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(19) **United States**(12) **Patent Application Publication**
GENG(10) **Pub. No.: US 2022/0073591 A1**(43) **Pub. Date: Mar. 10, 2022**(54) **RECOMBINANT CLUSTERIN AND USE
THEREOF IN THE TREATMENT AND
PREVENTION OF DISEASE**(60) Provisional application No. 62/087,364, filed on Dec.
4, 2014.(71) Applicant: **BOARD OF REGENTS OF THE
UNIVERSITY OF TEXAS SYSTEM,**
Austin, TX (US)**Publication Classification**(51) **Int. Cl.**
C07K 14/775 (2006.01)
A61P 9/10 (2006.01)(72) Inventor: **Yong-Jian GENG,** Houston, TX (US)(52) **U.S. Cl.**
CPC **C07K 14/775** (2013.01); **A61P 9/10**
(2018.01); **A61K 9/19** (2013.01); **C07K**
2319/00 (2013.01); **C07K 2319/50** (2013.01);
C07K 2319/21 (2013.01)(73) Assignee: **BOARD OF REGENTS OF THE
UNIVERSITY OF TEXAS SYSTEM,**
Austin, TX (US)(21) Appl. No.: **17/406,051**(57) **ABSTRACT**(22) Filed: **Aug. 18, 2021****Related U.S. Application Data**(63) Continuation of application No. 16/597,400, filed on
Oct. 9, 2019, now Pat. No. 11,104,719, which is a
continuation of application No. 14/957,030, filed on
Dec. 2, 2015, now Pat. No. 10,464,994.Recombinant clusterin polypeptides and compositions com-
prising the same are provided. In some aspects, recombinant
clusterin or nucleic acids encoding the same may be used for
treating and preventing an abnormality of morphology and
function in a mammal with disease (e.g., cardiovascular
diseases or alcoholism).**Specification includes a Sequence Listing.**

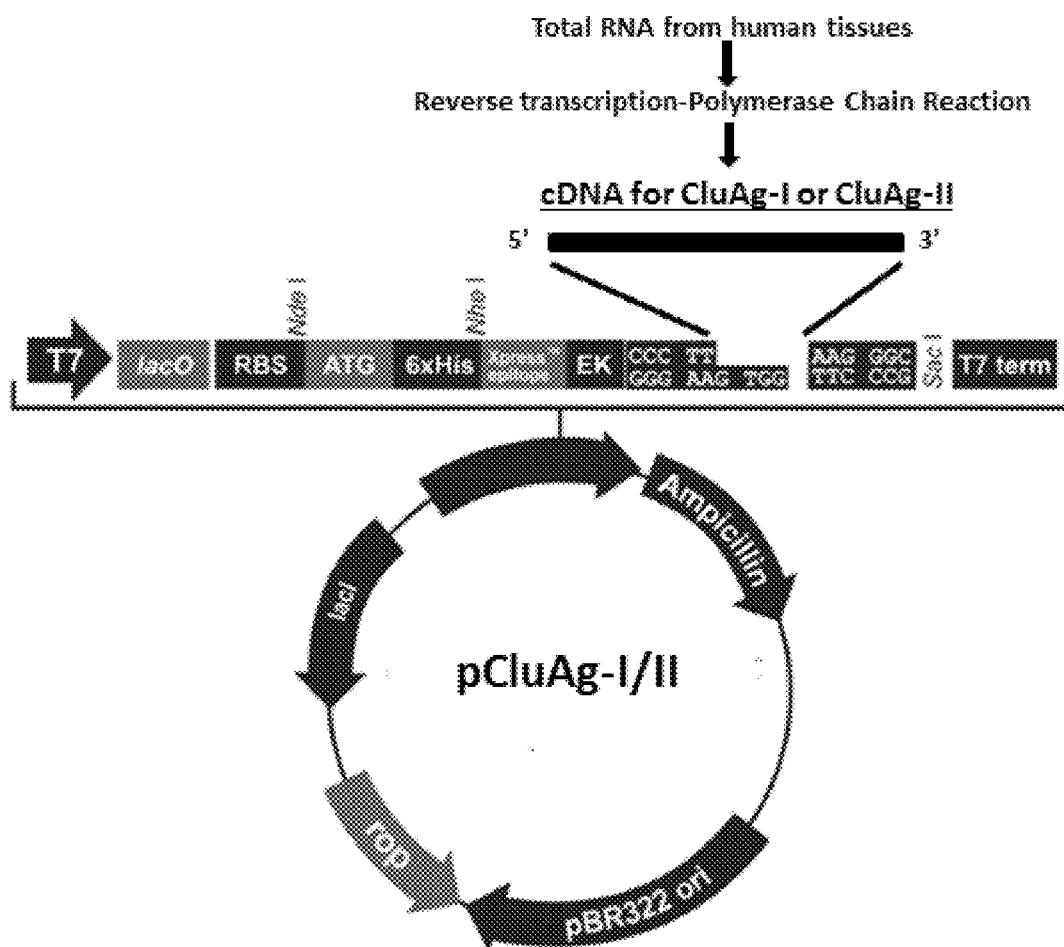


FIG. 1

CluAg-a (Clusterin) – SEQ ID NO: 1

MMKTL~~LLFVGLLT~~WESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSN~~LEEA~~KKKKEDALN
ETRESETKLKELPGVNCNETMMALWEECKPCLKQTCMKFYARVCRSGGLVGRQLEEFNQSSPFYFWMNGDRIDSLENDRQQTH
MLDVMQDHFSSRASSIIDELFQDRFFTREPDQTYHYLPFSLPHRRPHFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQA
MDIHFSAPAFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSDCSTNNPSQAKLRREDESQVAERLTKYNEL
LKSQWKMLNTSSILLEQNEQFNWVSRLANLTQGEDQY~~YLRVTT~~VASHTSDSDVPSGVTEVV~~KL~~FSDSDPITVTPVEVSRKNPKF
METVAEKALQEYRKKHREE

CluAg-b (Clusterin- Δ TMD) – SEQ ID NO: 2

TWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSN~~LEEA~~KKKKEDALNRESETKLKELPG
VCNETMMALWEECKPCLKQTCMKFYARVCRSGGLVGRQLEEFNQSSPFYFWMNGDRIDSLENDRQQTHMLDVMQDHFSSR
ASSIIDELFQDRFFTREPDQTYHYLPFSLPHRRPHFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQAMDIFHSPAFQHP
PTFEIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSDCSTNNPSQAKLRREDESQVAERLTKYNELLKSYQWKMLNTSS
LLEQLNEQFNWVSRLANLTQGEDQY~~YLRVTT~~VASHTSDSDVPSGVTEVV~~KL~~FSDSDPITVTPVEVSRKNPKFMETVAEKALQEYR
KKHREE

CluAg-c (Clusterin- Δ TMD- Δ NLS) – SEQ ID NO: 3

TWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNEDALNRESETKLKELPGVCNETM
MALWEECKPCLKQTCMKFYARVCRSGGLVGRQLEEFNQSSPFYFWMNGDRIDSLENDRQQTHMLDVMQDHFSSRASSIIDEL
FQDRFFTREPDQTYHYLPFSLPHRRPHFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQAMDIFHSPAFQHPPTFEIRE
GDDDRTVCREIRHNSTGCLRMKDQCDKCREILSDCSTNNPSQAKLRREDESQVAERLTKYNELLKSYQWKMLNTSSLLEQLNE
QFNWVSRLANLTQGEDQY~~YLRVTT~~VASHTSDSDVPSGVTEVV~~KL~~FSDSDPITVTPVEVSRKNPKFMETVAEKALQEYRKKHREE

FIG. 2A

CluAg-Ia (HisTR-Clusterin) – SEQ ID NO: 4
HHHHHHLVPRGSMIMKTLLFVGLLTTWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSN
LEFAKKKKEDALNETRESETKLKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMINGDR
IDSLLENDRQQTHMLDVMQDHFSSRASSIIDELFQDRFFTRPQDITYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMIFQ
PFLEMIHEAQQAMDIHFHSPAFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSVDCSTNNPSQAKLRRELD
QVAERLTRKYNELLKSYQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITV
TVPVEVSRKNPKFMETVAEKALQEYRKKHREE

CluAg-Ib (HisTR-Clusterin-ΔTMD) – SEQ ID NO: 5
HHHHHHLVPRGSTWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNLEFAKKKKEDALN
ETRESETKLKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLENDRQQTH
MLDVMQDHFSSRASSIIDELFQDRFFTRPQDITYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMIFQPFLEMIHEAQQ
MDIHFHSPAFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSVDCSTNNPSQAKLRRELD
LKSQYQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITVTPVEVSRKNPKF
METVAEKALQEYRKKHREE

CluAg-Ic (HisTR-Clusterin-ΔTMD-ΔNLS) – SEQ ID NO: 6
HHHHHHLVPRGSTWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNEDALNETRESETK
LKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLENDRQQTHMLDVMQ
DHFSSRASSIIDELFQDRFFTRPQDITYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMIFQPFLEMIHEAQQAMDIHFHSP
AFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSVDCSTNNPSQAKLRRELD
MLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITVTPVEVSRKNPKFMETVAEK
ALQEYRKKHREE

FIG. 2B

CluAg-IIa (Clusterin-TRHis) – SEQ ID NO: 7
MMKTLLFVGLLTWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNEEAKKKKEDALN
ETRESETKLKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNQSSPFYFWMNGDRIDSLLENDRQQTH
MLDVMQDHFSAQHPPTFIREGDDRTVCREIRHNSTGCLRMKDQCCKREILSDCSTNNPSQAKLRREDESQVAERLTRKYNEL
LKSQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFSDDPITVTPVEVSRKNPKF
METVAEKALQEYRKKHREELVPRGSHHHHHH

CluAg-IIb (Clusterin-ΔTMD-TRHis) – SEQ ID NO: 8
TWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNEEAKKKKEDALNETRESETKLKELPG
VCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNQSSPFYFWMNGDRIDSLLENDRQQTHMLDVMQDHFSAQ
ASSIIDELFQDRFFFTREPQDTHYLPFSLPHRRPHFFPKSRIVRSLMPFSPYEPNLFHAMFQPFLEMIHEAQQAMDIFHSPAFQHP
PTEFIREGDDRTVCREIRHNSTGCLRMKDQCCKREILSDCSTNNPSQAKLRREDESQVAERLTRKYNELLSYQWKMLNTSS
LLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFSDDPITVTPVEVSRKNPKFMETVAEKALQEYR
KKHREELVPRGSHHHHHH

CluAg-IIc (Clusterin-ΔTMD-ΔNLS-TRHis) – SEQ ID NO: 9
MMKTTWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNEDALNETRESETKLKELPGVC
NETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNQSSPFYFWMNGDRIDSLLENDRQQTHMLDVMQDHFSAQ
SIIDELFQDRFFFTREPQDTHYLPFSLPHRRPHFFPKSRIVRSLMPFSPYEPNLFHAMFQPFLEMIHEAQQAMDIFHSPAFQHPPT
EFIREGDDRTVCREIRHNSTGCLRMKDQCCKREILSDCSTNNPSQAKLRREDESQVAERLTRKYNELLSYQWKMLNTSSLE
QLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFSDDPITVTPVEVSRKNPKFMETVAEKALQEYRKKH
REELVPRGSHHHHHH

FIG. 2C

CluAg-IIIa (HisEK-Clusterin) – SEQ ID NO: 10
HHHHHHDDDDKMMKTLLEFVGILLTWESGQVLGDQTVSDNELQEMSNQGSKYVNGVKQIKTLIEKTNEERKTLNLS
LEEAKKKKEDALNETRESETKLKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDR
IDSLLENDRQQTHMLDVMQDHFSSASSIIDELFQDRFFTRPQDITYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMFQ
PFLEMIHEAQQAMDIHFHSPAFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSVDCSTNNPSQAKLRRELD
ESLQVAERLTRKYNELLKSYQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITV
TVPVEVSRKNPKFMETVAEKALQEYRKKHREE

CluAg-IIIb (HisEK-Clusterin-ΔTMD) – SEQ ID NO: 11
HHHHHHDDDDKTTWESGQVLGDQTVSDNELQEMSNQGSKYVNGVKQIKTLIEKTNEERKTLNLSLEEAKKKKEDALN
ETRESETKLKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLLENDRQQTH
MLDVMQDHFSSASSIIDELFQDRFFTRPQDITYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQ
AMDIHFHSPAFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSVDCSTNNPSQAKLRRELD
ESLQVAERLTRKYNELLKSYQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITV
TVPVEVSRKNPKFMETVAEKALQEYRKKHREE

CluAg-IIIc (HisEK-Clusterin-ΔNLS-ΔTMD) – SEQ ID NO: 12
HHHHHHDDDDKTTWESGQVLGDQTVSDNELQEMSNQGSKYVNGVKQIKTLIEKTNEERKTLNLSNEDALNETRESETKL
KELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLLENDRQQTHMLDVMQ
DHFSSASSIIDELFQDRFFTRPQDITYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQAMDIHFHSP
AFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCDKCREILSVDCSTNNPSQAKLRRELD
ESLQVAERLTRKYNELLKSYQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITV
TVPVEVSRKNPKFMETVAEKALQEYRKKHREE

FIG. 2D

CluAg-IVa (Clusterin-EKHis) – SEQ ID NO: 13
MMKTL~~LL~~FPVGL~~LL~~TWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSN~~EEAKKKK~~EDALN
ETRESETKLKELPGVCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLLENDRQQTH
MLDVMQDHFSSASSIIDELFQDRFFTREPDQTYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQA
MDIHFSAPAFQHPPTFEFIREGDDRTVCREIRHNSTGCLRMKDQCCKREILSDCSTNPSQAKLRRELDLSQVAERLTRKYNEL
LKSQWKMLNTSSLLEQLNEQFNWVSRLANLTQGEDQYYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITVTPVEVSRKNPKF
METVAEKALQEYRKKHREELDDDDKKHHHHHH

CluAg-IVb (Clusterin-ΔTMD-EKHis) – SEQ ID NO: 14
TWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSN~~EEAKKKK~~EDALNETRESETKLKELPG
VCNETMMALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLLENDRQQTHMLDVMQDHFSS
ASSIIDELFQDRFFTREPDQTYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQA~~MDIHFS~~APAFQHP
PTEFIREGDDRTVCREIRHNSTGCLRMKDQCCKREILSDCSTNPSQAKLRRELDLSQVAERLTRKYNELLSYQWKMLNTSS
LLEQLNEQFNWVSRLANLTQGEDQYYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITVTPVEVSRKNPKFMETVAEKALQEYR
KKHREELDDDDKKHHHHHH

CluAg-IVc (Clusterin-ΔTMD-ΔNLS-EKHis) – SEQ ID NO: 15
TWESGQVLGDQTVSDNELQEMSNQGSKYVNKEIQNAVNGVKQIKTLIEKTNEERKTLLSNEDALNETRESETKLKELPGVCNETM
MALWEECKPCLKQTCMKFYARVCRSGSLVGRQLEEFNLQSSPFYFWMNGDRIDSLLENDRQQTHMLDVMQDHFSSASSIIDEL
FQDRFFTREPDQTYHYLPFSLPHRRPHFFFPKSRIVRSLMPFSPYEPLNFHAMFQPFLEMIHEAQQA~~MDIHFS~~APAFQHPPTFEFIRE
GDDRTVCREIRHNSTGCLRMKDQCCKREILSDCSTNPSQAKLRRELDLSQVAERLTRKYNELLSYQWKMLNTSSLLEQLNE
QFNWVSRLANLTQGEDQYYLRVTTVASHTSDSDVPSGVTEVVVKLFDSDPITVTPVEVSRKNPKFMETVAEKALQEYRKKHREEL
DDDDKKHHHHHH

FIG. 2E

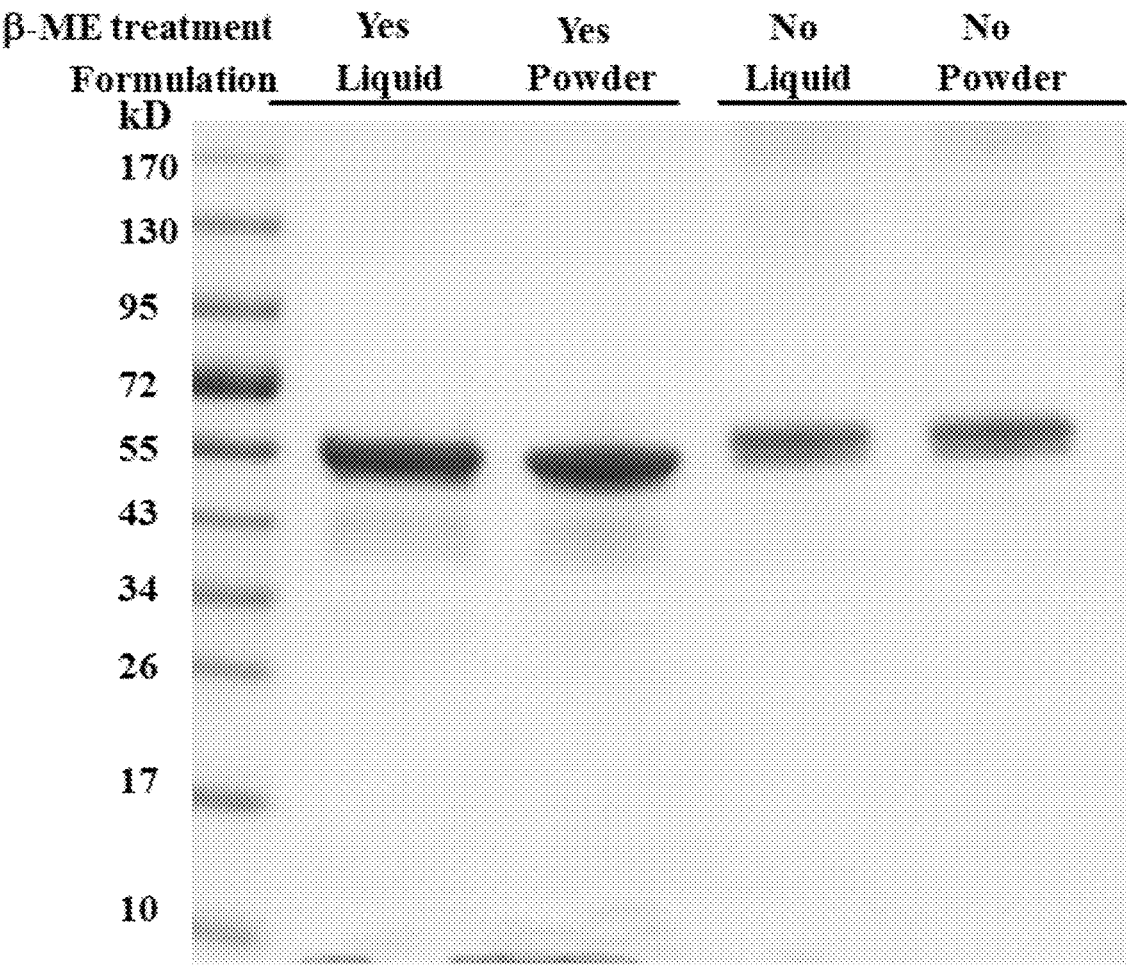


FIG. 3

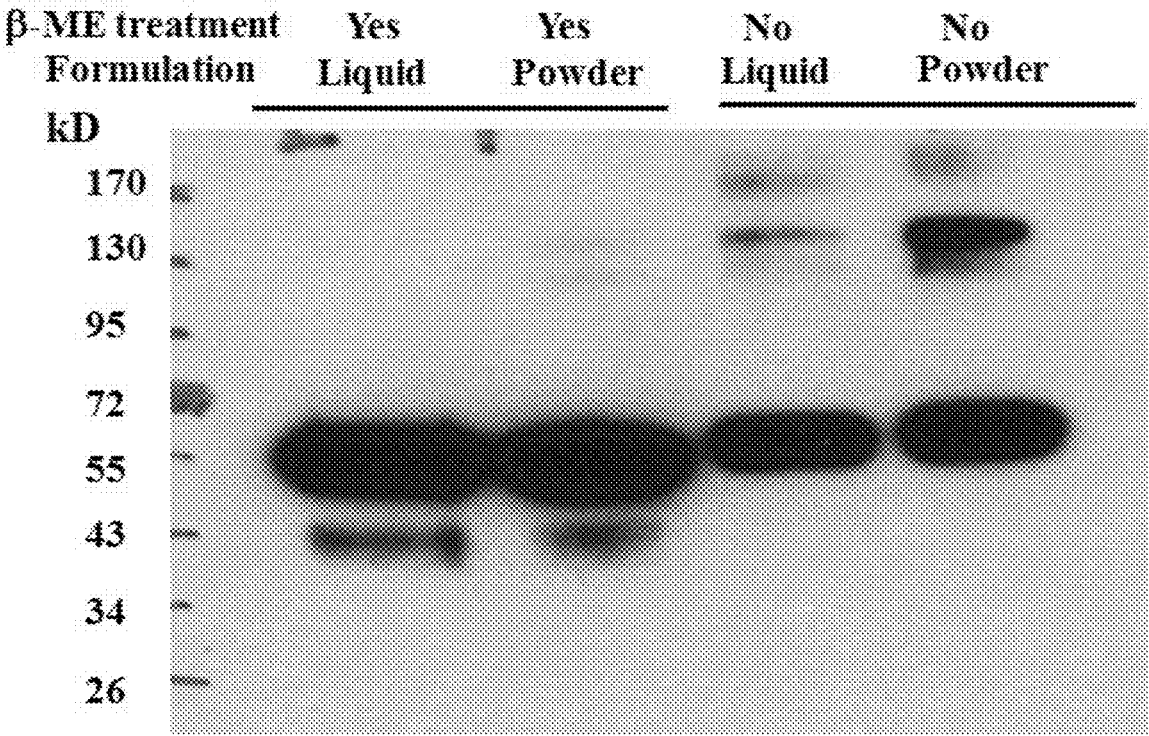
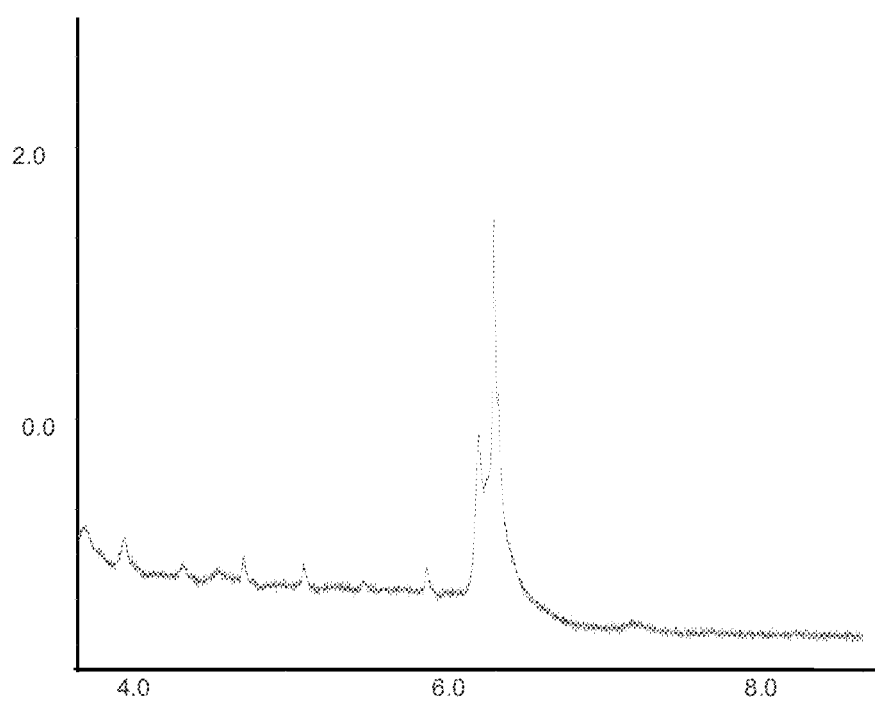
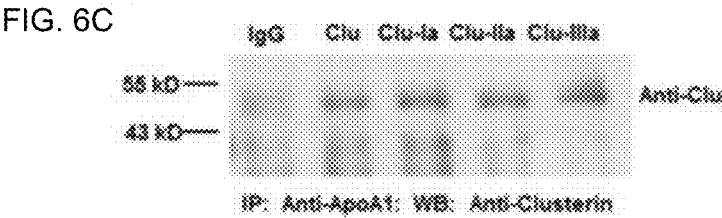
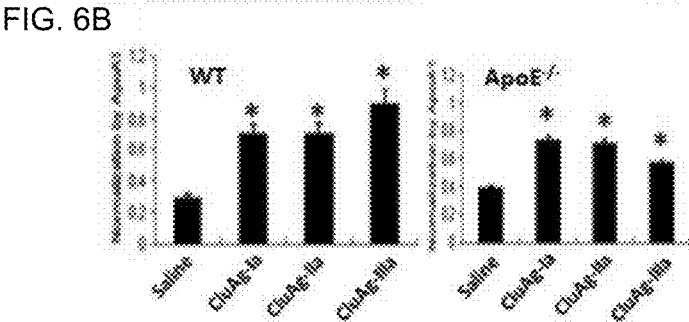
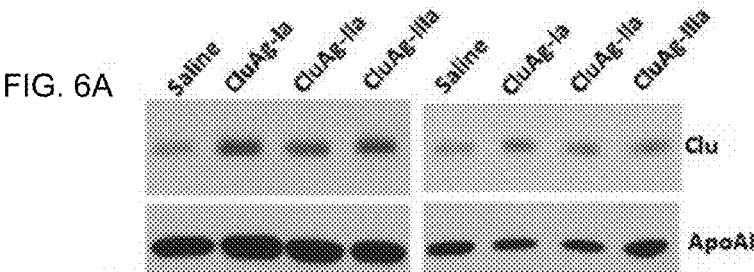


FIG. 4

**FIG. 5**



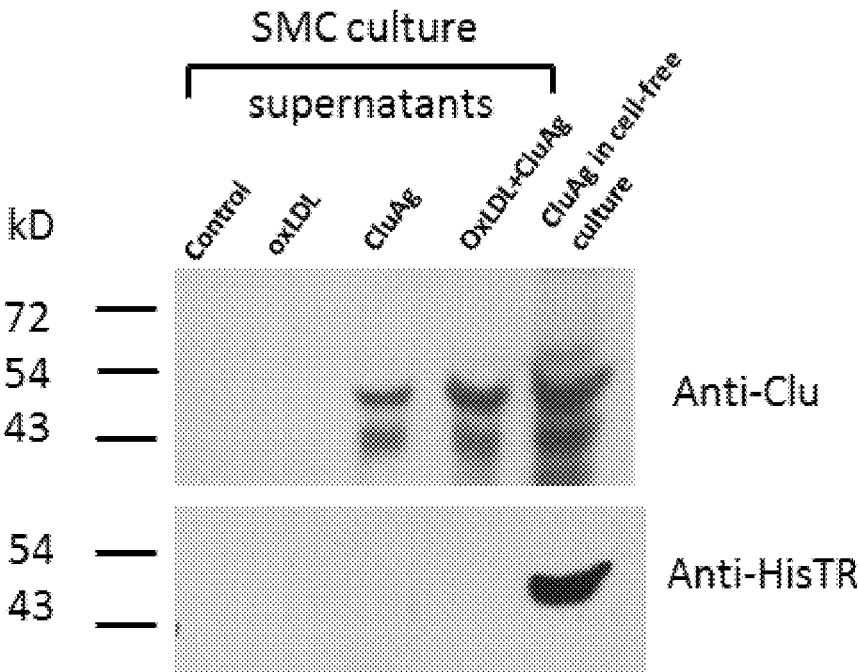


FIG. 7

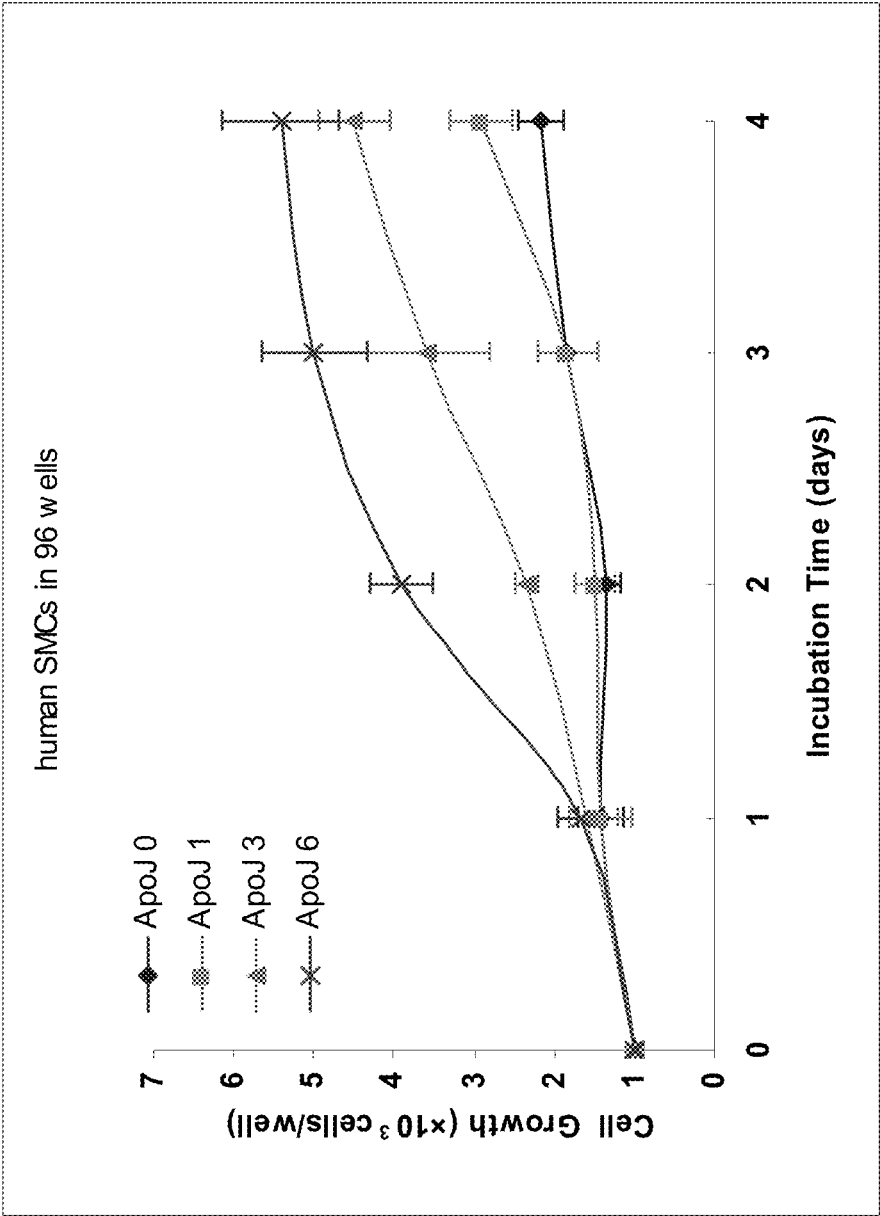


FIG. 8

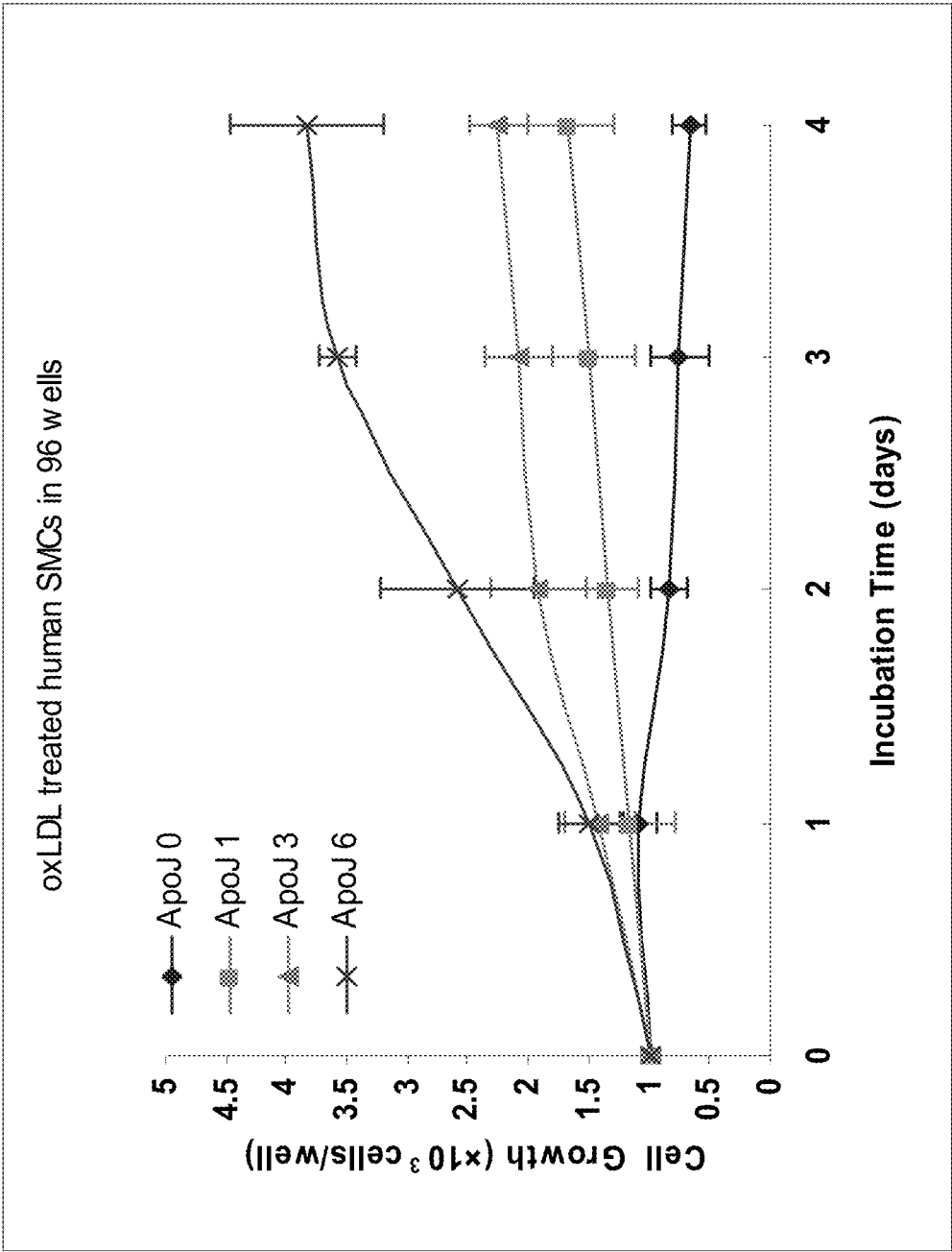


FIG. 9

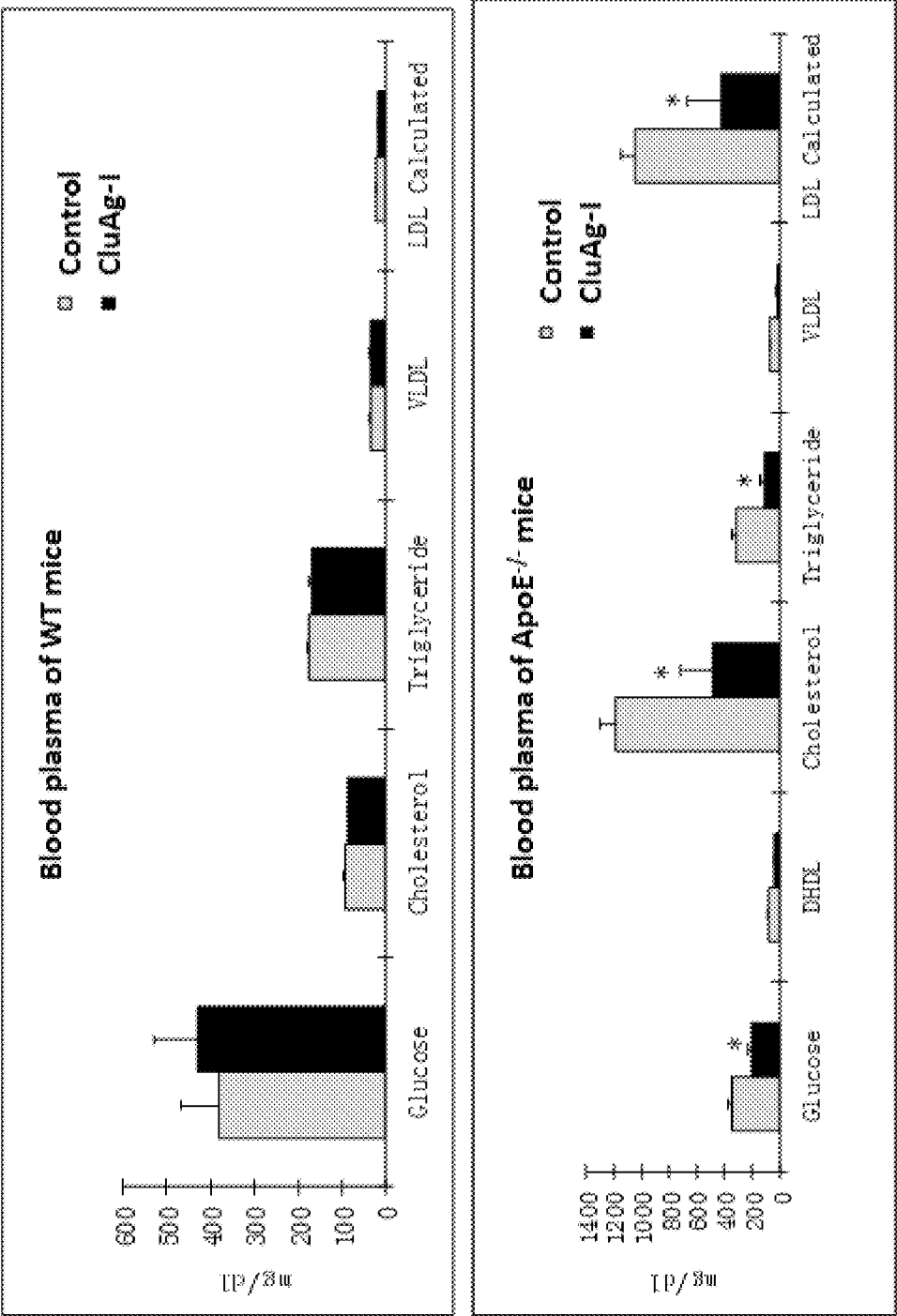


FIG. 10

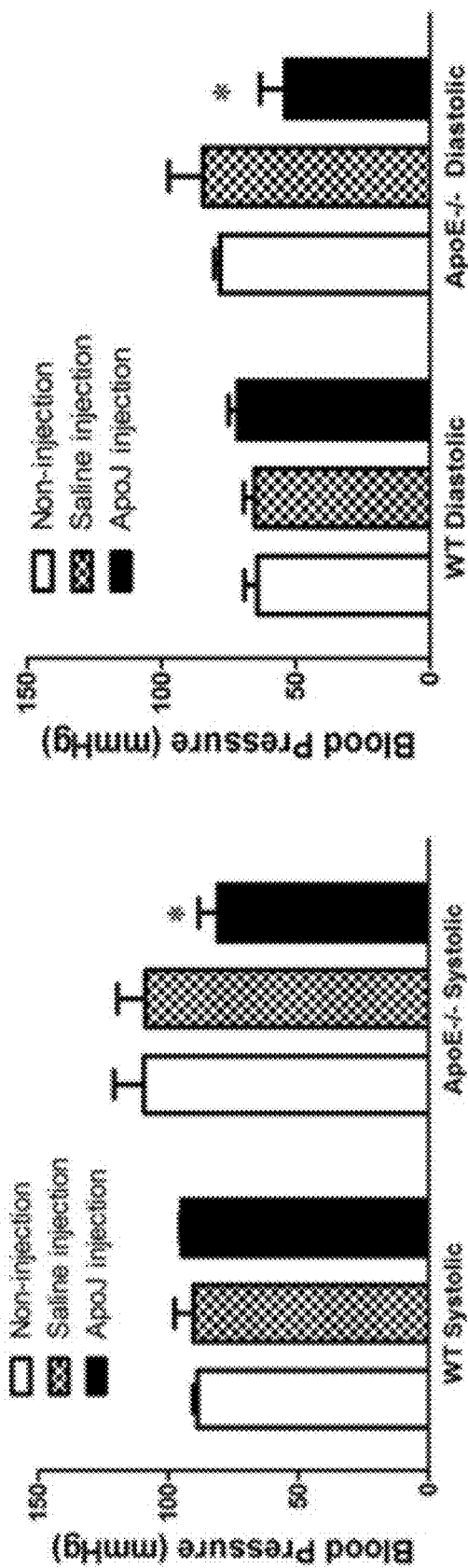


FIG. 11

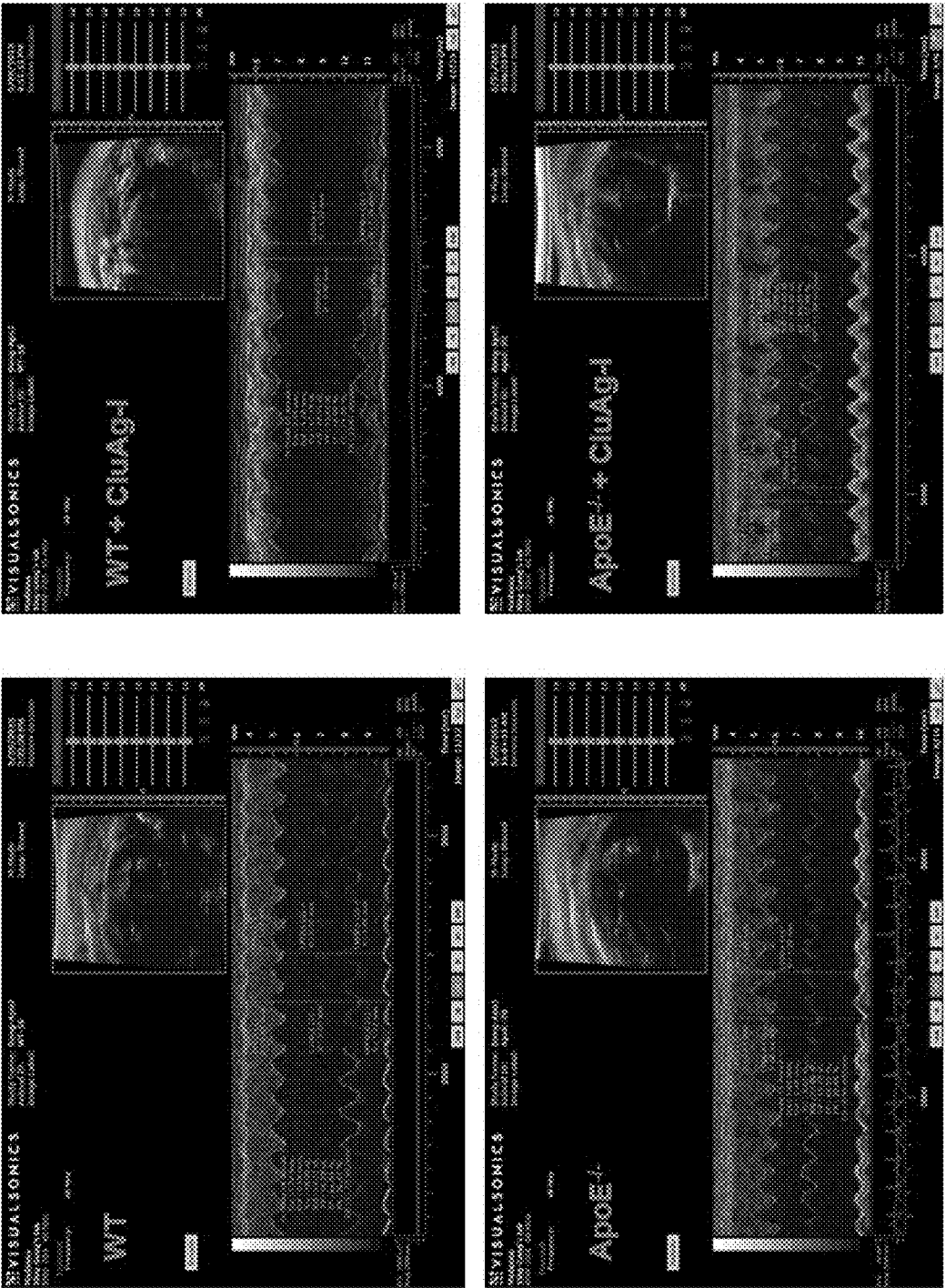


FIG. 12

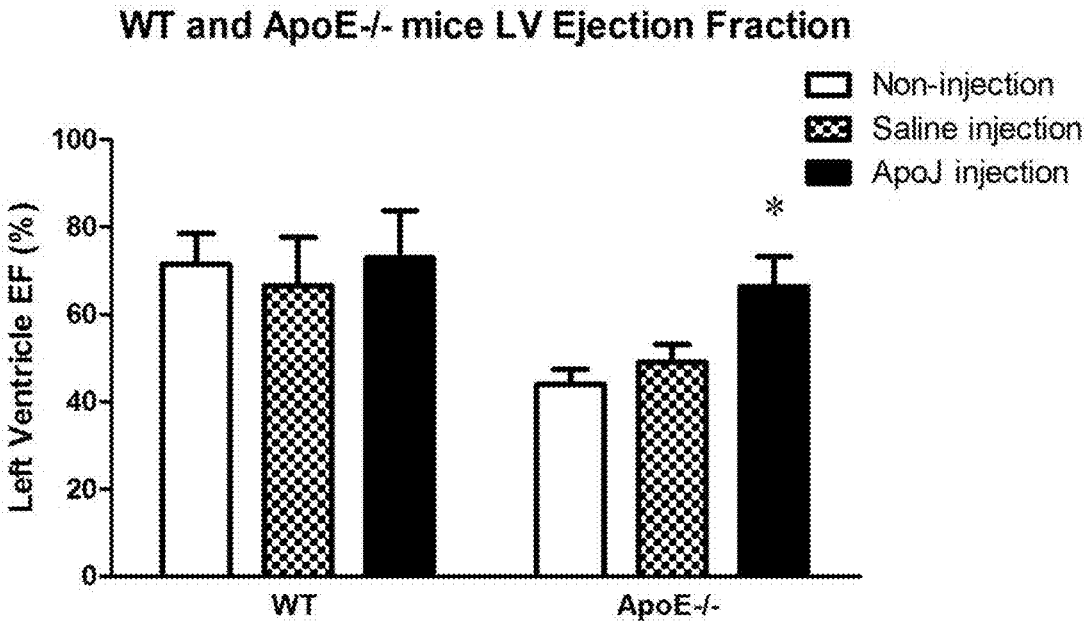
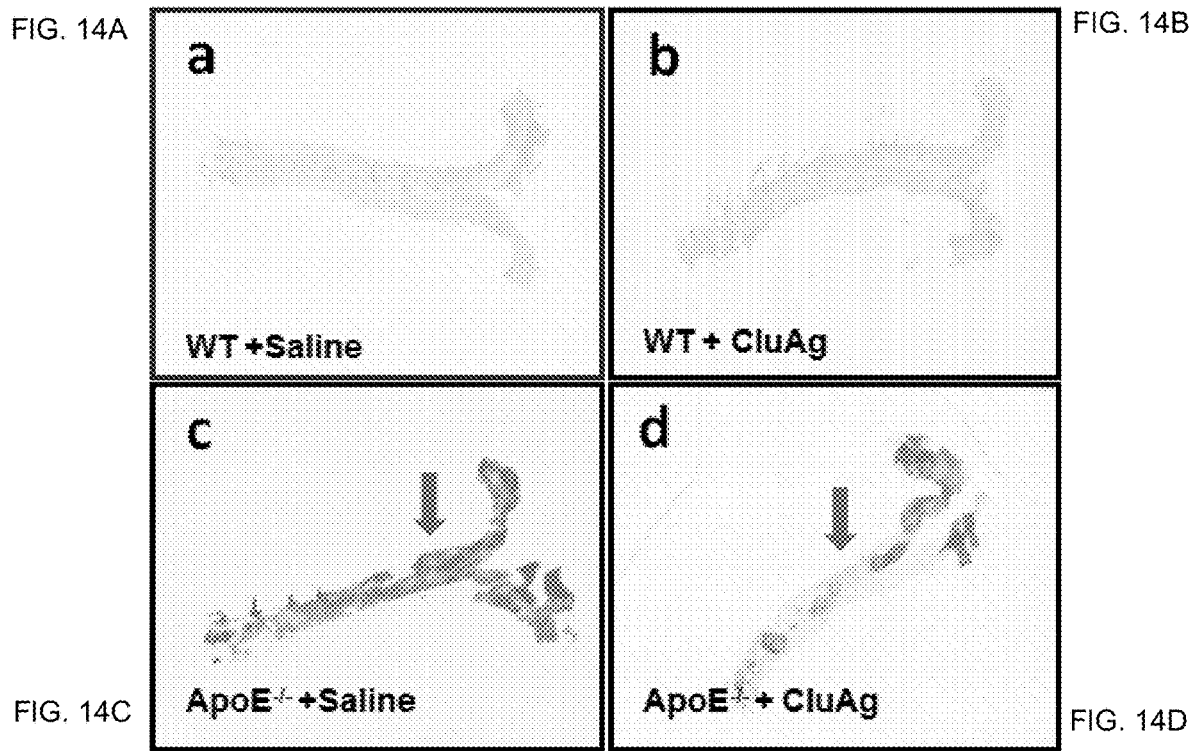


FIG. 13



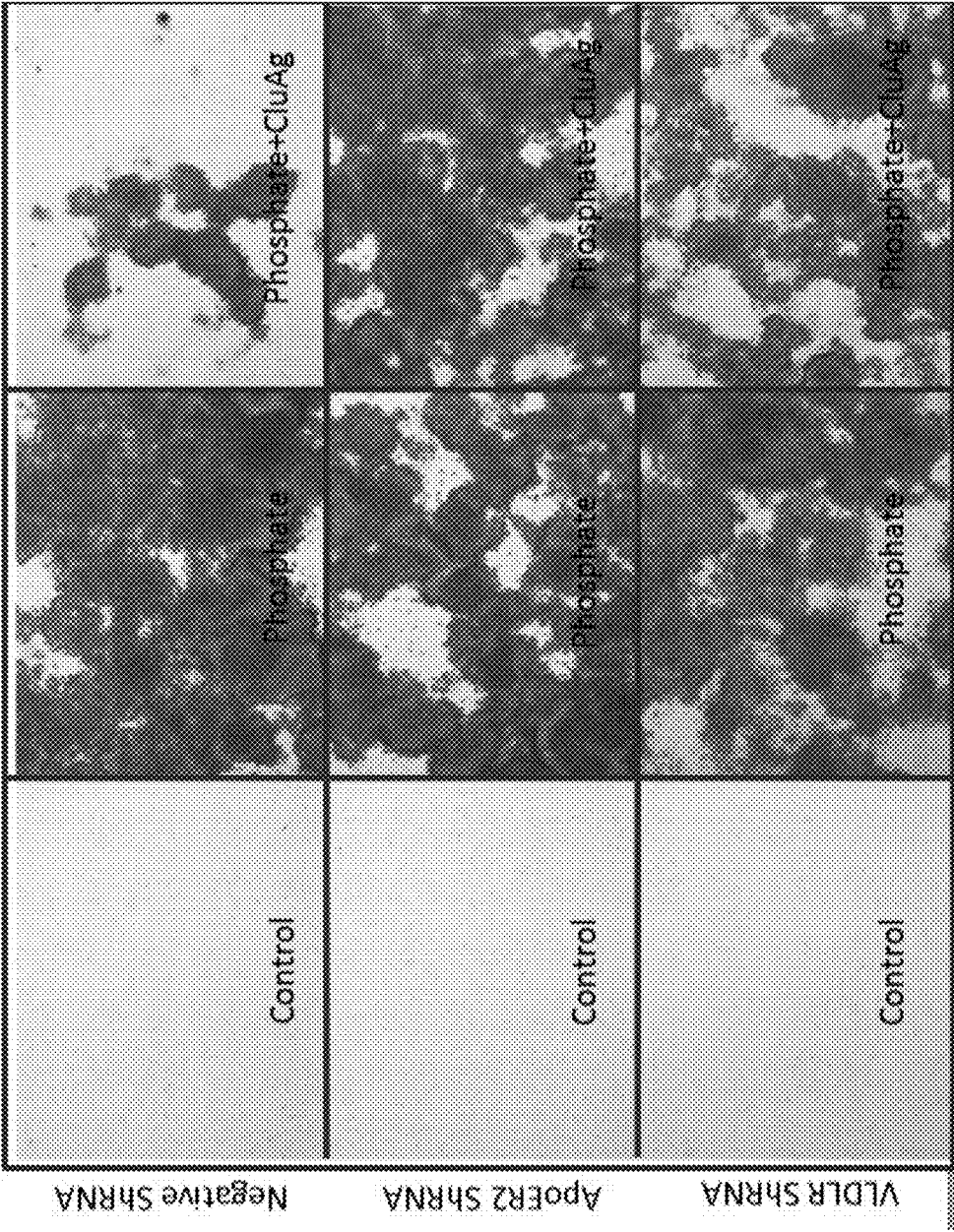
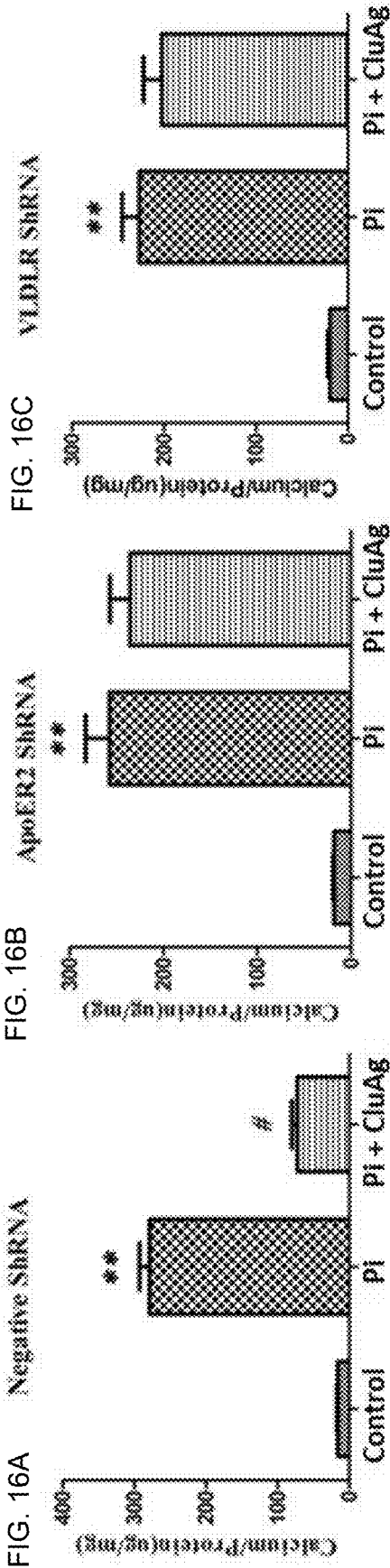
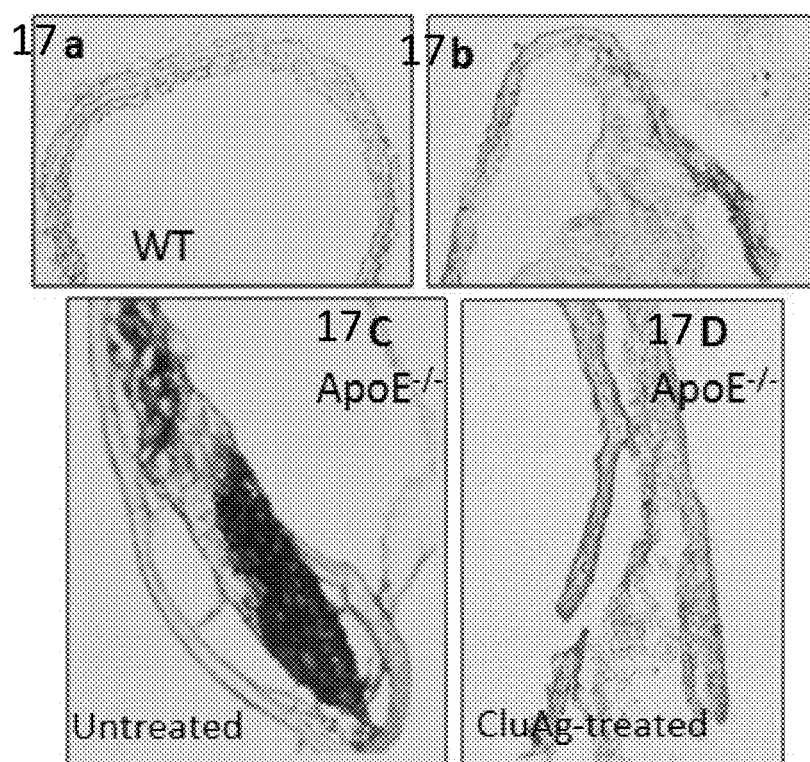
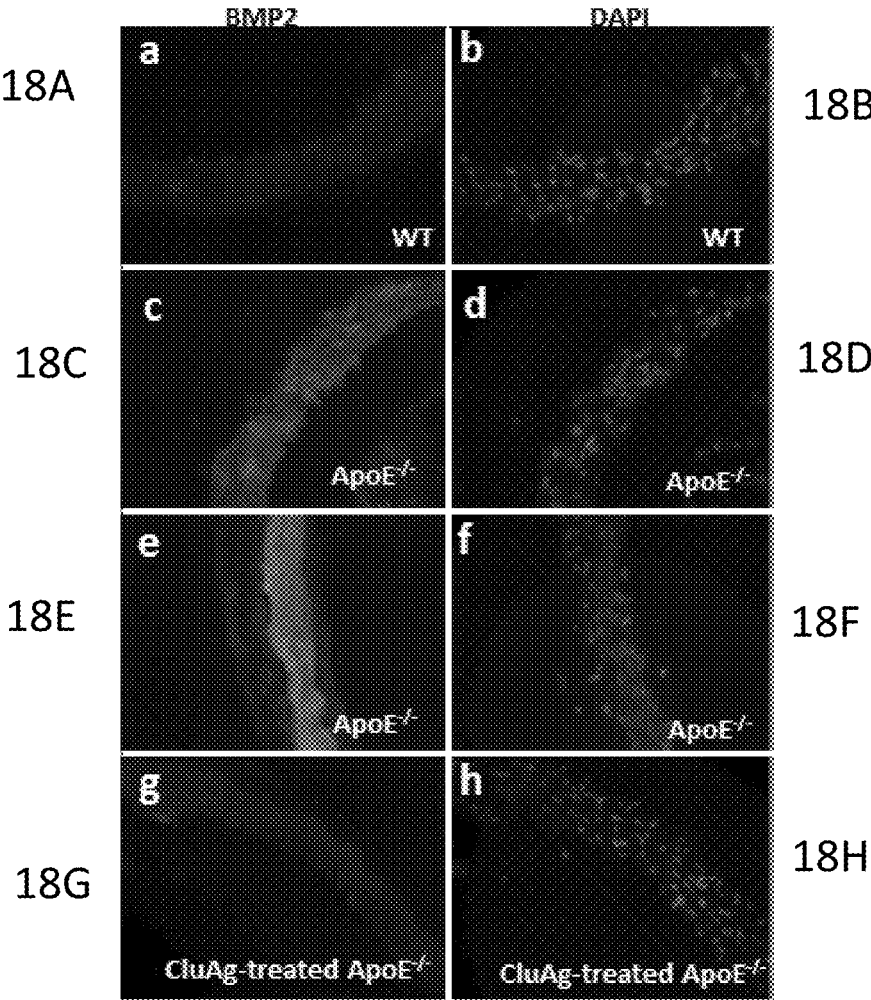


FIG. 15





FIGS. 17A-17D



FIGS. 18A-18H

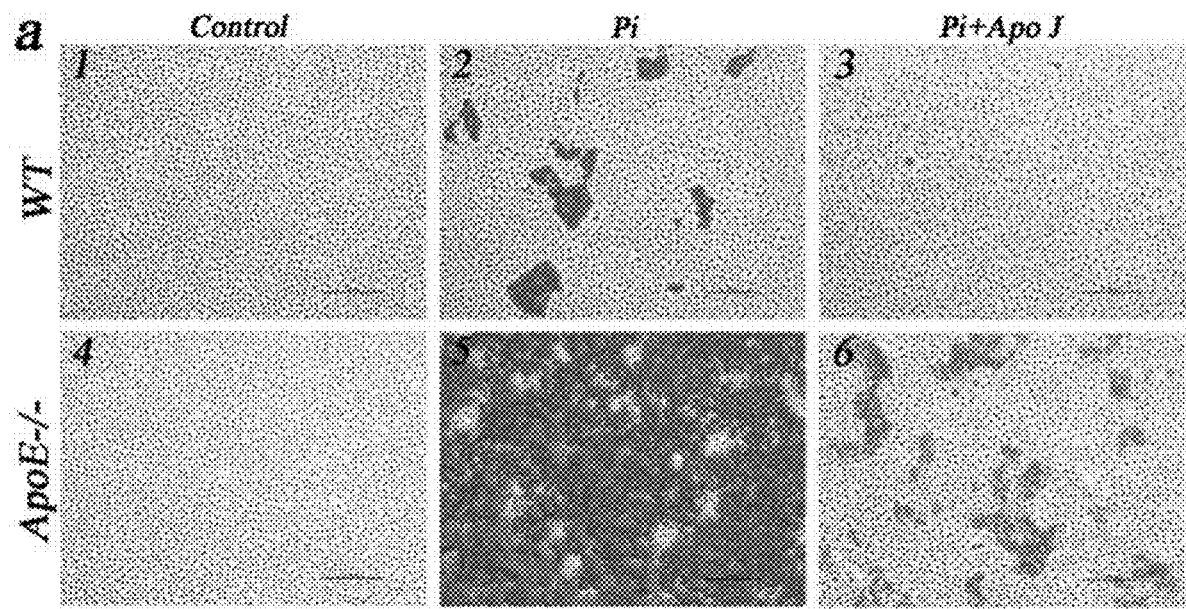


FIG. 19A

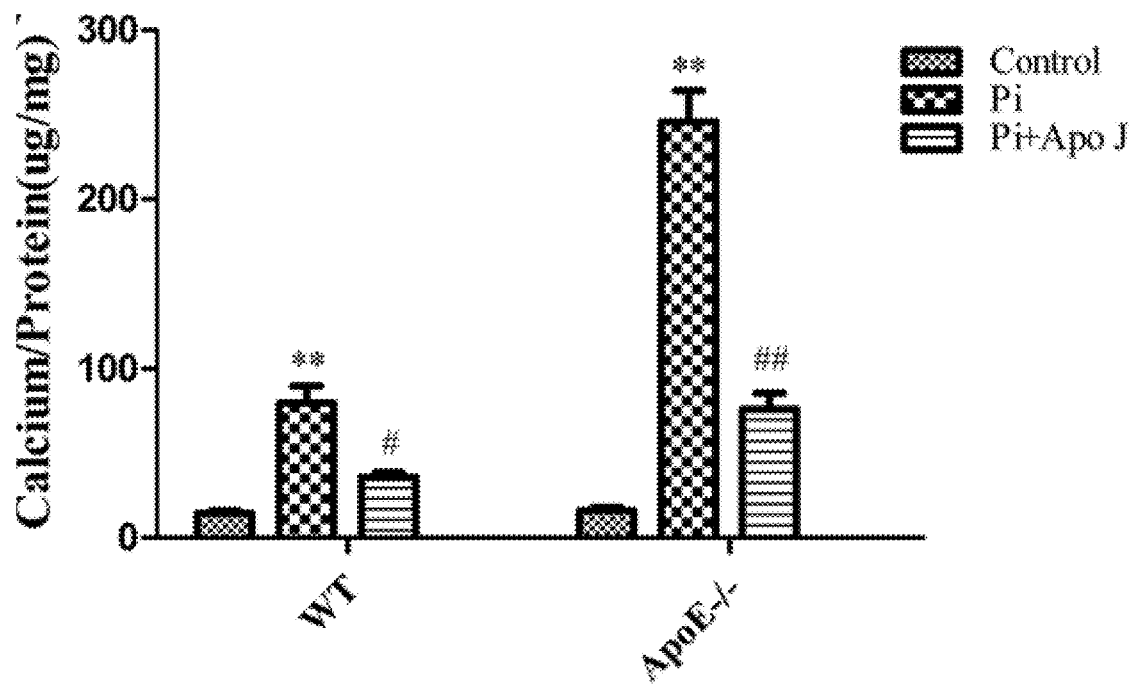


FIG. 19B

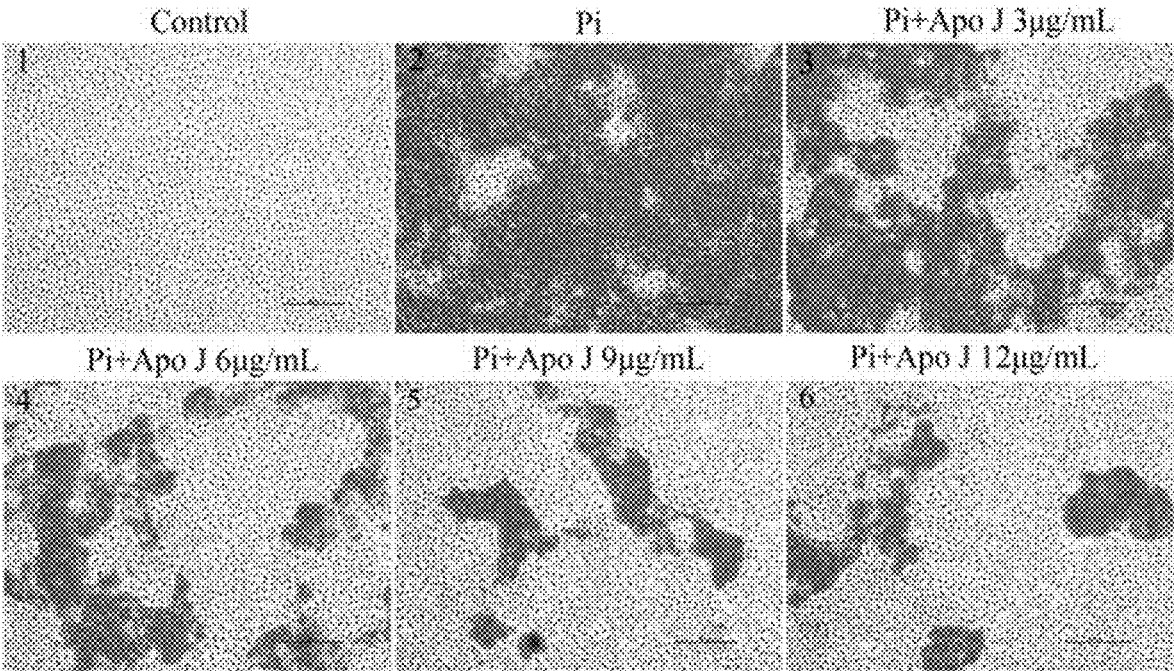


FIG. 20A

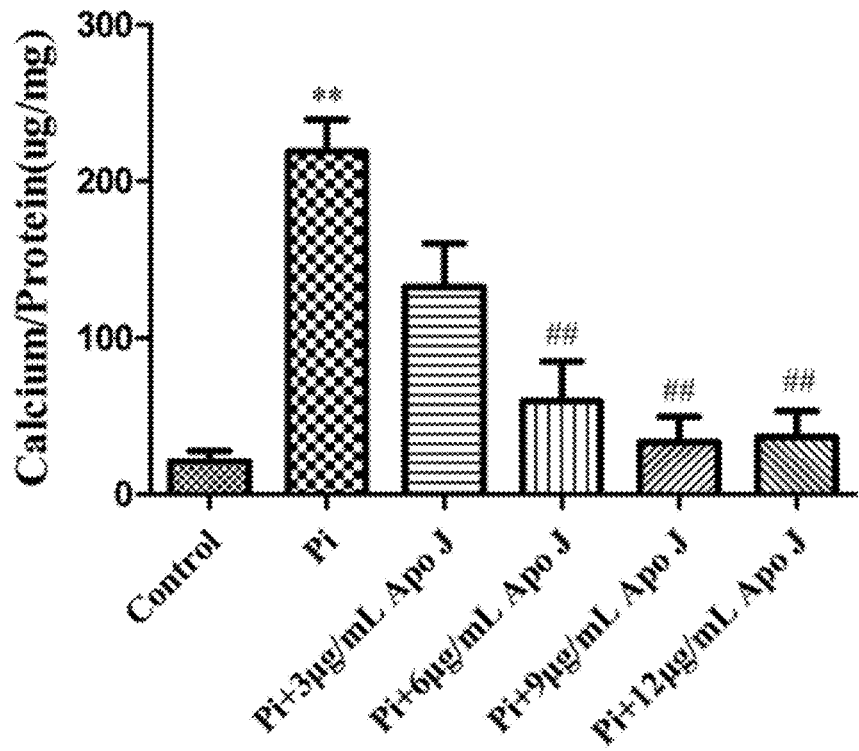
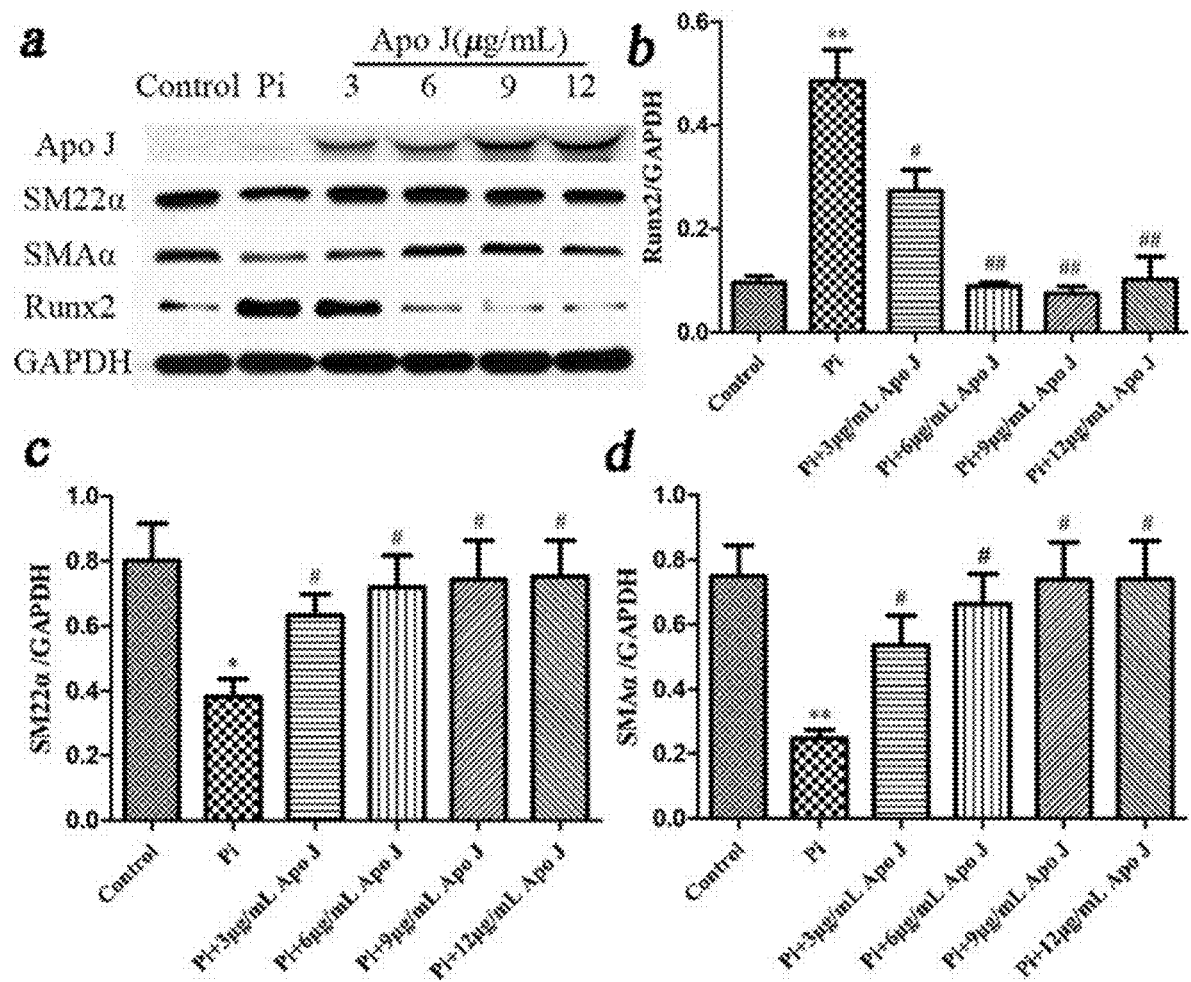
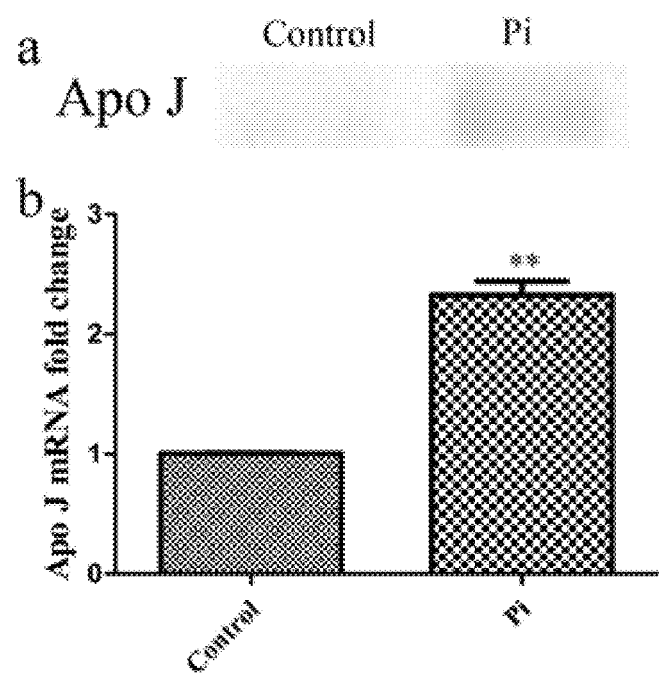


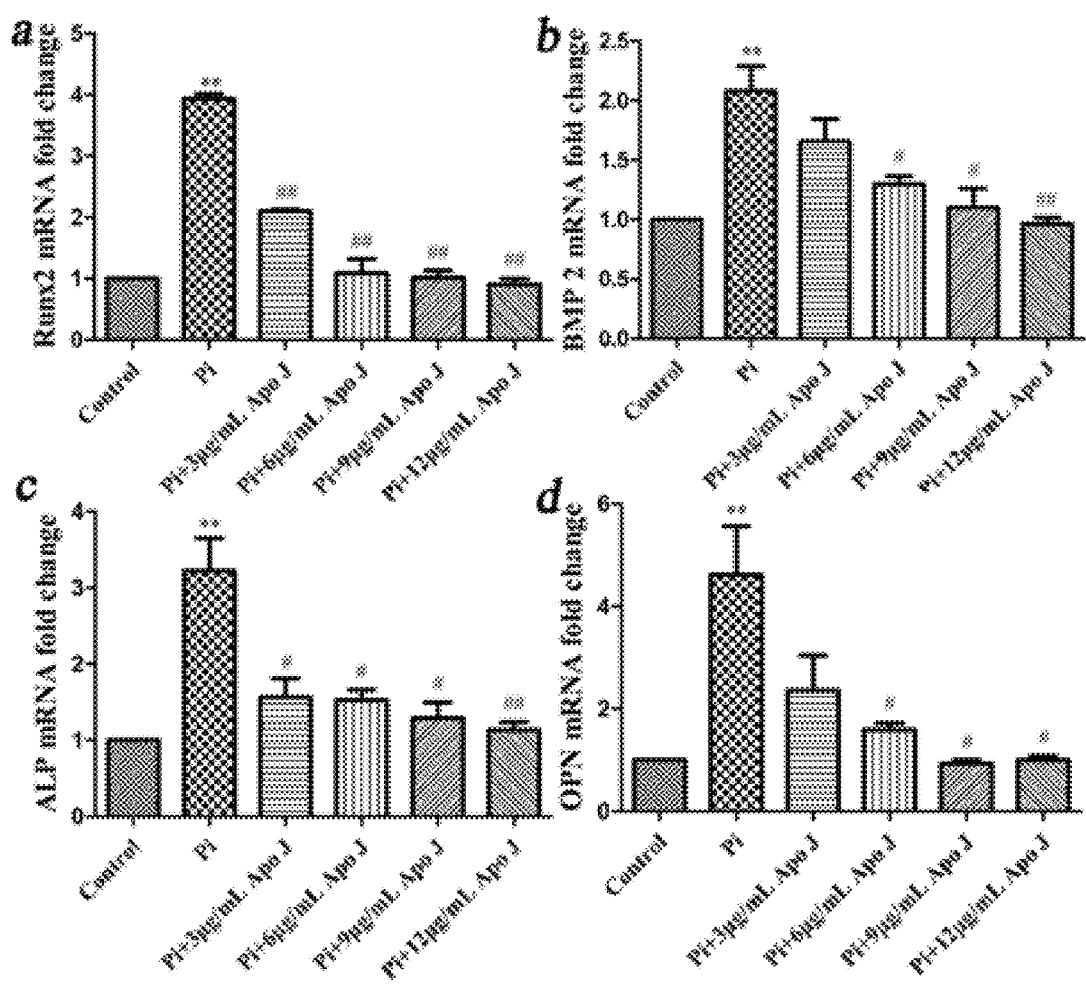
FIG. 20B



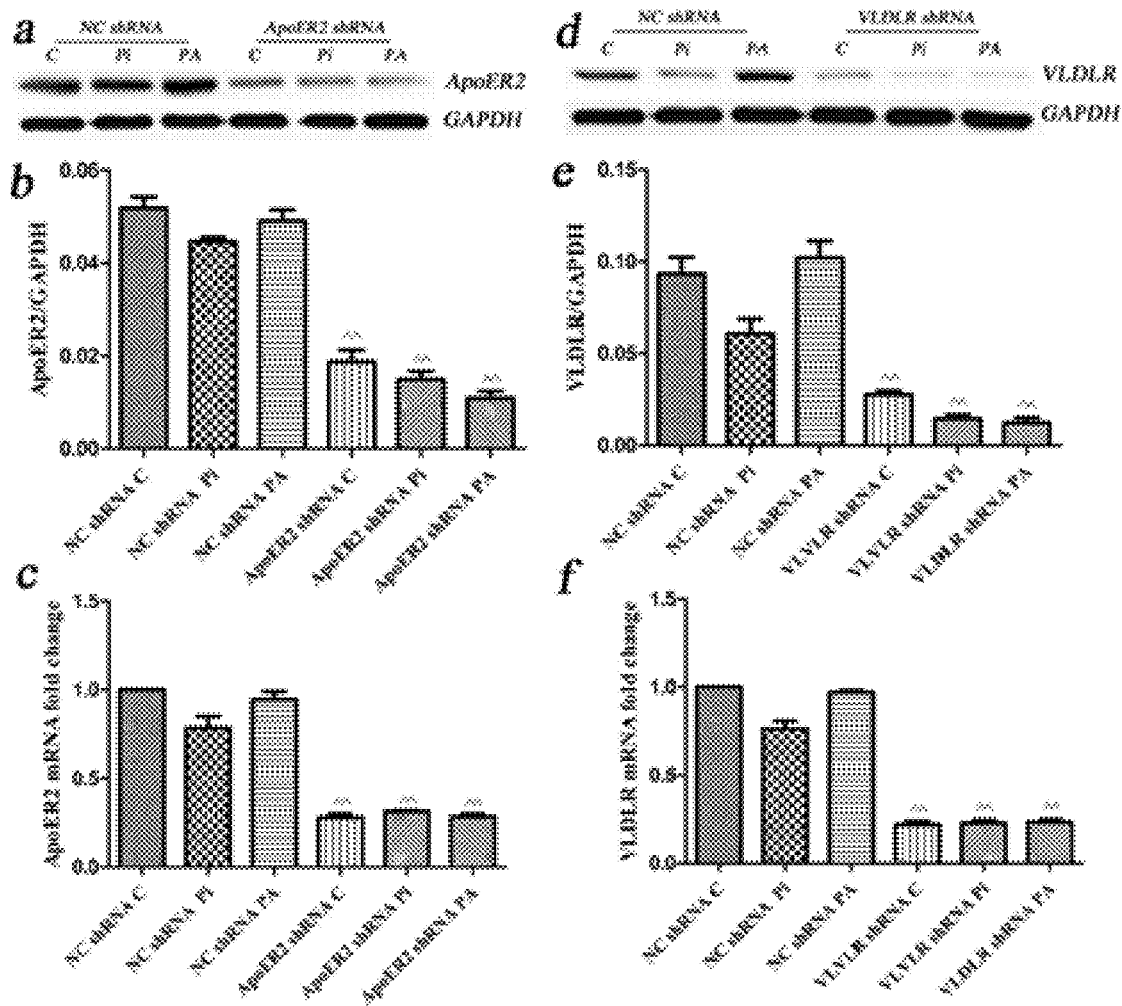
FIGS. 21A-21D



FIGS. 22A-22B



FIGS. 23A-23D



FIGS. 24A-24F

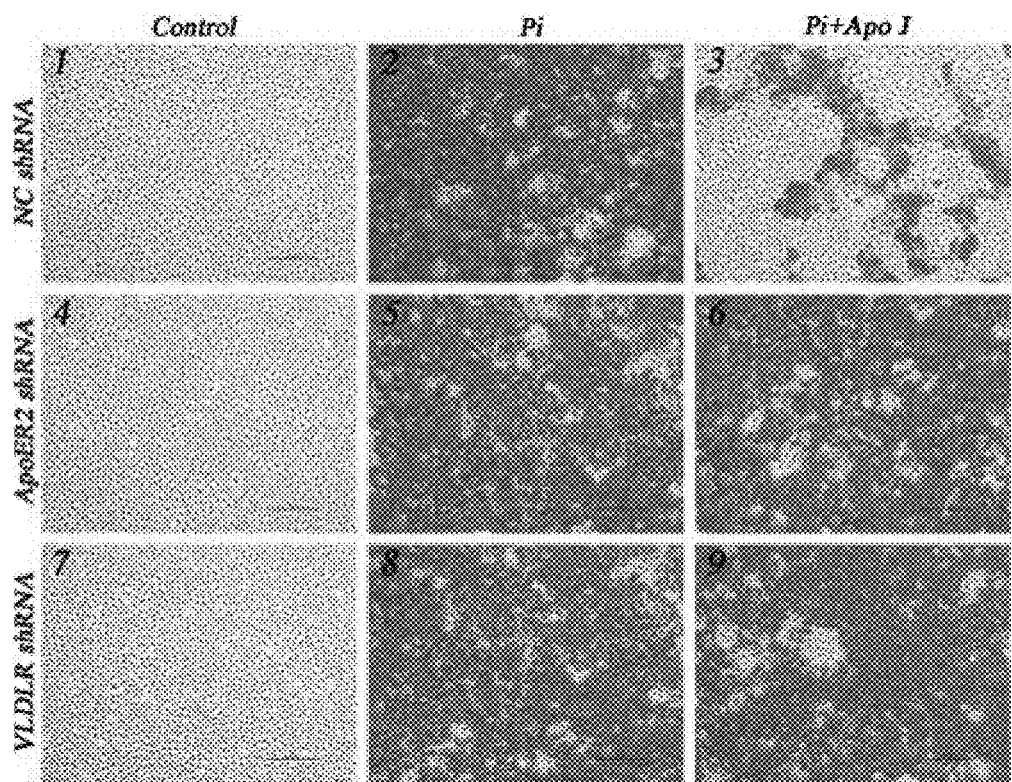


FIG. 25A

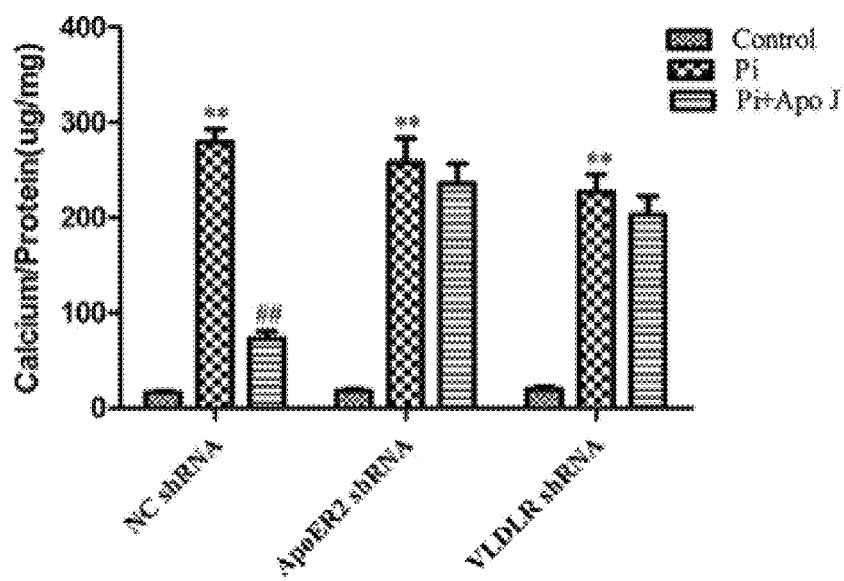
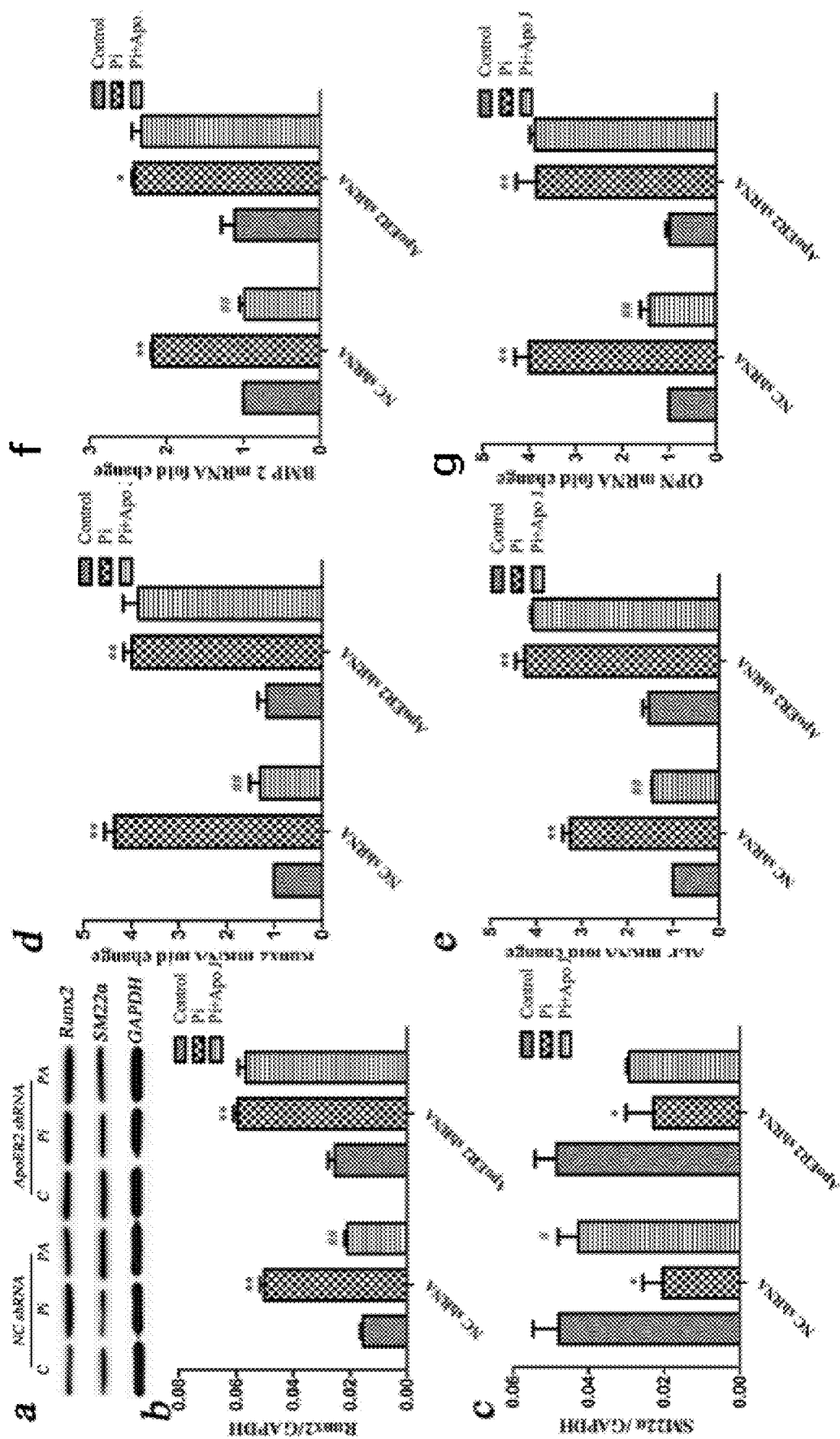
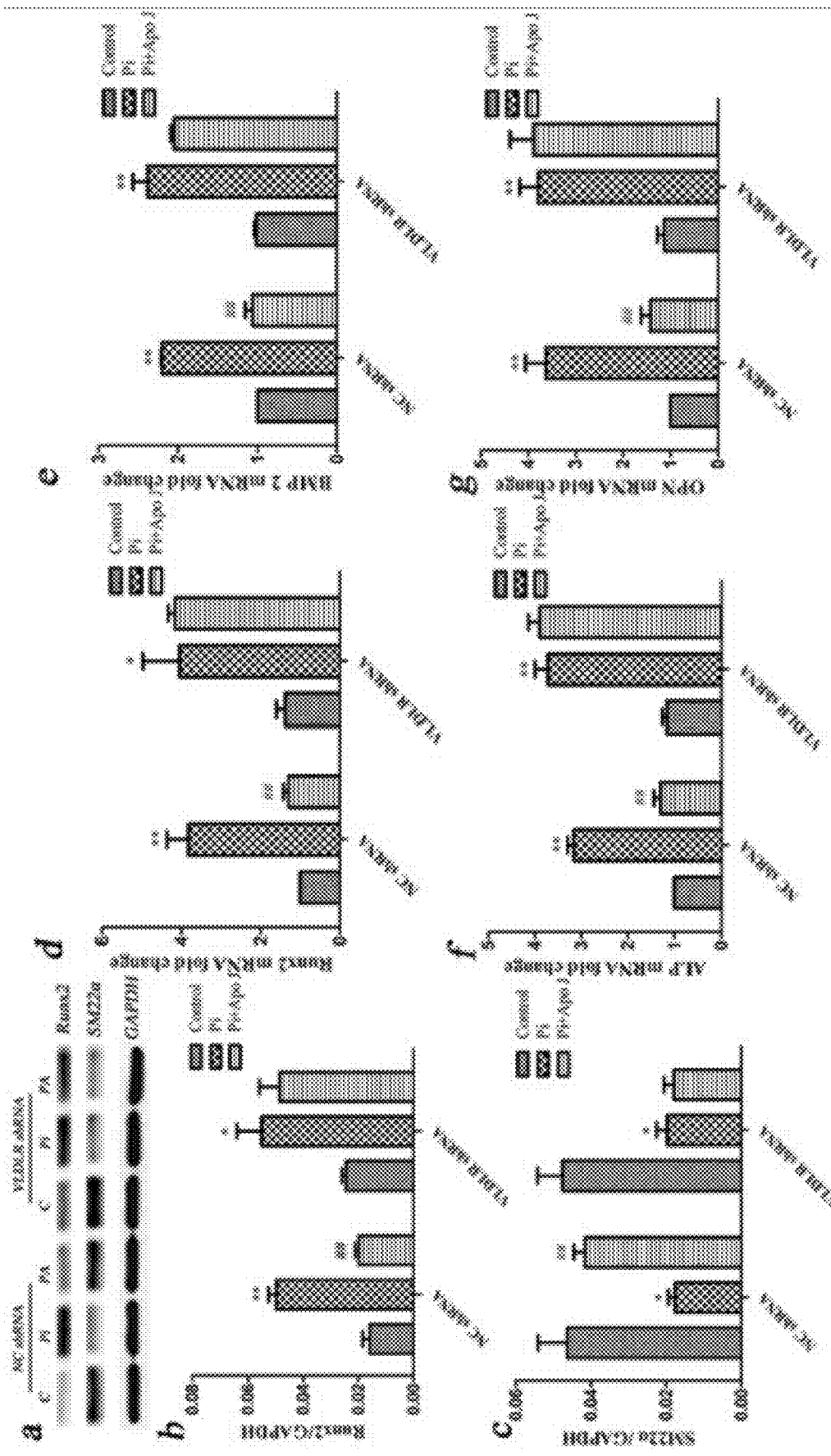
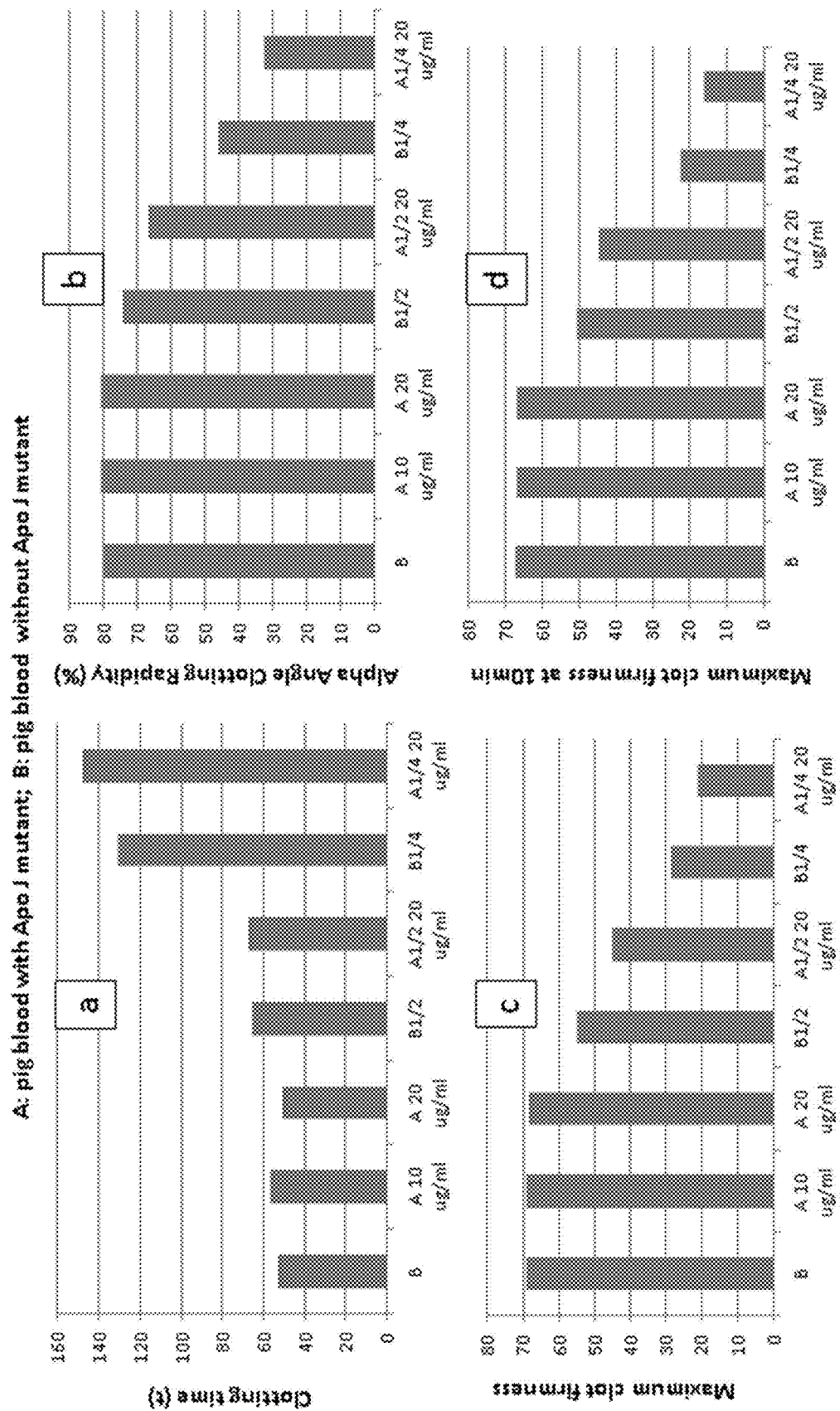


FIG. 25B





FIGS. 27A-27G



FIGS. 28a-28d

RECOMBINANT CLUSTERIN AND USE THEREOF IN THE TREATMENT AND PREVENTION OF DISEASE

[0001] This application is a continuation of U.S. patent application Ser. No. 16/597,400, now U.S. Pat. No. 11,104,719, filed Oct. 9, 2019, which is a continuation of U.S. patent application Ser. No. 14/957,030, now U.S. Pat. No. 10,464,994, filed Dec. 2, 2015, which claims the benefit of U.S. Provisional Patent Application No. 62/087,364, filed Dec. 4, 2014, the entirety of each of which is incorporated herein by reference.

INCORPORATION OF SEQUENCE LISTING

[0002] The sequence listing that is contained in the file named “UTSHP0306USC2_ST25.txt”, which is 61 KB (as measured in Microsoft Windows®) and was created on Aug. 17, 2021, is filed herewith by electronic submission and is incorporated by reference herein.

BACKGROUND OF THE INVENTION

1. Field of the Invention

[0003] The present invention generally relates production of recombinant clusterin and the use thereof for prevention and treatment of pathological conditions.

2. Description of Related Art

[0004] Hypercholesterolemia represents the most defined risk factor for atherosclerosis, an arterial disease that causes myocardial and cerebral infarctions. Low density lipoprotein (LDL) carries cholesterol from the liver to peripheral tissues, and when elevated in blood, LDL deposits the lipid in the arterial wall, which in turn develops atherosclerotic plaques and increases the risk for thrombotic events in the arteries. In contrast, high density lipoprotein (HDL) functions as a reverse cholesterol-transporter that removes the lipid from the arterial wall to the liver where cholesterol is metabolized. In essence, LDL is pro-atherogenic while HDL anti-atherogenic (Nicholls et al., 2005; Ansell et al., 2004; von Eckardstein et al., 2005). LDL mainly contains ApoB 100 and HDL apoE and apoA-I. In spite of the success of lowering LDL-cholesterol therapy with statins, raising HDL levels with torcetrapib (an inhibitor of cholesterol ester transfer protein (CETP) has shown little benefit to patients with atherosclerosis (Kastelein et al., 2007; Nissen et al., 2007). The failure of torcetrapib therapy underscores the incompleteness of our fundamental understanding of HDL function. HDL particles are heterogeneous in shape, density, size, composition and have multiple functional properties such as reverse cholesterol transport, as well as anti-oxidant, anti-inflammatory, and anti-thrombotic activities. Indeed, dysfunctional, proinflammatory HDL has been found in several pathological conditions, including atherosclerosis (Smith et al., 2010), diabetes (Hoofnagle et al., 2010) and autoimmune disorders (McMahon et al., 2009; Volkmann et al., 2010). Thus, the development of an anti-atherogenic, anti-apoptotic, and anti-inflammatory agent that enables HDL beneficial action is of clinical significance.

[0005] Clusterin is a sulfated, heterodimeric glycoprotein containing two 40 kDa chains joined by a unique five disulfide bond motif 1). It contains several domains, such as amphipathic helix, heparin-binding domain, and lipid-binding domain. This protein was initially identified from ram

rete testes fluid and named for its ability to elicit clustering of Sertoli cells supporting sperm maturation and development (NCBI/GenBank Accession No. NM_203339, NM_001831). Thereafter, species homologues have been isolated and cloned by a number of groups working in widely divergent areas, generating various synonyms of clusterin, including testosterone repressed prostate message-2 (TRPM-2), sulfated glycoprotein-2 (SGP-2), apolipoprotein-J (Apo-J), SP-40, 40, complement cytotoxicity inhibitor (CLI), and dimeric acidic glycoprotein (DAG), gp 80, NA1/NA2, glycoprotein III, etc.

[0006] Encoded on a 2-kb mRNA, clusterin is transcribed from a single copy gene located on mouse chromosome 14 and human 8p21 (Fink et al., 1993), and translated as a 51 kD or so protein comprising 427 amino acid sequence (Jordan-Starck et al., 1994). In the blood stream, clusterin circulates mainly with HDL as one of apolipoproteins (Choi-Miura et al., 1992; Stuart et al., 1992) but a small portion of clusterin may exist in LDL (Karlsson et al., 2005). Clusterin expression is induced by stress responses (Wilson et al., 2000). Clusterin binds megalin/LRP-2 receptor, members of LDL receptor family. Increased expression of clusterin occurs in both human (Mackness et al., 1997; Ishikawa et al., 2001; Ishikawa et al., 1998) and experimental animal (Jordan-Starck et al., 1994; Navab et al., 1997) atherosclerotic lesions. Reported functions of clusterin include apoptosis inhibition (Kowolik et al. 2006), complement factor inactivation (Correa-Rotter et al., 1992), lipid recycling and transport (Gelissen et al., 1998), membrane protection, and maintenance of cell-cell or cell-substratum contacts. It can effectively bind to lipids and promote efflux of cholesterol and oxysterols from lipid-laden foam cells, a hall-marker of atherosclerosis (Gelissen et al., 1998). Clusterin has a high-affinity to a wide array of biological ligands. The presence of both hydrophilic and hydrophobic domains enables clusterin to act as a chaperone or a “biological detergent”.

[0007] Clusterin plays a role in regulation of metabolism and function of various tissues and organs, particularly in the cardiovascular system. HDL with decreased levels of clusterin has been found in association with a high incidence of myocardial infarction in patients with insulin-resistant metabolic syndromes (Hoofnagle et al., 2010). Administration of an oral clusterin peptide was reported to reduce atherosclerosis in ApoE-null mice (Navab et al., 2005), and intravenous injection of clusterin diminishes rat myocardial infarction (Van Dijk et al., 2010). Transduction of clusterin can restore the mitochondrial membrane potential and prevent the release of cytochrome-c from mitochondria into cytoplasm in cardiac myoblasts damaged by ethanol (Li et al., 2007). Furthermore, increased clusterin expression in myoblasts enhances the cell capacity of migration and homing through induction of CXCR4, a chemokine-receptor for stromal cell-derived factor (SDF) (Li et al., 2010).

[0008] Human clusterin gene located in chromosome 8 (location 8p21-p18) with 17876 bp long contains 10 exons in total. Exon one and exon two are alternative yielding two different transcript isoforms. Other exons (Ansell et al., 2004; von Eckardstein et al., 2005; Kastelein et al., 2007; Nissen et al., 2007; Smith et al., 2010; Hoofnagle et al. 2010; McMahon et al., 2009; Volkmann et al., 2010) are shared with both isoforms. clusterin transcripts contain 3 different translation start sites (ATG), all in-frame. The best characterized protein isoform is produced from transcript isoform 2, where translation starts at the second ATG present in exon

2, right before ER-targeting signal. clusterin protein precursor (NP-976084) consists of 449 amino acids. There is evidence suggesting that two nuclear protein isoforms can be produced from this transcript isoform, one in which translation starts at ATG in exon 3 (417 aa), and another with translation starting from ATG in exon 1 (459 aa). Secreted clusterin is produced from the transcript isoform 2. The initial protein precursor, presecretory psCLU (~60 kDa), becomes heavily glycosylated and cleaved in the ER, and the resulting alpha and beta peptide chains are held together by 5 disulfide bonds in the mature secreted heterodimer protein form, sCLU (~75-80 kDa).

[0009] Under stimulation by ionic radiation and oxidative stress, the nuclear clusterin is first translated as a non-glycosylated protein precursor, pnCLU (~49 kDa), that is then translocated into nucleus. There is evidence of two distinct sized nuclear clusterin proteins (~50 kDa and ~60 kDa) (Pajak et al., 2007), that could result from translation started either at ATG present in exon 3 or in exon 1, respectively. Secreted clusterin is cytoprotective but nuclear clusterin cytotoxic. The controversy of clusterin functions mainly results from the not well-established role of the two different protein isoforms with distinct subcellular localization and somewhat opposing functionalities. Some known functions include involvement in apoptosis through complexing with Ku70 autoantigen (nCLU, proapoptotic) or interfering with Bax-activation (sCLU, antiapoptotic) (Araki et al., 2005; Klovov et al., 2004; Leskov et al., 2003; Yang et al., 2000). Clusterin has also been linked to spermatogenesis, epithelial cell differentiation, TGF-beta signaling through Smad2/Smad3 (Shin et al., 2008; Ahn et al., 2008; Lee et al., 1992), complement activation (Dietzsch et al., 1992; O'Bryan et al., 1990). Secreted native clusterin contains the sequence domains of nuclear clusterin critical for nuclear translocation and binding to nuclear death signaling proteins such as Ku70.

[0010] Despite the various roles in cellular regulation ascribed to clusterin, there remains a need for the development of recombinant clusterin and clusterin analogs as potential therapeutics. Embodiments of this invention disclose technology of producing recombinant clusterin with a high homology to the secreted form of native clusterin with a protective function, and compositions of recombinant clusterin that can be used for prevention and treatment of diseases in a mammal.

SUMMARY OF THE INVENTION

[0011] In a first embodiment, a recombinant polypeptide is provided that comprises a mammalian clusterin coding sequence. In various aspects, the clusterin coding sequence may have a deletion of a nuclear localization signal and/or transmembrane domain (TMD). In some aspects, the polypeptide may be a fusion protein comprising the clusterin coding sequence and a heterologous polypeptide sequence. For example, the polypeptide may further comprise a tag sequence. In further aspects, the polypeptide may comprise a protease cleavage site (e.g., a thrombin cleavage site (Leu-Val-Pro-Arg-Gly-Ser; SEQ ID NO: 16) or enteropeptidase cleavage site (Asp-Asp-Asp-Asp-Lys; SEQ ID NO: 17)). For example, protease cleavage site can be positioned between the tag sequence and the clusterin coding sequence. In various aspects, the tag sequence may be a polyhistidine tag. In some aspects, the tag sequence may be positioned N-terminally relative to the Clusterin coding sequence,

while in other aspects the tag sequence may be positioned C-terminally relative to the Clusterin coding sequence.

[0012] In a further embodiment, a composition is provided that comprises a clusterin polypeptide of the present embodiments in a pharmaceutically acceptable carrier. In various aspects, the composition may be frozen or lyophilized.

[0013] In yet a further embodiment, an isolated polynucleotide molecule is provided that comprises a nucleic acid sequence encoding a clusterin polypeptide of the present embodiments. In some aspects, the nucleic acid sequence encoding the polypeptide may be operably linked to a promoter. In certain aspects, the promoter may be a promoter functional in mammalian, bacterial or insect cells. In some aspects, the polynucleotide molecule may be part of an expression vector, such as, a plasmid, an episomal expression vector or a viral expression vector.

[0014] In a further embodiment, a host cell is provided that comprises a polynucleotide molecule encoding a clusterin polypeptide of the present embodiments. In some aspect, the host cell may be a bacterial cell, an insect cell, or a mammalian cell. In some specific aspects, the host cell is a human cell, such as a pluripotent cell, a cardiac cell, and endothelial cell, or a cardiac or endothelial precursor cell.

[0015] In yet a further embodiment, a method of manufacture a recombinant clusterin polypeptide is provided that comprises (a) expressing a polynucleotide molecule encoding a clusterin polypeptide of the present embodiments in a cell; and (b) purifying the polypeptide from the cell. In various aspects, the polypeptide may comprise a purification tag, and purifying the polypeptide may comprise use of a matrix having an affinity for the purification tag. In some aspects, the purification tag may be a polyhistidine tag, and purifying the polypeptide may comprise purifying the polypeptide using a metal affinity column. In certain aspects, the purification tag may further comprise a protease cleavage site positioned between the tag sequence and the clusterin coding sequence, and purifying the polypeptide may comprise contacting the polypeptide with a protease that cleaves at the cleavage site.

[0016] In certain aspects, methods of the embodiments concern construction and/or transfection of a nucleotide encoding clusterin into cells of a mammalian cell or a non-mammalian cell causes sufficient expression of clusterin polypeptides. In certain embodiments, the step of causing the expression of an amount of a nucleotide encoding clusterin includes transfecting cells of the tissue with a DNA sequence encoding the entire clusterin polypeptide sequence, or a biologically active portion of the clusterin sequence, operably linked to a promoter and capable of being expressed in the cells to provide an amount of clusterin sufficient to be identified, concentrated, extracted, and purified.

[0017] In still a further embodiment, a method of treating or preventing a cardiovascular disease in a subject is provided that comprises administering an effective amount of a clusterin composition comprising (a) a recombinant clusterin polypeptide, (b) a polynucleotide (e.g., an expression vector) encoding a clusterin polypeptide, and/or (c) cells expressing exogenous clusterin polypeptide of the present embodiments. In some aspects, the cardiovascular disease may be hypertension, hyperlipidemia, hypercholesterolemia, hyperglycemia, hypertension, atherosclerosis and atherosclerosis-associated ischemic heart failure, stenosis, cal-

cification of cardiovascular tissues, stroke, myocardial infarction or cerebral infarction. In some aspects, the cardiovascular disease is hyperlipidemia, hypercholesterolemia or atherosclerosis. In still further aspects, the cardiovascular disease may be diabetes. In various aspects, an effective amount of a clusterin composition may be an amount effective to reduce blood cholesterol, reduce blood glucose, reduce blood triglyceride, increase efflux of intracellular cholesterol, and/or increase vascular or cardiac cell survival. In some aspects an effective amount of a clusterin composition provides enhancement or promotion of cell survival and growth against cytotoxic or cytostatic factors, including but not limited to, oxysterols, oxidized lipoproteins, and proinflammatory cytokines.

[0018] In accordance with certain embodiments, a method of treating or preventing atherosclerosis, or a complication thereof in a mammal, is provided. For the purposes of this disclosure, the term “preventing” atherosclerosis has its usual meaning in the art and includes “detering” and “reducing the risk of” atherosclerosis. This method comprises carrying out an above-described method wherein the tissue is a cardiac or vascular or brain region comprising an atherosclerotic lesion, or an area that is at risk of forming an atherosclerotic lesion, and wherein the contacting of cells in the tissue with clusterin polypeptides deters or prevents apoptotic cell death sufficiently to prevent, or reduce the risk of, formation of an atherosclerotic lesion. In some embodiments, the contacting of cells in the tissue with clusterin polypeptides deters or prevents tissue injury or degeneration sufficiently to prevent, or reduce the risk of, rupture of an atherosclerotic lesion.

[0019] In certain aspects, the atherosclerotic lesion comprises calcification in a vessel or a valve with inflammation, which causes vascular tissue stiffness and cardiac or aortic valve malfunction, comprising stenosis or insufficient closure. In certain of the above-described methods, the amounts of clusterin compositions are effective to deter or prevent calcification and/or protect against inflammatory injury, induced by at least one condition chosen from the group consisting of: hypercholesterolemia, hyperglycemia, hyperphosphatemia, and/or hypertension.

[0020] In some aspects, the atherosclerotic lesion or plaque comprises an unstable plaque caused by hyperlipidemia and inflammation, and the amount of clusterin contacting a treatment site are effective to stabilize the plaque (e.g., reduce the risk of rupture of the plaque, thrombus formation, or other complications).

[0021] Another embodiment of the present invention provides a method of treating acute vascular syndromes and heart failure in a mammal, which comprises delivery of an amount of clusterin composition into the heart of the mammal to protect and improve heart function.

[0022] In yet a further embodiment, a method of treating or preventing alcoholism in a subject is provided that comprises administering an effective amount of a clusterin composition comprising (a) a recombinant clusterin polypeptide, (b) a polynucleotide (e.g., an expression vector) encoding a clusterin polypeptide, and/or (c) cells expressing exogenous clusterin polypeptide of the present embodiments. In some aspects, the effective amount of the clusterin composition is an amount effective to reduce withdrawal symptoms, alcohol intake or markers of liver damage in a subject.

[0023] In some aspects, a clusterin composition of the embodiments (e.g., recombinant clusterin polypeptide, a polynucleotide encoding a clusterin polypeptide, and/or cells expressing exogenous clusterin polypeptide) may be administered by intravenous injection or catheter delivery. In some aspects, clusterin compositions are delivered into one or more tissue or organs of a mammal suffering from, or at risk of being subjected to, physical and/or chemical injury. In some embodiments the tissue is heart or vascular or brain tissue. In certain embodiments, the vascular or cardiac tissue is affected with atherosclerosis or heart failure. In other embodiments, the vascular or cardiac tissue is not affected with atherosclerosis, acute vascular syndromes, or heart failure. In some embodiments, the cells are one or more of the cell types: vascular endothelial cells, smooth muscle cells, cardiac myocytes and brain cells.

[0024] Thus, in certain aspects, there is provided a method of administering a mammalian cell, such as a stem cell, with enhanced expression of recombinant clusterin, or that have been treated with recombinant clusterin, into a tissue or organ. In some embodiments, the tissue or organ comprises a failing heart or an atherosclerotic blood vessel. In some embodiments, the stem cells are administered by intravenous injection, intra-arterial catheter, or by intramuscular or intratissue injection. In certain embodiments, the stem cells are delivered or injected together with an agent that causes vascular dilation and/or are co-administered with an anti-thrombotic agent. These and other embodiments, features and advantages of the present invention will be recognized by those of skill in the art from the following detailed description and drawings.

[0025] In further aspects, delivering an amount of recombinant clusterin comprises dissolving the polypeptide in a solution or buffer and injecting the clusterin containing solution into blood stream or tissues of a mammal or apply said clusterin solution on the surface of tissues or organs with injury.

[0026] As used herein, the terms “clusterin”, “Apolipoprotein J”, and “Apo J” are used interchangeably.

[0027] As used herein, “essentially free,” in terms of a specified component, is used herein to mean that none of the specified component has been purposefully formulated into a composition and/or is present only as a contaminant or in trace amounts. The total amount of the specified component resulting from any unintended contamination of a composition is therefore well below 0.05%. In some aspects, most preferred is a composition in which no amount of the specified component can be detected with standard analytical methods.

[0028] As used herein in the specification and claims, “a” or “an” may mean one or more. As used herein in the specification and claims, when used in conjunction with the word “comprising”, the words “a” or “an” may mean one or more than one. As used herein, in the specification and claim, “another” or “a further” may mean at least a second or more.

[0029] As used herein in the specification and claims, the term “about” is used to indicate that a value includes the inherent variation of error for the device, the method being employed to determine the value, or the variation that exists among the study subjects.

[0030] Other objects, features and advantages of the present invention will become apparent from the following detailed description. It should be understood, however, that

the detailed description and the specific examples, while indicating certain embodiments of the invention, are given by way of illustration only, since various changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art from this detailed description.

BRIEF DESCRIPTION OF THE DRAWINGS

[0031] The following drawings form part of the present specification and are included to further demonstrate certain aspects of the present invention. The invention may be better understood by reference to one or more of these drawings in combination with the detailed description of specific embodiments presented herein.

[0032] FIG. 1—A schematic representation of the clusterin analog plasmid pCluAg. Representation of clusterin cDNA subcloning and construction of an expression vector with an insert of clusterin analog-encoding cDNA is shown (SEQ ID NO: 26).

[0033] FIGS. 2A-2E—Polypeptide sequences and structures of various clusterin polypeptides of the embodiments.

[0034] FIG. 3—Coomassie blue (G250) dye-stained polyacrylamide gel (SDS-PAGE) of recombinant human clusterin protein stored in solution and in dry powder for 3 months. The results show long term stability of the polypeptides.

[0035] FIG. 4.—Immunoblotting analysis of the recombinant human clusterin polypeptide. Clusterin polypeptides (1 μ g/lane) were loaded into SDS-PAGE (7%). After electrophoresis, protein bands were transferred onto PVDF membrane, probed with rabbit anti-human clusterin antibodies (1:200), and immunostained bands were developed by chemiluminescence. The results again show long term stability of the polypeptides.

[0036] FIG. 5. —Ion exchange fast protein liquid chromatography (FPLC) of recombinant human clusterin proteins. Clusterin polypeptides were separated from other components in an aqueous solution, or buffer. The buffer flow rate was controlled by a positive-displacement pump and is normally kept constant, while the composition of the buffer can be varied by drawing fluids in different proportions from two or more external reservoirs. Ion Exchange FPLC (BioRad UnoQ12 column); Buffer A: 20 mM Tris (pH8.0) 0.5 mM EDTA; Buffer B: 20 mM Tris (pH8.0) 0.5 mM EDTA+1N NaCl; Flow rate: 2 ml/min; Injected amount: 250 μ g; Result: peak eluted after 80 minutes.

[0037] FIGS. 6A-6C—Immunoblotting analysis of levels of recombinant clusterin injected intravenously into the blood stream of wild type and ApoE-null mice. Immunoblotting for clusterin and ApoA1 proteins in the plasma of WT and ApoE-null mice with different recombinant clusterin analog (CluAg) or saline control injection (7 days) (FIG. 6A). Densitometry showed increased clusterin levels in Clu injected mice as compared to saline injected one (FIG. 6B). Co-precipitation of clusterin (Clu) in the plasma of C57BL/6 mice injected with recombinant human clusterin analogs (FIG. 6C). Apolipoprotein-A1, a known HDL component, was pulled-down with anti-ApoA1 antibody, and Western blot analysis of the ApoA1 pull-down was conducted with anti-clusterin antibodies.

[0038] FIG. 7—Immunoblotting analysis of clusterin and HisTR tag in proteins extracted from the supernatants of

human smooth muscle cell (SMC) cultures treated with oxLDL and recombinant clusterin analog (CluAg) with HisTR tag.

[0039] FIG. 8—Growth curves evidencing that incubation with recombinant clusterin analog stimulates proliferation of human vascular smooth muscle cells. Human smooth muscle cells (SMC) were incubated in DMEM media containing CluAg-I (1-6 μ g/ml).

[0040] FIG. 9—Growth curves evidencing that incubation with recombinant clusterin analog blocks inhibitory effect of oxidized lipoprotein and stimulates proliferation of human vascular smooth muscle cells (SMCs). Human SMCs were incubated in DMEM media containing CluAg-I (1-6 μ g/ml) in the presence of 50 μ g/ml oxidized low density lipoprotein (oxLDL).

[0041] FIG. 10—Blood plasma comparison of WT mice (upper panel) and ApoE^{-/-} mice (lower panel). Graphs show injection of human recombinant clusterin, CluAg-I, reducing blood levels of glucose, cholesterol, triglyceride, and LDL in ApoE^{-/-} atherosclerosis-prone mice but not in age and sex-matched normal wild type (WT) control mice.

[0042] FIG. 11—Blood pressure comparison of WT mice and ApoE^{-/-} mice. Graphs show intravenous injection of human recombinant clusterin analog, CluAg-I, for 3 months reducing both systolic and blood pressure in ApoE^{-/-} atherosclerosis-prone mice but has no effect on that in age (6-8 months old) and sex (male)-matched normal wild type (WT) control mice.

[0043] FIG. 12—Echocardiography of normal wild type (WT) and ApoE^{-/-} mice with CluAg-I injection. WT and ApoE^{-/-} mice were injected intravenously with CluAg for 3 months and then subjected to ultrasound examination using the VisualsonicsTM echo device.

[0044] FIG. 13—Left ventricle ejection fraction comparison of WT mice and ApoE^{-/-} mice. Intravenous injection of human recombinant clusterin analog, CluAg-I, for 3 months reduces blood pressure in ApoE^{-/-} atherosclerosis-prone mice but not in age (6-8 months old) and sex (male)-matched normal wild type (WT) control mice.

[0045] FIGS. 14A-14D—Oil Red O staining of WT and ApoE^{-/-} aortas, evidencing that weekly intravenous injection of human recombinant clusterin analog, CluAg-I, for 3 months reduces plaque sizes in ApoE^{-/-} atherosclerosis-prone mice, but not in age (6-8 months old) and sex (male)-matched normal wild type (WT) control mice. Reduced Oil red O staining in ApoE^{-/-} mice treated with CluAg is observed (FIG. 14D).

[0046] FIG. 15—Alizarin Red staining of murine vascular smooth muscle cells evidence that CluAg reduces phosphate-induced calcification of ApoE^{-/-} SMCs in a manner dependent upon expression of ApoER2 and VLDLR. SMCs treated with ApoER2 or VLDLR shRNA were incubated with Pi (3.6 mmol/L)+/-CluAg in DMEM containing 5% FBS for 6 days. At the end of culture cells were washed in PBS, fixed in 10% formalin, and stained in 2% Alizarin Red at 37° C. for 10 min. Reduced Alizarin Red staining could be partly blocked by ApoER2 and VLDLR shRNA.

[0047] FIGS. 16A-16C—Graphs show that CluAg treatment reduces phosphate-induced (Pi) calcification in ApoE^{-/-} vascular smooth muscle cells (SMCs) in a manner dependent upon expression of ApoER2 and VLDLR. SMCs treated with ApoER2 or VLDLR shRNA were incubated with Pi (3.6 mmol/L) in the presence or absence of CluAg (6 μ g/ml) in DMEM containing 5% FBS for 6 days. In the

end of culture cells were washed in PBS, incubated in 0.6M HCl overnight. Cell lysates were mixed with 0.1N NaOH was used to lysis cells and concentrations of proteins were measured. Supernatants were collected, and calcium contents measured using a calcium assay; were mixed with 90 μ l of the Chromogenic Reagent, and 60 μ l of the Calcium Assay Buffer, mix gently; Incubated the reaction for 5-10 mins at room temperature, protected from light; Measured OD at 575 nm. **16A**, SMCs treated with negative control ShRNA; **16B**, with ApoER2 ShRNA; and **16C**, with VLDLR ShRNA. Data represent means $\pm$ S.D., **, $p<0.01$ and # $p<0.05$.

[0048] FIGS. **17A-17D**—Alizarin Red staining showing that weekly intravenous injection of human recombinant clusterin analog, CluAg for 3 months inhibits atherosclerosis as well as calcification in ApoE^{-/-} atherosclerosis-prone mice. **17A** shows wild type (WT) control mice. Aortas were stained in 2% Alizarin Red at 37° C. for 10 min. Images were taken using $\times 4$ objective for **17A** and **17B**, and $\times 10$ objective for **17C** and **17D**. Reduced plaque size and Alizarin staining in ApoE^{-/-} mice treated with CluAg was observed.

[0049] FIGS. **18A-18H**—Immunofluorescence of bone morphogenic protein-2 (BMP2) in aortas of wild type (WT) mice and ApoE^{-/-} mice treated with or without CluAg. Again, results show that CluAg treatment is able to reduce BMP2 expression in ApoE^{-/-} mice.

[0050] FIGS. **19A-19B**—Calcium deposit in smooth muscle cells (SMC) isolated from aorta of WT and ApoE^{-/-} null mice. ApoE^{-/-} VSMCs are more prone to Pi-induced calcification. Alizarin red staining (**19A**) and calcium assay (**19B**) of VSMCs isolated from aorta of WT and ApoE^{-/-} mice. Scale bar=200 μ m. VSMCs from WT and ApoE^{-/-} mice were treated with inorganic phosphate (Pi) with or without Apo J (6 μ g/mL). Bars represent means $\pm$ SD, $n=5$, * $P<0.05$ vs. control; ** $P<0.01$ vs. control; # $P<0.05$ vs. Pi group; ## $P<0.01$ vs. Pi group.

[0051] FIGS. **20A-20B**—Calcium deposit in SMC treating with or without Apo J. Inhibitory effect of Apo J on calcification in ApoE^{-/-} VSMCs is correlated to concentration of Apo J. a: Alizarin S staining of VSMCs from ApoE^{-/-} mice. VSMCs were treated with different concentrations of Apo J (0-12 μ g/mL) during calcification. Scale bar=200 μ m. b: Calcium assay of ApoE^{-/-} VSMCs treated with various concentrations of Apo J (0-12 μ g/mL) and Pi. Bars represent means $\pm$ SD; $n=5$ per group. ** $P<0.01$ vs. control; ## $P<0.01$ vs. Pi group.

[0052] FIGS. **21A-21D**—Apo J modulates the protein expression of smooth muscle lineage-specific markers and calcification-related genes. VSMCs were treated with Pi and Apo J (0-12 μ g/mL). **21A**: 40 μ L cell culture medium was used to detect the level of Apo J in culture medium. Levels of SM22 α , α SMA and Runx2 protein expressions were assessed by western blot. GAPDH was used as the loading control. **21B-21D**: Densitometry analysis shows the quantification of SM22 α , α SMA and Runx2. Bars represent means $\pm$ SD; $n=5$ per group. * $P<0.05$ vs. control; ** $P<0.01$ vs. control; # $P<0.05$ vs. Pi group; ## $P<0.01$ vs. Pi group.

[0053] FIGS. **22A-22B**—Expression of Apo J during calcification. **22A**: 40 μ L cell culture medium was used to detect the expression of Apo J by Western blot. **22B**: Apo J mRNA expression was detected by qRT-PCR. GAPDH was used as an internal control. Bars represent means $\pm$ SD; $n=3$ per group. * $P<0.05$ vs. control; ** $P<0.01$ vs. control.

[0054] FIGS. **23A-23D**—Apo J affects osteogenetic markers on mRNA levels during calcification. VSMCs were treated with Pi in combination with Apo J (0-12 μ g/mL). Runx2 (**23A**), BMP-2 (**23B**), ALP (**23C**), and OPN (**23D**) mRNA expressions were quantified by qRT-PCR and normalized to GAPDH as an internal control. Bars represent means $\pm$ SD; $n=5$ per group. ** $P<0.01$ vs. control; # $P<0.05$ vs. Pi group; ## $P<0.01$ vs. Pi group.

[0055] FIGS. **24A-24F**—Lentiviral shRNA knockdown of ApoER2 and VLDLR receptors. **24A-24B**: ApoE^{-/-} cells were transfected with ApoER2 shRNA, VLDLR shRNA or negative control shRNA to collect whole-cell lysates for western blot analysis of ApoER2 or VLDLR. **24C-24D**: Densitometry analysis shows the quantification of ApoER2 or VLDLR expression. **24E-24F**: Bar graph illustrating real-time PCR data demonstrating that ApoER2 or VLDLR was successfully knocked down on mRNA level. Relative mRNA expressions were normalized to GAPDH as an internal control. Bars represent means $\pm$ SD; $n=3$ per group. * $P<0.05$ vs. control, ** $P<0.01$ vs. control; # $P<0.05$ vs. Pi group, ## $P<0.01$ vs. Pi group.

[0056] FIGS. **25A-25B**—Knockdown of ApoER2 or VLDLR gene abolishes the inhibitory effect of Apo J on calcification. Alizarin red staining (**25A**) and calcium assay (**25B**) of receptor knockdown VSMCs under the treatment of Pi with or without Apo J (6 μ g/mL). Scale bar=200 μ m. Bars represent means $\pm$ SD; $n=3$ per group. ** $P<0.01$ vs. control; ## $P<0.01$ vs. Pi group.

[0057] FIGS. **26A-26G**—Knockdown of ApoER2 gene abolishes the effect of Apo J on calcification markers and smooth muscle lineage-specific markers. ApoE^{-/-} VSMCs were treated with Pi and Apo J (6 μ g/mL). **26A**: SM22 α and Runx2 protein expressions in ApoER2 knockdown cells were detected by western blot. GAPDH was used as the loading control. **26A-C**: Densitometry analysis shows the quantification of SM22 α and Runx2 in ApoER2 knockdown cells. **26D-G**: Bar graph illustrating real-time PCR data showing the mRNA expression of Runx2, BMP-2, ALP and OPN. Relative mRNA expressions were normalized to GAPDH as an internal control. Bars represent means $\pm$ SD; $n=3$ per group. + $P<0.05$ means negative control shRNA infected cells vs. ApoER2 shRNA infected cells, ++ $P<0.01$; * $P<0.05$ vs. control, ** $P<0.01$ vs. control; # $P<0.05$ vs. Pi group, ## $P<0.01$ vs. Pi group.

[0058] FIGS. **27A-27G**—Knockdown of VLDLR gene abolishes the effect of Apo J on calcification markers and smooth muscle lineage-specific markers. ApoE^{-/-} VSMCs were treated with Pi and Apo J (6 μ g/mL). **27A**: SM22 α and Runx2 protein expressions in VLDLR knockdown cells were detected by western blot. GAPDH was used as the loading control. **27B-C**: Densitometry analysis shows the quantification of SM22 α and Runx2 in VLDLR knockdown cells. **27D-G**: Bar graph illustrating real-time PCR data showing the mRNA expression of Runx2, BMP-2, ALP and OPN. Bars represent means $\pm$ SD; $n=3$ per group. + $P<0.05$ means negative control shRNA infected cells vs. VLDLR shRNA infected cells, ++ $P<0.01$; * $P<0.05$ vs. control, ** $P<0.01$ vs. control; # $P<0.05$ vs. Pi group, ## $P<0.01$ vs. Pi group.

[0059] FIGS. **28a-28d**—ApoJ mutants with a peptide comprising a thrombin-specific cleavage site exert an anticoagulation effect by competitively inhibiting the conversion of fibrinogen into time of pig blood with (A) or without (B) ApoJ mutant. Fresh blood (0.3 ml) from adult pig at

different dilutions (0, $\frac{1}{2}$, and $\frac{1}{4}$ in PBS) was mixed with recombinant ApoJ mutant at 10 or 10 mg/ml), incubated at 37° C., and then subjected to rotational thromboelastometry (ROTEM™). a, clotting time; b, Alpha Angle clotting rapidity; c, maximum clot firmness; and d, maximum clot firmness at 10 minutes. Data represented from average of two separate experiments with the ApoJ mutant, Clusterin-ATMD-TRHis (SEQ ID NO: 8).

DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

[0060] Clusterin is a multifunctional protein that may play an important role in regulation of survival, proliferation and differentiation of a diversity of cell types in a mammal. The present disclosure provides recombinant clusterin polypeptides and demonstrates the therapeutic efficacy of such recombinant polypeptides. In particular, data presented herein demonstrates that recombinant clusterin can be effectively synthesized and purified and that the clusterin polypeptide preparations are highly stable. The recombinant polypeptides appear to be essentially non-toxic, when administered to animals. However, in murine models for cardiovascular disease the clusterin polypeptides are able to reduce markers of cardiovascular disease such as hyperglycemia, hypertension and calcification as well as to normalize serum lipid levels. These data indicate that the recombinant clusterin polypeptides of the embodiments (and cells and nucleic acids encoding the same) could be used as safe and effective therapeutics for treatment and prevention of cardiovascular diseases and alcoholism.

I. RECOMBINANT CLUSTERIN POLYPEPTIDES

[0061] For the purposes of this disclosure, the terms “clusterin” or “recombinant clusterin” refers to proteins, whose sequence is based on a mammalian clusterin sequence. In preferred aspects a recombinant clusterin polypeptide comprises a deletion of a nuclear localization signal, a transmembrane domain and/or is fused with a heterologous polypeptide sequence (e.g., a purification tag). A skilled artisan will recognize that deletions of the clusterin TMD likewise can disrupt the endoplasmic reticulum (ER)-targeting of recombinant clusterin. The terms “ApoJ” and “Clusterin” as used interchangeably herein. Examples of specific clusterin polypeptides include, without limitation, polypeptides provided as NCBI Acc. No. NM_203339, NM_001831, NM_013492, NM_053021, NM_012679, each of which is incorporated herein by reference. In certain aspects, the recombinant clusterin is about or at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the clusterin polypeptide sequence of SEQ ID NOs: 1-3. For examples, in some aspects, the recombinant clusterin polypeptide about or at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to SEQ ID NO:1, but comprises a deletion of a nuclear localization signal, ER-targeting sequence and/or a transmembrane domain. In yet further aspects, the recombinant clusterin is about or at least about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the clusterin polypeptide sequence of SEQ ID NOs: 4-15.

[0062] Clusterin polypeptides and fragments, mutated, truncated or deleted forms of the clusterin and/or clusterin fusion proteins can be prepared for a variety of uses,

including but not limited to the generation of antibodies, as reagents in diagnostic assays, as reagents in assays for the identification of other cellular gene products involved in the regulation of clusterin mediated disorders, as reagents in assays for screening for compounds that can be used in the treatment of clusterin mediated disorders, and as pharmaceutical reagents, useful in the treatment of disorders related to clusterin.

[0063] Embodiments of the present invention also encompasses proteins that are functionally equivalent to the clusterin encoded by the nucleotide sequences described, as judged by any of a number of criteria, including but not limited to resulting in the biological effect of clusterin, a change in phenotype when the clusterin equivalent is present in an appropriate cell type. Such functionally equivalent clusterin proteins include but are not limited to additions or substitutions of amino acid residues within the amino acid sequence encoded by the clusterin nucleotide sequences described, but which result in a silent change, thus producing a functionally equivalent gene product.

[0064] In additional aspects, clusterin polypeptides may be further modified by one or more other amino substitutions while maintaining their biological activity. For example, amino acid substitutions can be made at one or more positions wherein the substitution is for an amino acid having a similar hydrophilicity. The importance of the hydropathic amino acid index in conferring interactive biologic function on a protein is generally understood in the art (Kyte and Doolittle, 1982). It is accepted that the relative hydropathic character of the amino acid contributes to the secondary structure of the resultant protein, which in turn defines the interaction of the protein with other molecules, for example, enzymes, substrates, receptors, DNA, antibodies, antigens, and the like. Thus such conservative substitution can be made in GrB and will likely only have minor effects on their activity. As detailed in U.S. Pat. No. 4,554, 101, the following hydrophilicity values have been assigned to amino acid residues: arginine (+3.0); lysine (+3.0); aspartate (+3.0 $\pm$ 1); glutamate (+3.0 $\pm$ 1); serine (+0.3); asparagine (+0.2); glutamine (+0.2); glycine (0); threonine (−0.4); proline (−0.5 $\pm$ 1); alanine (0.5); histidine (−0.5); cysteine (−1.0); methionine (−1.3); valine (−1.5); leucine (−1.8); isoleucine (−1.8); tyrosine (−2.3); phenylalanine (−2.5); tryptophan (−3.4). These values can be used as a guide and thus substitution of amino acids whose hydrophilicity values are within ± 2 are preferred, those that are within ± 1 are particularly preferred, and those within ± 0.5 are even more particularly preferred. Thus, any of the GrB polypeptides described herein may be modified by the substitution of an amino acid, for different, but homologous amino acid with a similar hydrophilicity value. Amino acids with hydrophilicities within ± 1.0 , or ± 0.5 points are considered homologous. Furthermore, it is envisioned that clusterin sequences may be modified by amino acid deletions, substitutions, additions or insertions while retaining its biological activity.

[0065] In some aspects, a clusterin polypeptide is fused with a heterologous polypeptide sequence. For example, heterologous polypeptide sequences may be included to aid production or purification of a cell targeting construct. Some specific examples of amino acid sequences that may be attached to clusterin include, but are not limited to, purification tags (e.g., a T7, MBP, GST, HA, or polyHis tag), proteolytic cleavage sites, such as a thrombin or furin

cleavage site, intracellular localization signals or secretion signals. In some aspects, a clusterin further comprises a cell-penetrating peptide (CPP). As used herein the terms CPP and membrane translocation peptide (MTP) as used interchangeably to refer to peptide sequences that enhance the ability of a protein to be internalized by a cell. Examples for CPPs for use according to the embodiments include, without limitation, peptide segments derived from HIV Tat, herpes virus VP22, the *Drosophila* Antennapedia homeobox gene product, protegrin I, as well as the T1, T2, and INF7 peptides.

[0066] Other mutations to the coding sequences described above can be made to generate polypeptides that are better suited for expression, scale up, etc. in the host cells chosen. For example, the triplet code for each amino acid can be modified to conform more closely to the preferential codon usage of the host cell's translational machinery, or, for example, to yield a messenger RNA molecule with a longer half-life. Those skilled in the art would readily know what modifications of the nucleotide sequence would be desirable to conform the nucleotide sequence to preferential codon usage or to make the messenger RNA more stable. Such information would be obtainable, for example, through use of computer programs, through review of available research data on codon usage and messenger RNA stability, and through other means known to those of skill in the art.

[0067] Polypeptides corresponding to one or more portions of clusterin, truncated or deleted clusterin as well as fusion proteins in which the full length clusterin or truncated clusterin is fused to an unrelated protein are also within the scope of the invention and can be designed on the basis of the clusterin nucleotide and clusterin amino acid sequences disclosed above. Such fusion proteins include but are not limited to IgFc fusions which stabilize the clusterin polypeptide and prolong half-life in vivo or in in vitro assays; fusions to any amino acid sequence that allows the fusion protein to be anchored to the cell membrane; or fusions to an enzyme, fluorescent protein, or luminescent protein which provide a marker function.

[0068] Additionally, the clusterin gene can be subcloned into a recombinant plasmid such that the gene's open reading frame is translationally fused to an amino-terminal tag consisting of multiple (generally about six) histidine residues. Extracts from cells infected or transfected with such constructs are loaded onto Ni^{2+} nitriloacetic acid-agarose columns and histidine-tagged proteins are selectively eluted with imidazole-containing buffers.

[0069] While clusterin polypeptides can be chemically synthesized (e.g., see Creighton, 1983), large polypeptides derived from the clusterin and the full length clusterin itself may advantageously be produced by recombinant DNA technology using techniques well known in the art for expressing nucleic acids. Such methods can be used to construct expression vectors containing the clusterin nucleotide sequences and appropriate transcriptional and translational control signals. These methods include, for example, in vitro recombinant DNA techniques, synthetic techniques, and in vivo genetic recombination. See, for example, the techniques described in Sambrook et al., 1989, supra, and Ausubel et al., 1989, supra. Alternatively, RNA capable of encoding Clusterin nucleotide sequences may be chemically synthesized using, for example, synthesizers. See, for example, the techniques described in "Oligonucleotide Syn-

thesis", 1984, Gait, M. J., ed., IRL Press, Oxford, which is incorporated by reference herein in its entirety.

[0070] A variety of host-expression vector systems can be utilized to express the clusterin nucleotide sequences of embodiments of the invention. Where clusterin polypeptide is a soluble derivative (e.g., with a deleted TMD), the polypeptide can be recovered from the culture, i.e., from the host cell in cases where clusterin polypeptide is not secreted, and from the culture media in cases where clusterin polypeptide is secreted by the cells. However, the expression systems also encompass engineered host cells that express clusterin or functional equivalents in situ, i.e., anchored in the cell membrane. Purification or enrichment of clusterin from such expression systems can be accomplished using appropriate detergents and lipid micelles and methods well known to those skilled in the art. However, such engineered host cells themselves can be used in situations where it is important not only to retain the structural and functional characteristics of clusterin, but to assess biological activity, e.g., in drug screening assays.

[0071] For long-term, high-yield production of recombinant proteins, stable expression is preferred. For example, cell lines which stably express clusterin sequences can be engineered, for example, as described in SEQ ID NOs: 4-15 and in the examples below. Rather than using expression vectors which contain viral origins of replication, host cells can be transformed with DNA controlled by appropriate expression control elements (e.g., promoter, enhancer sequences, transcription terminators, polyadenylation sites, etc.), and a selectable marker. Following the introduction of the foreign DNA, engineered cells may be allowed to grow for 1-2 days in an enriched media, and then are switched to a selective media. The selectable marker in the recombinant plasmid confers resistance to the selection and allows cells to stably integrate the plasmid into their chromosomes and grow to form foci which in turn can be cloned and expanded into cell lines. This method may advantageously be used to engineer cell lines which express the Clusterin gene product. Such engineered cell lines may be particularly useful in screening and evaluation of compounds that affect the endogenous activity of the Clusterin gene product. A number of selection systems can be used, including but not limited to the herpes simplex virus thymidine kinase (Wigler, et al., 1977), hypoxanthine-guanine phosphoribosyltransferase (Szybalska & Szybalski, 1962), and adenine phosphoribosyltransferase (Lowy, et al., 1980) genes can be employed in tk-, hgpri- or apri-cells, respectively. Also, anti-metabolite resistance can be used as the basis of selection for the following genes: dhfr, which confers resistance to methotrexate (Wigler, et al., 1980; O'Hare, et al., 1981); gpt, which confers resistance to mycophenolic acid (Mulligan & Berg, 1981); neo, which confers resistance to the aminoglycoside G-418 (Colberre-Garapin, et al., 1981); and hygri, which confers resistance to hygromycin (Santerre, et al., 1984).

[0072] The expression systems that can be used for purposes of the embodiments include but are not limited to microorganisms such as bacteria (e.g., *E. coli*, *B. subtilis*) transformed with recombinant bacteriophage DNA, plasmid DNA or cosmid DNA expression vectors containing clusterin nucleotide sequences; yeast (e.g., *Saccharomyces*, *Pichia*) transformed with recombinant yeast expression vectors containing the clusterin nucleotide sequences; insect cell systems infected with recombinant virus expression vectors (e.g., baculovirus) containing the Clusterin

sequences; plant cell systems infected with recombinant virus expression vectors (e.g., cauliflower mosaic virus, CaMV; tobacco mosaic virus, TMV) or transformed with recombinant plasmid expression vectors (e.g., Ti plasmid) containing Clusterin nucleotide sequences; or mammalian cell systems (e.g., COS, CHO, BHK, 293, 3T3) harboring recombinant expression constructs containing promoters derived from the genome of mammalian cells (e.g., metallothionein promoter) or from mammalian viruses (e.g., the adenovirus late promoter; the vaccinia virus 7.5K promoter).

[0073] In bacterial systems, a number of expression vectors may be advantageously selected depending upon the use intended for the clusterin gene product being expressed. For example, when a large quantity of such a protein is to be produced, for the generation of pharmaceutical compositions of clusterin protein or for raising antibodies to the clusterin protein, for example, vectors which direct the expression of high levels of fusion protein products that are readily purified may be desirable. Such vectors include, but are not limited, to the *E. coli* expression vector pUR278 (Ruther et al., 1983), in which the clusterin coding sequence may be ligated individually into the vector in frame with the lacZ coding region so that a fusion protein is produced; pIN vectors (Inouye & Inouye, 1985; Van Heeke & Schuster, 1989); and the like. pGEX vectors may also be used to express foreign polypeptides as fusion proteins with glutathione S-transferase (GST). In general, such fusion proteins are soluble and can easily be purified from lysed cells by adsorption to glutathione-agarose beads followed by elution in the presence of free glutathione. The pGEX vectors are designed to include thrombin or factor Xa protease cleavage sites so that the cloned target gene product can be released from the GST moiety.

[0074] In an insect system, *Autographa californica* nuclear polyhydrosis virus (AcNPV) is used as a vector to express foreign sequences. The virus grows in *Spodoptera frugiperda* cells. The clusterin gene coding sequence may be cloned individually into non-essential regions (for example the polyhedrin gene) of the virus and placed under control of an AcNPV promoter (for example the polyhedrin promoter). Successful insertion of clusterin coding sequence will result in inactivation of the polyhedrin gene and production of non-occluded recombinant virus, (i.e., virus lacking the proteinaceous coat coded for by the polyhedrin gene). These recombinant viruses are then used to infect *Spodoptera frugiperda* cells in which the inserted polynucleotide is expressed (e.g., see Smith et al., 1983 and U.S. Pat. No. 4,215,051).

[0075] In mammalian host cells, a number of viral-based expression systems can be utilized. In cases where an adenovirus is used as an expression vector, the clusterin nucleotide sequence of interest may be ligated to an adenovirus transcription/translation control complex, e.g., the late promoter and tripartite leader sequence. This chimeric gene may then be inserted in the adenovirus genome by in vitro or in vivo recombination. Insertion in a non-essential region of the viral genome (e.g., region E1 or E3) will result in a recombinant virus that is viable and capable of expressing the clusterin gene product in infected hosts (e.g., See Logan & Shenk, 1984). Specific initiation signals may also be important for efficient translation of inserted clusterin nucleotide sequences. These signals include the ATG initiation codon and adjacent sequences. In cases where an entire clusterin gene or cDNA, including its own initiation codon

and adjacent sequences, is inserted into the appropriate expression vector, no additional translational control signals may be needed. However, in cases where only a portion of the clusterin coding sequence is inserted, exogenous translational control signals, including, perhaps, the ATG initiation codon, must be provided. Furthermore, the initiation codon must be in phase with the reading frame of the desired coding sequence to ensure translation of the entire insert. These exogenous translational control signals and initiation codons can be of a variety of origins, both natural and synthetic. The efficiency of expression may be enhanced by the inclusion of appropriate transcription enhancer elements, transcription terminators, etc. (See Bitter, et al., 1987).

[0076] In yeast, a number of vectors containing constitutive or inducible promoters may be used. For a review, see Current Protocols in Molecular Biology, 1988; Grant, et al., 1987; Wu & Grossman, 1987; Bitter, 1987; and "The Molecular Biology of the Yeast *Saccharomyces*", 1982.

[0077] In cases where plant expression vectors are used, the expression of the coding sequence may be driven by any of a number of promoters. For example, viral promoters such as the 35S RNA and 19S RNA promoters of CaMV (Brisson et al., 1984), or the coat protein promoter of TMV (Takamatsu et al., 1987) may be used; alternatively, plant promoters such as the small subunit of RUBISCO (Coruzzi et al., 1984; Broglie et al., 1984); or heat shock promoters, e.g., soybean hsp17.5-E or hsp17.3-B (Gurley et al., 1986) may be used. These constructs can be introduced into plant cells using Ti plasmids, Ri plasmids, plant virus vectors, direct DNA transformation, microinjection, electroporation, etc. For reviews of such techniques see, for example, Methods for Plant Molecular Biology 1988; and Grierson & Corey, 1988.

[0078] In cases where an adenovirus is used as an expression vector, the nucleotide sequence of interest can be ligated to an adenovirus transcription/translation control complex, e.g., the late promoter and tripartite leader sequence. This chimeric gene can then be inserted in the adenovirus genome by in vitro or in vivo recombination. Insertion in a non-essential region of the viral genome (e.g., region E1 or E3) will result in a recombinant virus that is viable and capable of expressing the gene product of interest in infected hosts (e.g., See Logan & Shenk, 1984). Specific initiation signals such as those described above can also be important for efficient translation of inserted nucleotide sequences of interest.

[0079] In addition, a host cell strain can be chosen that modulates the expression of the inserted sequences, or modifies and processes the gene product in the specific fashion desired. Such modifications (e.g., glycosylation) and processing (e.g., cleavage) of protein products can be important for the function of the protein. Different host cells have characteristic and specific mechanisms for the post-translational processing and modification of proteins and gene products. Appropriate cell lines or host systems can be chosen to ensure the correct modification and processing of the foreign protein expressed. To this end, eukaryotic host cells which possess the cellular machinery for proper processing of the primary transcript, glycosylation, and phosphorylation of the gene product can be used. Such mammalian host cells include but are not limited to CHO, VERO, BHK, HeLa, COS, MDCK, 293, 3T3, WI38 and U937 cells.

II. THERAPEUTIC FORMULATION AND ADMINISTRATION

[0080] Therapeutics comprising clusterin polypeptides and/or nucleic acids encoding clusterin polypeptides, such as those described in SEQ ID NOs: 4-15 can be administered to a patient at therapeutically effective doses of pharmaceutical preparations used to treat or ameliorate conditions such as, but not limited to, cardiovascular disease may be hypertension, hyperlipidemia, hypercholesterolemia, hyperglycemia, hypertension, atherosclerosis and atherosclerosis-associated ischemic heart failure, stenosis, calcification of cardiovascular tissues, stroke, myocardial infarction or cerebral infarction. In some aspects, the cardiovascular disease is hyperlipidemia, hypercholesterolemia or atherosclerosis. In still further aspects, the cardiovascular disease may be diabetes. In various aspects, an effective amount of a clusterin composition may be an amount effective to reduce blood cholesterol, reduce blood glucose, reduce blood triglyceride, increase efflux of intracellular cholesterol, and/or increase vascular or cardiac cell survival. In some aspects an effective amount of a clusterin composition provides enhancement or promotion of cell survival and growth against cytotoxic or cytostatic factors, including but not limited to, oxysterols, oxidized lipoproteins, and proinflammatory cytokines, for examples those induced by alcoholism. In further aspects a therapeutically effective dose refers to that amount of the compound sufficient to result in any amelioration or retardation of disease symptoms or progression in a mammal with acute vascular syndromes, to prevent and treat degeneration, stenosis and calcification of cardiovascular tissues, comprising valve tissues. In further aspects, clusterin derived polypeptides can be used to prevent or treat male infertility by, for example, supporting sperm maturation and development, as has been described for native clusterin.

[0081] Toxicity and therapeutic efficacy of such clusterin derived compositions can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD_{50} (the dose lethal to 50% of the population) and the ED_{50} (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD_{50}/ED_{50} . Compositions which exhibit large therapeutic indices are preferred. While compositions that exhibit toxic side effects may be used, care should be taken to design a delivery system that targets such compounds to the site of affected tissue in order to minimize potential damage to uninfected cells and, thereby, reduce side effects.

[0082] Data obtained from the cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compositions are preferably within a range of circulating concentrations that include the ED_{50} with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. For any compositions used in the methods of the embodiments, the therapeutically effective dose can be estimated initially from cell culture assays. A dose may be formulated in animal models to achieve a circulating plasma concentration range that includes the IC_{50} (i.e., the concentration of the test compound which achieves a half-maximal inhibition of symptoms) as determined in cell culture. Such information can be used to more accurately determine useful doses in

humans. Levels in plasma may be measured, for example, by high performance liquid chromatography.

[0083] When the therapeutic treatment of disease is contemplated, the appropriate dosage can also be determined using animal studies to determine the maximal tolerable dose, or MTD, of a bioactive agent per kilogram weight of the test subject. In general, at least one animal species tested is mammalian. Those skilled in the art regularly extrapolate doses for efficacy and avoiding toxicity to other species, including human. Before human studies of efficacy are undertaken, Phase I clinical studies in normal subjects help establish safe doses.

[0084] Additionally, the bioactive agent (e.g., a clusterin polypeptide) may be complexed with a variety of well-established compounds or structures that, for instance, enhance the stability of the bioactive agent, or otherwise enhance its pharmacological properties (e.g., increase in vivo half-life, reduce toxicity, etc.).

[0085] Pharmaceutical compositions for use in accordance with the present embodiments can be formulated in conventional manner using one or more physiologically acceptable carriers or excipients.

[0086] The above therapeutic agents will be administered by any number of methods known to those of ordinary skill in the art including, but not limited to, administration by inhalation; by subcutaneous (sub-q), intravenous (I.V.), intraperitoneal (I.P.), intramuscular (I.M.), or intrathecal injection; or as a topically applied agent (transderm, ointments, creams, salves, eye drops, and the like). Thus, the compositions can be formulated for administration by inhalation or insufflation (either through the mouth or the nose) or oral, buccal, parenteral or rectal administration.

[0087] For oral administration, the pharmaceutical compositions may take the form of, for example, tablets or capsules prepared by conventional means with pharmaceutically acceptable excipients such as binding agents (e.g., pregelatinized maize starch, polyvinylpyrrolidone or hydroxypropyl methylcellulose); fillers (e.g., lactose, microcrystalline cellulose or calcium hydrogen phosphate); lubricants (e.g., magnesium stearate, talc or silica); disintegrants (e.g., potato starch or sodium starch glycolate); or wetting agents (e.g., sodium lauryl sulphate). The tablets can be coated by methods well known in the art. Liquid preparations for oral administration may take the form of, for example, solutions, syrups or suspensions, or they may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations may be prepared by conventional means with pharmaceutically acceptable additives such as suspending agents (e.g., sorbitol syrup, cellulose derivatives or hydrogenated edible fats); emulsifying agents (e.g., lecithin or acacia); non-aqueous vehicles (e.g., almond oil, oily esters, ethyl alcohol or fractionated vegetable oils); and preservatives (e.g., methyl or propyl-p-hydroxybenzoates or sorbic acid). The preparations may also contain buffer salts, flavoring, coloring and sweetening agents as appropriate. Preparations for oral administration can be suitably formulated to give controlled release of the active composition.

[0088] For buccal administration the compositions can take the form of tablets or lozenges formulated in conventional manner.

[0089] For administration by inhalation, the compositions for use according to the embodiments are conveniently delivered in the form of an aerosol spray presentation from

pressurized packs or a nebulizer, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of e.g. gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch.

[0090] The compositions can be formulated for parenteral administration by injection, e.g., by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g., in ampules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

[0091] The compositions can also be formulated for rectal administration such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glycerides.

[0092] In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt. The compositions may, if desired, be presented in a pack or dispenser device which may contain one or more unit dosage forms containing the active ingredient. The pack may for example comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration.

III. EXAMPLES

[0093] The following examples are included to demonstrate preferred embodiments of the invention. It should be appreciated by those of skill in the art that the techniques disclosed in the examples which follow represent techniques discovered by the inventor to function well in the practice of the invention, and thus can be considered to constitute preferred modes for its practice. However, those of skill in the art should, in light of the present disclosure, appreciate that many changes can be made in the specific embodiments which are disclosed and still obtain a like or similar result without departing from the spirit and scope of the invention.

Example 1—General Methods and Results

[0094] Preparation of native and recombinant clusterin analogs. To treat stem cells with clusterin, two forms of clusterin were prepared. Native clusterin (NCBI Accession No. NM_0134921, incorporated herein by reference). Clusterin was prepared from blood plasma using affinity chromatography with anti-clusterin antibody. In addition, a plasmid was constructed in which mouse or human clusterin cDNA is fused with a His-tag and inserted under a promoter that will be activated in a mammalian (e.g., human cells) or

non-mammalian (e.g., bacteria) cell. Schematic presentation of the plasmid pCluAg with a clusterin cDNA-His insert is shown in FIG. 1. pCluAg was constructed by inserting clusterin-TR/EK-His cDNA using a TOPO plasmid (Invitrogen). Recombinant clusterin produced by transfected mammalian cells (e.g., human 293 cells) or bacterial cells (*E. coli*) was purified. Different clusterin analog cDNAs were generated by RT-PCR and sequences predicting the encoding of different clusterin analog polypeptide sequences (FIGS. 2A-D and Table 1, below).

TABLE 1

Sequence Descriptions	
Description:	SEQ ID NO:
CluAg-a (recombinant Clusterin)	1
CluAg-b (recombinant Clusterin-ATMD)	2
CluAg-c (recombinant Clusterin-ATMD-ANLS)	3
CluAg-Ia (HisTR-Clusterin)	4
CluAg-Ib (HisTR-Clusterin-ATMD)	5
CluAg-Ic (HisTR-Clusterin-ATMD-ANLS)	6
CluAg-IIa (Clusterin-TR-His)	7
CluAg-IIb (Clusterin-TR-His-ATMD)	8
CluAg-IIc (Clusterin-TR-His-ATMD-ANLS)	9
CluAg-IIIa (HisEK-Clusterin)	10
CluAg-IIIb (HisEK-Clusterin-ATMD)	11
CluAg-IIIc (HisEK-Clusterin-ATMD-ANLS)	12
CluAg-IVa (HisEK-Clusterin)	13
CluAg-IVb (HisEK-Clusterin-ATMD)	14
CluAg-IVc (HisEK-Clusterin-ATMD-ANLS)	15

[0095] Analysis of recombinant clusterin analog protein properties. The purity of recombinant clusterin analogs was examined by polyacrylamide gel electrophoresis. Clusterin analog proteins are isolated from pCluAg transfected cells, stored in solution or in dry powder, and loaded into SDS-PAGE. After electrophoresis, fractions of proteins will be electrotransferred onto a PVDF membrane, and probed with anti-clusterin antibodies. The membrane will be developed by using a chemiluminescence kit. Single bands of clusterin analog CluAg-I are visualized in SDS-PAGE stained with Commassia blue dye G250 (FIG. 3) and confirmed by immunoblotting with anti-Clu antibody (FIG. 4). Furthermore, the analogs were examined by ion exchange chromatography, showing a clear eluent of clusterin analogs (FIG. 5).

[0096] Clusterin analogs in the HDL fraction in mice. Apolipoprotein E deficient (ApoE^{-/-}) mice are widely used murine mode for atherosclerosis. To evaluate the levels of clusterin association with ApoAI in the blood, an immunoprecipitation method was developed with monoclonal antibody against apoAI and clusterin. In the plasma of blood from CluAg injected mice, ApoAI antibody co-precipitates ApoAI and CluAg, indicating that ApoAI is binding to CluAg (FIG. 6). To further confirm the presence of CluAg and compare them to native clusterin, CluAg was added to SMC culture and incubated for 2 days with 5% serum containing native {00939480}-29-clusterin in the presence of oxLDL. Supernatants of the cultures were subjected to analysis of poly-His tag in CluAg. In the culture with or without oxLDL, CluAg apparently are free of the poly-His tag while CluAg (FIG. 7) in cell free cultures contains the poly-His tag (FIG. 7).

[0097] Clusterin analog regulation of vascular cell proliferation in culture. An in vitro system was first employed to

examine the protective effects of clusterin analogs on survival and growth of vascular cells. Cells were treated with clusterin analogs at different concentrations in the cultures with or without oxidized LDL. After 2-4 days of stimulation, cell survival and proliferation were examined using a combination of techniques including flow cytometry, fluorescent microscopy, and radioactive isotope labeling, as described below. Control experiments were set up using other types of proteins, such as bovine albumin. FIGS. 8 and 9 show growth curves of SMCs treated with or without CluAg (1-6 $\mu\text{g}/\text{ml}$) in the presence or absence of oxLDL (50 $\mu\text{g}/\text{ml}$). Treatment with the analog increased SMC proliferation even under the condition of oxLDL exposure. CluAg dose-dependently increases SMC proliferation even in the presence of oxLDL (FIGS. 8 and 9).

[0098] Analysis of lipid profiles, cholesterol, glucose, and triglyceride in wild type and atherosclerosis prone mice injected with clusterin. Weekly injection of CluAg (30-40 μg) for 3 months was conducted in wild type (WT) and apoE-null mice. The blood samples were collected, during tail DNA sampling, from the mice injected with CluAg. Serum was prepared from the blood samples. Cholesterol levels, lipoprotein profiles and clusterin concentrations were determined respectively. In brief, serum diluted in PBS was incubated in a 96 well plate coated with a rabbit polyclonal antibody to clusterin. After incubation and washing in PBS, bound clusterin was detected by incubating with mouse monoclonal antibody to clusterin. Goat anti-mouse IgG conjugated with peroxidase was used as the second antibody. Cholesterol and HDL was determined in the laboratory of Department of Laboratory Medicine. The ratio of clusterin vs. HDL was calculated after normalization with the lipid content. In addition to ELISA, immunoblotting assays were performed to verify the results. CluAg injection significantly reduced blood levels of total cholesterol, glucose, triglyceride, and LDL in apoE-null mice but no changes were found in WT mice (FIG. 10).

[0099] Analysis of blood pressure. Blood pressure measurement was performed in the tail artery using a tail-cuff method. Each measurement was repeated 3 times to ensure reproducibility. Weekly injection of CluAg (30-40 μg) for 3 months led to reduction in the tail arterial blood pressure of both systolic and diastolic phases was conducted in apoE-null mice (FIG. 11, left panel) but no changes in blood pressure were found in WT mice (FIG. 11, right panel).

[0100] Echocardiography of mice injected with CluAg. After weekly injection of CluAg, morphological and functional changes were monitored using echocardiography. B- and M-Mode echocardiography was performed one, 6, 10, and 18 weeks after injection. The echocardiography studies were conducted actually using ultrasonography as the murine heart is small. Mice were anesthetized with ketamine and xylazine, chests shaved and a layer of acoustic coupling gel will be applied to the thorax. A dynamically focused 9-MHz annual array transducer was applied using a warmed saline bag as a standoff. All echo studies were performed using a state of the art echo machine (HP Model Sonos 5500 HP). Area fractions were determined by planimetry of diastolic and systolic volumes in parasternal short axis. The LV end-diastolic and end-systolic dimensions were measured using the M-Mode from >3 beats by two independent investigators blinded to the research animals. LVEF (left ventricular ejection fraction) was calculated as follows: $\text{LVEF} = [(\text{LVIDd}) - (\text{LVIDS})] / (\text{LVIDd})$, LVIDd: end-diastolic

left ventricular internal diameter; LVIDS: end-systolic left ventricular diameter. FIG. 12 shows ultrasound images of murine hearts with or without CluAg injection in both 2D and M-mode. FIG. 13 demonstrates that weekly administration of CluAg (30-40 $\mu\text{g}/\text{mouse}$) increases ejection fraction in the hearts of ApoE^{-/-} atherosclerosis-prone mice but no changes in wild type mice with the same dose CluAg. **[0101]** Oil red O staining of aortic wall. Mice were sacrificed 3 months after CluAg treatment. Aortas were opened across the long axis and fixed in 10% buffered formalin for histological evaluation. Aortas were stained with Oil Red O solution for assessing neutral lipid contents. Little staining was detected in WT aortas. However, compared to WT aortas, the ApoE-null aortas were stained very intensively with Oil Red O, which visualizes atherosclerotic plaques. Decreased Oil Red O stains were found in ApoE-null mice injected with CluAg protein (FIG. 14).

[0102] Analysis of aortic tissue and cell calcification. Aortic smooth muscle cells (SMCs) were incubated with CluAg in the presence of sodium phosphate (3.6 mmol/L) for 6 days. Alizarin red S (Sigma, St. Louis, Mo., USA) staining assesses calcium deposition in a reaction in which Alizarin red S dye binds with Ca ions in cell layer matrix. Cells were fixed with 2% paraformaldehyde and stained with 1% Alizarin red S (pH 4.2). The culture plates were photographed under a light microscope and assessed for the mineralized nodules which shown as red (FIG. 15). CluAg treatment markedly reduces calcification in SMCs. To determine if clusterin receptors mediate the inhibitory effect of CluAg on phosphate induced calcification in SMCs, snRNA for ApoER2 and VLDLR, two known receptors for clusterin, was constructed and used to knock down ApoER2 and VLDLR. CluAg treatment had no inhibitory effect on calcification in SMCs with the receptor knockdown with the snRNA (FIG. 15). This result was confirmed by direct calcium assays (FIGS. 16A-C). For assessing in vivo calcification of aortic tissue, Alizarin red S staining was performed in the aortas of WT and apoE-null mice with CluAg injection. Sections of aortas from apoE-null mice but not wild type control mice show the development of atherosclerotic plaques. Calcium deposits were highly abundant in ApoE-null mice. Treatment with CluAg reduces calcification in ApoE-null mice (FIGS. 18A-H).

[0103] Immunohistochemistry of aortic tissue. In order to assess whether CluAg injection alters expression of calcification-regulatory proteins, such as bone morphogenic protein (BMP)-2, in the aortic tissue, aortic sections were stained with anti-BMP-2 antibodies. Immunofluorescence for BMP-2 was developed with rhodamine-conjugated second antibodies (Sigma, St. Louis, Mo.). The slides were mounted in the Vectashield mounting medium with 4',6-diamidino-2-phenylindole (DAPI) (Vector, Burlingame, Calif.), and examined under an Olympus fluorescence microscope. Intensive BMP-2 immunofluorescence was found in ApoE-null aorta but CluAg injection reduced the intensity of the BMP-2 fluorescence.

Example 2—Further Analysis of Clusterin In Vivo Methods and Materials

[0104] Cell culture. Aortic VSMCs were isolated from 5- to 6-week-old male ApoE^{-/-} or WT mice with C57BL/6 background. The cells were maintained in Dulbecco's Modified Eagle's Medium (DMEM; Invitrogen, Carlsbad, Calif., USA) supplemented with 10% fetal bovine serum (FBS)

with 100 ng/mL penicillin and streptomycin (Invitrogen) at 37° C. in a humidified atmosphere with 5% CO₂. Calcification medium was made by adding NaH₂PO₄ (pH adjusted to 7.4) into 5% FBS medium to obtain a final concentration of 3.6 mM inorganic phosphate. VSMCs from passage 5-10 were used. Apo J Medium was replaced every 2 or 3 days for up to 9 days. Cells maintained in regular culture medium with 0.9 mM phosphate were used as controls.

[0105] Lentivirus infection and selection. To generate a stable (long-term) knockdown of ApoER2 or VLDLR gene expression in VSMCs, the VSMCs were infected with lentivirus particles containing a pLKO.1 vector with the information to express a shRNA against mouse ApoER2 or VLDLR. This plasmid also has a puromycin resistance gene, thereby allowing for the selection of cells stably expressing desired shRNA by addition of puromycin into culture medium. The optimal puromycin concentration for VSMCs before initiating the experiments (titration assay) was determined to be 1 µg/mL. To perform lentivirus infection, cells approximately 80% confluent were used. 1.5 ml of fresh culture media containing virus was added onto cells. Fresh media was changed every 3 days 48 h after infection. To select the infected cells, selection media containing puromycin was used for culture 48 h after infection. ApoER2 and VLDLR knockdown were monitored by western blot analysis and was achieved after four passages. One plate of cells infected with pLKO.1 vector with shRNA verified to contain no homology to known mammalian genes was maintained in parallel. This plate served as a negative control for following experiments.

[0106] Calcium determination. Calcium content in VSMCs was determined by colorimetric calcium detection kit (Abcam). Cells were washed with PBS and then incubated with 0.6M HCl under 37° C. overnight to be decalcified. The calcium content in the supernatants was measured using spectrometer. Then cells were lysed with 0.1 mol/L NaOH/0.1% SDS. Protein content was determined with BCA protein assay kit (Thermo Scientific) and calcium content was normalized to total protein content.

[0107] Alizarin Red S staining. Cells were fixed in 2.5% glutaraldehyde and incubated with 2% Alizarin Red S under 37° C. for 15 minutes. Then VSMCs were rinsed with PBS three times. Calcium mineralization visualized by red staining was observed under microscope.

[0108] Real-time RT-PCR. Total RNA was extracted from VSMCs using Trizol (Invitrogen). 4 µg total RNA was used for cDNA synthesis in a reaction mixture of 20 µL with SuperScript III First-Strand Synthesis SuperMix (Invitrogen). Real-time PCR amplification was performed with IQTM SYBR Green Supermix (Bio-Rad) in a ICYCLERIQTM thermocycler (Bio-Rad). The following primers sets were used: ALP, 5'-CACAAATATCAAGGATATCGACGTGA-3' (sense; SEQ ID NO: 18) and 5'-ACATCAGTCTCTTCTTCGGGTACA-3' (antisense; SEQ ID NO: 19); BMP-2, 5'-TTGTATGTGGACTTCAGTGATGTG-3' (sense; SEQ ID NO: 20) and 5'-AGTTCAGGTGGTCAGCAAGG-3' (antisense; SEQ ID NO: 21); Osteopontin, 5'-TGGCTATAGGATCTGGGTGC-3' (sense; SEQ ID NO: 22) and 5'-ATTGCTTTTGCTGTTTGG-3' (antisense; SEQ ID NO: 23); and Runx2, 5'-TTACCTACACCCGCCAGTC-3' (sense; SEQ ID NO: 24) and 5'-TGCTGGTCTGGAAGGGTCC-3' (antisense; SEQ ID NO: 25).

[0109] Western blot. Proteins were isolated from VSMCs using RIPA buffer (Pierce) containing protease and phosphatase inhibitors (Sigma-Aldrich). Proteins separated on 10% SDS-Polyacrylamide gel were transferred to polyvinylidenedifluoride (PVDF) membranes (Millipore) and Western blot was performed using the standard protocol. Membranes were blocked with 5% bovine serum albumin (BSA) in TBS containing 0.1% Tween-20 (TBST). Primary antibodies were diluted in 3% BSA [goat anti-ApoJ 1:1000 (Santa Cruz Biotechnology), rabbit anti-Runx2 1:1000 (Cell Signaling), goat anti-SM22α 1:5000 (Abcam), rabbit anti-αSMA 1:1600 (Abcam), rabbit anti-GAPDH 1:30000 (Abcam)] and were detected using HRP-conjugated secondary antibodies [donkey anti-goat 1:5000 (Santa Cruz Biotechnology), goat anti-rabbit 1:20000 (Santa Cruz Biotechnology)] diluted in 3% BSA in TBST.

[0110] Data analysis and statistics. Western blot results were analyzed by densitometry using Scion Image (Scion Corp). Real-time polymerase chain reaction data was quantified using EXCEL[®] software (Microsoft Corp). Values were graphed as mean±SD of at least triplicates determinations. Statistics (t test and ANOVA) were performed using Graphpad software (Graphpad Software Inc). A value of P<0.05 was considered statistically significant.

[0111] Results

[0112] Apo J attenuates calcification in both ApoE^{-/-} and WT smooth muscle cells. Because ApoE^{-/-} mice are more prone to vascular calcification than WT ones, the responses of ApoE^{-/-} and WT VSMCs to inorganic phosphate (Pi) were examined as well as the effect of Apo J on calcification of these two groups of cells. The study shows that when induced with Pi, ApoE^{-/-} VSMCs exhibited much higher levels of calcification on day 6 of treatment than WT cells. The addition of Apo J (6 m/mL) into culture medium reduced calcium level in both groups, with a more dramatic inhibition on calcification in ApoE^{-/-} VSMCs, shown by both Alizarin S staining (FIGS. 19A and 20A) and calcium assay (FIGS. 19B and 20B). Therefore, additional experiments were conducted using ApoE^{-/-} VSMCs. Various concentrations (3 m/mL, 6 m/mL, 9 m/mL, and 12 m/mL) of Apo J were then applied to test if the suppressive effect of the Apo J on vascular cell calcification is dose dependent. The data demonstrated that 6 m/mL Apo J in culture is sufficient to significantly decrease calcium deposition level in ApoE^{-/-} VSMCs, while 12 µg/mL is incapable to further attenuate calcification compared to 9 m/mL group (FIG. 20B), possibly due to saturation of Apo J receptors.

[0113] Apo J modulates osteogenesis-related genes during calcification in ApoE^{-/-} VSMCs. To confirm the effect of Apo J on calcification, the change of osteogenesis regulator Runx2 was assessed, as well as the smooth muscle lineage markers SM22α and αSMA, with or without added Apo J during calcification of ApoE^{-/-} VSMCs, which were treated with Pi for 6 days. Concentrations from 3 m/mL to 12 µg/mL of Apo J were used. Equal volumes (40 µL) of culture medium were immunoblotted with Apo J antibody at the endpoint of the experiment to verify the existence of Apo J protein in medium. The data suggested that both the mRNA level and the native secreted form of Apo J increased in calcifying cells (FIGS. 21A and 22A-B); addition of Apo J resulted in a much stronger signal detected by Apo J antibody, indicating Apo J in medium was sustained on a potent level compared to control group and not subject to bulk degradation through the period of experiment. It was

found Runx2 was increased in response to Pi and this increase was attenuated in all Apo J treated groups, with 6 mM Apo J sufficient to keep Runx2 protein expression down to approximately the same level in uncalcified cells. No significant differences of Runx2 levels were observed between 6 μ g/mL, 9 μ g/mL, and 12 μ g/mL groups (FIGS. 21A-21B). SM22 α and α SMA protein expressions were downregulated by Pi compared to control group; Apo J rescued the decreased level of SM22 α and α SMA at a concentration as small as 3 μ g/mL (FIGS. 21A, 21C and 21D). mRNA expressions of osteogenic genes including Runx2, BMP-2, OPN and ALP were increased during calcification while Apo J weakened this effect (FIGS. 23A-23D).

[0114] Knockdown of apolipoprotein-E receptor 2 (ApoER2 or very low density lipoprotein receptor (VLDLR) abolishes the inhibitory effect of Apo J on calcification. Because 6 μ g/mL of Apo J has been shown to inhibit calcium mineralization as well as activation of calcification-associated genes, this same concentration of Apo J was used to treat VSMCs in remaining experiments. Apo J receptors ApoER2 and VLDLR were knocked down in ApoE $^{-/-}$ VSMCs by lentiviral shRNA. Knockdown effect was confirmed by western blots and qRT-PCR. The data demonstrated that the knockdown of VLDLR or ApoER2 itself didn't change calcification level under the influence of Pi. The negative control shRNA didn't alter the sensitivity of VSMCs to Apo J in calcification and calcium amount still decreased in Apo J treated negative control group. In contrast, VLDLR or ApoER2 shRNA mediated knockdown of these receptors eliminated the repressive impact of Apo J on calcification, shown by Alizarin S and quantitative calcium assay (FIGS. 24A-24F), indicating Apo J functions through ApoER2 and VLDLR to regulate calcification process.

[0115] Knockdown of ApoER2 or VLDLR partially abolishes the effect of Apo J on osteogenesis related genes in calcification. Knockdown of ApoER2 or VLDLR weakened the negative regulation of Apo J on the enhanced expression of Runx2 in calcifying ApoE $^{-/-}$ VSMCs on protein level (FIGS. 25A and 26C and mRNA level (FIG. 27C). SM22 α protein expression decreased in Pi treated cells with ApoER2 or VLDLR knockdown despite the presence of Apo J in culture medium (FIGS. 25B and 26D). Osteogenic markers alkaline phosphatase (ALP), bone morphogenic protein (BMP)-2, and Osteopontin (OPN) remained upregulated compared to uncalcified control group in ApoER2 or VLDLR knockdown cells; no significant differences were found in mRNA expressions of these markers between Apo J treated group and group without Apo J (FIGS. 27D-27F). This implies that knockdown of ApoER2 or VLDLR partially abolished the effect of Apo J on osteogenesis related genes in calcification in ApoE $^{-/-}$ VSMCs.

[0116] Studies were undertaken to assess the effect recombinant Clusterin on blood coagulation. ApoJ mutants with a peptide comprising a thrombin-specific cleavage site (e.g., Clusterin-TRHis, Clusterin- Δ TMD-TRHis, Clusterin- Δ TMD- Δ NLS-TRHis, HisTR-Clusterin- Δ TMD- Δ NLS, or HisTR-Clusterin- Δ TMD) exert an anti-coagulation effect by competitively inhibiting the conversion of fibrinogen into time of pig blood with (A) or without (B) ApoJ mutant. Fresh blood (0.3 ml) from adult pig at different dilutions (0, $\frac{1}{2}$, and $\frac{1}{4}$ in PBS) was mixed with recombinant ApoJ mutant at 10 or 10 mg/ml, incubated at 37° C., and then subjected to rotational thromboelastometry (ROTEM™). a, clotting

time; b, Alpha Angle clotting rapidity; c, maximum clot firmness; and d, maximum clot firmness at 10 minutes. Data represented from average of two separate experiments with the ApoJ mutant, Clusterin- Δ TMD-TRHis (SEQ ID NO: 8). Thus, these data indicate that recombinant Clusterin with thrombin-cleavable peptide target, such as Clusterin-TRHis, Clusterin- Δ TMD-TRHis, Clusterin- Δ TMD- Δ NLS-TRHis, HisTR-Clusterin- Δ TMD- Δ NLS, or HisTR-Clusterin- Δ TMD can exert an anti-coagulation effect by competitively inhibiting the conversion of fibrinogen into fibrin and thus decreasing clot firmness and increasing the clotting time of blood. Thus, recombinant Clusterin may serve as a multi-functional agent for the treatment of patients who suffer from a heart disease, such as heart disease complicated by hypercholesterolemia, hypertension, hyperglycemia and thrombogenesis.

* * *

[0117] All of the methods disclosed and claimed herein can be made and executed without undue experimentation in light of the present disclosure. While the compositions and methods of this invention have been described in terms of preferred embodiments, it will be apparent to those of skill in the art that variations may be applied to the methods and in the steps or in the sequence of steps of the method described herein without departing from the concept, spirit and scope of the invention. More specifically, it will be apparent that certain agents which are both chemically and physiologically related may be substituted for the agents described herein while the same or similar results would be achieved. All such similar substitutes and modifications apparent to those skilled in the art are deemed to be within the spirit, scope and concept of the invention as defined by the appended claims.

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<223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 1

Met Met Lys Thr Leu Leu Leu Phe Val Gly Leu Leu Leu Thr Trp Glu
1             5             10             15

Ser Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn Glu Leu Gln
20            25            30

Glu Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu Ile Gln Asn
35            40            45

Ala Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu Lys Thr Asn
50            55            60

Glu Glu Arg Lys Thr Leu Leu Ser Asn Leu Glu Glu Ala Lys Lys Lys
65            70            75            80

Lys Glu Asp Ala Leu Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys
85            90            95

Glu Leu Pro Gly Val Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu
100           105           110

Cys Lys Pro Cys Leu Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val
115           120           125

Cys Arg Ser Gly Ser Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu
130           135           140

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Asn	Gln	Ser	Ser	Pro	Phe	Tyr	Phe	Trp	Met	Asn	Gly	Asp	Arg	Ile	Asp
145					150					155					160
Ser	Leu	Leu	Glu	Asn	Asp	Arg	Gln	Gln	Thr	His	Met	Leu	Asp	Val	Met
				165					170					175	
Gln	Asp	His	Phe	Ser	Arg	Ala	Ser	Ser	Ile	Ile	Asp	Glu	Leu	Phe	Gln
			180						185					190	
Asp	Arg	Phe	Phe	Thr	Arg	Glu	Pro	Gln	Asp	Thr	Tyr	His	Tyr	Leu	Pro
		195					200					205			
Phe	Ser	Leu	Pro	His	Arg	Arg	Pro	His	Phe	Phe	Phe	Pro	Lys	Ser	Arg
	210					215					220				
Ile	Val	Arg	Ser	Leu	Met	Pro	Phe	Ser	Pro	Tyr	Glu	Pro	Leu	Asn	Phe
225					230					235					240
His	Ala	Met	Phe	Gln	Pro	Phe	Leu	Glu	Met	Ile	His	Glu	Ala	Gln	Gln
			245						250					255	
Ala	Met	Asp	Ile	His	Phe	His	Ser	Pro	Ala	Phe	Gln	His	Pro	Pro	Thr
			260					265					270		
Glu	Phe	Ile	Arg	Glu	Gly	Asp	Asp	Asp	Arg	Thr	Val	Cys	Arg	Glu	Ile
		275					280					285			
Arg	His	Asn	Ser	Thr	Gly	Cys	Leu	Arg	Met	Lys	Asp	Gln	Cys	Asp	Lys
	290					295					300				
Cys	Arg	Glu	Ile	Leu	Ser	Val	Asp	Cys	Ser	Thr	Asn	Asn	Pro	Ser	Gln
305					310					315					320
Ala	Lys	Leu	Arg	Arg	Glu	Leu	Asp	Glu	Ser	Leu	Gln	Val	Ala	Glu	Arg
			325					330						335	
Leu	Thr	Arg	Lys	Tyr	Asn	Glu	Leu	Leu	Lys	Ser	Tyr	Gln	Trp	Lys	Met
			340					345					350		
Leu	Asn	Thr	Ser	Ser	Leu	Leu	Glu	Gln	Leu	Asn	Glu	Gln	Phe	Asn	Trp
		355					360					365			
Val	Ser	Arg	Leu	Ala	Asn	Leu	Thr	Gln	Gly	Glu	Asp	Gln	Tyr	Tyr	Leu
	370					375					380				
Arg	Val	Thr	Thr	Val	Ala	Ser	His	Thr	Ser	Asp	Ser	Asp	Val	Pro	Ser
385					390					395					400
Gly	Val	Thr	Glu	Val	Val	Val	Lys	Leu	Phe	Asp	Ser	Asp	Pro	Ile	Thr
			405					410						415	
Val	Thr	Val	Pro	Val	Glu	Val	Ser	Arg	Lys	Asn	Pro	Lys	Phe	Met	Glu
			420					425					430		
Thr	Val	Ala	Glu	Lys	Ala	Leu	Gln	Glu	Tyr	Arg	Lys	Lys	His	Arg	Glu
		435					440					445			

Glu

<210> SEQ ID NO 2
 <211> LENGTH: 436
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 2

Thr	Trp	Glu	Ser	Gly	Gln	Val	Leu	Gly	Asp	Gln	Thr	Val	Ser	Asp	Asn
1			5						10					15	
Glu	Leu	Gln	Glu	Met	Ser	Asn	Gln	Gly	Ser	Lys	Tyr	Val	Asn	Lys	Glu
		20					25						30		
Ile	Gln	Asn	Ala	Val	Asn	Gly	Val	Lys	Gln	Ile	Lys	Thr	Leu	Ile	Glu

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35	40	45
Lys Thr Asn Glu Glu Arg	Lys Thr Leu Leu Ser	Asn Leu Glu Glu Ala
50	55	60
Lys Lys Lys Lys Glu Asp	Ala Leu Asn Glu Thr	Arg Glu Ser Glu Thr
65	70	75
Lys Leu Lys Glu Leu Pro	Gly Val Cys Asn Glu Thr	Met Met Ala Leu
85	90	95
Trp Glu Glu Cys Lys Pro	Cys Leu Lys Gln Thr	Cys Met Lys Phe Tyr
100	105	110
Ala Arg Val Cys Arg Ser	Gly Ser Gly Leu Val	Gly Arg Gln Leu Glu
115	120	125
Glu Phe Leu Asn Gln Ser	Ser Pro Phe Tyr Phe	Trp Met Asn Gly Asp
130	135	140
Arg Ile Asp Ser Leu Leu	Glu Asn Asp Arg Gln	Gln Thr His Met Leu
145	150	155
Asp Val Met Gln Asp His	Phe Ser Arg Ala Ser	Ser Ile Ile Asp Glu
165	170	175
Leu Phe Gln Asp Arg Phe	Phe Thr Arg Glu Pro	Gln Asp Thr Tyr His
180	185	190
Tyr Leu Pro Phe Ser Leu	Pro His Arg Arg Pro	His Phe Phe Phe Pro
195	200	205
Lys Ser Arg Ile Val Arg	Ser Leu Met Pro Phe	Ser Pro Tyr Glu Pro
210	215	220
Leu Asn Phe His Ala Met	Phe Gln Pro Phe Leu	Glu Met Ile His Glu
225	230	235
Ala Gln Gln Ala Met Asp	Ile His Phe His Ser	Pro Ala Phe Gln His
245	250	255
Pro Pro Thr Glu Phe Ile	Arg Glu Gly Asp Asp	Asp Arg Thr Val Cys
260	265	270
Arg Glu Ile Arg His Asn	Ser Thr Gly Cys Leu	Arg Met Lys Asp Gln
275	280	285
Cys Asp Lys Cys Arg Glu	Ile Leu Ser Val Asp	Cys Ser Thr Asn Asn
290	295	300
Pro Ser Gln Ala Lys Leu	Arg Arg Glu Leu Asp	Glu Ser Leu Gln Val
305	310	315
Ala Glu Arg Leu Thr Arg	Lys Tyr Asn Glu Leu	Leu Lys Ser Tyr Gln
325	330	335
Trp Lys Met Leu Asn Thr	Ser Ser Leu Leu Glu	Gln Leu Asn Glu Gln
340	345	350
Phe Asn Trp Val Ser Arg	Leu Ala Asn Leu Thr	Gln Gly Glu Asp Gln
355	360	365
Tyr Tyr Leu Arg Val Thr	Thr Val Ala Ser His	Thr Ser Asp Ser Asp
370	375	380
Val Pro Ser Gly Val Thr	Glu Val Val Val Lys	Leu Phe Asp Ser Asp
385	390	395
Pro Ile Thr Val Thr Val	Pro Val Glu Val Ser	Arg Lys Asn Pro Lys
405	410	415
Phe Met Glu Thr Val Ala	Glu Lys Ala Leu Gln	Glu Tyr Arg Lys Lys
420	425	430
His Arg Glu Glu		
435		

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<210> SEQ ID NO 3
<211> LENGTH: 428
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 3

Thr Trp Glu Ser Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn
1      5      10      15

Glu Leu Gln Glu Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu
20     25     30

Ile Gln Asn Ala Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu
35     40     45

Lys Thr Asn Glu Glu Arg Lys Thr Leu Leu Ser Asn Glu Asp Ala Leu
50     55     60

Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys Glu Leu Pro Gly Val
65     70     75     80

Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu Cys Lys Pro Cys Leu
85     90     95

Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val Cys Arg Ser Gly Ser
100    105    110

Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu Asn Gln Ser Ser Pro
115    120    125

Phe Tyr Phe Trp Met Asn Gly Asp Arg Ile Asp Ser Leu Leu Glu Asn
130    135    140

Asp Arg Gln Gln Thr His Met Leu Asp Val Met Gln Asp His Phe Ser
145    150    155    160

Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln Asp Arg Phe Phe Thr
165    170    175

Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro Phe Ser Leu Pro His
180    185    190

Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg Ile Val Arg Ser Leu
195    200    205

Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe His Ala Met Phe Gln
210    215    220

Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln Ala Met Asp Ile His
225    230    235    240

Phe His Ser Pro Ala Phe Gln His Pro Pro Thr Glu Phe Ile Arg Glu
245    250    255

Gly Asp Asp Asp Arg Thr Val Cys Arg Glu Ile Arg His Asn Ser Thr
260    265    270

Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys Cys Arg Glu Ile Leu
275    280    285

Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln Ala Lys Leu Arg Arg
290    295    300

Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg Leu Thr Arg Lys Tyr
305    310    315    320

Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met Leu Asn Thr Ser Ser
325    330    335

Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp Val Ser Arg Leu Ala
340    345    350

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Asn Leu Thr Gln Gly Glu Asp Gln Tyr Tyr Leu Arg Val Thr Thr Val
   355                               360                               365

Ala Ser His Thr Ser Asp Ser Asp Val Pro Ser Gly Val Thr Glu Val
   370                               375                               380

Val Val Lys Leu Phe Asp Ser Asp Pro Ile Thr Val Thr Val Pro Val
   385                               390                               395                               400

Glu Val Ser Arg Lys Asn Pro Lys Phe Met Glu Thr Val Ala Glu Lys
   405                               410                               415

Ala Leu Gln Glu Tyr Arg Lys Lys His Arg Glu Glu
   420                               425

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<210> SEQ ID NO 4
<211> LENGTH: 461
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide

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<400> SEQUENCE: 4

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His His His His His His Leu Val Pro Arg Gly Ser Met Met Lys Thr
 1      5      10      15

Leu Leu Leu Phe Val Gly Leu Leu Leu Thr Trp Glu Ser Gly Gln Val
 20      25      30

Leu Gly Asp Gln Thr Val Ser Asp Asn Glu Leu Gln Glu Met Ser Asn
 35      40      45

Gln Gly Ser Lys Tyr Val Asn Lys Glu Ile Gln Asn Ala Val Asn Gly
 50      55      60

Val Lys Gln Ile Lys Thr Leu Ile Glu Lys Thr Asn Glu Glu Arg Lys
 65      70      75      80

Thr Leu Leu Ser Asn Leu Glu Glu Ala Lys Lys Lys Lys Glu Asp Ala
 85      90      95

Leu Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys Glu Leu Pro Gly
100     105     110

Val Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu Cys Lys Pro Cys
115     120     125

Leu Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val Cys Arg Ser Gly
130     135     140

Ser Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu Asn Gln Ser Ser
145     150     155     160

Pro Phe Tyr Phe Trp Met Asn Gly Asp Arg Ile Asp Ser Leu Leu Glu
165     170     175

Asn Asp Arg Gln Gln Thr His Met Leu Asp Val Met Gln Asp His Phe
180     185     190

Ser Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln Asp Arg Phe Phe
195     200     205

Thr Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro Phe Ser Leu Pro
210     215     220

His Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg Ile Val Arg Ser
225     230     235     240

Leu Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe His Ala Met Phe
245     250     255

Gln Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln Ala Met Asp Ile
260     265     270

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His Phe His Ser Pro Ala Phe Gln His Pro Pro Thr Glu Phe Ile Arg
 275 280 285
 Glu Gly Asp Asp Asp Arg Thr Val Cys Arg Glu Ile Arg His Asn Ser
 290 295 300
 Thr Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys Cys Arg Glu Ile
 305 310 315 320
 Leu Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln Ala Lys Leu Arg
 325 330 335
 Arg Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg Leu Thr Arg Lys
 340 345 350
 Tyr Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met Leu Asn Thr Ser
 355 360 365
 Ser Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp Val Ser Arg Leu
 370 375 380
 Ala Asn Leu Thr Gln Gly Glu Asp Gln Tyr Tyr Leu Arg Val Thr Thr
 385 390 395 400
 Val Ala Ser His Thr Ser Asp Ser Asp Val Pro Ser Gly Val Thr Glu
 405 410 415
 Val Val Val Lys Leu Phe Asp Ser Asp Pro Ile Thr Val Thr Val Pro
 420 425 430
 Val Glu Val Ser Arg Lys Asn Pro Lys Phe Met Glu Thr Val Ala Glu
 435 440 445
 Lys Ala Leu Gln Glu Tyr Arg Lys Lys His Arg Glu Glu
 450 455 460

<210> SEQ ID NO 5
 <211> LENGTH: 448
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 5

His His His His His His Leu Val Pro Arg Gly Ser Thr Trp Glu Ser
 1 5 10 15
 Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn Glu Leu Gln Glu
 20 25 30
 Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu Ile Gln Asn Ala
 35 40 45
 Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu Lys Thr Asn Glu
 50 55 60
 Glu Arg Lys Thr Leu Leu Ser Asn Leu Glu Glu Ala Lys Lys Lys Lys
 65 70 75 80
 Glu Asp Ala Leu Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys Glu
 85 90 95
 Leu Pro Gly Val Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu Cys
 100 105 110
 Lys Pro Cys Leu Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val Cys
 115 120 125
 Arg Ser Gly Ser Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu Asn
 130 135 140
 Gln Ser Ser Pro Phe Tyr Phe Trp Met Asn Gly Asp Arg Ile Asp Ser
 145 150 155 160

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Leu Leu Glu Asn Asp Arg Gln Gln Thr His Met Leu Asp Val Met Gln
 165 170 175
 Asp His Phe Ser Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln Asp
 180 185 190
 Arg Phe Phe Thr Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro Phe
 195 200 205
 Ser Leu Pro His Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg Ile
 210 215 220
 Val Arg Ser Leu Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe His
 225 230 235 240
 Ala Met Phe Gln Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln Ala
 245 250 255
 Met Asp Ile His Phe His Ser Pro Ala Phe Gln His Pro Pro Thr Glu
 260 265 270
 Phe Ile Arg Glu Gly Asp Asp Arg Thr Val Cys Arg Glu Ile Arg
 275 280 285
 His Asn Ser Thr Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys Cys
 290 295 300
 Arg Glu Ile Leu Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln Ala
 305 310 315 320
 Lys Leu Arg Arg Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg Leu
 325 330 335
 Thr Arg Lys Tyr Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met Leu
 340 345 350
 Asn Thr Ser Ser Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp Val
 355 360 365
 Ser Arg Leu Ala Asn Leu Thr Gln Gly Glu Asp Gln Tyr Tyr Leu Arg
 370 375 380
 Val Thr Thr Val Ala Ser His Thr Ser Asp Ser Asp Val Pro Ser Gly
 385 390 395 400
 Val Thr Glu Val Val Val Lys Leu Phe Asp Ser Asp Pro Ile Thr Val
 405 410 415
 Thr Val Pro Val Glu Val Ser Arg Lys Asn Pro Lys Phe Met Glu Thr
 420 425 430
 Val Ala Glu Lys Ala Leu Gln Glu Tyr Arg Lys Lys His Arg Glu Glu
 435 440 445

<210> SEQ ID NO 6
 <211> LENGTH: 440
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 6

His His His His His His Leu Val Pro Arg Gly Ser Thr Trp Glu Ser
 1 5 10 15
 Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn Glu Leu Gln Glu
 20 25 30
 Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu Ile Gln Asn Ala
 35 40 45
 Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu Lys Thr Asn Glu
 50 55 60

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Glu	Arg	Lys	Thr	Leu	Leu	Ser	Asn	Glu	Asp	Ala	Leu	Asn	Glu	Thr	Arg	65	70	75	80
Glu	Ser	Glu	Thr	Lys	Leu	Lys	Glu	Leu	Pro	Gly	Val	Cys	Asn	Glu	Thr	85	90	95	
Met	Met	Ala	Leu	Trp	Glu	Glu	Cys	Lys	Pro	Cys	Leu	Lys	Gln	Thr	Cys	100	105	110	
Met	Lys	Phe	Tyr	Ala	Arg	Val	Cys	Arg	Ser	Gly	Ser	Gly	Leu	Val	Gly	115	120	125	
Arg	Gln	Leu	Glu	Glu	Phe	Leu	Asn	Gln	Ser	Ser	Pro	Phe	Tyr	Phe	Trp	130	135	140	
Met	Asn	Gly	Asp	Arg	Ile	Asp	Ser	Leu	Leu	Glu	Asn	Asp	Arg	Gln	Gln	145	150	155	160
Thr	His	Met	Leu	Asp	Val	Met	Gln	Asp	His	Phe	Ser	Arg	Ala	Ser	Ser	165	170	175	
Ile	Ile	Asp	Glu	Leu	Phe	Gln	Asp	Arg	Phe	Phe	Thr	Arg	Glu	Pro	Gln	180	185	190	
Asp	Thr	Tyr	His	Tyr	Leu	Pro	Phe	Ser	Leu	Pro	His	Arg	Arg	Pro	His	195	200	205	
Phe	Phe	Phe	Pro	Lys	Ser	Arg	Ile	Val	Arg	Ser	Leu	Met	Pro	Phe	Ser	210	215	220	
Pro	Tyr	Glu	Pro	Leu	Asn	Phe	His	Ala	Met	Phe	Gln	Pro	Phe	Leu	Glu	225	230	235	240
Met	Ile	His	Glu	Ala	Gln	Gln	Ala	Met	Asp	Ile	His	Phe	His	Ser	Pro	245	250	255	
Ala	Phe	Gln	His	Pro	Pro	Thr	Glu	Phe	Ile	Arg	Glu	Gly	Asp	Asp	Asp	260	265	270	
Arg	Thr	Val	Cys	Arg	Glu	Ile	Arg	His	Asn	Ser	Thr	Gly	Cys	Leu	Arg	275	280	285	
Met	Lys	Asp	Gln	Cys	Asp	Lys	Cys	Arg	Glu	Ile	Leu	Ser	Val	Asp	Cys	290	295	300	
Ser	Thr	Asn	Asn	Pro	Ser	Gln	Ala	Lys	Leu	Arg	Arg	Glu	Leu	Asp	Glu	305	310	315	320
Ser	Leu	Gln	Val	Ala	Glu	Arg	Leu	Thr	Arg	Lys	Tyr	Asn	Glu	Leu	Leu	325	330	335	
Lys	Ser	Tyr	Gln	Trp	Lys	Met	Leu	Asn	Thr	Ser	Ser	Leu	Leu	Glu	Gln	340	345	350	
Leu	Asn	Glu	Gln	Phe	Asn	Trp	Val	Ser	Arg	Leu	Ala	Asn	Leu	Thr	Gln	355	360	365	
Gly	Glu	Asp	Gln	Tyr	Tyr	Leu	Arg	Val	Thr	Thr	Val	Ala	Ser	His	Thr	370	375	380	
Ser	Asp	Ser	Asp	Val	Pro	Ser	Gly	Val	Thr	Glu	Val	Val	Val	Lys	Leu	385	390	395	400
Phe	Asp	Ser	Asp	Pro	Ile	Thr	Val	Thr	Val	Pro	Val	Glu	Val	Ser	Arg	405	410	415	
Lys	Asn	Pro	Lys	Phe	Met	Glu	Thr	Val	Ala	Glu	Lys	Ala	Leu	Gln	Glu	420	425	430	
Tyr	Arg	Lys	Lys	His	Arg	Glu	Glu	435	440										

<210> SEQ ID NO 7

<211> LENGTH: 461

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<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 7

Met Met Lys Thr Leu Leu Leu Phe Val Gly Leu Leu Leu Thr Trp Glu
1 5 10 15

Ser Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn Glu Leu Gln
20 25 30

Glu Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu Ile Gln Asn
35 40 45

Ala Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu Lys Thr Asn
50 55 60

Glu Glu Arg Lys Thr Leu Leu Ser Asn Leu Glu Glu Ala Lys Lys Lys
65 70 75 80

Lys Glu Asp Ala Leu Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys
85 90 95

Glu Leu Pro Gly Val Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu
100 105 110

Cys Lys Pro Cys Leu Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val
115 120 125

Cys Arg Ser Gly Ser Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu
130 135 140

Asn Gln Ser Ser Pro Phe Tyr Phe Trp Met Asn Gly Asp Arg Ile Asp
145 150 155 160

Ser Leu Leu Glu Asn Asp Arg Gln Gln Thr His Met Leu Asp Val Met
165 170 175

Gln Asp His Phe Ser Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln
180 185 190

Asp Arg Phe Phe Thr Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro
195 200 205

Phe Ser Leu Pro His Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg
210 215 220

Ile Val Arg Ser Leu Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe
225 230 235 240

His Ala Met Phe Gln Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln
245 250 255

Ala Met Asp Ile His Phe His Ser Pro Ala Phe Gln His Pro Pro Thr
260 265 270

Glu Phe Ile Arg Glu Gly Asp Asp Asp Arg Thr Val Cys Arg Glu Ile
275 280 285

Arg His Asn Ser Thr Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys
290 295 300

Cys Arg Glu Ile Leu Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln
305 310 315 320

Ala Lys Leu Arg Arg Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg
325 330 335

Leu Thr Arg Lys Tyr Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met
340 345 350

Leu Asn Thr Ser Ser Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp
355 360 365

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Val	Ser	Arg	Leu	Ala	Asn	Leu	Thr	Gln	Gly	Glu	Asp	Gln	Tyr	Tyr	Leu
370						375					380				
Arg	Val	Thr	Thr	Val	Ala	Ser	His	Thr	Ser	Asp	Ser	Asp	Val	Pro	Ser
385					390					395					400
Gly	Val	Thr	Glu	Val	Val	Val	Lys	Leu	Phe	Asp	Ser	Asp	Pro	Ile	Thr
			405						410					415	
Val	Thr	Val	Pro	Val	Glu	Val	Ser	Arg	Lys	Asn	Pro	Lys	Phe	Met	Glu
			420					425					430		
Thr	Val	Ala	Glu	Lys	Ala	Leu	Gln	Glu	Tyr	Arg	Lys	Lys	His	Arg	Glu
		435					440					445			
Glu	Leu	Val	Pro	Arg	Gly	Ser	His	His	His	His	His	His			
450						455						460			

<210> SEQ ID NO 8
 <211> LENGTH: 448
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 8

Thr	Trp	Glu	Ser	Gly	Gln	Val	Leu	Gly	Asp	Gln	Thr	Val	Ser	Asp	Asn
1				5					10					15	
Glu	Leu	Gln	Glu	Met	Ser	Asn	Gln	Gly	Ser	Lys	Tyr	Val	Asn	Lys	Glu
			20					25					30		
Ile	Gln	Asn	Ala	Val	Asn	Gly	Val	Lys	Gln	Ile	Lys	Thr	Leu	Ile	Glu
		35				40						45			
Lys	Thr	Asn	Glu	Glu	Arg	Lys	Thr	Leu	Leu	Ser	Asn	Leu	Glu	Glu	Ala
		50				55					60				
Lys	Lys	Lys	Lys	Glu	Asp	Ala	Leu	Asn	Glu	Thr	Arg	Glu	Ser	Glu	Thr
65					70					75				80	
Lys	Leu	Lys	Glu	Leu	Pro	Gly	Val	Cys	Asn	Glu	Thr	Met	Met	Ala	Leu
			85					90						95	
Trp	Glu	Glu	Cys	Lys	Pro	Cys	Leu	Lys	Gln	Thr	Cys	Met	Lys	Phe	Tyr
			100					105					110		
Ala	Arg	Val	Cys	Arg	Ser	Gly	Ser	Gly	Leu	Val	Gly	Arg	Gln	Leu	Glu
		115				120					125				
Glu	Phe	Leu	Asn	Gln	Ser	Ser	Pro	Phe	Tyr	Phe	Trp	Met	Asn	Gly	Asp
		130				135					140				
Arg	Ile	Asp	Ser	Leu	Leu	Glu	Asn	Asp	Arg	Gln	Gln	Thr	His	Met	Leu
145				150						155				160	
Asp	Val	Met	Gln	Asp	His	Phe	Ser	Arg	Ala	Ser	Ser	Ile	Ile	Asp	Glu
			165					170						175	
Leu	Phe	Gln	Asp	Arg	Phe	Phe	Thr	Arg	Glu	Pro	Gln	Asp	Thr	Tyr	His
		180					185						190		
Tyr	Leu	Pro	Phe	Ser	Leu	Pro	His	Arg	Arg	Pro	His	Phe	Phe	Phe	Pro
		195				200						205			
Lys	Ser	Arg	Ile	Val	Arg	Ser	Leu	Met	Pro	Phe	Ser	Pro	Tyr	Glu	Pro
		210				215					220				
Leu	Asn	Phe	His	Ala	Met	Phe	Gln	Pro	Phe	Leu	Glu	Met	Ile	His	Glu
225				230						235				240	
Ala	Gln	Gln	Ala	Met	Asp	Ile	His	Phe	His	Ser	Pro	Ala	Phe	Gln	His
			245					250						255	

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Pro	Pro	Thr	Glu	Phe	Ile	Arg	Glu	Gly	Asp	Asp	Asp	Arg	Thr	Val	Cys
			260					265					270		
Arg	Glu	Ile	Arg	His	Asn	Ser	Thr	Gly	Cys	Leu	Arg	Met	Lys	Asp	Gln
		275					280					285			
Cys	Asp	Lys	Cys	Arg	Glu	Ile	Leu	Ser	Val	Asp	Cys	Ser	Thr	Asn	Asn
	290					295					300				
Pro	Ser	Gln	Ala	Lys	Leu	Arg	Arg	Glu	Leu	Asp	Glu	Ser	Leu	Gln	Val
305					310					315					320
Ala	Glu	Arg	Leu	Thr	Arg	Lys	Tyr	Asn	Glu	Leu	Leu	Lys	Ser	Tyr	Gln
			325						330					335	
Trp	Lys	Met	Leu	Asn	Thr	Ser	Ser	Leu	Leu	Glu	Gln	Leu	Asn	Glu	Gln
			340					345					350		
Phe	Asn	Trp	Val	Ser	Arg	Leu	Ala	Asn	Leu	Thr	Gln	Gly	Glu	Asp	Gln
		355					360					365			
Tyr	Tyr	Leu	Arg	Val	Thr	Thr	Val	Ala	Ser	His	Thr	Ser	Asp	Ser	Asp
	370					375					380				
Val	Pro	Ser	Gly	Val	Thr	Glu	Val	Val	Val	Lys	Leu	Phe	Asp	Ser	Asp
385					390					395					400
Pro	Ile	Thr	Val	Thr	Val	Pro	Val	Glu	Val	Ser	Arg	Lys	Asn	Pro	Lys
			405					410						415	
Phe	Met	Glu	Thr	Val	Ala	Glu	Lys	Ala	Leu	Gln	Glu	Tyr	Arg	Lys	Lys
			420					425					430		
His	Arg	Glu	Glu	Leu	Val	Pro	Arg	Gly	Ser	His	His	His	His	His	His
		435					440					445			

<210> SEQ ID NO 9
 <211> LENGTH: 444
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 9

Met	Met	Lys	Thr	Thr	Trp	Glu	Ser	Gly	Gln	Val	Leu	Gly	Asp	Gln	Thr
1				5					10					15	
Val	Ser	Asp	Asn	Glu	Leu	Gln	Glu	Met	Ser	Asn	Gln	Gly	Ser	Lys	Tyr
			20					25					30		
Val	Asn	Lys	Glu	Ile	Gln	Asn	Ala	Val	Asn	Gly	Val	Lys	Gln	Ile	Lys
		35					40					45			
Thr	Leu	Ile	Glu	Lys	Thr	Asn	Glu	Glu	Arg	Lys	Thr	Leu	Leu	Ser	Asn
	50					55					60				
Glu	Asp	Ala	Leu	Asn	Glu	Thr	Arg	Glu	Ser	Glu	Thr	Lys	Leu	Lys	Glu
65				70					75						80
Leu	Pro	Gly	Val	Cys	Asn	Glu	Thr	Met	Met	Ala	Leu	Trp	Glu	Glu	Cys
			85					90					95		
Lys	Pro	Cys	Leu	Lys	Gln	Thr	Cys	Met	Lys	Phe	Tyr	Ala	Arg	Val	Cys
		100						105					110		
Arg	Ser	Gly	Ser	Gly	Leu	Val	Gly	Arg	Gln	Leu	Glu	Glu	Phe	Leu	Asn
		115					120					125			
Gln	Ser	Ser	Pro	Phe	Tyr	Phe	Trp	Met	Asn	Gly	Asp	Arg	Ile	Asp	Ser
	130					135					140				
Leu	Leu	Glu	Asn	Asp	Arg	Gln	Gln	Thr	His	Met	Leu	Asp	Val	Met	Gln
145					150					155					160

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Asp	His	Phe	Ser	Arg 165	Ala	Ser	Ser	Ile	Ile 170	Asp	Glu	Leu	Phe	Gln 175	Asp
Arg	Phe	Phe	Thr 180	Arg	Glu	Pro	Gln	Asp 185	Thr	Tyr	His	Tyr	Leu 190	Pro	Phe
Ser	Leu	Pro	His 195	Arg	Arg	Pro	His 200	Phe	Phe	Phe	Pro	Lys 205	Ser	Arg	Ile
Val	Arg 210	Ser	Leu	Met	Pro	Phe 215	Ser	Pro	Tyr	Glu	Pro 220	Leu	Asn	Phe	His
Ala 225	Met	Phe	Gln	Pro	Phe 230	Leu	Glu	Met	Ile	His 235	Glu	Ala	Gln	Gln	Ala 240
Met	Asp	Ile	His 245	Phe	His	Ser	Pro	Ala	Phe 250	Gln	His	Pro	Pro	Thr 255	Glu
Phe	Ile	Arg	Glu 260	Gly	Asp	Asp	Asp	Arg 265	Thr	Val	Cys	Arg	Glu 270	Ile	Arg
His	Asn	Ser 275	Thr	Gly	Cys	Leu	Arg 280	Met	Lys	Asp	Gln	Cys 285	Asp	Lys	Cys
Arg	Glu 290	Ile	Leu	Ser	Val	Asp 295	Cys	Ser	Thr	Asn	Asn 300	Pro	Ser	Gln	Ala
Lys 305	Leu	Arg	Arg	Glu	Leu 310	Asp	Glu	Ser	Leu	Gln 315	Val	Ala	Glu	Arg	Leu 320
Thr	Arg	Lys	Tyr 325	Asn	Glu	Leu	Leu	Lys	Ser 330	Tyr	Gln	Trp	Lys	Met 335	Leu
Asn	Thr	Ser 340	Ser	Leu	Leu	Glu	Gln	Leu 345	Asn	Glu	Gln	Phe	Asn 350	Trp	Val
Ser	Arg	Leu 355	Ala	Asn	Leu	Thr	Gln 360	Gly	Glu	Asp	Gln	Tyr 365	Tyr	Leu	Arg
Val	Thr 370	Thr	Val	Ala	Ser	His 375	Thr	Ser	Asp	Ser	Asp 380	Val	Pro	Ser	Gly
Val 385	Thr	Glu	Val	Val	Val 390	Lys	Leu	Phe	Asp	Ser 395	Asp	Pro	Ile	Thr	Val 400
Thr	Val	Pro	Val 405	Glu	Val	Ser	Arg	Lys	Asn 410	Pro	Lys	Phe	Met	Glu 415	Thr
Val	Ala	Glu	Lys 420	Ala	Leu	Gln	Glu	Tyr 425	Arg	Lys	Lys	His	Arg 430	Glu	Glu
Leu	Val	Pro 435	Arg	Gly	Ser	His	His 440	His	His	His	His				

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<210> SEQ ID NO 10
<211> LENGTH: 460
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide
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<400> SEQUENCE: 10

His	His	His	His	His	His	Asp	Asp	Asp	Asp	Lys	Met	Met	Lys	Thr	Leu
1				5					10					15	
Leu	Leu	Phe	Val	Gly	Leu	Leu	Leu	Thr	Trp	Glu	Ser	Gly	Gln	Val	Leu
			20					25					30		
Gly	Asp	Gln	Thr	Val	Ser	Asp	Asn	Glu	Leu	Gln	Glu	Met	Ser	Asn	Gln
		35					40					45			
Gly	Ser	Lys	Tyr	Val	Asn	Lys	Glu	Ile	Gln	Asn	Ala	Val	Asn	Gly	Val
	50					55					60				

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Lys	Gln	Ile	Lys	Thr	Leu	Ile	Glu	Lys	Thr	Asn	Glu	Glu	Arg	Lys	Thr	
65					70					75					80	
Leu	Leu	Ser	Asn	Leu	Glu	Glu	Ala	Lys	Lys	Lys	Lys	Glu	Asp	Ala	Leu	
			85					90						95		
Asn	Glu	Thr	Arg	Glu	Ser	Glu	Thr	Lys	Leu	Lys	Glu	Leu	Pro	Gly	Val	
			100					105					110			
Cys	Asn	Glu	Thr	Met	Met	Ala	Leu	Trp	Glu	Glu	Cys	Lys	Pro	Cys	Leu	
		115					120					125				
Lys	Gln	Thr	Cys	Met	Lys	Phe	Tyr	Ala	Arg	Val	Cys	Arg	Ser	Gly	Ser	
	130					135					140					
Gly	Leu	Val	Gly	Arg	Gln	Leu	Glu	Glu	Phe	Leu	Asn	Gln	Ser	Ser	Pro	
145					150					155					160	
Phe	Tyr	Phe	Trp	Met	Asn	Gly	Asp	Arg	Ile	Asp	Ser	Leu	Leu	Glu	Asn	
				165					170					175		
Asp	Arg	Gln	Gln	Thr	His	Met	Leu	Asp	Val	Met	Gln	Asp	His	Phe	Ser	
			180					185					190			
Arg	Ala	Ser	Ser	Ile	Ile	Asp	Glu	Leu	Phe	Gln	Asp	Arg	Phe	Phe	Thr	
		195					200					205				
Arg	Glu	Pro	Gln	Asp	Thr	Tyr	His	Tyr	Leu	Pro	Phe	Ser	Leu	Pro	His	
	210					215					220					
Arg	Arg	Pro	His	Phe	Phe	Phe	Pro	Lys	Ser	Arg	Ile	Val	Arg	Ser	Leu	
225					230					235					240	
Met	Pro	Phe	Ser	Pro	Tyr	Glu	Pro	Leu	Asn	Phe	His	Ala	Met	Phe	Gln	
				245					250					255		
Pro	Phe	Leu	Glu	Met	Ile	His	Glu	Ala	Gln	Gln	Ala	Met	Asp	Ile	His	
			260					265					270			
Phe	His	Ser	Pro	Ala	Phe	Gln	His	Pro	Pro	Thr	Glu	Phe	Ile	Arg	Glu	
		275					280					285				
Gly	Asp	Asp	Asp	Arg	Thr	Val	Cys	Arg	Glu	Ile	Arg	His	Asn	Ser	Thr	
	290					295					300					
Gly	Cys	Leu	Arg	Met	Lys	Asp	Gln	Cys	Asp	Lys	Cys	Arg	Glu	Ile	Leu	
305					310					315					320	
Ser	Val	Asp	Cys	Ser	Thr	Asn	Asn	Pro	Ser	Gln	Ala	Lys	Leu	Arg	Arg	
				325					330					335		
Glu	Leu	Asp	Glu	Ser	Leu	Gln	Val	Ala	Glu	Arg	Leu	Thr	Arg	Lys	Tyr	
			340					345					350			
Asn	Glu	Leu	Leu	Lys	Ser	Tyr	Gln	Trp	Lys	Met	Leu	Asn	Thr	Ser	Ser	
		355					360					365				
Leu	Leu	Glu	Gln	Leu	Asn	Glu	Gln	Phe	Asn	Trp	Val	Ser	Arg	Leu	Ala	
		370				375					380					
Asn	Leu	Thr	Gln	Gly	Glu	Asp	Gln	Tyr	Tyr	Leu	Arg	Val	Thr	Thr	Val	
385					390					395					400	
Ala	Ser	His	Thr	Ser	Asp	Ser	Asp	Val	Pro	Ser	Gly	Val	Thr	Glu	Val	
				405				410						415		
Val	Val	Lys	Leu	Phe	Asp	Ser	Asp	Pro	Ile	Thr	Val	Thr	Val	Pro	Val	
			420					425					430			
Glu	Val	Ser	Arg	Lys	Asn	Pro	Lys	Phe	Met	Glu	Thr	Val	Ala	Glu	Lys	
			435				440					445				
Ala	Leu	Gln	Glu	Tyr	Arg	Lys	Lys	His	Arg	Glu	Glu					
	450					455					460					

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<210> SEQ ID NO 11
<211> LENGTH: 447
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 11
His His His His His His Asp Asp Asp Asp Lys Thr Trp Glu Ser Gly
1          5          10          15
Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn Glu Leu Gln Glu Met
20          25          30
Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu Ile Gln Asn Ala Val
35          40          45
Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu Lys Thr Asn Glu Glu
50          55          60
Arg Lys Thr Leu Leu Ser Asn Leu Glu Glu Ala Lys Lys Lys Lys Glu
65          70          75          80
Asp Ala Leu Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys Glu Leu
85          90          95
Pro Gly Val Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu Cys Lys
100         105         110
Pro Cys Leu Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val Cys Arg
115         120         125
Ser Gly Ser Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu Asn Gln
130         135         140
Ser Ser Pro Phe Tyr Phe Trp Met Asn Gly Asp Arg Ile Asp Ser Leu
145         150         155         160
Leu Glu Asn Asp Arg Gln Gln Thr His Met Leu Asp Val Met Gln Asp
165         170         175
His Phe Ser Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln Asp Arg
180         185         190
Phe Phe Thr Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro Phe Ser
195         200         205
Leu Pro His Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg Ile Val
210         215         220
Arg Ser Leu Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe His Ala
225         230         235         240
Met Phe Gln Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln Ala Met
245         250         255
Asp Ile His Phe His Ser Pro Ala Phe Gln His Pro Pro Thr Glu Phe
260         265         270
Ile Arg Glu Gly Asp Asp Asp Arg Thr Val Cys Arg Glu Ile Arg His
275         280         285
Asn Ser Thr Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys Cys Arg
290         295         300
Glu Ile Leu Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln Ala Lys
305         310         315         320
Leu Arg Arg Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg Leu Thr
325         330         335
Arg Lys Tyr Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met Leu Asn
340         345         350
Thr Ser Ser Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp Val Ser

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355					360					365					
Arg	Leu	Ala	Asn	Leu	Thr	Gln	Gly	Glu	Asp	Gln	Tyr	Tyr	Leu	Arg	Val
370						375					380				
Thr	Thr	Val	Ala	Ser	His	Thr	Ser	Asp	Ser	Asp	Val	Pro	Ser	Gly	Val
385					390					395					400
Thr	Glu	Val	Val	Val	Lys	Leu	Phe	Asp	Ser	Asp	Pro	Ile	Thr	Val	Thr
				405					410					415	
Val	Pro	Val	Glu	Val	Ser	Arg	Lys	Asn	Pro	Lys	Phe	Met	Glu	Thr	Val
			420					425					430		
Ala	Glu	Lys	Ala	Leu	Gln	Glu	Tyr	Arg	Lys	Lys	His	Arg	Glu	Glu	
		435					440					445			
<210> SEQ ID NO 12															
<211> LENGTH: 439															
<212> TYPE: PRT															
<213> ORGANISM: Artificial Sequence															
<220> FEATURE:															
<223> OTHER INFORMATION: Synthetic peptide															
<400> SEQUENCE: 12															
His	His	His	His	His	His	Asp	Asp	Asp	Asp	Lys	Thr	Trp	Glu	Ser	Gly
1				5					10					15	
Gln	Val	Leu	Gly	Asp	Gln	Thr	Val	Ser	Asp	Asn	Glu	Leu	Gln	Glu	Met
		20						25					30		
Ser	Asn	Gln	Gly	Ser	Lys	Tyr	Val	Asn	Lys	Glu	Ile	Gln	Asn	Ala	Val
		35				40						45			
Asn	Gly	Val	Lys	Gln	Ile	Lys	Thr	Leu	Ile	Glu	Lys	Thr	Asn	Glu	Glu
	50					55				60					
Arg	Lys	Thr	Leu	Leu	Ser	Asn	Glu	Asp	Ala	Leu	Asn	Glu	Thr	Arg	Glu
65					70					75					80
Ser	Glu	Thr	Lys	Leu	Lys	Glu	Leu	Pro	Gly	Val	Cys	Asn	Glu	Thr	Met
			85					90						95	
Met	Ala	Leu	Trp	Glu	Glu	Cys	Lys	Pro	Cys	Leu	Lys	Gln	Thr	Cys	Met
			100					105					110		
Lys	Phe	Tyr	Ala	Arg	Val	Cys	Arg	Ser	Gly	Ser	Gly	Leu	Val	Gly	Arg
		115					120					125			
Gln	Leu	Glu	Glu	Phe	Leu	Asn	Gln	Ser	Ser	Pro	Phe	Tyr	Phe	Trp	Met
	130					135					140				
Asn	Gly	Asp	Arg	Ile	Asp	Ser	Leu	Leu	Glu	Asn	Asp	Arg	Gln	Gln	Thr
145					150					155					160
His	Met	Leu	Asp	Val	Met	Gln	Asp	His	Phe	Ser	Arg	Ala	Ser	Ser	Ile
			165						170					175	
Ile	Asp	Glu	Leu	Phe	Gln	Asp	Arg	Phe	Phe	Thr	Arg	Glu	Pro	Gln	Asp
		180						185					190		
Thr	Tyr	His	Tyr	Leu	Pro	Phe	Ser	Leu	Pro	His	Arg	Arg	Pro	His	Phe
		195					200					205			
Phe	Phe	Pro	Lys	Ser	Arg	Ile	Val	Arg	Ser	Leu	Met	Pro	Phe	Ser	Pro
	210					215					220				
Tyr	Glu	Pro	Leu	Asn	Phe	His	Ala	Met	Phe	Gln	Pro	Phe	Leu	Glu	Met
225					230					235				240	
Ile	His	Glu	Ala	Gln	Gln	Ala	Met	Asp	Ile	His	Phe	His	Ser	Pro	Ala
			245						250					255	
Phe	Gln	His	Pro	Pro	Thr	Glu	Phe	Ile	Arg	Glu	Gly	Asp	Asp	Asp	Arg

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260					265					270					
Thr	Val	Cys	Arg	Glu	Ile	Arg	His	Asn	Ser	Thr	Gly	Cys	Leu	Arg	Met
	275						280					285			
Lys	Asp	Gln	Cys	Asp	Lys	Cys	Arg	Glu	Ile	Leu	Ser	Val	Asp	Cys	Ser
	290					295					300				
Thr	Asn	Asn	Pro	Ser	Gln	Ala	Lys	Leu	Arg	Arg	Glu	Leu	Asp	Glu	Ser
	305					310					315				320
Leu	Gln	Val	Ala	Glu	Arg	Leu	Thr	Arg	Lys	Tyr	Asn	Glu	Leu	Leu	Lys
			325						330					335	
Ser	Tyr	Gln	Trp	Lys	Met	Leu	Asn	Thr	Ser	Ser	Leu	Leu	Glu	Gln	Leu
			340					345					350		
Asn	Glu	Gln	Phe	Asn	Trp	Val	Ser	Arg	Leu	Ala	Asn	Leu	Thr	Gln	Gly
		355					360					365			
Glu	Asp	Gln	Tyr	Tyr	Leu	Arg	Val	Thr	Thr	Val	Ala	Ser	His	Thr	Ser
	370					375					380				
Asp	Ser	Asp	Val	Pro	Ser	Gly	Val	Thr	Glu	Val	Val	Val	Lys	Leu	Phe
	385					390					395				400
Asp	Ser	Asp	Pro	Ile	Thr	Val	Thr	Val	Pro	Val	Glu	Val	Ser	Arg	Lys
			405						410					415	
Asn	Pro	Lys	Phe	Met	Glu	Thr	Val	Ala	Glu	Lys	Ala	Leu	Gln	Glu	Tyr
			420					425					430		
Arg	Lys	Lys	His	Arg	Glu	Glu									
			435												

<210> SEQ ID NO 13
 <211> LENGTH: 461
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 13

Met	Met	Lys	Thr	Leu	Leu	Leu	Phe	Val	Gly	Leu	Leu	Leu	Thr	Trp	Glu
1				5					10					15	
Ser	Gly	Gln	Val	Leu	Gly	Asp	Gln	Thr	Val	Ser	Asp	Asn	Glu	Leu	Gln
		20						25					30		
Glu	Met	Ser	Asn	Gln	Gly	Ser	Lys	Tyr	Val	Asn	Lys	Glu	Ile	Gln	Asn
		35					40					45			
Ala	Val	Asn	Gly	Val	Lys	Gln	Ile	Lys	Thr	Leu	Ile	Glu	Lys	Thr	Asn
		50				55					60				
Glu	Glu	Arg	Lys	Thr	Leu	Leu	Ser	Asn	Leu	Glu	Glu	Ala	Lys	Lys	Lys
	65				70					75				80	
Lys	Glu	Asp	Ala	Leu	Asn	Glu	Thr	Arg	Glu	Ser	Glu	Thr	Lys	Leu	Lys
			85						90					95	
Glu	Leu	Pro	Gly	Val	Cys	Asn	Glu	Thr	Met	Met	Ala	Leu	Trp	Glu	Glu
			100					105					110		
Cys	Lys	Pro	Cys	Leu	Lys	Gln	Thr	Cys	Met	Lys	Phe	Tyr	Ala	Arg	Val
		115					120					125			
Cys	Arg	Ser	Gly	Ser	Gly	Leu	Val	Gly	Arg	Gln	Leu	Glu	Glu	Phe	Leu
	130					135					140				
Asn	Gln	Ser	Ser	Pro	Phe	Tyr	Phe	Trp	Met	Asn	Gly	Asp	Arg	Ile	Asp
	145				150					155				160	
Ser	Leu	Leu	Glu	Asn	Asp	Arg	Gln	Gln	Thr	His	Met	Leu	Asp	Val	Met

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165          170          175
Gln Asp His Phe Ser Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln
      180          185          190
Asp Arg Phe Phe Thr Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro
      195          200          205
Phe Ser Leu Pro His Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg
      210          215          220
Ile Val Arg Ser Leu Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe
      225          230          235          240
His Ala Met Phe Gln Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln
      245          250          255
Ala Met Asp Ile His Phe His Ser Pro Ala Phe Gln His Pro Pro Thr
      260          265          270
Glu Phe Ile Arg Glu Gly Asp Asp Asp Arg Thr Val Cys Arg Glu Ile
      275          280          285
Arg His Asn Ser Thr Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys
      290          295          300
Cys Arg Glu Ile Leu Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln
      305          310          315          320
Ala Lys Leu Arg Arg Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg
      325          330          335
Leu Thr Arg Lys Tyr Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met
      340          345          350
Leu Asn Thr Ser Ser Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp
      355          360          365
Val Ser Arg Leu Ala Asn Leu Thr Gln Gly Glu Asp Gln Tyr Tyr Leu
      370          375          380
Arg Val Thr Thr Val Ala Ser His Thr Ser Asp Ser Asp Val Pro Ser
      385          390          395          400
Gly Val Thr Glu Val Val Val Lys Leu Phe Asp Ser Asp Pro Ile Thr
      405          410          415
Val Thr Val Pro Val Glu Val Ser Arg Lys Asn Pro Lys Phe Met Glu
      420          425          430
Thr Val Ala Glu Lys Ala Leu Gln Glu Tyr Arg Lys Lys His Arg Glu
      435          440          445
Glu Leu Asp Asp Asp Asp Lys His His His His His His
      450          455          460

<210> SEQ ID NO 14
<211> LENGTH: 448
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 14
Thr Trp Glu Ser Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn
1          5          10          15
Glu Leu Gln Glu Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu
      20          25          30
Ile Gln Asn Ala Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu
      35          40          45
Lys Thr Asn Glu Glu Arg Lys Thr Leu Leu Ser Asn Leu Glu Glu Ala

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50					55					60					
Lys 65	Lys	Lys	Lys	Glu	Asp 70	Ala	Leu	Asn	Glu	Thr 75	Arg	Glu	Ser	Glu	Thr 80
Lys	Leu	Lys	Glu	Leu 85	Pro	Gly	Val	Cys	Asn 90	Glu	Thr	Met	Met	Ala 95	Leu
Trp	Glu	Glu	Cys 100	Lys	Pro	Cys	Leu	Lys 105	Gln	Thr	Cys	Met	Lys 110	Phe	Tyr
Ala	Arg	Val 115	Cys	Arg	Ser	Gly	Ser 120	Gly	Leu	Val	Gly 125	Arg	Gln	Leu	Glu
Glu	Phe 130	Leu	Asn	Gln	Ser 135	Ser	Pro	Phe	Tyr	Phe 140	Trp	Met	Asn	Gly	Asp
Arg 145	Ile	Asp	Ser	Leu 150	Leu	Glu	Asn	Asp	Arg 155	Gln	Gln	Thr	His	Met	Leu 160
Asp	Val	Met	Gln 165	Asp	His	Phe	Ser	Arg 170	Ala	Ser	Ser	Ile	Ile 175	Asp	Glu
Leu	Phe	Gln 180	Asp	Arg	Phe	Phe	Thr	Arg 185	Glu	Pro	Gln	Asp	Thr 190	Tyr	His
Tyr	Leu 195	Pro	Phe	Ser	Leu	Pro	His 200	Arg	Arg	Pro	His 205	Phe	Phe	Phe	Pro
Lys 210	Ser	Arg	Ile	Val	Arg 215	Ser	Leu	Met	Pro	Phe 220	Ser	Pro	Tyr	Glu	Pro
Leu 225	Asn	Phe	His	Ala 230	Met	Phe	Gln	Pro	Phe 235	Leu	Glu	Met	Ile	His	Glu 240
Ala	Gln	Gln 245	Ala	Met	Asp	Ile	His	Phe 250	His	Ser	Pro	Ala	Phe 255	Gln	His
Pro	Pro	Thr 260	Glu	Phe	Ile	Arg	Glu	Gly 265	Asp	Asp	Asp	Arg	Thr 270	Val	Cys
Arg	Glu 275	Ile	Arg	His	Asn	Ser	Thr 280	Gly	Cys	Leu	Arg	Met 285	Lys	Asp	Gln
Cys 290	Asp	Lys	Cys	Arg	Glu 295	Ile	Leu	Ser	Val	Asp 300	Cys	Ser	Thr	Asn	Asn
Pro 305	Ser	Gln	Ala	Lys 310	Leu	Arg	Arg	Glu	Leu 315	Asp	Glu	Ser	Leu	Gln	Val 320
Ala	Glu	Arg	Leu 325	Thr	Arg	Lys	Tyr	Asn 330	Glu	Leu	Leu	Lys	Ser	Tyr 335	Gln
Trp	Lys	Met 340	Leu	Asn	Thr	Ser	Ser	Leu 345	Leu	Glu	Gln	Leu	Asn 350	Glu	Gln
Phe	Asn 355	Trp	Val	Ser	Arg	Leu	Ala 360	Asn	Leu	Thr	Gln 365	Gly	Glu	Asp	Gln
Tyr 370	Tyr	Leu	Arg	Val	Thr 375	Thr	Val	Ala	Ser	His 380	Thr	Ser	Asp	Ser	Asp
Val 385	Pro	Ser	Gly	Val	Thr 390	Glu	Val	Val	Val 395	Lys	Leu	Phe	Asp	Ser	Asp 400
Pro	Ile	Thr 405	Val	Thr	Val	Pro	Val	Glu 410	Val	Ser	Arg	Lys	Asn 415	Pro	Lys
Phe	Met	Glu 420	Thr	Val	Ala	Glu	Lys 425	Ala	Leu	Gln	Glu	Tyr	Arg 430	Lys	Lys
His	Arg 435	Glu	Glu	Leu	Asp	Asp 440	Asp	Lys	His 445	His	His	His	His	His	His

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<211> LENGTH: 440
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic peptide

<400> SEQUENCE: 15
Thr Trp Glu Ser Gly Gln Val Leu Gly Asp Gln Thr Val Ser Asp Asn
1          5          10          15
Glu Leu Gln Glu Met Ser Asn Gln Gly Ser Lys Tyr Val Asn Lys Glu
          20          25          30
Ile Gln Asn Ala Val Asn Gly Val Lys Gln Ile Lys Thr Leu Ile Glu
          35          40          45
Lys Thr Asn Glu Glu Arg Lys Thr Leu Leu Ser Asn Glu Asp Ala Leu
          50          55          60
Asn Glu Thr Arg Glu Ser Glu Thr Lys Leu Lys Glu Leu Pro Gly Val
          65          70          75          80
Cys Asn Glu Thr Met Met Ala Leu Trp Glu Glu Cys Lys Pro Cys Leu
          85          90          95
Lys Gln Thr Cys Met Lys Phe Tyr Ala Arg Val Cys Arg Ser Gly Ser
          100         105         110
Gly Leu Val Gly Arg Gln Leu Glu Glu Phe Leu Asn Gln Ser Ser Pro
          115         120         125
Phe Tyr Phe Trp Met Asn Gly Asp Arg Ile Asp Ser Leu Leu Glu Asn
          130         135         140
Asp Arg Gln Gln Thr His Met Leu Asp Val Met Gln Asp His Phe Ser
          145         150         155         160
Arg Ala Ser Ser Ile Ile Asp Glu Leu Phe Gln Asp Arg Phe Phe Thr
          165         170         175
Arg Glu Pro Gln Asp Thr Tyr His Tyr Leu Pro Phe Ser Leu Pro His
          180         185         190
Arg Arg Pro His Phe Phe Phe Pro Lys Ser Arg Ile Val Arg Ser Leu
          195         200         205
Met Pro Phe Ser Pro Tyr Glu Pro Leu Asn Phe His Ala Met Phe Gln
          210         215         220
Pro Phe Leu Glu Met Ile His Glu Ala Gln Gln Ala Met Asp Ile His
          225         230         235         240
Phe His Ser Pro Ala Phe Gln His Pro Pro Thr Glu Phe Ile Arg Glu
          245         250         255
Gly Asp Asp Asp Arg Thr Val Cys Arg Glu Ile Arg His Asn Ser Thr
          260         265         270
Gly Cys Leu Arg Met Lys Asp Gln Cys Asp Lys Cys Arg Glu Ile Leu
          275         280         285
Ser Val Asp Cys Ser Thr Asn Asn Pro Ser Gln Ala Lys Leu Arg Arg
          290         295         300
Glu Leu Asp Glu Ser Leu Gln Val Ala Glu Arg Leu Thr Arg Lys Tyr
          305         310         315         320
Asn Glu Leu Leu Lys Ser Tyr Gln Trp Lys Met Leu Asn Thr Ser Ser
          325         330         335
Leu Leu Glu Gln Leu Asn Glu Gln Phe Asn Trp Val Ser Arg Leu Ala
          340         345         350
Asn Leu Thr Gln Gly Glu Asp Gln Tyr Tyr Leu Arg Val Thr Thr Val
          355         360         365

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Ala Ser His Thr Ser Asp Ser Asp Val Pro Ser Gly Val Thr Glu Val
370 375 380

Val Val Lys Leu Phe Asp Ser Asp Pro Ile Thr Val Thr Val Pro Val
385 390 395 400

Glu Val Ser Arg Lys Asn Pro Lys Phe Met Glu Thr Val Ala Glu Lys
405 410 415

Ala Leu Gln Glu Tyr Arg Lys Lys His Arg Glu Glu Leu Asp Asp Asp
420 425 430

Asp Lys His His His His His His
435 440

<210> SEQ ID NO 16
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Protease cleavage site

<400> SEQUENCE: 16

Leu Val Pro Arg Gly Ser
1 5

<210> SEQ ID NO 17
<211> LENGTH: 5
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Protease cleavage site

<400> SEQUENCE: 17

Asp Asp Asp Asp Lys
1 5

<210> SEQ ID NO 18
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 18

cacaatatca aggatatcga cgtga 25

<210> SEQ ID NO 19
<211> LENGTH: 25
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 19

acatcagttc tgttcttcgg gtaca 25

<210> SEQ ID NO 20
<211> LENGTH: 24
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 20

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ttgtatgtgg acttcagtga tgtg	24
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<210> SEQ ID NO 21
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 21

agttcaggtg gtcagcaagg	20
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<210> SEQ ID NO 22
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 22

tggctatagg atctgggtgc	20
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<210> SEQ ID NO 23
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 23

atttgctttt gcctgtttgg	20
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<210> SEQ ID NO 24
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 24

ttacctacac cccgccagtc	20
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<210> SEQ ID NO 25
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic primer

<400> SEQUENCE: 25

tgctgggtctg gaagggtcc	19
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<210> SEQ ID NO 26
<211> LENGTH: 15
<212> TYPE: DNA
<213> ORGANISM: Artificial sequence
<220> FEATURE:
<223> OTHER INFORMATION: Synthetic oligonucleotide

<400> SEQUENCE: 26

gcccttggtg aaggg	15
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<210> SEQ ID NO 27

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<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
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<400> SEQUENCE: 27
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Leu Glu Glu Ala Lys Lys Lys Lys
1                      5
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<210> SEQ ID NO 28
<211> LENGTH: 9
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens
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<400> SEQUENCE: 28
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Leu Leu Leu Phe Val Gly Leu Leu Leu
1                      5
```

1. (canceled)
2. The method of claim 47, comprising a Clusterin coding sequence having a deletion of the nuclear localization signal and/or transmembrane domain.
3. The method of claim 47, comprising a Clusterin coding sequence and a heterologous polypeptide sequence fused to said Clusterin coding sequence.
4. The method of claim 3, wherein the heterologous polypeptide sequence comprises a protease cleavage site.
5. The method of claim 3, wherein the protease cleavage site is a thrombin cleavage site.
6. The method of claim 47, wherein the polypeptide is a fusion protein comprising the Clusterin coding sequence and a heterologous polypeptide sequence.
7. The method of claim 47, wherein the polypeptide is aglycosylated.
8. The method of claim 47, wherein the Clusterin coding sequence has a deletion of the nuclear localization signal.
9. The method of claim 47, wherein the Clusterin coding sequence has a deletion of the transmembrane domain.
10. The method of claim 9, wherein the deletion of the transmembrane domain disrupts endoplasmic reticulum (ER)-targeting of the polypeptide.
11. The method of claim 47, further comprising a tag sequence.
12. The method of claim 11, further comprising a protease cleavage site positioned between the tag sequence and the clusterin coding sequence.
13. The method of claim 11, wherein the protease cleavage site is a thrombin or enteropeptidase cleavage site.
14. The method of claim 11, wherein the tag sequence is a polyhistidine tag.
15. The method of claim 11, wherein the tag sequence is positioned N-terminally relative to the Clusterin coding sequence.
16. The method of claim 11, wherein the tag sequence is positioned C-terminally relative to the Clusterin coding sequence.
- 17-46. (canceled)
47. A method of treating or preventing alcoholism in a subject comprising administering an effective amount of a composition comprising a polypeptide comprising:
 - (i) a Clusterin coding sequence, said coding sequence having a deletion of the nuclear localization signal and/or transmembrane domain; or
 - (ii) a Clusterin coding sequence and a heterologous polypeptide sequence fused to said Clusterin coding sequence.
48. The method of claim 47, wherein the composition is administered by intravenous injection, intratissue injection and/or catheter delivery.
49. The method of claim 47, wherein the nucleic acid sequence encoding the Clusterin polypeptide is operably linked to a promoter.
50. The method of claim 49, wherein the promoter is a gene promoter functional in mammalian cells.
51. The method of claim 50, wherein the polynucleotide molecule is part of a gene expression vector, such as a viral expression vector.

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