereof; a moiety selected from a detectable label, a cytotoxic agent or a therapeutic agent; and the nanocarrier components.

Embodiments of the disclosed compounds include a plurality of therapeutic agents, detectable labels of cytotoxic agents. In such embodiments, the compounds include different therapeutic agents, imaging agents and cytotoxic agents. In certain embodiments compounds having two or more therapeutic agents have increased therapeutic efficiency
5 due to multivalent effects

The present invention further provides for a pharmaceutical composition comprising the disclosed compounds and conjugates, formulated for administration to a subject. An additional aspect of the present invention provides for methods of treating a subject having a proliferative disease including cancer, metastatic disease and fibrosis, using the
10 disclosed compounds, and hence pharmaceutical compositions are provided herein for this purpose.

In preferred embodiments the therapeutic agent is a chemically modified siRNA compound that inhibits expression of a target gene set forth in Table A.

In some embodiments the compositions comprise a mixture of two or more different
15 therapeutic agents including two or more siRNA that target a single gene or multiple genes.

The invention further provides a pharmaceutical composition comprising at least one compound of the invention covalently or non-covalently bound to one or more compounds of the invention in an amount effective to inhibit target gene expression or activity; and a
20 pharmaceutically acceptable carrier.

The pharmaceutically "effective amount" for purposes herein is thus determined by such considerations as are known in the art. The amount must be effective to achieve improvement including but not limited to improved survival rate or more rapid recovery, or improvement or elimination of symptoms and other indicators as are selected as
25 appropriate measures by those skilled in the art. The compounds of the present invention can be administered by any of the conventional routes of administration. It should be noted that the compound can be administered as the compound or as pharmaceutically acceptable salt and can be administered alone or as an active ingredient in combination with pharmaceutically acceptable carriers, solvents, diluents, excipients, adjuvants and
30 vehicles. The compounds can be administered orally, subcutaneously or parenterally including intravenous, intraarterial, intramuscular, intraperitoneally, and intranasal administration as well as intrathecal and infusion techniques. Implants of the compounds

are also useful. Liquid forms may be prepared for injection, the term including subcutaneous, transdermal, intravenous, intramuscular, intrathecal, and other parental routes of administration. The liquid compositions include aqueous solutions, with and without organic cosolvents, aqueous or oil suspensions, emulsions with edible oils, as well as similar pharmaceutical vehicles. In addition, under certain circumstances the compositions for use in the novel treatments of the present invention may be formed as aerosols, for intranasal and like administration. The patient being treated is a warm-blooded animal and, in particular, mammals including man. The pharmaceutically acceptable carriers, solvents, diluents, excipients, adjuvants and vehicles as well as implant carriers generally refer to inert, non-toxic solid or liquid fillers, diluents or encapsulating material not reacting with the active ingredients of the invention and they include liposomes, lipidated glycosaminoglycans and microspheres. Examples of delivery systems useful in the present invention include US Patent Nos. 5,225,182; 5,169,383; 5,167,616; 4,959,217; 4,925,678; 4,487,603; 4,486,194; 4,447,233; 4,447,224; 4,439,196; and 4,475,196. Many other such implants, delivery systems, and modules are well known to those skilled in the art.

In general, the active dose of compound for humans is in the range of from 1ng/kg to about 20-100 mg/kg body weight per day, preferably about 0.01 mg to about 2-10 mg/kg body weight per day, in a regimen of one dose per day or twice or three or more times per day for a period of 1-2 weeks or longer, preferably for 24-to 48 hrs or by continuous infusion during a period of 1-2 weeks or longer.

Additionally, the invention provides a method of inhibiting the expression of the genes of the present invention by at least 50% as compared to a control comprising contacting an mRNA transcript of the gene of the present invention with one or more of the compounds of the invention.

In one embodiment the therapeutic agent inhibits a target gene, whereby the inhibition is selected from the group comprising inhibition of gene function, inhibition of polypeptide and inhibition of mRNA expression.

The pharmaceutical composition is formulated to provide for a single dosage administration or a multi-dosage administration.

In various embodiments the pharmaceutical composition comprising a conjugate or mixture of the invention is administered intravenously, intramuscularly, locally, or subcutaneously to the subject.

5 The pharmaceutical composition according to the present invention can also be used in a method for preventing and/or treating a disease as disclosed herein, whereby the method comprises the administration of a conjugate according to the present invention, a mixture according to the present invention or a pharmaceutical composition or medicament according to the present invention for any of the diseases described herein.

Diagnostics

10 The compounds of the invention are useful in diagnosing ENDO180 expressing cells in biological samples.

Delivery

In some embodiments the antibodies, antigen-binding fragments and/or conjugates of the present invention are delivered to the target tissue by direct application of the naked
15 molecules prepared with a carrier or a diluent.

The term "naked molecule" refers to antibodies, antigen-binding fragments or conjugates that are free from any delivery vehicle that acts to assist, promote or facilitate entry into the cell, including viral sequences, viral particles, lipid particles, liposome formulations, lipofectin or precipitating agents and the like. For example, siRNA in PBS is "naked
20 siRNA". However, in some embodiments the antibodies, antigen-binding fragments or conjugates of the invention are delivered with lipid particles, polysaccharide particles or combinations thereof, liposome formulations, or lipofectin formulations and the like and can be prepared by methods well known to those skilled in the art. Such methods are described, for example, in US Patent Nos. 5,593,972, 5,589,466, and 5,580,859.

25 In other embodiments the antibodies, antigen-binding fragments or conjugates of the invention are attached to or are entrapped within a delivery particle. In some embodiments the ENDO180 binding domain of the conjugate molecule is exposed on the external surface of delivery particle. In some embodiments the delivery particle is a liposome. In specific preferred embodiment the delivery particle is a lipidated glycosaminoglycan
30 particle. Such particles are described, for example in US Patent Application Serial No. 10/487,022 (Publication No. 20040241248), US Patent Application Serial No. 11/718,485

(Publication No. 20080248092), US Patent Application Serial No. 11/ 632,647

(Publication No. 20090022656). Without wishing to be bound to theory, the antibody or antigen-binding fragment thereof is exposed on the surface of the delivery particle and homes in on or targets the target cell 5 expressing an ENDO180 polypeptide on its surface.

5 The pharmaceutically acceptable carriers, solvents, diluents, excipients, adjuvants and vehicles as well as implant carriers generally refer to inert, non-toxic solid or liquid fillers, diluents or encapsulating material not reacting with the active ingredients of the invention and they include liposomes and microspheres. Examples of delivery systems useful in the 10 present invention include US Patent Nos. 5,225,182; 5,169,383; 5,167,616; 4,959,217; 10 4,925,678; 4,487,603; 4,486,194; 4,447,233; 4,447,224; 4,439,196; and 4,475,196. Many other such implants, delivery systems, and modules are well known to those skilled in the art. In one specific embodiment of this invention topical and transdermal formulations may be selected. The siRNAs or pharmaceutical compositions of the present invention are 15 administered and dosed in accordance with good medical practice, taking into account the 15 clinical condition of the individual patient, the disease to be treated, the site and method of administration, scheduling of administration, patient age, sex, body weight and other factors known to medical practitioners.

The "therapeutically effective dose" for purposes herein is thus determined by such 20 considerations as are known in the art. The dose must be effective to achieve improvement 20 including but not limited to improved survival rate or more rapid recovery, or improvement or elimination of symptoms and other indicators as are selected as appropriate measures by those skilled in the art.

In general, the active dose of compound for humans is in the range of from 1ng/kg to 25 about 20-100 mg/kg body weight per day, preferably about 0.01 mg to about 2-10 mg/kg 25 body weight per day, in a regimen of one dose per day or twice or three or more times per day for a period of 1-4 weeks or longer.

The compounds of the present invention can be administered by any of the conventional routes of administration. It should be noted that the compound can be administered as the 30 compound or as pharmaceutically acceptable salt and can be administered alone or as an 30 active ingredient in combination with pharmaceutically acceptable carriers, solvents, diluents, excipients, adjuvants and vehicles. The compounds can be administered orally, subcutaneously or parenterally including intravenous, intraarterial, intramuscular,

intraperitoneally, and intranasal, inhalation, transtympanic administration as well as intrathecal and infusion techniques. Implants of the compounds are also useful. Liquid forms may be prepared for injection, the term including subcutaneous, transdermal, intravenous, intramuscular, intrathecal, intranasal and other parental routes of administration. The liquid compositions include aqueous solutions, with and without organic co-solvents, aqueous or oil suspensions, emulsions with edible oils, as well as similar pharmaceutical vehicles. In a particular embodiment, the administration comprises intravenous administration. In another embodiment the administration comprises topical or local administration. In addition, in certain embodiments the compositions for use in the novel treatments of the present invention may be formed as aerosols, for example for intranasal administration.

In certain embodiments, oral compositions (such as tablets, suspensions, solutions) may be effective for local delivery to the oral cavity such as oral composition suitable for mouthwash for the treatment of oral mucositis.

Liquid forms are prepared for drops or spray. The liquid compositions include aqueous solutions, with and without organic co-solvents, aqueous or oil suspensions, emulsions with oils, as well as similar pharmaceutical vehicles. In some embodiments administration comprises topical or local administration.

These compounds are administered to humans and other animals for therapy by any suitable route of administration to the eye, as by, for example, a spray or drops, and topically, as by ointments, suspensions or drops.

In preferred embodiments the subject being treated is a warm-blooded animal and, in particular, mammals including human.

Suitable methods for delivery of the compounds of present invention include, among others, systemic delivery, transfection, lipofection, and electroporation. In a further aspect the present invention is related to a pharmaceutical composition comprising a delivery molecule-therapeutic agent conjugate or an anti-ENDO180 antibody or anti-ENDO180 antibody-therapeutic agent mixture according to the present invention and, a pharmaceutically acceptable carrier, diluent or adjuvants or other vehicle(s).

Preferably, such carrier, diluents, adjuvants and vehicles are inert, and non-toxic. The pharmaceutical composition is in its various embodiments adapted for administration in various ways. Such administration comprises systemic and local administration as well as

oral, subcutaneous, parenteral, intravenous, intraarterial, intramuscular, intraperitoneal, intranasal, intrathecal, transtympanic and intraocular.

In some embodiments the vehicle is selected from a lipid particle, a polysaccharide particle or a combination thereof and a lipidated glycosaminoglycan particle (gagomer). In various embodiments the delivery molecule-therapeutic agent conjugate or antibody-therapeutic mixture is at least partially exposed on the external surface of the lipid particle or the lipidated glycosaminoglycan particle.

It will be acknowledged by the one skilled in the art that the amount of the pharmaceutical composition and the respective siRNA depends on the clinical condition of the individual patient, the site and method of administration, scheduling of administration, patient age, sex, bodyweight and other factors known to medical practitioners. The pharmaceutically effective amount for purposes of prevention and/or treatment is thus determined by such considerations as are known in the medical arts. Preferably, the amount is effective to achieve improvement including but limited to improve the diseased condition or to provide for a more rapid recovery, improvement or elimination of symptoms and other indicators as are selected as appropriate measures by those skilled in the medical arts.

Combination Therapy

In various embodiments the present invention relates to combination therapy. In one embodiment, the co-administration of two or more therapeutic agents achieves a synergistic effect, i.e., a therapeutic affect that is greater than the sum of the therapeutic effects of the individual components of the combination. In another embodiment, the co-administration of two or more therapeutic agents achieves an additive effect.

The active ingredients that comprise a combination therapy may be administered together via a single dosage form or by separate administration of each active agent. In certain embodiments, the first and second therapeutic agents are administered in a single dosage form. The agents may be formulated into a single tablet, pill, capsule, or solution for parenteral administration and the like. Alternatively, the first therapeutic agent and the second therapeutic agents may be administered as separate compositions. The first active agent may be administered at the same time as the second active agent or the first active agent may be administered intermittently with the second active agent. The length of time between administration of the first and second therapeutic agent may be adjusted to achieve the desired therapeutic effect. For example, the second therapeutic agent may be

administered only a few minutes (e.g., 1, 2, 5, 10, 30, or 60 min) or several hours (e.g., 2, 4, 6, 10, 12, 24, or 36 hr) after administration of the first therapeutic agent. In certain embodiments, it may be advantageous to administer more than one dosage of one of the therapeutic agents between administrations of the second therapeutic agent. For example, the second therapeutic agent may be administered at 2 hours and then again at 10 hours following administration of the first therapeutic agent. Alternatively, it may be advantageous to administer more than one dosage of the first therapeutic agent between administrations of the second therapeutic agent. Importantly, it is preferred that the therapeutic effects of each active ingredient overlap for at least a portion of the duration of each therapeutic agent so that the overall therapeutic effect of the combination therapy is attributable in part to the combined or synergistic effects of the combination therapy.

The present invention relates to compounds and the use of compounds, which down-regulate the expression of the genes of the invention particularly to conjugates comprising small interfering RNAs (siRNAs). Methods, molecules and compositions useful for inhibition of target genes are discussed herein at length, and any of said molecules and/or compositions may be beneficially employed in the treatment of a subject suffering from a proliferative or fibrotic disease.

Methods of Treatment

An additional aspect of the present invention provides for methods of treating a proliferative disease including cancer, metastatic disease and fibrosis. Methods for therapy of diseases or disorders associated with uncontrolled, pathological cell growth, e.g. cancer, psoriasis, autoimmune diseases, *inter alia*, and methods for therapy of diseases associated with ischemia and lack of proper blood flow, e.g. myocardial infarction (MI) and stroke, are provided. In particular, the compounds and compositions of the invention are useful in treating proliferative diseases in which ENDO180 is expressed in at least a portion of the diseased cells and or tissue.

“Cancer” or “Tumor” refers to an abnormal proliferation of cells. These terms include both primary tumors, which may be benign or malignant, as well as secondary tumors, or metastases which have spread to other sites in the body. Examples of proliferative diseases include, *inter alia*: carcinoma (e.g.: breast, colon and lung), leukemia such as B cell leukemia, lymphoma such as B-cell lymphoma, blastoma such as neuroblastoma and melanoma and sarcoma. It will be acknowledged that the pharmaceutical composition according to the present invention can be used for any disease which involves undesired

development or growth of vasculature including angiogenesis, as well as any of the diseases and conditions described herein.

The present invention provides methods and compositions for treating a patient suffering from a cancerous proliferative disease, (e.g. lung cancer, breast cancer, cervical cancer, colon cancer, gastric cancer, kidney cancer, leukemia, liver cancer, lymphoma, ovarian cancer, pancreatic cancer, prostate cancer, rectal cancer, sarcoma, skin cancer, testicular cancer, and uterine cancer) which the cancer cell expresses ENDO180 polypeptide. In one particular embodiment, the cancer is renal cancer including RCC and TCC.

“Cancer and "cancerous disease" are used interchangeably and refer to a disease that is caused by or results in inappropriately high levels of cell division, inappropriately low levels of apoptosis, or both. Examples of cancerous diseases include, without limitation, leukemias (e. g., acute leukemia, acute lymphocytic leukemia, acute myelocytic leukemia, acute myeloblastic leukemia, acute promyelocytic leukemia, acute myelomonocytic leukemia, acute monocytic leukemia, acute erythroleukemia, chronic leukemia, chronic myelocytic leukemia, chronic lymphocytic leukemia), polycythemia vera, lymphoma (Hodgkin's disease, non-Hodgkin's disease), Waldenstrom's macroglobulinemia, heavy chain disease, and solid tumors such as sarcomas and carcinomas (e.g., fibrosarcoma, myxosarcoma, liposarcoma, chondrosarcoma, osteogenic sarcoma, chordoma, angiosarcoma, endotheliosarcoma, lymphangio sarcoma, lymphangioendotheliosarcoma, synovioma, mesothelioma, Ewing's tumor, leiomyo sarcoma, rhabdomyosarcoma, colon carcinoma, pancreatic cancer, breast cancer, ovarian cancer, prostate cancer, squamous cell carcinoma, basal cell carcinoma, adenocarcinoma, sweat gland carcinoma, sebaceous gland carcinoma, papillary carcinoma, papillary adenocarcinomas, cystadenocarcinoma, medullary carcinoma, bronchogenic carcinoma, renal cell carcinoma, hepatoma, bile duct carcinoma, choriocarcinoma, seminoma, embryonal carcinoma, Wilm's tumor, cervical cancer, uterine cancer, testicular cancer, lung carcinoma, small cell lung carcinoma, bladder carcinoma, epithelial carcinoma, glioma, astrocytoma, medulloblastoma, craniopharyngioma, ependymoma, pinealoma, hemangioblastoma, acoustic neuroma, oligodendroglioma, schwannoma, meningioma, melanoma, neuroblastoma, and retinoblastoma). Metastases of a primary cancer is included. In some preferred embodiments the compounds of the present invention are useful in treating renal cancer, breast cancer, ovarian cancer and metastases thereof in various organs including lung and bone.

As used herein, the term "proliferative disease " refers to any disease in which cellular proliferation, either malignant or benign, contributes to the pathology of the condition. Such unwanted proliferation is the hallmark of cancer and many chronic inflammatory diseases, thus examples of "proliferative disease" include the cancers listed *supra* and
5 chronic inflammatory proliferative diseases such as psoriasis, inflammatory bowel disease and rheumatoid arthritis; proliferative cardiovascular diseases such as restenosis; proliferative ocular disorders such as diabetic retinopathy; and benign hyperproliferative diseases such as hemangiomas.

Fibrotic Disease

10 Fibrotic diseases are a group of chronic disease characterized by the excess production of a fibrous material called the extracellular matrix, which contributes to abnormal changes in tissue architecture and interferes with normal organ function. Millions of people worldwide suffer from these chronic diseases, that are often life threatening. Unfortunately, although fibrosis is widely prevalent, debilitating and often life threatening,
15 there is no effective treatment currently available.

The human body responds to trauma and injury by scarring. Fibrosis, a type of disorder characterized by excessive scarring, occurs when the normal wound healing response is disturbed. During fibrosis, the wound healing response continues causing an excessive production and deposition of collagen.

20 Although fibrotic disorders can be acute or chronic, the disorders share a common characteristic of excessive collagen accumulation and an associated loss of function when normal tissue is replaced with scar tissue.

Fibrosis results from diverse causes, and may be established in various organs. Cirrhosis, pulmonary fibrosis, sarcoidosis, keloids, hypertension and kidney fibrosis, are all chronic
25 diseases that induce a progressive fibrosis which causing a continuous loss of tissue function.

Acute fibrosis (usually with a sudden and severe onset and of short duration) occurs as a common response to various forms of trauma including accidental injuries (particularly injuries to the spine and central nervous system), infections, surgery (cardiac scarring
30 following heart attack), burns, environmental pollutants, alcohol and other types of toxins, acute respiratory distress syndrome, radiation and chemotherapy treatments. All tissues damaged by trauma are prone to scar and become fibrotic, particularly if the damage is

repeated. Deep organ fibrosis is often extremely serious because the progressive loss of organ function leads to morbidity, hospitalization, dialysis, disability and even death. Fibrotic diseases or diseases in which fibrosis is evident include pulmonary fibrosis, interstitial lung disease, human fibrotic lung disease, liver fibrosis, cardiac fibrosis, macular degeneration, retinal and vitreal retinopathy, myocardial fibrosis, Grave's ophthalmopathy, drug induced ergotism, cardiovascular disease, atherosclerosis / restenosis, keloids and hypertrophic scars, Hansen's disease and inflammatory bowel disease, including collagenous colitis.

Further information on different types of fibrosis may be found for example in Yu et al (2002), "Therapeutic strategies to halt renal fibrosis", *Curr Opin Pharmacol.* 2(2):177-81; Keane and Lyle (2003), "Recent advances in management of type 2 diabetes and nephropathy: lessons from the RENAAL study", *Am J Kidney Dis.* 41(3 Suppl 2): S22-5; Bohle et al (1989), "The pathogenesis of chronic renal failure", *Pathol Res Pract.* 185(4):421-40; Kikkawa et al (1997), "Mechanism of the progression of diabetic nephropathy to renal failure", *Kidney Int Suppl.* 62:S39-40; Bataller and Brenner (2001), "Hepatic stellate cells as a target for the treatment of liver fibrosis", *Semin Liver Dis.* 21(3):437-51; Gross and Hunninghake (2001) "Idiopathic pulmonary fibrosis", *N Engl J Med.* 345(7):517-25; Frohlich (2001) "Fibrosis and ischemia: the real risks in hypertensive heart disease", *Am J Hypertens*;14(6 Pt 2):194S-199S.

20 Diabetic nephropathy

Diabetic nephropathy, hallmarks of which are glomerulosclerosis and kidney fibrosis, is the single most prevalent cause of end-stage renal disease in the modern world, and diabetic patients constitute the largest population on dialysis. Such therapy is costly and far from optimal. Transplantation offers a better outcome but suffers from a severe shortage of donors. More targeted therapies against diabetic nephropathy (as well as against other types of kidney pathologies) are not developed, since molecular mechanisms underlying these pathologies are largely unknown. Identification of an essential functional target gene that is modulated in the disease and affects the severity of the outcome of diabetes nephropathy has a high diagnostic as well as therapeutic value.

30 It is known in the art that many pathological processes in the kidney eventually culminate in similar or identical morphological changes, namely glomerulosclerosis and fibrosis. Human kidney disease may evolve from various origins including glomerular nephritis, nephritis associated with systemic lupus, cancer, physical obstructions, toxins, metabolic

disease and immunological diseases, all of which culminate in kidney fibrosis. The meaning of this phenomenon is that different types of insults converge on the same single genetic program resulting in two hallmarks of fibrosis: the proliferation of fibroblasts and overproduction by them of various protein components of connective tissue. In addition, thickening of the basal membrane in the glomeruli accompanies interstitial fibrosis and culminates in glomerulosclerosis. Genes encoding proteins that are involved in kidney fibrosis and glomerulosclerosis may be roughly divided into two groups:

1. Genes, the expression of which leads to the triggering of proliferation of fibroblasts and overproduction by them of various protein components of connective tissue. These may be specific to different pathological conditions; and
2. Genes, the expression of which leads to the execution of the “fibrotic or sclerotic programs”. These may be common to all renal pathologies leading to fibrosis and glomerulosclerosis.

The identification of genes that belong to the second group should contribute to the understanding of molecular mechanisms that accompany fibroblast and mesangial cell proliferation and hypersecretion, and may constitute genetic targets for drug development, aimed at preventing renal failure. Application of such drugs is expected to suppress, retard, prevent, inhibit or attenuate progression of fibrosis and glomerulosclerosis.

Combination therapy

The present invention provides for combination therapy for proliferative disease as disclosed herein and in particular cancer. In said combination therapy, one or more genes are targeted to ameliorate symptoms of the disease being treated. These genes are inhibited the antibody-nucleotide complex of the present invention.

Kits

Kits comprising at least one anti-ENDO180 monoclonal antibody of the invention are further provided. A "kit" refers to any manufacture (e.g., a package or a container) comprising at least one reagent, i.e., an antibody, for specific binding to ENDO180. The kit may be used for performing the methods of the present invention, including therapeutic treatment and diagnostics. Additionally, the kit may contain a package insert describing the kit, its content and methods for use.

In one embodiment, a kit of the invention comprises at least composition comprising an anti-ENDO180 antibody or antigen binding fragment thereof selected from

a) the monoclonal antibody produced by the hybridoma cell line E3-8D8 (BCCM Accession Number LMBP 7203CB);

5 b) an antibody or fragment thereof that binds to the same epitope as the antibody in (a);

c) a fragment of an antibody comprising a polypeptide substantially similar to SEQ ID NO: 6; and

10 d) a recombinant polypeptide comprising CDRs having an amino acid sequence substantially similar to amino acid sequences set forth in SEQ ID NO:7 and 8.

The invention has been described in an illustrative manner, and it is to be understood that the terminology which has been used is intended to be in the nature of words of description rather than of limitation.

15 Citation of any document herein is not intended as an admission that such document is pertinent prior art, or considered material to the patentability of any claim of the present application. Any statement as to content or a date of any document is based on the information available to applicant at the time of filing and does not constitute an admission as to the correctness of such a statement.

EXAMPLES

20 General methods in molecular biology

Standard molecular biology techniques known in the art and not specifically described were generally followed as in Sambrook et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory Press, New York (1989), and in Ausubel et al., *Current Protocols in Molecular Biology*, John Wiley and Sons, Baltimore, Maryland (1989) and in
 25 Perbal, *A Practical Guide to Molecular Cloning*, John Wiley & Sons, New York (1988), and in Watson et al., *Recombinant DNA*, Scientific American Books, New York and in Birren et al (eds) *Genome Analysis: A Laboratory Manual Series*, Vols. 1-4 Cold Spring Harbor Laboratory Press, New York (1998) and methodology as set forth in United States patents 4,666,828; 4,683,202; 4,801,531; 5,192,659 and 5,272,057. Polymerase chain reaction
 30 (PCR) was carried out generally as in PCR

Protocols: A Guide To Methods And Applications, Academic Press, San Diego, CA (1990). In situ (In cell) PCR in combination with Flow Cytometry can be used for detection of cells containing specific DNA and mRNA sequences (Testoni et al., 1996, Blood 87:3822.)

- 5 General methods in immunology: Standard methods in immunology known in the art and not specifically described are generally followed as in Stites et al (eds), Basic and Clinical Immunology (8th Edition), Appleton & Lange, Norwalk, CT (1994) and Mishell and Shiigi (eds), Selected Methods in Cellular Immunology, W.H. Freeman and Co., New York (1980).
- 10 Immunoassays: ELISA immunoassays are well known to those skilled in the art. Both polyclonal and monoclonal antibodies can be used in the assays. Where appropriate, other immunoassays such as radioimmunoassays (RIA) can be used as are known to those skilled in the art. Available immunoassays are extensively described in the patent and scientific literature. See, for example, United States Patent Nos. 3,791,932; 3,839,153;
- 15 3,850,752; 3,850,578; 3,853,987; 3,867,517; 3,879,262; 3,901,654; 3,935,074; 3,984,533; 3,996,345; 4,034,074; 4,098,876; 4,879,219; 5,011,771 and 5,281,521 as well as Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Springs Harbor, New York, 1989.

Antibody Production

- 20 By the term “antibody” as used in the present invention is meant both polyclonal and monoclonal complete antibodies as well as fragments thereof, such as Fab, F(ab')₂, scFv and Fv, which are capable of binding the epitope determinant. These antibody fragments retain the ability to selectively bind with its antigen or receptor and are exemplified as follows, inter alia:
- 25 A Fab, the fragment which contains a monovalent antigen-binding fragment of an antibody molecule can be produced by digestion of whole antibody with the enzyme papain to yield a light chain and a portion of the heavy chain;
- A (Fab')₂, the fragment of the antibody that can be obtained by treating whole antibody with the enzyme pepsin without subsequent reduction; F(ab')₂ is a dimer of two Fab
- 30 fragments held together by two disulfide bonds;
- A Fv, defined as a genetically engineered fragment containing the variable region of the light chain and the variable region of the heavy chain expressed as two chains; and

A scFv fragment (i.e. a single chain antibody ("SCA")), defined as a genetically engineered molecule containing the variable region of the light chain and the variable region of the heavy chain linked by a suitable polypeptide linker as a genetically fused single chain molecule.

5 Such fragments having antibody functional activity can be prepared by methods known to those skilled in the art (Bird et al. (1988) Science 242:423-426)

Conveniently, antibodies may be prepared against an immunogen or portion thereof, for example, a synthetic peptide based on the sequence, or prepared recombinantly by cloning techniques or the natural gene product and/or portions thereof may be isolated and used as
10 the immunogen. Immunogens can be used to produce antibodies by standard antibody production technology well known to those skilled in the art, as described generally in Harlow and Lane (1988), Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY, and Borrebaeck (1992), Antibody Engineering - A Practical Guide, W.H. Freeman and Co., NY.

15 For producing polyclonal antibodies a host, such as a rabbit or goat, is immunized with the immunogen or immunogen fragment, generally with an adjuvant and, if necessary, coupled to a carrier; antibodies to the immunogen are collected from the sera. Further, the polyclonal antibody can be absorbed such that it is monospecific; that is, the sera can be absorbed against related immunogens so that no cross-reactive antibodies remain in the
20 sera, rendering it monospecific.

For producing monoclonal antibodies the technique involves hyperimmunization of an appropriate donor with the immunogen, generally a mouse, and isolation of splenic antibody-producing cells. These cells are fused to an immortal cell, such as a myeloma cell, to provide a fused cell hybrid that is immortal and secretes the required antibody.
25 The cells are then cultured, in bulk, and the monoclonal antibodies harvested from the culture media for use.

For producing recombinant antibody see generally Huston et al. (1991) "Protein engineering of single-chain Fv analogs and fusion proteins" in Methods in Enzymology (JJ Langone, ed., Academic Press, New York, NY) 203:46-88; Johnson and Bird (1991)
30 "Construction of single-chain Fvb derivatives of monoclonal antibodies and their production in Escherichia coli in Methods in Enzymology (JJ Langone, ed.; Academic Press, New York, NY) 203:88-99; Mernaugh and Mernaugh (1995) "An overview of

phage-displayed recombinant antibodies" in *Molecular Methods In Plant Pathology* (RP Singh and US Singh, eds.; CRC Press Inc., Boca Raton, FL:359-365). Additionally, messenger RNAs from antibody-producing B-lymphocytes of animals, or hybridoma can be reverse-transcribed to obtain complementary DNAs (cDNAs). Antibody cDNA, which
5 can be full or partial length, is amplified and cloned into a phage or a plasmid. The cDNA can be a partial length of heavy and light chain cDNA, separated or connected by a linker. The antibody, or antibody fragment, is expressed using a suitable expression system to obtain recombinant antibody. Antibody cDNA can also be obtained by screening pertinent expression libraries.

10 The antibody can be bound to a solid support substrate or conjugated with a detectable moiety or be both bound and conjugated as is well known in the art. (For a general discussion of conjugation of fluorescent or enzymatic moieties see Johnstone & Thorpe (1982.), *Immunochemistry in Practice*, Blackwell Scientific Publications, Oxford). The binding of antibodies to a solid support substrate is also well known in the art (for a
15 general discussion, see Harlow & Lane (1988) *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory Publications, New York; and Borrebaeck (1992), *Antibody Engineering - A Practical Guide*, W.H. Freeman and Co.). The detectable moieties or label contemplated with the present invention include, but are not limited to, fluorescent, metallic, enzymatic and radioactive markers such as biotin, gold, ferritin, alkaline
20 phosphatase, β -galactosidase, peroxidase, urease, fluorescein, rhodamine, tritium, ^{14}C and iodine.

Recombinant Protein Purification

For standard purification, see Marshak et al. (1996), "Strategies for Protein Purification and Characterization. A laboratory course manual." CSHL Press.

25 The polynucleotide sequence of human ENDO180 mRNA is set forth in accession number NM_006039: 5641 bases, of that the open reading frame (ORF) is 4439 bases (from 117-4441); the polypeptide sequence of 1479 amino acids (aa) is set forth in accession number NP_006030 with gene identifier number: GI:110624774. The mouse mRNA sequence is 5818 bases, accession number MMU56734 with ORF of 1479 aa.

30 ENDO180 comprises several protein domains, as follows: 1-31 aa SP (signal peptide); 41-161 aa cysteine rich N-terminal domain; 180-228 aa FNII (fibronectin type II) domain; 8 CDR (carbohydrate recognition domain) domains 1CRD-8CRD (235-360 aa 1CRD, 382-

521-645 aa 3CRD, 669-809 aa 4CRD, 825-951 aa 5CRD, 972-1108 aa 6CRD, 1161-1244 aa 7CRD, 1261-1394 aa 8CRD); 1413-1435 aa 1 TM (transmembrane domain), 1437-1479 aa-cytoplasmic domain.

EXAMPLE 1 : Identification of ENDO180 overexpression by microarray hybridization study

In accordance with the present invention, the microarray hybridization approach was utilized in order to discover genes that are differentially regulated in diabetic nephropathy and kidney fibrosis.

Microarray-based analysis of gene expression was based on the analysis of human fibroblasts subject to selected stimuli resulting in changes in extracellular collagen accumulation and proliferation - the hallmarks of fibrosis. According to the present invention, a specific "Fibrosis" DNA chip was first prepared followed by a microarray hybridization experiments with 19 different types of probes. Analysis of the results was carried out by proprietary algorithms, and analysis of the selected set of genes was performed by the inventors using bioinformatics and the scientific literature.

EXAMPLE 2: Production of Human anti-ENDO180 Antibodies

The aim was to generate anti-ENDO180 antibodies that bind to the extracellular portion of ENDO180 and internalize an anti-ENDO180 antibody-cargo complex/conjugate.

Antigen production: The structural considerations in selecting an antigen for antibody production included the information that amino acids 1-522 (SEQ ID NO:9) spatially create the ligand binding structure.

Recombinant ENDO180 antigen was produced by cloning nucleotides 1-1566 of human ENDO180 polynucleotide into an expression vector comprising the FLAG epitope. The polynucleotide sequence of the recombinant clone is set forth in SEQ ID NO:9, the amino acid sequence is set forth in SEQ ID NO:4. The vectors were transfected into 293T cells and a clones expressing the 59KD partial extracellular domain of human ENDO180 were

identified. The ECDhENDO180-FLAG protein was isolated as follows: about 2.2 liters of conditioned medium were filtered through a 0.22um filter. Medium was loaded on a pre-equilibrated (with TBS) M2 agarose (5ml, Sigma) at a flow rate of 1ml/min. Resin was washed with 10 column volumes using TBS and then 10 volumes with 50 mM Tris pH 7.5, 1M NaCl and finally with 10 volumes of TBS. Elution was done with 10 ml of 0.5mg/ml FLAG peptide in TBS pH 7.5 (final pH). Resin was incubated for 20 min with elution buffer before starting the flow out of column. Sample was concentrated and depleted of FLAG peptide using VivaSpin™ (cut-off 10Kd). Glycerol was added to 10% final and protein was flash frozen in liquid nitrogen.

10 Identification of minibodies: minibody antibodies were identified according to the methods disclosed in Di Niro et al, 2007. Construction of miniantibodies for the in vivo study of human autoimmune diseases in animal models. BMC Biotechnology 7:46.

Certain preferred antibodies according to the present invention are recombinant polypeptides comprising heavy chain and light chain CDR3 domains having amino acid sequences set forth in SEQ ID NO:7 and in SEQ ID NO:8.

EXAMPLE 3: Anti-ENDO180 monoclonal antibodies

Methods and summary:

1. Labeling of mAbs and MB with CypHer5e was performed according to manufacturer's directions. (GE Healthcare).
- 20 2. Labeled vs. unlabeled mAbs and MB were tested for binding to purified ENDO180 extra-cellular domain by standard ELISA.

Clones 6D6, 8D8, 8E7 and 9G10 displayed saturated binding to endo180DCTLD3-8-FLAG even after labeling with CypHer5E. In contrast, binding of clone 8D2 and MB (minibody) was significantly impaired upon labeling.

3. Internalization assays were performed according to methods to those skilled in the art.

After 1 hr at 37°C, ENDO180 expressing cells that were incubated with labeled mAbs 6D6, 8D8, 8E7 and 9G10, displayed some increase in fluorescence. Most notably, the same cells that were incubated with labeled MB, showed strong fluorescence. This increase in fluorescence was not seen

in control cells or in ENDO180 cells at 40C. In addition, control mAb (10F12) had no effect

4. Kinetics experiments

Based on the results of previous experiments, mAb 6D6, 8D8 and MB were tested. In the 10 min-1 hr kinetics experiment, mAb 8D8 and MB showed the best internalization. All negative controls, control cells, (4°C; shown as 40C in some figures) and control mAb-were negative.

Details of Experiments and Results

Anti-ENDO180 monoclonal antibodies (mAb) were generated against the most N-terminal domain (1-522 aa) of human ENDO180 (SEQ ID NO:9) and were screened for internalization in an ENDO180-specific manner per se and conjugated to the fluorophore, CypHer5E.

mAbs production: About 8 liters of each hybridoma were grown in DMEM medium supplemented with 5% FBS IgG FREE, 1% Penstrep, 1% L-Glutamine, 50µg/ml Gentamycin, 2.5µg/ml Amphotricin B (Fungizone). About 60 ml was obtained from cell line flasks, with an antibody concentration of about 200µg/ml. The duration of the growth was 1 month. The purification was done using Protein A Sepharose column followed by two cycles of sizing column.

The mAb 8D8 (E3-8D8), 6D6, 8E7, 9G10, 8D2 were selected for conjugation to labeling moieties.

The mAbs were covalently linked to CypHer5E (Cat# pA15405, Amersham) a red excited fluorescent pH sensitive cyanine dye according to manufacturer's instructions in a molar ratio of 20:1 CypHer5E:Antibody. The fluorophore is excited in acidic pH, as found within the endosome. Therefore, those antibodies that get internalized will be seen as fluorescent signals in cellular vesicles. The mAbs were covalently linked to biotin using EZ-Sulfo-NHS-biotin (Pierce, Cat # 21217, Lot # CE49927). Biotinylation was performed according to manufacturer's instructions using a molar ratio of 20:1 Biotin:Antibody at room temperature for 30 min. Following covalently binding of mAbs to CypHer5E or biotin, the antibody solution was dialyzed overnight at 4°C against solution of PBS following additional two hours at room temperature against a fresh PBS

solution. Labeling of mAbs and scFv (SEQ ID NO. 6, also referred to as “minibody”) with CypHer5E was performed according to manufacture’s instructions (GE Healthcare).

Internalization of the conjugated receptor was performed in the absence of collagen (ligand independent internalization). The following clones were shown to secrete
5 antibodies that bind specifically to ENDO180: Clones 6D6, 8D8, 8E7 and 9G10 displayed saturated binding to endo180DCTLD3-8-FLAG per se and when labeled with CypHer5E. Clone 8D2 showed limited uptake into cells following labeling while clone 10F12 exhibited significant and saturated binding to endo180DCTLD3-8-FLAG which was diminished after labeling.

10 Cells expressing ENDO180 (NRK52E-ENDO180) and control cells (NRK52E) were incubated at 37 °C with the indicated anti-ENDO180 mAbs or control mAbs, covalently linked to CypHer5E. The mAbs were also incubated at 4°C with ENDO180 cells. The cells were plated in a 96-well plate and fluorescence was measured by Analyst AD &HT, Biosystems (excitation 530nm, emission 590nm, dichroic 560nm). A steady increase in
15 fluorescence at 37°C in ENDO180 cells was seen with one mAb (E3-8D8). In contrast, fluorescence was not seen in control cells, at 4°C or with control mAbs. Antibody binding and internalization was tested in ENDO180 expressing NRK52 cells at permissive (37oC) and non-permissive (4oC) temperature and tested after one hour. Figure 2A shows level of fluorescence in ENDO180 expressing calls at permissive (37oC) and non-permissive
20 (4oC). Clones 6D6 and 8D8 and the scFv (SEQ ID NO:6) show highest level of internalization. Fig. 2B shows that 8D8 exhibits no non-specific binding.

The ENDO180 receptor was shown to be an internalization and recycling receptor (Howard and Isacke, 2002, JBC 277, 35:32320-31) yet not all antibodies produced are internalized at the same rate or in the same amount. mAb 8D8 and the G7V scFv showed
25 unexpected internalization. No mAb 10F2 was internalized, even after 8 hours.

Kinetics of internalization was studied: NRK52 cells stably expressing human ENDO180-FLAG or empty vector, were seeded in TC-grade black 96-well plates at a density of 6000 cells/well. At 24 or 48 hrs later, mAbs (5ug/ml in growth medium) were added to the wells, 100ul/well at either room temperature or on ice (for control plates). The plates were
30 immediately incubated at 37oC for various times. Control plates were kept on ice for the required times. At each time point, the plates were washed 3X in 200ul ice-cold PBS. After last wash, 100ul ice-cold PBS was added and plates read using Analyst at Ex 610nm/Em 670nm.

Figures 2C-2E show a 10-minute to 1-hour time course of internalization of anti-ENDO180-CypHer5E conjugates by ENDO180 expressing cells. Fig. 2C shows internalization of CypHer5E labeled 8D8. Fig. 2D shows internalization of CypHer5E labeled scFv G7V (SEQ ID NO:6). Fig. 2E shows internalization of CypHer5E labeled 10F2.

Figures 2F-2H shows a 1-hour to 8-hour time course of internalization of anti-ENDO180-CypHer5E conjugates by ENDO180 expressing cells. Fig. 2F shows internalization of CypHer5E labeled 8D8. Fig. 2G shows internalization of CypHer5E labeled scFv G7V (SEQ ID NO:6). Fig. 2E shows internalization of CypHer5E labeled 10F2.

10 The hybridoma cell line E3-8D8 that secretes monoclonal antibody E3-8D8, also referred to as 8D8, was deposited as per the Budapest Treaty in the Belgian Co-ordinated Collections of Micro-organisms (BCCM); Department of Biomedical Molecular Biology; Ghent University with Accession Number LMBP 7203CB. The deposit was made on 9-March-2010 and tested and shown to be viable on 18-March-2010.

15 A composition for the systemic uptake of a drug across a mucosal membrane comprising a polyethylene glycol-chitosan conjugate, wherein the polyethylene glycol-chitosan conjugate comprises a chitosan or chitosan derivative moiety and a polyethylene glycol or a polyethylene glycol derivative moiety, and the composition is formulated for delivery to a mucosal membrane.

20 The general method of preparing a substrate-agent conjugate according to the invention involves covalently binding at least one therapeutic or diagnostic agent to a substrate. Certain cytotoxic drugs that are useful for anticancer therapy are relatively insoluble in serum. Some are also quite toxic in unconjugated form and their toxicity is considerably reduced by conversion to conjugates. Conversion of a relatively poorly soluble drug to a
25 more soluble conjugate, e.g., a glucuronide, will improve its solubility in the aqueous phase of serum and its ability to pass through venous, arterial or capillary cell walls and reach the interstitial fluid bathing the tumor. In fact, conversion of certain toxic substances such as aromatic or alicyclic alcohols, thiols, phenols and amines to glucuronides in the liver is the body's method of detoxifying them and making them more easily excreted in
30 the urine.

EXAMPLE 4: in vitro and in vivo internalization of antibody conjugates

Figure 3 shows internalization of Biotin by anti-ENDO180 mAbs to mice Unilateral Ureter Obstructed kidney.

The following experiment was designed in order to assay ENDO180 antibody
5 accumulation in ENDO180 expressing tissue. Unilateral ureter obstructed (UUO) was performed in mice. The level of ENDO180 in kidneys at day 7 of UUO surgery, was higher than in the contra-lateral kidney (Data not shown).

Mice were injected with 3 mg/kg of E3- 8D8-Biotin conjugate or NMIgG-Biotin conjugate at day 7 post UUO surgery, 24 hours later the kidney. The level of E3-8D8-
10 Biotin conjugate and NMIgG-Biotin (normal mouse IgG control) conjugate uptake in the Unilateral ureter obstructed (UUO) kidney and Contra-lateral kidney was examined by Western blot (WB). The same amount of kidney total protein extract was examined by WB using Goat-anti-Biotin HRP (Cell signaling #7075).

Figure 4 shows internalization of anti-ENDO180-CypHer5E conjugate in Myelo-
15 Monocytoid human leukemia MonoMac cell line expressing ENDO180. MonoMac cells were incubated with E3 8D8-CypHer5E or E3 8D2-CypHer5E. Cells were washed twice with cold PBS and internalization was measured by FACS using FL-1 or FL-4 filter. E3 8D8 bound ENDO180 with higher affinity than E3-8D2 (Data not shown). 8D2 is a mAb that binds ENDO180 with lower affinity than 8D8.

20 EXAMPLE 5: Linking antibody to therapeutic agent

- a. MB: Full human protamine (~ 50 a.a) is cloned directly downstream to the constant region of the heavy chain. A similar strategy was taken using anti HIV-1gp 120 recombinant Fab fragment with a bicistronic vector expressing VH-CH1-protamine from one promoter and VKCK light chain from another
25 (Chen et al., Gene Therapy (1995) 2, 116-123). This construct was shown to have *in vivo* anti-tumor activity (Song et al., Nature Biotech. 2005. 23(6), pg. 709-717,). In another study, protamine was fused downstream to single chain antibody to ErbB (Li et al., Cancer Gene Therapy 8(8), 2001, pg. 555-565).
- b. mAb: The CDR domains of the monoclonal antibody 8D8 are sequenced and
30 cloned so that protamine can be engineered in fusion with it as with the MB.

- c. Standard methods are used to link the antibody or antigen-binding fragment thereof (MB, isolated mAb, scFv, F'ab etc.) to the therapeutic agent using one or more of a peptide, nucleic acid, chemical or lipid linker

5 EXAMPLE 6: in situ hybridization in cancer tissue samples

Samples of human tissue from cancer patients were tested for expression of ENDO180 using in situ hybridization techniques. The tissue samples were analyzed by a skilled pathologist. The results showed the following expression patterns:

1. Renal Cell Carcinoma (RCC)

10 High level of ENDO180 mRNA expression appeared in cells in all five sarcomatoid areas studied, in two different renal cell carcinoma types – clear cell (four cases) and papillary (one case) carcinomas. Sarcomatoid carcinomas develop in all main types of renal cell carcinomas (clear cell, papillary, chromophobe and collecting duct carcinomas). They appear in approximately 1- 1.5% of all adult
15 renal tumors and are associated with an aggressive clinical course and poor prognosis.

ENDO180 mRNA expression appears also in intratumoral stromal cells, in non tumoral stromal cells and in glomerular cells, with some sample to sample variation of the amount of cells, and signal intensity.

20 2. Ovarian cancer:

In most borderline serous papillary tumors that are represented in this study (7 out of 8), ENDO180 expression appeared in subsets of tumor cells, in various intensities.

25 ENDO180 mRNA expression in ovarian cancer cells appears in 8 out of 21 cases with some sample-to-sample variation of both amount and signal intensity of expressing cells.

High intensity signals of ENDO180 mRNA expression appeared in subsets of peritumor stromal cells and in subsets of ovarian stromal cells (94% and 100%). ENDO180 mRNA expression was also detected in single cells in normal
30 epithelium.

3. Transitional Cell Carcinoma (TCC):

In most primary transitional cell (urothelial) carcinoma of bladder (18 out of 27 cases), and 4 cases of metastatic (in lymph nodes) tumors, that are represented in this study, ENDO180 expression was weak. High intensity signals of ENDO180 mRNA expression appeared in subsets of peritumor stromal cells.

4. Breast cancer:

Expression of ENDO180 mRNA was seen in the peritumoral cells in most of the cases of invasive carcinoma (84%). No consistent expression pattern in epithelial cells.

10 EXAMPLE 7: Animal models for testing compounds in treatment of fibrosis

The following animal models are presented as non-limiting examples for use in testing exemplary molecules and conjugates of the present invention for efficacy in treating a subject suffering from fibrosis and fibrotic diseases. Other animal models are also considered.

15 A useful way to assess the development of renal diseases involving fibrosis and glomerulosclerosis is to characterize gene expression in established animal models of kidney diseases. Examples of such models include without limitation: (i) fa/fa rats - animals genetically deficient in leptin receptor that develop insulin resistant diabetes (type II diabetes) with progressive diabetic nephropathy, and (ii) GK rats - which are genetically manipulated, NIDDM phenotype rats. Another animal model in which mainly kidney fibrosis is evident, but without a background of diabetes, is unilateral ureteral obstruction (UUO) in which interstitial fibrosis is rapid and occurs within days following the obstruction. 5/6 nephrectomy is another useful animal model for chronic renal insufficiency (CRI) in which fibrosis is evident.

25 Additional aspects of research may be based on an in vitro model system involving culture of human fibroblasts in vitro under conditions mimicking various parameters of the cell microenvironment existing in CRI and fibrosis. These include treatment with high concentrations of glucose (modeling hyperglycemia), low concentrations of glucose, hypoxia (both modeling ischemic conditions that develop in the kidney following fibrosis and glomerulosclerosis), and TGF- β - one of the recognized pathogenic factors in fibrosis. Such in vitro model systems may complement the animal models in several important aspects: First, the system is fibroblast-specific; accordingly, none of the interferences

often found in complex tissues that contain many cell types are present. Second, the cells are of human origin, unlike the animal models. Furthermore, the insults are specific and of various concentrations and duration, thus enabling the investigation of both acute and chronic responses.

5 EXAMPLE 8: Animal models for testing the compounds in Cancer therapy

The following animal models are presented as non-limiting examples for use in testing exemplary molecules and conjugates of the present invention for efficacy in treating a subject suffering from cancer and other proliferative and metastatic diseases. Other animal models are also considered.

10 Transplantation in immunodeficient mice

The NOD/SCID mouse is defective in both lymphoid and myeloid function and readily accepts the long-term survival of human hematopoietic cells. Transplantation of human bone marrow into NOD/SCID mice to human/mouse chimeras, is well documented.

15 The NOD/SCID mice were used by Bertolini et al (2000. Blood 96-282) as a model high-grade non-Hodgkin lymphoma. The Namalwa cell line was used, which is derived from an Epstein-Barr virus-positive Burkitt non-Hodgkin lymphoma. The cells (10×10^6) were injected intraperitoneally into 6-8 weeks old mice. Intraperitoneal tumors were formed in the injection site which could be measured by calipers. The formula: “width² x length x 0.52” was applied to approximate the volume of a spheroid. (see Bohem et al (1997) for
20 further reference).

The model used in the following studies is based on transgenic SCID mice expressing human GM-CSF. The expression of this cytokine enabled the successful grafting of relevant cells lines in the SCID mice. Miyakawa et al (1996) details the production of the hGM-CSF SCID transgenic mice. Fukuchi et al (1998) shows that a retinoic-acid resistant
25 leukemia can be established in these transgenic mice. The model consisted of UF-1 cells, an RA-resistant APL cell line established in that laboratory, which are transplanted into these transgenic SCID mice and cause the appearance of subcutaneous tumors. Kinjo et al (2000) uses this model to test a specific treatment, arsenic trioxide, to be used in cases of RA resistant acute promyelocytic leukemia (APL).

30 Lewis et al (1998) used the more profoundly immunodeficient mouse strain NOD/SCID in which both T-cell and B-cells are functionally defective, and there is marked impairment of macrophage, natural killer cell, and hemolytic complement activity. These mice can be

engrafted with cells taken from cancer patients leading to a relatively high success rate and thus form a good model for the disease.

CLAIMS

1. An anti-ENDO180 antibody or antigen-binding fragment thereof, wherein the anti-ENDO180 antibody is selected from the group consisting of:
 - a. the isolated monoclonal antibody produced by the hybridoma cell line E3-8D8 deposited with the BCCM with Accession Number LMBP 7203CB;
 - b. a humanized version of the antibody of (a), or an antigen binding fragment thereof; and
 - c. a chimeric version of the antibody of (a), or an antigen binding fragment thereof;wherein upon contact with a cell expressing ENDO180 the antibody or antigen-binding fragment thereof is internalized into the cell.
2. A composition comprising the anti-ENDO180 antibody or the antigen-binding fragment thereof of claim 1, and a pharmaceutically acceptable carrier.
3. The composition of claim 2, further comprising a moiety selected from the group consisting of a detectable label, a cytotoxic agent and a therapeutic agent.
4. A conjugate comprising:
 - a) the antibody or antigen-binding fragment of claim 1;
 - b) a moiety selected from the group consisting of a detectable label, a cytotoxic agent and a therapeutic agent; and
 - c) optionally a linker which links a) to b).
5. The conjugate of claim 4, wherein the linker is selected from a polypeptide linker, peptide linker, lipid linker and nucleic acid linker.
6. The composition of claim 3 or the conjugate of claim 4, wherein the moiety is a therapeutic agent which is an inhibitory oligonucleotide selected from the group consisting of an antisense compound, a chemically modified siRNA compound, an unmodified siRNA compound, a chemically modified shRNA compound, an unmodified shRNA compound, a chemically modified miRNA compound, and an unmodified miRNA compound.
7. The composition or conjugate of claim 6, wherein the inhibitory oligonucleotide is a chemically modified siRNA compound.

8. A composition comprising the conjugate of any one of claims 4 to 7; and a pharmaceutically acceptable carrier.
9. The composition of claim 3, wherein the carrier comprises a lipid particle, a polysaccharide particle, a liposome or a combination thereof.
10. The composition of claim 3, wherein the carrier comprises a lipidated polysaccharide particle.
11. The composition of claim 10, wherein the lipidated polysaccharide comprises lipidated glycosaminoglycan.
12. The composition of claim 9 or 10, wherein the anti-ENDO180 antibody or antigen-binding fragment thereof is immobilized on the particle or liposome.
13. The composition of any one of claims 9, 10 or 12, wherein the moiety is encapsulated within the particle or liposome.
14. The composition of any one of claims 3 and 8 to 13, for the treatment of a disease selected from the group consisting of cancer, fibrosis and macrophage-associated disease.
15. The anti-ENDO180 antibody or antigen binding fragment of claim 1, wherein the anti-ENDO180 antibody is full IgG or wherein the antigen binding fragment is selected from the group consisting of: an Fab fragment, an F(ab')₂ fragment, an Fab miniantibody and an scFv.
16. The hybridoma cell line E3-8D8, deposited with the BCCM with Accession Number LMBP 7203CB.
17. The anti-ENDO180 antibody or antigen binding fragment of claim 1 or 15, wherein the antibody or antigen binding fragment is attached to a surface of a delivery particle, the delivery particle comprising at least one moiety selected from the group consisting of: a detectable label, a cytotoxic agent and a therapeutic agent.
18. Use of the conjugate of claim 4 for delivering the moiety to an ENDO180 expressing cell in a subject.
19. Use of the composition of claim 13 for delivering the moiety to an ENDO180 expressing cell in a subject.

FIGURE 1A

1 CGGAGGAGGA CGCGAGCCCC UUGC GGGCGG UCAUCACAGC CCAGCCUCGG GGCUGCCACA
 61 GCGCGUUGCG CCUGUGCGCC CUCGGUCCCC GCGUCCACUG AGCGCCGCGC UCGGGGAUGG
 121 GGCCCGGCCG GCCGGCCCCC GCGCCUGGC CUCGUCACCU GCUGCGCUGC GUCCUGCUCC
 181 UCGGGUGCCU GCACCUCGGC CGUCCCGGCG CCCUGGGGA CGCCGCCUC CCGGAACCCA
 241 ACGUCUUCU CAUCUUCAGC CAUGGACUGC AGGGCUGCCU GGAGGCCAG GCGGGCAGG
 301 UCAGAGUCAC CCCGGCUUGC AAUACCAGCC UCCUGCCA GCGCUGGAAG UGGGUCUCCC
 361 GAAACCGGCU AUUCAACCUG GGUACCAUGC AGUGCCUGGG CACAGGCUGG CCAGGCACCA
 421 ACACCACGGC CUCCUGGGC AUGUAUGAGU GUGACCGGGA AGCACUGAAU CUUCGCUGGC
 481 AUUGUCGUAC ACUGGGUGAC CAGCUGUCCU UGCUCCUGGG GGCCCGCACC AGCAACAUAU
 541 CCAAGCCUGG CACCCUUGAG CGUGGUGACC AGACCCGCAG UGGCCAGUGG CGCAUCUACG
 601 GCAGCGAGGA GGACCUAUGU GCUCUGCCCU ACCACGAGGU CUACACCAUC CAGGGAAACU
 661 CCCACGGAAA GCCGUGCACC AUCCCUUCA AAUAUGACAA CCAGUGGUUC CACGGCUGCA
 721 CCAGCACGGG CCGCGAGGAU GGUCACCUGU GGUGUGCCAC CACCCAGGAC UACGGCAAAG
 781 ACGAGCGCUG GGGCUUCUGC CCAUCAAGA GUAACGACUG CGAGACCUUC UGGGACAAGG
 841 ACCAGCUGAC UGACAGCUGC UACCAGUUUA ACUUCAGUC CACGCUGUCG UGGAGGGAGG
 901 CCUGGGCCAG CUGCGAGCAG CAGGGUGCGG AUCUGCUGAG CAUCACGGAG AUCCACGAGC
 961 AGACCUACAU CAACGGCCUC CUCACUGGGU ACAGCUCCAC CCUGUGGAUC GGCUUGAAUG
 1021 ACUUGGACAC GAGCGGAGGC UGGCAGUGGU CGGACAACUC GCCCCUCAAG UACCUCAACU
 1081 GGGAGAGUGA CCAGCCGGAC AACCCAGUG AGGAGAACUG UGGAGUGAUC CGCACUGAGU
 1141 CCUCGGGCGG CUGGCAGAAC CGUGACUGCA GCAUCGCGCU GCCCUAUGUG UGCAAGAAGA
 1201 AGCCCAACGC CACGGCCGAG CCCACCCUC CAGACAGGUG GGCCAAUGUG AAGGUGGAGU
 1261 GCGAGCCGAG CUGGCAGCCC UUCAGGGCC ACUGCUACCG CCUGCAGGCC GAGAAGCGCA
 1321 GCUGGCAGGA GUCCAAGAAG GCAUGUCUAC GGGGCGGUGG CGACCUGGUC AGCAUCCACA
 1381 GCAUGGCGGA GCUGGAAUUC AUCACCAAGC AGAUAAGCA AGAGGUGGAG GAGCUGUGGA
 1441 UCGGCCUCAA CGAUUUGAAA CUGCAGAUGA AUUUUGAGUG GUCUGACGGG AGCCUUGUGA
 1501 GCUUCACCCA CUGGCACCCC UUUGAGCCCA ACAACUUCG GGACAGUCUG GAGGACUGUG
 1561 UCACCAUCUG GGGCCCGGAA GGCCGCUGGA ACGACAGUCC CUGUAACCAG UCCUUGCCA
 1621 CCAUCUGCAA GAAGGCAGGC CAGCUGAGCC AGGGGGCCGC CGAGGAGGAC CAUGGCUGCC
 1681 GGAAGGGUUG GACGUGGCAC AGCCAUCCU GCUACUGGCU GGGAGAAGAC CAAGUGACCU
 1741 ACAGUGAGGC CCGGCGCCUG UGCACUGACC AUGGCUCUCA GCUGGUCACC AUCACCAACA
 1801 GGUUCGAGCA GGCCUUCGUC AGCAGCCUCA UCUACAACUG GGAGGGCGAG UACUUCUGGA
 1861 CGGCCUGCA GGACCUCAAC AGCACCGGCU CCUUCUUCUG GCUCAGUGGG GAUGAAGUCA
 1921 UGUACACCCA CUGGAACCGG GACCAGCCCG GGUACAGCCG UGGGGGCGC GUGGCGCUGG
 1981 CCACUGGCAG CGCCAUGGGG CUGUGGGAGG UGAAGAACUG UACCUCGUUC CGGGCCCGCU
 2041 ACAUCUGCCG GCAGAGCCUG GGCACUCCAG UGACGCCGGA GCUGCCGGGG CCAGAUCCCA
 2101 CGCCAGCCU CACUGGCUCU UGUCCCAGG GCUGGGCCUC GGACACCAA CUCCGGUAUU
 2161 GCUAUAAGGU GUUCAGCUCA GAGCGGCUGC AGGACAAGAA GAGCUGGGUC CAGGCCAGG
 2221 GGGCCUGCCA GGAGCUGGGG GCCCAGCUGC UGAGCCUGGC CAGCUACGAG GAGGAGCACU

FIGURE 1A cont.

2281 UUGUGGCCAA CAUGCUC AAC AAGAUUCUUCG GUGAAUCAGA ACCCGAGAUC CACGAGCAGC
 2341 ACUGGUUCUG GAUCGGCCUG AACCGUCGGG AUCCCAGAGG GGGUCAGAGU UGGCGCUGGA
 2401 GCGACGGCGU AGGGUUCUCU UACCACAAUU UCGACCGGAG CCGGCACGAC GACGACGACA
 2461 UCCGAGGCUG UGCGGUGCUG GACCUGGCCU CCCUGCAGUG GGUGGCCAUG CAGUGCGACA
 2521 CACAGCUGGA CUGGAUCUGC AAGAUCCCCA GAGGUACGGA CGUGCAGGAG CCCGACGACA
 2581 GCCUCAAGG CCGACGGGAA UGGCUGCGCU UCCAGGAGGC CGAGUACAAG UUCUUUGAGC
 2641 ACCACUCCAC GUGGGCGCAG GCGCAGCGCA UCUGCACGUG GUUCCAGGCC GAGCUGACCU
 2701 CCGUGCACAG CCAGGCAGAG CUAGACUUC UGAGCCACAA CUUGCAGAAG UUCUCCCGGG
 2761 CCCAGGAGCA GCACUGGUGG AUCGGCCUGC ACACCUCUGA GAGCGAUGGG CGCUUCAGAU
 2821 GGACAGAUGG UCCAUUAUA AACUUCAUCU CCUGGGCACC AGGCAAACCU CGGCCUGUCG
 2881 GCAAGGACAA GAAGUGCGUG UACAUGACAG CCAGCCGAGA GGACUGGGGG GACCAGAGGU
 2941 GCCUGACAGC CUUGCCCUAC AUCUGCAAGC GCAGCAACGU CACCAAAGAA ACGCAGCCCC
 3001 CAGACCUGCC AACUACAGCC CUGGGGGGCU GCCCUCUGA CUGGAUCCAG UUCCUCAACA
 3061 AGUGUUUUA GGUCCAGGGC CAGGAACCCC AGAGCCGGGU GAAGUGGUCA GAGGCACAGU
 3121 UCUCUGUGA ACAGCAAGAG GCCCAGCUGG UCACCAUCAC AAACCCCUUA GAGCAAGCAU
 3181 UCAUCACAGC CAGCCUGCCC AAUGUGACCU UUGACCUUUG GAUUGGCCUC CAUGCCUCGC
 3241 AGAGGGACUU CCAGUGGGUG GAGCAGGAGC CUUUGAUGUA UGCCAACUGG GCACCUGGGG
 3301 AGCCUCUGG CCCUAGCCCU GCUCCAGUG GCAACAAACC GACCAGCUGU GCGGUGGUCC
 3361 UGCACAGCCC CUCAGCCCAC UUCACUGGCC GCUGGGACGA UCGGAGCUGC ACGGAGGAGA
 3421 CCAUGGCUU CAUCUGCCAG AAGGGCACGG ACCCUCCCU GAGCCCGUCC CCAGCAGCGC
 3481 UGCCCCCGC CCCGGGCACU GAGCUCUCCU ACCUCAACGG CACCUUCCGG CUGCUCAGGA
 3541 AGCCGCUGCG CUGGCACGAU GCCCUCUGC UGUGUGAGAG CCACAAUGCC AGCCUGGCCU
 3601 ACGUGCCCGA CCCUACACC CAGGCCUUC UCACGCAGGC UGCCGAGGG CUGCGCACGC
 3661 CGCUCUGGAU UGGCUGGCCU GCGGAGGAG GCUCUCGGCG GUACUCCUGG GUCUCAGAGG
 3721 AGCCGCUGAA CUACGUGGGC UGGCAGGACG GGGAGCCGCA GCAGCCGGGG GGCUGUACCU
 3781 ACGUAGAUGU GGACGGGGCC UGGCGCACCA CCAGCUGUGA CACCAAGCUG CAGGGGGCUG
 3841 UGUGUGGGGU UAGCAGUGGG CCCCUCUCU CCCGAAGAAU AAGCUACCAU GGCAGCUGUC
 3901 CCCAGGGACU GGCAGACUCC GCGUGGAUUC CCUCCGGGA GCACUGCUAU UCUUCCACA
 3961 UGGAGCUGCU GCUGGGCCAC AAGGAGGCGC GACAGCGCUG CCAGAGAGCG GGUGGGGCGG
 4021 UCCUGUCUAU CCUGGAUGAG AUGGAGAAUG UGUUUGUCUG GGAGCACCUG CAGAGCUAUG
 4081 AGGGCCAGAG UCGGGGCGCC UGGCUGGGCA UGAACUUC AA CCCC AAAGGA GGCACUCUGG
 4141 UCUGGCAGGA CAACACAGCU GUGAACUACU CCAACUGGGG GCCCCCGGGC UUGGGCCCCA
 4201 GCAUGCUGAG CCACAACAGC UGCUACUGGA UUCAGAGCAA CAGCGGGCUA UGGCGCCCCG
 4261 GCGCUUGCAC CAACAUCACC AUGGGUGUCG UCUGCAAGCU UCCUCGUGCU GAGCAGAGCA
 4321 GCUUCUCCCC AUCAGCGCUU CCAGAGAACC CAGCGGCCCU GGUGGUGGUG CUGAUGGCGG
 4381 UGCUGCUGCU CCUGGCCUUG CUGACCGCAG CCCUCAUCCU UUACCGGAGG CGCCAGAGCA
 4441 UCGAGCGCGG GGCCUUUGAG GGUGCCCGCU ACAGCCGCAG CAGCUCCAGC CCCACCGAGG
 4501 CCACUGAGAA GAACAUCUG GUGUCAGACA UGGAAAUGAA UGAGCAACAA GAAUAGAGCC

FIGURE 1A cont.

4561 AGGCGCGUGG GCAGGGCCAG GCGGGGAGGA GCUGGGGAGC UGGGGCCCUG GGUCAGUCUG
 4621 GCCCCCACC AGCUGCCUGU CCAGUUGGCC UAUGGAAGGG UGCCCUUGGG AGUCGCUGUU
 4681 GGGAGCCGGA GCUGGGCAGA GCCUGGGCUG GUGGGGUGCC ACCCUCCCAC AAGGGCUGGG
 4741 CUGAGACCCA GCUGAGUGCA GCGUGGCGUU UCCCUUUCUG GGGGGGCCUG AGGUCUUGUC
 4801 ACCUGGUCCU GUGCCCCCAC AGGAACCAGA GGUAGGAUGG GAGGGGGAAC GAGAGCCUCU
 4861 UUCUCCCCAG AGCCCCCGGC CCAGGCCUGU UGAUCCGCGC CCCAGGACCC CCUUCUUUGC
 4921 AGAGCCCGAG GAGCCUCCCC UGUCCCCUCG GGCAGAUUCG UUGUGUCUCU CUUCCCACCU
 4981 GGCAGCCUCA GCUCUGUGCC CCUCACCCUG CUCCCUCUCG CCCCUUCUCU CCCACCCUU
 5041 CCUUCUGAGC CGGGCCCUGG GGAUUGGGGA GCCUCUUGU UCCUGAUGAG GGUCAGCUGA
 5101 GGGGGCUGAG CAUCCAUCAC UCCUGUGCCU GCUGGGGUGG CUGUGGGGCG UGGCAGGAGG
 5161 GGCCUAGGUG GGUUGGGCCU GAGAACCAGG GCACGGGUGU GGUGUCUGCU GGGCUGGAGA
 5221 UAAGACUGGG GAGAGACACC CCAACCUCUCC AGGGUGGGAG CUGGGCCGGG CUGGGAUGUC
 5281 AUCUCCUGCC GGGCGGGGGA GGGCUCUGCC CCUGGAAGAG UCCCCUGUGG GGACCAAAAU
 5341 AAGUUCCCUA ACAUCUCCAG CUCCUGGCUC UGGUUUGGAG CAAGGGGAAG GGUUGCCAGA
 5401 GUCCUGGGGG CCCAGAGGA GCAGGAGUCU GGGAGGGCCC AGAGUUCACC CUCUAGUGGA
 5461 UCCAGGAGGA GCAGCACCCG AGCCUGGAG UGGCCAGUA CCCUCCAAG AGGCCACAGU
 5521 CCCAGCCAGG ACAAAGUAUG CGGCCCAUCC UGGUGCGACA GCGUGGGACA AUGUGAACAU
 5581 GGACUCGAAG ACAUGGCCCU UUCUCUGUAG UUGAUUUUUU AAAUGUGCCA UUAUUGUUUU
 5641 U

FIGURE 1B

1 MGPGRPAPAP WPRHLLRCVL LLGCLHLGRP GAPGDAALPE PNVFLIFSHG LQGCLEAQQG
 61 QVRVTPACNT SLPAQRWKWV SRNRLFNLGT MQCLGTGWPG TTTASLGM Y ECDREALNLR
 121 WHCRTLGDL SLLLGARTSN ISKPGTLERG DQTRSGQWRI YGSEEDLCAL PYHEVYTIQG
 181 NSHGKPTIP FKYDNQWFHG CTSTGREDGH LWCATTQDYG KDERWGFCEP I KSNDCETFWD
 241 KDQLTDSCYQ FNFQSTLSWR EAWASCEQQG ADLLSITEIH EQTYINGLLT GYSSTLWIGL
 301 NDLDTSGGWQ WSDNSPLKYL NWESDQPDNP SEENCGVIRT ESSGGWQNRD CSIALPYVCK
 361 KKPNTAEPT PPDRWANVKV ECEPSWQPFQ GHCYRLQAEK RSWQESKAC LRGGDLVSI
 421 HSMAELEFIT KQIKQEVEEL WIGLNDLKLQ MNFEWSDGSL VSFTHWHPFE PNNFRDSLED
 481 CVTIWGPEGR WNDSPCNQSL PSICKKAGQL SQGAAEEDHG CRKGWTWHSP SCYWLGEDQV
 541 TYSEARRLCT DHGSQLVTIT NRFEQAFVSS LIYNWEGEYF WTALQDLNST GSFFWLSGDE
 601 VMYTHWNRDQ PGYSRGGCVA LATGSAMGLW EVKNCTSFRA RYICRQSLGT PVTPELPGPD
 661 PTPSLTGSCP QGWASDTKLR YCYKVFSSER LQDKKSWVQA QGACQELGAQ LLSLASYEEE
 721 HFVANMLNKI FGESEPEIHE QHFWWIGLNR RDPRGGQSWR WSDGVGFSYH NFDRSRHDDD
 781 DIRGCAVLDL ASLQWVAMQC DTQLDWICKI PRGTDVREPD DSPQGRREW L RFQEA EYKFF
 841 EHHSTWAQAQ RICTWFQ AEL TSVHSQAELD FLSHNLQKFS RAQE QHWWIG LHTSESDGRF
 901 RWTDGSIINF ISWAPGKPRP VGKDKKCVYM TASREDWGDQ RCLTALPYIC KRSNVTKETQ
 961 PPDLPPTALG GCPDWIQFL NKCFQVQGQE PQSRVKWSEA QFSCEQQEAQ LVTITNPLEQ

FIGURE 1B cont

1021 AFITASLPNV TFDLWIGLHA SQRDFQWVEQ EPLMYANWAP GEPSGPSPAP SGNKPTSCAV
 1081 VLHSPSAHFT GRWDDRSCTE ETHGFICQKG TDPSSLSPSPA ALPPAPGTEL SYLNGTFRLL
 1141 QKPLRWHDAL LLCESHNASL AYVPDPYTQA FLTQAARGLR TPLWIGLAGE EGSRRYSWVS
 1201 EEPLNYVGWQ DGEPQQPGGC TYVDVDGAWR TTSCDTKLQG AVCGVSSGPP PRRRISYHGS
 1261 CPQGLADSAW IPFREHCYSF HMELLLGHKE ARQRCQRAGG AVLSILDEME NVFVWEHLQS
 1321 YEGQSRGAWL GMNFPKGGT LVWQDNTAVN YSNWGPPLG PSMLSHNSCY WIQSNSGLWR
 1381 PACTNITMG VVCKLPRAEQ SSFSPSALPE NPAALVVVLM AVLLLLLALLT AALILYRRRQ
 1441 SIERGAFEGA RYSRSSSSPT EATEKNILVS DMEEMNEQQE

FIGURE 1C

ATGGGGCCCGGCCGGCCGGCCCGCCCGCGCCCTGGCCTCGTCACCTGCTGCGCTGCGTCCTG
 CTCCTCGGGTGCCTGCACCTCGGCCGTCCCGGCCCGCCCTGGGGACGCCGCCCTCCCGGAA
 CCCAACGTCTTCCTCATCTTCAGCCATGGACTGCAGGGCTGCCTGGAGGCCAGGGCGGG
 CAGGTCAGAGTCACCCCGGCTTGCAATACCAGCCTCCCTGCCAGCGCTGGAAGTGGGTC
 TCCCGAAACCGGCTATTCAACCTGGGTACCATGCAGTGCCTGGGCACAGGCTGGCCAGGC
 ACCAACACCACGGCCTCCCTGGGCATGTATGAGTGTGACCGGGAAGCACTGAATCTTCGC
 TGGCATTGTCTGACTACTGGGTGACCAGCTGTCTTGCTCCTGGGGGCCCGCACCAGCAAC
 ATATCCAAGCCTGGCACCCTTGAGCGTGGTGACCAGACCCGCAGTGGCCAGTGGCGCATC
 TACGGCAGCGAGGAGGACCTATGTGCTCTGCCCTACCACGAGGTCTACACCATCCAGGGA
 AACTCCCACGGAAAGCCGTGCACCATCCCCTTCAAATATGACAACCAGTGGTTCACGGC
 TGCACCAGCACGGGCCGCGAGGATGGTACCTGTGGTGTGCCACCACCAGGACTACGGC
 AAAGACGAGCGCTGGGGCTTCTGCCCATCAAGAGTAACGACTGCGAGACCTTCTGGGAC
 AAGGACCAGCTGACTGACAGCTGTACCAGTTTAACTTCCAGTCCACGCTGTCTGGAGG
 GAGGCTGGGCCAGCTGCGAGCAGCAGGGTGCAGGATCTGCTGAGCATCACGGAGATCCAC
 GAGCAGACCTACATCAACGGCCTCCTCACTGGGTACAGCTCCACCCTGTGGATCGGCTTG
 AATGACTTGGACACGAGCGGAGGCTGGCAGTGGTCCGACAACCTCGCCCTCAAGTACCTC
 AACTGGGAGAGTGACCAGCCGGACAACCCAGTGAGGAGAACTGTGGAGTGATCCGCACT
 GAGTCTCGGGCGGCTGGCAGAACCGTGACTGCAGCATCGCGCTGCCCTATGTGTGCAAG
 AAGAAGCCCAACGCCACGGCCGAGCCACCCCTCCAGACAGGTGGGCCAATGTGAAGGTG
 GAGTGCAGCCGAGCTGGCAGCCCTTCCAGGGCCACTGCTACCGCCTGCAGGCCGAGAAG
 CGCAGCTGGCAGGAGTCCAAGAAGGCATGTCTACGGGGCGGTGGCGACCTGGTCCAGCATC
 CACAGCATGGCGGAGCTGGAATTCATCACCAGCAGATCAAGCAAGAGGTGGAGGAGCTG
 TGGATCGGCCTCAACGATTTGAAACTGCAGATGAATTTTGTAGTGGTCTGACGGGAGCCTT
 GTGAGCTTCACCACTGGCACCCTTTGAGCCCAACAACCTTCCGGGACAGTCTGGAGGAC
 TGTGTACCATCTGGGGCCCGGAAGGCCGCTGGAACGACAGTCCCTGTAACCAGTCCCTTG
 CCATCCATCTGCAAGAAGGCAGGCCAGCTGAGCCAGGGGGCCCGCAGGAGGACCATGGC
 TGCCGGGATTACAAGGACGACGACGATAAGTGA

FIGURE 1D

MGPGRPAPAPWPRHLLRCVLLLGCLHLGRPGAPGDAALPEPNVFLIFSHGLQGCLEAQQG
 QVRVTPACNTSLPAQRWKWVSRNRLFNLGTMQCLGTGWPGTNTTASLGMYECDREALNLR
 WHCRTLGDQLSLLLGARTSNISKPGTLERGDQTRSGQWRIYGSEEDLCALPYHEVYTIQG
 NSHGKPCITPFKYDNQWFHGCTSTGREDGHLWCATTQDYGKDERWGFPCPKSNDCEFTWD
 KDQLTDSCYQFNQSTLSWREAWASCEQQGADLLSITEIHEQTYINGLLTGYSSTLWIGL
 NDLDTSGGWQWSDNSPLKYLNWESDQPDNPSEENCGVIRTESSGGWQNRDCSIALPYVCK
 KKPATAEPTPPDRWANVKVECEPSWQPFQGHQCYRLQAEKRSWQESKKA CLRGGDLVSI
 HSMAELEFITKQIKQEVEELWIGLNDLKLQMNFEWSDGSLVSFTHWHPFEPNNFRDSLED
 CVTIWGPEGRWNSPCNQSLPSICKKAGQLSQGAAEEDHGCRDYKDDDDK

FIGURE 1E

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PCT/US2010/028200

TAATGTGAGTTAGCTACTCTTAGGCACCCCAGGCTTTACACTTTATGCTTCCGGCTCGTATGTTGTGTGG
 AATTGTGAGCGGATAACAATTTACACACAGGAAACAGCTATGACCATGATTACGCCAAGCTTGCCAAATTC
 TATTTCAAGGAGACAGTCATAATGAAATACCTATTGCCTACGGCAGCCGCTGGATTGTTATTACTCGCAG
 CAAGCGGCGCGCATGCCCAGGTGCAGCTGGTGCAGTCTGGGGGAGGCCCTGGTCAAGCCTGGGGGGTCCCT
 GAGACTCTCCTGTGCAGCCTCTGGATTACCTTCAGTAGCTATAGCATGAACTGGGTCCGCCAGGCTCCA
 GGAAGGGGCTGGAGTGGGTGGCCAACATAAAGCCAGATGGAAGTGAGAGACACTCTGTGGACTCTGTGA
 AGGGCCGATTCACCATCTCCAGAGACAACTCCAAGAACTCACTGTATCTGCAAATGAACAGCCTGAGAGC
 CGAGGACACGGCTGTTTATTACTGTGCGCGACCCGGGGCTGGGCGACTTGACTACTGGGGCCAGGGCACC
 CTGGTCACCGTCTCCTCAGGTGGAGGCGGTTTCAGGCGGAGGTGGCTCTGGCGGTGGCGGATCGAATTTTA
 TGCTGACTCAGGACGCTGCTGTGTCTGTGGCCTTGGGACAGACAGTCAGGATCACATGCCAAGGAGACAG
 CCTCAGAAGCTATTATGCAAGCTGGTACCAACAGAAGCCAGGACAGGCCCTGTACTTGTCTGTCTATGGT
 AAAACAACCGACCCCTCAGGGATCCCAGACCGATTCTCTGGCTCCAGCTCAGGAAACACAGCTTCCTTGA
 CCATCACTGGGGCTCAGGCGGAAGATGAGGCTGACTATTACTGTAACCTCCCGGGACAGCAGTGGTAACCC
 CTGGGCGTTTCGGCGGAGGGACCAAGGTGACCGTCTTAGCTAGCGGCAAACCAATCCCAAACCCACTGCTG
 GGCCTGGATAGTACTCACCATCACCATCACCATTAGGCGGCCGCTACTGTTGAAAGTTGTTTACAAAAC
 CTCATACAGAAAATTCATTTACTAACGTCTGGAAAGACGACAAAACCTTTAGATCGTTACGCTAACTATGA
 GGGCTTG

FIGURE 1F

M K Y L L P T A A A G L L L L A A S G A H A Q V Q L V Q S G G G L V K
 P G G S L R L S C A A S G F T F S S Y S M N W V R Q A P G K G L E W V
 A N I K P D G S E R H S V D S V K G R F T I S R D N S K N S L Y L Q M
 N S L R A E D T A V Y Y C A R P G A G R L D Y W G Q G T L V T V S S G
 G G G S G G G S G G G S N F M L T Q D A A V S V A L G Q T V R I T
 C Q G D S L R S Y Y A S W Y Q Q K P G Q A P V L V V Y G K N N R P S G
 I P D R F S G S S S G N T A S L T I T G A Q A E D E A D Y Y C N S R D
S S G N P W A F G G G T K V T V L A S G K P I P N P L L G L D S T H H
 H H H H

FIGURE 1G

CDR3 - H CARPGAGRLDYW

FIGURE 1H

CDR3 - L CNSRDSSGNPWAF

FIGURE 1J

MGPGRPAPAPWPRHLLRCVLLLGCLHLGRPGAPGDAALPEPNVFLIFSHGLQGCLEAQQG
 QVRVTPACNTSLPAQRWKWVSRNRLFNLGTMQCLGTGWPGTNTTASLGMYECDREALNLR
 WHCRTLGDLQSLLLGARTSNISKPGTLERGDQTRSGQWRIYGSEEDLCALPYHEVYTIQG
 NSHGKPTIPFKYDNQWFHGCTSTGREDGHLWCATTQDYGKDERWGFPCPKSNDCETFWD
 KDQLTDSCYQFNQSTLSWREAWASCEQQGADLLSITEIHEQTYINGLLTGYSSTLWIGL
 NDLDTSGGWQWSDNSPLKYLNWESDQPDNPSEENCGVIRTESSGGWQNRDCSIALPYVCK
 KKP NATAEPTPPDRWANVKVECEPSWQPFQGHYRLQAEKRSWQESKKACLRGGDLVSI
 HSMAELEFITKQIKQEVEELWIGLNDLKLQMNFEWSDGSLVSFTHWHPFEPNNFRDSLED
 CVTIWGPGRWNSPCNQSLPSICKKAGQLSQGAAEEDHGCR

FIGURE 2A

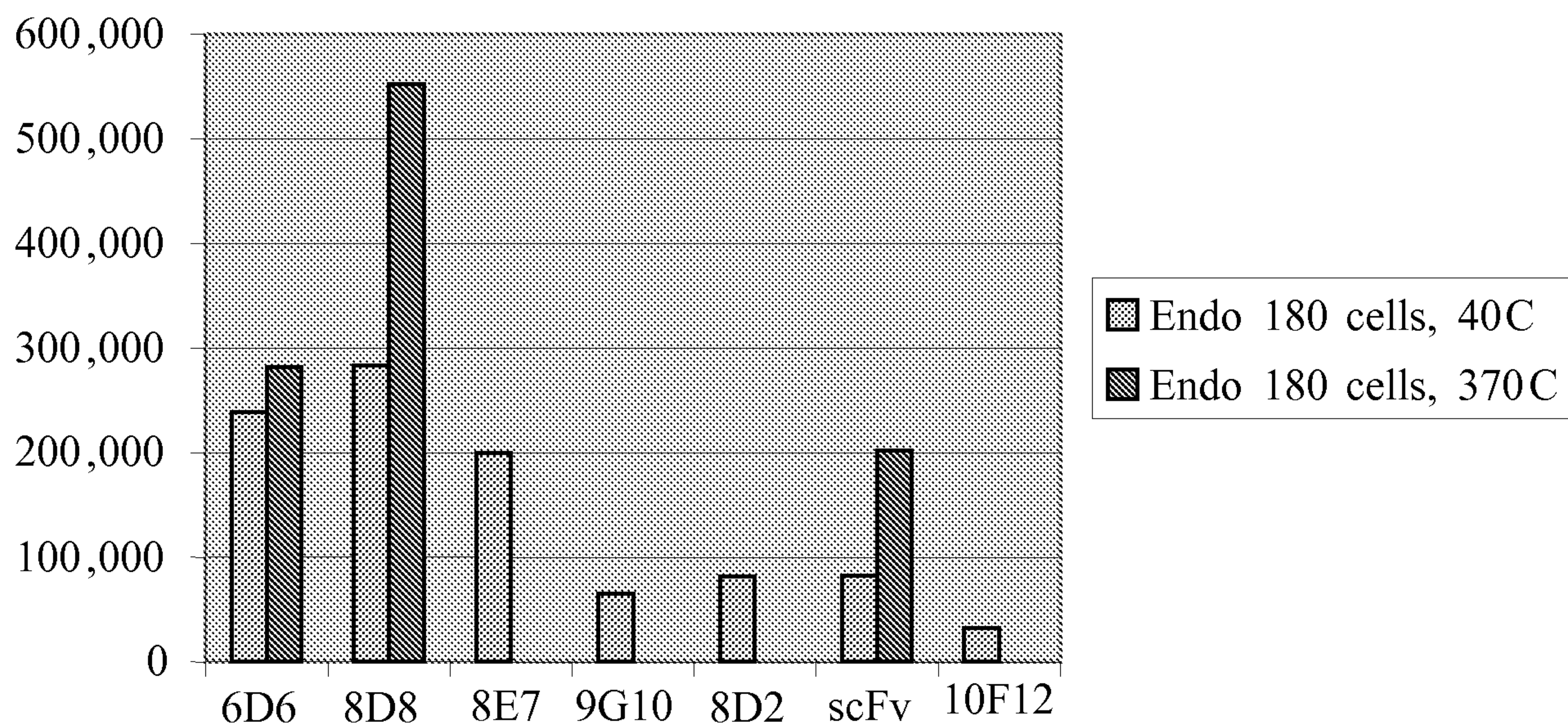


FIGURE 2B

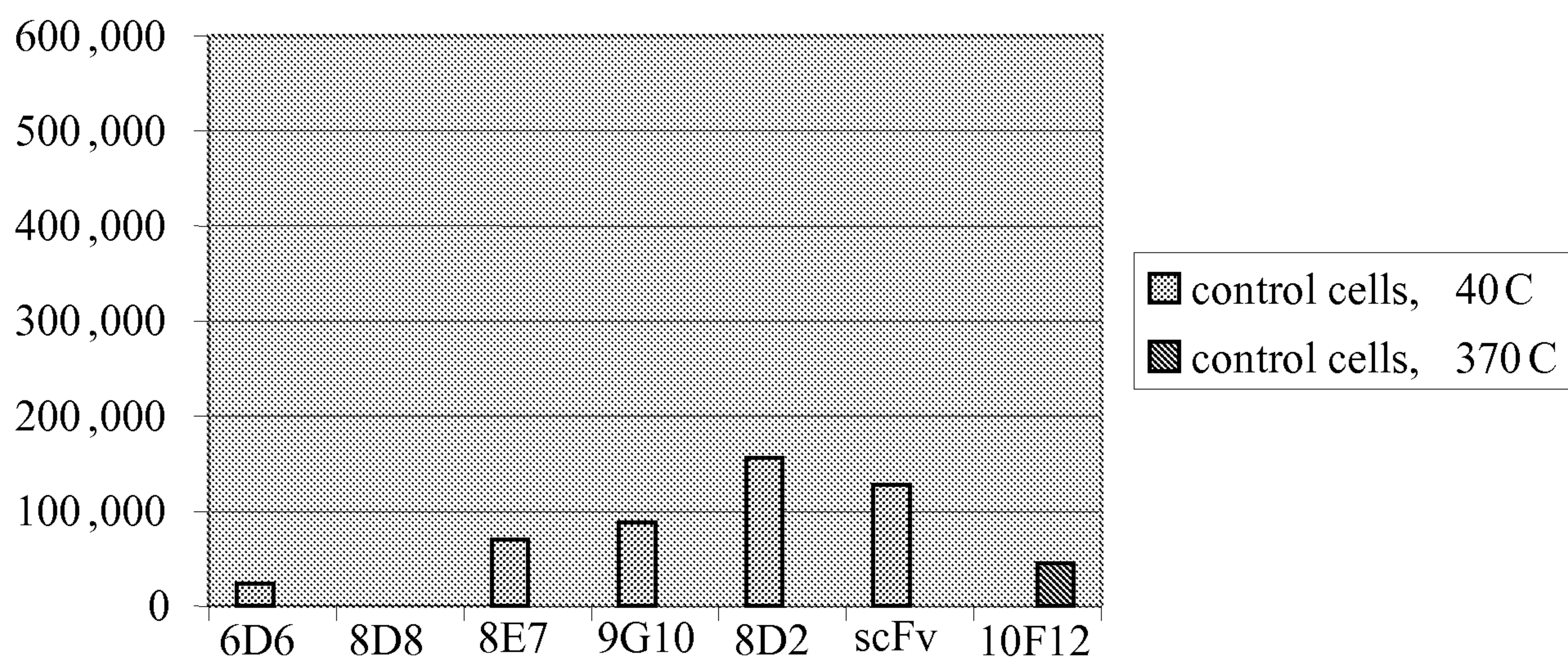


FIGURE 2C

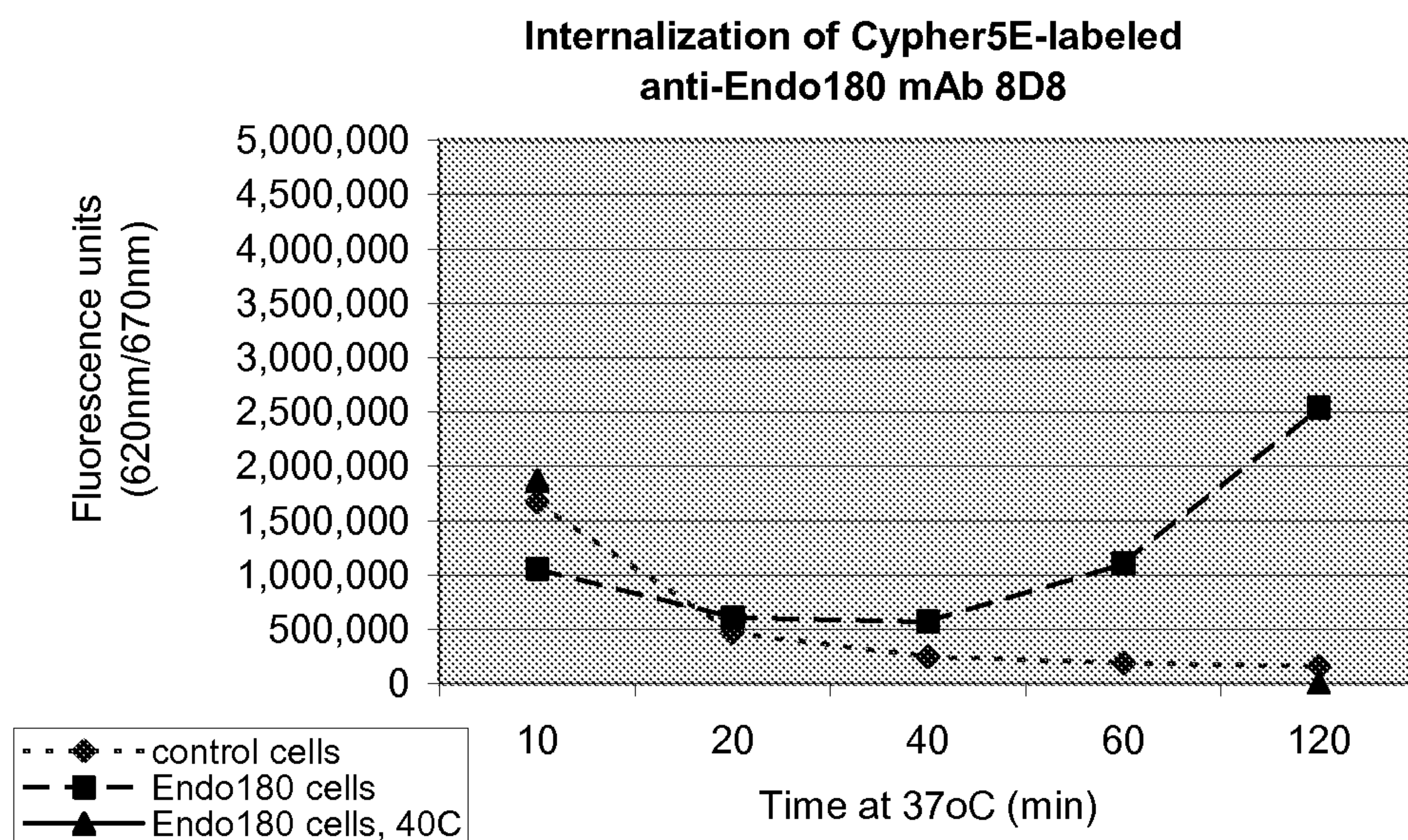


FIGURE 2D

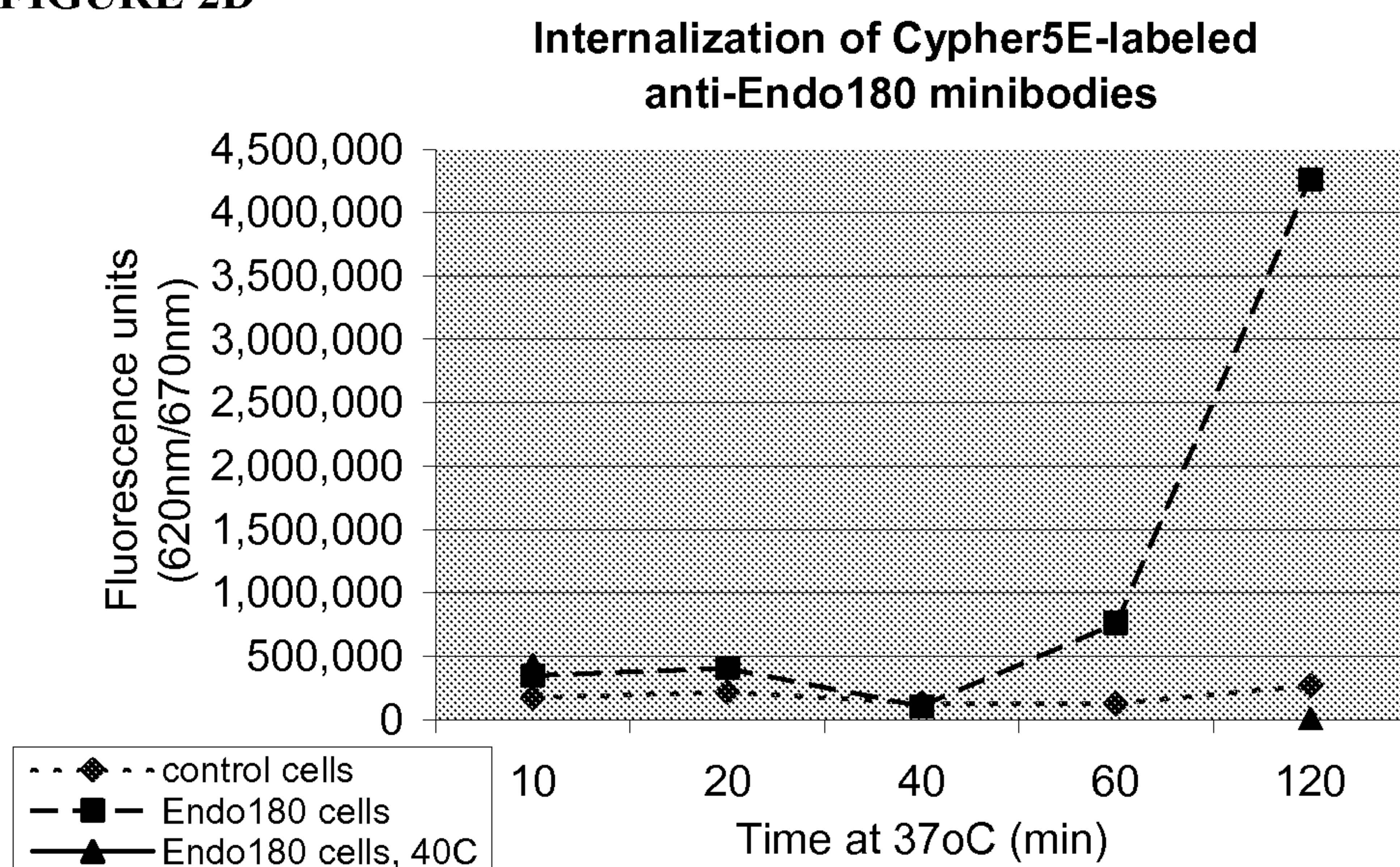


FIGURE 2E

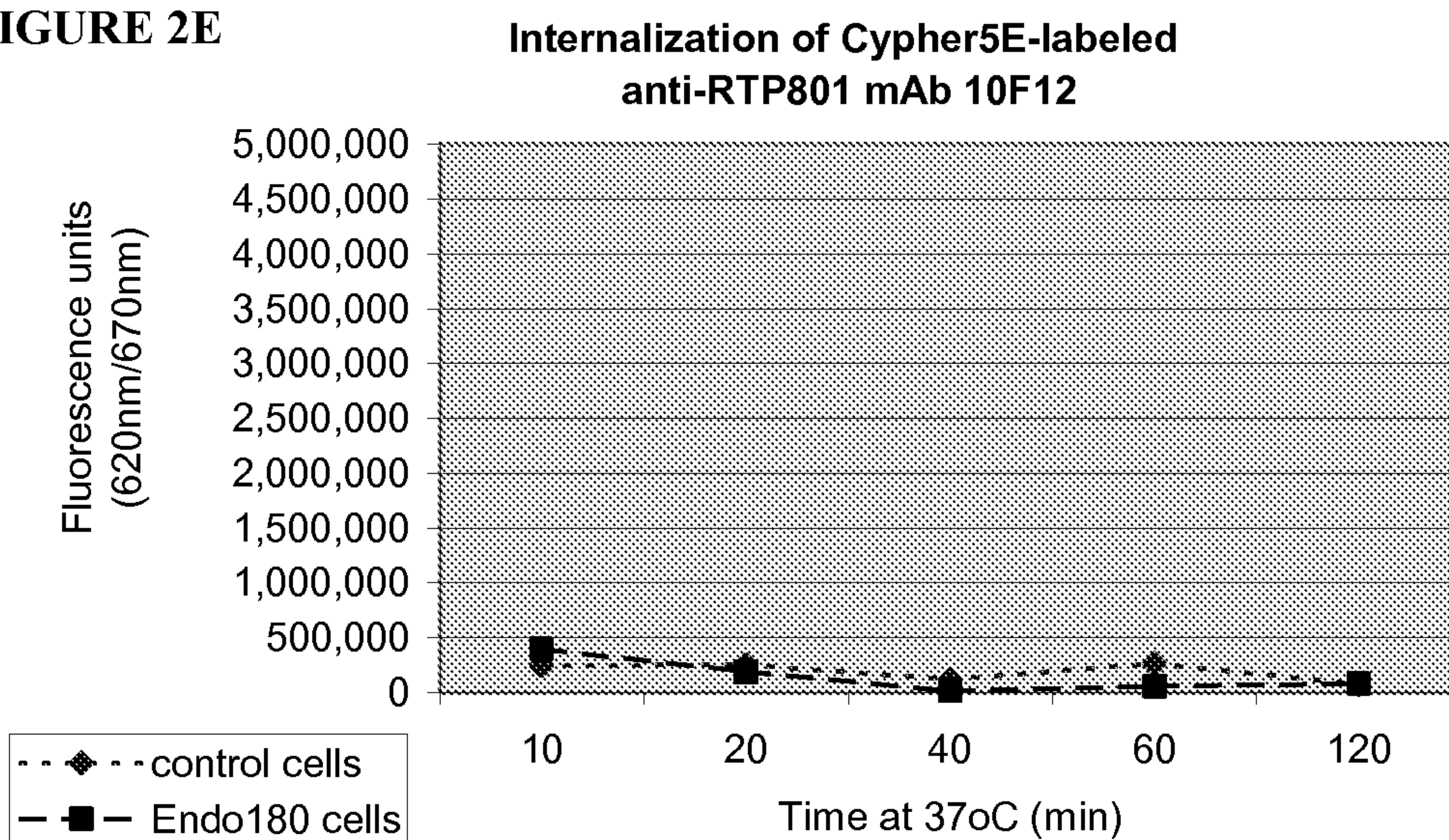


FIGURE 2F

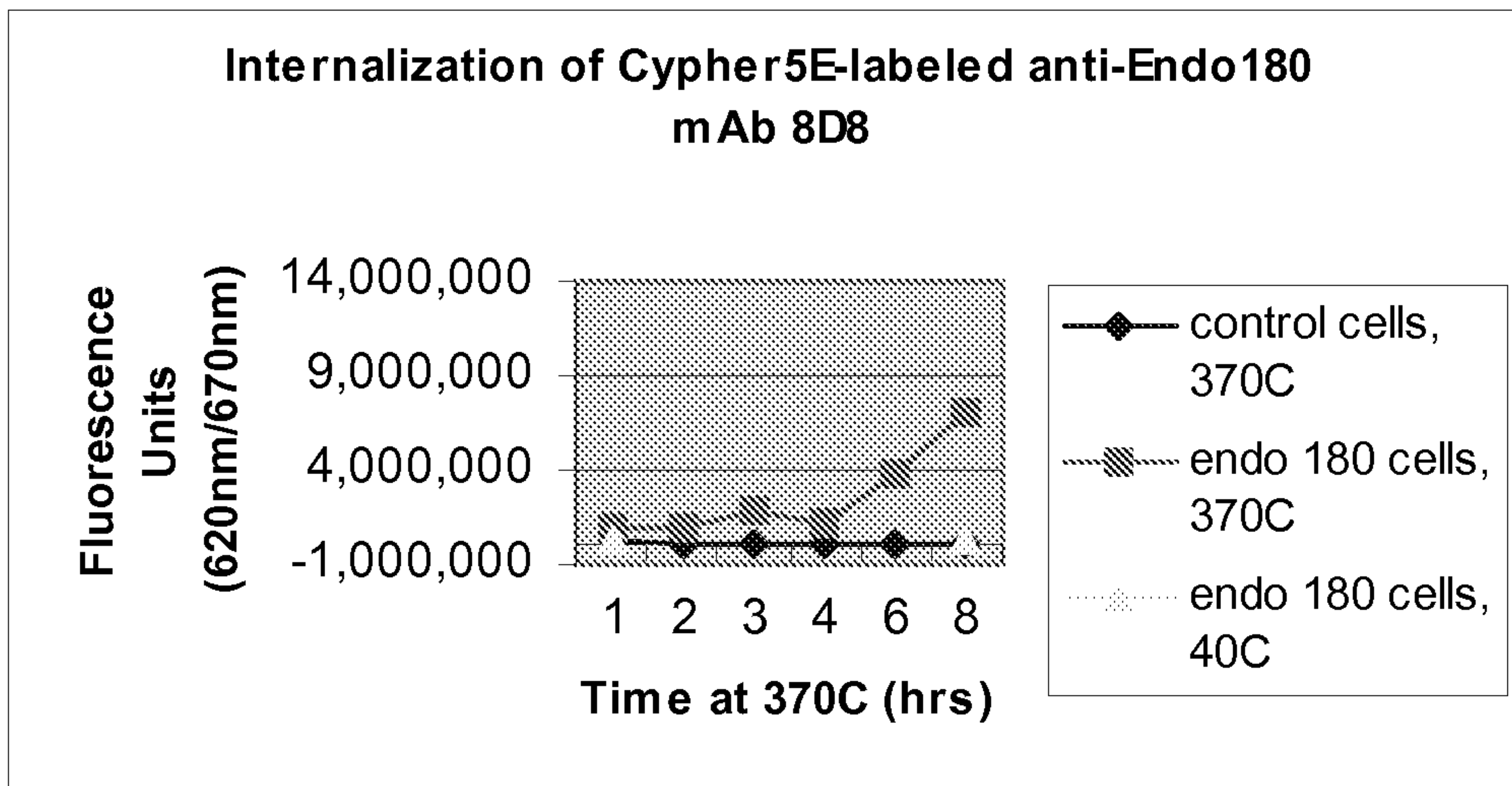


FIGURE 2G

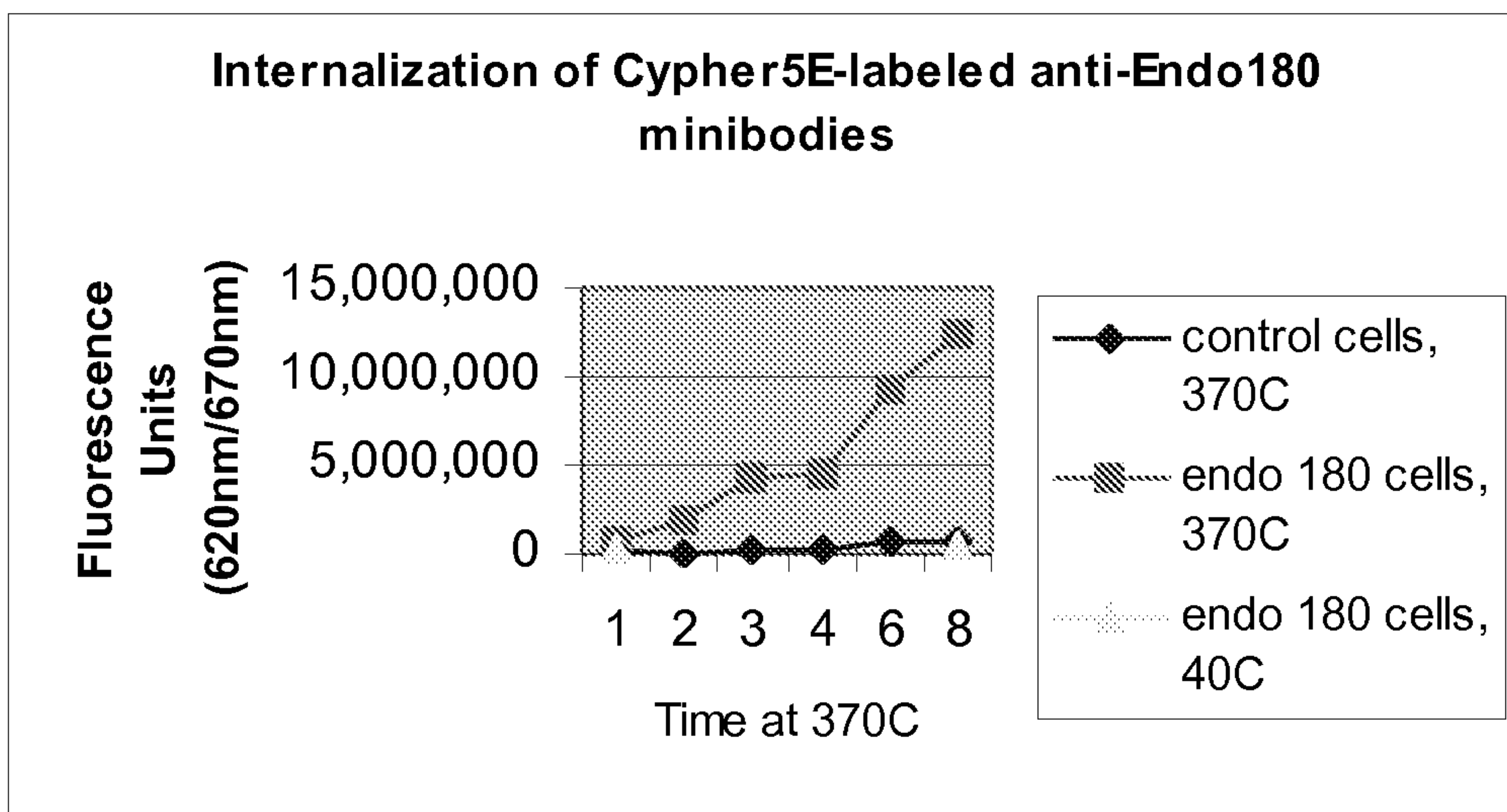


FIGURE 2H

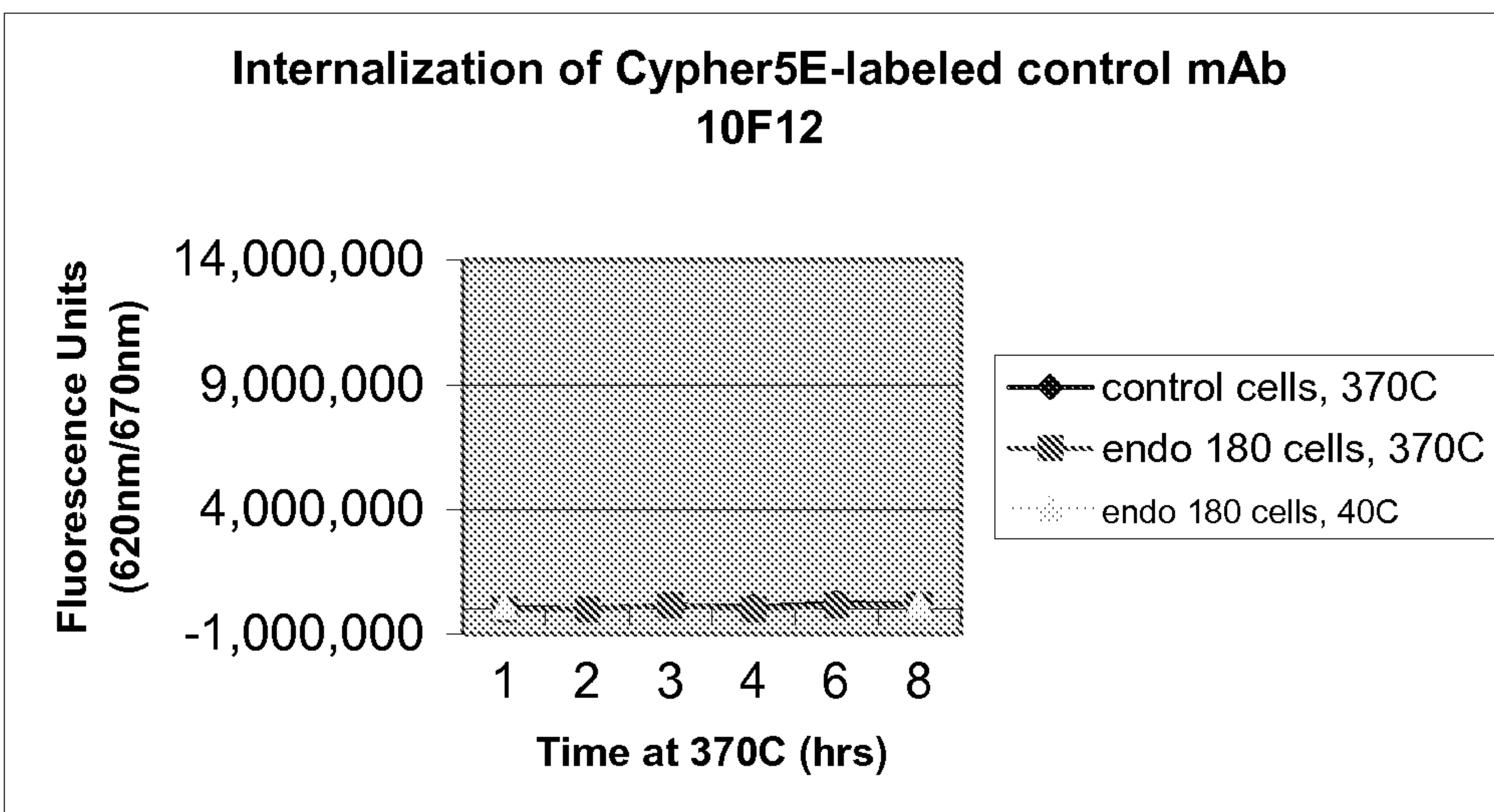
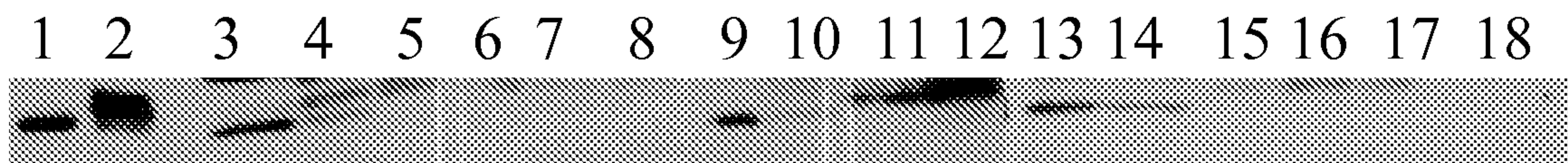


FIGURE 3:



1. E3 8D8-Biotin
2. NMIgG-Biotin
3. Mouse #26 injected with E3 8D8-Biotin, UUO kidney
4. Mouse #26 injected with E3 8D8-Biotin, Contra-lateral kidney
5. Mouse #38 injected with NMIgG-Biotin, UUO kidney
6. Mouse #38 injected with NMIgG-Biotin, Contra-lateral kidney
7. Mouse #50 injected with PBS, UUO kidney
8. Mouse #50 injected with PBS, Contra-lateral kidney
9. Mouse #33 injected with E3 8D8-Biotin, UUO kidney
10. Mouse #33 injected with E3 8D8-Biotin, Contra-lateral kidney
11. Mouse #40 injected with NMIgG-Biotin, UUO kidney
12. Mouse #40 injected with NMIgG-Biotin, Contra-lateral kidney
13. Mouse #27 injected with E3 8D8-Biotin, UUO kidney
14. Mouse #27 injected with E3 8D8-Biotin, Contra-lateral kidney
15. Mouse #36 injected with NMIgG-Biotin, UUO kidney
16. Mouse #36 injected with NMIgG-Biotin, Contra-lateral kidney
17. Mouse #37 injected with NMIgG-Biotin, UUO kidney
18. Mouse #37 injected with NMIgG-Biotin, Contra-lateral kidney

FIGURE 4A

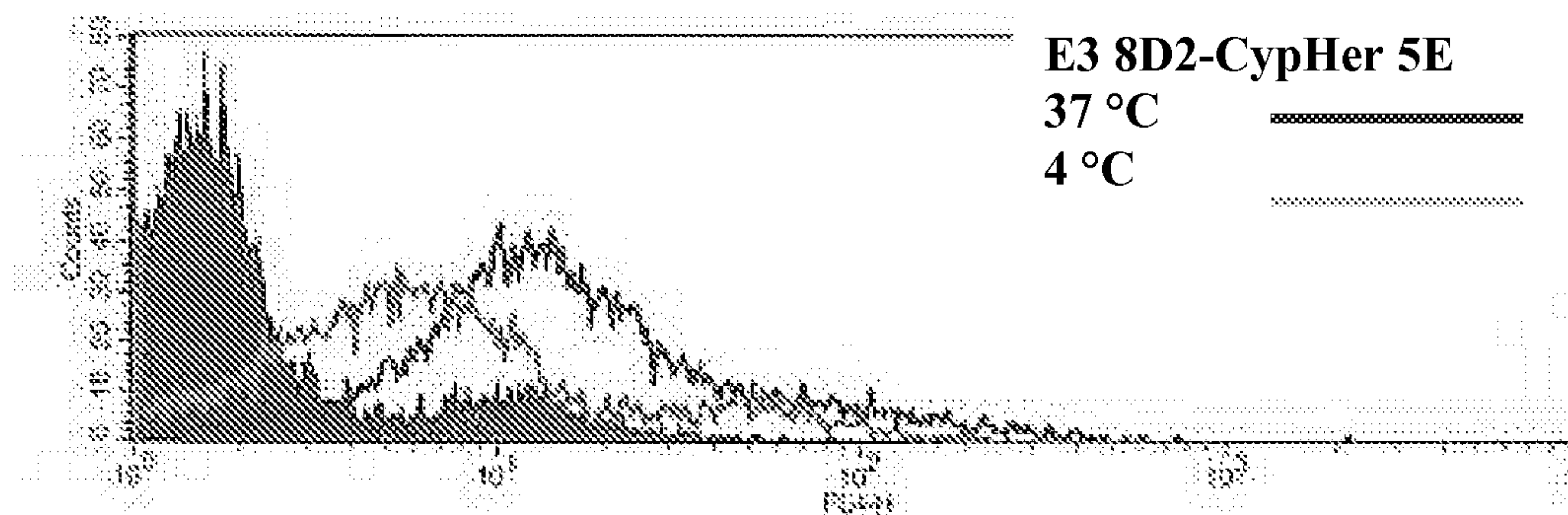
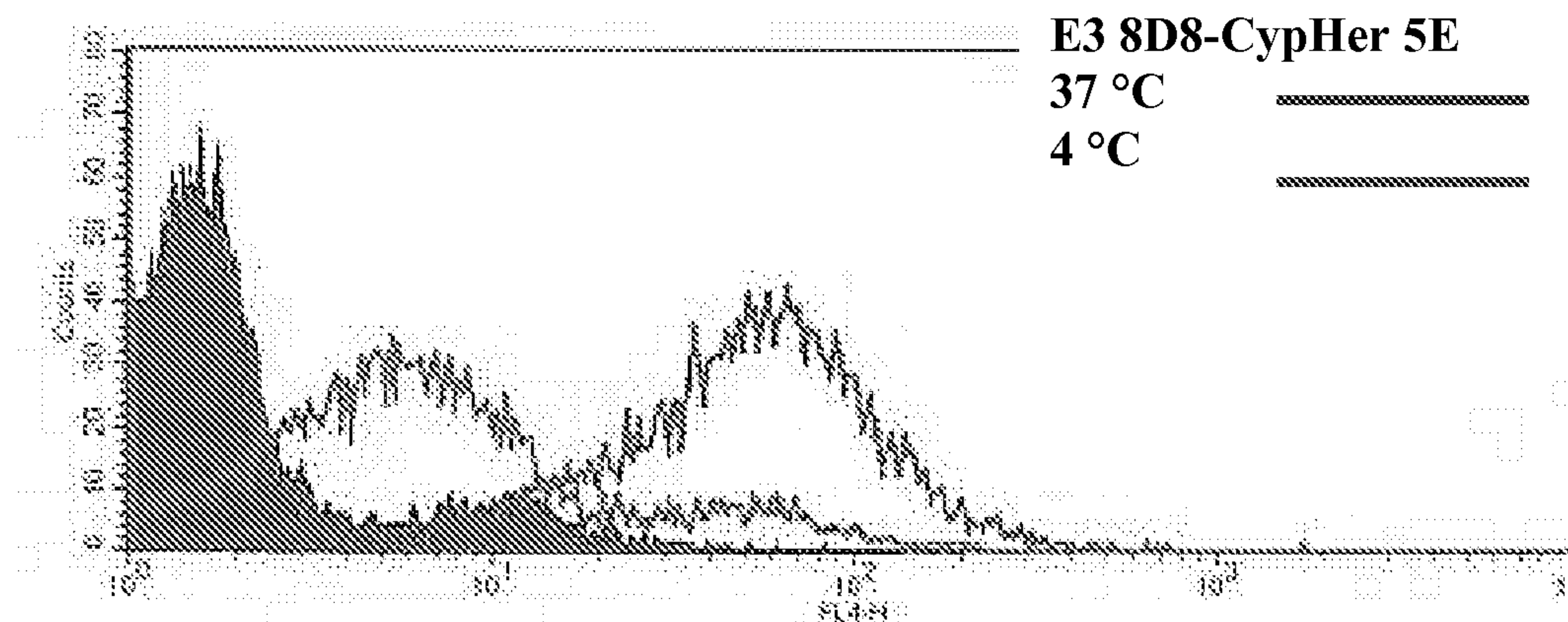
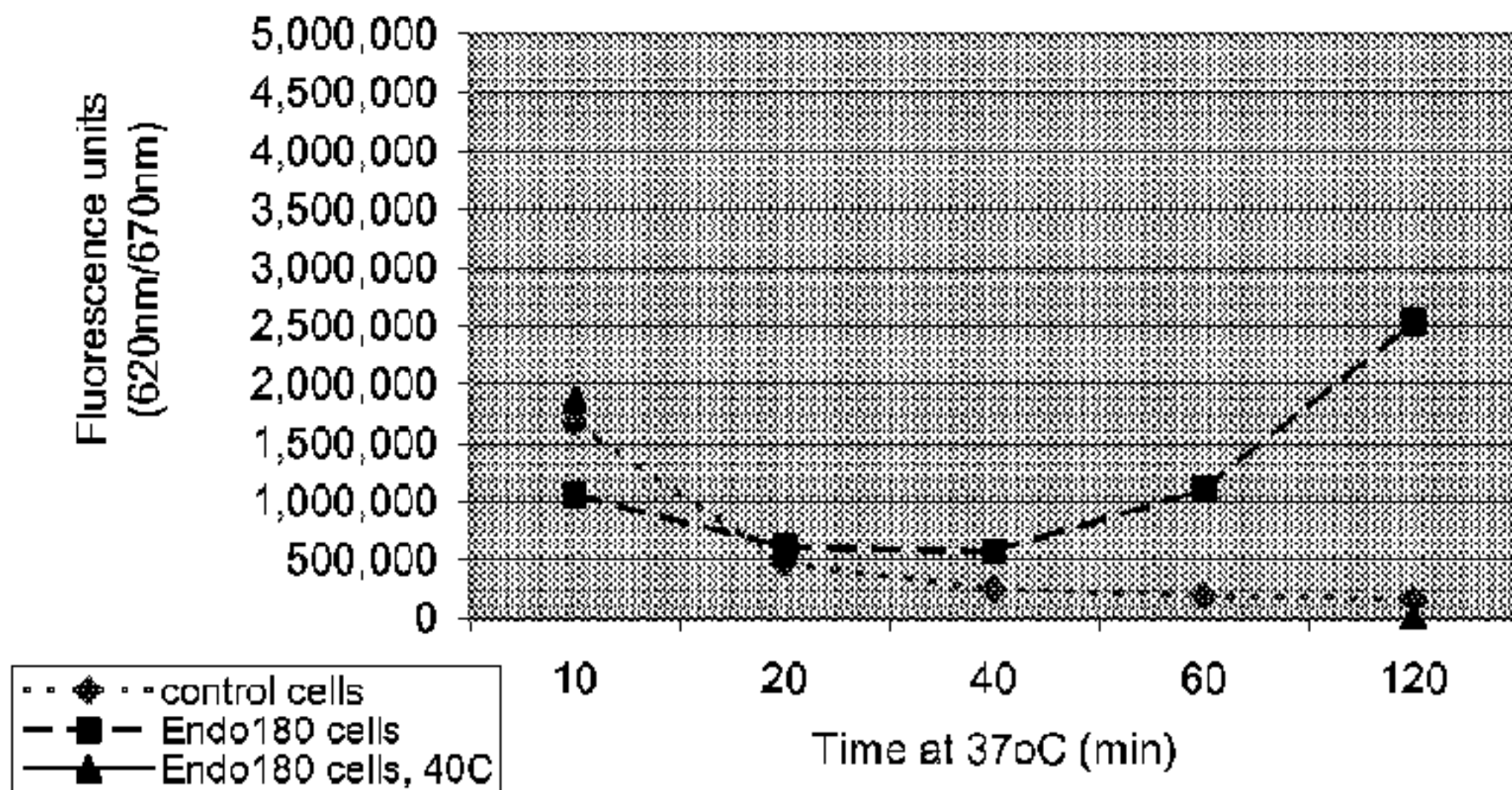


FIGURE 4B



C**Internalization of Cypher5E-labeled
anti-Endo180 mAb 8D8****D****Internalization of Cypher5E-labeled
anti-Endo180 minibodies**