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**Wang ET AL "CRISPR-Cas9 Targeting of PCSK9 in Human Hepatocytes In Vivo—Brief Report", ARTERIOSCLER THROMB VASC BIOL., vol. 36, no. 5, 2016, pages 783-6**



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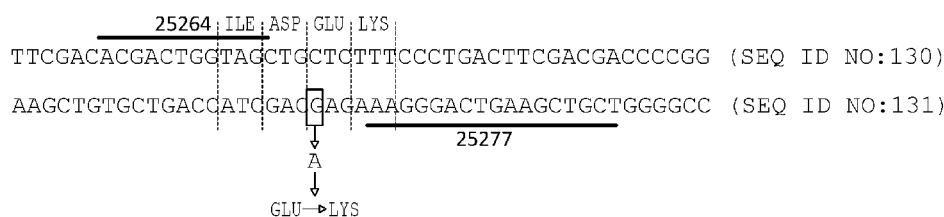


Figure 1A

(57) Abstract: Described herein are compositions and methods for modulation of gene expression in the liver including modulation of PCSK9, TTR, SERPINA1, KLKB1 and/or HA01. Specifically, the disclosure provides methods for modulating the expression of the PCSK9, TTR, SERPINA1, KLKB1 and/or HA01 genes in the liver using one or more engineered nucleases.

## MODULATION OF LIVER GENES

### CROSS-REFERENCE TO RELATED APPLICATIONS

- 5    **[0001]**       The present application claims the benefit of U.S. Provisional Application No. 62/384,428 filed September 7, 2016, the disclosure of which is hereby incorporated by reference in its entirety.

### TECHNICAL FIELD

- 10   **[0002]**       The present disclosure is in the field of gene therapy, particularly knock- out of disease related endogenous genes and targeted delivery of transgene-encoding constructs to the liver for expression of inhibitory proteins.

### BACKGROUND

- 15   **[0003]**       Gene therapy can be used to genetically engineer a cell to have one or more inactivated genes and/or to cause that cell to express a product not previously being produced in that cell (*e.g.*, via transgene insertion and/or via correction of an endogenous sequence). Examples of uses of transgene insertion include the insertion of one or more genes encoding one or more novel therapeutic proteins, insertion of a  
20   coding sequence encoding a protein that is lacking or dysfunctional in the cell or in the individual, insertion of a wild type gene or fragment thereof in a cell containing a mutated gene sequence, and/or insertion of a sequence that encodes a structural nucleic acid such as a microRNA or siRNA. Examples of useful applications of ‘correction’ of an endogenous gene sequence include alterations of disease-associated  
25   gene mutations, alterations in sequences encoding splice sites, alterations in regulatory sequences and/or targeted alterations of sequences encoding structural characteristics of a protein.

- [0004]**       Hepatic gene transfer provides an effective means of delivering transgenes to a subject for treatment and/or prevention of various disorders, including  
30   hemophilias and lysosomal storage disorders. *See, e.g.*, U.S. Patent No. 9,150,847 and U.S. Publication Nos. 20170119906, 20130177983 and 20140017212. Vectors specific for liver-directed gene therapy have also been described. *See, e.g.*, WO 2014064277; WO 2009130208; PCT Publication No. WO 2017/074526; EP 2451474B1, Chuah *et al.*, (2014) *Molecular Therapy*, 22, 1605-1613; and Nair *et al.*

(2014) *Blood* 123:3195-3199. These vectors can include the wild-type mouse minute virus (MVM) intron sequence. *See, e.g.,* Haut and Pintel (1998) *J. Virol.* 72:1834-1843; Haut and Pintel (1998) *Virol.* 258:84-94. Thus, liver-directed gene therapy holds promise for treatment or prevention of a number of diseases.

5     **[0005]**         Artificial transcription factors and nucleases, such as engineered zinc finger protein transcription factors (ZFP-TFs), transcription-activator like effector transcription factors (TALE-TFs), CRISPR/Cas transcription factors (CRISPR-TFs), zinc finger nucleases (ZFN), transcription-activator like effector nucleases (TALENs), the CRISPR/Cas system with an engineered crRNA/tracr RNA ('single guide RNA'),  
 10     also referred to as RNA guided nucleases, and/or nucleases based on the Argonaute system (*e.g.,* from *T. thermophilus*, known as 'TtAgo', (Swarts *et al* (2014) *Nature* 507(7491): 258-261), comprise DNA binding domains (nucleotide or polypeptide) associated with or operably linked to transcriptional regulatory domains (for  
 15     transcription factors) or cleavage domains (for nucleases), and have been used for modulation of gene expression and targeted alteration of genomic sequences. For  
 20     example, artificial nucleases have been used to insert exogenous sequences, inactivate one or more endogenous genes, create organisms (*e.g.,* crops) and cell lines with altered gene expression patterns, and the like. *See, e.g.,* U.S. Patent Nos. 9,255,250; 9,200,266; 9,045,763; 9,005,973; 8,956,828; 8,945,868; 8,703,489; 8,586,526;  
 25     6,534,261; 6,599,692; 6,503,717; 6,689,558; 7,067,317; 7,262,054; 7,888,121; 7,972,854; 7,914,796; 7,951,925; 8,110,379; 8,409,861; U.S. Patent Publications 20030232410; 20050208489; 20050026157; 20050064474; 20060063231; 20080159996; 201000218264; 20120017290; 20110265198; 20130137104; 20130122591; 20130177983 and 20130177960 and 20150056705. Similarly,  
 30     artificial transcription factors targeted to particular sequences have been used to activate or repress endogenous gene expression. *See, e.g.,* U.S. Patent Nos. 9,234,016; 8,563,314 and 8,841,260; Perez-Pinera *et al.* (2013) *Nature Methods* 10:973-976. Clinical trials using these engineered transcription factors containing zinc finger proteins have shown that these novel transcription factors are capable of  
 treating various conditions. (*see, e.g.,* Yu *et al.* (2006) *FASEB J.* 20:479-481).

**[0006]**         However, there remains a need for the treatment or prevention of a number of diseases that can be addressed through liver-directed gene therapy. Such diseases include TTR-Mediated Amyloidosis, A1AT Deficiency, Hereditary

Angioedema, Familial Hypercholesterolemia/Static resistant hypercholesterolemia and Hyperxoluria.

#### SUMMARY

**[0007]** The present invention describes compositions and methods for modulating the expression of the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 genes in the liver and/or for modulating the amount and/or activity of their gene products in the serum following expression of a specific inhibitor of the gene product from a liver cell. Modulation of the gene expression for these genes can be accomplished via genetic modification (*e.g.*, cleavage which results in sequence modifications to the gene resulting in gene knock-out) of the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 genes using one or more engineered nucleases and/or via introduction of a modulator (activator or inhibitor) of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene expression such as a transcription factor that regulates (turns off or downregulates, or turns on or up-regulates) expression of the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 genes. The modulator of gene expression may either act by direct action on the PCSK9, TTR, SERPINA1, KLKB1 and HAO1 genes and/or by indirect action (*e.g.*, activation of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 inhibitors).

**[0007a]** In one embodiment, the present invention provides a liver cell comprising a zinc finger protein that modulates expression of an endogenous Serpin Family A Member 1 (SERPINA1) gene at the Z mutation locus and further wherein the zinc finger protein that modulates SERPINA1 gene expression comprises 4 or 5 zinc finger domains ordered F1 to F4 or F1 to F5, each zinc finger domain comprising a recognition helix region as shown in a single row of the following Table:

F1	F2	F3	F4	F5
QSGSLTR (SEQ ID NO:67)	QSADRTK (SEQ ID NO:68)	RSDHLST (SEQ ID NO:69)	QSAHRIT (SEQ ID NO:70)	N/A
QSSDLRR (SEQ ID NO:71)	QSSDLSR (SEQ ID NO:15)	QSGNLAR (SEQ ID NO:14)	QSGHLAR (SEQ ID NO:72)	RLDNRT A (SEQ ID NO:73)

and wherein the liver cell comprises an oligonucleotide donor that corrects the G->A mutation at the Z mutation locus and wherein expression of an endogenous SERPINA1 gene is altered as compared to wild-type.

[0008] Additionally, provided herein are methods and compositions for deleting (inactivating) or repressing the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 genes to produce a PCSK9, TTR, SERPINA1, KLKB1 or HAO1 null cell, stem cell, tissue or whole organism. Modulation of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 genes can also be achieved by inhibition of the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene products, for example by introduction of a transgene into a liver cell that encodes the inhibitor, which may be expressed extra-chromosomally (episomally) or may be integrated into the genome of the liver cell (*e.g.*, via nuclease-mediated targeted integration, for example into an albumin locus). In some embodiments, the transgene encodes an antibody or polypeptide capable of inhibiting the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene products. In some aspects, the inhibitor is an inhibitory nucleic acid such as an RNAi.

[0009] Thus, in one aspect, described herein are cells in which the expression of a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene is modulated. In some embodiments, the cells comprise a knock-out of a PCSK9, TTR, SERPINA1, KLKB1

[Text continued on page 4]

and/or HAO1 gene, for example a cell in which PCSK9, TTR, SERPINA1, KLKB1 or HAO1 is inactivated (partially or fully) using one or more engineered nucleases to knockout a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene. In other embodiments, cells are described that comprise an engineered transcription factor (TF) such that the expression of a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene is modulated. In some embodiments, the cells are liver cells. Further described are cells in which the expression of a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene is modulated (*e.g.*, via nuclease-mediated inactivation and/or using one or more engineered TFs) and wherein the cells are further engineered to comprise a least one exogenous transgene or an additional knock out of at least one endogenous gene or combinations thereof. The exogenous transgene may be integrated into a PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene and/or may be integrated into a safe harbor locus. In some cases, the exogenous transgene encodes a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 inhibitor (*e.g.*, a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 polypeptide inhibitors such as an antibody and/or a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 inhibitor RNA molecule).

**[0010]** In certain embodiments, the cells described herein comprise a modification (*e.g.*, deletion and/or insertion) to a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene sequence in which the modification is made using a nuclease that binds to a sequence within the selected target gene. In certain embodiments, the DNA-binding domain (*e.g.*, ZFP, TALE, single guide RNA, etc.) of the nuclease used for modification of the indicated target gene binds to a target site of 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21 or even more base pairs) as shown in herein (*e.g.*, Tables 1, 3, 5, 7, 11, 13, 14, and 16). The nuclease target sites may be contiguous or non-contiguous sequences. In certain embodiments, paired nucleases used. Nuclease-mediated modification as described herein can result in modifications (insertions and/or deletions) that are within or near nuclease(s) binding and/or cleavage site(s), including but not limited to, modifications to sequences within 1-300 (or any number of base pairs therebetween) base pairs upstream, downstream and/or including 1 or more base pairs of the site(s) of cleavage and/or binding site; modifications within 1-100 base pairs (or any number of base pairs therebetween) of including and/or on either side of the binding and/or cleavage site(s); modifications within 1 to 50 base pairs (or any number of base pairs therebetween) including and/or on either side of the binding and/or cleavage site(s); and/or modifications to one or more base pairs within

the nuclease binding site and/or cleavage site. In certain embodiments, the modification is within or near to the target sites shown herein, including but not limited to modifications in the genome of the cell within, between (*e.g.*, for paired target sites) or near (*e.g.*, 1-50 nucleotides or more (or any number of nucleotides)) the target sites shown herein (*e.g.*, Tables 1, 3, 5, 7, 11, 13, 14, and 16).

**[0011]** In another aspect, the compositions (modified cells) and methods described herein can be used, for example, in the treatment and/or prevention or amelioration of a disorder. The methods typically comprise (a) cleaving or down regulating expression of an endogenous PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene in a cell (*e.g.* a hepatocyte) using a nuclease (*e.g.*, ZFN or TALEN) or RNA-guided nuclease system such as CRISPR/Cas with an engineered crRNA/tracr RNA, or using an engineered transcription factor (*e.g.* ZFP-TF, TALE-TF or Cas9-TF) such that the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene is inactivated or down modulated; thereby treating or preventing the disorder.

**[0012]** Also described herein are artificial nucleases and/or transcription factors comprising a DNA-binding domain (*e.g.*, ZFP, TALE, sgRNA, etc.) that binds to a target site within a liver-specific gene (*e.g.*, PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1) and a functional domain (*e.g.*, nuclease/cleavage domain in the case of nuclease and transcriptional activation or repression domain in the case of transcription factors). The DNA-binding domain may bind to any sequence within the target sequence to effect modulation of the gene. In certain embodiments, the DNA-binding domain binds to a target site of 12 or more (12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22 or more) nucleotides of the target sites as shown in the appended Tables. The bound nucleotides may be contiguous or non-contiguous. Binding of the DNA-binding domain of an artificial transcription factor as described herein to its target site then modulates expression of the target gene via up-regulation or down-regulation. Similarly, binding of the DNA-binding domain of an artificial nuclease as described herein (*e.g.*, a DNA-binding domain in association with a cleavage/nuclease domain) causes a break (single- or double-stranded) in the target gene that results in modification via NHEJ mediated repair (insertions and/or deletions known as “indels”), NHEJ-mediated integration of a donor (*e.g.*, via end capture) and/or homology-directed repair (*e.g.*, integration of a donor with homology arms into the break). The site of cleavage and/or modification may be within or adjacent to (*e.g.*, 1-



50 nucleotides, including, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10 or more) the nuclease target site or may be between paired target sites.

**[0013]** The nuclease(s), transcription factor(s) and/or transgenes can be introduced as mRNA, in protein form and/or as a DNA sequence encoding one or more components of the nuclease(s), TFs and/or transgenes. In one aspect, provided herein are zinc finger nucleases (ZFNs), TALENs and/or CRISPR/Cas systems that cleave a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene. In other embodiments, provided herein are ZFP-TFs, TALE-TFs and Cas9-TFs that modulate expression of a PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene. In certain embodiments, the ZFPs, TALEs or single guide RNAs (sgRNA) of a CRISPR/Cas system of the nucleases or transcription factors bind to target sites in a human PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene. The zinc finger proteins may include 1, 2, 3, 4, 5, 6 or more zinc fingers, each zinc finger having a recognition helix that specifically contacts a target subsite in the target gene. In certain embodiments, the zinc finger proteins comprise 4 or 5 or 6 fingers (designated F1, F2, F3, F4, F5 and F6 and ordered F1 to F4 or F5 or F6 from N-terminus to C-terminus). In other embodiments, the single guide RNAs may bind to a target site in the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene.

**[0014]** Any of the nucleases described herein may further comprise a cleavage domain and/or a cleavage half-domain (*e.g.*, a wild-type or engineered *FokI* cleavage half-domain). Thus, in any of the ZFNs and/or TALENs described herein, the nuclease domain may comprise a wild-type nuclease domain or nuclease half-domain (*e.g.*, a *FokI* cleavage half domain). In other embodiments, the ZFNs and/or TALENs comprise engineered nuclease domains or half-domains, for example engineered *FokI* cleavage half domains that form obligate heterodimers. See, *e.g.*, U.S. Patent Publication No. 20080131962. In some embodiments, the ZFNs may further comprise modification to the ZFP backbone to decrease non-specific DNA interactions. In further embodiments, the *FokI* domains in the engineered ZFNs, TALENs, or dCas-*FokI* fusions comprise mutations to disrupt non-specific interactions between the *FokI* domain and the DNA molecule (see U.S. Application No. 15/685,580). In still further embodiments, the ZFNs, TALENs and/or RNA-guided CRISPR/Cas systems act as nickases. In some instances, double strand cleavage is achieved using two pairs of nickases (*e.g.* U.S. Patent No 9,200,266).

Any of the transcription factors described herein may further comprise a transcriptional activation or repression domain.

**[0015]** In another aspect, described herein are methods of inactivating a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene in a cell by introducing one or more proteins, polynucleotides, systems and/or vectors into the cell as described herein. In any of the methods described herein the nucleases may induce targeted mutagenesis, deletions of cellular DNA sequences, and/or facilitate targeted recombination at a predetermined chromosomal locus. Thus, in certain embodiments, the nucleases delete or insert one or more nucleotides of the target gene. In some embodiments the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene is inactivated by nuclease cleavage followed by non-homologous end joining. In other embodiments, a genomic sequence in the target gene is replaced, for example using a nuclease (or vector encoding said nuclease) as described herein and a “donor” sequence that is inserted into the gene following targeted cleavage with the nuclease. The donor sequence may be present in the nuclease vector, present in a separate vector (*e.g.*, AAV, Ad or LV vector) or, alternatively, may be introduced into the cell using a different nucleic acid delivery mechanism.

**[0016]** In certain embodiments, the cell comprising the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 modulations described herein (*e.g.*, down-regulation via a PCSK9, TTR, SERPINA1, KLKB1 or HAO1 targeted nuclease and/or transcription factor) further comprises one or more additional genomic modification, for example, an integrated exogenous sequence (into a cleaved PCSK9, TTR, SERPINA1, KLKB1 or HAO1, or other gene, for example a safe harbor gene or locus). The exogenous sequence may be introduced via a vector (*e.g.* Ad, AAV, LV), or by using a technique such as electroporation. In further embodiments, the additional modification comprises introducing a wildtype cDNA copy of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 into the cell where an endogenous mutant copy in the cell has been knocked out by any of the methods described herein. In some aspects, the wildtype cDNA copy of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 may comprise silent gene modifications such that the cDNA is not subject to cleavage by the targeted nucleases. In some embodiments, the cDNA copy is integrated into the genome, while in other embodiments, the cDNA copy is maintained extrachromosomally.

**[0017]** In some aspects, the PCSK9, TTR, SERPINA1, KLKB1 or HAO1 modulated cells further comprise a gene encoding an exogenous transgene that inhibits PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 activity. In other aspects, described herein are cells lacking PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene modification(s) but comprising a construct for the expression of a transgene that encodes a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 inhibitory activity. In some embodiments, the inhibitory transgene encodes an antibody that inhibits the proteins encoded by PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1. In other embodiments, the inhibitory transgene encodes an inhibitory nucleic acid (e.g. RNAi, shRNA) that inhibits PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 expression. In some embodiments, the inhibitory RNA is a double stranded RNA (U.S. Patent No. 9,249,415). Also described are cells produced by the methods described herein, including cells descended from these cells (e.g., genetically modified cells descended from the cells modified as described herein). In certain embodiments, the genetic modifications comprise one or more insertions and/or deletions within a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene, including, but not limited to, insertions and/or deletions within the target sites disclosed herein and/or between paired target sites as described herein.

**[0018]** In one aspect, described herein is a polynucleotide expression construct comprising a sequence encoding a transgene that encodes one or more modulators of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene expression (e.g., the engineered nuclease and/or engineered transcription factor targeted to PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1). In certain embodiments, the polynucleotide expression construct further comprises an enhancer sequence (e.g., a wild-type or mutated Serpin1 enhancer), a promoter sequence (e.g., a transthyretin minimal promoter (TTRm) promoter), and the transgene and, optionally, a polyadenylation sequence (e.g., a synthetic polyadenylation sequence (SPA) and/or a signal peptide (SP) (see U.S. Patent Publication No. US-2017-0119906-A1). In certain embodiments, the expression construct further comprises an intron sequence (e.g., wild-type MVM or a mutated MVM sequence and/or chimeric intron). In certain embodiments, the expression constructs comprise in 5' to 3' orientation, an enhancer sequence, a promoter sequence, an intronic sequence, a transgene (optionally comprising a signal peptide), and a polyadenylation signal.

[0019] The expression cassette encoding the gene modulators as described herein may be included in any viral or non-viral vector, including but not limited to plasmid vectors, adenovirus vector, retroviral vectors and adeno associated vector (AAV). In a preferred embodiments, the expression construct is carried on an AAV construct and further comprises 5' and 3' ITRs flanking the expression constructs as described herein. Optionally, spacer molecules are also included between one or more of the components of the expression construct, for example, between the 5' ITR and the enhancer and/or between the polyadenylation signal and the 3' ITR.

[0020] In some embodiments, the one or more transgenes include sequences encoding gene modulators such as engineered nucleases (*e.g.* ZFNs, TALENs, TtAgo and CRISPR/Cas systems). In other embodiments, the gene modulating transgenes include sequences encoding engineered transcription factors (*e.g.* ZFP-TFs, TALE-TFs, CRISPR/Cas-TF systems). In certain embodiments, the gene modulators (*e.g.*, nucleases and/or transcription factors) are targeted to PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1.

[0021] In certain aspects, the polynucleotides as described herein are introduced into a cell such that they are maintained episomally while driving expression of the PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene-modulating transgene. In other aspects, the expression constructs are randomly integrated into the genome of the cell into which they are introduced. In further aspects the expression constructs driving transgene expression are integrated into a genome by nuclease-mediated targeted integration.

[0022] In still further aspects, described herein are cells comprising any of the gene modulators disclosed herein (polynucleotides and/or proteins) as well as cells comprising any genetic modifications made by the gene modulators described herein. Also provided are cells descended from these cells, for example cells descended (including but not limited to cells differentiated from stem or progenitor cells) from genetically modified cells as described herein, in which the cells no longer comprise the gene modulator(s). The isolated cells may be introduced into the subject (*ex vivo* cell therapy) or cells as described herein (*e.g.*, genetically modified cells) may be modified when it is part of the subject (*in vivo*).

[0023] In further aspects, described herein are methods for expressing one or more transgenes (modulators of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1) in a liver cell, the methods comprising introducing one or more expression constructs as

described herein into the cell such that the transgene is expressed in the cell. In certain embodiments, the expression construct is carried on a viral or non-viral vector, preferably an AAV vector (for example AAV2, AAV6, AAV8, AAV2/6, or AAV2/8).

5     **[0024]**           In another aspect, provided herein is a method of expressing one or more transgenes (modulators of PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1) in a live animal, the methods comprising administering one or more expression cassettes as described herein to the live animal. In certain embodiments, the expression cassettes are administered to the liver of the live animal. In certain embodiments, the  
10    expression construct is carried on a viral or non-viral vector, preferably an AAV vector (for example AAV2 or AAV2/6). In some embodiments, the expression construct is administered systemically via a peripheral vein (*e.g.* intravenously).

**[0025]**           In another aspect, pharmaceutical compositions comprising one or more of cells, expression constructs, transcription factors and/or nucleases described  
15    herein are provided.

**[0026]**           In certain aspects, described herein are compositions, methods and systems for targeted integration of a liver-specific expression cassette. The methods and systems comprise administering one or more expression cassettes as described herein and administering one or more nucleases specific for a target gene (*e.g.*,  
20    PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 and/or safe harbor gene) to a cell. Following nuclease-mediated cleavage of the target gene, the expression cassette is integrated into the gene via homology-dependent or homology-independent mechanisms. In certain embodiments, the target gene is an endogenous albumin gene.

**[0027]**           For nuclease-mediated targeted integration of the expression  
25    constructs of the present invention, any nuclease can be used, including but not limited to, one or more zinc finger nucleases (ZFNs), TALENs, CRISPR/Cas nucleases and/or TtAgo nucleases, such that the expression construct is integrated into the region (gene) cleaved by the nuclease(s). In certain embodiments, one or more pairs of nucleases are employed. The nucleases may be introduced in mRNA form or  
30    may be administered to the cell using non-viral or viral vectors. In some aspects, the nuclease polynucleotides may be delivered by lentivirus or by non-integrating lentivirus. In other aspects, the expression cassette may be delivered by AAV and/or DNA oligos.

**[0028]** In a further aspect, methods and compositions are described herein for providing a therapeutic protein for treating a disorder where the therapeutic protein is a single chain antibody. In certain embodiments, the methods comprise administering an expression cassette (*e.g.*, AAV vector) as described herein to the liver of a subject in need thereof. In other embodiments, the method comprises administering a modified cell (expressing a functional version of a protein that is aberrantly expressed in a subject from an expression cassette as described) to the subject.

**[0029]** In any of the compositions and methods described, expression cassettes and/or nucleases may be carried on an AAV vector, including but not limited to AAV1, AAV3, AAV4, AAV5, AAV6, AAV8, AAV9 and AAVrh10 or pseudotyped AAV such as AAV2/8, AAV8.2, AAV2/5 and AAV2/6 and the like. In certain embodiments, the polynucleotides (expression constructs and/or nucleases) are delivered using the same AAV vector types. In other embodiments, the polynucleotides are delivered using different AAV vector types. The polynucleotides may be delivered using one or more vectors. In certain embodiments, the polynucleotides are delivered via intravenous (*e.g.*, intra-portal vein or peripheral vein) administration into the liver of an intact animal.

**[0030]** The compositions and methods described herein can be used to treat or prevent disorders including *e.g.*, TTR-Mediated Amyloidosis, A1AT Deficiency, Hereditary Angioedema, Familial Hypercholesterolemia/Static resistant hypercholesterolemia and Hyperxoluria.

**[0031]** The methods described herein can be practiced *in vitro*, *ex vivo* or *in vivo*. In certain embodiments, the compositions are introduced into a live, intact mammal. The mammal may be at any stage of development at the time of delivery, *e.g.*, embryonic, fetal, neonatal, infantile, juvenile or adult. Additionally, targeted cells may be healthy or diseased. In certain embodiments, one or more of the compositions are delivered intravenously (*e.g.*, to the liver via the intraportal vein, for example tail vein injection or systemically via a peripheral vein), intra-arterially, intraperitoneally, intramuscularly, into liver parenchyma (*e.g.*, via injection), into the hepatic artery (*e.g.*, via injection), and/or through the biliary tree (*e.g.*, via injection).

**[0032]** For targeting the compositions to a particular type of cell, *e.g.*, platelets, fibroblasts, hepatocytes, etc., one or more of the administered compositions may be associated with a homing agent that binds specifically to a surface receptor of

the cell. For example, the vector may be conjugated to a ligand (*e.g.*, galactose) for which certain hepatic system cells have receptors. The conjugation may be covalent, *e.g.*, a crosslinking agent such as glutaraldehyde, or noncovalent, *e.g.*, the binding of an avidinated ligand to a biotinylated vector. Another form of covalent conjugation is provided by engineering the AAV helper plasmid used to prepare the vector stock so that one or more of the encoded coat proteins is a hybrid of a native AAV coat protein and a peptide or protein ligand, such that the ligand is exposed on the surface of the viral particle.

**[0033]** A kit, comprising the cells and/or expression constructs described herein, is also provided. The kit may further comprise nucleic acids encoding nucleases, (*e.g.* RNA molecules encoding ZFNs, TALENs or Cas and modified Cas proteins, and guide RNAs), transcription factors or aliquots of the nucleases, transcription factors, cells, instructions for performing the methods of the invention, and the like.

**[0034]** These and other aspects will be readily apparent to the skilled artisan in light of disclosure as a whole.

#### BRIEF DESCRIPTION OF THE DRAWINGS

**[0035]** **Figures 1A and 1B** depict partial sequence of the region in exon 5 of the human SERPINA1 gene near where the Z mutation is located. Figure 1A (SEQ ID NO:130 and 131) shows the site of the G->A mutation in the Z mutation gene sequence and also shows where the SERPINA1-specific ZFNs bind. Figure 1B (SEQ ID NO:132) shows the G->A mutation and also indicates the location of the sites of the altered nucleotides following integration of a gene correction donor.

**[0036]** **Figures 2A and 2B** are graphs depicting the amount of NHEJ and TI in mice that have been treated with ZFN and the gene-correction donor where the animals were sacrificed at two weeks. Figure 2A depicts the amount of ZFN-mediated NHEJ detected in the livers of the mice who were treated with either AAV8-ZFNs alone (25264 & 25277) at a low dose (7.5e10 vg per ZFN per mouse) or a high dose (1.5 e11 vg per ZFN per mouse) while Figure 2B depicts the amount of targeted integration of the gene correction donor (1.5e12 vg per mouse AAV8 comprising the gene correction donor).

**[0037]**        **Figures 3A through 3C** are graphs showing the NHEJ and TI data in mice that had been treated with ZFN and/or the gene correction donor where the animals were sacrificed at 6 months. Figure 3A depicts the amount of ZFN-mediated NHEJ detected in the livers of the mice who were treated with either AAV8-ZFNs alone (25264 & 25277) at a low dose ( $7.5 \times 10^{10}$  vg per ZFN per mouse AAV8) or a high dose ( $1.5 \times 10^{11}$  vg per ZFN per mouse) and Figure 3B depicts the amount of targeted integration of the gene correction donor ( $1.5 \times 10^{12}$  vg per mouse AAV8 comprising the gene correction donor). Figure 3C shows a direct comparison in the amount of detected NHEJ for the 2 week samples as compared to the 6 month samples.

**[0038]**        **Figures 4A through 4E** (SEQ ID NO:133-138) show partial sequences of TTR gene including potential areas to target with ZFNs. The heaviness of the box around the letter name of each target matches the heaviness of the target lines. Figures 4A through 4C depict the Exon 1 sequence while Figure 4D shows sequence in Exon 2 and Figure 4E shows sequence in Exon 3. For example, target 'A' is indicated with the light lines in Figure 4A. Target 'B' is indicated by the heavier lines where the B pair will both bind to the sense (Watson) strand of the gene, while the 'C' target is bound by a ZFN pair that binds to the 5' sense target of B, and the C target on the antisense strand (also in a heavy line) in Figure 4A. Figure 4B shows the 'E' target which corresponds to the two heavy lines on the sense strand (shown under the sequence). The 'F' target uses the heavy E target line at the 5' end of the sense strand and the heavy target line on the antisense strand near F. The G target is depicted as the light lines on the sense strand of Figure 4B. For Figure 4C, the 'D' target is indicated by the light lines under the sense and antisense strands. Figure 4D shows the 'H', 'I' and 'J' target sequences where all letter boxes are shown in the center of the two ZFN binding sequences. Figure 4E depicts two ZFN targets, 'L' and 'M' where the L pair binds to the heavy lines and the M pair binds to the light lines.

**[0039]**        **Figures 5A and 5B** are graphs depicting the percent gene modification ("indels" indicating insertions and/or deletions characteristic of NHEJ following nuclease-mediated cleavage) in mouse hepatocytes treated with the indicated gene modulators (see Table 11). Figure 5A shows results with mRNA dose per ZFN in B16-F10 cells when the ZFN-encoding mRNAs are introduced via BTX electroporation, three days post transfection. Figure 5B shows results in mouse



hepatocytes three days post transfection when the ZFN-encoding mRNAs are introduced via lipofection.

**[0040]**       **Figures 6A and 6B** are graphs depicting the activity of TTR ZFNs in mice. Figure 6A shows the cleavage activity (% indels) in livers from mice treated with the indicated ZFNs, either at a high ( $1.5 \times 10^{11}$  vg per ZFN per mouse) or low ( $2.5 \times 10^{10}$  vg per ZFN per mouse) doses, harvested 28 days post treatment. Figure 6B shows the plasma mTTR concentration under the same treatment conditions. The data demonstrates that the ZFN cleave their targets *in vivo*, and cause a reduction in plasma TTR.

**[0041]**       **Figure 7** is a schematic of the mouse PCSK9 gene, showing the exons and the sites selected to target.

**[0042]**       **Figures 8A and 8B** are graphs depicting cleavage of the murine TTR targets by ZFN. The ZFNs tested in this analysis comprise mutations in their backbones and in the FokI cleavage domain to remove specific phosphate contacting residues that may be related to off-target cleavage. Figure 8A shows the activity results at the on-target location of each of the test proteins, and Figure 8B shows the activity at the sum of the 3 highest off target loci. The mutant proteins have increased on-target activity and decreased off target activity as compared to the parent proteins.

#### DETAILED DESCRIPTION

**[0043]**       Disclosed herein are methods and compositions to modulate the expression of PCSK9, TTR, SERPINA1, KLKB1 or HAO1 genes as well as cassettes for production of PCSK9, TTR, SERPINA1, KLKB1 or HAO1 inhibitory transgenes, particularly in liver cells. Modulation of PCSK9, TTR, SERPINA1, KLKB1 or HAO1 expression may be achieved through the use of targeted nucleases (ZFN, TALEN, Ttago, CRISPR/Cas) to knock out the gene, and/or transcription factors (ZFP-TF, TALE-TF, RNA guided Cas-TF) to inhibit endogenous PCSK9, TTR, SERPINA1, KLKB1 or HAO1 expression. Cells comprising the gene modulators (polynucleotides and/or proteins) and/or comprising genetic modifications made by the gene modulators (but not comprising the gene modulators themselves) as well as cells descended from these cells are also provided. The methods and compositions of the invention can be used to express PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 inhibitory transgenes via liver-specific expression constructs. The compositions (constructs comprising gene modulators, proteins, and/or cells) as described herein may be delivered any transgene(s) to liver cells, *in vivo* or *in vitro*

and can be used for the treatment and/or prevention of any disease or disorder which can be ameliorated by the provision of one or more of the transgenes via *in vivo* and/or *ex vivo* therapies.

**[0044]** Provided herein is a liver cell in which expression of an endogenous PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene is altered as compared to wild-type, for example by cleaving the gene using at least one artificial nuclease comprising a DNA-binding domain (that binds to a target site in the endogenous gene) and a cleavage domain or using an artificial transcription factor (repressor or activator) comprising a DNA-binding domain (that binds to a target site in the endogenous gene) and a transcriptionally regulatory domain (activator or repressor) such that expression is altered. The target site can comprise at least 12 (*e.g.*, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22 or more) contiguous or non-contiguous nucleotides as shown in the target sites of any of the appended Tables (*e.g.*, Tables 1, 3, 5, 7, 11, 13, 14 and 16). An exogenous sequence (*e.g.*, a transgene; a sequence that introduces a mutation into the gene, or a sequence that corrects a mutation in the endogenous gene) may be integrated into the endogenous gene following cleavage and/or one or more nucleotides may be inserted and/or deleted following cleavage. Also provided are fusion molecules comprising a DNA-binding domain (*e.g.*, zinc finger protein, TAL-effector domain protein or single-guide RNA (sgRNA)) that binds to a target site in an endogenous PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene and a functional domain (*e.g.*, cleavage domain, transcriptional activation domain or transcriptional repression domain) and polynucleotides (*e.g.*, mRNA, viral or non-viral vectors, etc.) encoding these fusion molecules. Pharmaceutical compositions comprising cells, fusion molecules and/or polynucleotides as described herein are also provided as are methods of making and using the cells, fusion molecules and/or pharmaceutical compositions for the alteration of endogenous gene expression and/or the treatment of disorders such as TTR-Mediated Amyloidosis, A1AT Deficiency, Hereditary Angioedema, Familial Hypercholesterolemia/Static resistant hypercholesterolemia and Hyperoxaluria (via administration thereof to a subject in need).

**[0045]** Hereditary angioedema (HAE) is an autosomal dominant disease that affects 1 in 50,000 people and is a result of decreased levels of the C1 inhibitor. Patients experience recurrent episodes of swelling in any part of the body where swelling localized to the oropharynx, larynx or abdomen carry the highest risk of

morbidity and death (see Tse and Zuraw, (2013) *Clev Clin J of Med* 80(5):297). The disease occurs from extravasation of plasma into tissues as a result of the over production of bradykinin. The mechanism seems to involve the cleavage of prekallikrein (also known as PKK, KLK3, PKKD, Fletcher factor and Kininogenin) by activate factor XII, releasing active plasma kallikrein (which activates more factor XII). Plasma kallikrein then cleaves kininogen, releasing bradykinin. The bradykinin then binds to the B2 bradykinin receptor on endothelial cells, increasing the permeability of the endothelium. Normally, the C1 inhibitor (encoded by SERPING1) controls bradykinin production by inhibiting plasma kallikrein and the activation of factor XII. HAE occurs in three types, Type I and II that are distinguished by the amount and type of C1 inhibitor present, and Type III which is tied to a Thr309Lys mutation in factor XII (Prieto *et al* (2009) *Allergy* 64(2):284). Type I HAE has low levels of C1 inhibitor that appear to be a result of poor expression and destruction of the small amount of C1 inhibitor protein that is made. Type 1 accounts for approximately 85% of HAE patients. Type II patients have normal levels of C1 inhibitor, but the C1 inhibitor protein is ineffectual due to mutations (Tse and Zuraw, *ibid*). More than 250 mutations in SERPING1 have been characterized that lead to Type I HAE including small and large insertions and deletions as well as duplications (Rijavec *et al* (2013) *PLoS One* 8(2): e56712). Due to this high variability in the genetic basis of HAE, the methods and compositions of the invention can be used to prevent or treat HAE by targeting downstream players in the manifestation of HAE. For example, targeting the gene encoding prekallikrein (KLKB1, expressed in hepatocytes) to effect a decrease in prekallikrein (abbreviated PKK) expression can result in a decrease in bradykinin production without regard to the type of mutation upstream that is causing the HAE, and thus result in a decrease in plasma extravasation. Thus, the methods and compositions of the invention may be used to cause a decrease in the expression of KLKB1 to prevent or treat HAE. Engineered nucleases can be used to knock out KLKB1 in a subset of hepatocytes that will reduce bradykinin production, and/or engineered transcription factors can be used to down regulate KLKB1 expression.

**[0046]** PCSK9 is a gene (also known as FH3; HCHOLA3; LDLCQ1; NARC-1; NARC1; PC9) encoding a protein that plays a major regulatory role in cholesterol homeostasis. The PCSK9 protein binds to the epidermal growth factor-like repeat A (EGF-A) domain of LDLR, and induces LDLR degradation. Autosomal dominant,

toxic gain of function mutations in PCSK9 (*e.g.* S127R, P216L, D374Y and N157K) have been described and are associated with hyperlipidemia and Familial hypercholesterolemia (FH) as a result of an increased rate of LDLR degradation leading to a corresponding increase in plasma LDL cholesterol (Abifadel *et al* (2003)

5 *Nat Gen* 34(2):154). In addition, loss of function PCSK9 mutations have been identified (*e.g.* Y142X, C679X and R46L) that cause an increase in hepatic LDLR levels, with an associated substantial decrease in the amount of plasma LDL cholesterol, leading to an 88% reduction in the incidence of coronary heart disease (Cohen *et al* (2003) *New Eng J Med* 354(12):1264). Thus the methods and  
10 compositions of the invention can be used to treat or prevent hyperlipidemia and/or FH. Engineered nucleases can be designed to knock out a PCSK9 gene comprising a mutation that is associated with a toxic gain of function. Additionally, a wild type PCSK9 gene may be knocked out in a number of cells in the liver to treat FH caused by mutations in other genes such as LDLR or APOB. Alternatively, engineered  
15 transcription factors can be used to repress expression from a mutant or wild type PCSK9 gene.

**[0047]** Transthyretin Amyloidoses (TTRA) is one of several degenerative diseases suspected to be linked to misfolded and aggregated protein (amyloids).

Transthyretin (TTR, also known as CTS; CTS1; HEL111; HsT2651; PALB; TBPA) is  
20 a tetramer produced in the liver and secreted into the bloodstream that serves to transport holo-retinal binding protein. However, upon conformational changes, it becomes amyloidogenic. Partial unfolding exposes stretches of hydrophobic residues in an extended conformation that efficiently misassemble into largely unstructured spherical aggregates that ultimately before cross- $\beta$  sheet amyloid structures (see  
25 Johnson *et al* (2012) *J Mol Biol* 421(2-3):183). TTRA can occur in patients in both sporadic and autosomal dominant inherited forms which include familial amyloid polyneuropathy (FAP) and familial amyloid cardiomyopathy (FAC). These inherited forms are usually earlier onset and relate to over 100 point mutations described in the TTR gene. Generally, the more destabilizing of the protein that the mutation is, the  
30 more likely it is to have some amount of amyloid pathology. The amyloid formed causes selective destruction of cardiac tissue in FAC or peripheral and central nervous tissue in FAP. Some new therapeutic strategies for treating these diseases such as inhibitory RNA strategies center on trying to decrease the amount of TTR to decrease the aggregation potential of the protein (Johnson *et al, ibid*). Thus the methods and

compositions of the invention can be used to target specific TTR mutants, and/or target wild type TTR in an effort to reduce the quantity of the pathological forms of the TTR protein and/or to decrease TTR concentration in general. Engineered nucleases may be used to knock out TTR in a subset of hepatocytes and/or engineered transcription factors specific for TTR may also be used to down regulate its expression.

**[0048]** Alpha-1-antitrypsin (A1AT) deficiency occurs in about 1 in 1500-3000 people of European ancestry but is rare in individuals of Asian descent. The alpha-1-antitrypsin protein is a protease inhibitor that is encoded by the SERPINA1 gene and serves to protect cells from the activity of proteases released by inflammatory cells, including neutrophil elastase, trypsin, metalloproteinase 9 (MMP-9), myeloperoxidase (MPO) and proteinase-3 (PR-3). Deficiency is an autosomal co-dominant or a recessive disorder caused by mutant SERPINA1 genes in heterozygous individuals where reduced expression from the mutant allele or the expression of a mutant A1AT protein with poor inhibitory activity leads to chronic lack of inhibition of neutrophil elastase resulting in tissue damage. The most common SERPINA1 mutation comprises a Glu342Lys substitution (also referred to as the Z allele) that causes the protein to form ordered polymers in the endoplasmic reticulum of patient hepatocytes. These inclusions ultimately cause liver cirrhosis which can only be treated by liver transplantation (Yusa *et al* (2011) *Nature* 478 p.391). The polymerization within the hepatocytes results in a severe decrease in plasma A1AT levels, leading to increased risk of this inflammatory disease. In addition, A1AT deficiency is linked to pulmonary diseases including chronic obstructive pulmonary disease (COPD), emphysema and chronic bronchitis (Tuder *et al* (2010) *Proc Am Thorac Soc* 7(6): p. 381) and potentially may have a far broader reach into the inhibition of the progression of other diseases including type 1 and type 2 diabetes, acute myocardial infarction, rheumatoid arthritis, inflammatory bowel disease, cystic fibrosis, transplant rejection, graft versus host disease and multiple sclerosis (Lewis (2012) *Mol Med* 18(1) p. 957). In infancy and early childhood onset liver disease, the pathology of the disease presents as neonatal jaundice and cholestasis, which can be followed by progression to advanced fibrosis or cirrhosis. In adults, liver disease manifests as slowly progressive fibrosis which is typically diagnosed when the patient is in their fifth decade, where these patients have an increased risk of cirrhosis and hepatocellular carcinoma (Guo *et al* (2014) *J Clin Invest* 124(1):251). Population

studies have suggested a minimum A1AT plasma threshold of approximately 0.5 mg/mL (normal plasma levels are approximately 0.9- 1.75 mg/mL in a non-inflammatory state) to avoid these diseases, and current therapies mostly act to reduce symptoms through the use of bronchodilators and the like, although the use of weekly infusions of A1AT (Zemaira®, Prolastin) is also an option for emphysema patients with a demonstrated severe lack of plasma A1AT (Koepke *et al* (2015) *PLoS One* 10(1): e0117497). Severe lung disease associated with A1AT also is ultimately treated by transplant. Clinical trials for the treatment of A1AT deficiency involve a variety of approaches including the delivery of concentrated A1AT protein, use of an AAV construct comprising an A1AT gene by IM injection, and the use of A1AT in HIV, to list just a few. Thus, the compositions and methods of the invention can be used to treat or prevent diseases related to A1AT deficiency. Transcription factors and systems as described herein that are specific for the mutant A1AT allele (*e.g.* the Z allele) can be made to silence the gene and prevent expression, thereby eliminating the hepatic aggregates that can lead to cirrhosis. In addition, a wild type SERPINA1 gene may be introduced into the genome of the cell for expression, may be introduced via a non-integrating vector system (*e.g.* carrying a cDNA copy of the wild type gene) for extracellular expression, or may be introduced into the albumin locus *in vivo* for increased hepatic secretion while specific SERPINA1 nucleases are introduced to knock out an endogenous mutant SERPINA1 allele (*e.g.* the gene comprising the Z point mutation). In some embodiments, the Z point mutation is corrected by nuclease-driven insertion of a correcting oligonucleotide or partial cDNA such that the point mutation is corrected.

**[0049]** Primary hyperoxaluria type 1 (PH1), an inherited rare autosomal recessive disorder occurring in 0.11 to 0.26 per 100,000 births, is disease of glyoxylate metabolism, and arises from mutations in the enzyme alanine-glyoxylate aminotransferase (AGT, Siegal *et al* (2011) *Int J Organ Transpl Med*). The resulting deficiency in this enzyme leads to abnormally high oxalate production resulting in calcium oxalate crystal formation and deposition in the kidney and many other tissues, with systemic oxalosis and end-stage renal disease (ESRD) being a common outcome. Normally in the cell, glyoxylate present in cytoplasm is converted to glycolate by the enzyme glyoxylate reductase, and the glycolate can be taken up in the peroxisome and converted into glycine by AGT. When AGT is deficient, the accumulating glyoxylate is converted to oxalate by the liver-specific enzyme

glycolate oxidase, or hydroxyacid oxidase-1 (HAO1, also known as HAO1X1, GOX1, GO and GOX). This increased accumulation of oxalate leads to supersaturation of urine with oxalate, and in turn leads to oxalate urolithiasis, nephrocalcinosis, renal tubular damage, renal failure and even death. ESRD in hyperoxaluria is accompanied by calcium oxalate deposition in the skin, bone marrow, bone (causing recurrent bone fractures), myocardium, nervous system, skeletal muscle, blood vessels and retina. Treatment is best initiated in children prior to kidney damage and can involve diet considerations (avoiding vitamin C and high oxalate foods) and daily dialysis. Unfortunately, dialysis is often able to significantly reduce the oxalate load. Liver transplant can be performed, but the therapeutics needed to avoid rejection can compound the kidney disorders associated with the PH1. In one study (Nolkemper *et al* (2000) *Pediatr Transplant* 4(3):177-81), there was an 82% survival when patients were given a liver transplant at 10 years of age, while when the transplant was performed at 20 years of age, survival dropped to 72%. In contrast, when a hepato-renal transplant was performed, there was an 80% patient survival rate when the transplant was performed 5 years of age, but this reduced to only 66% when performed at 20 years old (Eytan and Weismann (2009) *Pediatr Transplant* 13(7):805-807). RNAi approaches may offer some help as injection of GO-specific siRNAs decreased the expression of GO and reduced urinary oxalate excretion in a mouse disease model (Li *et al* (2016) *Biochim Biophys Acta* 1862(2):233-239).

## General

**[0050]** Practice of the methods, as well as preparation and use of the compositions disclosed herein employ, unless otherwise indicated, conventional techniques in molecular biology, biochemistry, chromatin structure and analysis, computational chemistry, cell culture, recombinant DNA and related fields as are within the skill of the art. These techniques are fully explained in the literature. *See*, for example, Sambrook *et al.* MOLECULAR CLONING: A LABORATORY MANUAL, Second edition, Cold Spring Harbor Laboratory Press, 1989 and Third edition, 2001; Ausubel *et al.*, CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, John Wiley & Sons, New York, 1987 and periodic updates; the series METHODS IN ENZYMOLOGY, Academic Press, San Diego; Wolffe, CHROMATIN STRUCTURE AND FUNCTION, Third edition, Academic Press, San Diego, 1998; METHODS IN ENZYMOLOGY, Vol. 304,

“Chromatin” (P.M. Wassarman and A. P. Wolffe, eds.), Academic Press, San Diego, 1999; and METHODS IN MOLECULAR BIOLOGY, Vol. 119, “Chromatin Protocols” (P.B. Becker, ed.) Humana Press, Totowa, 1999.

#### Definitions

**[0051]** Throughout the specification and claims, unless the context requires otherwise, the word “comprise” or variations such as “comprises” or “comprising”, will be understood to imply the inclusion of a stated integer or group of integers but not the exclusion of any other integer or group of integers.

**[0051a]** The terms "nucleic acid," "polynucleotide," and "oligonucleotide" are used interchangeably and refer to a deoxyribonucleotide or ribonucleotide polymer, in linear or circular conformation, and in either single- or double-stranded form. For the purposes of the present disclosure, these terms are not to be construed as limiting with respect to the length of a polymer. The terms can encompass known analogues of natural nucleotides, as well as nucleotides that are modified in the base, sugar and/or phosphate moieties (e.g., phosphorothioate backbones). In general, an analogue of a particular nucleotide has the same base-pairing specificity; i.e., an analogue of A will base-pair with T.

**[0052]** The terms "polypeptide," "peptide" and "protein" are used interchangeably to refer to a polymer of amino acid residues. The term also applies to amino acid polymers in which one or more amino acids are chemical analogues or modified derivatives of a corresponding naturally-occurring amino acids.

**[0053]** "Recombination" refers to a process of exchange of genetic information between two polynucleotides, including but not limited to, capture by non-homologous end joining (NHEJ) and homologous recombination. For the purposes of this disclosure, "homologous recombination (HR)" refers to the specialized form of such exchange that takes place, for example, during repair of double-strand breaks in cells via homology-directed repair mechanisms.

**[0054]** In certain methods of the disclosure, one or more targeted nucleases as described herein create a double-stranded break (DSB) in the target sequence (e.g., cellular chromatin) at a predetermined site (e.g., PCSK9 gene). The DSB mediates integration of a construct as described herein. Optionally, the construct has homology to the nucleotide sequence in the region of the break. The expression construct may be physically integrated or, alternatively, the expression cassette is used as a template for repair of the break via homologous recombination, resulting in the introduction of all or part of the nucleotide sequence as in the expression cassette into the cellular chromatin. Thus, a first sequence in cellular chromatin can be altered and, in certain



embodiments, can be converted into a sequence present in an expression cassette.

Thus, the use of the terms "replace" or "replacement" can be understood to represent replacement of one nucleotide sequence by another, (*i.e.*, replacement of a sequence in the informational sense), and does not necessarily require physical or chemical

5 replacement of one polynucleotide by another.

**[0055]** In any of the methods described herein, the exogenous nucleotide sequence (the "expression construct" or "expression cassette" or "vector") can contain sequences that are homologous, but not identical, to genomic sequences in the region of interest, thereby stimulating homologous recombination to insert a non-identical

10 sequence in the region of interest. Thus, in certain embodiments, portions of the expression cassette sequence that are homologous to sequences in the region of interest exhibit between about 80 to 99% (or any integer therebetween) sequence identity to the genomic sequence that is replaced. In other embodiments, the homology between the expression cassette and genomic sequence is higher than 99%,  
15 for example if only 1 nucleotide differs as between the homology regions of the expression cassette and genomic sequences of over 100 contiguous base pairs. In certain cases, a non-homologous portion of the expression cassette can contain sequences not present in the region of interest, such that new sequences are introduced into the region of interest. In these instances, the non-homologous sequence is  
20 generally flanked by sequences of 50-1,000 base pairs (or any integral value therebetween) or any number of base pairs greater than 1,000, that are homologous or identical to sequences in the region of interest.

**[0056]** The term "sequence" refers to a nucleotide sequence of any length, which can be DNA or RNA; can be linear, circular or branched and can be either  
25 single-stranded or double stranded. The term "transgene" refers to a nucleotide sequence that is inserted into a genome. A transgene can be of any length, for example between 2 and 100,000,000 nucleotides in length (or any integer value therebetween or thereabove), preferably between about 100 and 100,000 nucleotides in length (or any integer therebetween), more preferably between about 2000 and  
30 20,000 nucleotides in length (or any value therebetween) and even more preferable, between about 5 and 15 kb (or any value therebetween).

**[0057]** A "chromosome," is a chromatin complex comprising all or a portion of the genome of a cell. The genome of a cell is often characterized by its karyotype,

which is the collection of all the chromosomes that comprise the genome of the cell.

The genome of a cell can comprise one or more chromosomes.

**[0058]** "Chromatin" is the nucleoprotein structure comprising the cellular genome. Cellular chromatin comprises nucleic acid, primarily DNA, and protein, including histones and non-histone chromosomal proteins. The majority of eukaryotic cellular chromatin exists in the form of nucleosomes, wherein a nucleosome core comprises approximately 150 base pairs of DNA associated with an octamer comprising two each of histones H2A, H2B, H3 and H4; and linker DNA (of variable length depending on the organism) extends between nucleosome cores. A molecule of histone H1 is generally associated with the linker DNA. For the purposes of the present disclosure, the term "chromatin" is meant to encompass all types of cellular nucleoprotein, both prokaryotic and eukaryotic. Cellular chromatin includes both chromosomal and episomal chromatin.

**[0059]** An "accessible region" is a site in cellular chromatin in which a target site present in the nucleic acid can be bound by an exogenous molecule which recognizes the target site. Without wishing to be bound by any particular theory, it is believed that an accessible region is one that is not packaged into a nucleosomal structure. The distinct structure of an accessible region can often be detected by its sensitivity to chemical and enzymatic probes, for example, nucleases.

**[0060]** A "target site" or "target sequence" is a nucleic acid sequence that defines a portion of a nucleic acid to which a binding molecule will bind, provided sufficient conditions for binding exist. For example, the sequence 5' GAATTC 3' is a target site for the Eco RI restriction endonuclease.

**[0061]** An "episome" is a replicating nucleic acid, nucleoprotein complex or other structure comprising a nucleic acid that is not part of the chromosomal karyotype of a cell. Examples of episomes include plasmids, minicircles and certain viral genomes. The liver specific constructs described herein may be episomally maintained or, alternatively, may be stably integrated into the cell.

**[0062]** An "exogenous" molecule is a molecule that is not normally present in a cell, but can be introduced into a cell by one or more genetic, biochemical or other methods. "Normal presence in the cell" is determined with respect to the particular developmental stage and environmental conditions of the cell. Thus, for example, a molecule that is present only during embryonic development of muscle is an exogenous molecule with respect to an adult muscle cell. Similarly, a molecule

induced by heat shock is an exogenous molecule with respect to a non-heat-shocked cell. An exogenous molecule can comprise, for example, a functioning version of a malfunctioning endogenous molecule or a malfunctioning version of a normally-functioning endogenous molecule.

5    **[0063]**       An exogenous molecule can be, among other things, a small molecule, such as is generated by a combinatorial chemistry process, or a macromolecule such as a protein, nucleic acid, carbohydrate, lipid, glycoprotein, lipoprotein, polysaccharide, any modified derivative of the above molecules, or any complex comprising one or more of the above molecules. Nucleic acids include DNA and  
10   RNA, can be single- or double-stranded; can be linear, branched or circular; and can be of any length. Nucleic acids include those capable of forming duplexes, as well as triplex-forming nucleic acids. See, for example, U.S. Patent Nos. 5,176,996 and 5,422,251. Proteins include, but are not limited to, DNA-binding proteins, transcription factors, chromatin remodeling factors, methylated DNA binding  
15   proteins, polymerases, methylases, demethylases, acetylases, deacetylases, kinases, phosphatases, ligases, deubiquitinases, integrases, recombinases, ligases, topoisomerases, gyrases and helicases.

**[0064]**       An exogenous molecule can be the same type of molecule as an endogenous molecule, *e.g.*, an exogenous protein or nucleic acid. For example, an  
20   exogenous nucleic acid can comprise an infecting viral genome, a plasmid or episome introduced into a cell, or a chromosome that is not normally present in the cell. Methods for the introduction of exogenous molecules into cells are known to those of skill in the art and include, but are not limited to, lipid-mediated transfer (*i.e.*, liposomes, including neutral and cationic lipids), electroporation, direct injection, cell  
25   fusion, particle bombardment, calcium phosphate co-precipitation, DEAE-dextran-mediated transfer and viral vector-mediated transfer. An exogenous molecule can also be the same type of molecule as an endogenous molecule but derived from a different species than the cell is derived from. For example, a human nucleic acid sequence may be introduced into a cell line originally derived from a mouse or hamster.

30   **[0065]**       By contrast, an "endogenous" molecule is one that is normally present in a particular cell at a particular developmental stage under particular environmental conditions. For example, an endogenous nucleic acid can comprise a chromosome, the genome of a mitochondrion, chloroplast or other organelle, or a naturally-

occurring episomal nucleic acid. Additional endogenous molecules can include proteins, for example, transcription factors and enzymes.

**[0066]** As used herein, the term "product of an exogenous nucleic acid" includes both polynucleotide and polypeptide products, for example, transcription products (polynucleotides such as RNA) and translation products (polypeptides).

**[0067]** A "fusion" molecule is a molecule in which two or more subunit molecules are linked, preferably covalently. The subunit molecules can be the same chemical type of molecule, or can be different chemical types of molecules.

Examples of fusion molecules include, but are not limited to, fusion proteins (for example, a fusion between a protein DNA-binding domain and a cleavage domain), fusions between a polynucleotide DNA-binding domain (*e.g.*, sgRNA) operatively associated with a cleavage domain, and fusion nucleic acids (for example, a nucleic acid encoding the fusion protein).

**[0068]** Expression of a fusion molecule in a cell can result from delivery of the fusion molecule to the cell or by delivery of a polynucleotide encoding one or more components of the fusion molecule to a cell, wherein the polynucleotide is transcribed, and the transcript is translated, to generate the fusion molecule. In the case of CRISPR/Cas transcription factors and/or nucleases, the sgRNA DNA binding domain associates with the functional domain(s) (*e.g.*, nuclease domain(s) or transcriptional regulatory domain) upon introduction of the sgRNA and functional domain encoding-sequences into the cell. Trans-splicing, polypeptide cleavage and polypeptide ligation can also be involved in expression of a protein in a cell. Methods for polynucleotide and polypeptide delivery to cells are presented elsewhere in this disclosure.

**[0069]** A "gene," for the purposes of the present disclosure, includes a DNA region encoding a gene product (see *infra*), as well as all DNA regions which regulate the production of the gene product, whether or not such regulatory sequences are adjacent to coding and/or transcribed sequences. Accordingly, a gene includes, but is not necessarily limited to, promoter sequences, terminators, translational regulatory sequences such as ribosome binding sites and internal ribosome entry sites, enhancers, silencers, insulators, boundary elements, replication origins, matrix attachment sites and locus control regions.

**[0070]** "Gene expression" refers to the conversion of the information, contained in a gene, into a gene product. A gene product can be the direct

transcriptional product of a gene (*e.g.*, mRNA, tRNA, rRNA, antisense RNA, ribozyme, structural RNA or any other type of RNA) or a protein produced by translation of an mRNA. Gene products also include RNAs which are modified, by processes such as capping, polyadenylation, methylation, and editing, and proteins modified by, for example, methylation, acetylation, phosphorylation, ubiquitination, ADP-ribosylation, myristilation, and glycosylation.

**[0071]** "Modulation" of gene expression refers to a change in the activity of a gene. Modulation of expression can include, but is not limited to, gene activation and gene repression. Genome editing (*e.g.*, cleavage, alteration, inactivation, random mutation) can be used to modulate expression. Gene inactivation refers to any reduction in gene expression as compared to a cell that does not include a ZFP, TALE or CRISPR/Cas system as described herein. Thus, gene inactivation may be partial or complete. Thus, a "gene modulator" is any molecule that modulates expression of a target gene, including but not limited to modification of the gene sequence and/or modification of gene expression.

**[0072]** A "region of interest" is any region of cellular chromatin, such as, for example, a gene or a non-coding sequence within or adjacent to a gene, in which it is desirable to bind an exogenous molecule. Binding can be for the purposes of targeted DNA cleavage and/or targeted recombination. A region of interest can be present in a chromosome, an episome, an organellar genome (*e.g.*, mitochondrial, chloroplast), or an infecting viral genome, for example. A region of interest can be within the coding region of a gene, within transcribed non-coding regions such as, for example, leader sequences, trailer sequences or introns, or within non-transcribed regions, either upstream or downstream of the coding region. A region of interest can be as small as a single nucleotide pair or up to 2,000 nucleotide pairs in length, or any integral value of nucleotide pairs.

**[0073]** A "safe harbor" locus is a locus within the genome wherein a gene may be inserted without any deleterious effects on the host cell. Most beneficial is a safe harbor locus in which expression of the inserted gene sequence is not perturbed by any read-through expression from neighboring genes. Non-limiting examples of safe harbor loci that are targeted by nuclease(s) include CCR5, HPRT, AAVS1, Rosa, ATPA1, CLYBL, and Albumin. See, *e.g.*, U.S. Patent Nos. 7,951,925; 8,771,985; 8,110,379; 7,951,925; U.S. Publication Nos. 20100218264; 20110265198;

20130137104; 20130122591; 20130177983; 20130177960; 20150056705 and 20150159172.

**[0074]** A "reporter gene" or "reporter sequence" refers to any sequence that produces a protein product that is easily measured, preferably although not necessarily in a routine assay. Suitable reporter genes include, but are not limited to, sequences encoding proteins that mediate antibiotic resistance (*e.g.*, ampicillin resistance, neomycin resistance, G418 resistance, puromycin resistance), sequences encoding colored or fluorescent or luminescent proteins (*e.g.*, green fluorescent protein, enhanced green fluorescent protein, red fluorescent protein, luciferase), and proteins which mediate enhanced cell growth and/or gene amplification (*e.g.*, dihydrofolate reductase). Epitope tags include, for example, one or more copies of FLAG, His, myc, Tap, HA or any detectable amino acid sequence. "Expression tags" include sequences that encode reporters that may be operably linked to a desired gene sequence in order to monitor expression of the gene of interest.

**[0075]** "Eukaryotic" cells include, but are not limited to, fungal cells (such as yeast), plant cells, animal cells, mammalian cells and human cells (*e.g.*, T-cells), including stem cells (pluripotent and multipotent).

**[0076]** "Binding" refers to a sequence-specific, non-covalent interaction between macromolecules (*e.g.*, between a protein and a nucleic acid). Not all components of a binding interaction need be sequence-specific (*e.g.*, contacts with phosphate residues in a DNA backbone), as long as the interaction as a whole is sequence-specific. Such interactions are generally characterized by a dissociation constant ( $K_d$ ) of  $10^{-6}$  M<sup>-1</sup> or lower. "Affinity" refers to the strength of binding: increased binding affinity being correlated with a lower  $K_d$ . "Non-specific binding" refers to generalized, non-covalent interactions that occur between any molecule of interest (*e.g.* an engineered nuclease) and a macromolecule (*e.g.* DNA) that can occur at any generalized location in one molecule wherein such interaction is not limited to a target sequence.

**[0077]** A "binding protein" is a protein that is able to bind non-covalently to another molecule. A binding protein can bind to, for example, a DNA molecule (a DNA-binding protein), an RNA molecule (an RNA-binding protein) and/or a protein molecule (a protein-binding protein). In the case of a protein-binding protein, it can bind to itself (to form homodimers, homotrimers, *etc.*) and/or it can bind to one or more molecules of a

different protein or proteins. A binding protein can have more than one type of binding activity. For example, zinc finger proteins have DNA-binding, RNA-binding and protein-binding activity. In the case of an RNA-guided nuclease system, the RNA guide is heterologous to the nuclease component (Cas9 or Cfp1) and both may be engineered.

5   **[0078]**           A “DNA binding molecule” is a molecule that can binding to DNA. Such DNA binding molecule can be a polypeptide, a domain of a protein, a domain within a larger protein or a polynucleotide. In some embodiments, the polynucleotide is DNA, while in other embodiments, the polynucleotide is RNA. In some embodiments, the DNA binding molecule is a protein domain of a nuclease (*e.g.* the *FokI* domain), while in other  
10   embodiments, the DNA binding molecule is a guide RNA component of an RNA-guided nuclease (*e.g.* Cas9 or Cfp1).

**[0079]**           A “DNA binding protein” (or binding domain) is a protein, or a domain within a larger protein, that binds DNA in a sequence-specific manner through one or more zinc fingers or through interaction with one or more RVDs in a zinc finger protein or  
15   TALE, respectively. The term zinc finger DNA binding protein is often abbreviated as zinc finger protein or ZFP.

**[0080]**           A “zinc finger DNA binding protein” (or binding domain) is a protein, or a domain within a larger protein, that binds DNA in a sequence-specific manner through one or more zinc fingers, which are regions of amino acid sequence within the binding domain  
20   whose structure is stabilized through coordination of a zinc ion. The term zinc finger DNA binding protein is often abbreviated as zinc finger protein or ZFP.

**[0081]**           A “TALE DNA binding domain” or “TALE” is a polypeptide comprising one or more TALE repeat domains/units. The repeat domains are involved in binding of the TALE to its cognate target DNA sequence. A single “repeat unit” (also referred to as a  
25   “repeat”) is typically 33-35 amino acids in length and exhibits at least some sequence homology with other TALE repeat sequences within a naturally occurring TALE protein. *See, e.g.*, U.S. Patent No. 8,586,526 incorporated by reference herein in its entirety.

**[0082]**           Zinc finger and TALE binding domains can be “engineered” to bind to a predetermined nucleotide sequence, for example via engineering (altering one or  
30   more amino acids) of the recognition helix region of a naturally occurring zinc finger protein or by engineering of the amino acids involved in DNA binding (the “repeat variable diresidue” or RVD region) . Therefore, engineered DNA binding proteins (zinc fingers or TALEs) are proteins that are non-naturally occurring. Non-limiting examples of methods for engineering DNA-binding proteins are design and selection.

A designed DNA binding protein is a protein not occurring in nature whose design/composition results principally from rational criteria. Rational criteria for design include application of substitution rules and computerized algorithms for processing information in a database storing information of existing ZFP and/or

5 TALE designs and binding data. See, for example, U.S. Patents 8,586,526; 6,140,081; 6,453,242; 6,746,838; 7,241,573; 6,866,997; 7,241,574 and 6,534,261; see also WO 98/53058; WO 98/53059; WO 98/53060; WO 02/016536 and WO 03/016496.

**[0083]** A "selected" zinc finger protein, TALE or CRISPR/Cas system is not found  
10 in nature whose production results primarily from an empirical process such as phage display, interaction trap, rational design or hybrid selection. See *e.g.*, U.S. Patent Nos. 8,586,526; 5,789,538; 5,925,523; 6,007,988; 6,013,453; 6,200,759; 6,242,568; 6,733,970; 7,297,491; WO 98/53057; WO 02/099084.

**[0084]** A "CRISPR/Cas" system refers to a nuclease or transcription factors system  
15 comprising a single guide RNA (sgRNA) DNA-binding domain and one or more cleavage domains (for nucleases) or transcriptional regulator domains (for transcription factors). The sgRNA can be designed to bind to any DNA sequence. The sgRNA, when associated with the cleavage or transcriptional regulatory domain(s) then mediates modulation of gene expression. See, *e.g.*, U.S. Patent Nos. 9,267,135 and 8,841,260 and U.S. Patent  
20 Publication Nos. 20150056705 and 20150031134.

**[0085]** "TtAgo" is a prokaryotic Argonaute protein thought to be involved in gene silencing. TtAgo is derived from the bacteria *Thermus thermophilus*. See, *e.g.*, Swarts *et al.* (2014) *Nature* 507(7491):258-61; G. Sheng *et al.*, (2013) *Proc. Natl. Acad. Sci. U.S.A.* 111, 652). A "TtAgo system" is all the components required  
25 including, for example, guide DNAs for cleavage by a TtAgo enzyme.

"Recombination" refers to a process of exchange of genetic information between two polynucleotides, including but not limited to, donor capture by non-homologous end joining (NHEJ) and homologous recombination. For the purposes of this disclosure, "homologous recombination (HR)" refers to the specialized form of such exchange  
30 that takes place, for example, during repair of double-strand breaks in cells via homology-directed repair mechanisms. This process requires nucleotide sequence homology, uses a "donor" molecule to template repair of a "target" molecule (*i.e.*, the one that experienced the double-strand break), and is variously known as "non-crossover gene conversion" or "short tract gene conversion," because it leads to the



transfer of genetic information from the donor to the target. Without wishing to be bound by any particular theory, such transfer can involve mismatch correction of heteroduplex DNA that forms between the broken target and the donor, and/or "synthesis-dependent strand annealing," in which the donor is used to resynthesize genetic information that will become part of the target, and/or related processes. Such specialized HR often results in an alteration of the sequence of the target molecule such that part or all of the sequence of the donor polynucleotide is incorporated into the target polynucleotide.

**[0086]** "Recombination" refers to a process of exchange of genetic information between two polynucleotides, including but not limited to, capture by non-homologous end joining (NHEJ) and homologous recombination. For the purposes of this disclosure, "homologous recombination (HR)" refers to the specialized form of such exchange that takes place, for example, during repair of double-strand breaks in cells via homology-directed repair mechanisms. This process requires nucleotide sequence homology, uses a "donor" molecule to template repair of a "target" molecule (i.e., the one that experienced the double-strand break), and is variously known as "non-crossover gene conversion" or "short tract gene conversion," because it leads to the transfer of genetic information from the donor to the target. Without wishing to be bound by any particular theory, such transfer can involve mismatch correction of heteroduplex DNA that forms between the broken target and the donor, and/or "synthesis-dependent strand annealing," in which the donor is used to resynthesize genetic information that will become part of the target, and/or related processes. Such specialized HR often results in an alteration of the sequence of the target molecule such that part or all of the sequence of the donor polynucleotide is incorporated into the target polynucleotide.

**[0087]** In certain methods of the disclosure, one or more targeted nucleases as described herein create a double-stranded break (DSB) in the target sequence (e.g., cellular chromatin) at a predetermined site (e.g., a gene or locus of interest). The DSB mediates integration of a construct (e.g. donor) as described herein. Optionally, the construct has homology to the nucleotide sequence in the region of the break. An expression construct may be physically integrated or, alternatively, the expression cassette is used as a template for repair of the break via homologous recombination, resulting in the introduction of all or part of the nucleotide sequence as in the expression cassette into the cellular chromatin. Thus, a first sequence in cellular

chromatin can be altered and, in certain embodiments, can be converted into a sequence present in an expression cassette. Thus, the use of the terms “replace” or “replacement” can be understood to represent replacement of one nucleotide sequence by another, (i.e., replacement of a sequence in the informational sense), and does not necessarily require physical or chemical replacement of one polynucleotide by another.

**[0088]** In any of the methods described herein, additional engineered nucleases can be used for additional double-stranded cleavage of additional target sites within the cell.

**[0089]** In certain embodiments of methods for targeted recombination and/or replacement and/or alteration of a sequence in a region of interest in cellular chromatin, a chromosomal sequence is altered by homologous recombination with an exogenous “donor” nucleotide sequence. Such homologous recombination is stimulated by the presence of a double-stranded break in cellular chromatin, if sequences homologous to the region of the break are present.

**[0090]** In any of the methods described herein, the first nucleotide sequence (the “donor sequence”) can contain sequences that are homologous, but not identical, to genomic sequences in the region of interest, thereby stimulating homologous recombination to insert a non-identical sequence in the region of interest. Thus, in certain embodiments, portions of the donor sequence that are homologous to sequences in the region of interest exhibit between about 80 to 99% (or any integer therebetween) sequence identity to the genomic sequence that is replaced. In other embodiments, the homology between the donor and genomic sequence is higher than 99%, for example if only 1 nucleotide differs as between donor and genomic sequences of over 100 contiguous base pairs. In certain cases, a non-homologous portion of the donor sequence can contain sequences not present in the region of interest, such that new sequences are introduced into the region of interest. In these instances, the non-homologous sequence is generally flanked by sequences of 50-1,000 base pairs (or any integral value therebetween) or any number of base pairs greater than 1,000, that are homologous or identical to sequences in the region of interest. In other embodiments, the donor sequence is non-homologous to the first sequence, and is inserted into the genome by non-homologous recombination mechanisms.

**[0091]** Any of the methods described herein can be used for partial or complete inactivation of one or more target sequences in a cell by targeted integration of donor sequence or via cleavage of the target sequence(s) followed by error-prone NHEJ-mediated repair that disrupts expression of the gene(s) of interest. Cell lines with partially or completely inactivated genes are also provided.

**[0092]** Furthermore, the methods of targeted integration as described herein can also be used to integrate one or more exogenous sequences. The exogenous nucleic acid sequence can comprise, for example, one or more genes or cDNA molecules, or any type of coding or noncoding sequence, as well as one or more control elements (e.g., promoters). In addition, the exogenous nucleic acid sequence may produce one or more RNA molecules (e.g., small hairpin RNAs (shRNAs), inhibitory RNAs (RNAis), microRNAs (miRNAs), etc.).

**[0093]** "Cleavage" refers to the breakage of the covalent backbone of a DNA molecule. Cleavage can be initiated by a variety of methods including, but not limited to, enzymatic or chemical hydrolysis of a phosphodiester bond. Both single-stranded cleavage and double-stranded cleavage are possible, and double-stranded cleavage can occur as a result of two distinct single-stranded cleavage events. DNA cleavage can result in the production of either blunt ends or staggered ends. In certain embodiments, fusion polypeptides are used for targeted double-stranded DNA cleavage.

**[0094]** A "cleavage half-domain" is a polypeptide sequence which, in conjunction with a second polypeptide (either identical or different) forms a complex having cleavage activity (preferably double-strand cleavage activity). The terms "first and second cleavage half-domains," "+ and – cleavage half-domains" and "right and left cleavage half-domains" are used interchangeably to refer to pairs of cleavage half-domains that dimerize.

**[0095]** An "engineered cleavage half-domain" is a cleavage half-domain that has been modified so as to form obligate heterodimers with another cleavage half-domain (e.g., another engineered cleavage half-domain).

**[0096]** The terms "operative linkage" and "operatively linked" (or "operably linked") are used interchangeably with reference to a juxtaposition of two or more components (such as sequence elements), in which the components are arranged such that both components function normally and allow the possibility that at least one of the components can mediate a function that is exerted upon at least one of the other

components. By way of illustration, a transcriptional regulatory sequence, such as a promoter, is operatively linked to a coding sequence if the transcriptional regulatory sequence controls the level of transcription of the coding sequence in response to the presence or absence of one or more transcriptional regulatory factors. A

5 transcriptional regulatory sequence is generally operatively linked in *cis* with a coding sequence, but need not be directly adjacent to it. For example, an enhancer is a transcriptional regulatory sequence that is operatively linked to a coding sequence, even though they are not contiguous.

[0097] A "functional fragment" of a protein, polypeptide or nucleic acid is a  
10 protein, polypeptide or nucleic acid whose sequence is not identical to the full-length protein, polypeptide or nucleic acid, yet retains the same function as the full-length protein, polypeptide or nucleic acid. A functional fragment can possess more, fewer, or the same number of residues as the corresponding native molecule, and/or can contain one or more amino acid or nucleotide substitutions. Methods for determining  
15 the function of a nucleic acid (*e.g.*, coding function, ability to hybridize to another nucleic acid) are well-known in the art. Similarly, methods for determining protein function are well-known.

[0098] A polynucleotide "vector" or "construct" is capable of transferring gene sequences to target cells. Typically, "vector construct," "expression vector,"  
20 "expression construct," "expression cassette," and "gene transfer vector," mean any nucleic acid construct capable of directing the expression of a gene of interest and which can transfer gene sequences to target cells. Thus, the term includes cloning, and expression vehicles, as well as integrating vectors.

[0099] The terms "subject" and "patient" are used interchangeably and refer to  
25 mammals such as human patients and non-human primates, as well as experimental animals such as rabbits, dogs, cats, rats, mice, and other animals. Accordingly, the term "subject" or "patient" as used herein means any mammalian patient or subject to which the expression cassettes of the invention can be administered. Subjects of the present invention include those with a disorder.

30 [0100] The terms "treating" and "treatment" as used herein refer to reduction in severity and/or frequency of symptoms, elimination of symptoms and/or underlying cause, prevention of the occurrence of symptoms and/or their underlying cause, and improvement or remediation of damage. Cancer, monogenic diseases and graft versus

host disease are non-limiting examples of conditions that may be treated using the compositions and methods described herein.

### **Liver-specific expression constructs**

- 5   **[0101]**        Described herein are expression cassettes (constructs) for use in directing expression of a transgene (*e.g.*, one or more modulators of a PCSK9, a TTR, a SERPINA1, a KLKB1 and/or a HAO1 gene) in a liver cell, including *in vivo* following administration of the expression cassette(s) to the subject (*e.g.*, peripheral or hepatic vein delivery). Also described here are expression cassettes (constructs) comprising a wildtype cDNA copy of a PCSK9, a TTR, a SERPINA1, a KLKB1 and/or a HAO1 gene for use in a cell where an endogenous mutant version of PCSK9, a TTR, a SERPINA1, a KLKB1 and/or a HAO1 gene has been knocked out by one or more modulators as described above. The expression construct may be maintained episomally and drive expression of the transgene extrachromosomally (see U.S. Publication No. 20170119906) or, alternatively, the expression construct may be integrated into the genome of a liver cell, for example by nuclease-mediated targeted integration.

- 15   **[0102]**        The polynucleotide expression construct comprises an enhancer sequence, a promoter sequence, and one or more transgenes. Optionally included are one or more of the following: an intronic sequence, a polyadenylation sequence and/or a signal peptide. Any enhancer sequence may be used in the expression constructs described herein. In certain embodiments, the enhancer is a wild-type or modified Serpin1 enhancer (Chuah et al., (2014) *Molecular Therapy*, 22, 1605-1613; Nair et al. (2014) *Blood* 123, 3195-3199).

- 20   **[0103]**        In preferred embodiments, the Serpin1 enhancer comprises one or more mutations (*e.g.*, point mutations) as compared to wild-type, for example a Serpin1 enhancer containing one or more nucleotide modifications as shown, namely modifications of nucleotides at one or more of locations within the enhancer. *See, e.g.*, U.S. Publication No. US-2017-0119906-A1.

- 25   **[0104]**        Similarly, any promoter sequence can be used in the expression cassettes of the invention. In certain embodiments, the promoter is a constitutive promoter. In other embodiments, the promoter is an inducible or tissue specific promoter. In some embodiments, the promoter is a liver-specific promoter. In certain

embodiments, the promoter is a transthyretin minimal promoter (TTRm) promoter. In other embodiments, the promoter is an alpha-1 anti-trypsin (hAAT) promoter.

**[0105]** Any of the polynucleotides described herein may further optionally comprise an intronic sequence. In certain embodiments, the expression construct includes a truncated chimeric intron (T-chimeric intron) sequence. The T-chimeric intron is a truncated version of the chimeric intron in pCI-neo (GenBank U47120). The chimeric intron in pCI-neo is the 5' splice donor site from the human  $\beta$ -globin gene, and the branchpoint and 3' acceptor site of an immunoglobulin gene heavy chain variable region. The T-chimeric intron contains a 45 bp deletion between the 5' splice donor and the branchpoint. In yet other embodiments, the expression constructs include a mutated MVM intron sequence (*e.g.*, as shown in U.S. Publication No. US-2017-0119906-A1).

**[0106]** Alternatively, the expression constructs as described herein may lack an intronic sequence, for example as shown in U.S. Publication No. US-2017-0119906-A1.

**[0107]** The constructs described herein may be contained within any viral or non-viral vector. The constructs may be maintained episomally or may be integrated into the genome of the cell (*e.g.*, via nuclease-mediated targeted integration).

**[0108]** Non-viral vectors include DNA or RNA plasmids, DNA MCs, naked nucleic acid, and nucleic acid complexed with a delivery vehicle such as a liposome, nanoparticle or poloxamer. Viral vectors that may be used to carry the expression cassettes described herein include, but are not limited to, retroviral, lentivirus, adenoviral, adeno-associated viral vectors, vaccinia and herpes simplex virus vectors. Integration in the host genome is possible with the retrovirus, lentivirus, and adeno-associated virus gene transfer methods, and as described herein may be facilitated by nuclease-mediated integration.

**[0109]** In certain preferred embodiments, the constructs are included in an adeno-associated virus ("AAV") vector or vector system that may be maintained episomally or integrated into the genome of a liver cell (*e.g.*, via nuclease-mediated targeted integration). Construction of recombinant AAV vectors is in a number of publications, including U.S. Pat. No. 5,173,414; Tratschin *et al.*, *Mol. Cell. Biol.* 5:3251-3260 (1985); Tratschin, *et al.*, *Mol. Cell. Biol.* 4:2072-2081 (1984); Hermonat

& Muzyczka, *PNAS* 81:6466-6470 (1984); and Samulski *et al.*, *J. Virol.* 63:03822-3828 (1989).

**[0110]** Thus, in certain embodiments, the expression construct is carried on an AAV construct and further comprises 5' and 3' ITRs flanking the expression constructs elements (*e.g.*, enhancer, promoter, optional intron, transgene, etc.) as described herein. Optionally, spacer molecules are also included between one or more of the components of the expression construct, for example, between the 5' ITR and the enhancer and/or between the polyadenylation signal and the 3' ITR. The spacers may function as homology arms to facilitate recombination into a safe-harbor locus (*e.g.* albumin). In certain embodiments, the construct is a construct as shown in U.S. Publication No. US-2017-0119906-A1.

**[0111]** In certain embodiments, the AAV vectors as described herein can be derived from any AAV. In certain embodiments, the AAV vector is derived from the defective and nonpathogenic parvovirus adeno-associated type 2 virus. All such vectors are derived from a plasmid that retains only the AAV 145 bp inverted terminal repeats flanking the transgene expression cassette. Efficient gene transfer and stable transgene delivery due to integration into the genomes of the transduced cell are key features for this vector system. (Wagner *et al.*, *Lancet* 351:9117 1702-3 (1998), Kearns *et al.*, *Gene Ther.* 9:748-55 (1996)). Other AAV serotypes, including AAV1, AAV2, AAV3, AAV4, AAV5, AAV6, AAV7, AAV8, AAV9 and AAVrh.10 and any novel AAV serotype can also be used in accordance with the present invention. In some embodiments, chimeric AAV is used where the viral origins of the LTR sequences of the viral nucleic acid are heterologous to the viral origin of the capsid sequences. Non-limiting examples include chimeric virus with LTRs derived from AAV2 and capsids derived from AAV5, AAV6, AAV8 or AAV9 (*i.e.* AAV2/5, AAV2/6, AAV2/8 and AAV2/9, respectively).

**[0112]** Retroviral vectors include those based upon murine leukemia virus (MuLV), gibbon ape leukemia virus (GaLV), Simian Immunodeficiency virus (SIV), human immunodeficiency virus (HIV), and combinations thereof (*see, e.g.*, Buchscher *et al.*, *J. Virol.* 66:2731-2739 (1992); Johann *et al.*, *J. Virol.* 66:1635-1640 (1992); Sommerfelt *et al.*, *Virol.* 176:58-59 (1990); Wilson *et al.*, *J. Virol.* 63:2374-2378 (1989); Miller *et al.*, *J. Virol.* 65:2220-2224 (1991); PCT/US94/05700).

**[0113]** The constructs described herein may also be incorporated into an adenoviral vector system. Adenoviral based vectors are capable of very high

transduction efficiency in many cell types and do not require cell division. With such vectors, high titer and high levels of expression have been obtained. This vector can be produced in large quantities in a relatively simple system.

[0114] pLASN and MFG-S are examples of retroviral vectors that have been used in clinical trials (Dunbar *et al.*, *Blood* 85:3048-305 (1995); Kohn *et al.*, *Nat. Med.* 1:1017-102 (1995); Malech *et al.*, *PNAS* 94:22 12133-12138 (1997)).

PA317/pLASN was the first therapeutic vector used in a gene therapy trial. (Blaese *et al.*, *Science* 270:475-480 (1995)). Transduction efficiencies of 50% or greater have been observed for MFG-S packaged vectors. (Ellem *et al.*, *Immunol Immunother.* 44(1):10-20 (1997); Dranoff *et al.*, *Hum. Gene Ther.* 1:111-2 (1997)).

[0115] Replication-deficient recombinant adenoviral vectors (Ad) can also be used with the polynucleotides described herein. Most adenovirus vectors are engineered such that a transgene replaces the Ad E1a, E1b, and/or E3 genes; subsequently the replication defective vector is propagated in human 293 cells that supply deleted gene function in *trans*. Ad vectors can transduce multiple types of tissues *in vivo*, including nondividing, differentiated cells such as those found in liver, kidney and muscle. Conventional Ad vectors have a large carrying capacity. An example of the use of an Ad vector in a clinical trial involved polynucleotide therapy for antitumor immunization with intramuscular injection (Stermann *et al.*, *Hum. Gene Ther.* 7:1083-9 (1998)). Additional examples of the use of adenovirus vectors for gene transfer in clinical trials include Rosenecker *et al.*, *Infection* 24:1 5-10 (1996); Stermann *et al.*, *Hum. Gene Ther.* 9:7 1083-1089 (1998); Welsh *et al.*, *Hum. Gene Ther.* 2:205-18 (1995); Alvarez *et al.*, *Hum. Gene Ther.* 5:597-613 (1997); Topf *et al.*, *Gene Ther.* 5:507-513 (1998); Stermann *et al.*, *Hum. Gene Ther.* 7:1083-1089 (1998).

[0116] Packaging cells are used to form virus particles that are capable of infecting a host cell. Such cells include 293 cells, which package AAV and adenovirus, and  $\psi$ 2 cells or PA317 cells, which package retrovirus. Viral vectors used in gene therapy are usually generated by a producer cell line that packages a nucleic acid vector into a viral particle. The vectors typically contain the minimal viral sequences required for packaging and subsequent integration into a host (if applicable), other viral sequences being replaced by an expression cassette encoding the protein to be expressed. The missing viral functions are supplied in *trans* by the packaging cell line. For example, AAV vectors used in gene therapy typically only possess inverted terminal repeat (ITR) sequences from the AAV genome which are



required for packaging and integration into the host genome. Viral DNA is packaged in a cell line, which contains a helper plasmid encoding the other AAV genes, namely *rep* and *cap*, but lacking ITR sequences. The cell line is also infected with adenovirus as a helper. The helper virus promotes replication of the AAV vector and expression of AAV genes from the helper plasmid. The helper plasmid is not packaged in significant amounts due to a lack of ITR sequences. Contamination with adenovirus can be reduced by, *e.g.*, heat treatment to which adenovirus is more sensitive than AAV. In some embodiments, AAV is produced using a baculovirus expression system.

[0117] In many gene therapy applications, it is desirable that the gene therapy vector be delivered with a high degree of specificity to a particular tissue type. Accordingly, a viral vector can be modified to have specificity for a given cell type by expressing a ligand as a fusion protein with a viral coat protein on the outer surface of the virus. The ligand is chosen to have affinity for a receptor known to be present on the cell type of interest. For example, Han *et al.*, *Proc. Natl. Acad. Sci. USA* 92:9747-9751 (1995), reported that Moloney murine leukemia virus can be modified to express human heregulin fused to gp70, and the recombinant virus infects certain human breast cancer cells expressing human epidermal growth factor receptor. This principle can be extended to other virus-target cell pairs, in which the target cell expresses a receptor and the virus expresses a fusion protein comprising a ligand for the cell-surface receptor. For example, filamentous phage can be engineered to display antibody fragments (*e.g.*, FAB or Fv) having specific binding affinity for virtually any chosen cellular receptor. Although the above description applies primarily to viral vectors, the same principles can be applied to nonviral vectors. Such vectors can be engineered to contain specific uptake sequences which favor uptake by specific target cells.

[0118] The polynucleotides described herein may include one or more non-natural bases and/or backbones. In particular, an expression cassette as described herein may include methylated cytosines to achieve a state of transcriptional quiescence in a region of interest.

[0119] Furthermore, the expression constructs as described herein may also include additional transcriptional or translational regulatory or other sequences, for example, additional promoters, enhancers, insulators, internal ribosome entry sites, sequences encoding 2A peptides and/or polyadenylation signals. Further, the control

elements of the genes of interest can be operably linked to reporter genes to create chimeric genes (*e.g.*, reporter expression cassettes).

## 5 Transgenes

[0120] The constructs described herein can be used for hepatic delivery of any transgene.

[0121] In certain embodiments, the transgene encodes a sequence comprising one or more modulators of a PCSK9, a TTR, a SERPINA1, a KLKB1 and/or a HAO1 gene, including but not limited to, one or more transcription factors that up-regulate or  
10 down-regulate expression of one or more these gene, one or more nucleases that cleave one or more of these genes, and/or one or more sequences that inhibit or activate one or more of these genes. Exemplary artificial transcription factors and/or nucleases that bind to target sites in one or more of these genes are described herein.

[0122] In other embodiments, the transgene encodes a cDNA sequence of a PCSK9, a TTR, a SERPINA1, a KLKB1 and/or a HAO1 gene. These transgenes may be used in cells in which the endogenous mutant copy of a PCSK9, a TTR, a  
15 SERPINA1, a KLKB1 and/or a HAO1 gene has been knocked out by a nuclease as described herein. The cDNA sequences may be integrated in the genome or may be  
20 maintained extrachromosomally (*e.g.* episomally).

[0123] The transgenes may also include promoter sequences, enhancer sequences, epitope tags, marker genes, cleavage enzyme recognition sites and various types of expression constructs. Marker genes that may be provided include, but are not limited to, sequences encoding proteins that mediate antibiotic resistance (*e.g.*,  
25 ampicillin resistance, neomycin resistance, G418 resistance, puromycin resistance), sequences encoding colored or fluorescent or luminescent proteins (*e.g.*, green fluorescent protein, enhanced green fluorescent protein, red fluorescent protein, luciferase), and proteins which mediate enhanced cell growth and/or gene amplification (*e.g.*, dihydrofolate reductase). Epitope tags include, for example, one  
30 or more copies of FLAG, His, myc, Tap, HA or any detectable amino acid sequence.

[0124] In a preferred embodiment, the transgene comprises a polynucleotide encoding any polypeptide of which expression in the cell is desired, including, but not limited to antibodies, antigens, enzymes, receptors (cell surface or nuclear), hormones, lymphokines, cytokines, reporter polypeptides, growth factors, and

functional fragments of any of the above. The coding sequences may be, for example, cDNAs.

**[0125]** In other embodiments, the transgene(s) encodes functional versions of proteins lacking or deficient in any genetic disease, including but not limited to, lysosomal storage disorders (*e.g.*, Gaucher's, Fabry's, Hunter's, Hurler's, Neimann-Pick's, etc.), metabolic disorders, and/or blood disorders such as hemophilias and hemoglobinopathies, etc. *See, e.g.*, U.S. Publication No. 20140017212 and 20140093913; U.S. Patent Nos. 9,255,250 and 9,175,280.

**[0126]** For example, the transgene may comprise a sequence encoding a polypeptide that is lacking or non-functional in the subject having a genetic disease, including but not limited to any of the following genetic diseases: achondroplasia, achromatopsia, acid maltase deficiency, adenosine deaminase deficiency (OMIM No.102700), adrenoleukodystrophy, aicardi syndrome, alpha-1 antitrypsin deficiency, alpha-thalassemia, androgen insensitivity syndrome, apert syndrome, arrhythmogenic right ventricular, dysplasia, ataxia telangiectasia, Barth syndrome, beta-thalassemia, blue rubber bleb nevus syndrome, Canavan disease, chronic granulomatous diseases (CGD), cri du chat syndrome, cystic fibrosis, Dercum's disease, ectodermal dysplasia, Fanconi anemia, fibrodysplasia ossificans progressive, fragile X syndrome, galactosemia, Gaucher's disease, generalized gangliosidosis (*e.g.*, GM1), hemochromatosis, the hemoglobin C mutation in the 6<sup>th</sup> codon of beta-globin (HbC), hemophilia, Huntington's disease, Hurler Syndrome, hypophosphatasia, Klinefelter syndrome, Krabbe Disease, Langer-Giedion Syndrome, leukocyte adhesion deficiency (LAD, OMIM No. 116920), leukodystrophy, long QT syndrome, Marfan syndrome, Moebius syndrome, mucopolysaccharidosis (MPS), nail patella syndrome, nephrogenic diabetes insipidus, neurofibromatosis, Neimann-Pick disease, osteogenesis imperfecta, porphyria, Prader-Willi syndrome, progeria, Proteus syndrome, retinoblastoma, Rett syndrome, Rubinstein-Taybi syndrome, Sanfilippo syndrome, severe combined immunodeficiency (SCID), Shwachman syndrome, sickle cell disease (sickle cell anemia), Smith-Magenis syndrome, Stickler syndrome, Tay-Sachs disease, Thrombocytopenia Absent Radius (TAR) syndrome, Treacher Collins syndrome, trisomy, tuberous sclerosis, Turner's syndrome, urea cycle disorder, von Hippel-Landau disease, Waardenburg syndrome, Williams syndrome, Wilson's disease, Wiskott-Aldrich syndrome, X-linked lymphoproliferative syndrome (XLP, OMIM No. 308240), acquired immunodeficiencies, lysosomal storage diseases (*e.g.*,

Gaucher's disease, GM1, Fabry disease and Tay-Sachs disease), mucopolysaccharidosis (*e.g.* Hunter's disease, Hurler's disease), hemoglobinopathies (*e.g.*, sickle cell diseases, HbC,  $\alpha$ -thalassemia,  $\beta$ -thalassemia) and hemophilias.

[0127] Non-limiting examples of proteins (including functional fragments thereof such as truncated versions) that may be expressed as described herein include fibrinogen, prothrombin, tissue factor, Factor V, Factor VII, Factor IX, Factor X, Factor XI, Factor XII (Hageman factor), Factor XIII (fibrin-stabilizing factor), von Willebrand factor, prekallikrein, high molecular weight kininogen (Fitzgerald factor), fibronectin, antithrombin III, heparin cofactor II, protein C, protein S, protein Z, protein Z-related protease inhibitor, plasminogen, alpha 2-antiplasmin, tissue plasminogen activator, urokinase, plasminogen activator inhibitor-1, plasminogen activator inhibitor-2, glucocerebrosidase (GBA),  $\alpha$ -galactosidase A (GLA), iduronate sulfatase (IDS), iduronidase (IDUA), acid sphingomyelinase (SMPD1), MMAA, MMAB, MMACHC, MMADHC (C2orf25), MTRR, LMBRD1, MTR, propionyl-CoA carboxylase (PCC) (PCCA and/or PCCB subunits), a glucose-6-phosphate transporter (G6PT) protein or glucose-6-phosphatase (G6Pase), an LDL receptor (LDLR), ApoB, LDLRAP-1, a PCSK9, a mitochondrial protein such as NAGS (N-acetylglutamate synthetase), CPS1 (carbamoyl phosphate synthetase I), and OTC (ornithine transcarbamylase), ASS (argininosuccinic acid synthetase), ASL (argininosuccinase acid lyase) and/or ARG1 (arginase), and/or a solute carrier family 25 (SLC25A13, an aspartate/glutamate carrier) protein, a UGT1A1 or UDP glucuronosyltransferase polypeptide A1, a fumarylacetoacetate hydrolyase (FAH), an alanine-glyoxylate aminotransferase (AGXT) protein, a glyoxylate reductase/hydroxypyruvate reductase (GRHPR) protein, a transthyretin gene (TTR) protein, an ATP7B protein, a phenylalanine hydroxylase (PAH) protein, a lipoprotein lyase (LPL) protein and/or a therapeutic single chain antibody.

[0128] In certain embodiments, the transgene can comprise a marker gene (described above), allowing selection of cells that have undergone targeted integration, and a linked sequence encoding an additional functionality. Non-limiting examples of marker genes include GFP, drug selection marker(s) and the like.

[0129] The constructs described herein may also be used for delivery of non-coding transgenes. Sequences encoding antisense RNAs, RNAi, shRNAs and micro RNAs (miRNAs) may also be used for targeted insertions.

**[0130]** In certain embodiments, the transgene includes sequences (*e.g.*, coding sequences, also referred to as transgenes) greater than 1 kb in length, for example between 2 and 200 kb, between 2 and 10 kb (or any value therebetween). The transgene may also include one or more nuclease target sites.

- 5 **[0131]** When integrated (*e.g.*, via nuclease-mediate integration), the transgene may be inserted into an endogenous gene such that all, some or none of the endogenous gene is expressed.

### **Nucleases/Transcription Factors**

- 10 **[0132]** In certain embodiments, the gene modulators described herein comprise one or more nucleases that inactivate a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene. Furthermore, nucleases may also be used to integrated one or more transgenes into a target gene, for instance into a PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene and/or a safe harbor gene (*e.g.*, albumin). Preferably,
- 15 integration of the transgene construct(s) is targeted following cleavage of the target gene by one or more nucleases (*e.g.*, zinc finger nucleases (“ZFNs”), TALENs, TtAgo, CRISPR/Cas nuclease systems, and homing endonucleases) and the construct integrated by either homology directed repair (HDR) or by end capture during non-homologous end joining (NHEJ) driven processes. *See, e.g.*, U.S. Patent Nos.
- 20 9,255,250; 9,200,266; 9,045,763; 9,005,973; 9,150,847; 8,956,828; 8,945,868; 8,703,489; 8,586,526; 6,534,261; 6,599,692; 6,503,717; 6,689,558; 7,067,317; 7,262,054; 7,888,121; 7,972,854; 7,914,796; 7,951,925; 8,110,379; 8,409,861; U.S. Patent Publications 20030232410; 20050208489; 20050026157; 20050064474; 20060063231; 20080159996; 201000218264; 20120017290; 20110265198;
- 25 20130137104; 20130122591; 20130177983; 20130196373 and 20150056705, the disclosures of which are incorporated by reference in their entireties for all purposes.

**[0133]** Any nuclease can be used for targeted integration of a transgene (*e.g.*, expression construct).

- [0134]** In certain embodiments, the nuclease comprises a zinc finger nuclease (ZFN), which comprises a zinc finger DNA-binding domain and a cleavage (nuclease) domain. *See, e.g.*, U.S. Patent Nos. 9,255,250; 9,200,266; 9,045,763; 9,005,973; 9,150,847; 8,956,828; 8,945,868; 8,703,489; 8,586,526; 6,534,261; 6,599,692; 6,503,717; 6,689,558; 7,067,317; 7,262,054; 7,888,121; 7,972,854; 7,914,796; 7,951,925; 8,110,379; 8,409,861.
- 30

[0135] In other embodiments, the nuclease comprises a TALEN, which comprises a TAL-effector DNA binding domain and a cleavage (nuclease) domain. *See, e.g.*, U.S. Patent No. 8,586,526 and U.S. Publication No. 20130196373.

5 [0136] In still further embodiments, the nuclease comprises a CRISPR/Cas nuclease system, which includes a single guide RNA for recognition of the target site and one or more cleavage domains. *See, e.g.*, U.S. Patent Publication No. 20150056705.

10 [0137] The cleavage domains of the nucleases may be wild-type or mutated, including non-naturally occurring (engineered) cleavage domains that form obligate heterodimers. *See, e.g.*, U.S. Patent Nos. 8,623,618; 7,888,121; 7,914,796; and 8,034,598 and U.S. Publication No. 20110201055.

[0138] The nuclease(s) may make one or more double-stranded and/or single-stranded cuts in the target site. The sites of cleavage and/or modification may be within the target site and/or between the two target sites when using a nuclease that  
15 functions as a dimer. In certain embodiments, the nuclease comprises a catalytically inactive cleavage domain (*e.g.*, FokI and/or Cas protein). *See, e.g.*, U.S. Patent No. 9,200,266; 8,703,489 and Guillinger *et al.* (2014) *Nature Biotech.* 32(6):577-582. The catalytically inactive cleavage domain may, in combination with a catalytically active domain act as a nickase to make a single-stranded cut. Therefore, two nickases  
20 can be used in combination to make a double-stranded cut in a specific region. Additional nickases are also known in the art, for example, McCaffery *et al.* (2016) *Nucleic Acids Res.* 44(2):e11. doi: 10.1093/nar/gkv878. Epub 2015 Oct 19.

[0139] In certain embodiments, the nuclease cleaves a safe harbor gene (*e.g.*, CCR5, HPRT, AAVS1, ATPA1, CLYBL, Rosa, Albumin, etc). *See, e.g.*, U.S. Patent  
25 Nos. 7,888,121; 7,972,854; 7,914,796; 7,951,925; 8,110,379; 8,409,861; 8,586,526; U.S. Patent Publications 20030232410; 20050208489; 20050026157; 20060063231; 20080159996; 201000218264; 20120017290; 20110265198; 20130137104; 20130122591; 20130177983 and 20130177960. In preferred embodiments, the nuclease cleaves an endogenous albumin gene such that the expression cassette is  
30 integrated into the endogenous albumin locus of a liver cell. Albumin-specific nucleases are described, for example, in U.S. Patent No. 9,150,847; and U.S. Publication Nos. 20130177983 and 20150056705.

[0140] In addition to, or instead of, nucleases as described herein, a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene may be modulated (down-regulated)

by the use of one or more engineered transcription factors. As with engineered nucleases, engineered transcription factors typically comprise at least one DNA-binding domain (*e.g.*, that binds the targeted gene) and a functional domain (*e.g.*, a transcriptional regulatory domain). Any engineered transcription factor that

5 modulated PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene expression may be used.

#### A. DNA-binding domains

[0141] Any DNA-binding domain can be used in the nuclease and

10 transcription factors described herein, including but not limited to a zinc finger DNA-binding domain, a TALE DNA binding domain, or a DNA-binding domain from a meganuclease, or a CRISPR/Cas DNA binding complex (*e.g.*, single guide RNA). In certain embodiments, the DNA-binding domain of the artificial transcription factor or artificial nuclease binds to a target site of 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20,

15 21, 22, 23, 24 or more nucleotides as shown in any of the target sites disclosed herein.

[0142] In certain embodiments, the DNA-binding domain comprises a naturally occurring or engineered (non-naturally occurring) TAL effector DNA binding domain. *See, e.g.*, U.S. Patent No. 8,586,526, incorporated by reference in its entirety herein. The plant pathogenic bacteria of the genus *Xanthomonas* are known

20 to cause many diseases in important crop plants. Pathogenicity of *Xanthomonas* depends on a conserved type III secretion (T3S) system which injects more than 25 different effector proteins into the plant cell. Among these injected proteins are transcription activator-like effectors (TALE) which mimic plant transcriptional activators and manipulate the plant transcriptome (see Kay *et al* (2007) *Science*

25 318:648-651). These proteins contain a DNA binding domain and a transcriptional activation domain. One of the most well characterized TALEs is AvrBs3 from *Xanthomonas campestris* pv. *Vesicatoria* (see Bonas *et al* (1989) *Mol Gen Genet* 218: 127-136 and WO2010079430). TALEs contain a centralized domain of tandem repeats, each repeat containing approximately 34 amino acids, which are key to the

30 DNA binding specificity of these proteins. In addition, they contain a nuclear localization sequence and an acidic transcriptional activation domain (for a review see Schornack *et al* (2006) *J Plant Physiol* 163(3): 256-272). In addition, in the phytopathogenic bacteria *Ralstonia solanacearum* two genes, designated brg11 and hpx17 have been found that are homologous to the AvrBs3 family of *Xanthomonas* in

the *R. solanacearum* biovar 1 strain GMI1000 and in the biovar 4 strain RS1000 (See Heuer *et al* (2007) *Appl and Envir Micro* 73(13): 4379-4384). These genes are 98.9% identical in nucleotide sequence to each other but differ by a deletion of 1,575 base pairs in the repeat domain of hpx17. However, both gene products have less than 40% sequence identity with AvrBs3 family proteins of *Xanthomonas*.

**[0143]** Thus, in some embodiments, the DNA binding domain that binds to a target site in a target locus is an engineered domain from a TAL effector similar to those derived from the plant pathogens *Xanthomonas* (see Boch *et al*, (2009) *Science* 326: 1509-1512 and Moscou and Bogdanove, (2009) *Science* 326: 1501) and *Ralstonia* (see Heuer *et al* (2007) *Applied and Environmental Microbiology* 73(13): 4379-4384); U.S. Patent Nos. 8,586,526; 8,420,782 and 8,440,431.

**[0144]** In certain embodiments, the DNA binding domain comprises a zinc finger protein. Preferably, the zinc finger protein is non-naturally occurring in that it is engineered to bind to a target site of choice. See, for example, Beerli *et al*. (2002) *Nature Biotechnol.* 20:135-141; Pabo *et al*. (2001) *Ann. Rev. Biochem.* 70:313-340; Isalan *et al*. (2001) *Nature Biotechnol.* 19:656-660; Segal *et al*. (2001) *Curr. Opin. Biotechnol.* 12:632-637; Choo *et al*. (2000) *Curr. Opin. Struct. Biol.* 10:411-416; U.S. Patent Nos. 7,888,121; 7,972,854; 6,453,242; 6,534,261; 6,599,692; 6,503,717; 6,689,558; 7,030,215; 6,794,136; 7,067,317; 7,262,054; 7,070,934; 7,361,635; 7,253,273, all incorporated herein by reference in their entireties.

**[0145]** An engineered zinc finger binding or TALE domain can have a novel binding specificity, compared to a naturally-occurring zinc finger or TALE protein. Engineering methods include, but are not limited to, rational design and various types of selection. Rational design includes, for example, using databases comprising triplet (or quadruplet) nucleotide sequences and individual zinc finger amino acid sequences, in which each triplet or quadruplet nucleotide sequence is associated with one or more amino acid sequences of zinc fingers which bind the particular triplet or quadruplet sequence. See, for example, U.S. Patents 8,586,526; 6,453,242 and 6,534,261, incorporated by reference herein in their entireties.

**[0146]** Exemplary selection methods, including phage display and two-hybrid systems, are disclosed in U.S. Patents 5,789,538; 5,925,523; 6,007,988; 6,013,453; 6,410,248; 6,140,466; 6,200,759; and 6,242,568; as well as WO 98/37186; WO 98/53057; WO 00/27878; WO 01/88197 and GB 2,338,237. In addition,



enhancement of binding specificity for zinc finger binding domains has been described, for example, in WO 02/077227.

[0147] In addition, as disclosed in these and other references, DNA domains (*e.g.*, multi-fingered zinc finger proteins or TALE domains) may be linked together using any suitable linker sequences, including for example, linkers of 5 or more amino acids in length. See, also, U.S. Patent Nos. 6,479,626; 6,903,185; 7,153,949; 9,567,609; and U.S. Patent Publication Nos. 20170218349 and 20170211075 for non-limiting examples of linker sequences. The DNA binding proteins described herein may include any combination of suitable linkers between the individual zinc fingers of the protein. In addition, enhancement of binding specificity for zinc finger binding domains has been described, for example, in WO 02/077227.

[0148] Selection of target sites; DNA-binding domains and methods for design and construction of fusion molecules (and polynucleotides encoding same) are known to those of skill in the art and described in detail in U.S. Patent Nos. 6,140,081; 5,789,538; 6,453,242; 6,534,261; 5,925,523; 6,007,988; 6,013,453; 6,200,759; WO 95/19431; WO 96/06166; WO 98/53057; WO 98/54311; WO 00/27878; WO 01/60970 WO 01/88197; WO 02/099084; WO 98/53058; WO 98/53059; WO 98/53060; WO 02/016536 and WO 03/016496 and U.S. Publication No. 20110301073.

[0149] In still further embodiments, the DNA binding domain comprises a DNA-binding single-guide RNA in combination with a functional domain (*e.g.*, CRISPR/Cas transcription factor). See, *e.g.*, U.S. Patent No. 8,697,359.

#### B. Functional Domains

[0150] The DNA-binding domains may be operably linked to any functional domain useful in gene modulation (*e.g.*, repression) as described herein. Thus, artificial nucleases and transcription factors comprising DNA-binding domains (*e.g.*, ZFPs or TALEs, CRISPR/Cas components such as single guide RNAs) as described herein associated with a heterologous regulatory (functional) domain (or functional fragment thereof) are also provided. Common domains include, *e.g.*, transcription factor domains (activators, repressors, co-activators, co-repressors), silencers, oncogenes (*e.g.*, myc, jun, fos, myb, max, mad, rel, ets, bcl, myb, mos family members etc.); DNA repair enzymes and their associated factors and modifiers; DNA rearrangement enzymes and their associated factors and modifiers; chromatin

associated proteins and their modifiers (*e.g.* kinases, acetylases and deacetylases); and DNA modifying enzymes (*e.g.*, methyltransferases, topoisomerases, helicases, ligases, kinases, phosphatases, polymerases, endonucleases) and their associated factors and modifiers. Thus, the invention provides artificial transcription factors comprising the DNA-binding domains described herein and a transcriptional regulatory domain as well as artificial nucleases comprising the DNA-binding domains and one or more nuclease domains.

**[0151]** In certain embodiments, the functional domain comprises a transcriptional regulatory domain, for example a repression domain. Exemplary repression domains include, but are not limited to, KRAB A/B, KOX, TGF-beta-inducible early gene (TIEG), v-erbA, SID, MBD2, MBD3, members of the DNMT family (*e.g.*, DNMT1, DNMT3A, DNMT3B), Rb, and MeCP2. See, for example, Bird *et al.* (1999) *Cell* 99:451-454; Tyler *et al.* (1999) *Cell* 99:443-446; Knoepfler *et al.* (1999) *Cell* 99:447-450; and Robertson *et al.* (2000) *Nature Genet.* 25:338-342. Additional exemplary repression domains include, but are not limited to, ROM2 and AtHD2A. See, for example, Chem *et al.* (1996) *Plant Cell* 8:305-321; and Wu *et al.* (2000) *Plant J.* 22:19-27.

**[0152]** It will be clear to those of skill in the art that, in the formation of a fusion molecule (or a nucleic acid encoding same) between a DNA-binding domain as described herein and a functional domain, either a repression domain or a molecule that interacts with a repression domain is suitable as a functional domain. Essentially any molecule capable of recruiting a repressor (*e.g.*, protein or complex) and/or repressing activity (such as, for example, histone demethylation) to the target gene is useful as a repression domain, and consequently can be used in the practice of the claimed invention. Insulator domains, localization domains, and chromatin remodeling proteins such as ISWI-containing domains and/or methyl binding domain proteins suitable for use as functional domains in fusion molecules are described, for example, in U.S. Patent Nos. 6,919,204 and 7,053,264.

**[0153]** Non-limiting examples of transcriptional activation domains. Suitable domains for achieving activation (transcriptional activation domains) include the HSV VP16 activation domain; nuclear hormone receptors; the p65 subunit of nuclear factor kappa B or artificial chimeric functional domains such as VP64 and degron. Additional exemplary activation domains include, Oct 1, Oct-2A, Sp1, AP-2, and

CTF1 as well as p300, CBP, PCAF, SRC1 P<sub>v</sub>ALF, AtHD2A, ERF-2, OsGAI, HALF-1, C1, AP1, ARF-5,-6,-7, and -8, CPRF1, CPRF4, MYC-RP/GP, and TRAB1.

**[0154]** In certain embodiments, the target site bound by the DNA-binding domain is present in an accessible region of cellular chromatin. Accessible regions can be determined as described, for example, in U.S. Patent No. 6,511,808. If the target site is not present in an accessible region of cellular chromatin, one or more accessible regions can be generated as described in WO 01/83793. In additional embodiments, the DNA-binding domain of a fusion molecule is capable of binding to cellular chromatin regardless of whether its target site is in an accessible region or not. For example, such DNA-binding domains are capable of binding to linker DNA and/or nucleosomal DNA. Examples of this type of "pioneer" DNA binding domain are found in certain steroid receptor and in hepatocyte nuclear factor 3 (HNF3). Cordingley *et al.* (1987) *Cell* 48:261-270; Pina *et al.* (1990) *Cell* 60:719-731; and Cirillo *et al.* (1998) *EMBO J.* 17:244-254.

**[0155]** In certain embodiments, at least one component of the proteins and systems described herein is naturally occurring (*e.g.*, a naturally occurring functional domain). In other embodiments, the compositions described herein are comprised of all non-naturally occurring components, *i.e.*, engineered in the DNA-binding domain and the functional domain. For example, the DNA-binding domain of a naturally-occurring domain may be altered to bind to a selected target site.

**[0156]** In certain embodiments, the PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene modulator comprises a CRISPR/Cas transcription factor system. The CRISPR (clustered regularly interspaced short palindromic repeats) locus, which encodes RNA components of the system, and the cas (CRISPR-associated) locus, which encodes proteins (Jansen *et al.*, 2002. *Mol. Microbiol.* 43: 1565-1575; Makarova *et al.*, 2002. *Nucleic Acids Res.* 30: 482-496; Makarova *et al.*, 2006. *Biol. Direct* 1: 7; Haft *et al.*, 2005. *PLoS Comput. Biol.* 1: e60) make up the gene sequences of the CRISPR/Cas system. CRISPR loci in microbial hosts contain a combination of CRISPR-associated (Cas) genes as well as non-coding RNA elements capable of programming the specificity of the CRISPR-mediated nucleic acid cleavage.

**[0157]** The Type II CRISPR, initially described in *S. pyogenes*, is one of the most well characterized systems and carries out targeted DNA double-strand break in four sequential steps. First, two non-coding RNA, the pre-crRNA array and tracrRNA, are transcribed from the CRISPR locus. Second, tracrRNA hybridizes to

the repeat regions of the pre-crRNA and mediates the processing of pre-crRNA into mature crRNAs containing individual spacer sequences where processing occurs by a double strand-specific RNase III in the presence of the Cas9 protein. Third, the mature crRNA:tracrRNA complex directs Cas9 to the target DNA via Watson-Crick base-pairing between the spacer on the crRNA and the protospacer on the target DNA next to the protospacer adjacent motif (PAM), an additional requirement for target recognition. In addition, the tracrRNA must also be present as it base pairs with the crRNA at its 3' end, and this association triggers Cas9 activity. Finally, Cas9 mediates cleavage of target DNA to create a double-stranded break within the protospacer. Activity of the CRISPR/Cas system comprises of three steps: (i) insertion of alien DNA sequences into the CRISPR array to prevent future attacks, in a process called 'adaptation', (ii) expression of the relevant proteins, as well as expression and processing of the array, followed by (iii) RNA-mediated interference with the alien nucleic acid. Thus, in the bacterial cell, several of the so-called 'Cas' proteins are involved with the natural function of the CRISPR/Cas system and serve roles in functions such as insertion of the alien DNA etc.

**[0158]** Type II CRISPR systems have been found in many different bacteria. BLAST searches on publically available genomes by Fonfara *et al* ((2013) *Nuc Acid Res* 42(4):2377-2590) found Cas9 orthologs in 347 species of bacteria. Additionally, this group demonstrated *in vitro* CRISPR/Cas cleavage of a DNA target using Cas9 orthologs from *S. pyogenes*, *S. mutans*, *S. therophilus*, *C. jejuni*, *N. meningitides*, *P. multocida* and *F. novicida*. Thus, the term "Cas9" refers to an RNA guided DNA nuclease comprising a DNA binding domain and two nuclease domains, where the gene encoding the Cas9 may be derived from any suitable bacteria.

**[0159]** The wild-type Cas9 protein has at least two nuclease domains: one nuclease domain is similar to a HNH endonuclease, while the other resembles a Ruv endonuclease domain. The HNH-type domain appears to be responsible for cleaving the DNA strand that is complementary to the crRNA while the Ruv domain cleaves the non-complementary strand. The Cas 9 nuclease can be engineered such that only one of the nuclease domains is functional, creating a Cas nickase (see Jinek *et al*, *ibid*). Nickases can be generated by specific mutation of amino acids in the catalytic domain of the enzyme, or by truncation of part or all of the domain such that it is no longer functional. Since Cas 9 comprises two nuclease domains, this approach may be taken on either domain. A double strand break can be achieved in the target DNA

by the use of two such Cas 9 nickases. The nickases will each cleave one strand of the DNA and the use of two will create a double strand break.

**[0160]** The requirement of the crRNA-tracrRNA complex can be avoided by use of an engineered “single-guide RNA” (sgRNA) that comprises the hairpin

5 normally formed by the annealing of the crRNA and the tracrRNA (see Jinek *et al* (2012) *Science* 337:816 and Cong *et al* (2013) *Scienceexpress*/10.1126/science.1231143). In *S. pyrogenes*, the engineered tracrRNA:crRNA fusion, or the sgRNA, guides Cas9 to cleave the target DNA when a double strand RNA:DNA heterodimer forms between the Cas associated RNAs and the target DNA. This system comprising the Cas9 protein and an engineered sgRNA containing a PAM sequence has been used for RNA guided genome editing (see Ramalingam *ibid*) and has been useful for zebrafish embryo genomic editing *in vivo* (see Hwang *et al* (2013) *Nature Biotechnology* 31 (3):227) with editing efficiencies similar to ZFNs and TALENs. In addition, CRISPR/Cas transcription factors have also been described. See, e.g., U.S. Patent No. 8,697,359; Platek *et al.* (2014) *Plant Biotechnology J.* doi: 10.1111/pbi.12284 and Perez-Pinera *et al.* (2013) *Nature Methods* 10:973–976).

**[0161]** Chimeric or sgRNAs can be engineered to comprise a sequence complementary to any desired target for transcriptional regulation as described herein.

20 In some embodiments, a guide sequence is about or more than about 5, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 35, 40, 45, 50, 75, or more nucleotides in length. In some embodiments, a guide sequence is less than about 75, 50, 45, 40, 35, 30, 25, 20, 15, 12, or fewer nucleotides in length. In certain embodiments, the RNAs comprise 22 bases of complementarity to a target and of the form G[n19], followed by a protospacer-adjacent motif (PAM) of the form NGG. Thus, in one method, sgRNAs can be designed by utilization of a known target of a DNA-binding domain (e.g., ZFP, TALE) in a gene of interest. In addition, sgRNAs can be designed to known paired nuclease sites by (i) aligning the target sequences of the ZFN heterodimer with the reference sequence of the relevant genome (human, mouse, or of a particular plant species); (ii) identifying the spacer region between the ZFN half-sites; (iii) identifying the location of the motif G[N20]GG that is closest to the spacer region (when more than one such motif overlaps the spacer, the motif that is centered relative to the spacer is chosen); (iv) using that motif as the core of the sgRNA. This method advantageously relies on proven nuclease targets.

Alternatively, sgRNAs can be designed to target any region of interest simply by identifying a suitable target sequence the conforms to the G[n20]GG formula. Along with the complementarity region, an sgRNA may comprise additional nucleotides to extend to tail region of the tracrRNA portion of the sgRNA (see Hsu *et al* (2013)

5 *Nature Biotech* doi:10.1038/nbt.2647). Tails may be of +67 to +85 nucleotides, or any number therebetween with a preferred length of +85 nucleotides. Truncated sgRNAs may also be used, “tru-gRNAs” (see Fu *et al*, (2014) *Nature Biotech* 32(3): 279). In tru-gRNAs, the complementarity region is diminished to 17 or 18 nucleotides in length.

10 **[0162]** Further, alternative PAM sequences may also be utilized, where a PAM sequence can be NAG as an alternative to NGG (Hsu 2014, *ibid*) using a *S. pyogenes* Cas9. Additional PAM sequences may also include those lacking the initial G (Sander and Joung (2014) *Nature Biotech* 32(4):347). In addition to the *S. pyogenes* encoded Cas9 PAM sequences, other PAM sequences can be used that are  
15 specific for Cas9 proteins from other bacterial sources. For example, the PAM sequences shown below (adapted from Sander and Joung, *ibid*, and Esvelt *et al*, (2013) *Nat Meth* 10(11):1116) are specific for these Cas9 proteins:

	Species	PAM
20	<i>S. pyogenes</i>	NGG
	<i>S. pyogenes</i>	NAG
	<i>S. mutans</i>	NGG
	<i>S. thermophilus</i>	NGGNG
	<i>S. thermophilus</i>	NNAAAW
25	<i>S. thermophilus</i>	NNAGAA
	<i>S. thermophilus</i>	NNNGATT
	<i>C. jejuni</i>	NNNNACA
	<i>N. meningitides</i>	NNNNGATT
	<i>P. multocida</i>	GNNNCNNA
30	<i>F. novicida</i>	NG

**[0163]** Thus, a suitable target sequence for use with a *S. pyogenes* CRISPR/Cas system can be chosen according to the following guideline: [n17, n18, n19, or n20](G/A)G. Alternatively the PAM sequence can follow the guideline G[n17, n18, n19, n20](G/A)G. For Cas9 proteins derived from non-*S. pyogenes*

bacteria, the same guidelines may be used where the alternate PAMs are substituted in for the *S. pyogenes* PAM sequences.

**[0164]** Most preferred is to choose a target sequence with the highest likelihood of specificity that avoids potential off target sequences. These undesired off target sequences can be identified by considering the following attributes: i) similarity in the target sequence that is followed by a PAM sequence known to function with the Cas9 protein being utilized; ii) a similar target sequence with fewer than three mismatches from the desired target sequence; iii) a similar target sequence as in ii), where the mismatches are all located in the PAM distal region rather than the PAM proximal region (there is some evidence that nucleotides 1-5 immediately adjacent or proximal to the PAM, sometimes referred to as the ‘seed’ region (Wu *et al* (2014) *Nature Biotech* doi:10.1038/nbt2889) are the most critical for recognition, so putative off target sites with mismatches located in the seed region may be the least likely to be recognized by the sg RNA); and iv) a similar target sequence where the mismatches are not consecutively spaced or are spaced greater than four nucleotides apart (Hsu 2014, *ibid*). Thus, by performing an analysis of the number of potential off target sites in a genome for whichever CRISPR/Cas system is being employed, using these criteria above, a suitable target sequence for the sgRNA may be identified.

**[0165]** In some embodiments, the CRISPR-Cpf1 system is used. The CRISPR-Cpf1 system, identified in *Francisella* spp, is a class 2 CRISPR-Cas system that mediates robust DNA interference in human cells. Although functionally conserved, Cpf1 and Cas9 differ in many aspects including in their guide RNAs and substrate specificity (see Fagerlund *et al*, (2015) *Genom Bio* 16:251). A major difference between Cas9 and Cpf1 proteins is that Cpf1 does not utilize tracrRNA, and thus requires only a crRNA. The FnCpf1 crRNAs are 42–44 nucleotides long (19-nucleotide repeat and 23–25-nucleotide spacer) and contain a single stem-loop, which tolerates sequence changes that retain secondary structure. In addition, the Cpf1 crRNAs are significantly shorter than the ~100-nucleotide engineered sgRNAs required by Cas9, and the PAM requirements for FnCpf1 are 5′ -TTN-3′ and 5′ -CTA-3′ on the displaced strand. Although both Cas9 and Cpf1 make double strand breaks in the target DNA, Cas9 uses its RuvC- and HNH-like domains to make blunt-ended cuts within the seed sequence of the guide RNA, whereas Cpf1 uses a RuvC-like domain to produce staggered cuts outside of the seed. Because Cpf1 makes

staggered cuts away from the critical seed region, NHEJ will not disrupt the target site, therefore ensuring that Cpf1 can continue to cut the same site until the desired HDR recombination event has taken place. Thus, in the methods and compositions described herein, it is understood that the term “Cas” includes both Cas9 and Cpf1 proteins. Thus, as used herein, a “CRISPR/Cas system” refers both CRISPR/Cas and/or CRISPR/Cpf1 systems, including both nuclease and/or transcription factor systems.

**[0166]** In certain embodiments, Cas protein may be a "functional derivative" of a naturally occurring Cas protein. A "functional derivative" of a native sequence polypeptide is a compound having a qualitative biological property in common with a native sequence polypeptide. "Functional derivatives" include, but are not limited to, fragments of a native sequence and derivatives of a native sequence polypeptide and its fragments, provided that they have a biological activity in common with a corresponding native sequence polypeptide. A biological activity contemplated herein is the ability of the functional derivative to hydrolyze a DNA substrate into fragments. The term "derivative" encompasses both amino acid sequence variants of polypeptide, covalent modifications, and fusions thereof. In some aspects, a functional derivative may comprise a single biological property of a naturally occurring Cas protein. In other aspects, a function derivative may comprise a subset of biological properties of a naturally occurring Cas protein. Suitable derivatives of a Cas polypeptide or a fragment thereof include but are not limited to mutants, fusions, covalent modifications of Cas protein or a fragment thereof. Cas protein, which includes Cas protein or a fragment thereof, as well as derivatives of Cas protein or a fragment thereof, may be obtainable from a cell or synthesized chemically or by a combination of these two procedures. The cell may be a cell that naturally produces Cas protein, or a cell that naturally produces Cas protein and is genetically engineered to produce the endogenous Cas protein at a higher expression level or to produce a Cas protein from an exogenously introduced nucleic acid, which nucleic acid encodes a Cas that is same or different from the endogenous Cas. In some case, the cell does not naturally produce Cas protein and is genetically engineered to produce a Cas protein.

**[0167]** Exemplary CRISPR/Cas single guide RNAs targeted to specific genes are disclosed for example, in U.S. Publication No. 20150056705.

**[0168]** Thus, the compositions and systems described herein comprises one or more DNA-binding domains that specifically bind to one or more target sites in



selected gene and at least one functional domain such that, when introduced into a cell, the composition (or system) modulates PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene expression.

## 5 Target Sites

[0169] As described in detail above, DNA-binding domains of the artificial nucleases and/or transcription factors as described herein can be engineered to bind to any sequence of choice. An engineered DNA-binding domain can have a novel binding specificity, compared to a naturally-occurring DNA-binding domain. In  
10 certain embodiments, the DNA-binding domains bind to a sequence within a liver-specific gene as described herein, for example a target site (typically 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21 or even more base pairs), including DNA-binding domains that bind to a sequence or 12 or more base pairs (contiguous or non-contiguous) of the target sites shown herein (*e.g.*, Tables 1, 3, 5, 7, 11, 13, 14, and  
15 16). Engineering methods include, but are not limited to, rational design and various types of selection. Rational design includes, for example, using databases comprising triplet (or quadruplet) nucleotide sequences and individual zinc finger amino acid sequences, in which each triplet or quadruplet nucleotide sequence is associated with one or more amino acid sequences of zinc fingers which bind the particular triplet or  
20 quadruplet sequence. *See*, for example, co-owned U.S. Patent Nos. 6,453,242 and 6,534,261, incorporated by reference herein in their entireties. Rational design of TAL-effector domains and single guide RNAs is also known. *See, e.g.*, U.S. Patent Nos. 8,771,985 and U.S. Patent Publication No. 20150056705.

## 25 Delivery

[0170] The nucleases, transcription factors and/or transgenes (*e.g.*, PCSK9, TTR, SERPINA1, KLKB1 or HAO1 inhibitors) may be delivered *in vivo* or *ex vivo* by any suitable means into any cell type, preferably to the liver (systemically or via hepatic delivery). Similarly, when used in combination with nucleases for targeted  
30 integration, the nucleases may be delivered in polynucleotide and/or protein form, for example using non-viral vector(s), viral vectors(s) and/or in RNA form, *e.g.*, as mRNA.

[0171] Methods of non-viral delivery of nucleic acids include electroporation, lipofection, microinjection, biolistics, virosomes, liposomes, immunoliposomes, other

nanoparticle, polycation or lipid:nucleic acid conjugates, naked DNA, artificial virions, and agent-enhanced uptake of DNA. Sonoporation using, *e.g.*, the Sonitron 2000 system (Rich-Mar) can also be used for delivery of nucleic acids. Additional exemplary nucleic acid delivery systems include those provided by AmaxaBiosystems (Cologne, Germany), Maxcyte, Inc. (Rockville, Maryland), BTX Molecular Delivery Systems (Holliston, MA) and Copernicus Therapeutics Inc., (*see* for example US6008336).

[0172] In preferred embodiments, the expression constructs are AAV vectors. The optional nucleases may be administered in mRNA form or using one or more viral vectors (AAV, Ad, etc.). Administration can be by any means in which the polynucleotides are delivered to the desired target cells. Both *in vivo* and *ex vivo* methods are contemplated. Intravenous injection to the portal vein is a preferred method of administration. Other *in vivo* administration modes include, for example, direct injection into the lobes of the liver or the biliary duct and intravenous injection distal to the liver, including through the hepatic artery, direct injection in to the liver parenchyma, injection via the hepatic artery, and/or retrograde injection through the biliary tree. *Ex vivo* modes of administration include transduction *in vitro* of resected hepatocytes or other cells of the liver, followed by infusion of the transduced, resected hepatocytes back into the portal vasculature, liver parenchyma or biliary tree of the human patient, *see e.g.*, Grossman *et al.*, (1994) *Nature Genetics*, 6:335-341.

[0173] In systems involving delivery of more than one polynucleotides (*e.g.*, construct as described herein and nuclease in polynucleotide form), the two or more polynucleotide(s) are delivered using one or more of the same and/or different vectors. For example, the nuclease in polynucleotide form may be delivered in mRNA form and the liver-specific constructs as described herein may be delivered via other modalities such as viral vectors (*e.g.*, AAV), minicircle DNA, plasmid DNA, linear DNA, liposomes, nanoparticles and the like.

[0174] Pharmaceutically acceptable carriers are determined in part by the particular composition being administered, as well as by the particular method used to administer the composition. Accordingly, there is a wide variety of suitable formulations of pharmaceutical compositions available, as described below (*see, e.g.*, *Remington's Pharmaceutical Sciences*, 17th ed., 1989).

[0175] The effective amount of expression cassette (and optional nuclease(s), and/or modified cells) to be administered will vary from patient to patient.

Accordingly, effective amounts are best determined by the physician administering the compositions (*e.g.*, cells) and appropriate dosages can be determined readily by one of ordinary skill in the art. Analysis of the serum or other tissue levels of the therapeutic polypeptide and comparison to the initial level prior to administration can determine whether the amount being administered is too low, within the right range or too high. Suitable regimes for initial and subsequent administrations are also variable, but are typified by an initial administration followed by subsequent administrations if necessary. Subsequent administrations may be administered at variable intervals, ranging from daily to annually to every several years. One of skill in the art will appreciate that appropriate immunosuppressive techniques may be recommended to avoid inhibition or blockage of transduction by immunosuppression of the delivery vectors, see *e.g.*, Vilquin *et al.*, (1995) *Human Gene Ther.*, 6:1391-1401.

[0176] Formulations for both *ex vivo* and *in vivo* administrations include suspensions (*e.g.*, of genetically modified cells, liposomes or nanoparticles) in liquid or emulsified liquids. The active ingredients often are mixed with excipients which are pharmaceutically acceptable and compatible with the active ingredient. Suitable excipients include, for example, water, saline, dextrose, glycerol, ethanol or the like, and combinations thereof. In addition, the composition may contain minor amounts of auxiliary substances, such as, wetting or emulsifying agents, pH buffering agents, stabilizing agents or other reagents that enhance the effectiveness of the pharmaceutical composition.

### Applications

[0177] The methods and compositions disclosed herein are useful in providing therapies for any PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 -related disorder by modulating the expression of the endogenous PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene, including via the provision of a PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 -inhibitory transgene (transcription factor, nuclease, etc.). In addition, the methods and compositions provide methods for knock out of a mutant PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene and the provision of a wild type PCSK9, TTR, SERPINA1, KLKB1 or HAO1 cDNA for treatment or prevention of such a disorder. The cell may be modified *in vivo* or may be modified *ex vivo* and subsequently administered to a subject. Thus, the methods and compositions provide for the treatment and/or prevention of such PCSK9, TTR, SERPINA1, KLKB1 and/or

HAO1 related disorders. Thus, the compositions and methods described herein can be used to treat or prevent disorders including *e.g.*, TTR-Mediated Amyloidosis, A1AT Deficiency, Hereditary Angioedema, Familial Hypercholesterolemia/Static resistant hypercholesterolemia and Hyperxoluria.

5    **[0178]**           The following Examples include exemplary embodiments of the present disclosure in which the optionally used nuclease comprises a zinc finger nuclease (ZFN) or CRISPR/Cas system. It will be appreciated that this is for purposes of exemplification only and that other nucleases can be used, for example TALENs, homing endonucleases (meganucleases) with engineered DNA-binding  
10   domains that bind to target sites as described herein and/or fusions of naturally occurring of engineered homing endonucleases (meganucleases) DNA-binding domains and heterologous cleavage domains and/or fusions of meganucleases and TALE proteins. In addition, it will be appreciated that expression constructs as described herein can be carried on other vectors (other than AAV) to produce the  
15   same results in the treatment and/or prevention of disorders caused by deficient protein production.

### EXAMPLES

**Example 1: Design, Construction and general characterization of compositions that modulate the PCSK9, TTR, SERPINA1, KLKB1 or HAO1 genes**

20   **[0179]**           Zinc finger, Cas and TALE proteins that bind to either the mouse or human PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene operably linked to transcriptional regulatory or nuclease domains are designed and incorporated into plasmids, AAV or adenoviral vectors or made into mRNA essentially as described in  
25   Urnov *et al.* (2005) *Nature* 435(7042):646-651, Perez *et al* (2008) *Nature Biotechnology* 26(7):808-816, and as described in U.S. Patent Nos. 8,586,526 and 6,534,261.

**[0180]**           sgRNAs for use in the CRISPR/Cas system are made synthetically by methods known in the art (see Hsu *et al*, (2013) *Nature Biotech* doi:10.1038/nbt.2647,  
30   or Sternberg *et al*, (2014) *Nature* 507: 62). The sgRNAs are engineered as described above and are designed to target a sequence in the endogenous PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 gene (*e.g.*, a target sequences as set forth in herein).

[0181] The human or mouse PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 -specific nucleases are introduced into human (*e.g.* K562) or mouse cells as mRNA and are analyzed for cleavage activity. Briefly, the cells are transfected with two doses of mRNA (2 or 6 µg in total of the two ZFNs) using a BTX 96 well electroporator (BTX) according to standard protocols. Cells are then expanded for an additional 7 days. Cells are removed at day 7 and analyzed for on target PCSK9, TTR, SERPINA1, KLKB1 or HAO1 modification using deep sequencing (Miseq, Illumina).

[0182] For PCSK9, TTR, SERPINA1, KLKB1 and/or HAO1 -specific CRISPR/Cas experiments, Cas9 is supplied on a pVAX plasmid, and the sgRNA are supplied on a plasmid under the control of the U6 promoter. The plasmids are mixed at either 100 ng of each or 400 ng of each and are mixed with 2e5 cells per run. The cells are transfected using the Amaxa system. Briefly, an Amaxa transfection kit is used and the nucleic acids are transfected using a standard Amaxa shuttle protocol. Following transfection, the cells are left to rest for 10 minutes at room temperature and then resuspended in prewarmed RPMI. The cells are then grown in standard conditions at 37 °C. Genomic DNA is isolated 7 days after transfection and subject to MiSeq analysis.

[0183] All nucleases are found bind to their target sites and to be functionally active (in cleaving the target gene).

#### **Example 2: Design, Construction and general characterization of compositions that modulate PCSK9, TTR, SERPINA1, KLKB1 or HAO1 gene expression**

[0184] Zinc finger proteins are targeted to PCSK9, TTR, SERPINA1, KLKB1 or HAO1 were engineered essentially as described in Zhang *et al* (2000) *J Biol Chem* 275(43):33850-33860). The ZFPs were evaluated and shown to be bind and cleave their target sites. Linkers and ZFN architecture are as previously described (U.S. Publication Nos. 20170218349 and 20170211075). For example, linker L0 is LRGSQLVKS (SEQ ID NO:139), linker N7a is SGTPHEVGVYTL (SEQ ID NO:140), linker N6a is SGAQGSTLDF (SEQ ID NO:141), and linker L8c4 is LRGSYAPMPPLALASP (SEQ ID NO:142).

# I. Nuclease targeting of PCSK9

[0185] PCSK9-specific zinc finger proteins were made as described above.

Exemplary proteins (including the linker) are shown below in Table 1.

**Table 1: Human PCSK9-specific zinc finger proteins**

	Design						
SBS #/ Target	F1	F2	F3	F4	F5	F6	linker
SBS#60045 5' taCGTGGTGGTGC TGAAGGAggagacc c (SEQ ID NO:1)	QSSHLTR (SEQ ID NO:13)	QSGNLAR (SEQ ID NO:14)	QSSDLNR (SEQ ID NO:15)	WHSSLHQ (SEQ ID NO:16)	LRHHLTR (SEQ ID NO:17)	HKQHRDA (SEQ ID NO:18)	L0
SBS#60043 5' agGATCCGtGGAG GTTGCTGgcaccta (SEQ ID NO:2)	RSDVLSE (SEQ ID NO:19)	TRNGLKY (SEQ ID NO:20)	TSGHLSR (SEQ ID NO:21)	QSGHLSR (SEQ ID NO:22)	NNRDLIN (SEQ ID NO:23)	TSSNLSR (SEQ ID NO:24)	N7a*
SBS#60134 5' ttACCGGGGGGCT GGTAttcatcgccc (SEQ ID NO:3)	QSGALAR (SEQ ID NO:25)	RSDVLSE (SEQ ID NO:19)	RSAHLSR (SEQ ID NO:26)	RSDHLSR (SEQ ID NO:27)	DRSVLAR (SEQ ID NO:28)	N/A	N7a*
SBS#60133 5' ccGCCCGGTACCG TGGAGGGgtaatccg (SEQ ID NO:4)	RSDHLSR (SEQ ID NO:27)	QSGHLSR (SEQ ID NO:22)	HKQHRDA (SEQ ID NO:18)	DNSNRIK (SEQ ID NO:29)	RSDHLSE (SEQ ID NO:30)	HSRTRTK (SEQ ID NO:31)	L0
SBS#60137 5' agATGGGGGTCTT ACCGGGGggtcgta (SEQ ID NO:5)	RSDHLSR (SEQ ID NO:27)	QQWDRKQ (SEQ ID NO:32)	TPSYLPT (SEQ ID NO:33)	DRSALAR (SEQ ID NO:34)	RSDHLSE (SEQ ID NO:30)	RKDARIT (SEQ ID NO:35)	N6a*
SBS#60334 5' ggTACCGGGCGG ATGAAtaccagccccc (SEQ ID NO:6)	QSGNLAR (SEQ ID NO:14)	TSGNLTR (SEQ ID NO:36)	RSDDLTR (SEQ ID NO:37)	RSDHLSE (SEQ ID NO:30)	DKSNRKK (SEQ ID NO:38)	N/A	N7a*
SBS#60259 5' gaGGCTGGGGAG TAGAGGCaggatc gt (SEQ ID NO:7)	QSGDLTR (SEQ ID NO:39)	RSDNLTR (SEQ ID NO:40)	QSGALAR (SEQ ID NO:25)	QSGHLSR (SEQ ID NO:22)	RSDHLSQ (SEQ ID NO:41)	DSSHRTR (SEQ ID NO:42)	N7a*
SBS#60254 5' cgCTGCCGGCAAC TTCCGGGacgatgcc (SEQ ID NO:8)	RSAHLSR (SEQ ID NO:26)	DSSDRKK (SEQ ID NO:43)	QHQLVLR (SEQ ID NO:44)	QNATRTK (SEQ ID NO:45)	RSDTLSE (SEQ ID NO:46)	RSPGRMG (SEQ ID NO:47)	N6a*
SBS#60256 5' cgCTGCCGGCAAC TTCCGGGacgatgcc (SEQ ID NO:8)	RSAHLSR (SEQ ID NO:26)	DSSDRKK (SEQ ID NO:43)	QHQLVLR (SEQ ID NO:44)	QNATRTK (SEQ ID NO:45)	RSDTLSE (SEQ ID NO:46)	RSPGRMG (SEQ ID NO:47)	N7a*
SBS#60289 5' agAGAAGTGGAT CAGTCTCTgcctcaa c (SEQ ID NO:9)	DGYLLPT (SEQ ID NO:48)	DRSALAR (SEQ ID NO:34)	ERQTLIK (SEQ ID NO:49)	QSGHLSR (SEQ ID NO:22)	HRWHLQT (SEQ ID NO:50)	AQCCLFH (SEQ ID NO:51)	N6a*
SBS#60287 5' cgGAGCTACCCCT GGCCGAGtgaggca (SEQ ID NO:10)	RSDNLAR (SEQ ID NO:52)	DRSVLHR (SEQ ID NO:53)	RSDTLA (SEQ ID NO:54)	DKSTRTK (SEQ ID NO:55)	PCRYRLD (SEQ ID NO:56)	RSANLTR (SEQ ID NO:57)	N6a*

SBS#60309 5' taTCCCCGGCGGG CAGCCTGggcctgca (SEQ ID NO:11)	RSDVLSE (SEQ ID NO:19)	QKCCLRS (SEQ ID NO:58)	DRSHLTR (SEQ ID NO:59)	RSDDLTR (SEQ ID NO:37)	RSDTLN (SEQ ID NO:60)	TNSDRTK (SEQ ID NO:61)	N6a*
SBS#60308 5' tgCAGGCGGCGG GCaGTGCGCtctgac t (SEQ ID NO:12)	HRQRLEE (SEQ ID NO:62)	RNASRTR (SEQ ID NO:63)	DRSHLTR (SEQ ID NO:59)	RSDDLTR (SEQ ID NO:37)	RSDTLSE (SEQ ID NO:46)	KPYNLQQ (SEQ ID NO:64)	L0

\* indicates that the DNA binding domain is attached to the FokI nuclease domain at the DNA binding domain's N-terminus (see U.S. Publication Nos. 20170218349 and 20170211075)

**[0186]** The nucleases were tested for activity in K562 cells where 2 µg of each mRNA encoding a partner nuclease was added in a 100 µL transfection volume. All proteins bound to their targets and induced cleavage. Exemplary results of pairwise combinations are shown below in Table 2, where “%indel” indicates the activity. %indels were measured by deep sequencing (MiSeq, Illumnia) and then analyzed for the percent of alleles comprising insertions and/or deletions at the cleavage site. Table 2 demonstrates that these exemplary ZFN pairs were all active.

**Table 2: PCSK9-specific nuclease activity**

Pair	%Indel
60045_60043	65.40
60134_60133	72.63
60137_60334	67.90
60259_60254	75.69
60259_60256	74.32
60289_60287	73.07
60309_60308	71.77

**[0187]** Zinc finger nucleases were also designed against the *Macacca mulatta* PCSK9 gene (mmPCSK9), and are shown below in Table 3.

**Table 3: *M. mulatta* PCSK9 zinc finger protein designs**

	Design						
SBS #/ Target/exon	F1	F2	F3	F4	F5	F6	Linker- Fok domain
SBS# 69903 5'gtGGAGG CTGCCCCG CACCTacgtg gtg (SEQ ID	TGQTLRG (SEQ ID NO:147)	QNATRTK (SEQ ID NO:45)	RSDNLSE (SEQ ID NO:95)	SKQYLIK (SEQ ID NO:148)	DRSHLTR (SEQ ID NO:59)	QSGHLR (SEQ ID NO:22)	N7a* ELD

NO:143), exon 2							
SBS# 69912 5'gtGGTGC TGAAGGA GGAGACcc accgct (SEQ ID NO:144), exon 2	DRSNLSR (SEQ ID NO:91)	QSGHLSR (SEQ ID NO:22)	QSGHLQR (SEQ ID NO:149)	QSGNLAR (SEQ ID NO:14)	QSSDLSR (SEQ ID NO:15)	TSGHLSR (SEQ ID NO:21)	L0 KKR
SBS# 69908 5'gtGGAGG CTGCCCGG CACCTacgtg gtg (SEQ ID NO:143), exon 2	YKWDLNN (SEQ ID NO:150)	QNATRTK (SEQ ID NO:45)	RSDTLSE (SEQ ID NO:46)	QKRNRTK (SEQ ID NO:151)	DRSHLTR (SEQ ID NO:59)	QSGHLSR (SEQ ID NO:22)	N7a* ELD
SBS# 69909 5'gtGGAGG CTGCCCGG CACCTacgtg gtg (SEQ ID NO:143), exon 2	TGQTLRG (SEQ ID NO:147)	QNATRTK (SEQ ID NO:45)	RSDTLSE (SEQ ID NO:46)	QKRNRTK (SEQ ID NO:151)	DRSHLTR (SEQ ID NO:59)	QSGHLSR (SEQ ID NO:22)	N7a* ELD
SBS# 69913 5'gtGGTGC TGAAGGA GGAGACcc accgct (SEQ ID NO:144), exon 2	DRSNLSR (SEQ ID NO:91)	QSGHLSR (SEQ ID NO:22)	QSGHLAR (SEQ ID NO:72)	QSGNLAR (SEQ ID NO:14)	QSSDLSR (SEQ ID NO:15)	TSGHLSR (SEQ ID NO:21)	L0 KKR
SBS# 69915 5'gtGGGAC ATCGCAGG CTGCTgccca cgt (SEQ ID NO:145), exon 7	QSSDLSR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLSA (SEQ ID NO:152)	RNNDRKT (SEQ ID NO:153)	DRSDLSR (SEQ ID NO:154)	RSHHLKA (SEQ ID NO:155)	N7a* ELD
SBS# 69922 5'ggGGTGG TGAATTAC CAGCCacgt gggc (SEQ ID NO:146), exon 7	DRSTRTK (SEQ ID NO:156)	RRDTLLD (SEQ ID NO:157)	QSAVLPG (SEQ ID NO:158)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR
SBS# 69925 5'ggGGTGG TGAATTAC CAGCCacgt gggc (SEQ ID NO:146), exon 7	DRSDLSR (SEQ ID NO:154)	QSGDLTR (SEQ ID NO:39)	QSAVLPG (SEQ ID NO:158)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR
SBS# 69928 5'ggGGTGG TGAATTAC CAGCCacgt gggc (SEQ ID NO:146), exon 7	ERGTLAR (SEQ ID NO:160)	QSADRTK (SEQ ID NO:68)	QSAVLPG (SEQ ID NO:158)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR
SBS# 69926 5'ggGGTGG TGAATTAC CAGCCacgt gggc (SEQ ID NO:146), exon 7	DRSDLSR (SEQ ID NO:154)	QSGDLTR (SEQ ID NO:39)	SHLGLTI (SEQ ID NO:161)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR



SBS# 69916 5'gtGGGAC ATCGCAGG CTGCTgccca cgt (SEQ ID NO:145), exon 7	QSSDLSR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLSA (SEQ ID NO:152)	RSNDRKK (SEQ ID NO:162)	DRSDLSR (SEQ ID NO:154)	RSHHLKA (SEQ ID NO:155)	N7a* ELD
SBS# 69918 5'gtGGGAC ATCGCAGG CTGCTgccca cgt (SEQ ID NO:145), exon 7	QSSDLSR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLSA (SEQ ID NO:152)	RNDRKKT (SEQ ID NO:153)	QNATRIN (SEQ ID NO:163)	RSAHLRS (SEQ ID NO:26)	N7a* ELD
SBS# 69919 5'gtGGGAC ATCGCAGG CTGCTgccca cgt (SEQ ID NO:145), exon 7	QSSDLSR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLSA (SEQ ID NO:152)	RSNDRKK (SEQ ID NO:162)	QNATRIN (SEQ ID NO:163)	RSAHLRS (SEQ ID NO:26)	N7a* ELD
SBS# 69920 5'gtGGGAC ATCGCAGG CTGCTgccca cgt (SEQ ID NO:145), exon 7	QSSDLSR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLST (SEQ ID NO:108)	RSNDRIK (SEQ ID NO:164)	QNATRIN (SEQ ID NO:163)	RSAHLRS (SEQ ID NO:26)	N7a* ELD
SBS# 69929 5'ggGGTGG TGACTTAC CAGCCacgt gggc (SEQ ID NO:146), exon 7	ERGTLAR (SEQ ID NO:160)	QSADRTK (SEQ ID NO:68)	SHLGLTI (SEQ ID NO:161)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR

\* indicates that the DNA binding domain is attached to the FokI nuclease domain at the DNA binding domain's N-terminus (see U.S. Publication Nos. 20170218349 and 20170211075)

**[0188]** Nucleases targeting the *M. mulatta* PCSK9 gene were tested for

- 5 activity in the Rhesus macaque cell line MK2, and exemplary pairwise combination results are shown below in Table 4. All proteins bound their targets and induced cleavage. The total dose of mRNA encoding each partner of the nuclease pair is indicated, and was added in a 100  $\mu$ L transfection volume. Activity is shown as % indels measured by deep sequencing (MiSeq, Illumina) and then analyzed for the
- 10 percent of alleles comprising insertions and/or deletions at the cleavage site.

**Table 4: Activity of *M. mulatta* PCSK9 ZFN pairs**

Indels (%)	total ug ZFN mRNA				
	0	0.003	0.01	0.03	0.1
69903/69912	0.05	0.17	1.93	9.08	33.74
69908/69912	0.05	0.22	1.73	11.82	42.54
69909/69912	0.05	0.48	4.19	21.30	58.71
69909/69913	0.05	0.56	3.63	18.28	53.39
69915/69922	0.03	0.21	3.93	31.72	77.97

69915/69925	0.03	0.26	1.26	25.27	71.53
69915/69928	0.03	0.20	1.16	17.96	63.41
69915/69926	0.03	0.16	1.26	7.28	37.05
69916/69922	0.03	0.46	5.88	27.89	81.41
69916/69925	0.03	0.56	3.28	17.00	76.97
69916/69928	0.03	0.25	3.15	23.90	69.22
69918/69922	0.03	0.28	2.02	21.78	71.24
69918/69925	0.03	0.14	1.75	15.82	68.68
69918/69928	0.03	0.09	0.79	14.92	57.86
69919/69922	0.03	0.16	2.62	26.07	76.11
69919/69925	0.03	0.25	1.82	22.85	66.23
69920/69922	0.03	0.51	2.94	20.56	72.11
69920/69925	0.03	0.43	2.17	18.13	71.79
69920/69928	0.03	0.27	1.51	12.35	54.45
69920/69929	0.03	0.30	1.45	10.77	43.34

**[0189]** ZFNs were designed against the mouse PCSK9 (mPCSK9) gene, and are shown below in Table 5.

**Table 5: Mouse PCSK9 zinc finger protein designs**

SBS #/ Target/exon (site)	Design						Linker- Fok domain
	F1	F2	F3	F4	F5	F6	
SBS# 60422 5'tcGGGAGAT TGAGGGCAG GGtcaccatc (SEQ ID NO:165), exon 4 (A)	RSAHLSR (SEQ ID NO:26)	QSGDLTR (SEQ ID NO:39)	RSDHLSA (SEQ ID NO:180)	SYWSRTV (SEQ ID NO:181)	QNAHRKT (SEQ ID NO:182)	RSAHLSR (SEQ ID NO:26)	N7a* ELD
SBS# 64793 5'tcGGGAGAT TGAGGGCAG GGtcaccatc (SEQ ID NO:165), exon 4 (A)	RSDHLSR (SEQ ID NO:27)	QSGDLTR (SEQ ID NO:39)	RSDHLSA (SEQ ID NO:180)	SYWSRTV (SEQ ID NO:181)	QSAHRKN (SEQ ID NO:183)	RSAHLSR (SEQ ID NO:26)	N7a* ELD
SBS# 64802 5'tcGGGAGAT TGAGGGCAG GGtcaccatc (SEQ ID NO:165), exon 4 (A)	RSDHLSR (SEQ ID NO:27)	QSSDLTR (SEQ ID NO:184)	RSDHLSA (SEQ ID NO:180)	SYWSRTV (SEQ ID NO:181)	QSAHRKN (SEQ ID NO:183)	RSAHLSR (SEQ ID NO:26)	N7a* ELD
SBS# 64791 5'tcGGGAGAT TGAGGGCAG GGtcaccatc (SEQ ID NO:165), exon 4 (A)	RSDHLSR (SEQ ID NO:27)	QSGDLTR (SEQ ID NO:39)	RSDHLSA (SEQ ID NO:180)	SYWSRTV (SEQ ID NO:181)	QNAHRKT (SEQ ID NO:182)	RSAHLSR (SEQ ID NO:26)	N7a* ELD
SBS# 60423 5'ggCACGCTG TTGAAGTCGG	TSGHLSR (SEQ ID NO:21)	DRSALAR (SEQ ID NO:34)	QSSNLAR (SEQ ID NO:94)	TGSLTR (SEQ ID NO:185)	QSSDLR (SEQ ID NO:15)	DRSNRNQ (SEQ ID NO:186)	N7a* KKR

Tgatggtga (SEQ ID NO:166), exon 4 (A)							
SBS# 64842 5'ggCACGCTG TTGAAGTCGG Tgatggtga (SEQ ID NO:166), exon 4 (A)	TSGHLSR (SEQ ID NO:21)	TSGSLTR (SEQ ID NO:185)	QSSNLAR (SEQ ID NO:94)	QSGSLTR (SEQ ID NO:67)	QSSDLSR (SEQ ID NO:15)	DRSNRNQ (SEQ ID NO:186)	N7a* KKR
SBS# 64844 5'ggCACGCTG TTGAAGTCGG Tgatggtga (SEQ ID NO:166), exon 4 (A)	TSGHLSR (SEQ ID NO:21)	TSGSLTR (SEQ ID NO:185)	QSSNLAR (SEQ ID NO:94)	TSGSLTR (SEQ ID NO:185)	QSSDLSR (SEQ ID NO:15)	DRSNRNQ (SEQ ID NO:186)	N7a* KKR
SBS# 64836 5'ggCACGCTG TTGAAGTCGG Tgatggtga (SEQ ID NO:166), exon 4 (A)	TSGHLSR (SEQ ID NO:21)	TSGSLTR (SEQ ID NO:185)	QSSNLAR (SEQ ID NO:94)	TSGSLTR (SEQ ID NO:185)	QSSDLSR (SEQ ID NO:15)	DRSNRNQ (SEQ ID NO:186)	N7a* KKR
SBS# 60416 5'caGAGCATC CCATGGAACC Tggagcgaa (SEQ ID NO:167) exon 3 (B)	HGQTLNE (SEQ ID NO:187)	QSGNLAR (SEQ ID NO:14)	RSDVLSN (SEQ ID NO:188)	DRSTRIT (SEQ ID NO:189)	LSWNLLT (SEQ ID NO:190)	RSANLTR (SEQ ID NO:57)	N7a* ELD
SBS# 64859 5'caGAGCATC CCATGGAACC Tggagcgaa (SEQ ID NO:167) exon 3 (B)	LQQTLAD (SEQ ID NO:191)	QSGNLAR (SEQ ID NO:14)	RSDVLSN (SEQ ID NO:188)	DRSTRIT (SEQ ID NO:189)	LKQNLDA (SEQ ID NO:159)	RSANLTR (SEQ ID NO:57)	N7a* ELD
SBS# 64848 5'caGAGCATC CCATGGAACC Tggagcgaa (SEQ ID NO:167) exon 3 (B)	HGQTLNE (SEQ ID NO:187)	QSGNLAR (SEQ ID NO:14)	RSDVLSN (SEQ ID NO:188)	DRSTRIT (SEQ ID NO:189)	LKQNLDA (SEQ ID NO:159)	RSANLTR (SEQ ID NO:57)	N7a* ELD
SBS# 64853 5'caGAGCATC CCATGGAACC Tggagcgaa (SEQ ID NO:167) exon 3 (B)	LQQTLAD (SEQ ID NO:191)	QSGNLAR (SEQ ID NO:14)	RSDVLSN (SEQ ID NO:188)	DRSTRIT (SEQ ID NO:189)	LKQNLDA (SEQ ID NO:159)	RSANLTR (SEQ ID NO:57)	N7a* ELD
SBS# 60417 5'ctGGTGCCAT GCTGGGATAA ttcgtcc (SEQ ID NO:168) exon 3 (B)	QSANRTK (SEQ ID NO:192)	DSSHRTR (SEQ ID NO:42)	RSDALSE (SEQ ID NO:193)	RSSTRKT (SEQ ID NO:194)	ERGTLAR (SEQ ID NO:160)	TSGSLTR (SEQ ID NO:185)	N7a* KKR
SBS# 64866 5'ctGGTGCCAT GCTGGGATAA ttcgtcc (SEQ ID NO:168) exon 3 (B)	QSANRTK (SEQ ID NO:192)	DSSHRTR (SEQ ID NO:42)	RSDVLSE (SEQ ID NO:19)	RKYSRLV (SEQ ID NO:195)	ERGTLAR (SEQ ID NO:160)	TSGSLTR (SEQ ID NO:185)	N7a* KKR
SBS# 64876 5'ctGGTGCCAT GCTGGGATAA ttcgtcc (SEQ ID NO:168) exon 3 (B)	QSANRTK (SEQ ID NO:192)	DSSHRTR (SEQ ID NO:42)	RSDALSE (SEQ ID NO:193)	RSSTRKT (SEQ ID NO:194)	ERGTLAR (SEQ ID NO:160)	TSGSLTR (SEQ ID NO:185)	N7a* KKR
SBS# 64869 5'ctGGTGCCAT GCTGGGATAA	QSANRTK (SEQ ID NO:192)	QSGHLSR (SEQ ID NO:22)	RSDALSE (SEQ ID NO:193)	RSSTRKT (SEQ ID NO:194)	ERGTLAR (SEQ ID NO:160)	TSGSLTR (SEQ ID NO:185)	N7a* KKR

ttcgtcc (SEQ ID NO:168) exon 3 (B)							
SBS# 60474 5'gaGCTGCGG CAGAGGCTGA Tccacttct (SEQ ID NO:169) exon 8 (D)	TSGNLTR (SEQ ID NO:36)	LSQDLNR (SEQ ID NO:196)	RSDNLAR (SEQ ID NO:52)	QNVSRPR (SEQ ID NO:197)	RSDDLTR (SEQ ID NO:37)	QSSDLRR (SEQ ID NO:71)	N6a* ELD
SBS# 64912 5'gaGCTGCGG CAGAGGCTGA Tccacttct (SEQ ID NO:169) exon 8 (D)	TSGNLTR (SEQ ID NO:36)	LSQDLNR (SEQ ID NO:196)	RSDNLAR (SEQ ID NO:52)	QSGDLTR (SEQ ID NO:39)	RSDSLSV (SEQ ID NO:125)	RSADLSR (SEQ ID NO:105)	N6a*
SBS# 60475 5'ttGATGACGT CTTTGGTAGA gaagtgga (SEQ ID NO:170) exon 8 (D)	QNAHRKT (SEQ ID NO:182)	LRHHLTR (SEQ ID NO:17)	TPSYLPT (SEQ ID NO:33)	DRSALAR (SEQ ID NO:34)	DRSNLSR (SEQ ID NO:91)	TSGNLTR (SEQ ID NO:36)	N6a* KKR
SBS# 64928 5'ttGATGACGT CTTTGGTAGA gaagtgga (SEQ ID NO:170) exon 8 (D)	QNAHRKT (SEQ ID NO:182)	LRHHLTR (SEQ ID NO:17)	SLTYLPT (SEQ ID NO:198)	DRSALAR (SEQ ID NO:34)	DRSNLSR (SEQ ID NO:91)	TSANLSR (SEQ ID NO:199)	N6a* KKR
SBS# 60412 5'gcTAAGTGc ATGGCTGTCT GGttctgta (SEQ ID NO:171) exon 3 (E)	RWQYLPT (SEQ ID NO:200)	DRSALAR (SEQ ID NO:34)	QSSDLR (SEQ ID NO:15)	RKDALVA (SEQ ID NO:201)	RSHSLR (SEQ ID NO:202)	QSANRTK (SEQ ID NO:192)	N7a* ELD
SBS# 64946 5'gcTAAGTGc ATGGCTGTCT GGttctgta (SEQ ID NO:171) exon 3 (E)	RSDHLST (SEQ ID NO:69)	DRSALAR (SEQ ID NO:34)	QSSDLR (SEQ ID NO:15)	RRDALLM (SEQ ID NO:203)	RSHSLR (SEQ ID NO:202)	QSANRTK (SEQ ID NO:192)	N7a* ELD
SBS# 64944 5'gcTAAGTGc ATGGCTGTCT GGttctgta (SEQ ID NO:171) exon 3 (E)	RSDHLST (SEQ ID NO:69)	DRSALAR (SEQ ID NO:34)	QSSDLR (SEQ ID NO:15)	RKDALVA (SEQ ID NO:201)	RSHSLR (SEQ ID NO:202)	QSANRTK (SEQ ID NO:192)	N7a* ELD
SBS# 64947 5'gcTAAGTGc ATGGCTGTCT GGttctgta (SEQ ID NO:171) exon 3 (E)	RSDHLSA (SEQ ID NO:180)	DRSALAR (SEQ ID NO:34)	QSSDLR (SEQ ID NO:15)	RKDALVA (SEQ ID NO:201)	RSHSLR (SEQ ID NO:202)	QSANRTK (SEQ ID NO:192)	N7a* ELD
SBS# 60413 5'caCATGGGG CAACTiCAGG GCctacaga (SEQ ID NO:172) exon 3 (E)	DSSHRTR (SEQ ID NO:42)	AKWNLDA (SEQ ID NO:204)	QHQLVLR (SEQ ID NO:44)	QNATRTK (SEQ ID NO:45)	RSDHLSR (SEQ ID NO:27)	TSSNRKT (SEQ ID NO:126)	N7a* KKR
SBS# 64951 5'caCATGGGG CAACTiCAGG GCctacaga (SEQ ID NO:172) exon 3 (E)	DSSHRTR (SEQ ID NO:42)	AKWNLDA (SEQ ID NO:204)	HASTLQN (SEQ ID NO:205)	QNATRTK (SEQ ID NO:45)	RSDHLSR (SEQ ID NO:27)	TSSNRKT (SEQ ID NO:126)	N7a* KKR
SBS# 64950 5'caCATGGGG CAACTiCAGG	DSSHRTR (SEQ ID NO:42)	AKWNLDA (SEQ ID NO:204)	HASTLQN (SEQ ID NO:205)	QNATRTK (SEQ ID NO:45)	RSDHLSR (SEQ ID NO:27)	TSSNRKT (SEQ ID NO:126)	N7a* KKR

GCctacaga (SEQ ID NO:172) exon 3 (E)							
SBS# 64952 5'caCATGGGG CAACTiCAGG GCctacaga (SEQ ID NO:172) exon 3 (E)	DRSHLTR (SEQ ID NO:59)	RSDNLR (SEQ ID NO:206)	QHQLVLR (SEQ ID NO:44)	QNATRTK (SEQ ID NO:45)	RSDHLSR (SEQ ID NO:27)	TSSNRKT (SEQ ID NO:126)	N7a* KKR
SBS# 60460 5'gtGGGACCT CACAGGCTGC Tgcccacgt (SEQ ID NO:173) exon 7 (F)	QSSDLR (SEQ ID NO:15)	HRSTRNR (SEQ ID NO:207)	RSDNLSQ (SEQ ID NO:208)	ASNDRKK (SEQ ID NO:209)	ASKTRTN (SEQ ID NO:210)	RSAHLR (SEQ ID NO:26)	N7a* ELD
SBS# 64971 5'gtGGGACCT CACAGGCTGC Tgcccacgt (SEQ ID NO:173) exon 7 (F)	QSSDLR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLSQ (SEQ ID NO:208)	ASNDRKK (SEQ ID NO:209)	DRSDLSR (SEQ ID NO:154)	RSHHLKA (SEQ ID NO:155)	N7a* ELD
SBS# 64969 5'gtGGGACCT CACAGGCTGC Tgcccacgt (SEQ ID NO:173) exon 7 (F)	QSSDLR (SEQ ID NO:15)	HRSTRNR (SEQ ID NO:207)	RSDNLSQ (SEQ ID NO:208)	ASNDRKK (SEQ ID NO:209)	DRSDLSR (SEQ ID NO:154)	RSHHLKA (SEQ ID NO:155)	N7a* ELD
SBS# 64968 5'gtGGGACCT CACAGGCTGC Tgcccacgt (SEQ ID NO:173) exon 7 (F)	QSSDLR (SEQ ID NO:15)	QSSDLRR (SEQ ID NO:71)	RSDNLSQ (SEQ ID NO:208)	ASNDRKK (SEQ ID NO:209)	DRSDLSR (SEQ ID NO:154)	RSHHLKA (SEQ ID NO:155)	N7a* ELD
SBS# 60461 5'ggGGTGGTG ACTCACCggc Cacgtgggc (SEQ ID NO:175) exon 7 (F)	DRSTRTK (SEQ ID NO:156)	RRDTLLD (SEQ ID NO:157)	QSADRTK (SEQ ID NO:68)	DRSNLTR (SEQ ID NO:99)	LRHHLTR (SEQ ID NO:17)	LKQHLTR (SEQ ID NO:211)	N7a* KKR
SBS# 64983 5'ggGGTGGTG ACTCACCggc Cacgtgggc (SEQ ID NO:175) exon 7 (F)	DRSTRTK (SEQ ID NO:156)	RRDTLLD (SEQ ID NO:157)	QSSDLR (SEQ ID NO:15)	DRSNLTR (SEQ ID NO:99)	HKQHRDA (SEQ ID NO:18)	LRHHLTR (SEQ ID NO:17)	N7a* KKR
SBS# 64982+ 5'ggGGTGGTG ACTCACCggc Cacgtgggc (SEQ ID NO:175) exon 7 (F)	DRSTRTK (SEQ ID NO:156)	RRDTLLD (SEQ ID NO:157)	QSSDLR (SEQ ID NO:15)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR
SBS# 64972+ 5'ggGGTGGTG ACTCACCggc Cacgtgggc (SEQ ID NO:175) exon 7 (F)	DRSTRTK (SEQ ID NO:156)	RRDTLLD (SEQ ID NO:157)	QSSDLR (SEQ ID NO:15)	DRSNLTR (SEQ ID NO:99)	LKQNLDA (SEQ ID NO:159)	LRHHLTR (SEQ ID NO:17)	N7a* KKR
SBS#60392 5'agGATGGAG ATTATGAAGA Gctgatgct (SEQ ID NO:176) exon 1 (G)	RSANLAR (SEQ ID NO:111)	QSANRTK (SEQ ID NO:192)	TSSNRKT (SEQ ID NO:126)	TSSNLSR (SEQ ID NO:24)	DRSHLSR (SEQ ID NO:212)	TSGNLTR (SEQ ID NO:36)	N7a* ELD
SBS# 64989 5'agGATGGAG ATTATGAAGA	RSANLAR (SEQ ID NO:111)	QSGNLAR (SEQ ID NO:14)	TSSNRKT (SEQ ID NO:126)	TSSNLSR (SEQ ID NO:24)	QSGHLQR (SEQ ID NO:149)	TSGNLTR (SEQ ID NO:36)	N7a* ELD

Gctgatgct (SEQ ID NO:176) exon 1 (G)							
SBS# 64992 5'agGATGGAG ATTATGAAGA Gctgatgct (SEQ ID NO:176) exon 1 (G)	RSANLAR (SEQ ID NO:111)	TSANLSR (SEQ ID NO:199)	TSSNRKT (SEQ ID NO:126)	TSSNLSR (SEQ ID NO:24)	QSGHLQR (SEQ ID NO:149)	TSGNLTR (SEQ ID NO:36)	N7a* ELD
SBS# 60393 5'tcCTCCTGGG ACGGgAGGGC Gagcatca (SEQ ID NO:177) exon 1 (G)	RSADLTR (SEQ ID NO:213)	RSDHLTQ (SEQ ID NO:214)	RSDHLSE (SEQ ID NO:30)	QSHHRKT (SEQ ID NO:215)	RSDVLSE (SEQ ID NO:19)	SPSSRRT (SEQ ID NO:216)	N7a* KKR
SBS# 65021 5'tcCTCCTGGG ACGGgAGGGC Gagcatca (SEQ ID NO:177) exon 1 (G)	RSADLTR (SEQ ID NO:213)	RSDHLTQ (SEQ ID NO:214)	RSDHLSE (SEQ ID NO:30)	QSGHLRS (SEQ ID NO:22)	RSDVLSE (SEQ ID NO:19)	SPSSRRT (SEQ ID NO:216)	N7a* KKR
SBS# 65022 5'tcCTCCTGGG ACGGgAGGGC Gagcatca (SEQ ID NO:177) exon 1 (G)	RSADLTR (SEQ ID NO:213)	RSDHLTQ (SEQ ID NO:214)	RSDHLSE (SEQ ID NO:30)	NSSSRIK (SEQ ID NO:217)	RSDVLSE (SEQ ID NO:19)	SPSSRRT (SEQ ID NO:216)	N7a* KKR
SBS# 58781 5'gtGGTGCTG ATGGAGGAG ACccagaggc (SEQ ID NO:178) exon 2 (J)	DRSNLSR (SEQ ID NO:91)	QSGHLRS (SEQ ID NO:22)	DRSHLSR (SEQ ID NO:212)	TSGNLTR (SEQ ID NO:36)	QSSDLSR (SEQ ID NO:15)	WHSSLH Q (SEQ ID NO:16)	L0 KKR
SBS# 65098 5'gtGGTGCTG ATGGAGGAG ACccagaggc (SEQ ID NO:178) exon 2 (J)	DRSNLSR (SEQ ID NO:91)	QSGHLRS (SEQ ID NO:22)	QSSHLTR (SEQ ID NO:13)	TSANLSR (SEQ ID NO:199)	QSSDLSR (SEQ ID NO:15)	TSGHLRS (SEQ ID NO:21)	L0 KKR
SBS# 65094 5'gtGGTGCTG ATGGAGGAG ACccagaggc (SEQ ID NO:178) exon 2 (J)	DRSNLSR (SEQ ID NO:91)	QSGHLRS (SEQ ID NO:22)	DRSHLSR (SEQ ID NO:212)	TSANLSR (SEQ ID NO:199)	QSSDLSR (SEQ ID NO:15)	TSGHLRS (SEQ ID NO:21)	L0 KKR
SBS# 65095 5'gtGGTGCTG ATGGAGGAG ACccagaggc (SEQ ID NO:178) exon 2 (J)	DRSNLSR (SEQ ID NO:91)	QSGHLRS (SEQ ID NO:22)	DRSHLSR (SEQ ID NO:212)	TSANLSR (SEQ ID NO:199)	QSSDLSR (SEQ ID NO:15)	HRHHLIR (SEQ ID NO:218)	L0 KKR
SBS# 58780 5'ctGGAGGCT GCCAGGAACC Tacattgtg (SEQ ID NO:179) exon 2 (J)	HGQTLNE (SEQ ID NO:187)	QSGNLAR (SEQ ID NO:14)	RSDNLSE (SEQ ID NO:95)	SKQYLIK (SEQ ID NO:148)	DRSHLTR (SEQ ID NO:59)	QSGHLRS (SEQ ID NO:22)	N7a* ELD
SBS# 65086 5'ctGGAGGCT GCCAGGAACC Tacattgtg (SEQ ID NO:191)	LQQTAD (SEQ ID NO:191)	QSGNLAR (SEQ ID NO:14)	RSDNLSE (SEQ ID NO:95)	SKQYLIK (SEQ ID NO:148)	DRSHLTR (SEQ ID NO:59)	QSGHLRS (SEQ ID NO:22)	N7a* ELD

ID NO:179) exon 2 (J)							
SBS# 65085 5'ctGGAGGCT GCCAGGAACC Tactattgtg (SEQ ID NO:179) exon 2 (J)	LQQTAD (SEQ ID NO:191)	QSGNLAR (SEQ ID NO:14)	RSDNLSE (SEQ ID NO:95)	RGDRRNK (SEQ ID NO:219)	DRSHLTR (SEQ ID NO:59)	QSGHLRS (SEQ ID NO:22)	N7a* ELD

\* indicates that the DNA binding domain is attached to the *FokI* nuclease domain at the DNA binding domain's N-terminus (see U.S. Publication Nos. 20170218349 and 20170211075). \*SBS# 64982 and SBS# 64972 have the same helices and *FokI* domains, but differ in the finger linkers; specifically in the linker between F2 and F3.

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**[0190]** Nucleases targeting the murine PCSK9 gene (mPCSK9) shown above were tested for activity in the murine liver cell line Hepa1-6 at 0.1 ug total ZFN mRNA dose, where half of the dose is each individual ZFN. ZFNs were added in a 100 µL transfection volume. All proteins bound to their targets and induced cleavage.

10 Exemplary activity (% indel) results of pairwise combinations are shown below in Tables 6A through 6G at the specific sites shown in Figure 6. % Indels were measured by deep sequencing (MiSeq, Illumina) and then analyzed for the percent of alleles comprising insertions and/or deletions at the cleavage site.

**Table 6A: mPCSK9 ZFN, Site A**

Site A	60423	64842	64844	64836
60422	16.4	35.0	26.8	26.7
64793	35.8	44.9	42.9	37.3
64802	30.6	43.0	41.6	37.0
64791	34.0	41.8	41.7	38.0

15

**Table 6B: mPCSK9 ZFN, Site B**

Site B	60417	64866	64876	64869
60416	13.9	35.7	32.4	30.9
64859	53.0	79.0	71.2	82.4
64848	36.2	73.6	64.8	73.0
64853	31.9	66.5	58.6	64.3

**Table 6C: mPCSK9 ZFN, Site D**

Site D	60475	64928
60474	0.1	0.1
64912	0.0	0.1

**Table 6D: mPCSK9 ZFN, Site E**

Site E	60413	64951	64950	64952
60412	22.8	28.6	20.6	23.5
64946	35.4	51.3	45.2	45.3
64944	41.4	48.8	38.7	49.1
64947	28.8	38.3	29.9	42.0

**Table 6E: mPCSK9 ZFN, Site F****G**

Site F	60461	64983	64982	64972
60460	27.2	50.5	53.8	23.7
64971	85.8	95.2	95.9	89.0
64969	69.1	79.4	86.1	69.8
64968	70.7	78.4	86.3	67.4

**Table 6F: mPCSK9 ZFN, Site**

Site G	60393	65021	65022
60392	53.7	68.1	69.2
64989	60.9	83.3	76.3
64992	51.4	79.8	69.1

**Table 6G: mPCSK9 ZFN, Site J**

Site J	58780	65086	65085
58781	40.5	62.3	54.1
65098	67.4	77.2	73.5
65094	69.6	77.8	73.3
65095	64.2	76.7	72.1

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**[0191]** Correlation between % indels and knockdown of secreted mPCSK9 protein in the transduced Hepa1-6 cell supernatant is calculated. Two doses are evaluated (Low dose is 0.5 ug total ZFN mRNA dose and high dose is 4 ug total ZFN mRNA dose), and mPCSK9 protein concentration is measured by ELISA at 3 days post transduction (“3DPT”). The analysis demonstrates a decrease in mPCSK9 concentration in the supernatant that correlates to the percent indels induced by the ZFN.

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## II. Nuclease targeting of SERPINA

**[0192]** Zinc finger nucleases were designed to cleave the SERPINA gene flanking the location of the Z mutation (Figure 1), Table 7, (see also Yusa *et al* (2011) *Nature* 478(7369):391-394). The zinc finger proteins were tested in HepG2 cells and K562 cells and the results are shown below in Table 8. “Linker” indicates that the DNA binding domain is attached to the *FokI* nuclease domain at the DNA binding domain’s N-terminus.

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**Table 7: SERPINA1-specific ZFN**

SBS #/ Target	F1	F2	F3	F4	F5	Linker
SBS#25264 5' gtCGATGGTCAGCAca	QSGSLTR (SEQ ID NO:67)	QSADRTK (SEQ ID NO:68)	RSDHLST (SEQ ID NO:69)	QSAHRIT (SEQ ID NO:70)	N/A	L0



gccttatgcacg (SEQ ID NO:65)						
SBS#25277 5'gaAAGGGActGAAGCT GCTggggccatg (SEQ ID NO:66)	QSSDLRR (SEQ ID NO:71)	QSSDLRS (SEQ ID NO:15)	QSGNLR (SEQ ID NO:14)	QSGHLAR (SEQ ID NO:72)	RLDNRT A (SEQ ID NO:73)	L0

**Table 8: Activity of SERPINA1-specific ZFN *in vitro***

Pair used, concentration	% Indels, HepG2 cells	% Indels, K562 cells
25264_25277_100ng	4.46%	23.03%
25264_25277_200ng	5.31%	40.91%
25264_25277_400ng	6.09%	40.68%
25264_25277_800ng	3.15%	27.02%

[0193] Exemplary ZFNs were then used in combination with an

- 5 oligonucleotide donor that would correct the G->A mutation responsible for the 'Z' mutation. In addition, the oligonucleotide comprised silent nucleotide alterations in the ZFN target sequences that would prevent cleavage of the oligonucleotide once integrated (Figure 1B). In particular, the nucleotides recognized by Fingers 2 and 4 (TCA and CGA, respectively) of SBS#25264 are altered such that these triplets are
- 10 TGA and CAA (variant SMS24). These new sequences will no longer be targeted by the SBS#25264 ZFN, and so once the oligonucleotide is integrated, cleavage will no longer occur. The SMS24 SERPINA1 variant oligonucleotide also comprised
- homology arms that were homologous to the SERPINA1 sequence flanking the cleavage site. When the SMS24 variant oligonucleotide was used with the
- 15 SERPINA1-specific ZFNs, targeted integration of the oligonucleotide occurred at approximately 4% of alleles in HepG2 cells and 25% of alleles in K562 cells (see Table 9).

**Table 9: ZFN mediated integration of corrective oligonucleotide *in vitro***

	HepG2		K562	
	% NHEJ	% TI	% NHEJ	% TI

SMS24 +ZFNs	15.42	4.36	28.71	24.88
ZFNs	12.75	0.00	41.13	0
SMS24	0.44	0.07	0.12	0.08

**[0194]** Next, the corrective oligonucleotide and SERPINA1-specific ZFNs were tested *in vivo* in PiZ mice, a strain comprising approximately 10 copies of the PiZ variant of the human SERPINA1 gene integrated into its germline (Carlson *et al* (1989) *J. Clin Invest.* 83: 1183-1190). PiZ mice are transgenic for the mutant human SERPINA1 (“SA1-ATZ”) and exhibit ATZ accumulation in hepatocytes and liver fibrosis. The SMS24 oligonucleotide and the ZFN pair were both delivered by AAV8 vectors via the intravenous route and groups of mice were sacrificed 2 weeks and 6 months after treatment for molecular, histological, and biochemical analyses. Untreated age/sex matched PiZ mice were used as controls. Table 10 shows the experimental outline and the injection schedules.

**Table 10: *In vivo* testing of ZFN-mediated integration of a corrective oligonucleotide donor**

Group	Test Article #1	Dose #1 (vg/mouse)	Volume/dose #1 (uL)	Test Article #1 Injections, mice	Test Article #2	Dose #2 (vg/mouse)	Volume/dose #2 (uL)	Test Article #2 Injections, mice	Sacrifice		Total
				Day 0				Day 0	2 weeks	6 months	
1	SMS24 AAV8	1.5E+12	200	6					3	3	6
2					25264/25277 AAV8	7.5E+10	200	6	3	3	6
3					25264/25277 AAV8	1.5E+11	200	6	3	3	6
4	SMS24 AAV8	1.5E+12	200	6	25264/25277 AAV8	7.5E+10	200	6	3	3	6
5	SMS24 AAV8	1.5E+12	200	6	25264/25277 AAV8	1.5E+11	200	6	3	3	6

**[0195]** Half of each cohort in the experiment was sacrificed two weeks following injection and the liver tissues were examined for the percent of alleles with NHEJ or targeted integration (TI) at the cleavage site. The results demonstrated that higher doses of ZFNs lead to increased cleavage activity (Figure 2A), and higher doses of ZFNs in the presence of the oligonucleotide donor lead to higher amounts of

TI (Figure 2B). At six months following injection, the second half of the cohorts were sacrificed. Liver analysis showed that there was a larger percent of alleles displaying NHEJ activity in all cohorts at six months than at two weeks (Figure 3C). The clear increased in the percent of NHEJ bearing cells may be indicative of a selective advantage for those cells bearing a knock out of the PiZ alleles. In contrast, the amount of TI detected comprising the corrective oligo was not significantly different than the two week samples (Figure 3B). More specifically, two weeks after treatment, deep sequencing of the hepatic SA1-ATZ gene pool showed 8+4% or 23+8% non-homologous end joining (NHEJ) respectively in mice receiving low dose or high dose rAAV-ZFN. When the rAAV-TI was co-administered with low dose or high dose of rAAV-ZFN, gene repair by targeted insertion (TI) of the normal AAT sequence occurred in 0.25+0.2% and 0.5+0.4 of SA1-ATZ genes. ZFN treatment reduced the number of PiZ globule-containing hepatocytes in liver at 6 months, indicating liver repopulation by genome-edited hepatocytes, as assayed by Diastase/periodic acid Schiff stain. At this time point, serum human ATZ levels declined by 30+6 and 40+5% in the low dose and high dose groups, respectively, compared with controls. Six months after treatment, the percentage of cells with NHEJ in the SA1-ATZ genes increased to 64+8% and 58+20% of recipients of low dose or high dose rAAV-ZFNs, respectively. In mice receiving high dose rAAV-ZFN plus rAAV-TI, up to 1.7% of SA1-ATZ genes showed gene correction. In parallel, serum ATZ levels declined by 47% and 70% in the low dose and high dose rAAV-ZFN recipients, respectively, and liver fibrosis, as measured by Sirius red staining, was greatly reduced compared with controls.

**[0196]** Thus, nuclease-mediated editing of the SA1-ATZ transgene *in vivo* appears to provide a proliferative advantage to PiZ mouse hepatocytes, allowing them to massively repopulate the liver and reverse hepatic fibrosis, indicating its use as a therapeutic for ATD.

### III. Nuclease targeting of TTR

**[0197]** TTR-specific zinc finger proteins were made to target murine and human TTR as described above. Several regions in the TTR gene were identified for potentially targeting (*see, e.g.*, Figure 4). Exemplary proteins as well as target sites and exemplary linkers and *FokI* mutations are shown below in Table 11.

Table 11: TTR-Specific ZFN designs

SBS #/ Target	F1	F2	F3	F4	F5	F6	Link er / Fok	reg ion
Mouse TTR								
SBS#59642 5'ctTTGCCTcG CTGGACTGgta tttgtgt (SEQ ID NO:74)	RSDVLSE (SEQ ID NO:19)	QSGHLSR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	QSSDLR (SEQ ID NO:15)	RLYTLHK (SEQ ID NO:90)	N/A	L0 / N- ELD	C Ex 1
SBS#59199 5'ccGCGGGGC CAGCTTCAG ACacaaatac (SEQ ID NO:75)	DRSNLSR (SEQ ID NO:91)	QSADRTK (SEQ ID NO:68)	QSSDLR (SEQ ID NO:15)	QRSTLKS (SEQ ID NO:92)	RSAHLSR (SEQ ID NO:26)	RSDDLTR (SEQ ID NO:37)	N7a / N- KKR	C Ex 1
SBS#59667 5'ccGCGGGGC CAGCTTCAG ACacaaatac (SEQ ID NO:75)	DRSNLSR( SEQ ID NO:91)	QSADRTK (SEQ ID NO:68)	QSSDLR (SEQ ID NO:15)	QRSTLKS (SEQ ID NO:92)	RSAHLSR (SEQ ID NO:26)	RSDDLTR (SEQ ID NO:37)	N7a / N- KKR	C Ex 1
SBS#61046 5'ctTTGCCTcG CTGGACTGgta tttgtgt (SEQ ID NO:74)	RSDVLSE (SEQ ID NO:19)	QSGHLSR (SEQ ID NO:22)	LSQDLNR( SEQ ID NO:93)	QSSDLR (SEQ ID NO:15)	RLYTLHK (SEQ ID NO:90)	N/A	N6a / N- ELD	C Ex 1
SBS#59192 5'ctTTGCCTcG CTGGACTGgta tttgtgt (SEQ ID NO:74)	RSDTLSE (SEQ ID NO:46)	QSGHLSR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	QSSDLR (SEQ ID NO:15)	RLYTLHK (SEQ ID NO:90)	N/A	N6a / N- ELD	C Ex 1
SBS#59771 5'gtGCCCAGG GTGCTGGAG AAtccaaatg (SEQ ID NO:76)	QSSNLAR( SEQ ID NO:94)	QSGHLSR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	TSGHLSR (SEQ ID NO:21)	RSDNLSE (SEQ ID NO:95)	ASKTRKN (SEQ ID NO:96)	N7a / N- ELD	H Ex 2
SBS#59245 5'agGACTTTG ACCATcAGAG GAcattgg (SEQ ID NO:77)	QSGHLAR( SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	GTQGLGI (SEQ ID NO:130)	DRSNLTR (SEQ ID NO:99)	N7a / N- KKR	H Ex 2
SBS#59244 5'gtGCCCAGG GTGCTGGAG AAtccaaatg (SEQ ID NO:76)	QSSNLAR( SEQ ID NO:94)	QSGHLSR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	WHSSLHQ (SEQ ID NO:16)	RSDNLSE (SEQ ID NO:95)	ASKTRKN (SEQ ID NO:96)	N7a / N- ELD	H Ex 2
SBS#59790 5'agGACTTTG ACCATcAGAG GAcattgg (SEQ ID NO:77)	QSGHLAR( SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	YRWLRN S (SEQ ID NO:100)	DRSNLTR (SEQ ID NO:99)	N7a / N- KKR	H Ex 2
SBS#59316 5'aaAAAGACc TCTGAGGGA TCCtgggagc (SEQ ID NO:78)	DSGGLSK( SEQ ID NO:101)	QSGHLSR (SEQ ID NO:22)	RSDNLAR( SEQ ID NO:52)	WRGDRVK (SEQ ID NO:102)	DRSNLSR (SEQ ID NO:91)	QRQNLVN (SEQ ID NO:103)	N7a / N- ELD	J

SBS#59317 5'ctTACCCAG AGGCAAAGgg ctcccagga (SEQ ID NO:79)	RSDNLSV (SEQ ID NO:104)	RSADLSR (SEQ ID NO:105)	RSDNLAR (SEQ ID NO:52)	QQQDRHK (SEQ ID NO:106)	DNSNRIK (SEQ ID NO:29)	N/A	N7a / N- KKR	J
SBS#61135 5'aaAAAGACc TCTGAGGGA TCCTgggagc (SEQ ID NO:78)	DSGGLSK (SEQ ID NO:101)	QSGHLSR (SEQ ID NO:22)	RSDNLAR (SEQ ID NO:52)	RSDNLAR (SEQ ID NO:52)	DRSNLSR (SEQ ID NO:91)	QRQNLVN (SEQ ID NO:103)	N7a / N- ELD	J
SBS#61137 5'aaAAAGACc TCTGAGGGA TCCTgggagc (SEQ ID NO:78)	DSSDRKK (SEQ ID NO:43)	QSGHLSR (SEQ ID NO:22)	RSDNLAR (SEQ ID NO:52)	WRGDRVK (SEQ ID NO:102)	DRSNLSR (SEQ ID NO:91)	QRQNLVN (SEQ ID NO:103)	N7a / N- ELD	J
Human TTR								
SBS#60489 5'gcAGAGGA GGAGCAGAC gatgagaagcc (SEQ ID NO:80)	DRSNLSR (SEQ ID NO:91)	QSGDLTR (SEQ ID NO:39)	QSGHLSR (SEQ ID NO:22)	QSGHLAR( SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	N/A	N7a / N- KKR	Ex 1
SBS#60488 5'atTCTTGGC AGGATGGCttc tcctcgtc (SEQ ID NO:81)	QYCCLTN (SEQ ID NO:107)	TSGNLTR (SEQ ID NO:36)	RSDNLST (SEQ ID NO:108)	FHSCLSA (SEQ ID NO:109)	RNSDRTK (SEQ ID NO:110)	N/A	N7a / N- ELD	Ex 1
SBS#57730 5'gaGGAGGA GCAGACGAT GAGaagccatc (SEQ ID NO:82)	RSANLAR (SEQ ID NO:111)	TSGNLTR (SEQ ID NO:36)	DRSNLSR (SEQ ID NO:91)	QSGDLTR (SEQ ID NO:39)	QSGHLAR (SEQ ID NO:72)	DRSHLAR (SEQ ID NO:112)	L0 / N- ELD	Ex 1
SBS#57731 5'ccTTGCTGG ACTGGTATTT Gtgtctgag (SEQ ID NO:83)	RPYTLRL (SEQ ID NO:113)	HRSNLNK (SEQ ID NO:114)	VSNLAC (SEQ ID NO:115)	DRSNLTR (SEQ ID NO:99)	RSDVLSE (SEQ ID NO:19)	RNFSLTMT (SEQ ID NO:116)	L0 / C- KKR	Ex 1
SBS#60602 5'caCATGCAC GGCCACATT GATggcagga (SEQ ID NO:84)	TSGNLTR (SEQ ID NO:36)	HKSARA A (SEQ ID NO:117)	YDYGRT (SEQ ID NO:118)	DRSHLAR (SEQ ID NO:112)	QSGDLTR (SEQ ID NO:39)	TSHNRNA (SEQ ID NO:119)	N7a / N- KKR	Ex 2
SBS#60601 5'taGATGCTgT CCGAGGCAgt cctgccat (SEQ ID NO:85)	QSGDLTR (SEQ ID NO:39)	RSDNLAR (SEQ ID NO:52)	DSSDRKK (SEQ ID NO:43)	QSSDLSR (SEQ ID NO:15)	TSGNLTR (SEQ ID NO:36)	N/A	N7a / N- ELD	Ex 2
SBS#60751 5'tgCATGCTC ATGGAATGG GGagatgccca (SEQ ID NO:86)	RSAHLSR (SEQ ID NO:26)	RSDALTQ (SEQ ID NO:120)	QSGHLAR (SEQ ID NO:72)	IRSNLLA (SEQ ID NO:121)	QSSDLSR (SEQ ID NO:15)	LRHNLRA (SEQ ID NO:122)	N6a / N- KKR	Ex 3
SBS#60750 5'ctTACTGGA AGGCACTTgg catctcccc (SEQ ID NO:87)	ARSTRIT (SEQ ID NO:123)	QSGSLTR (SEQ ID NO:67)	RSDNLSV (SEQ ID NO:104)	RNAHRIN (SEQ ID NO:124)	DNSNRIK (SEQ ID NO:29)	N/A	N6a / ELD	Ex 3
SBS#60764 5'gcTCATGGA ATGGGGAGA TGccaagtgc	RSDLSLV (SEQ ID NO:125)	RSANLTR (SEQ ID NO:57)	RSDHLSR (SEQ ID NO:27)	TSSNRKT (SEQ ID NO:126)	RSDHLSQ (SEQ ID NO:41)	QSADRTK (SEQ ID NO:68)	L0 / N- ELD	Ex 3

(SEQ ID NO:88)								
SBS#60765 5'gcAGAGGTG AGTATACAG ACcttcgagg (SEQ ID NO:89)	DRSNLSR (SEQ ID NO:91)	QKVTLA A (SEQ ID NO:127)	TSSNRKT (SEQ ID NO:126)	RSDDLRSR (SEQ ID NO:128)	LRHHLTR (SEQ ID NO:17)	QSAHLKA (SEQ ID NO:129)	L0 / C- KKR	Ex 3
SBS# 60502 5'ctGCCTTGC TGGACTGGtatt tgtgtct (SEQ ID NO:220)	VSNLAC (SEQ ID NO:115)	DRSNLTR (SEQ ID NO:99)	RSDVLSE (SEQ ID NO:19)	RNFSLTM (SEQ ID NO:116)	HRKSLSR (SEQ ID NO:223)	N/A	L8c4 / KKR	Ex 1
SBS# 64066 5'gaGCAGACG ATGAGAAAGcc atcctgcc (SEQ ID NO:221)	AHGARWN (SEQ ID NO:224)	RSANLTR (SEQ ID NO:57)	TSGNLTR (SEQ ID NO:36)	DRSNLSR (SEQ ID NO:91)	QSGDLTR (SEQ ID NO:39)	N/A	L8c4 / ELD	Ex 1
SBS# 67451 5'gaGCAGACG ATGAGAAAGcc atcctgcc (SEQ ID NO:221)	AHGARWN (SEQ ID NO:224)	RSANLTR (SEQ ID NO:57)	TSANLSR (SEQ ID NO:199)	DRSNLSR (SEQ ID NO:91)	QSSDLTR (SEQ ID NO:184)	N/A	L8c4 / ELD	Ex 1
SBS# 67458 5'ctGCCTTGC TGGACTGGtatt tgtgtct (SEQ ID NO:220)	VSNLAC (SEQ ID NO:115)	DRSNLTR (SEQ ID NO:99)	RSDTLSE (SEQ ID NO:46)	RRWSLSV (SEQ ID NO:225)	DRSTRTK (SEQ ID NO:156)	N/A	L8c4 / KKR	Ex 1
SBS# 60501 5'gaGCAGACG ATGAGAAAGcc atcctgcc (SEQ ID NO:221)	AHGARWN (SEQ ID NO:224)	RSANLTR (SEQ ID NO:57)	TSGNLTR (SEQ ID NO:36)	DRSNLTR (SEQ ID NO:99)	QSGDLTR (SEQ ID NO:39)	N/A	L8c4 / ELD	Ex 1
SBS# 64080 5'ctGCCTTGC TGGACTGGtatt tgtgtct (SEQ ID NO:220)	VSNLAC (SEQ ID NO:115)	DRSNLTR (SEQ ID NO:99)	RSDVLSE (SEQ ID NO:19)	RNFSLTM (SEQ ID NO:116)	DRSTRTK (SEQ ID NO:156)	N/A	L8c4 / KKR	Ex 1
SBS# 67495 5'tgCATGCTC ATGGAATGG GGagatgcc (SEQ ID NO:86)	RSDHLST (SEQ ID NO:69)	RSDARTN (SEQ ID NO:226)	QSGHLAR (SEQ ID NO:72)	IRSNLLA (SEQ ID NO:121)	QSSDLRSR (SEQ ID NO:15)	LKWNLRT (SEQ ID NO:227)	N6a* / KKR	Ex 3
SBS# 60750 5'ctTACTGGA AGGCACTTgg catctccc (SEQ ID NO:87)	ARSTRIT (SEQ ID NO:123)	QSGSLTR (SEQ ID NO:67)	RSDNLSV (SEQ ID NO:104)	RNAHRIN (SEQ ID NO:124)	DNSNRIK (SEQ ID NO:29)	N/A	N6a* / ELD	Ex 3
SBS# 67493 5'tgCATGCTC ATGGAATGG GGagatgcc (SEQ ID NO:86)	RSDHLST (SEQ ID NO:69)	RSDARTN (SEQ ID NO:226)	QSGHLAR (SEQ ID NO:72)	IRSNLLA (SEQ ID NO:121)	QSSDLRSR (SEQ ID NO:15)	LRHNLRA (SEQ ID NO:122)	N6a* / KKR	Ex 3
SBS# 64347 5'tcTACTGgA AGGCACTTgg catctccc (SEQ ID NO:222)	YTYSLSE (SEQ ID NO:228)	QSGDLTR (SEQ ID NO:39)	RKDQLVA (SEQ ID NO:229)	RSDVLSE (SEQ ID NO:19)	QRTPRAK (SEQ ID NO:230)	N/A	N6a* / ELD	Ex 3
SBS# 60751 5'tgCATGCTC ATGGAATGG GGagatgcc (SEQ ID NO:86)	RAHLSR (SEQ ID NO:26)	RSDALTQ (SEQ ID NO:231)	QSGHLAR (SEQ ID NO:72)	IRSNLLA (SEQ ID NO:121)	QSSDLRSR (SEQ ID NO:15)	LRHNLRA (SEQ ID NO:122)	N6a* / KKR	Ex 3

SBS# 67489 5'tcTTACTGgA AGGCACTTgg catctccc (SEQ ID NO:222)	YTYSLSE (SEQ ID NO:228)	QSGDLTR (SEQ ID NO:39)	RKDQLVA (SEQ ID NO:229)	RSDVLSE (SEQ ID NO:19)	QRTPRAK (SEQ ID NO:230)	N/A	N6a* / ELD	Ex 3
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**[0198]** All murine specific nucleases were tested in B16-F10 cells (Figure 5A), primary mouse hepatocytes (Figure 5B), or Hepa1-6 cells (Figures 8A and 8B) and found to be active. Lead ZFNs targeting mTTR exon 1, Site C (59919/59642) and exon 2, Site H (59771/59245), were cloned into individual AAV expression cassette vectors containing a human ApoE enhancer and human SERPINA1 promoter and then packaged into serotype AAV8. AAV ZFNs were then injected intravenously into wildtype C57BL6 mice at a high dose (1.5e11 vg per ZFN per mouse) or a low dose (2.5e10 vg per ZFN per mouse) and livers were harvested for gene modification and plasma was analyzed for mTTR protein knockdown 28 days later. Figure 6A shows robust editing within the liver, while Figure 6B shows high levels of circulating mTTR protein knockdown which is specific to the mTTR-targeted ZFN AAV vectors

**[0199]** All human specific ZFNs were tested in the human liver cell line HepG2 and found to be active (see Table 12 below for activity of exemplary pairs).

In brief, total ZFN mRNA dose, where half of the dose is each individual ZFN, is shown in Table 12 and was as added in a 100  $\mu$ L transfection volume. All proteins bound to their targets and induced cleavage. Exemplary activity (% indel) results of pairwise combinations of ZFNs which bind exon 1 and exon 3 of the gene. % Indels were measured by deep sequencing (MiSeq, Illumnia) and then analyzed for the percent of alleles comprising insertions and/or deletions at the cleavage site.

**Table 12: Activity of Human TTR reagents**

<u>Sample</u>	<u>mRNA Dose (ug/mL)</u>		<u>Exon</u>
	<u>5</u>	<u>40</u>	
64066/60502	19.0	37.3	1
67451/60502	17.9	36.6	1
67458/60501	27.4	47.6	1
64080/60501	24.2	47.0	1
67495/60750	30.0	61.8	3
67493/60750	24.8	59.2	3
64347/60751	25.7	63.5	3
67489/60751	19.3	62.2	3

**[0200]** To determine the specificity of the ZFNs which bind the murine TTR gene (mTTR), ZFN pairs were subjected to unbiased identification of candidate off-target sites using methods similar to those previously described (Tsai *et al* (2015) *Nat Biotechnol* 33:187-197) in B16-F10 cells. Briefly, B16-F10 cells were electroporated with mRNA encoding the ZFNs as well as barcoded ssDNA oligos using the BTX electroporation device to allow for unbiased identification of sites which have undergone double-stranded DNA cleavage and NHEJ-mediated integration of the ssDNA oligos. The top 24 sites found identified by Miseq next-generation sequencing (NGS) to contain integrated oligos were then confirmed in primary mouse hepatocyte cells which were transduced with the ZFN mRNA via lipofection (Figure 5B). Genomic DNA from ZFN-treated mouse hepatocytes was amplified by PCR generating amplicons of approximately 200 bp surrounding the potential ZFN binding site. In order to build an equimolar library, PCR products were quantified with KAPA Library Quantification Kit for Illumina sequencing platforms (KAPABIOSYSTEMS) on C1000 Thermal Cycler (BIO-RAD) and sequenced on MiSeq Illumina Platform using MiSeq Reagent v.3 (Illumina). Quantification of insertions and deletions and insertions (indels) was then performed. Briefly, raw paired-end reads were joined and aligned to the specific genomic target sequences. Sequences with indels of  $\geq 1$  bp located within a 40 bp region encompassing the ZFN target site were considered as nuclease-induced genome modifications. All sites that produced a Bonferroni p-value  $\leq 0.05$  in comparison to a GFP-encoding mRNA electroporated control, were deemed off-target sites.

**[0201]** Modifications to residues within the ZFP which undergo non-specific binding to genomic DNA, in both the zinc finger backbone and in the *FokI* domain, were mutated in several ZFNs (R to Q changes at 1 to 6 residues) as well as mutations to the *FokI* which either affect catalytic activity or potentially non-specific DNA binding (U.S. Patent Application 15/685,580). The mutations made are shown below in Table 13 below.

**Table 13: Backbone/*FokI* variants of SBS#59771/SBS#59790**

SBS # (target site, 5'- 3')	Design [Helix Sequence, SEQ ID]						Linker
	[Mutations to finger backbone]						Fok mutants
	F1	F2	F3	F4	F5	F6	



<b>59771</b> 5'gtGCCCAGGG TGCTGGAGAA ccaaatg (SEQ ID NO:76)	QSSNLAR (SEQ ID NO:94)	QSGHLR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	TSGHLR (SEQ ID NO:21)	RSDNLSE (SEQ ID NO:95)	ASKTRKN (SEQ ID NO:96)	N7a
	none	none	none	none	none	none	ELD
<b>69063</b> 5'gtGCCCAGGG TGCTGGAGAA ccaaatg (SEQ ID NO:76)	QSSNLAR (SEQ ID NO:94)	QSGHLR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	TSGHLR (SEQ ID NO:21)	RSDNLSE (SEQ ID NO:95)	ASKTRKN (SEQ ID NO:96)	N7a
	Qm5	none	Qm5	none	Qm5	Qm5	ELD G480S
<b>69052</b> 5'gtGCCCAGGG TGCTGGAGAA ccaaatg (SEQ ID NO:76)	QSSNLAR (SEQ ID NO:94)	QSGHLR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	TSGHLR (SEQ ID NO:21)	RSDNLSE (SEQ ID NO:95)	ASKTRKN (SEQ ID NO:96)	N7a
	none	Qm5	none	Qm5	none	Qm5	ELD D421S
<b>69121</b> 5'gtGCCCAGGG TGCTGGAGAA ccaaatg (SEQ ID NO:76)	QSSNLAR (SEQ ID NO:94)	QSGHLR (SEQ ID NO:22)	QSSDLR (SEQ ID NO:15)	TSGHLR (SEQ ID NO:21)	RSDNLSE (SEQ ID NO:95)	ASKTRKN (SEQ ID NO:96)	N7a
	none	none	none	none	none	none	ELD D421S, Q531R
<b>59790</b> 5'agGACTTTGA CCATcAGAGG Acatttg (SEQ ID NO:77)	QSGHLAR (SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	YRWLRN S (SEQ ID NO:100)	DRSNLTR (SEQ ID NO:99)	N7a
	none	none	none	none	none	none	KKR
<b>69110</b> 5'agGACTTTGA CCATcAGAGG Acatttg (SEQ ID NO:77)	QSGHLAR (SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	YRWLRN S (SEQ ID NO:100)	DRSNLTR (SEQ ID NO:99)	N7a
	Qm5	none	Qm5	Qm5	Qm5	Qm5	KKR S418P
<b>69107</b> 5'agGACTTTGA CCATcAGAGG Acatttg (SEQ ID NO:77)	QSGHLAR (SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	YRWLRN S (SEQ ID NO:100)	DRSNLTR (SEQ ID NO:99)	N7a
	Qm5	none	Qm5	Qm5	Qm5	Qm5	KKR K387S
<b>69102</b> 5'agGACTTTGA CCATcAGAGG Acatttg (SEQ ID NO:77)	QSGHLAR (SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	YRWLRN S (SEQ ID NO:100)	DRSNLTR (SEQ ID NO:99)	N7a
	none	Qm5	none	Qm5	none	Qm5	KKR S418P
<b>69128</b> 5'agGACTTTGA CCATcAGAGG Acatttg (SEQ ID NO:77)	QSGHLAR (SEQ ID NO:72)	QLTHLNS (SEQ ID NO:97)	SKLYLNN( SEQ ID NO:98)	DRSNLTR( SEQ ID NO:99)	YRWLRN S (SEQ ID NO:100)	DRSNLTR (SEQ ID NO:99)	N7a
	none	none	none	none	none	none	KKR D421S, Q481H

**[0202]** These variants were then tested for on target and off target cleavage activity. Figure 8A shows the on-target activity in Hepa1-6 cells of the lead mutant ZFNs originating from the 59771/59790 pair. Figure 8B shows the indel sum of the top 3 identified off-target sites for these lead ZFNs, demonstrating a decrease in off-target cleavage for the mutant ZFNs.

## IV. Nuclease targeting of HAO1

**[0203]** HAO1-specific zinc finger proteins were made to target *Macaca mulatta* HAO1 as described above. Exemplary proteins are shown below in Table 14.

5

**Table 14: HAO1 ZFN designs for *M. mulatta***

SBS #/ Target	F1	F2	F3	F4	F5	F6	Linker / Fok
SBS# 66075 5'acTTGCTGC ATATGtGGCT AAagcaata (SEQ ID NO:232)	QLGNLHT (SEQ ID NO:234)	DRSHLAR (SEQ ID NO:112)	RSDVLST (SEQ ID NO:235)	ASGNLLN (SEQ ID NO:236)	RSDVLSE (SEQ ID NO:19)	RKYSLRV (SEQ ID NO:195)	N6a* ELD
SBS# 66079 5'acTTGCTGC ATATGtGGCT AAagcaata (SEQ ID NO:232)	QLGNLHT (SEQ ID NO:234)	DRSHLAR (SEQ ID NO:112)	RSDVLST (SEQ ID NO:235)	ASGNLLN (SEQ ID NO:236)	RSDTLSE (SEQ ID NO:46)	RRWSLSV (SEQ ID NO:225)	N6a* ELD
SBS# 64555 5'tcCCAGCTG ATAGATGGG TCtattgctt (SEQ ID NO:233)	DRSALAR (SEQ ID NO:34)	RKYYLAK (SEQ ID NO:237)	QSAHLVC (SEQ ID NO:238)	TSGNLTR (SEQ ID NO:36)	QSSDLSE (SEQ ID NO:15)	QRSTLKS (SEQ ID NO:92)	N6a* KKR
SBS# 66083 5'tcCCAGCTG ATAGATGGG TCtattgctt (SEQ ID NO:233)	DRSALAR (SEQ ID NO:34)	RKYYLAK (SEQ ID NO:237)	QSAHLVC (SEQ ID NO:238)	TSGNLTR (SEQ ID NO:36)	QSSDLSE (SEQ ID NO:15)	QRSTLKS (SEQ ID NO:92)	N7a* KKR
SBS# 66084 5'tcCCAGCTG ATAGATGGG TCtattgctt (SEQ ID NO:233)	DRSALAR (SEQ ID NO:34)	RKYYLAK (SEQ ID NO:237)	QSAHLVC (SEQ ID NO:238)	TSGNLTR (SEQ ID NO:36)	QSSDLSE (SEQ ID NO:15)	QRSTLKS (SEQ ID NO:92)	N6a* KKR
SBS# 64552 5'acTTGCTGC ATATGtGGCT AAagcaata (SEQ ID NO:232)	QLGNLHT (SEQ ID NO:234)	DRSHLAR (SEQ ID NO:112)	RSDVLST (SEQ ID NO:235)	ASGNLLN (SEQ ID NO:236)	RSDVLSE (SEQ ID NO:19)	RNFSLTM (SEQ ID NO:116)	N6a* ELD

**[0204]** Activity was analyzed as described above, and is shown below in Table 15. Total ZFN mRNA dose means that half of the dose is each individual ZFN, and was as added in a 100  $\mu$ L transfection volume. All proteins bound to their targets and induced cleavage. Exemplary activity (% indel) results of pairwise combinations of ZFNs which bind exon 4 of the gene. % Indels were measured by deep sequencing (MiSeq, Illumina) and then analyzed for the percent of alleles comprising insertions and/or deletions at the cleavage site.

10

**Table 15: Activity of *M. mulatta* HAO1 ZFNs**

	% Indels	
<b>Sample (mRNA Dose (ug/mL))</b>	<b>5</b>	<b>20</b>
66075/64555	86.9	74.8
66079/64555	84.2	75.7
66083/64552	80.6	76.9
66084/64552	80.6	78.2

**V. Nuclease targeting of KLKB1**

**[0205]** KLKB1-specific zinc finger proteins were made to target murine

- 5 KLKB1 as described above, shown below in Table 16. Several exons in the KLKB1 gene were selected for targeting in mouse liver cell line Hepa1-6.

**Table 16: Design of murine KLKB1 ZFNs**

SBS #/ Target	F1	F2	F3	F4	F5	F6	Linker / Fok	Exon
SBS# 63342 5'ctGCTCCTT GCACGAAGG TCacattcag (SEQ ID NO:239)	DRSALSR (SEQ ID NO:251)	RLDNRTA (SEQ ID NO:73)	RSDTLSE (SEQ ID NO:46)	QKRNRTK (SEQ ID NO:151)	IRSTLRD (SEQ ID NO:252)	HRSSLRR (SEQ ID NO:253)	N7a* KKR	Ex 9
SBS# 63341 5'ttGACTTTGA AGGGGAAGA Actgaatgt (SEQ ID NO:240)	QSANRTK (SEQ ID NO:192)	QSGNLAR (SEQ ID NO:14)	RSDHLSE (SEQ ID NO:30)	QSGNLAR (SEQ ID NO:14)	GTQGLGI (SEQ ID NO:130)	DRSNLTR (SEQ ID NO:99)	N7a* ELD	Ex 9
SBS# 63238 5'gcAGAGTGC TGGAAAATatc catggggc (SEQ ID NO:241)	TSSNRKT (SEQ ID NO:126)	QSGHLSR (SEQ ID NO:22)	RSDVLSE (SEQ ID NO:19)	RLYTLHK (SEQ ID NO:254)	QNAHRKT (SEQ ID NO:182)	N/A	N7a* KKR	Ex 7
SBS# 63236 5'ccGCTGAGtT CACAGGTTG CCcattgga (SEQ ID NO:242)	ERGTLAR (SEQ ID NO:160)	TSGSLTR (SEQ ID NO:185)	RSDNLSQ (SEQ ID NO:208)	ASNDRKK (SEQ ID NO:209)	RSANLAR (SEQ ID NO:111)	QSSDLRR (SEQ ID NO:71)	N7a* ELD	Ex 7
SBS# 63195 5'ttGTCGGTCT TAGAGATATT aaagtgg (SEQ ID NO:243)	YSWTLRD (SEQ ID NO:255)	TSGNLTR (SEQ ID NO:36)	QNAHRKT (SEQ ID NO:182)	STAALSY (SEQ ID NO:256)	TSGHLSR (SEQ ID NO:21)	DRSALAR (SEQ ID NO:34)	N7a* KKR	Ex 5
SBS# 63194 5'aaGGACTTG ATATGaGAGG GTccaactt (SEQ ID NO:244)	TSGHLSR (SEQ ID NO:21)	RSDNLTR (SEQ ID NO:40)	RSDTLSQ (SEQ ID NO:257)	ASANRTK (SEQ ID NO:258)	YTYSLSE (SEQ ID NO:228)	QSGHLSR (SEQ ID NO:22)	N7a* ELD	Ex 5
SBS# 63411 5'ttCACTTGCA GGCTGACctgc	DRSNLSR (SEQ ID NO:91)	LRQDLKR (SEQ ID NO:259)	RSDNLST (SEQ ID NO:108)	RQWSLRI (SEQ ID NO:260)	DRSNRTT (SEQ ID NO:261)	N/A	N6a* KKR	Ex 11

catggcc (SEQ ID NO:245)								
SBS# 63410 5'tcTTTAGGG GAGTGGCCatg gcaggcca (SEQ ID NO:246)	DRSTRTK (SEQ ID NO:156)	RSDSLLR (SEQ ID NO:262)	QRCHLTK (SEQ ID NO:263)	RSDHLSQ (SEQ ID NO:41)	STAALSY (SEQ ID NO:256)	N/A	N6a* ELD	Ex 11
SBS# 63375 5'tgCATGCCA TAGGTGATCC Tagttgggg (SEQ ID NO:247)	HNHDLRN (SEQ ID NO:264)	TSGNLTR (SEQ ID NO:36)	LRHHLTR (SEQ ID NO:17)	QKWPRDS (SEQ ID NO:265)	DRSDLSR (SEQ ID NO:154)	LRFNLNRN (SEQ ID NO:266)	N7a* KKR	Ex 10
SBS# 63374 5'ggGAGCCAT CTGTGGATA ACcttaagga (SEQ ID NO:248)	DRSNRTT (SEQ ID NO:261)	TNSNRKR (SEQ ID NO:267)	RSDSLLR (SEQ ID NO:262)	LKQDRRK (SEQ ID NO:268)	IKHDLHR (SEQ ID NO:269)	RSANLTR (SEQ ID NO:57)	L0 ELD	Ex 10
SBS# 63107 5'acGGCGAGA AAGCTGAACA gcagccacc (SEQ ID NO:249)	QSGNLAR (SEQ ID NO:14)	QSSDLR (SEQ ID NO:15)	QSGNRTT (SEQ ID NO:270)	RSANLAR (SEQ ID NO:111)	DRSHLAR (SEQ ID NO:112)	N/A	N7a* KKR	Ex 3
SBS# 63106 5'ggCACCTGg GGTGAAaAGT GCacattt (SEQ ID NO:250)	QSGDLTR (SEQ ID NO:39)	LRHQLKS (SEQ ID NO:271)	QRSNLVR (SEQ ID NO:272)	TSGHLR (SEQ ID NO:21)	RSDVLST (SEQ ID NO:235)	DTRNLRA (SEQ ID NO:273)	L0 ELD	Ex 3

**[0206]** All nucleases shown bound to their target sites and were active. Table 17 below shows the activity of the indicated pairs, indicating the individual ZFN mRNA dose, where each ZFN mRNA was used in a 1:1 mass ratio that was as added in a 100  $\mu$ L transfection volume. All proteins bound to their targets and induced cleavage. Exemplary activity (% indel) results of pairwise combinations of ZFNs which bind various coding regions of the gene. % Indels were measured by deep sequencing (MiSeq, Illumina) and then analyzed for the percent of alleles comprising insertions and/or deletions at the cleavage site.

**Table 17: Activity of murine KLKB1 ZFN**

Activity (% Indels)			
		mRNA per ZFN ( $\mu$ g/mL)	
Locus	Sample	5	40
Exon 9	63342/63341	64.6129	96.7541
Exon 7	63238/63236	45.2949	92.497
Exon 5	63195/63194	17.8121	89.8445
Exon 11	63411/63410	15.8158	84.6924
Exon 10	63375/63374	11.8068	93.6912

**[0207]** All patents, patent applications and publications mentioned herein are hereby incorporated by reference in their entirety.

**[0208]** Although disclosure has been provided in some detail by way of illustration and example for the purposes of clarity of understanding, it will be  
5 apparent to those skilled in the art that various changes and modifications can be practiced without departing from the spirit or scope of the disclosure. Accordingly, the foregoing descriptions and examples should not be construed as limiting.

CLAIMS

What we claim is:

1. A liver cell comprising a zinc finger protein that modulates expression of an endogenous Serpin Family A Member 1 (SERPINA1) gene at the Z mutation locus and further wherein

the zinc finger protein that modulates SERPINA1 gene expression comprises 4 or 5 zinc finger domains ordered F1 to F4 or F1 to F5, each zinc finger domain comprising a recognition helix region as shown in a single row of the following Table:

F1	F2	F3	F4	F5
QSGSLTR (SEQ ID NO:67)	QSADRTK (SEQ ID NO:68)	RSDHLST (SEQ ID NO:69)	QSAHRIT (SEQ ID NO:70)	N/A
QSSDLRR (SEQ ID NO:71)	QSSDLSR (SEQ ID NO:15)	QSGNLA R (SEQ ID NO:14)	QSGHLA R (SEQ ID NO:72)	RLDNRT A (SEQ ID NO:73)

and wherein the liver cell comprises an oligonucleotide donor that corrects the G->A mutation at the Z mutation locus and wherein expression of an endogenous SERPINA1 gene is altered as compared to wild-type.

2. The liver cell of claim 1, wherein the sequence of the endogenous gene is altered by cleaving the gene using at least one zinc finger nuclease comprising a pair of zinc finger binding proteins.

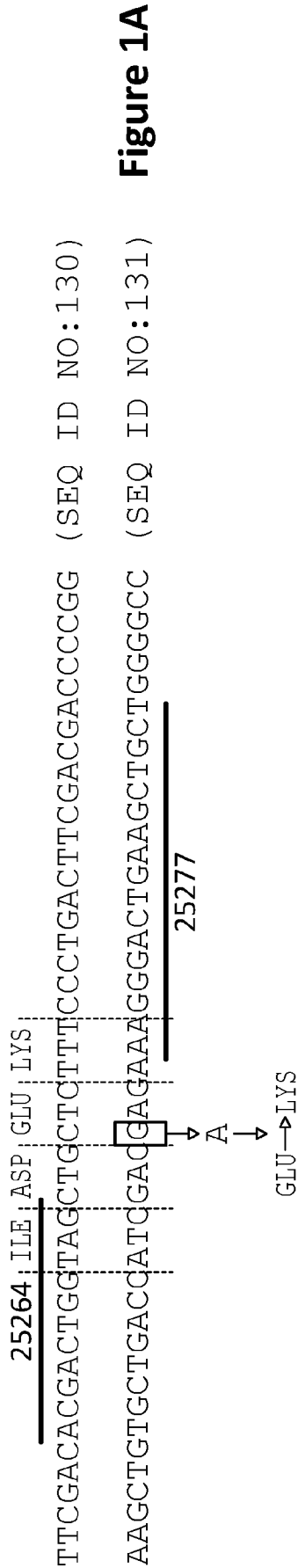
3. The liver cell of claim 2, wherein the zinc finger nuclease cleaves a human or mouse gene.

4. A pharmaceutical composition comprising a liver cell according to any one of claims 1 to 3.

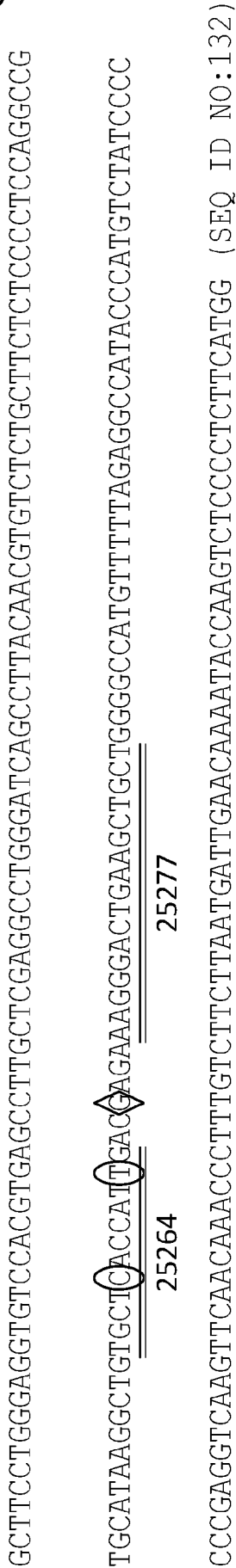
5. The pharmaceutical composition according to claim 4 when used to treat A1AT Deficiency and/or Hereditary Angioedema.

6. A kit comprising the liver cell according to any one of claims 1 to 3.

7. A genetically modified cell descended from the liver cell according to any one of claims 1 to 3, and that retains the zinc finger protein and the genetic modification at the Z mutation locus in the SERPINA1 gene.
8. The liver cell of any one of claims 1 to 3, wherein the zinc finger protein is encoded by a polynucleotide, wherein the polynucleotide is mRNA form or is carried on a viral or non-viral vector.
9. A method of treating A1AT Deficiency and/or Hereditary Angioedema in a subject in need thereof, comprising administering to the subject a liver cell according to any one of claims 1 to 3 or the pharmaceutical composition of claim 4.
10. Use of a liver cell according to any one of claims 1 to 3 in the preparation of a medicament for the treatment of A1AT Deficiency and/or Hereditary Angioedema.



1/9



O = NUCLEOTIDES ALTERED IN OLIGONUCLEOTIDE DONOR

X = ZFN TARGET SITE

◇ = SITE OF Z MUTATION

**Figure 1B**



2/9

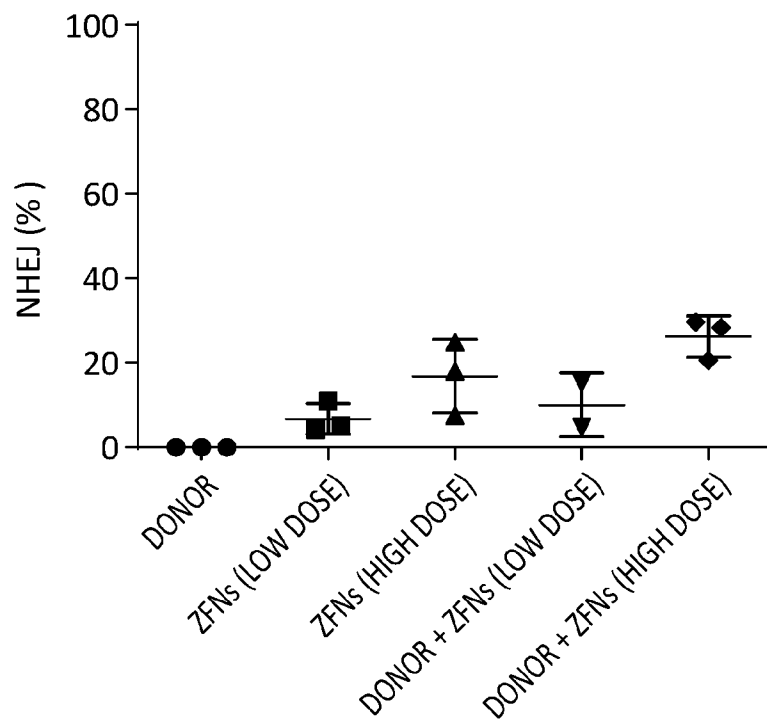
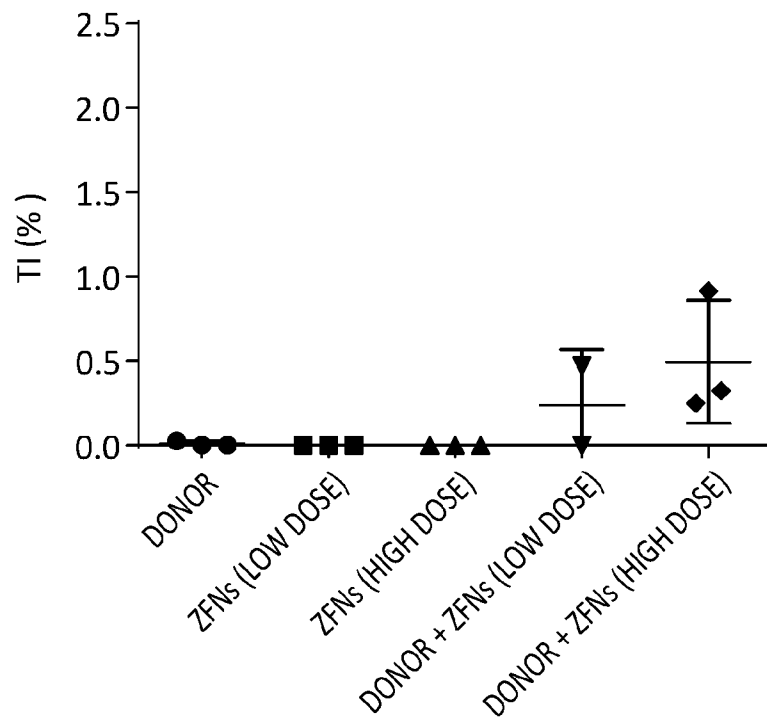
**Figure 2A****Figure 2B**

Figure 3C

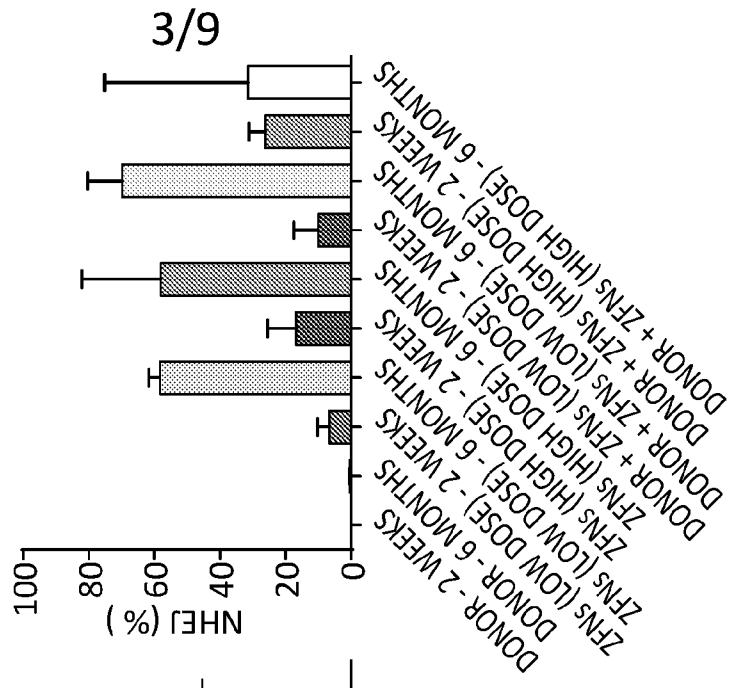


Figure 3B

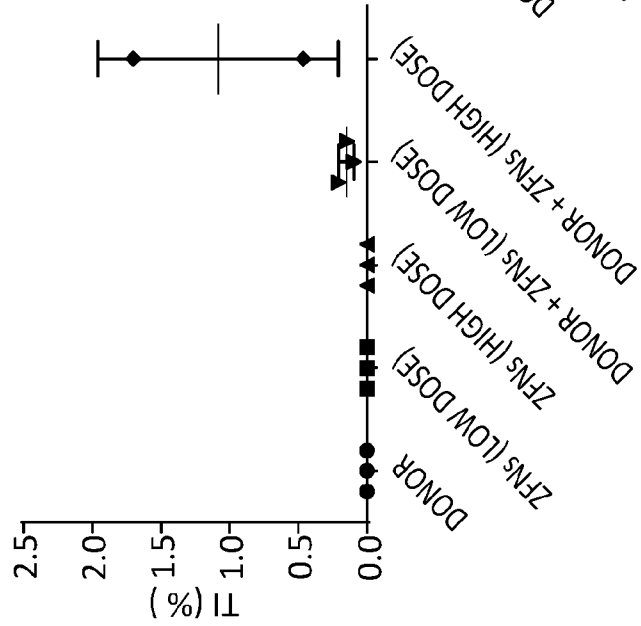
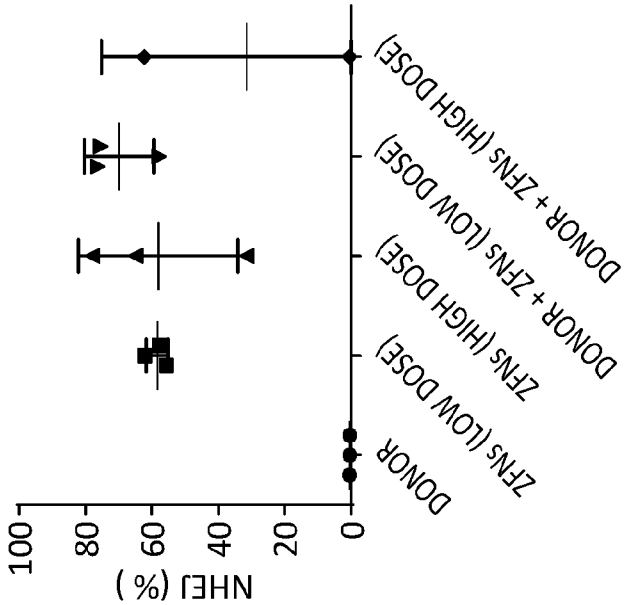


Figure 3A



**Exon 1**  
(69-bp)

**B**

gacaggATGGCTTCCCTTCCTTCCTGCCTCGCTGGACTGGTATTTGTGTCTGAAGCTGGCCCCGGtgagtgatcctgtgagcga

ctgtccTACCGAAGGAAGCTGAGAGGAGGAACGGAGCGACCTGACCATAAACACAGACTTCGACCGGGGGCCactcactaggaactcgct

gacaggATGGCTTCCCTTCGACTCTTCCCTCCTTGCCTCGCTGGACTGGTATTGTGCTCTGAAGCTGGCCCCCGGTgagtgatcctgtgagcga

ctgtccTACCGAAGGAAGCTGAGAAGGAGGAAACGGAGCGACCTGACCATAAACACAGACTTCGACCGGGCGGCactcactaqqacactcgct

**pacaggATGGCTTCCCTTCGACTCTTCCCTCCTTTGCCCTCGCTGGACTGGTATTGTGTC<sup>T</sup>GAAAGCTGGCCCCCGCGGTgagtgatcctgtgagcgga**

ctgtccTACCGAAGGAAGCTGAGAAGGAGAAACGGAGCGACCTGACCATAAACACAGACTTCGACCGGGGGCactcactaggaactcgct

### Figure 4C

**Exon 2**  
**(131-bp)**

**H** **I**  
gtgccagGGTGTGGAGAAATCCAAATGTCTCTGATGGTCAAAGTCCCTGGATGCTGCCGAGGCAGCCCTGCTGTAGACGTGGCTGTAAAAGTGTT  
cacgggtcccACGACCCTCTTAGGTTTACAGGAGACTACCAGTTTCAGGACCTACGACAGGCTCCGTCCGGACGACATCTGCACCGACATTTTCACAA

SUBSTITUTE SHEET (RULE 26)

**Figure 4D** 5/9

**J**  
CAAAAAGACCCTCTGAGGGATCCTGGGAGCCCTTTGCCCTCTGGgtaagctt  
GTTTTCTGGAGACTCCCTAGGACCCCTCGGAAACGGAGACCCcattcgaa

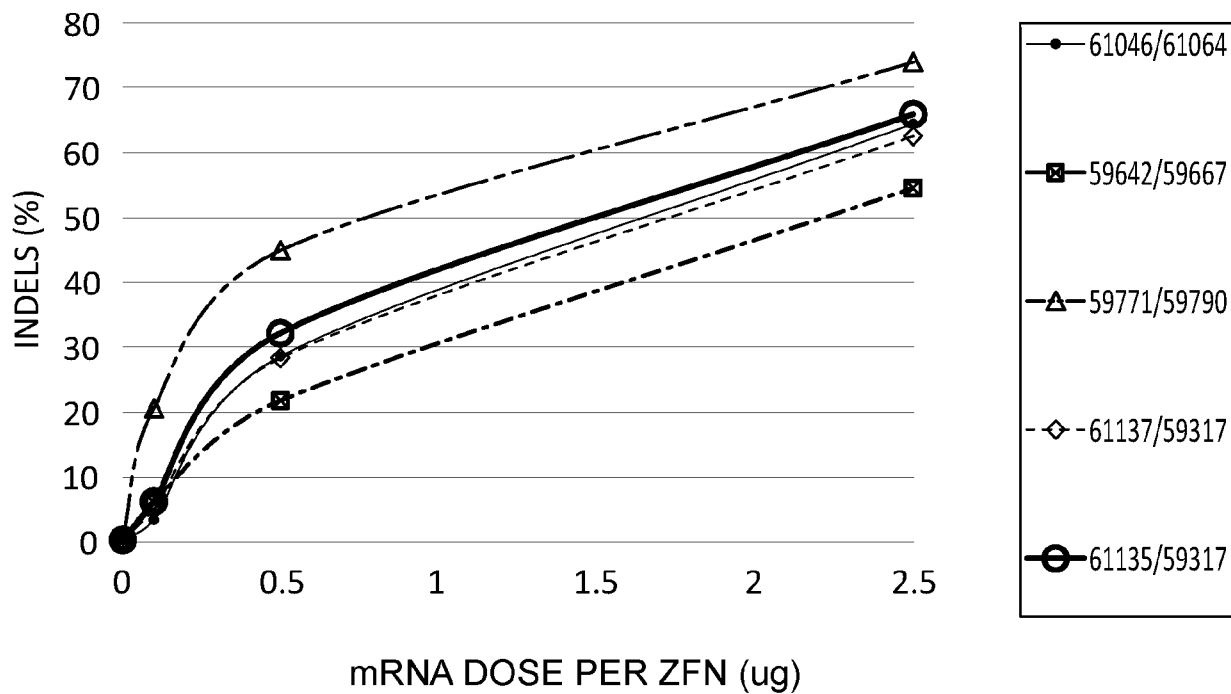
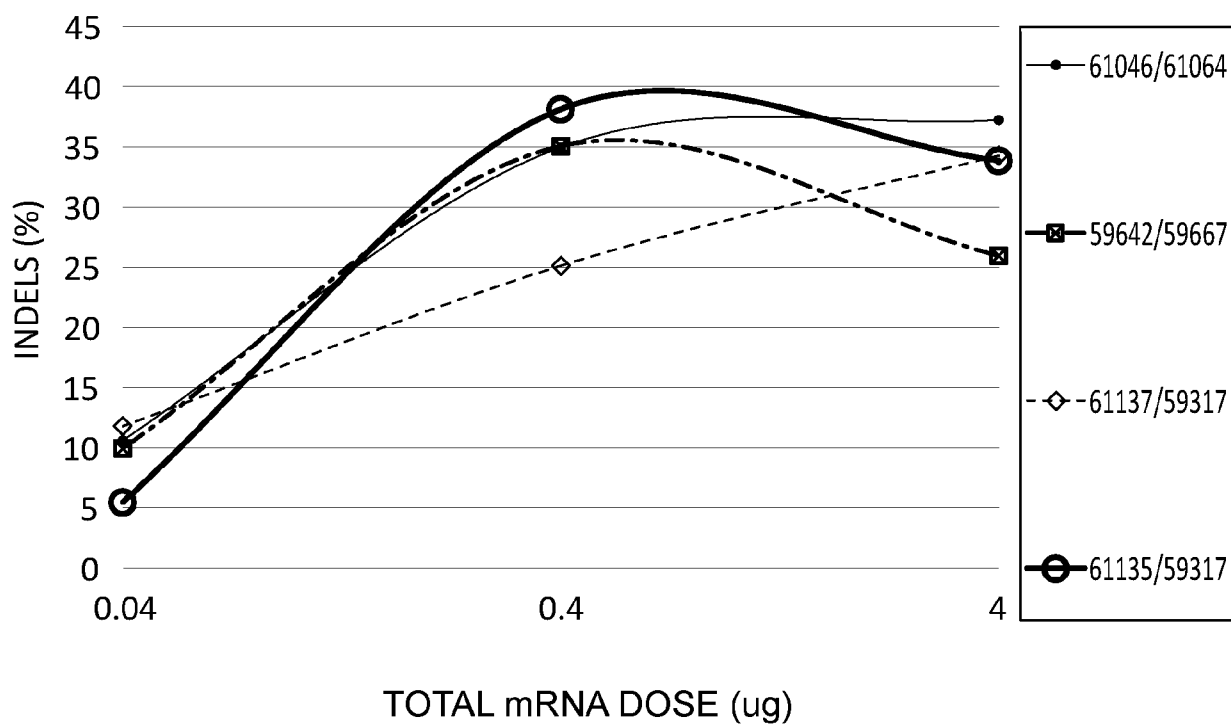
**Exon 3**  
**(136-bp)**

**L** **M**  
ccctccagGAAGACCGCGGAGTCTGGAGAGCTGCACGGGCTCACCACAGATGAGAAGTTTGTAGAAGGAGTGTACAGAGTAGAACTGGACACCCAAAT<sup>\*</sup>  
gggaggctcCTTCTGGCGCCCTCAGACCCTCTCGACGTGCCCGAGTGGTGCTACTCTTCAAACATCTTCCCTCACATGTCTCATCTTGACCTGTGGTTTA

**K (2 BOTTOM SITES)**  
CGTACTGGAAGACACTTGGCATTTCCTCCGTTCATGAATTTCGCGGATgtaagtgg  
GCATGACCTTCTGTGAACCGTAAAGGGGCAAGGTACTTAAGCGCCTAcattcacc

**Figure 4E**

6/9

**Figure 5A****Figure 5B**

7/9

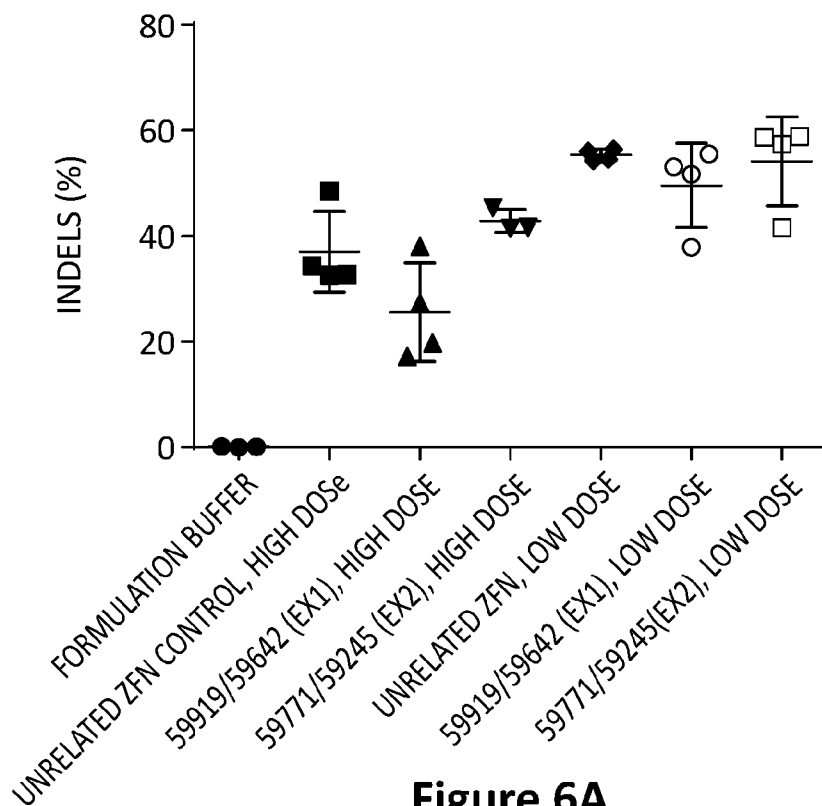


Figure 6A

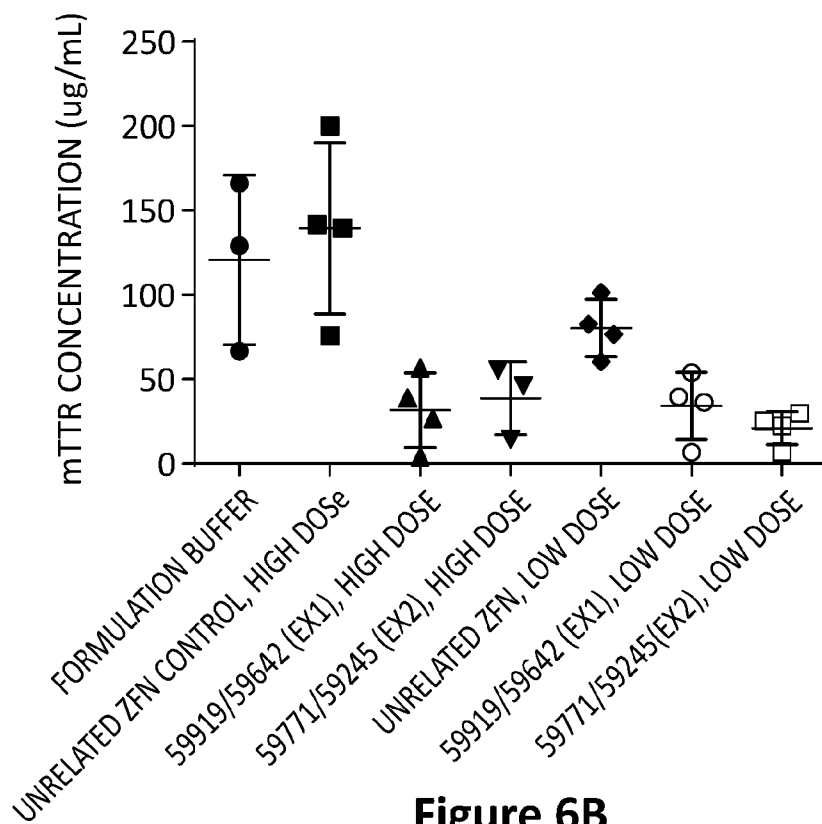


Figure 6B

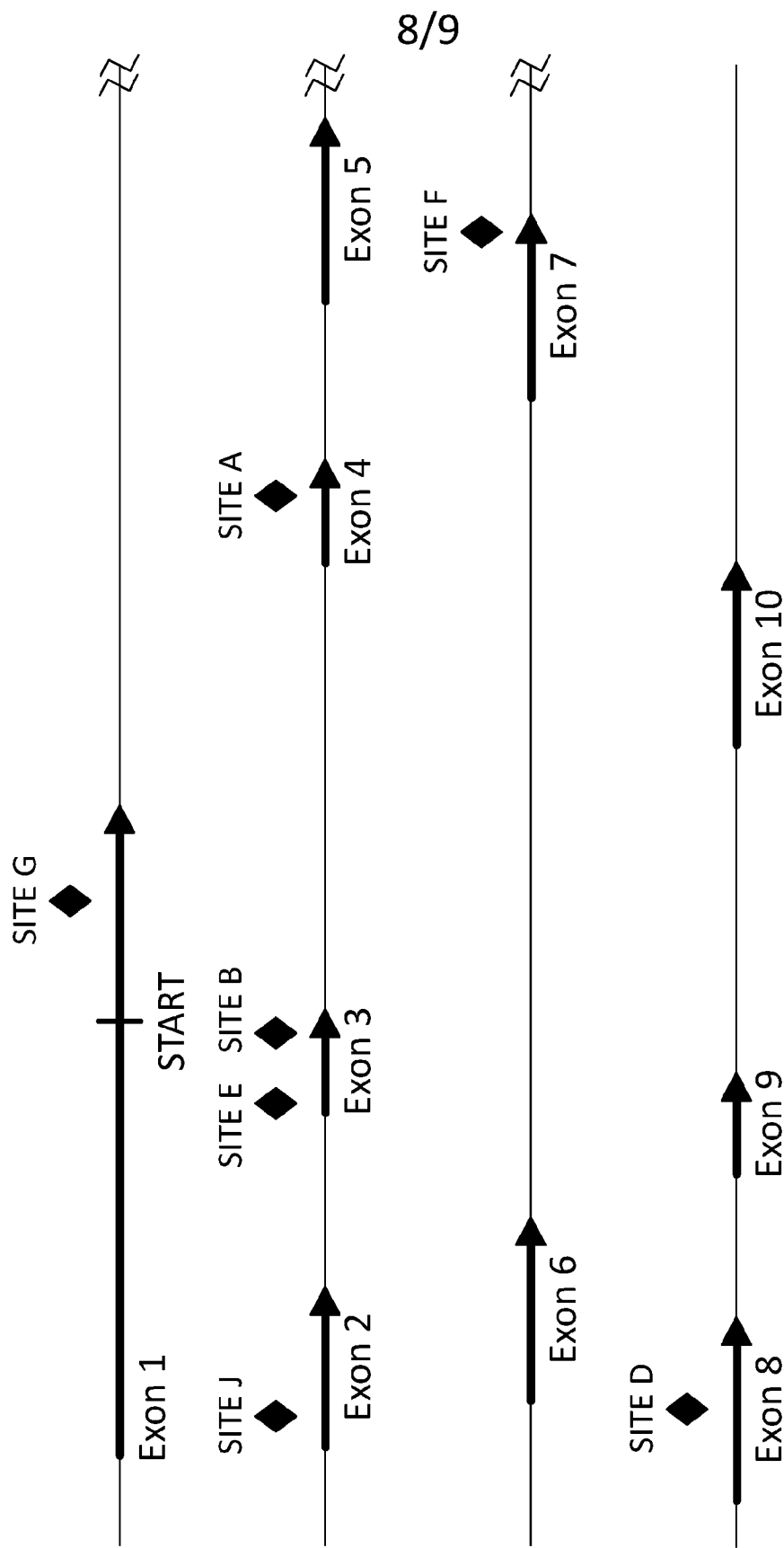


Figure 7

9/9

