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(54) Title: POLYNUCLEOTIDES ENCODING GLUCOSE-6-PHOSPHATASE FOR THE TREATMENT OF GLYCOGEN STORAGE DISEASE

FIGURE 1A

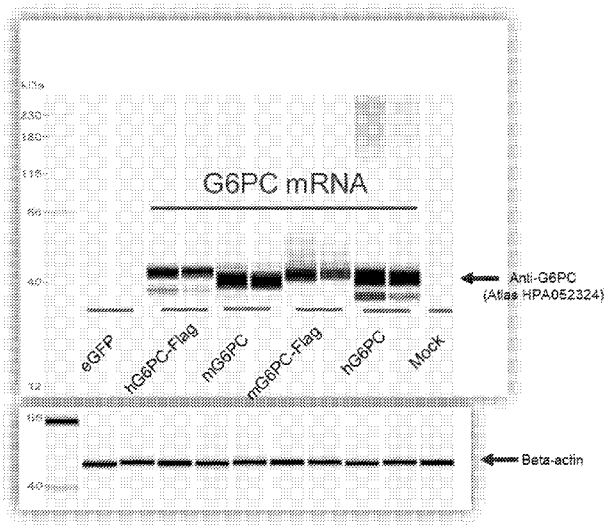
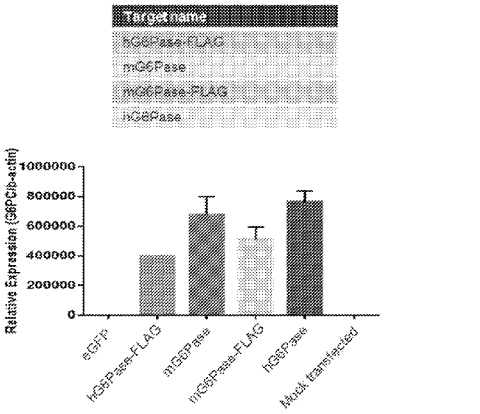


FIGURE 1B



Quantitative analysis of capillary electrophoresis (CE) signals (normalized by beta-actin)

(57) Abstract: This disclosure relates to mRNA therapy for the treatment of glycogen storage disease type 1a, (GSD-Ia), and related symptoms such as hypoglycemia. mRNAs for use in the invention, when administered *in vivo*, encode human glucose-6-phosphatase (G6Pase or G6PC), and functional fragments and variants thereof. mRNAs of the invention are preferably encapsulated in lipid nanoparticles (LNPs) to effect efficient delivery to cells and/or tissues in subjects, when administered thereto. mRNA therapies of the invention increase and/or restore deficient levels of G6PC expression and/or activity in subjects. mRNA therapies of the invention further increase the glucose production, and reduce the abnormal accumulation of glycogen and/or glucose-6-phosphate associated with GSD-Ia.

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[Continued on next page]

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POLYNUCLEOTIDES ENCODING GLUCOSE-6-PHOSPHATASE FOR THE TREATMENT OF GLYCOGEN STORAGE DISEASE

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority to U.S. Provisional Application No. 62/730,940, filed September 13, 2018, and U.S. Provisional Application No. 62/812,800, filed March 1, 2019. The content of both applications is incorporated by reference herein in their entirety.

SEQUENCE LISTING

[0002] The instant application contains a Sequence Listing which has been submitted electronically in ASCII format and is hereby incorporated by reference in its entirety. Said ASCII copy, created on September 10, 2019, is named 45817-0049WO1_SL.txt and is 239,349 bytes in size.

BACKGROUND

[0003] Glycogen storage disease type 1 (GSD-I), also known as von Gierke disease, is caused by an accumulation of glycogen in organs and tissues of the body. GSD-I can be divided into two major subtypes, including glycogen storage disease type 1a (GSD-Ia, MIM232200) and type Ib (GSD-Ib, MIM232220). Chou et al., Liver Research, 2017, 1:174-180. GSD-I has an incidence of about 1 in 100,000. GSD-Ia is an autosomal recessive metabolic disorder that is caused by a deficiency in glucose -6-phosphatase (G6Pase or G6PC; EC 3.1.3.9), and accounts for about 80% of GSD-I cases. G6PC is a key enzyme that is necessary for glucose production that is expressed mostly in the liver, kidney, and intestine, and functions inside of the endoplasmic reticulum (ER) lumen of cells. G6PC catalyzes the hydrolysis of glucose-6-phosphate (G6P) to glucose and phosphate late during gluconeogenesis and glycogenolysis. Normally, as blood glucose levels fall in between meals, G6P is produced during the terminal step of gluconeogenesis and glycogenolysis in the gluconeogenic organs (primarily liver, and also the kidneys and intestine) and it is hydrolyzed by G6PC to glucose and then released back into the blood.

[0004] GSD-Ia is caused by mutation of the gene encoding G6PC that impairs the enzyme's ability to hydrolyze G6P. After a short fast of a few hours, individuals with GSD-Ia become hypoglycemic. G6P becomes elevated in cellular cytoplasm, which leads to an accumulation of glycogen and fat in the liver and kidneys, and impaired blood glucose homeostasis. The accumulation of glycogen promotes progressive hepatomegaly and nephromegaly. GSD-Ia patients can also develop other metabolic complications, including hypercholesterolemia, hypertriglyceridemia, hyperuricemia, hyperlipidemia, and lactic academia. Individuals with GSD-Ia can also develop longer term problems, including growth retardation, osteoporosis, gout, pulmonary hypertension, and renal disease. One long-term complication is the development of hepatic tumors in 75% of GSD-Ia patients over the age of 25 years old, of which, approximately 10% transform into malignant tumors.

[0005] GSD-1a is lethal during childhood if left untreated. The standard of care for GSD-Ia is strict diet control that involves the frequent consumption of uncooked cornstarch, which often results in poor compliance with the dietary restrictions. Furthermore, dietary control fails to prevent the long-term metabolic complications associated with G6PC deficiency, and cannot prevent the long-term consequences of hepatocellular adenoma. Enzyme replacement therapy is not an option for subjects with GSD-Ia because of drug delivery challenges, and current virus-based gene therapy approaches to correct G6PC deficiency have yet to be successful. Currently, only a liver transplant is curative for GSD-Ia patients. As a result, new therapies to prevent or treat GSD-Ia are needed. In view of the significant problems associated with existing approaches for preventing or treating GSD-Ia, there is an unmet need in the art for improved GSD-Ia therapies.

SUMMARY

[0006] The present disclosure provides messenger RNA (mRNA) therapeutics for the treatment of glycogen storage disease type 1 (GSD-Ia). The mRNA therapeutics of the invention are particularly well-suited for the treatment of GSD-Ia as the technology provides for the intracellular delivery of mRNA encoding glucose-6-phosphatase (G6Pase or G6PC) followed by *de novo* synthesis of functional G6PC protein within target cells. The instant invention features the incorporation of modified nucleotides within therapeutic mRNAs to (1) minimize unwanted immune

activation (*e.g.*, the innate immune response associated with the *in vivo* introduction of foreign nucleic acids) and (2) optimize the translation efficiency of mRNA to protein. Exemplary aspects of the disclosure feature a combination of nucleotide modification to reduce the innate immune response and sequence optimization, in particular, within the open reading frame (ORF) of therapeutic mRNAs encoding G6PC to enhance protein expression.

[0007] In further embodiments, the mRNA therapeutic technology of the instant disclosure also features delivery of mRNA encoding G6PC *via* a lipid nanoparticle (LNP) delivery system. The instant disclosure features ionizable lipid-based LNPs, which have improved properties when combined with mRNA encoding G6PC and administered *in vivo*, for example, cellular uptake, intracellular transport and/or endosomal release or endosomal escape. The LNP formulations of the disclosure also demonstrate reduced immunogenicity associated with the *in vivo* administration of LNPs.

[0008] In certain aspects, the disclosure relates to compositions and delivery formulations comprising a polynucleotide, e.g., a ribonucleic acid (RNA), e.g., a mRNA, encoding G6PC and methods for treating GSD-Ia in a human subject in need thereof by administering the same.

[0009] The present disclosure provides a pharmaceutical composition comprising a lipid nanoparticle encapsulated mRNA that comprises an open reading frame (ORF) encoding a G6PC polypeptide, wherein the composition is suitable for administration to a human subject in need of treatment for GSD-Ia.

[0010] The present disclosure further provides a pharmaceutical composition comprising: (a) a mRNA that comprises (i) an open reading frame (ORF) encoding a G6PC polypeptide, wherein the ORF comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof and (ii) an untranslated region (UTR) comprising a microRNA (miRNA) binding site; and (b) a delivery agent, wherein the pharmaceutical composition is suitable for administration to a human subject in need of treatment for GSD-Ia.

[0011] In certain embodiments, the pharmaceutical composition or polynucleotide is administered intravenously. In some instances, the pharmaceutical composition or polynucleotide is administered at a dose of 0.1 mg/kg to 2.0 mg/kg. In some instances, the pharmaceutical composition or polynucleotide is administered at a dose of 0.1 mg/kg to 1.5 mg/kg. In some instances, the pharmaceutical composition or

polynucleotide is administered at a dose of 0.1 mg/kg to 1.0 mg/kg. In some instances, the pharmaceutical composition or polynucleotide is administered at a dose of 0.1 mg/kg to 0.5 mg/kg.

[0012] In one aspect, the disclosure features a pharmaceutical composition comprising an mRNA comprising an open reading frame (ORF) encoding a glucose-6-phosphatase (G6PC) polypeptide, wherein the composition when administered as a single intravenous dose to a human subject in need thereof is sufficient to: (i) increase the level of G6PC activity in liver tissue to within at least 2%, at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% of normal G6PC activity level for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (ii) increase the level of G6PC activity in liver tissue at least 1.5-fold, at least 2-fold, at least 3-fold, at least 4-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold compared to the human subject's baseline G6PC activity level or a reference G6PC activity level in a human subject having glycogen storage disease type 1a (GSD-Ia) for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (iii) reduce liver levels of glycogen at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (iv) reduce liver levels of glucose-6-phosphate (G6P) at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (v) reduce liver levels of glycogen at least 1.5-fold, at least 2-fold, at

least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold when the human subject is fasting as compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (vi) reduce liver levels of G6P at least 1.5-fold, at least 2-fold at least 5-fold, at least 10-fold, at least 20-fold or at least 50-fold as compared to the human subject's baseline liver G6P levels while fasting or a reference liver G6P level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (vii) reduce serum levels of triglycerides at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (viii) reduce serum levels of triglycerides at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold when the human subject is fasting as compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (ix) increase blood, plasma, and/or serum levels of glucose at least 2%, at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline blood, plasma, or serum glucose level while fasting or a reference blood, plasma, or serum glucose level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; (x) increase blood, plasma, and/or serum levels of glucose at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold

when the human subject is fasting as compared to the human subject's baseline blood, plasma, and/or serum glucose level while fasting or a reference blood, plasma, and/or serum glucose level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; and/or (xi) increase blood, plasma, or serum levels of glucose to at least 60 mg/dl, at least 65 mg/dl, at least 70 mg/dl, at least 75 mg/dl , at least 80 mg/dl , at least 85 mg/dl , at least 90 mg/dl , at least 95 mg/dl , at least 100 mg/dl, at least 105 mg/dl, at least 110 mg/dl, at least 115 mg/dl, at least 120 mg/dl, at least 125 mg/dl, at least 130 mg/dl, at least 135 mg/dl, at least 140 mg/dl, at least 145 mg/dl, or at least 150 mg/dl in a fasting patient with GSD-Ia for at least 6 hours, at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration.

[0013] In some embodiments, the pharmaceutical further comprises a delivery agent. In some embodiments, the delivery agent comprises a lipid nanoparticle comprising: (a) (i) Compound II, (ii) Cholesterol, and (iii) PEG-DMG or Compound I; (b) (i) Compound VI, (ii) Cholesterol, and (iii) PEG-DMG or Compound I; (c) (i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) PEG-DMG or Compound I; (d) (i) Compound VI, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) PEG-DMG or Compound I; (e) (i) Compound II, (ii) Cholesterol, and (iii) Compound I; or (f) (i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) Compound I.

[0014] In some embodiments, the G6PC polypeptide consists of the amino acid sequence set forth in SEQ ID NO:83, SEQ ID NO:84, SEQ ID NO:194-206, 208, or 209.

[0015] In some embodiments, the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to a nucleic acid sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207.

[0016] In some embodiments, the mRNA comprises a microRNA (miR) binding site. In some instances, the microRNA is expressed in an immune cell of hematopoietic lineage or a cell that expresses TLR7 and/or TLR8 and secretes pro-inflammatory cytokines and/or chemokines. In some instances, the microRNA binding site is for a

microRNA selected from the group consisting of miR-126, miR-142, miR-144, miR-146, miR-150, miR-155, miR-16, miR-21, miR-223, miR-24, miR-27, miR-26a, or any combination thereof. In some instances, the microRNA binding site is for a microRNA selected from the group consisting of miR126-3p, miR-142-3p, miR-142-5p, miR-155, or any combination thereof. In some instances, the microRNA binding site is a miR-142-3p binding site. In some instances, the microRNA binding site is located in the 3′ UTR of the mRNA.

[0017] In some embodiments, the mRNA comprises a 3' UTR comprising a nucleic acid sequence at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 3' UTR sequence of SEQ ID NO:150 or 175.

[0018] In some embodiments, the mRNA comprises a 3' UTR comprising a nucleic acid sequence at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 3' UTR sequence of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213.

[0019] In some embodiments, the mRNA comprises a 5' UTR comprising a nucleic acid sequence at least 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 5' UTR sequence of SEQ ID NO:3.

[0020] In some embodiments, the mRNA comprises a 5' UTR comprising a nucleic acid sequence at least 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 5' UTR sequence of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211.

[0021] In some embodiments, the mRNA comprises a 5′ terminal cap. In some instances, the 5′ terminal cap comprises a Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5′ methylG cap, or an analog thereof.

[0022] In some embodiments, the mRNA comprises a poly-A region. In some instances, the poly-A region is at least about 10, at least about 20, at least about 30, at least about 40, at least about 50, at least about 60, at least about 70, at least about 80, at least about 90 nucleotides in length, or at least about 100 nucleotides in length. In some instances, the poly-A region has about 10 to about 200, about 20 to about 180,

about 50 to about 160, about 70 to about 140, or about 80 to about 120 nucleotides in length.

[0023] In some embodiments, the mRNA comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof. In some instances, the at least one chemically modified nucleobase is selected from the group consisting of pseudouracil (ψ), N1-methylpseudouracil (m1ψ), 1-ethylpseudouracil, 2-thiouracil (s2U), 4'-thiouracil, 5-methylcytosine, 5-methyluracil, 5-methoxyuracil, and any combination thereof. In some instances, at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 99%, or 100% of the uracils are chemically modified to N1-methylpseudouracils.

[0024] In some embodiments, the human subject has GSD-Ia.

[0025] In some embodiments, the human subject has been fasting.

[0026] In some embodiments, the human subject has been fasting for at least 2 hours.

[0027] In another aspect, the disclosure features a polynucleotide comprising an mRNA comprising: (i) a 5' UTR; (ii) an open reading frame (ORF) encoding a human G6PC polypeptide, wherein the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to a nucleic acid sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207; (iii) a stop codon; and (iv) a 3' UTR.

[0028] In some embodiments,

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:8 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:205;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:11 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:204;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:12 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:195;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:13 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:197;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:14 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:196;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:15 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:198;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:16 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:194;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:17 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:200;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:18 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:201;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:19 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:202;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:20 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:203;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least

98%, at least 99%, or 100% sequence identity to SEQ ID NO:22 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:208;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:23 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:194;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:24 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:205;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:25 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:209;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:26 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:194;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:27 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:205;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:28 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:209;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:29 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:30 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:31 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:32 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:33 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:34 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:35 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:36 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:37 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:38 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84; or
the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least

98%, at least 99%, or 100% sequence identity to SEQ ID NO:207 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:206.

[0029] In some embodiments, the mRNA comprises a microRNA (miR) binding site. In some instances, the microRNA is expressed in an immune cell of hematopoietic lineage or a cell that expresses TLR7 and/or TLR8 and secretes pro-inflammatory cytokines and/or chemokines. In some instances, the microRNA binding site is for a microRNA selected from the group consisting of miR-126, miR-142, miR-144, miR-146, miR-150, miR-155, miR-16, miR-21, miR-223, miR-24, miR-27, miR-26a, or any combination thereof. In some instances, the microRNA binding site is for a microRNA selected from the group consisting of miR126-3p, miR-142-3p, miR-142-5p, miR-155, or any combination thereof. In some instances, the microRNA binding site is a miR-142-3p binding site. In some instances, the microRNA binding site is located in the 3′ UTR of the mRNA.

[0030] In some embodiments, the 3' UTR comprises a nucleic acid sequence at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 3' UTR of SEQ ID NO:150 or SEQ ID NO:175.

[0031] In some embodiments, the 3' UTR comprises a nucleic acid sequence at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 3' UTR of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213.

[0032] In some embodiments, the 5' UTR comprises a nucleic acid sequence at least 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 5' UTR sequence of SEQ ID NO:3.

[0033] In some embodiments, the mRNA comprises a 5' UTR comprising a nucleic acid sequence at least 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 5' UTR sequence of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211.

[0034] In some embodiments, the mRNA comprises a 5′ terminal cap. In some instances, the 5′ terminal cap comprises a Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5′ methylG cap, or an analog thereof.

[0035] In some embodiments, the mRNA comprises a poly-A region. In some instances, the poly-A region is at least about 10, at least about 20, at least about 30, at least about 40, at least about 50, at least about 60, at least about 70, at least about 80, at least about 90 nucleotides in length, or at least about 100 nucleotides in length. In some instances, the poly-A region has about 10 to about 200, about 20 to about 180, about 50 to about 160, about 70 to about 140, or about 80 to about 120 nucleotides in length.

[0036] In some embodiments, the mRNA comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof. In some instances, the at least one chemically modified nucleobase is selected from the group consisting of pseudouracil (ψ), N1-methylpseudouracil (m1ψ), 1-ethylpseudouracil, 2-thiouracil (s2U), 4'-thiouracil, 5-methylcytosine, 5-methyluracil, 5-methoxyuracil, and any combination thereof. In some instances, at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 99%, or 100% of the uracils are chemically modified to N1-methylpseudouracils.

[0037] In some embodiments, the polynucleotide comprises a nucleic acid sequence selected from the group consisting of SEQ ID NO:53-81, 214, 215, and 217.

[0038] In another aspect, the disclosure features a polynucleotide comprising an mRNA comprising: (i) a 5'-terminal cap; (ii) a 5' UTR comprising the nucleic acid sequence of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211; (iii) an open reading frame (ORF) encoding a G6PC polypeptide, wherein the ORF comprises a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207; (iv) a 3' UTR comprising the nucleic acid sequence of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213; and (vi) a poly-A-region.

[0039] In another aspect, the disclosure features a polynucleotide comprising an mRNA comprising: (i) a 5'-terminal cap; (ii) a 5' UTR comprising the nucleic acid sequence of SEQ ID NO:3; (iii) an open reading frame (ORF) encoding a G6PC polypeptide, wherein the ORF comprises a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207; (iv) a 3' UTR comprising the nucleic acid sequence of SEQ ID NO:150 or SEQ ID NO:175; and (vi) a poly-A-region.

[0040] In some embodiments, the 5′ terminal cap comprises a Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5′ methylG cap, or an analog thereof.

[0041] In some embodiments, the poly-A region is at least about 10, at least about 20, at least about 30, at least about 40, at least about 50, at least about 60, at least about 70, at least about 80, at least about 90 nucleotides in length, or at least about 100 nucleotides in length.

[0042] In some embodiments, the poly-A region has about 10 to about 200, about 20 to about 180, about 50 to about 160, about 70 to about 140, or about 80 to about 120 nucleotides in length.

[0043] In some embodiments, the mRNA comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof. In some instances, the at least one chemically modified nucleobase is selected from the group consisting of pseudouracil (ψ), N1-methylpseudouracil (m1ψ), 1-ethylpseudouracil, 2-thiouracil (s2U), 4'-thiouracil, 5-methylcytosine, 5-methyluracil, 5-methoxyuracil, and any combination thereof.

[0044] In some embodiments, the polynucleotide comprises a nucleic acid sequence selected from the group consisting of SEQ ID NO:53-81, 214, 215, and 217. In some instances of this embodiment, the 5′ terminal cap comprises Cap1 and all of the uracils of the polynucleotide are N1-methylpseudouracils. In some instances of this embodiment, the poly-A-region is 100 nucleotides in length.

[0045] In another aspect, the disclosure features a pharmaceutical composition comprising a polynucleotide disclosed herein, and a delivery agent. In some embodiments, the delivery agent comprises a lipid nanoparticle comprising: (a) (i) Compound II, (ii) Cholesterol, and (iii) PEG-DMG or Compound I; (b) (i) Compound VI, (ii) Cholesterol, and (iii) PEG-DMG or Compound I; (c) (i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) PEG-DMG or Compound I; (d) (i) Compound VI, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) PEG-DMG or Compound I; (e) (i) Compound II, (ii) Cholesterol, and (iii) Compound I; or (f) (i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) Compound I.

[0046] In another aspect, the disclosure features a method of expressing a glucose-6-phosphatase (G6PC) polypeptide in a human subject in need thereof, comprising

administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0047] In another aspect, the disclosure features a method of treating, preventing, or delaying the onset and/or progression of glycogen storage disease type 1a (GSD-Ia) in a human subject in need thereof, comprising administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0048] In another aspect, the disclosure features a method of increasing glucose blood, plasma or serum level in a human subject in need thereof, comprising administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0049] In another aspect, the disclosure features a method of reducing liver glycogen level in a human subject in need thereof, comprising administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0050] In another aspect, the disclosure features a method of reducing serum triglyceride level in a human subject in need thereof, comprising administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0051] In another aspect, the disclosure features a method of reducing glucose-6-phosphate (G6P) liver level in a human subject in need thereof, comprising administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0052] In some embodiments of the foregoing methods, (i) the liver glycogen level is reduced at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting as compared to the subject's baseline glycogen liver level while fasting or a reference glycogen liver level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; (ii) the liver G6P level is reduced at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% as compared to the subject's baseline liver G6P level while fasting or a reference liver G6P level in a

patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; (iii) the liver glycogen level is reduced to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting as compared to a normal liver glycogen level while fasting within at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks; (iv) the liver G6P level is reduced to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting, as compared to a normal liver G6P level while fasting, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; (v) the serum triglyceride level is reduced at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting as compared to the subject's baseline serum triglyceride level while fasting or a reference liver glycogen level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; (vi) the serum triglyceride level is reduced to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting as compared to a normal serum triglyceride level while fasting within at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; (vii) the blood, plasma or serum glucose liver is increased at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% as compared to the subject's baseline glucose blood, plasma or serum level while fasting or a reference glucose blood, plasma or serum level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; (viii) the glucose blood, plasma or serum level is increased to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting as compared to a

normal glucose blood, plasma or serum level while fasting within at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; and/or (ix) the glucose blood, plasma or serum level is increased to at least 70 mg/dl, at least 75 mg/dl , at least 80 mg/dl , at least 85 mg/dl , at least 90 mg/dl , at least 95 mg/dl , at least 100 mg/dl, at least 105 mg/dl, at least 110 mg/dl, at least 115 mg/dl, at least 120 mg/dl, at least 125 mg/dl, at least 130 mg/dl, at least 135 mg/dl, at least 140 mg/dl, at least 145 mg/dl, or at least 150 mg/dl in the subject when fasting for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration.

[0053] In another aspect, the disclosure features a method of increasing G6PC activity in a human subject in need thereof, comprising administering to the subject an effective amount of a pharmaceutical composition disclosed herein or a polynucleotide disclosed herein.

[0054] In some embodiments, (i) the level of G6PC activity in the subject is increased at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold as compared to a reference G6PC activity level in a subject having GSD-Ia for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; and/or (ii) 12 hours after a single administration of the pharmaceutical composition or polynucleotide is administered to the subject, the G6PC activity in the subject is increased at least 2%, at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, at least 100%, at least 150%, at least 200%, at least 300%, at least 400%, at least 500%, or at least 600% compared to the subject's baseline G6PC activity. In some embodiments, the G6PC activity is increased in the liver or blood of the subject.

[0055] In some embodiments of the foregoing methods, the human subject has been fasting (e.g., for at least 2 hours).

[0056] In some embodiments of the foregoing methods, the administration to the subject is about once a week, about once every two weeks, or about once a month.

[0057] In some embodiments of the foregoing methods, the pharmaceutical composition or polynucleotide is administered intravenously.

BRIEF DESCRIPTION OF THE DRAWINGS

[0058] **FIG. 1A** includes a capillary electrophoresis (CE)-based Western analysis showing the expression of human G6PC (hG6Pase), mouse G6PC (mG6Pase), and FLAG-tagged human and mouse G6PC proteins (hG6Pase-FLAG and mG6Pase-FLAG) in HEK 293 cells 24 hours post transfection.

[0059] **FIG. 1B** is a bar graph showing the quantitative analysis of expression of hG6PC, mG6PC, hG6PC-FLAG, and mG6PC-FLAG as determined by capillary electrophoresis (as shown in FIG. 1A), and normalized by beta-actin.

[0060] **FIG. 2A** is a schematic outlining the Taussky-Shorr Color Reagent (TSCR) assay used to detect G-6-phosphatase (G6Pase) activity.

[0061] **FIG. 2B** is a bar graph showing the activity of human G6PC (hG6Pase), mouse G6PC (mG6Pase), and FLAG-tagged human and mouse G6PC proteins (hG6Pase-FLAG and mG6Pase-FLAG) in HEK 293 cells 24 hours post transfection.

[0062] **FIG. 3A** is a bar graph showing the expression of mRNA encoding human G6PC Variant 1 (V1), G6PC Variant 2 (V2), and wild-type G6PC in wild-type CD-1 mice at 24 hours, 48 hours, and 96 hours after mRNA injection, in comparison to the hepatic G6PC protein expression in wild-type CD-1 mice treated with a control mRNA encoding eGFP.

[0063] **FIG. 3B** is a bar graph showing the activity of human G6PC Variant 1 (V1), G6PC Variant 2 (V2), and wild-type G6PC in wild-type CD-1 mice at 24 hours, 48 hours, and 96 hours after injection of mRNA encoding the G6PC proteins, relative to G6PC activity in control mice injected with an eGFP-encoding mRNA.

[0064] **FIG. 4** is a bar graph showing the results of a bioinformatics analysis of G6PC-related protein sequences from >100 mammalian species including human G6PC, which identified 36 consensus amino acids that differ from human G6PC. A subset of variants with the greatest difference between the consensus and reference scores were selected for further investigation.

[0065] **FIG. 5A** is a bar graph showing the activity of human G6PC proteins with single amino acid substitutions and G6PC Variant 2 in Hela cells 48 hours after transfection with G6PC-encoding mRNA.

[0066] **FIG. 5B** is a bar graph showing the activity of human G6PC proteins with single amino acid substitutions and G6PC proteins with multiple amino acid substitutions in Hela cells 48 hours after transfection with G6PC-encoding mRNA.

[0067] **FIG. 5C** is a bar graph showing the expression of human G6PC proteins with single amino acid substitutions and G6PC proteins with multiple amino acid substitutions in Hela cells 48 hours after transfection with G6PC-encoding mRNA, as determined by Western blot.

[0068] **FIG. 5D** is a bar graph showing the activity of human G6PC proteins with single amino acid substitutions and G6PC Variant 2 in Hela cells 48 hours after transfection with G6PC-encoding mRNA.

[0069] **FIG. 5E** is a bar graph showing the activity of human G6PC proteins with single amino acid substitutions in Hela cells 48 hours after transfection with G6PC-encoding mRNA.

[0070] **FIG. 5F** is a bar graph showing the expression of human G6PC proteins with single amino acid substitutions and G6PC Variant 2 in Hela cells 48 hours after transfection with G6PC-encoding mRNA, as determined by Western blot.

[0071] **FIG. 5G** is a bar graph showing the expression of human G6PC proteins with single amino acid substitutions in Hela cells 48 hours after transfection with G6PC-encoding mRNA, as determined by Western blot.

[0072] **FIG. 6A** is a bar graph showing the activity of human G6PC Variant 1 (CoV1) and G6PC Variant 2 (CoV2) expressed from codon-optimized mRNA constructs, and G6PC Variant 2 (V2) expressed from a non-codon-optimized mRNA construct, at 48 and 96 hours after injection of mRNA formulated in MC-3 lipid nanoparticles in wild-type mice.

[0073] **FIG. 6B** is a bar graph showing the activity of human G6PC Variant 1 (CoV1) and G6PC Variant 2 (CoV2) expressed from codon-optimized mRNA constructs, and G6PC Variant 2 (V2) expressed from a non-codon-optimized mRNA construct, at 48 and 96 hours after injection of mRNA formulated in Compound II lipid nanoparticles in wild-type mice.

[0074] **FIG. 6C** is a bar graph showing the activity of human G6PC Variant 1 and G6PC Variant 2 proteins expressed from codon-optimized mRNA constructs, and G6PC Variant 1 and Variant 2 proteins expressed from a non-codon-optimized mRNA construct, at 96 hours after injection of mRNA in wild-type mice.

[0075] **FIG. 6D** is a bar graph showing the expression of human G6PC Variant 1 and G6PC Variant 2 proteins expressed from codon-optimized mRNA constructs, and G6PC Variant 1 and Variant 2 proteins expressed from a non-codon-optimized mRNA construct, at 96 hours after injection of mRNA in wild-type mice.

[0076] **FIG. 7A** is a bar graph showing the activity of human G6PC variants with single or multiple amino acid substitutions expressed from codon-optimized or non-codon optimized mRNA constructs at 96 hours after mRNA injection in wild-type mice. The arrow denotes the activity of G6PC with an S298C substitution (S298C_CoV1) expressed from a codon-optimized mRNA.

[0077] **FIG. 7B** is a bar graph showing the expression of the human G6PC variants presented in FIG. 7A. The arrow denotes the expression of G6PC with an S298C substitution (S298C_CoV1) expressed from a codon-optimized mRNA.

[0078] **FIGs. 8A-8F** are bar graphs showing the glucose levels in the liver-specific G6PC knockout (L-G6PC(-/-)) mice injected with either 0.5 mg/kg or 1 mg/kg of mRNA encoding G6PC S298C or GFP (control) in mice that were fed or fasted for 2.5 hours or 6 hours at 1, 3, 5, 8, 11, and 15 days (FIGs. 8A-8F, respectively) after mRNA injection. The dotted horizontal line in each figure denotes a glucose level of 70 mg/dl, which is the threshold blood glucose level to maintain normal physiology.

[0079] **FIGs. 8G-8H** are graphs showing the duration of action of a single dose of 0.5 mg/kg of mRNA encoding G6PC S298C (FIG. 8G) or 1 mg/kg of mRNA encoding G6PC S298C (FIG. 8H) in the L-G6PC(-/-) mice.

[0080] **FIGs. 9A-9C** are graphs showing fasting glucose levels in the L-G6PC(-/-) mice injected with 1 mg/kg of mRNA encoding G6PC S298C (FIG. 9A), 0.5 mg/kg of mRNA encoding G6PC S298C (FIG. 9B), or mRNA encoding eGFP (FIG. 9C), at 1, 3, 5, 8, 11, and 15 days after mRNA injection, plotted as a paired comparison between pre- and post-treatment. $*P<0.05$, $**P<0.01$, $***P<0.001$, $****P<0.0001$ vs before (pre-treatment), paired *t*-test.

[0081] **FIG. 10A** is a bar graph showing the glucose levels in L-G6PC(-/-) mice injected with either 0.2 mg/kg, 0.5 mg/kg or 1 mg/kg of mRNA encoding G6PC S298C or mRNA encoding GFP (control) in mice that were fed or fasted for 2.5 hours, 6 hours, or 24 hours during the time course of 24 hours after mRNA injection.

[0082] **FIG. 10B** is a graph showing that injection of 0.2 mg/kg, 0.5 mg/kg, or 1.0 mg/kg of mRNA encoding G6PC S298C or mRNA resulted in fasting blood glucose levels above the therapeutic threshold of 70 mg/dL (indicated by the solid bar).

[0083] **FIG. 10C** are graphs showing fasting glucose levels in L-G6PC(-/-) mice injected with 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg of mRNA encoding G6PC S298C or mRNA encoding eGFP at 1 day after mRNA injection, plotted as a paired comparison between pre- (before) and post-treatment (after). $^{**}P<0.01$, $^{****}P<0.0001$ vs before (pre-treatment), paired *t*-test.

[0084] **FIGs. 11A-11B** are bar graphs showing the liver weights of L-G6PC(-/-) mice 24 hours after injection with a 0.2 mg/kg, 0.5 mg/kg, or 1.0 kg/mg dose of mRNA encoding G6PC S298C, or with mRNA encoding GFP (1 mg/kg), in total grams (FIG. 11A) and as % body weight (FIG. 11B). Results were compared to liver weight in L-G6PC(-/-) mice injected with GFP.

[0085] **FIG. 12** depicts representative images of livers in wild-type mice treated with PBS or in L-G6PC(-/-) mice 24 hours after injection with a 0.2 mg/kg, 0.5 mg/kg, or 1.0 kg/mg dose of mRNA encoding G6PC S298C, or with a 1 mg/kg dose of mRNA encoding GFP.

[0086] **FIG. 13A** is a graph showing hepatic G6P levels in wild-type mice treated with PBS or L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared to liver weight in L-G6PC(-/-) mice injected with GFP.

[0087] **FIG. 13B** is a graph showing hepatic glycogen levels in wild-type mice treated with PBS or L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared to L-G6PC(-/-) mice injected with GFP (eGFP).

[0088] **FIG. 13C** is a graph showing serum triglyceride levels in wild-type mice treated with PBS or in L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared to L-G6PC(-/-) mice injected with GFP (eGFP).

[0089] **FIG. 13D** is a graph showing serum ALT levels in wild-type mice treated with PBS or L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared to L-G6PC(-/-) mice injected with GFP (eGFP).

[0090] **FIG. 13E** is a graph showing serum lactate levels in wild-type mice treated with PBS or in L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared to L-G6PC(-/-) mice injected with GFP (eGFP).

[0091] **FIG. 13F** is a graph showing liver lactate levels in wild-type mice treated with PBS or in L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared to L-G6PC(-/-) mice injected with GFP (eGFP).

[0092] **FIG. 14A** is a graph showing hepatic G6Pase activity levels in wild-type mice treated with PBS or in L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection. Results were compared L-G6PC(-/-) mice injected with GFP (eGFP).

[0093] **FIG. 14B** includes a Western blot analysis showing the expression of G6PC protein in wild-type mice treated with PBS or in L-G6PC(-/-) mice injected with a 0.2 mg/kg, 0.5 mg/kg, or 1 mg/kg dose of mRNA encoding G6PC S298C, or mRNA encoding eGFP, at 1 day after mRNA injection, normalized to ERP72 protein expression. Results were compared L-G6PC(-/-) mice injected with GFP (eGFP).

[0094] **FIG. 14C** is a bar graph showing the results of the quantitative analysis of the Western blot analysis depicted in FIG. 14B.

[0095] **FIG. 15A** is a graph showing the amount of mRNA encoding human G6PC S298C and G6PC Q139R/Q247R/S298C in liver of wild-type mice at 6, 24, 72, 168, and 336 hours after injection of a 1 mg/kg dose of mRNA into the mice.

[0096] **FIGs. 15B-15C** are graphs showing G6PC protein expression in livers of wild type mice injected with mRNA encoding human G6PC S298C (S298C_CoV1) or human G6PC Q139R/Q247R/S298C (Q139R/Q247R/S298C_CoV1) over 336 hours post-injection.

[0097] **FIG. 15D** is a graph merging the graphs of FIGs. 12A-B.

[0098] **FIG. 15E** is a bar graph showing the area under the curve (AUC) values based on the expression of the G6PC S298C_CoV1 and G6PC Q139R/Q247R/S298C_CoV1proteins depicted in FIGs. 15A-15D.

[0099] **FIG. 15F** is a graph showing the introduced hG6PC protein activity in wild type mice injected with mRNA encoding human G6PC S298C_CoV1 or human G6PC Q139R/Q247R/S298C_CoV1 over 336 hours post-injection.

[0100] **FIG. 15G** is a bar graph showing the area under the curve (AUC) values based on the activity of the G6PC S298C_CoV1 and G6PC Q139R/Q247R/S298C_CoV1 proteins.

[0101] **FIG. 15H** is a graph showing hepatic G6PC protein expression in wild type mice injected with hG6PC S298C_CoV1 or G6PC_S298C_Astart_3XmiR142 mRNAs.

[0102] **FIG. 15I** is a graph showing hepatic G6Pase activity in wild type mice injected with hG6PC S298C_CoV1 or G6PC_S298C_Astart_3XmiR142 mRNAs.

[0103] **FIG. 16** is a graph showing fasting blood glucose levels in wild-type mice treated with PBS or L-G6PC(-/-) mice that were administered five doses of 0.25 mg/kg of mRNA encoding G6PC with the S298C substitution ("single"), mRNA encoding G6PC with the S298C/Q139R/Q247R substitutions ("triple"), or mRNA encoding eGFP.

[0104] **FIG. 17** is a graph showing liver weight in wild-type mice treated with PBS or mice that were administered 0.5 mg/kg of mRNA encoding G6PC with the S298C substitution ("single"), mRNA encoding G6PC with the S298C/Q139R/Q247R substitutions ("triple"), or mRNA encoding eGFP.

[0105] **FIG. 18** is a graph showing time-dependent fasting blood glucose levels (2.5 hours post-fasting) in wild-type mice (WT) treated with Tris-sucrose buffer or L-G6PC(-/-) mice that were administered mRNA encoding the G6PC S298C variant (G6PC_S298C_Astart) in two different formulations (CpdII/CpdI or CpdII/PEG-DMG) or eGFP mRNA in the CpdII/CpdI formulation.

[0106] **FIG. 19A** is a graph showing G6Pase protein expression (normalized to Erp72) in the liver of wild-type mice treated with mRNA encoding eGFP or mRNA encoding hG5PC with a S298C substitution and a 3′ UTR with (miR-142) or without (no miR-142) miR-142 target sites. **FIG. 19B** is a graph showing G6Pase protein expression (normalized to Erp72) in the spleen of wild-type mice treated with mRNA encoding eGFP or mRNA encoding hG6PC with a S298C substitution and a 3′ UTR with (miR-142) or without (no miR-142) miR-142 target sites.

[0107] **FIG. 20A** are graphs showing LNP uptake in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C

substitution in the presence of 10% fetal bovine serum (FBS; left) or 5% human serum (right); negative control represents non-transfected cells (triangles pointing down). Cells were transfected with Compound II/PEG-DMG (circles), Compound II/Compound I (squares), or MessengerMAX (ThermoFisher Scientific; triangle pointing up). **FIG. 20B** are graphs showing eGFP expression in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding GFP in the presence of 10% FBS (left) or 5% human serum (right). **FIG. 20C** are graphs showing hG6PC expression in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C substitution in the presence of 10% FBS (left) or 5% human serum (right). **FIG. 20D** are graphs showing the percentage of cells expressing transfected mRNA in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C substitution in the presence of 10% FBS (left) or 5% human serum (right). **FIGs. 20B-20D**: for each time point, from left to right: (i) Compound II/PEG-DMG, (ii) Compound II/Compound I, (iii) MessengerMAX; (iv) negative control.

[0108] **FIG. 21A** is a graph showing G6Pase expression, as a percentage of the level in wild-type mice, in wild-type mice administered PBS or K.G6PC $^{-/-}$ mice administered eGFP mRNA or hG6PC S298C mRNA. **FIG. 21B** is a graph showing G6Pase activity in wild-type mice administered PBS or K.G6PC -/- mice administered eGFP mRNA or hG6PC S298C mRNA. For **FIGs. 21A-21B**: For each time point, from left to right: (i) PBS, (ii) eGFP mRNA (K.G6PC$^{-/-}$ mice), (iii) hG6PC mRNA (K.G6PC$^{-/-}$ mice).

DETAILED DESCRIPTION

[0109] The present disclosure provides mRNA therapeutics for the treatment of glycogen storage disease type 1a (GSD-Ia). GSD-Ia is an autosomal recessive metabolic disorder affecting the ability to produce glucose, and individuals with GSD-Ia are typically hypoglycemic. GSD-Ia is caused by mutations in the *G6PC* gene, which codes for the enzyme glucose-6-phosphatase (G6Pase or G6PC). G6PC hydrolyzes glucose-6-phosphate (G6P) to glucose and phosphate late during gluconeogenesis and glycogenolysis. Without G6PC, G6P and glycogen abnormally accumulate in the liver and kidneys, and glucose production is impaired, particularly after the individual with GSD-Ia has fasted for a few hours. mRNA therapeutics are

particularly well-suited for the treatment of GSD-Ia as the technology provides for the intracellular delivery of mRNA encoding G6PC followed by *de novo* synthesis of functional G6PC protein within target cells. After delivery of mRNA to the target cells, the desired G6PC protein is expressed by the cells' own translational machinery, and hence, fully functional G6PC protein replaces the defective or missing protein.

[0110] One challenge associated with delivering nucleic acid-based therapeutics (*e.g.*, mRNA therapeutics) *in vivo* stems from the innate immune response which can occur when the body's immune system encounters foreign nucleic acids. Foreign mRNAs can activate the immune system *via* recognition through toll-like receptors (TLRs), in particular TLR7/8, which is activated by single-stranded RNA (ssRNA). In nonimmune cells, the recognition of foreign mRNA can occur through the retinoic acid-inducible gene I (RIG-I). Immune recognition of foreign mRNAs can result in unwanted cytokine effects including interleukin-1β (IL-1β) production, tumor necrosis factor-α (TNF-α) distribution and a strong type I interferon (type I IFN) response. This disclosure features the incorporation of different modified nucleotides within therapeutic mRNAs to minimize the immune activation and optimize the translation efficiency of mRNA to protein. Particular aspects feature a combination of nucleotide modification to reduce the innate immune response and sequence optimization, in particular, within the open reading frame (ORF) of therapeutic mRNAs encoding G6PC to enhance protein expression.

[0111] Certain embodiments of the mRNA therapeutic technology of the instant disclosure also feature delivery of mRNA encoding G6PC *via* a lipid nanoparticle (LNP) delivery system. Lipid nanoparticles (LNPs) are an ideal platform for the safe and effective delivery of mRNAs to target cells. LNPs have the unique ability to deliver nucleic acids by a mechanism involving cellular uptake, intracellular transport and endosomal release or endosomal escape. The instant invention features ionizable lipid-based LNPs combined with mRNA encoding G6PC which have improved properties when administered *in vivo*. Without being bound in theory, it is believed that the ionizable lipid-based LNP formulations of the invention have improved properties, for example, cellular uptake, intracellular transport and/or endosomal release or endosomal escape. LNPs administered by systemic route (*e.g.*, intravenous (IV) administration), for example, in a first administration, can accelerate the clearance of subsequently injected LNPs, for example, in further administrations. This phenomenon is known as accelerated blood clearance (ABC) and is a key challenge

when replacing deficient enzymes (*e.g.*, G6PC) in a therapeutic context. This is because repeat administration of mRNA therapeutics is in most instances essential to maintain necessary levels of enzyme in target tissues in subjects (*e.g.*, subjects suffering from GSD-Ia.) Repeat dosing challenges can be addressed on multiple levels. mRNA engineering and/or efficient delivery by LNPs can result in increased levels and or enhanced duration of protein (*e.g.*, G6PC) being expressed following a first dose of administration, which in turn, can lengthen the time between first dose and subsequent dosing. It is known that the ABC phenomenon is, at least in part, transient in nature, with the immune responses underlying ABC resolving after sufficient time following systemic administration. As such, increasing the duration of protein expression and/or activity following systemic delivery of an mRNA therapeutic of the disclosure in one aspect, combats the ABC phenomenon. Moreover, LNPs can be engineered to avoid immune sensing and/or recognition and can thus further avoid ABC upon subsequent or repeat dosing. An exemplary aspect of the disclosure features LNPs which have been engineered to have reduced ABC.

1. Glucose-6-Phosphatase (G6PC)

[0112] Glucose-6-phosphatase (G6Pase or G6PC, EC 3.1.3.9) catalyzes the hydrolysis of glucose-6-phosphate (G6P) to glucose and phosphate (Pi) in the terminal step of gluconeogenesis and glycogenolysis. G6PC is primarily present in the liver, and to a lesser extent in the kidneys and intestine. G6PC, encoded by the *G6PC* gene, is a multi-subunit integral membrane protein that resides on the endoplasmic reticulum (ER) membrane, and it hydrolyzes G6P to glucose and Pi in the ER lumen. The G6PC protein is 357 amino acids in length, with a molecular mass of about 40.5 kDa. It is composed of a catalytic subunit, and transporters for G6P, inorganic phosphate (Pi) and glucose. The amino acid sequence of human G6PC is provided in SEQ ID NO:199.

[0113] Normally, as blood glucose levels fall in between meals (fasting), G6P is produced during the terminal step of gluconeogenesis and glycogenolysis in the gluconeogenic organs (primarily liver, and also the kidneys and intestine) and it is hydrolyzed by G6PC to glucose and then released back into the blood. More than 80 mutations in *G6PC* have been identified, and several mutations are known in GSD-Ia patients to impair the phosphohydrolase activity of the enzyme, thereby impairing glucose production. See, Chou et al., Liver Research, 2017, 1:174-180, herein

incorporated by reference in its entirety. After a short fast of a few hours, individuals with GSD-Ia become hypoglycemic. G6P becomes elevated in cellular cytoplasm, which leads to an accumulation of glycogen and fat in the liver and kidneys, and impaired blood glucose homeostasis. The accumulation of glycogen promotes progressive hepatomegaly and nephromegaly. GSD-Ia patients can also develop other metabolic complications, including hypercholesterolemia, hypertriglyceridemia, hyperuricemia, hyperlipidemia, and lactic academia. Individuals with GSD-Ia can also develop longer term problems, including growth retardation, osteoporosis, gout, pulmonary hypertension, and renal disease. One long-term complication is the development of hepatic tumors in 75% of GSD-Ia patients over the age of 25 years old, of which, approximately 10% transform into malignant tumors.

[0114] GSD-1a is lethal if left untreated. The standard of care for GSD-1a is strict diet control that involves the frequent consumption of uncooked cornstarch. Currently, only a liver transplant is curative for GSD-Ia patients.

[0115] In certain aspects, the disclosure provides a polynucleotide (e.g., a RNA, e.g., a mRNA) comprising a nucleotide sequence (e.g., an open reading frame (ORF)) encoding a G6PC polypeptide. In some embodiments, the G6PC polypeptide of the invention is a wild type full length human G6PC protein. In some embodiments, the G6PC polypeptide of the invention is a variant, a peptide or a polypeptide containing a substitution, and insertion and/or an addition, a deletion and/or a covalent modification with respect to a wild-type G6PC sequence. In some embodiments, the G6PC polypeptide of the invention is a truncated human G6PC protein. In some embodiments, sequence tags or amino acids, can be added to the sequences encoded by the polynucleotides of the invention (e.g., at the N-terminal or C-terminal ends), e.g., for localization. In some embodiments, amino acid residues located at the carboxy, amino terminal, or internal regions of a polypeptide of the invention can optionally be deleted providing for fragments.

[0116] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) comprising a nucleotide sequence (e.g., an ORF) of the invention encodes a substitutional variant of a human G6PC sequence, which can comprise one, two, three, four, five, six, seven, eight, nine, tenor more than ten substitutions. In some embodiments, the substitutional variant can comprise one or more conservative amino acids substitutions. In other embodiments, the variant is an insertional variant. In other embodiments, the variant is a deletional variant.

[0117] G6PC protein fragments, functional protein domains, variants, and homologous proteins (orthologs) are also within the scope of the G6PC polypeptides of the disclosure. Nonlimiting examples of polypeptide encoded by the polynucleotides of the invention are shown in SEQ ID NO:83, SEQ ID NO:84, SEQ ID NO:194-206, 208, and 209.

2. Polynucleotides and Open Reading Frames (ORFs)

[0118] The instant invention features mRNAs for use in treating or preventing a glycogen storage disease type I (GSD-I) such as glycogen storage disease type Ia (GSD-Ia). The mRNAs featured for use in the invention are administered to subjects and encode human G6PC protein *in vivo*. Accordingly, the invention relates to polynucleotides, e.g., mRNA, comprising an open reading frame of linked nucleosides encoding human G6PC (SEQ ID NO:199), mutational variations thereof (e.g., SEQ ID NOs:194-198, 200-206, 208, and 209), truncated fragments thereof, and fusion proteins comprising G6PC. In some embodiments, the open reading frame is sequence-optimized. In particular embodiments, the invention provides sequence-optimized polynucleotides comprising nucleotides encoding the polypeptide sequence of human G6PC, or sequence having high sequence identity with those sequence optimized polynucleotides.

[0119] In certain aspects, the invention provides polynucleotides (e.g., a RNA such as an mRNA) that comprise a nucleotide sequence (e.g., an ORF) encoding one or more G6PC polypeptides. In some embodiments, the encoded G6PC polypeptide of the invention can be selected from:

(i) a full length G6PC polypeptide (e.g., having the same or essentially the same length as wild-type G6PC; e.g., wild type human G6PC);

(ii) a functional fragment of G6PC described herein (e.g., a truncated (e.g., deletion of carboxy, amino terminal, or internal regions) sequence shorter than G6PC; but still retaining G6PC enzymatic activity);

(iii) a variant thereof (e.g., full length or truncated G6PC proteins in which one or more amino acids (e.g., 1, 2, 3, 4, 5, 6, 7, 8, or more amino acids) have been replaced, e.g., variants that retain all or most of the G6PC activity of the polypeptide with respect to a reference protein (such as any natural or artificial variants known in the art)); or

(iv) a fusion protein comprising (i) a full length G6PC protein (e.g., SEQ ID NO:199), a functional fragment or a variant thereof (e.g., SEQ ID NOs:194-198, 200-206, 208, and 209), and (ii) a heterologous protein.

[0120] In certain embodiments, the encoded G6PC polypeptide is a mammalian G6PC polypeptide, such as a human G6PC polypeptide, a functional fragment or a variant thereof.

[0121] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention increases G6PC protein expression levels and/or detectable G6PC enzymatic activity levels in cells when introduced in those cells, e.g., by at least 2%, at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100%, compared to G6PC protein expression levels and/or detectable G6PC enzymatic activity levels in the cells prior to the administration of the polynucleotide of the invention. G6PC protein expression levels and/or G6PC enzymatic activity can be measured according to methods know in the art. In some embodiments, the polynucleotide is introduced to the cells *in vitro*. In some embodiments, the polynucleotide is introduced to the cells *in vivo*.

[0122] In some embodiments, the polynucleotides (e.g., a RNA, e.g., an mRNA) of the invention comprise a nucleotide sequence (e.g., an ORF) that encodes a wild-type human G6PC, e.g., (SEQ ID NO:199) or a variant thereof (e.g., SEQ ID NOs:194-198, 200-206, 208, and 209).

[0123] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a codon optimized nucleic acid sequence, wherein the open reading frame (ORF) of the codon optimized nucleic acid sequence is derived from a wild-type G6PC sequence (e.g., wild-type human G6PC). For example, for polynucleotides of invention comprising a sequence optimized ORF encoding G6PC, the corresponding wild type sequence is the native human G6PC. Similarly, for a sequence optimized mRNA encoding a functional variant or fragment of human G6PC, the corresponding wild type sequence is the corresponding variant or fragment from human G6PC.

[0124] In some embodiments, the polynucleotides (e.g., a RNA, e.g., an mRNA) of the invention comprise a nucleotide sequence encoding G6PC having the full-length sequence of human G6PC (i.e., including the initiator methionine).

[0125] In some embodiments, the polynucleotides (e.g., a RNA, e.g., an mRNA) of the invention comprise a nucleotide sequence (e.g., an ORF) encoding a mutant G6PC polypeptide. In some embodiments, the polynucleotides of the invention comprise an ORF encoding a G6PC polypeptide that comprises at least one-point mutation (e.g., 1, 2, 3, 4, 5, 6, 7, 8, or more point mutations) in the G6PC amino acid sequence and retains G6PC enzymatic activity. In some embodiments, the mutant G6PC polypeptide has a G6PC activity which is at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100% of the G6PC activity of the corresponding wild-type G6PC (i.e., the same G6PC polypeptide but without the mutation(s)). In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprising an ORF encoding a mutant G6PC polypeptide is sequence optimized.

[0126] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) that encodes a G6PC polypeptide with mutations that do not alter G6PC enzymatic activity. Such mutant G6PC polypeptides can be referred to as function-neutral. In some embodiments, the polynucleotide comprises an ORF that encodes a mutant G6PC polypeptide comprising one or more function-neutral point mutations.

[0127] In some embodiments, the mutant G6PC polypeptide has higher G6PC enzymatic activity than the corresponding wild-type G6PC. In some embodiments, the mutant G6PC polypeptide has a G6PC activity that is at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100% higher than the activity of the corresponding wild-type G6PC (i.e., the same G6PC protein but without the mutation(s)).

[0128] In some embodiments, the polynucleotides (e.g., a RNA, e.g., an mRNA) of the invention comprise a nucleotide sequence (e.g., an ORF) encoding a functional G6PC fragment, e.g., where one or more fragments correspond to a polypeptide subsequence of a wild type G6PC polypeptide and retain G6PC enzymatic activity. In some embodiments, the G6PC fragment has a G6PC activity which is at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least

40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100% of the G6PC activity of the corresponding full length G6PC. In some embodiments, the polynucleotides (e.g., a RNA, e.g., an mRNA) of the invention comprising an ORF encoding a functional G6PC fragment is sequence optimized.

[0129] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC fragment that has higher G6PC enzymatic activity than the corresponding full length G6PC. Thus, in some embodiments the G6PC fragment has a G6PC activity which is at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or at least 100% higher than the G6PC activity of the corresponding full length G6PC.

[0130] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC fragment that is at least 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 11%, 12%, 13%, 14%, 15%, 16%, 17%, 18%, 19%, 20%, 21%, 22%, 23%, 24% or 25% shorter than wild-type G6PC.

[0131] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the nucleotide sequence is at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% identical to the sequence of SEQ ID NOs:8, 11-20, 22-38, or 207.

[0132] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the nucleotide sequence has at least 70%, at least 71%, at least 72%, at least 73%, at least 74%, at least 75%, at least 76%, at least 77%, at least 78%, at least 79%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207.

[0133] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the nucleotide sequence has 70% to 100%, 75% to 100%, 80% to 100%, 85% to 100%, 70% to 95%, 80% to 95%, 70% to 85%, 75% to 90%, 80% to 95%, 70% to 75%, 75% to 80%, 80% to 85%, 85% to 90%, 90% to 95%, or 95% to 100%, sequence identity to a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207.

[0134] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises an ORF encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the polynucleotide comprises a nucleic acid sequence having 70% to 100%, 75% to 100%, 80% to 100%, 85% to 100%, 70% to 95%, 80% to 95%, 70% to 85%, 75% to 90%, 80% to 95%, 70% to 75%, 75% to 80%, 80% to 85%, 85% to 90%, 90% to 95%, or 95% to 100%, sequence identity to a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207.

[0135] In some embodiments the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the nucleotide sequence is between 70% and 90% identical; between 75% and 85% identical; between 76% and 84% identical; between 77% and 83% identical, between 77% and 82% identical, or between 78% and 81% identical to the sequence of SEQ ID NOs:8, 11-20, 22-38, or 207.

[0136] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises from about 900 to about 100,000 nucleotides (*e.g.,* from 900 to 1,000, from 900 to 1,100, from 900 to 1,200, from 900 to 1,300, from 900 to 1,400, from 900 to 1,500, from 1,000 to 1,100, from 1,000 to 1,100, from 1,000 to 1,200, from 1,000 to 1,300, from 1,000 to 1,400, from 1,000 to 1,500, from 1,187 to 1,200, from 1,187 to 1,400, from 1,187 to 1,600, from 1,187 to 1,800, from 1,187 to 2,000, from 1,187 to 3,000, from 1,187 to 5,000, from 1,187 to 7,000, from 1,187 to 10,000, from 1,187 to 25,000, from 1,187 to 50,000, from 1,187 to 70,000, or from 1,187 to 100,000).

[0137] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC

polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the length of the nucleotide sequence (e.g., an ORF) is at least 500 nucleotides in length (*e.g.,* at least or greater than about 500, 600, 700, 80, 900, 1,000, 1,050, 1,100, 1,187, 1,200, 1,300, 1,400, 1,500, 1,600, 1,700, 1,800, 1,900, 2,000, 2,100, 2,200, 2,300, 2,400, 2,500, 2,600, 2,700, 2,800, 2,900, 3,000, 3,100, 3,200, 3,300, 3,400, 3,500, 3,600, 3,700, 3,800, 3,900, 4,000, 4,100, 4,200, 4,300, 4,400, 4,500, 4,600, 4,700, 4,800, 4,900, 5,000, 5,100, 5,200, 5,300, 5,400, 5,500, 5,600, 5,700, 5,800, 5,900, 6,000, 7,000, 8,000, 9,000, 10,000, 20,000, 30,000, 40,000, 50,000, 60,000, 70,000, 80,000, 90,000 or up to and including 100,000 nucleotides).

[0138] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof) further comprises at least one nucleic acid sequence that is noncoding, e.g., a microRNA binding site. In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention further comprises a 5'-UTR (e.g., selected from the sequences of SEQ ID NOs: 3, 88-102, or 165-167 or selected from SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211) and a 3'UTR (e.g., selected from the sequences of SEQ ID NOs: 4, 104-112, 150, or 175 or selected from SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213). In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207. In a further embodiment, the polynucleotide (e.g., a RNA, e.g., an mRNA) comprises a 5' terminal cap (e.g., Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5' methylG cap, or an analog thereof) and a poly-A-tail region (e.g., about 100 nucleotides in length). In a further embodiment, the polynucleotide (e.g., a RNA, e.g., an mRNA) a comprises a 3' UTR comprising a nucleic acid sequence selected from the group consisting of SEQ ID NOs: 150 or 175, or any combination thereof. In some embodiments, the mRNA comprises a 3' UTR comprising a nucleic acid sequence of any of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213. In some embodiments, the mRNA comprises a polyA tail. In some instances, the poly A tail is 50-150 (SEQ ID NO:235), 75-150 (SEQ ID NO:218), 85-

150 (SEQ ID NO:219), 90-150 (SEQ ID NO:220), 90-120 (SEQ ID NO:221), 90-130 (SEQ ID NO:222), or 90-150 (SEQ ID NO:220) nucleotides in length. In some instances, the poly A tail is 100 nucleotides in length (SEQ ID NO:223).

[0139] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide is single stranded or double stranded.

[0140] In some embodiments, the polynucleotide of the invention comprising a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof) is DNA or RNA. In some embodiments, the polynucleotide of the invention is RNA. In some embodiments, the polynucleotide of the invention is, or functions as, a mRNA. In some embodiments, the mRNA comprises a nucleotide sequence (e.g., an ORF) that encodes at least one G6PC polypeptide, and is capable of being translated to produce the encoded G6PC polypeptide *in vitro, in vivo, in situ* or *ex vivo*.

[0141] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) comprises a sequence-optimized nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof, see e.g., SEQ ID NOs:8, 11-20, 22-38, or 207), wherein the polynucleotide comprises at least one chemically modified nucleobase, e.g., N1-methylpseudouracil or 5-methoxyuracil. In certain embodiments, all uracils in the polynucleotide are N1-methylpseudouracils. In other embodiments, all uracils in the polynucleotide are 5-methoxyuracils. In some embodiments, the polynucleotide further comprises a miRNA binding site, e.g., a miRNA binding site that binds to miR-142 and/or a miRNA binding site that binds to miR-126.

[0142] In some embodiments, the polynucleotide (e.g., a RNA, e.g., a mRNA) disclosed herein is formulated with a delivery agent comprising, e.g., a compound having the Formula (I), e.g., any of Compounds 1-232, e.g., Compound II; a compound having the Formula (III), (IV), (V), or (VI), e.g., any of Compounds 233-342, e.g., Compound VI; or a compound having the Formula (VIII), e.g., any of Compounds 419-428, e.g., Compound I, or any combination thereof. In some embodiments, the delivery agent comprises Compound II, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 50:10:38.5:1.5. In some embodiments, the delivery agent comprises Compound VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio in the range of about 30 to about

60 mol% Compound II or VI (or related suitable amino lipid) (e.g., 30-40, 40-45, 45-50, 50-55 or 55-60 mol% Compound II or VI (or related suitable amino lipid)), about 5 to about 20 mol% phospholipid (or related suitable phospholipid or "helper lipid") (e.g., 5-10, 10-15, or 15-20 mol% phospholipid (or related suitable phospholipid or "helper lipid")), about 20 to about 50 mol% cholesterol (or related sterol or "non-cationic" lipid) (e.g., about 20-30, 30-35, 35-40, 40-45, or 45-50 mol% cholesterol (or related sterol or "non-cationic" lipid)) and about 0.05 to about 10 mol% PEG lipid (or other suitable PEG lipid) (e.g., 0.05-1, 1-2, 2-3, 3-4, 4-5, 5-7, or 7-10 mol% PEG lipid (or other suitable PEG lipid)). An exemplary delivery agent can comprise mole ratios of, for example, 47.5:10.5:39.0:3.0 or 50:10:38.5:1.5. In certain instances, an exemplary delivery agent can comprise mole ratios of, for example, 47.5:10.5:39.0:3; 47.5:10:39.5:3; 47.5:11:39.5:2; 47.5:10.5:39.5:2.5; 47.5:11:39:2.5; 48.5:10:38.5:3; 48.5:10.5:39:2; 48.5:10.5:38.5:2.5; 48.5:10.5:39.5:1.5; 48.5:10.5:38.0:3; 47:10.5:39.5:3; 47:10:40.5:2.5; 47:11:40:2; 47:10.5:39.5:3; 48:10.5:38.5:3; 48:10:39.5:2.5; 48:11:39:2; or 48:10.5:38.5:3. In some embodiments, the delivery agent comprises Compound II or VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0. In some embodiments, the delivery agent comprises Compound II or VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 50:10:38.5:1.5.

[0143] In some embodiments, the polynucleotide of the disclosure is an mRNA that comprises a 5'-terminal cap (e.g., Cap 1), a 5'UTR (e.g., SEQ ID NO:3), an ORF sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207, a 3'UTR (e.g., SEQ ID NO:4, 150, 175, 177, or 178), and a poly A tail (e.g., about 100 nt in length), wherein all uracils in the polynucleotide are N1-methylpseudouracils. In some embodiments, the delivery agent comprises Compound II or Compound VI as the ionizable lipid and PEG-DMG or Compound I as the PEG lipid.

[0144] In some embodiments, the polynucleotide of the disclosure is an mRNA that comprises a 5'-terminal cap (e.g., Cap 1), a 5'UTR (e.g., SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211), an ORF sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207, a 3'UTR (e.g., SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213), and a poly A tail (e.g., about 100 nt in length), wherein all uracils in the polynucleotide are N1 methylpseudouracils. In

some embodiments, the delivery agent comprises Compound II or Compound VI as the ionizable lipid and PEG-DMG or Compound I as the PEG lipid.

3. Signal Sequences

[0145] The polynucleotides (e.g., a RNA, e.g., an mRNA) of the invention can also comprise nucleotide sequences that encode additional features that facilitate trafficking of the encoded polypeptides to therapeutically relevant sites. One such feature that aids in protein trafficking is the signal sequence, or targeting sequence. The peptides encoded by these signal sequences are known by a variety of names, including targeting peptides, transit peptides, and signal peptides. In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) comprises a nucleotide sequence (e.g., an ORF) that encodes a signal peptide operably linked to a nucleotide sequence that encodes a G6PC polypeptide described herein.

[0146] In some embodiments, the "signal sequence" or "signal peptide" is a polynucleotide or polypeptide, respectively, which is from about 30-210, e.g., about 45-80 or 15-60 nucleotides (e.g., about 20, 30, 40, 50, 60, or 70 amino acids) in length that, optionally, is incorporated at the 5' (or N-terminus) of the coding region or the polypeptide, respectively. Addition of these sequences results in trafficking the encoded polypeptide to a desired site, such as the endoplasmic reticulum or the mitochondria through one or more targeting pathways. Some signal peptides are cleaved from the protein, for example by a signal peptidase after the proteins are transported to the desired site.

[0147] In some embodiments, the polynucleotide of the invention comprises a nucleotide sequence encoding a G6PC polypeptide, wherein the nucleotide sequence further comprises a 5' nucleic acid sequence encoding a heterologous signal peptide.

4. Fusion Proteins

[0148] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) can comprise more than one nucleic acid sequence (e.g., an ORF) encoding a polypeptide of interest. In some embodiments, polynucleotides of the invention comprise a single ORF encoding a G6PC polypeptide, a functional fragment, or a variant thereof. However, in some embodiments, the polynucleotide of the invention can comprise more than one ORF, for example, a first ORF encoding a G6PC polypeptide (a first polypeptide of interest), a functional fragment, or a variant

thereof, and a second ORF expressing a second polypeptide of interest. In some embodiments, two or more polypeptides of interest can be genetically fused, i.e., two or more polypeptides can be encoded by the same ORF. In some embodiments, the polynucleotide can comprise a nucleic acid sequence encoding a linker (e.g., a G_4S (SEQ ID NO: 86) peptide linker or another linker known in the art) between two or more polypeptides of interest.

[0149] In some embodiments, a polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) can comprise two, three, four, or more ORFs, each expressing a polypeptide of interest.

[0150] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) can comprise a first nucleic acid sequence (e.g., a first ORF) encoding a G6PC polypeptide and a second nucleic acid sequence (e.g., a second ORF) encoding a second polypeptide of interest.

Linkers and Cleavable Peptides

[0151] In certain embodiments, the mRNAs of the disclosure encode more than one G6PC domain or a heterologous domain, referred to herein as multimer constructs. In certain embodiments of the multimer constructs, the mRNA further encodes a linker located between each domain. The linker can be, for example, a cleavable linker or protease-sensitive linker. In certain embodiments, the linker is selected from the group consisting of F2A linker, P2A linker, T2A linker, E2A linker, and combinations thereof. This family of self-cleaving peptide linkers, referred to as 2A peptides, has been described in the art (see for example, Kim, J.H. et al. (2011) PLoS ONE 6:e18556). In certain embodiments, the linker is an F2A linker. In certain embodiments, the linker is a GGGS (SEQ ID NO: 216) linker. In certain embodiments, the linker is a (GGGS)n (SEQ ID NO: 190) linker, wherein n =2, 3,4, or 5. In certain embodiments, the multimer construct contains three domains with intervening linkers, having the structure: domain-linker-domain-linker-domain e.g., G6PC domain-linker-G6PC domain-linker-G6PC domain.

[0152] In one embodiment, the cleavable linker is an F2A linker (e.g., having the amino acid sequence GSGVKQTLNFDLLKLAGDVESNPGP (SEQ ID NO:186)). In other embodiments, the cleavable linker is a T2A linker (e.g., having the amino acid sequence GSGEGRGSLLTCGDVEENPGP (SEQ ID NO:187)), a P2A linker (e.g., having the amino acid sequence GSGATNFSLLKQAGDVEENPGP (SEQ ID NO:188)) or an E2A linker (e.g., having the amino acid sequence

GSGQCTNYALLKLAGDVESNPGP (SEQ ID NO:189)). The skilled artisan will appreciate that other art-recognized linkers may be suitable for use in the constructs of the invention (e.g., encoded by the polynucleotides of the invention). The skilled artisan will likewise appreciate that other multi-cistronic constructs may be suitable for use in the invention. In exemplary embodiments, the construct design yields approximately equimolar amounts of intrabody and/or domain thereof encoded by the constructs of the invention.

[0153] In one embodiment, the self-cleaving peptide may be, but is not limited to, a 2A peptide. A variety of 2A peptides are known and available in the art and may be used, including e.g., the foot and mouth disease virus (FMDV) 2A peptide, the equine rhinitis A virus 2A peptide, the Thosea asigna virus 2A peptide, and the porcine teschovirus-1 2A peptide. 2A peptides are used by several viruses to generate two proteins from one transcript by ribosome-skipping, such that a normal peptide bond is impaired at the 2A peptide sequence, resulting in two discontinuous proteins being produced from one translation event. As a non-limiting example, the 2A peptide may have the protein sequence of SEQ ID NO: 188, fragments or variants thereof. In one embodiment, the 2A peptide cleaves between the last glycine and last proline. As another non-limiting example, the polynucleotides of the present invention may include a polynucleotide sequence encoding the 2A peptide having the protein sequence of fragments or variants of SEQ ID NO: 188. One example of a polynucleotide sequence encoding the 2A peptide is:GGAAGCGGAGCUACUAACUUCAGCCUGCUGAAGCAGGCUGGAGACGUGGAGGAGAACCCUGGACCU (SEQ ID NO:191). In one illustrative embodiment, a 2A peptide is encoded by the following sequence: 5'-UCCGGACUCAGAUCCGGGGAUCUCAAAAUUGUCGCUCCUGUCAAACAAACUCUUAACUUUGAUUUACUCAAACUGGCTGGGGAUGUAGAAAGCAAUCCAGGTCCACUC-3'(SEQ ID NO: 192). The polynucleotide sequence of the 2A peptide may be modified or codon optimized by the methods described herein and/or are known in the art.

[0154] In one embodiment, this sequence may be used to separate the coding regions of two or more polypeptides of interest. As a non-limiting example, the sequence encoding the F2A peptide may be between a first coding region A and a second coding region B (A-F2Apep-B). The presence of the F2A peptide results in the cleavage of the one long protein between the glycine and the proline at the end of the

F2A peptide sequence (NPGP (SEQ ID NO:224) is cleaved to result in NPG and P) thus creating separate protein A (with 21 amino acids of the F2A peptide attached, ending with NPG) and separate protein B (with 1 amino acid, P, of the F2A peptide attached). Likewise, for other 2A peptides (P2A, T2A and E2A), the presence of the peptide in a long protein results in cleavage between the glycine and proline at the end of the 2A peptide sequence (NPGP (SEQ ID NO:224) is cleaved to result in NPG and P). Protein A and protein B may be the same or different peptides or polypeptides of interest (e.g., a G6PC polypeptide such as full length human G6PCor a truncated version thereof).

5. Sequence Optimization of Nucleotide Sequence Encoding a G6PC Polypeptide

[0155] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention is sequence optimized. In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide, optionally, a nucleotide sequence (e.g., an ORF) encoding another polypeptide of interest, a 5'-UTR, a 3'-UTR, the 5' UTR or 3' UTR optionally comprising at least one microRNA binding site, optionally a nucleotide sequence encoding a linker, a polyA tail, or any combination thereof), in which the ORF(s) are sequence optimized.

[0156] A sequence-optimized nucleotide sequence, e.g., a codon-optimized mRNA sequence encoding a G6PC polypeptide, is a sequence comprising at least one synonymous nucleobase substitution with respect to a reference sequence (e.g., a wild type nucleotide sequence encoding a G6PC polypeptide).

[0157] A sequence-optimized nucleotide sequence can be partially or completely different in sequence from the reference sequence. For example, a reference sequence encoding polyserine uniformly encoded by UCU codons can be sequence-optimized by having 100% of its nucleobases substituted (for each codon, U in position 1 replaced by A, C in position 2 replaced by G, and U in position 3 replaced by C) to yield a sequence encoding polyserine which would be uniformly encoded by AGC codons. The percentage of sequence identity obtained from a global pairwise alignment between the reference polyserine nucleic acid sequence and the sequence-optimized polyserine nucleic acid sequence would be 0%. However, the protein products from both sequences would be 100% identical.

[0158] Some sequence optimization (also sometimes referred to codon optimization) methods are known in the art (and discussed in more detail below) and can be useful to achieve one or more desired results. These results can include, e.g., matching codon frequencies in certain tissue targets and/or host organisms to ensure proper folding; biasing G/C content to increase mRNA stability or reduce secondary structures; minimizing tandem repeat codons or base runs that can impair gene construction or expression; customizing transcriptional and translational control regions; inserting or removing protein trafficking sequences; removing/adding post translation modification sites in an encoded protein (*e.g.*, glycosylation sites); adding, removing or shuffling protein domains; inserting or deleting restriction sites; modifying ribosome binding sites and mRNA degradation sites; adjusting translational rates to allow the various domains of the protein to fold properly; and/or reducing or eliminating problem secondary structures within the polynucleotide. Sequence optimization tools, algorithms and services are known in the art, non-limiting examples include services from GeneArt (Life Technologies), DNA2.0 (Menlo Park CA) and/or proprietary methods.

[0159] Codon options for each amino acid are given in TABLE 1.

TABLE 1. Codon Options

Amino Acid	Single Letter Code	Codon Options
Isoleucine	I	AUU, AUC, AUA
Leucine	L	CUU, CUC, CUA, CUG, UUA, UUG
Valine	V	GUU, GUC, GUA, GUG
Phenylalanine	F	UUU, UUC
Methionine	M	AUG
Cysteine	C	UGU, UGC
Alanine	A	GCU, GCC, GCA, GCG
Glycine	G	GGU, GGC, GGA, GGG
Proline	P	CCU, CCC, CCA, CCG
Threonine	T	ACU, ACC, ACA, ACG
Serine	S	UCU, UCC, UCA, UCG, AGU, AGC
Tyrosine	Y	UAU, UAC
Tryptophan	W	UGG
Glutamine	Q	CAA, CAG
Asparagine	N	AAU, AAC
Histidine	H	CAU, CAC
Glutamic acid	E	GAA, GAG
Aspartic acid	D	GAU, GAC
Lysine	K	AAA, AAG
Arginine	R	CGU, CGC, CGA, CGG, AGA, AGG

Selenocysteine	Sec	UGA in mRNA in presence of Selenocysteine insertion element (SECIS)
Stop codons	Stop	UAA, UAG, UGA

[0160] In some embodiments, a polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a sequence-optimized nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide, a functional fragment, or a variant thereof, wherein the G6PC polypeptide, functional fragment, or a variant thereof encoded by the sequence-optimized nucleotide sequence has improved properties (e.g., compared to a G6PC polypeptide, functional fragment, or a variant thereof encoded by a reference nucleotide sequence that is not sequence optimized), e.g., improved properties related to expression efficacy after administration *in vivo*. Such properties include, but are not limited to, improving nucleic acid stability (e.g., mRNA stability), increasing translation efficacy in the target tissue, reducing the number of truncated proteins expressed, improving the folding or prevent misfolding of the expressed proteins, reducing toxicity of the expressed products, reducing cell death caused by the expressed products, increasing and/or decreasing protein aggregation.

[0161] In some embodiments, the sequence-optimized nucleotide sequence (e.g., an ORF) is codon optimized for expression in human subjects, having structural and/or chemical features that avoid one or more of the problems in the art, for example, features which are useful for optimizing formulation and delivery of nucleic acid-based therapeutics while retaining structural and functional integrity; overcoming a threshold of expression; improving expression rates; half-life and/or protein concentrations; optimizing protein localization; and avoiding deleterious bio-responses such as the immune response and/or degradation pathways.

[0162] In some embodiments, the polynucleotides of the invention comprise a nucleotide sequence (e.g., a nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide, a nucleotide sequence (e.g., an ORF) encoding another polypeptide of interest, a 5'-UTR, a 3'-UTR, a microRNA binding site, a nucleic acid sequence encoding a linker, or any combination thereof) that is sequence-optimized according to a method comprising:

(i) substituting at least one codon in a reference nucleotide sequence (e.g., an ORF encoding a G6PC polypeptide) with an alternative codon to increase or decrease uridine content to generate a uridine-modified sequence;

(ii) substituting at least one codon in a reference nucleotide sequence (e.g., an ORF encoding a G6PC polypeptide) with an alternative codon having a higher codon frequency in the synonymous codon set;

(iii) substituting at least one codon in a reference nucleotide sequence (e.g., an ORF encoding a G6PC polypeptide) with an alternative codon to increase G/C content; or

(iv) a combination thereof.

[0163] In some embodiments, the sequence-optimized nucleotide sequence (e.g., an ORF encoding a G6PC polypeptide) has at least one improved property with respect to the reference nucleotide sequence.

[0164] In some embodiments, the sequence optimization method is multiparametric and comprises one, two, three, four, or more methods disclosed herein and/or other optimization methods known in the art.

[0165] Features, which can be considered beneficial in some embodiments of the invention, can be encoded by or within regions of the polynucleotide and such regions can be upstream (5') to, downstream (3') to, or within the region that encodes the G6PC polypeptide. These regions can be incorporated into the polynucleotide before and/or after sequence-optimization of the protein encoding region or open reading frame (ORF). Examples of such features include, but are not limited to, untranslated regions (UTRs), microRNA sequences, Kozak sequences, oligo(dT) sequences, poly-A tail, and detectable tags and can include multiple cloning sites that can have XbaI recognition.

[0166] In some embodiments, the polynucleotide of the invention comprises a 5′ UTR, a 3′ UTR and/or a microRNA binding site. In some embodiments, the polynucleotide comprises two or more 5′ UTRs and/or 3′ UTRs, which can be the same or different sequences. In some embodiments, the polynucleotide comprises two or more microRNA binding sites, which can be the same or different sequences. Any portion of the 5’ UTR, 3’ UTR, and/or microRNA binding site, including none, can be sequence-optimized and can independently contain one or more different structural or chemical modifications, before and/or after sequence optimization.

[0167] In some embodiments, after optimization, the polynucleotide is reconstituted and transformed into a vector such as, but not limited to, plasmids, viruses, cosmids, and artificial chromosomes. For example, the optimized polynucleotide can be reconstituted and transformed into chemically competent *E. coli,* yeast, neurospora,

maize, drosophila, etc. where high copy plasmid-like or chromosome structures occur by methods described herein.

6. Sequence-Optimized Nucleotide Sequences Encoding G6PC Polypeptides

[0168] In some embodiments, the polynucleotide of the invention comprises a sequence-optimized nucleotide sequence encoding a G6PC polypeptide disclosed herein. In some embodiments, the polynucleotide of the invention comprises an open reading frame (ORF) encoding a G6PC polypeptide, wherein the ORF has been sequence optimized.

[0169] Exemplary sequence-optimized nucleotide sequences encoding human full length G6PC (wild-type and variants thereof) are set forth as SEQ ID NOs:8, 11-20, 22-38, or 207. In some embodiments, the sequence optimized G6PC sequences, fragments, and variants thereof are used to practice the methods disclosed herein.

[0170] In some embodiments, a polynucleotide of the present disclosure, for example a polynucleotide comprising an mRNA nucleotide sequence encoding a G6PC polypeptide, comprises from 5' to 3' end:

(i) a 5' cap provided herein, for example, Cap1;

(ii) a 5' UTR, such as the sequences provided herein, for example, SEQ ID NO: 3;

(iii) an open reading frame encoding a G6PC polypeptide, e.g., a sequence optimized nucleic acid sequence encoding G6PC set forth as SEQ ID NOs:8, 11-20, 22-38, or 207;

(iv) at least one stop codon (if not present at 5' terminus of 3'UTR);

(v) a 3' UTR, such as the sequences provided herein, for example, SEQ ID NO: 150, or 175; and

(vi) a poly-A tail provided above.

In some embodiments, a polynucleotide of the present disclosure, for example a polynucleotide comprising an mRNA nucleotide sequence encoding a G6PC polypeptide, comprises from 5' to 3' end:

(i) a 5' cap provided herein, for example, Cap1;

(ii) a 5' UTR, such as the sequences provided herein, for example, SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211;

(iii) an open reading frame encoding a G6PC polypeptide, e.g., a sequence optimized nucleic acid sequence encoding G6PC set forth as SEQ ID NOs:8, 11-20, 22-38, or 207;

(iv) at least one stop codon (if not present at 5' terminus of 3'UTR);

(v) a 3' UTR, such as the sequences provided herein, for example, SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213; and

(vi) a poly-A tail provided above.

In certain embodiments, all uracils in the polynucleotide are N1-methylpseudouracil (G5). In certain embodiments, all uracils in the polynucleotide are 5-methoxyuracil (G6).

[0171] The sequence-optimized nucleotide sequences disclosed herein are distinct from the corresponding wild type nucleotide acid sequences and from other known sequence-optimized nucleotide sequences, e.g., these sequence-optimized nucleic acids have unique compositional characteristics.

[0172] In some embodiments, the percentage of uracil or thymine nucleobases in a sequence-optimized nucleotide sequence (e.g., encoding a G6PC polypeptide, a functional fragment, or a variant thereof) is modified (e.g., reduced) with respect to the percentage of uracil or thymine nucleobases in the reference wild-type nucleotide sequence. Such a sequence is referred to as a uracil-modified or thymine-modified sequence. The percentage of uracil or thymine content in a nucleotide sequence can be determined by dividing the number of uracils or thymines in a sequence by the total number of nucleotides and multiplying by 100. In some embodiments, the sequence-optimized nucleotide sequence has a lower uracil or thymine content than the uracil or thymine content in the reference wild-type sequence. In some embodiments, the uracil or thymine content in a sequence-optimized nucleotide sequence of the invention is greater than the uracil or thymine content in the reference wild-type sequence and still maintain beneficial effects, e.g., increased expression and/or reduced Toll-Like Receptor (TLR) response when compared to the reference wild-type sequence.

[0173] Methods for optimizing codon usage are known in the art. For example, an ORF of any one or more of the sequences provided herein may be codon optimized. Codon optimization, in some embodiments, may be used to match codon frequencies in target and host organisms to ensure proper folding; bias GC content to increase

mRNA stability or reduce secondary structures; minimize tandem repeat codons or base runs that may impair gene construction or expression; customize transcriptional and translational control regions; insert or remove protein trafficking sequences; remove/add post translation modification sites in encoded protein (e.g., glycosylation sites); add, remove or shuffle protein domains; insert or delete restriction sites; modify ribosome binding sites and mRNA degradation sites; adjust translational rates to allow the various domains of the protein to fold properly; or reduce or eliminate problem secondary structures within the polynucleotide. Codon optimization tools, algorithms and services are known in the art - non-limiting examples include services from GeneArt (Life Technologies), DNA2.0 (Menlo Park CA) and/or proprietary methods. In some embodiments, the open reading frame (ORF) sequence is optimized using optimization algorithms.

7. Characterization of Sequence Optimized Nucleic Acids

[0174] In some embodiments of the invention, the polynucleotide (e.g., a RNA, e.g., an mRNA) comprising a sequence optimized nucleic acid disclosed herein encoding a G6PC polypeptide can be tested to determine whether at least one nucleic acid sequence property (e.g., stability when exposed to nucleases) or expression property has been improved with respect to the non-sequence optimized nucleic acid.

[0175] As used herein, "expression property" refers to a property of a nucleic acid sequence either *in vivo* (e.g., translation efficacy of a synthetic mRNA after administration to a subject in need thereof) or *in vitro* (e.g., translation efficacy of a synthetic mRNA tested in an *in vitro* model system). Expression properties include but are not limited to the amount of protein produced by an mRNA encoding a G6PC polypeptide after administration, and the amount of soluble or otherwise functional protein produced. In some embodiments, sequence optimized nucleic acids disclosed herein can be evaluated according to the viability of the cells expressing a protein encoded by a sequence optimized nucleic acid sequence (e.g., a RNA, e.g., an mRNA) encoding a G6PC polypeptide disclosed herein.

[0176] In a particular embodiment, a plurality of sequence optimized nucleic acids disclosed herein (e.g., a RNA, e.g., an mRNA) containing codon substitutions with respect to the non-optimized reference nucleic acid sequence can be characterized functionally to measure a property of interest, for example an expression property in an

in vitro model system, or *in vivo* in a target tissue or cell.

a. Optimization of Nucleic Acid Sequence Intrinsic Properties

[0177] In some embodiments of the invention, the desired property of the polynucleotide is an intrinsic property of the nucleic acid sequence. For example, the nucleotide sequence (e.g., a RNA, e.g., an mRNA) can be sequence optimized for *in vivo* or *in vitro* stability. In some embodiments, the nucleotide sequence can be sequence optimized for expression in a particular target tissue or cell. In some embodiments, the nucleic acid sequence is sequence optimized to increase its plasma half-life by preventing its degradation by endo and exonucleases.

[0178] In other embodiments, the nucleic acid sequence is sequence optimized to increase its resistance to hydrolysis in solution, for example, to lengthen the time that the sequence optimized nucleic acid or a pharmaceutical composition comprising the sequence optimized nucleic acid can be stored under aqueous conditions with minimal degradation.

[0179] In other embodiments, the sequence optimized nucleic acid can be optimized to increase its resistance to hydrolysis in dry storage conditions, for example, to lengthen the time that the sequence optimized nucleic acid can be stored after lyophilization with minimal degradation.

b. Nucleic Acids Sequence Optimized for Protein Expression

[0180] In some embodiments of the invention, the desired property of the polynucleotide is the level of expression of a G6PC polypeptide encoded by a sequence optimized sequence disclosed herein. Protein expression levels can be measured using one or more expression systems. In some embodiments, expression can be measured in cell culture systems, e.g., CHO cells or HEK293 cells. In some embodiments, expression can be measured using *in vitro* expression systems prepared from extracts of living cells, e.g., rabbit reticulocyte lysates, or *in vitro* expression systems prepared by assembly of purified individual components. In other embodiments, the protein expression is measured in an *in vivo* system, e.g., mouse, rabbit, monkey, etc.

[0181] In some embodiments, protein expression in solution form can be desirable. Accordingly, in some embodiments, a reference sequence can be sequence optimized to yield a sequence optimized nucleic acid sequence having optimized levels of

expressed proteins in soluble form. Levels of protein expression and other properties such as solubility, levels of aggregation, and the presence of truncation products (i.e., fragments due to proteolysis, hydrolysis, or defective translation) can be measured according to methods known in the art, for example, using electrophoresis (e.g., native or SDS-PAGE) or chromatographic methods (e.g., HPLC, size exclusion chromatography, etc.).

c. Optimization of Target Tissue or Target Cell Viability

[0182] In some embodiments, the expression of heterologous therapeutic proteins encoded by a nucleic acid sequence can have deleterious effects in the target tissue or cell, reducing protein yield, or reducing the quality of the expressed product (e.g., due to the presence of protein fragments or precipitation of the expressed protein in inclusion bodies), or causing toxicity.

[0183] Accordingly, in some embodiments of the invention, the sequence optimization of a nucleic acid sequence disclosed herein, e.g., a nucleic acid sequence encoding a G6PC polypeptide, can be used to increase the viability of target cells expressing the protein encoded by the sequence optimized nucleic acid.

[0184] Heterologous protein expression can also be deleterious to cells transfected with a nucleic acid sequence for autologous or heterologous transplantation. Accordingly, in some embodiments of the present disclosure the sequence optimization of a nucleic acid sequence disclosed herein can be used to increase the viability of target cells expressing the protein encoded by the sequence optimized nucleic acid sequence. Changes in cell or tissue viability, toxicity, and other physiological reaction can be measured according to methods known in the art.

d. Reduction of Immune and/or Inflammatory Response

[0185] In some cases, the administration of a sequence optimized nucleic acid encoding G6PC polypeptide or a functional fragment thereof can trigger an immune response, which could be caused by (i) the therapeutic agent (e.g., an mRNA encoding a G6PC polypeptide), or (ii) the expression product of such therapeutic agent (e.g., the G6PC polypeptide encoded by the mRNA), or (iv) a combination thereof. Accordingly, in some embodiments of the present disclosure the sequence optimization of nucleic acid sequence (e.g., an mRNA) disclosed herein can be used to decrease an immune or inflammatory response triggered by the administration of a

nucleic acid encoding a G6PC polypeptide or by the expression product of G6PC encoded by such nucleic acid.

[0186] In some aspects, an inflammatory response can be measured by detecting increased levels of one or more inflammatory cytokines using methods known in the art, e.g., ELISA. The term "inflammatory cytokine" refers to cytokines that are elevated in an inflammatory response. Examples of inflammatory cytokines include interleukin-6 (IL-6), CXCL1 (chemokine (C-X-C motif) ligand 1; also known as GROα, interferon-γ (IFNγ), tumor necrosis factor α (TNFα), interferon γ-induced protein 10 (IP-10), or granulocyte-colony stimulating factor (G-CSF). The term inflammatory cytokines includes also other cytokines associated with inflammatory responses known in the art, e.g., interleukin-1 (IL-1), interleukin-8 (IL-8), interleukin-12 (IL-12), interleukin-13 (Il-13), interferon α (IFN-α), etc.

8. Modified Nucleotide Sequences Encoding G6PC Polypeptides

[0187] In some embodiments, the polynucleotide (e.g., a RNA, e.g., an mRNA) of the invention comprises a chemically modified nucleobase, for example, a chemically modified uracil, e.g., pseudouracil, N1-methylpseudouracil, 5-methoxyuracil, or the like. In some embodiments, the mRNA is a uracil-modified sequence comprising an ORF encoding a G6PC polypeptide, wherein the mRNA comprises a chemically modified nucleobase, for example, a chemically modified uracil, e.g., pseudouracil, N1-methylpseudouracil, or 5-methoxyuracil.

[0188] In certain aspects of the invention, when the modified uracil base is connected to a ribose sugar, as it is in polynucleotides, the resulting modified nucleoside or nucleotide is referred to as modified uridine. In some embodiments, uracil in the polynucleotide is at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least 90%, at least 95%, at least 99%, or about 100% modified uracil. In one embodiment, uracil in the polynucleotide is at least 95% modified uracil. In another embodiment, uracil in the polynucleotide is 100% modified uracil.

[0189] In embodiments where uracil in the polynucleotide is at least 95% modified uracil overall uracil content can be adjusted such that an mRNA provides suitable protein expression levels while inducing little to no immune response. In some embodiments, the uracil content of the ORF is between about 100% and about 150%, between about 100% and about 110%, between about 105% and about 115%, between

about 110% and about 120%, between about 115% and about 125%, between about 120% and about 130%, between about 125% and about 135%, between about 130% and about 140%, between about 135% and about 145%, between about 140% and about 150% of the theoretical minimum uracil content in the corresponding wild-type ORF (%U_{TM}). In other embodiments, the uracil content of the ORF is between about 121% and about 136% or between 123% and 134% of the %U_{TM}. In some embodiments, the uracil content of the ORF encoding a G6PC polypeptide is about 115%, about 120%, about 125%, about 130%, about 135%, about 140%, about 145%, or about 150% of the %U_{TM}. In this context, the term "uracil" can refer to modified uracil and/or naturally occurring uracil.

[0190] In some embodiments, the uracil content in the ORF of the mRNA encoding a G6PC polypeptide of the invention is less than about 30%, about 25%, about 20%, about 15%, or about 10% of the total nucleobase content in the ORF. In some embodiments, the uracil content in the ORF is between about 10% and about 20% of the total nucleobase content in the ORF. In other embodiments, the uracil content in the ORF is between about 10% and about 25% of the total nucleobase content in the ORF. In one embodiment, the uracil content in the ORF of the mRNA encoding a G6PC polypeptide is less than about 20% of the total nucleobase content in the open reading frame. In this context, the term "uracil" can refer to modified uracil and/or naturally occurring uracil.

[0191] In further embodiments, the ORF of the mRNA encoding a G6PC polypeptide having modified uracil and adjusted uracil content has increased Cytosine (C), Guanine (G), or Guanine/Cytosine (G/C) content (absolute or relative). In some embodiments, the overall increase in C, G, or G/C content (absolute or relative) of the ORF is at least about 2%, at least about 3%, at least about 4%, at least about 5%, at least about 6%, at least about 7%, at least about 10%, at least about 15%, at least about 20%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, or at least about 100% relative to the G/C content (absolute or relative) of the wild-type ORF. In some embodiments, the G, the C, or the G/C content in the ORF is less than about 100%, less than about 90%, less than about 85%, or less than about 80% of the theoretical maximum G, C, or G/C content of the corresponding wild type nucleotide sequence encoding the G6PC polypeptide (%G_{TMX}; %C_{TMX}, or %G/C_{TMX}). In some embodiments, the increases in G and/or C content (absolute or relative) described

herein can be conducted by replacing synonymous codons with low G, C, or G/C content with synonymous codons having higher G, C, or G/C content. In other embodiments, the increase in G and/or C content (absolute or relative) is conducted by replacing a codon ending with U with a synonymous codon ending with G or C.

[0192] In further embodiments, the ORF of the mRNA encoding a G6PC polypeptide of the invention comprises modified uracil and has an adjusted uracil content containing less uracil pairs (UU) and/or uracil triplets (UUU) and/or uracil quadruplets (UUUU) than the corresponding wild-type nucleotide sequence encoding the G6PC polypeptide. In some embodiments, the ORF of the mRNA encoding a G6PC polypeptide of the invention contains no uracil pairs and/or uracil triplets and/or uracil quadruplets. In some embodiments, uracil pairs and/or uracil triplets and/or uracil quadruplets are reduced below a certain threshold, e.g., no more than 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 occurrences in the ORF of the mRNA encoding the G6PC polypeptide. In a particular embodiment, the ORF of the mRNA encoding the G6PC polypeptide of the invention contains less than 20, 19, 18, 17, 16, 15, 14, 13, 12, 11, 10, 9, 8, 7, 6, 5, 4, 3, 2, or 1 non-phenylalanine uracil pairs and/or triplets. In another embodiment, the ORF of the mRNA encoding the G6PC polypeptide contains no non-phenylalanine uracil pairs and/or triplets.

[0193] In further embodiments, the ORF of the mRNA encoding a G6PC polypeptide of the invention comprises modified uracil and has an adjusted uracil content containing less uracil-rich clusters than the corresponding wild-type nucleotide sequence encoding the G6PC polypeptide. In some embodiments, the ORF of the mRNA encoding the G6PC polypeptide of the invention contains uracil-rich clusters that are shorter in length than corresponding uracil-rich clusters in the corresponding wild-type nucleotide sequence encoding the G6PC polypeptide.

[0194] In further embodiments, alternative lower frequency codons are employed. At least about 5%, at least about 10%, at least about 15%, at least about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, at least about 95%, at least about 99%, or 100% of the codons in the G6PC polypeptide–encoding ORF of the modified uracil-comprising mRNA are substituted with alternative codons, each alternative codon having a codon frequency lower than the codon frequency of the substituted codon in the synonymous codon set.

The ORF also has adjusted uracil content, as described above. In some embodiments, at least one codon in the ORF of the mRNA encoding the G6PC polypeptide is substituted with an alternative codon having a codon frequency lower than the codon frequency of the substituted codon in the synonymous codon set.

[0195] In some embodiments, the adjusted uracil content, G6PC polypeptide-encoding ORF of the modified uracil-comprising mRNA exhibits expression levels of G6PC when administered to a mammalian cell that are higher than expression levels of G6PC from the corresponding wild-type mRNA. In some embodiments, the mammalian cell is a mouse cell, a rat cell, or a rabbit cell. In other embodiments, the mammalian cell is a monkey cell or a human cell. In some embodiments, the human cell is a HeLa cell, a BJ fibroblast cell, or a peripheral blood mononuclear cell (PBMC). In some embodiments, G6PC is expressed a level higher than expression levels of G6PC from the corresponding wild-type mRNA when the mRNA is administered to a mammalian cell *in vivo*. In some embodiments, the mRNA is administered to mice, rabbits, rats, monkeys, or humans. In one embodiment, mice are null mice. In some embodiments, the mRNA is administered to mice in an amount of about 0.01 mg/kg, about 0.05 mg/kg, about 0.1 mg/kg, or 0.2 mg/kg or about 0.5 mg/kg. In some embodiments, the mRNA is administered intravenously or intramuscularly. In other embodiments, the G6PC polypeptide is expressed when the mRNA is administered to a mammalian cell *in vitro*. In some embodiments, the expression is increased by at least about 2-fold, at least about 5-fold, at least about 10-fold, at least about 50-fold, at least about 500-fold, at least about 1500-fold, or at least about 3000-fold. In other embodiments, the expression is increased by at least about 10%, about 20%, about 30%, about 40%, about 50%, 60%, about 70%, about 80%, about 90%, or about 100%.

[0196] In some embodiments, adjusted uracil content, G6PC polypeptide-encoding ORF of the modified uracil-comprising mRNA exhibits increased stability. In some embodiments, the mRNA exhibits increased stability in a cell relative to the stability of a corresponding wild-type mRNA under the same conditions. In some embodiments, the mRNA exhibits increased stability including resistance to nucleases, thermal stability, and/or increased stabilization of secondary structure. In some embodiments, increased stability exhibited by the mRNA is measured by determining the half-life of the mRNA (e.g., in a plasma, serum, cell, or tissue sample) and/or determining the area under the curve (AUC) of the protein expression

by the mRNA over time (e.g., in vitro or in vivo). An mRNA is identified as having increased stability if the half-life and/or the AUC is greater than the half-life and/or the AUC of a corresponding wild-type mRNA under the same conditions.

[0197] In some embodiments, the mRNA of the present invention induces a detectably lower immune response (e.g., innate or acquired) relative to the immune response induced by a corresponding wild-type mRNA under the same conditions. In other embodiments, the mRNA of the present disclosure induces a detectably lower immune response (e.g., innate or acquired) relative to the immune response induced by an mRNA that encodes for a G6PC polypeptide but does not comprise modified uracil under the same conditions, or relative to the immune response induced by an mRNA that encodes for a G6PC polypeptide and that comprises modified uracil but that does not have adjusted uracil content under the same conditions. The innate immune response can be manifested by increased expression of pro-inflammatory cytokines, activation of intracellular PRRs (RIG-I, MDA5, etc.), cell death, and/or termination or reduction in protein translation. In some embodiments, a reduction in the innate immune response can be measured by expression or activity level of Type 1 interferons (e.g., IFN-α, IFN-β, IFN-κ, IFN-δ, IFN-ε, IFN-τ, IFN-ω, and IFN-ζ) or the expression of interferon-regulated genes such as the toll-like receptors (e.g., TLR7 and TLR8), and/or by decreased cell death following one or more administrations of the mRNA of the invention into a cell.

[0198] In some embodiments, the expression of Type-1 interferons by a mammalian cell in response to the mRNA of the present disclosure is reduced by at least 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, 99%, 99.9%, or greater than 99.9% relative to a corresponding wild-type mRNA, to an mRNA that encodes a G6PC polypeptide but does not comprise modified uracil, or to an mRNA that encodes a G6PC polypeptide and that comprises modified uracil but that does not have adjusted uracil content. In some embodiments, the interferon is IFN-β. In some embodiments, cell death frequency caused by administration of mRNA of the present disclosure to a mammalian cell is 10%, 25%, 50%, 75%, 85%, 90%, 95%, or over 95% less than the cell death frequency observed with a corresponding wild-type mRNA, an mRNA that encodes for a G6PC polypeptide but does not comprise modified uracil, or an mRNA that encodes for a G6PC polypeptide and that comprises modified uracil but that does not have adjusted uracil content. In some embodiments, the mammalian cell is a BJ fibroblast cell. In other embodiments, the mammalian cell

is a splenocyte. In some embodiments, the mammalian cell is that of a mouse or a rat. In other embodiments, the mammalian cell is that of a human. In one embodiment, the mRNA of the present disclosure does not substantially induce an innate immune response of a mammalian cell into which the mRNA is introduced.

9. Methods for Modifying Polynucleotides

[0199] The disclosure includes modified polynucleotides comprising a polynucleotide described herein (e.g., a polynucleotide, e.g. mRNA, comprising a nucleotide sequence encoding a G6PC polypeptide). The modified polynucleotides can be chemically modified and/or structurally modified. When the polynucleotides of the present invention are chemically and/or structurally modified the polynucleotides can be referred to as "modified polynucleotides."

[0200] The present disclosure provides for modified nucleosides and nucleotides of a polynucleotide (*e.g.*, RNA polynucleotides, such as mRNA polynucleotides) encoding a G6PC polypeptide. A "nucleoside" refers to a compound containing a sugar molecule (*e.g.*, a pentose or ribose) or a derivative thereof in combination with an organic base (*e.g.*, a purine or pyrimidine) or a derivative thereof (also referred to herein as "nucleobase"). A "nucleotide" refers to a nucleoside including a phosphate group. Modified nucleotides can be synthesized by any useful method, such as, for example, chemically, enzymatically, or recombinantly, to include one or more modified or non-natural nucleosides. Polynucleotides can comprise a region or regions of linked nucleosides. Such regions can have variable backbone linkages. The linkages can be standard phosphodiester linkages, in which case the polynucleotides would comprise regions of nucleotides.

[0201] The modified polynucleotides disclosed herein can comprise various distinct modifications. In some embodiments, the modified polynucleotides contain one, two, or more (optionally different) nucleoside or nucleotide modifications. In some embodiments, a modified polynucleotide, introduced to a cell can exhibit one or more desirable properties, e.g., improved protein expression, reduced immunogenicity, or reduced degradation in the cell, as compared to an unmodified polynucleotide.

[0202] In some embodiments, a polynucleotide of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) is structurally modified. As used herein, a "structural" modification is one in which two or more linked nucleosides are inserted, deleted, duplicated, inverted or randomized in

a polynucleotide without significant chemical modification to the nucleotides themselves. Because chemical bonds will necessarily be broken and reformed to affect a structural modification, structural modifications are of a chemical nature and hence are chemical modifications. However, structural modifications will result in a different sequence of nucleotides. For example, the polynucleotide "ATCG" can be chemically modified to "AT-5meC-G". The same polynucleotide can be structurally modified from "ATCG" to "ATCCCG". Here, the dinucleotide "CC" has been inserted, resulting in a structural modification to the polynucleotide.

[0203] Therapeutic compositions of the present disclosure comprise, in some embodiments, at least one nucleic acid (e.g., RNA) having an open reading frame encoding G6PC (e.g., SEQ ID NOs:8, 11-20, 22-38, or 207), wherein the nucleic acid comprises nucleotides and/or nucleosides that can be standard (unmodified) or modified as is known in the art. In some embodiments, nucleotides and nucleosides of the present disclosure comprise modified nucleotides or nucleosides. Such modified nucleotides and nucleosides can be naturally-occurring modified nucleotides and nucleosides or non-naturally occurring modified nucleotides and nucleosides. Such modifications can include those at the sugar, backbone, or nucleobase portion of the nucleotide and/or nucleoside as are recognized in the art.

[0204] In some embodiments, a naturally-occurring modified nucleotide or nucleotide of the disclosure is one as is generally known or recognized in the art. Non-limiting examples of such naturally occurring modified nucleotides and nucleotides can be found, inter alia, in the widely recognized MODOMICS database.

[0205] In some embodiments, a non-naturally occurring modified nucleotide or nucleoside of the disclosure is one as is generally known or recognized in the art. Non-limiting examples of such non-naturally occurring modified nucleotides and nucleosides can be found, inter alia, in published US application Nos. PCT/US2012/058519; PCT/US2013/075177; PCT/US2014/058897; PCT/US2014/058891; PCT/US2014/070413; PCT/US2015/36773; PCT/US2015/36759; PCT/US2015/36771; or PCT/IB2017/051367 all of which are incorporated by reference herein.

[0206] In some embodiments, at least one RNA (e.g., mRNA) of the present disclosure is not chemically modified and comprises the standard ribonucleotides consisting of adenosine, guanosine, cytosine and uridine. In some embodiments, nucleotides and nucleosides of the present disclosure comprise standard nucleoside

residues such as those present in transcribed RNA (e.g. A, G, C, or U). In some embodiments, nucleotides and nucleosides of the present disclosure comprise standard deoxyribonucleosides such as those present in DNA (e.g. dA, dG, dC, or dT).

[0207] Hence, nucleic acids of the disclosure (*e.g.*, DNA nucleic acids and RNA nucleic acids, such as mRNA nucleic acids) can comprise standard nucleotides and nucleosides, naturally-occurring nucleotides and nucleosides, non-naturally-occurring nucleotides and nucleosides, or any combination thereof.

[0208] Nucleic acids of the disclosure (*e.g.*, DNA nucleic acids and RNA nucleic acids, such as mRNA nucleic acids), in some embodiments, comprise various (more than one) different types of standard and/or modified nucleotides and nucleosides. In some embodiments, a particular region of a nucleic acid contains one, two or more (optionally different) types of standard and/or modified nucleotides and nucleosides.

[0209] In some embodiments, a modified RNA nucleic acid (*e.g.*, a modified mRNA nucleic acid), introduced to a cell or organism, exhibits reduced degradation in the cell or organism, respectively, relative to an unmodified nucleic acid comprising standard nucleotides and nucleosides.

[0210] In some embodiments, a modified RNA nucleic acid (*e.g.*, a modified mRNA nucleic acid), introduced into a cell or organism, may exhibit reduced immunogenicity in the cell or organism, respectively (*e.g.*, a reduced innate response) relative to an unmodified nucleic acid comprising standard nucleotides and nucleosides.

[0211] Nucleic acids (*e.g.*, RNA nucleic acids, such as mRNA nucleic acids), in some embodiments, comprise non-natural modified nucleotides that are introduced during synthesis or post-synthesis of the nucleic acids to achieve desired functions or properties. The modifications may be present on internucleotide linkages, purine or pyrimidine bases, or sugars. The modification may be introduced with chemical synthesis or with a polymerase enzyme at the terminal of a chain or anywhere else in the chain. Any of the regions of a nucleic acid may be chemically modified.

[0212] The present disclosure provides for modified nucleosides and nucleotides of a nucleic acid (*e.g.*, RNA nucleic acids, such as mRNA nucleic acids). A "nucleoside" refers to a compound containing a sugar molecule (*e.g.*, a pentose or ribose) or a derivative thereof in combination with an organic base (*e.g.*, a purine or pyrimidine) or a derivative thereof (also referred to herein as "nucleobase"). A "nucleotide" refers to a nucleoside, including a phosphate group. Modified nucleotides may by

synthesized by any useful method, such as, for example, chemically, enzymatically, or recombinantly, to include one or more modified or non-natural nucleosides. Nucleic acids can comprise a region or regions of linked nucleosides. Such regions may have variable backbone linkages. The linkages can be standard phosphodiester linkages, in which case the nucleic acids would comprise regions of nucleotides.

[0213] Modified nucleotide base pairing encompasses not only the standard adenosine-thymine, adenosine-uracil, or guanosine-cytosine base pairs, but also base pairs formed between nucleotides and/or modified nucleotides comprising non-standard or modified bases, wherein the arrangement of hydrogen bond donors and hydrogen bond acceptors permits hydrogen bonding between a non-standard base and a standard base or between two complementary non-standard base structures, such as, for example, in those nucleic acids having at least one chemical modification. One example of such non-standard base pairing is the base pairing between the modified nucleotide inosine and adenine, cytosine or uracil. Any combination of base/sugar or linker may be incorporated into nucleic acids of the present disclosure.

[0214] In some embodiments, modified nucleobases in nucleic acids (*e.g.*, RNA nucleic acids, such as mRNA nucleic acids) comprise N1-methyl-pseudouridine (m1ψ), 1-ethyl-pseudouridine (e1ψ), 5-methoxy-uridine (mo5U), 5-methyl-cytidine (m5C), and/or pseudouridine (ψ). In some embodiments, modified nucleobases in nucleic acids (*e.g.*, RNA nucleic acids, such as mRNA nucleic acids) comprise 5-methoxymethyl uridine, 5-methylthio uridine, 1-methoxymethyl pseudouridine, 5-methyl cytidine, and/or 5-methoxy cytidine. In some embodiments, the polyribonucleotide includes a combination of at least two (*e.g.*, 2, 3, 4 or more) of any of the aforementioned modified nucleobases, including but not limited to chemical modifications.

[0215] In some embodiments, a RNA nucleic acid of the disclosure comprises N1-methyl-pseudouridine (m1ψ) substitutions at one or more or all uridine positions of the nucleic acid.

[0216] In some embodiments, a RNA nucleic acid of the disclosure comprises N1-methyl-pseudouridine (m1ψ) substitutions at one or more or all uridine positions of the nucleic acid and 5-methyl cytidine substitutions at one or more or all cytidine positions of the nucleic acid.

[0217] In some embodiments, a RNA nucleic acid of the disclosure comprises pseudouridine (ψ) substitutions at one or more or all uridine positions of the nucleic acid.

[0218] In some embodiments, a RNA nucleic acid of the disclosure comprises pseudouridine (ψ) substitutions at one or more or all uridine positions of the nucleic acid and 5-methyl cytidine substitutions at one or more or all cytidine positions of the nucleic acid.

[0219] In some embodiments, a RNA nucleic acid of the disclosure comprises uridine at one or more or all uridine positions of the nucleic acid.

[0220] In some embodiments, nucleic acids (*e.g.*, RNA nucleic acids, such as mRNA nucleic acids) are uniformly modified (*e.g.*, fully modified, modified throughout the entire sequence) for a particular modification. For example, a nucleic acid can be uniformly modified with N1-methyl-pseudouridine, meaning that all uridine residues in the mRNA sequence are replaced with N1-methyl-pseudouridine. Similarly, a nucleic acid can be uniformly modified for any type of nucleoside residue present in the sequence by replacement with a modified residue such as those set forth above.

[0221] The nucleic acids of the present disclosure may be partially or fully modified along the entire length of the molecule. For example, one or more or all or a given type of nucleotide (*e.g.*, purine or pyrimidine, or any one or more or all of A, G, U, C) may be uniformly modified in a nucleic acid of the disclosure, or in a predetermined sequence region thereof (*e.g.*, in the mRNA including or excluding the polyA tail). In some embodiments, all nucleotides X in a nucleic acid of the present disclosure (or in a sequence region thereof) are modified nucleotides, wherein X may be any one of nucleotides A, G, U, C, or any one of the combinations A+G, A+U, A+C, G+U, G+C, U+C, A+G+U, A+G+C, G+U+C or A+G+C.

[0222] The nucleic acid may contain from about 1% to about 100% modified nucleotides (either in relation to overall nucleotide content, or in relation to one or more types of nucleotide, *i.e.*, any one or more of A, G, U or C) or any intervening percentage (*e.g.*, from 1% to 20%, from 1% to 25%, from 1% to 50%, from 1% to 60%, from 1% to 70%, from 1% to 80%, from 1% to 90%, from 1% to 95%, from 10% to 20%, from 10% to 25%, from 10% to 50%, from 10% to 60%, from 10% to 70%, from 10% to 80%, from 10% to 90%, from 10% to 95%, from 10% to 100%, from 20% to 25%, from 20% to 50%, from 20% to 60%, from 20% to 70%, from 20% to 80%, from 20% to 90%, from 20% to 95%, from 20% to 100%, from 50% to 60%,

from 50% to 70%, from 50% to 80%, from 50% to 90%, from 50% to 95%, from 50% to 100%, from 70% to 80%, from 70% to 90%, from 70% to 95%, from 70% to 100%, from 80% to 90%, from 80% to 95%, from 80% to 100%, from 90% to 95%, from 90% to 100%, and from 95% to 100%). It will be understood that any remaining percentage is accounted for by the presence of unmodified A, G, U, or C.

[0223] The nucleic acids may contain at a minimum 1% and at maximum 100% modified nucleotides, or any intervening percentage, such as at least 5% modified nucleotides, at least 10% modified nucleotides, at least 25% modified nucleotides, at least 50% modified nucleotides, at least 80% modified nucleotides, or at least 90% modified nucleotides. For example, the nucleic acids may contain a modified pyrimidine such as a modified uracil or cytosine. In some embodiments, at least 5%, at least 10%, at least 25%, at least 50%, at least 80%, at least 90% or 100% of the uracil in the nucleic acid is replaced with a modified uracil (*e.g.*, a 5-substituted uracil). The modified uracil can be replaced by a compound having a single unique structure, or can be replaced by a plurality of compounds having different structures (*e.g.*, 2, 3, 4 or more unique structures). In some embodiments, at least 5%, at least 10%, at least 25%, at least 50%, at least 80%, at least 90% or 100% of the cytosine in the nucleic acid is replaced with a modified cytosine (*e.g.*, a 5-substituted cytosine). The modified cytosine can be replaced by a compound having a single unique structure, or can be replaced by a plurality of compounds having different structures (*e.g.*, 2, 3, 4 or more unique structures).

10. Untranslated Regions (UTRs)

[0224] Translation of a polynucleotide comprising an open reading frame encoding a polypeptide can be controlled and regulated by a variety of mechanisms that are provided by various cis-acting nucleic acid structures. For example, naturally-occurring, cis-acting RNA elements that form hairpins or other higher-order (e.g., pseudoknot) intramolecular mRNA secondary structures can provide a translational regulatory activity to a polynucleotide, wherein the RNA element influences or modulates the initiation of polynucleotide translation, particularly when the RNA element is positioned in the 5′ UTR close to the 5’-cap structure (Pelletier and Sonenberg (1985) Cell 40(3):515-526; Kozak (1986) Proc Natl Acad Sci 83:2850-2854).

[0225] Untranslated regions (UTRs) are nucleic acid sections of a polynucleotide before a start codon (5' UTR) and after a stop codon (3' UTR) that are not translated. In some embodiments, a polynucleotide (e.g., a ribonucleic acid (RNA), e.g., a messenger RNA (mRNA)) of the invention comprising an open reading frame (ORF) encoding a G6PC polypeptide further comprises UTR (e.g., a 5′ UTR or functional fragment thereof, a 3′ UTR or functional fragment thereof, or a combination thereof).

[0226] Cis-acting RNA elements can also affect translation elongation, being involved in numerous frameshifting events (Namy et al., (2004) Mol Cell 13(2):157-168). Internal ribosome entry sequences (IRES) represent another type of cis-acting RNA element that are typically located in 5′ UTRs, but have also been reported to be found within the coding region of naturally-occurring mRNAs (Holcik et al. (2000) Trends Genet 16(10):469-473). In cellular mRNAs, IRES often coexist with the 5′-cap structure and provide mRNAs with the functional capacity to be translated under conditions in which cap-dependent translation is compromised (Gebauer et al., (2012) Cold Spring Harb Perspect Biol 4(7):a012245). Another type of naturally-occurring cis-acting RNA element comprises upstream open reading frames (uORFs). Naturally-occurring uORFs occur singularly or multiply within the 5′ UTRs of numerous mRNAs and influence the translation of the downstream major ORF, usually negatively (with the notable exception of GCN4 mRNA in yeast and ATF4 mRNA in mammals, where uORFs serve to promote the translation of the downstream major ORF under conditions of increased eIF2 phosphorylation (Hinnebusch (2005) Annu Rev Microbiol 59:407-450)). Additional exemplary translational regulatory activities provided by components, structures, elements, motifs, and/or specific sequences comprising polynucleotides (e.g., mRNA) include, but are not limited to, mRNA stabilization or destabilization (Baker & Parker (2004) Curr Opin Cell Biol 16(3):293-299), translational activation (Villalba et al., (2011) Curr Opin Genet Dev 21(4):452-457), and translational repression (Blumer et al., (2002) Mech Dev 110(1-2):97-112). Studies have shown that naturally-occurring, cis-acting RNA elements can confer their respective functions when used to modify, by incorporation into, heterologous polynucleotides (Goldberg-Cohen et al., (2002) J Biol Chem 277(16):13635-13640).

Modified Polynucleotides Comprising Functional RNA Elements

[0227] The present disclosure provides synthetic polynucleotides comprising a modification (e.g., an RNA element), wherein the modification provides a desired

translational regulatory activity. In some embodiments, the disclosure provides a polynucleotide comprising a 5' untranslated region (UTR), an initiation codon, a full open reading frame encoding a polypeptide, a 3' UTR, and at least one modification, wherein the at least one modification provides a desired translational regulatory activity, for example, a modification that promotes and/or enhances the translational fidelity of mRNA translation. In some embodiments, the desired translational regulatory activity is a cis-acting regulatory activity. In some embodiments, the desired translational regulatory activity is an increase in the residence time of the 43S pre-initiation complex (PIC) or ribosome at, or proximal to, the initiation codon. In some embodiments, the desired translational regulatory activity is an increase in the initiation of polypeptide synthesis at or from the initiation codon. In some embodiments, the desired translational regulatory activity is an increase in the amount of polypeptide translated from the full open reading frame. In some embodiments, the desired translational regulatory activity is an increase in the fidelity of initiation codon decoding by the PIC or ribosome. In some embodiments, the desired translational regulatory activity is inhibition or reduction of leaky scanning by the PIC or ribosome. In some embodiments, the desired translational regulatory activity is a decrease in the rate of decoding the initiation codon by the PIC or ribosome. In some embodiments, the desired translational regulatory activity is inhibition or reduction in the initiation of polypeptide synthesis at any codon within the mRNA other than the initiation codon. In some embodiments, the desired translational regulatory activity is inhibition or reduction of the amount of polypeptide translated from any open reading frame within the mRNA other than the full open reading frame. In some embodiments, the desired translational regulatory activity is inhibition or reduction in the production of aberrant translation products. In some embodiments, the desired translational regulatory activity is a combination of one or more of the foregoing translational regulatory activities.

[0228] Accordingly, the present disclosure provides a polynucleotide, e.g., an mRNA, comprising an RNA element that comprises a sequence and/or an RNA secondary structure(s) that provides a desired translational regulatory activity as described herein. In some aspects, the mRNA comprises an RNA element that comprises a sequence and/or an RNA secondary structure(s) that promotes and/or enhances the translational fidelity of mRNA translation. In some aspects, the mRNA comprises an RNA element that comprises a sequence and/or an RNA secondary structure(s) that

provides a desired translational regulatory activity, such as inhibiting and/or reducing leaky scanning. In some aspects, the disclosure provides an mRNA that comprises an RNA element that comprises a sequence and/or an RNA secondary structure(s) that inhibits and/or reduces leaky scanning thereby promoting the translational fidelity of the mRNA.

[0229] In some embodiments, the RNA element comprises natural and/or modified nucleotides. In some embodiments, the RNA element comprises of a sequence of linked nucleotides, or derivatives or analogs thereof, that provides a desired translational regulatory activity as described herein. In some embodiments, the RNA element comprises a sequence of linked nucleotides, or derivatives or analogs thereof, that forms or folds into a stable RNA secondary structure, wherein the RNA secondary structure provides a desired translational regulatory activity as described herein. RNA elements can be identified and/or characterized based on the primary sequence of the element (e.g., GC-rich element), by RNA secondary structure formed by the element (e.g. stem-loop), by the location of the element within the RNA molecule (e.g., located within the 5' UTR of an mRNA), by the biological function and/or activity of the element (e.g., "translational enhancer element"), and any combination thereof.

[0230] In some aspects, the disclosure provides an mRNA having one or more structural modifications that inhibits leaky scanning and/or promotes the translational fidelity of mRNA translation, wherein at least one of the structural modifications is a GC-rich RNA element. In some aspects, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising a sequence of linked nucleotides, or derivatives or analogs thereof, preceding a Kozak consensus sequence in a 5' UTR of the mRNA. In one embodiment, the GC-rich RNA element is located about 30, about 25, about 20, about 15, about 10, about 5, about 4, about 3, about 2, or about 1 nucleotide(s) upstream of a Kozak consensus sequence in the 5' UTR of the mRNA. In another embodiment, the GC-rich RNA element is located 15-30, 15-20, 15-25, 10-15, or 5-10 nucleotides upstream of a Kozak consensus sequence. In another embodiment, the GC-rich RNA element is located immediately adjacent to a Kozak consensus sequence in the 5' UTR of the mRNA.

[0231] In any of the foregoing or related aspects, the disclosure provides a GC-rich RNA element which comprises a sequence of 3-30, 5-25, 10-20, 15-20, about 20,

about 15, about 12, about 10, about 7, about 6 or about 3 nucleotides, derivatives or analogs thereof, linked in any order, wherein the sequence composition is 70-80% cytosine, 60-70% cytosine, 50%-60% cytosine, 40-50% cytosine, 30-40% cytosine bases. In any of the foregoing or related aspects, the disclosure provides a GC-rich RNA element which comprises a sequence of 3-30, 5-25, 10-20, 15-20, about 20, about 15, about 12, about 10, about 7, about 6 or about 3 nucleotides, derivatives or analogs thereof, linked in any order, wherein the sequence composition is about 80% cytosine, about 70% cytosine, about 60% cytosine, about 50% cytosine, about 40% cytosine, or about 30% cytosine.

[0232] In any of the foregoing or related aspects, the disclosure provides a GC-rich RNA element which comprises a sequence of 20, 19, 18, 17, 16, 15, 14, 13, 12, 11, 10, 9, 8, 7, 6, 5, 4, or 3 nucleotides, or derivatives or analogs thereof, linked in any order, wherein the sequence composition is 70-80% cytosine, 60-70% cytosine, 50%-60% cytosine, 40-50% cytosine, or 30-40% cytosine. In any of the foregoing or related aspects, the disclosure provides a GC-rich RNA element which comprises a sequence of 20, 19, 18, 17, 16, 15, 14, 13, 12, 11, 10, 9, 8, 7, 6, 5, 4, or 3 nucleotides, or derivatives or analogs thereof, linked in any order, wherein the sequence composition is about 80% cytosine, about 70% cytosine, about 60% cytosine, about 50% cytosine, about 40% cytosine, or about 30% cytosine.

[0233] In some embodiments, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising a sequence of linked nucleotides, or derivatives or analogs thereof, preceding a Kozak consensus sequence in a 5' UTR of the mRNA, wherein the GC-rich RNA element is located about 30, about 25, about 20, about 15, about 10, about 5, about 4, about 3, about 2, or about 1 nucleotide(s) upstream of a Kozak consensus sequence in the 5' UTR of the mRNA, and wherein the GC-rich RNA element comprises a sequence of 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 nucleotides, or derivatives or analogs thereof, linked in any order, wherein the sequence composition is >50% cytosine. In some embodiments, the sequence composition is >55% cytosine, >60% cytosine, >65% cytosine, >70% cytosine, >75% cytosine, >80% cytosine, >85% cytosine, or >90% cytosine.

[0234] In other aspects, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising a sequence of linked nucleotides, or derivatives or analogs thereof,

preceding a Kozak consensus sequence in a 5' UTR of the mRNA, wherein the GC-rich RNA element is located about 30, about 25, about 20, about 15, about 10, about 5, about 4, about 3, about 2, or about 1 nucleotide(s) upstream of a Kozak consensus sequence in the 5' UTR of the mRNA, and wherein the GC-rich RNA element comprises a sequence of about 3-30, 5-25, 10-20, 15-20 or about 20, about 15, about 12, about 10, about 6 or about 3 nucleotides, or derivatives or analogues thereof, wherein the sequence comprises a repeating GC-motif, wherein the repeating GC-motif is [CCG]n, wherein n = 1 to 10 (SEQ ID NO:225), n= 2 to 8 (SEQ ID NO:226), n= 3 to 6 (SEQ ID NO:227), or n= 4 to 5 (SEQ ID NO:228). In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 1, 2, 3, 4 or 5 (SEQ ID NO:229). In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 1, 2, or 3. In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 1. In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 2. In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 3. In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 4 (SEQ ID NO:230). In some embodiments, the sequence comprises a repeating GC-motif [CCG]n, wherein n = 5 (SEQ ID NO:231).

[0235] In another aspect, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising a sequence of linked nucleotides, or derivatives or analogs thereof, preceding a Kozak consensus sequence in a 5' UTR of the mRNA, wherein the GC-rich RNA element comprises any one of the sequences set forth in Table 2. In one embodiment, the GC-rich RNA element is located about 30, about 25, about 20, about 15, about 10, about 5, about 4, about 3, about 2, or about 1 nucleotide(s) upstream of a Kozak consensus sequence in the 5' UTR of the mRNA. In another embodiment, the GC-rich RNA element is located about 15-30, 15-20, 15-25, 10-15, or 5-10 nucleotides upstream of a Kozak consensus sequence. In another embodiment, the GC-rich RNA element is located immediately adjacent to a Kozak consensus sequence in the 5' UTR of the mRNA.

[0236] In other aspects, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising the sequence V1 [CCCCGGCGCC (SEQ ID NO: 43)] as set forth in Table 2, or derivatives or analogs thereof, preceding a Kozak consensus sequence in

the 5' UTR of the mRNA. In some embodiments, the GC-rich element comprises the sequence V1 as set forth in Table 2 located immediately adjacent to and upstream of the Kozak consensus sequence in the 5' UTR of the mRNA. In some embodiments, the GC-rich element comprises the sequence V1 as set forth in Table 2 located 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA. In other embodiments, the GC-rich element comprises the sequence V1 as set forth in Table 2 located 1-3, 3-5, 5-7, 7-9, 9-12, or 12-15 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA.

[0237] In other aspects, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising the sequence V2 [CCCCGGC (SEQ ID NO: 44)] as set forth in Table 2, or derivatives or analogs thereof, preceding a Kozak consensus sequence in the 5' UTR of the mRNA. In some embodiments, the GC-rich element comprises the sequence V2 as set forth in Table 2 located immediately adjacent to and upstream of the Kozak consensus sequence in the 5' UTR of the mRNA. In some embodiments, the GC-rich element comprises the sequence V2 as set forth in Table 2 located 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA. In other embodiments, the GC-rich element comprises the sequence V2 as set forth in Table 2 located 1-3, 3-5, 5-7, 7-9, 9-12, or 12-15 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA.

[0238] In other aspects, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising the sequence EK [GCCGCC (SEQ ID NO:42)] as set forth in Table 2, or derivatives or analogs thereof, preceding a Kozak consensus sequence in the 5' UTR of the mRNA. In some embodiments, the GC-rich element comprises the sequence EK as set forth in Table 2 located immediately adjacent to and upstream of the Kozak consensus sequence in the 5' UTR of the mRNA. In some embodiments, the GC-rich element comprises the sequence EK as set forth in Table 2 located 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA. In other embodiments, the GC-rich element comprises the sequence EK as set forth in Table 2 located 1-3, 3-5, 5-7, 7-9, 9-12, or 12-15 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA.

[0239] In yet other aspects, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element

comprising the sequence V1 [CCCCGGCGCC (SEQ ID NO: 43)] as set forth in Table 2, or derivatives or analogs thereof, preceding a Kozak consensus sequence in the 5' UTR of the mRNA, wherein the 5' UTR comprises the following sequence shown in Table 2:

[0240] GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGA (SEQ ID NO: 85). The skilled artisan will of course recognize that all Us in the RNA sequences described herein will be Ts in a corresponding template DNA sequence, for example, in DNA templates or constructs from which mRNAs of the disclosure are transcribed, e.g., via IVT.

[0241] In some embodiments, the GC-rich element comprises the sequence V1 as set forth in Table 2 located immediately adjacent to and upstream of the Kozak consensus sequence in the 5' UTR sequence shown in Table 2. In some embodiments, the GC-rich element comprises the sequence V1 as set forth in Table 2 located 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA, wherein the 5' UTR comprises the following sequence shown in Table 2:

[0242] GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGA (SEQ ID NO: 85).

[0243] In other embodiments, the GC-rich element comprises the sequence V1 as set forth in Table 2 located 1-3, 3-5, 5-7, 7-9, 9-12, or 12-15 bases upstream of the Kozak consensus sequence in the 5' UTR of the mRNA, wherein the 5' UTR comprises the following sequence shown in Table 2:

[0244] GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGA (SEQ ID NO: 85).

[0245] In some embodiments, the 5' UTR comprises the following sequence set forth in Table 2:

[0246] GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGACC CCGGCGCCGCCACC (SEQ ID NO: 39)

Table 2

	5' UTRs	**5' UTR Sequence**
	Standard	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAG AAAUAUAAGAGCCACC (SEQ ID NO: 3)

	V1-UTR	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAG AAAUAUAAGACCCCGGCGCCGCCACC (SEQ ID NO: 39)
	V2-UTR	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAG AAAUAUAAGACCCCGGCGCCACC (SEQ ID NO: 40)
	GC-Rich RNA Elements	**Sequence**
	K0 (Traditional Kozak consensus)	[GCCA/GCC] (SEQ ID NO: 41)
	EK	[GCCGCC] (SEQ ID NO:42)
	V1	[CCCCGGCGCC] (SEQ ID NO:43)
	V2	[CCCCGGC] (SEQ ID NO:44)
	$(CCG)_n$, where n=1-10	$[CCG]_n$ (SEQ ID NO:225)
	$(GCC)_n$, where n=1-10	$[GCC]_n$ (SEQ ID NO:232)

[0247] In another aspect, the disclosure provides a modified mRNA comprising at least one modification, wherein at least one modification is a GC-rich RNA element comprising a stable RNA secondary structure comprising a sequence of nucleotides, or derivatives or analogs thereof, linked in an order which forms a hairpin or a stem-loop. In one embodiment, the stable RNA secondary structure is upstream of the Kozak consensus sequence. In another embodiment, the stable RNA secondary structure is located about 30, about 25, about 20, about 15, about 10, or about 5 nucleotides upstream of the Kozak consensus sequence. In another embodiment, the stable RNA secondary structure is located about 20, about 15, about 10 or about 5 nucleotides upstream of the Kozak consensus sequence. In another embodiment, the stable RNA secondary structure is located about 5, about 4, about 3, about 2, about 1 nucleotides upstream of the Kozak consensus sequence. In another embodiment, the stable RNA secondary structure is located about 15-30, about 15-20, about 15-25, about 10-15, or about 5-10 nucleotides upstream of the Kozak consensus sequence. In another embodiment, the stable RNA secondary structure is located 12-15 nucleotides upstream of the Kozak consensus sequence. In another embodiment, the stable RNA secondary structure has a deltaG of about -30 kcal/mol, about -20 to -30 kcal/mol, about -20 kcal/mol, about -10 to -20 kcal/mol, about -10 kcal/mol, about -5 to -10 kcal/mol.

[0248] In another embodiment, the modification is operably linked to an open reading frame encoding a polypeptide and wherein the modification and the open reading frame are heterologous.

[0249] In another embodiment, the sequence of the GC-rich RNA element is comprised exclusively of guanine (G) and cytosine (C) nucleobases.

[0250] RNA elements that provide a desired translational regulatory activity as described herein can be identified and characterized using known techniques, such as ribosome profiling. Ribosome profiling is a technique that allows the determination of the positions of PICs and/or ribosomes bound to mRNAs (see e.g., Ingolia et al., (2009) Science 324(5924):218-23, incorporated herein by reference). The technique is based on protecting a region or segment of mRNA, by the PIC and/or ribosome, from nuclease digestion. Protection results in the generation of a 30-bp fragment of RNA termed a 'footprint'. The sequence and frequency of RNA footprints can be analyzed by methods known in the art (e.g., RNA-seq). The footprint is roughly centered on the A-site of the ribosome. If the PIC or ribosome dwells at a particular position or location along an mRNA, footprints generated at these positions would be relatively common. Studies have shown that more footprints are generated at positions where the PIC and/or ribosome exhibits decreased processivity and fewer footprints where the PIC and/or ribosome exhibits increased processivity (Gardin et al., (2014) eLife 3:e03735). In some embodiments, residence time or the time of occupancy of the PIC or ribosome at a discrete position or location along a polynucleotide comprising any one or more of the RNA elements described herein is determined by ribosome profiling.

[0251] A UTR can be homologous or heterologous to the coding region in a polynucleotide. In some embodiments, the UTR is homologous to the ORF encoding the G6PC polypeptide. In some embodiments, the UTR is heterologous to the ORF encoding the G6PC polypeptide. In some embodiments, the polynucleotide comprises two or more 5′ UTRs or functional fragments thereof, each of which has the same or different nucleotide sequences. In some embodiments, the polynucleotide comprises two or more 3′ UTRs or functional fragments thereof, each of which has the same or different nucleotide sequences.

[0252] In some embodiments, the 5′ UTR or functional fragment thereof, 3′ UTR or functional fragment thereof, or any combination thereof is sequence optimized.

[0253] In some embodiments, the 5′UTR or functional fragment thereof, 3′ UTR or functional fragment thereof, or any combination thereof comprises at least one chemically modified nucleobase, e.g., N1-methylpseudouracil or 5-methoxyuracil.

[0254] UTRs can have features that provide a regulatory role, e.g., increased or decreased stability, localization and/or translation efficiency. A polynucleotide comprising a UTR can be administered to a cell, tissue, or organism, and one or more regulatory features can be measured using routine methods. In some embodiments, a functional fragment of a 5' UTR or 3' UTR comprises one or more regulatory features of a full length 5' or 3' UTR, respectively.

[0255] Natural 5′UTRs bear features that play roles in translation initiation. They harbor signatures like Kozak sequences that are commonly known to be involved in the process by which the ribosome initiates translation of many genes. Kozak sequences have the consensus CCR(A/G)CCAUGG (SEQ ID NO:87), where R is a purine (adenine or guanine) three bases upstream of the start codon (AUG), which is followed by another 'G'. 5′ UTRs also have been known to form secondary structures that are involved in elongation factor binding.

[0256] By engineering the features typically found in abundantly expressed genes of specific target organs, one can enhance the stability and protein production of a polynucleotide. For example, introduction of 5′ UTR of liver-expressed mRNA, such as albumin, serum amyloid A, Apolipoprotein A/B/E, transferrin, alpha fetoprotein, erythropoietin, or Factor VIII, can enhance expression of polynucleotides in hepatic cell lines or liver. Likewise, use of 5′UTR from other tissue-specific mRNA to improve expression in that tissue is possible for muscle (e.g., MyoD, Myosin, Myoglobin, Myogenin, Herculin), for endothelial cells (e.g., Tie-1, CD36), for myeloid cells (e.g., C/EBP, AML1, G-CSF, GM-CSF, CD11b, MSR, Fr-1, i-NOS), for leukocytes (e.g., CD45, CD18), for adipose tissue (e.g., CD36, GLUT4, ACRP30, adiponectin) and for lung epithelial cells (e.g., SP-A/B/C/D).

[0257] In some embodiments, UTRs are selected from a family of transcripts whose proteins share a common function, structure, feature or property. For example, an encoded polypeptide can belong to a family of proteins (i.e., that share at least one function, structure, feature, localization, origin, or expression pattern), which are expressed in a particular cell, tissue or at some time during development. The UTRs from any of the genes or mRNA can be swapped for any other UTR of the same or different family of proteins to create a new polynucleotide.

[0258] In some embodiments, the 5' UTR and the 3' UTR can be heterologous. In some embodiments, the 5' UTR can be derived from a different species than the 3' UTR. In some embodiments, the 3' UTR can be derived from a different species than the 5' UTR.

[0259] Co-owned International Patent Application No. PCT/US2014/021522 (Publ. No. WO/2014/164253, incorporated herein by reference in its entirety) provides a listing of exemplary UTRs that can be utilized in the polynucleotide of the present invention as flanking regions to an ORF.

[0260] Exemplary UTRs of the application include, but are not limited to, one or more 5′UTR and/or 3′UTR derived from the nucleic acid sequence of: a globin, such as an α- or β-globin (e.g., a *Xenopus*, mouse, rabbit, or human globin); a strong Kozak translational initiation signal; a CYBA (e.g., human cytochrome b-245 α polypeptide); an albumin (e.g., human albumin7); a HSD17B4 (hydroxysteroid (17-β) dehydrogenase); a virus (e.g., a tobacco etch virus (TEV), a Venezuelan equine encephalitis virus (VEEV), a Dengue virus, a cytomegalovirus (CMV) (e.g., CMV immediate early 1 (IE1)), a hepatitis virus (e.g., hepatitis B virus), a sindbis virus, or a PAV barley yellow dwarf virus); a heat shock protein (e.g., hsp70); a translation initiation factor (e.g., elF4G); a glucose transporter (e.g., hGLUT1 (human glucose transporter 1)); an actin (e.g., human α or β actin); a GAPDH; a tubulin; a histone; a citric acid cycle enzyme; a topoisomerase (e.g., a 5′UTR of a TOP gene lacking the 5′ TOP motif (the oligopyrimidine tract)); a ribosomal protein Large 32 (L32); a ribosomal protein (e.g., human or mouse ribosomal protein, such as, for example, rps9); an ATP synthase (e.g., ATP5A1 or the β subunit of mitochondrial H^+-ATP synthase); a growth hormone e (e.g., bovine (bGH) or human (hGH)); an elongation factor (e.g., elongation factor 1 α1 (EEF1A1)); a manganese superoxide dismutase (MnSOD); a myocyte enhancer factor 2A (MEF2A); a β-F1-ATPase, a creatine kinase, a myoglobin, a granulocyte-colony stimulating factor (G-CSF); a collagen (e.g., collagen type I, alpha 2 (Col1A2), collagen type I, alpha 1 (Col1A1), collagen type VI, alpha 2 (Col6A2), collagen type VI, alpha 1 (Col6A1)); a ribophorin (e.g., ribophorin I (RPNI)); a low density lipoprotein receptor-related protein (e.g., LRP1); a cardiotrophin-like cytokine factor (e.g., Nnt1); calreticulin (Calr); a procollagen-lysine, 2-oxoglutarate 5-dioxygenase 1 (Plod1); and a nucleobindin (e.g., Nucb1).

[0261] In some embodiments, the 5' UTR is selected from the group consisting of a β-globin 5' UTR; a 5′UTR containing a strong Kozak translational initiation signal; a

cytochrome b-245 α polypeptide (CYBA) 5' UTR; a hydroxysteroid (17-β) dehydrogenase (HSD17B4) 5' UTR; a Tobacco etch virus (TEV) 5' UTR; a Venezuelan equine encephalitis virus (TEEV) 5' UTR; a 5' proximal open reading frame of rubella virus (RV) RNA encoding nonstructural proteins; a Dengue virus (DEN) 5' UTR; a heat shock protein 70 (Hsp70) 5' UTR; a eIF4G 5' UTR; a GLUT1 5' UTR; functional fragments thereof and any combination thereof.

[0262] In some embodiments, the 3' UTR is selected from the group consisting of a β-globin 3' UTR; a CYBA 3' UTR; an albumin 3' UTR; a growth hormone (GH) 3' UTR; a VEEV 3' UTR; a hepatitis B virus (HBV) 3' UTR; α-globin 3′UTR; a DEN 3' UTR; a PAV barley yellow dwarf virus (BYDV-PAV) 3' UTR; an elongation factor 1 α1 (EEF1A1) 3' UTR; a manganese superoxide dismutase (MnSOD) 3' UTR; a β subunit of mitochondrial H(+)-ATP synthase (β-mRNA) 3' UTR; a GLUT1 3' UTR; a MEF2A 3' UTR; a β-F1-ATPase 3' UTR; functional fragments thereof and combinations thereof.

[0263] Wild-type UTRs derived from any gene or mRNA can be incorporated into the polynucleotides of the invention. In some embodiments, a UTR can be altered relative to a wild type or native UTR to produce a variant UTR, e.g., by changing the orientation or location of the UTR relative to the ORF; or by inclusion of additional nucleotides, deletion of nucleotides, swapping or transposition of nucleotides. In some embodiments, variants of 5′ or 3′ UTRs can be utilized, for example, mutants of wild type UTRs, or variants wherein one or more nucleotides are added to or removed from a terminus of the UTR.

[0264] Additionally, one or more synthetic UTRs can be used in combination with one or more non-synthetic UTRs. *See*, e.g., Mandal and Rossi, Nat. Protoc. 2013 8(3):568-82, the contents of which are incorporated herein by reference in their entirety.

[0265] UTRs or portions thereof can be placed in the same orientation as in the transcript from which they were selected or can be altered in orientation or location. Hence, a 5′ and/or 3′ UTR can be inverted, shortened, lengthened, or combined with one or more other 5′ UTRs or 3′ UTRs.

[0266] In some embodiments, the polynucleotide comprises multiple UTRs, e.g., a double, a triple or a quadruple 5' UTR or 3' UTR. For example, a double UTR comprises two copies of the same UTR either in series or substantially in series. For

example, a double beta-globin 3′UTR can be used (see US2010/0129877, the contents of which are incorporated herein by reference in its entirety).

[0267] In certain embodiments, the polynucleotides of the invention comprise a 5' UTR and/or a 3' UTR selected from any of the UTRs disclosed herein. In some embodiments, the 5' UTR comprises:

5' UTR-001 (Upstream UTR) (GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.:3);

5' UTR-002 (Upstream UTR) (GGGAGAUCAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.:89);

5' UTR-003 (Upstream UTR) (See WO2016/100812);

5' UTR-004 (Upstream UTR) (GGGAGACAAGCUUGGCAUUCCGGUACUGUUGGUAAAGCCACC) (SEQ ID NO.:90);

5' UTR-005 (Upstream UTR) (GGGAGAUCAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.:91);

5' UTR-006 (Upstream UTR) (See WO2016/100812);

5' UTR-007 (Upstream UTR) (GGGAGACAAGCUUGGCAUUCCGGUACUGUUGGUAAAGCCACC) (SEQ ID NO.:92);

5' UTR-008 (Upstream UTR) (GGGAAUUAACAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.:93);

5' UTR-009 (Upstream UTR) (GGGAAAUUAGACAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.:94);

5' UTR-010, Upstream (GGGAAAUAAGAGAGUAAAGAACAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.:95);

5' UTR-011 (Upstream UTR) (GGGAAAAAAGAGAGAAAAGAAGACUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID NO.: 96);

5' UTR-012 (Upstream UTR) (GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAUAUAUAAGAGCCACC) (SEQ ID NO.:97);

5' UTR-013 (Upstream UTR)
(GGGAAAUAAGAGACAAAACAAGAGUAAGAAGAAAUAUAAGAGCCACC)
(SEQ ID NO.:98);

5' UTR-014 (Upstream UTR)
(GGGAAAUUAGAGAGUAAAGAACAGUAAGUAGAAUUAAAAGAGCCACC)
(SEQ ID NO.:99);

5' UTR-015 (Upstream UTR)
(GGGAAAUAAGAGAGAAUAGAAGAGUAAGAAGAAAUAUAAGAGCCACC)
(SEQ ID NO.:100);

5' UTR-016 (Upstream UTR)
(GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAAUUAAGAGCCACC)
(SEQ ID NO.:101);

5' UTR-017 (Upstream UTR); or
(GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUUUAAGAGCCACC)
(SEQ ID NO.:102);

5' UTR-018 (Upstream UTR) 5' UTR

(UCAAGCUUUUGGACCCUCGUACAGAAGCUAAUACGACUCACUAUAGGGAA
AUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC) (SEQ ID
NO.:88).

[0268] In some embodiments, the 3' UTR comprises:

142-3p 3' UTR (UTR including miR142-3p binding site)
(UGAUAAUAGUCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCC
AUGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACC
CGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC) (SEQ ID
NO.:104);

142-3p 3' UTR (UTR including miR142-3p binding site)
(UGAUAAUAGGCUGGAGCCUCGGUGGCUCCAUAAAGUAGGAAACACUACAC
AUGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACC
CGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC) (SEQ ID
NO.:105); or

142-3p 3' UTR (UTR including miR142-3p binding site)
(UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUCCAUAAA
GUAGGAAACACUACAUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCAC
CCGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC) (SEQ ID
NO.:106);

142-3p 3' UTR (UTR including miR142-3p binding site)
(UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUC
CCCCCAGUCCAUAAAGUAGGAAACACUACACCCCUCCUCCCCUUCCUGCAC
CCGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC) (SEQ ID NO.:
107);

142-3p 3' UTR (UTR including miR142-3p binding site) (UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUC CCCCCAGCCCCUCCUCCCCCUUCUCCAUAAAGUAGGAAACACUACACUGCAC CCGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC) (SEQ ID NO.:108);

142-3p 3' UTR (UTR including miR142-3p binding site) (UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUC CCCCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCCUCCAUAAAGUAGGA AACACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC) (SEQ ID NO.:109).

142-3p 3' UTR (UTR including miR142-3p binding site) (UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUC CCCCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUA AAGUUCCAUAAAGUAGGAAACACUACACUGAGUGGGCGGC) (SEQ ID NO.:110);

3' UTR-018 (See SEQ ID NO. 150);

UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCC CCCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUA AAGUCUGAGUGGGCGGC (SEQ ID NO: 150)

3' UTR (miR142 and miR126 binding sites variant 1)

(UGAUAAUAGUCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCC AUGCUUCUUGCCCCUUGGGCCUCCCCCCCAGCCCCUCCUCCCCCUUCCUGCACC CGUACCCCCCCGCAUUAUUACUCACGGUACGAGUGGUCUUUGAAUAAAGUCU GAGUGGGCGGC) (SEQ ID NO:111)

3' UTR (miR142 and miR126 binding sites variant 2)

(UGAUAAUAGUCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCC UAGCUUCUUGCCCCUUGGGCCUCCCCCCCAGCCCCUCCUCCCCCUUCCUGCACC CGUACCCCCCCGCAUUAUUACUCACGGUACGAGUGGUCUUUGAAUAAAGUCU GAGUGGGCGGC) (SEQ ID NO.:112); or

3'UTR (miR142-3p binding site variant 3)

UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCC CCCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGUAGGAA ACACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (SEQ ID NO:4).

3'UTR:

UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCC CCCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUA AAGUCUGAGUGGGCGGC (SEQ ID NO: 175)

[0269] In certain embodiments, the 5' UTR and/or 3' UTR sequence of the invention comprises a nucleotide sequence at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or about 100% identical to a sequence selected from the group consisting of 5' UTR sequences comprising any of SEQ ID NOs: 3, 88-102, or 165-167 and/or 3' UTR sequences comprises any of SEQ ID NOs:4, 104-112, 150, or 175, and any combination thereof.

[0270] In certain embodiments, the 5′ UTR and/or 3′ UTR sequence of the invention comprises a nucleotide sequence at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or about 100% identical to a sequence selected from the group consisting of 5′ UTR sequences comprising any of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211 and/or 3′ UTR sequences comprises any of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213, and any combination thereof.

[0271] In some embodiments, the 5′ UTR comprises an amino acid sequence set forth in Table 4B (SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211). In some embodiments, the 3′ UTR comprises an amino acid sequence set forth in Table 4B (SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213). In some embodiments, the 5′ UTR comprises an amino acid sequence set forth in Table 4B (SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211) and the 3′ UTR comprises an amino acid sequence set forth in Table 4B (SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213).

[0272] The polynucleotides of the invention can comprise combinations of features. For example, the ORF can be flanked by a 5′UTR that comprises a strong Kozak translational initiation signal and/or a 3′UTR comprising an oligo(dT) sequence for templated addition of a poly-A tail. A 5′UTR can comprise a first polynucleotide fragment and a second polynucleotide fragment from the same and/or different UTRs (see, e.g., US2010/0293625, herein incorporated by reference in its entirety).

[0273] Other non-UTR sequences can be used as regions or subregions within the polynucleotides of the invention. For example, introns or portions of intron sequences

can be incorporated into the polynucleotides of the invention. Incorporation of intronic sequences can increase protein production as well as polynucleotide expression levels. In some embodiments, the polynucleotide of the invention comprises an internal ribosome entry site (IRES) instead of or in addition to a UTR (see, e.g., Yakubov et al., Biochem. Biophys. Res. Commun. 2010 394(1):189-193, the contents of which are incorporated herein by reference in their entirety). In some embodiments, the polynucleotide comprises an IRES instead of a 5' UTR sequence. In some embodiments, the polynucleotide comprises an ORF and a viral capsid sequence. In some embodiments, the polynucleotide comprises a synthetic 5' UTR in combination with a non-synthetic 3' UTR.

[0274] In some embodiments, the UTR can also include at least one translation enhancer polynucleotide, translation enhancer element, or translational enhancer elements (collectively, "TEE," which refers to nucleic acid sequences that increase the amount of polypeptide or protein produced from a polynucleotide. As a non-limiting example, the TEE can be located between the transcription promoter and the start codon. In some embodiments, the 5' UTR comprises a TEE.

[0275] In one aspect, a TEE is a conserved element in a UTR that can promote translational activity of a nucleic acid such as, but not limited to, cap-dependent or cap-independent translation.

11. MicroRNA (miRNA) Binding Sites

[0276] Polynucleotides of the invention can include regulatory elements, for example, microRNA (miRNA) binding sites, transcription factor binding sites, structured mRNA sequences and/or motifs, artificial binding sites engineered to act as pseudo-receptors for endogenous nucleic acid binding molecules, and combinations thereof. In some embodiments, polynucleotides including such regulatory elements are referred to as including “sensor sequences”.

[0277] In some embodiments, a polynucleotide (e.g., a ribonucleic acid (RNA), e.g., a messenger RNA (mRNA)) of the invention comprises an open reading frame (ORF) encoding a polypeptide of interest and further comprises one or more miRNA binding site(s). Inclusion or incorporation of miRNA binding site(s) provides for regulation of polynucleotides of the invention, and in turn, of the polypeptides encoded therefrom, based on tissue-specific and/or cell-type specific expression of naturally-occurring miRNAs.

[0278] The present invention also provides pharmaceutical compositions and formulations that comprise any of the polynucleotides described above. In some embodiments, the composition or formulation further comprises a delivery agent.

[0279] In some embodiments, the composition or formulation can contain a polynucleotide comprising a sequence optimized nucleic acid sequence disclosed herein which encodes a polypeptide. In some embodiments, the composition or formulation can contain a polynucleotide (e.g., a RNA, e.g., an mRNA) comprising a polynucleotide (e.g., an ORF) having significant sequence identity to a sequence optimized nucleic acid sequence disclosed herein which encodes a polypeptide. In some embodiments, the polynucleotide further comprises a miRNA binding site, e.g., a miRNA binding site that binds

[0280] A miRNA, e.g., a natural-occurring miRNA, is a 19-25 nucleotide long noncoding RNA that binds to a polynucleotide and down-regulates gene expression either by reducing stability or by inhibiting translation of the polynucleotide. A miRNA sequence comprises a “seed” region, i.e., a sequence in the region of positions 2-8 of the mature miRNA. A miRNA seed can comprise positions 2-8 or 2-7 of the mature miRNA.

[0281] microRNAs derive enzymatically from regions of RNA transcripts that fold back on themselves to form short hairpin structures often termed a pre-miRNA (precursor-miRNA). A pre-miRNA typically has a two-nucleotide overhang at its 3' end, and has 3' hydroxyl and 5' phosphate groups. This precursor-mRNA is processed in the nucleus and subsequently transported to the cytoplasm where it is further processed by DICER (a RNase III enzyme), to form a mature microRNA of approximately 22 nucleotides. The mature microRNA is then incorporated into a ribonuclear particle to form the RNA-induced silencing complex, RISC, which mediates gene silencing. Art-recognized nomenclature for mature miRNAs typically designates the arm of the pre-miRNA from which the mature miRNA derives; "5p" means the microRNA is from the 5-prime arm of the pre-miRNA hairpin and "3p" means the microRNA is from the 3-prime end of the pre-miRNA hairpin. A miR referred to by number herein can refer to either of the two mature microRNAs originating from opposite arms of the same pre-miRNA (e.g., either the 3p or 5p microRNA). All miRs referred to herein are intended to include both the 3p and 5p arms/sequences, unless particularly specified by the 3p or 5p designation.

[0282] As used herein, the term "microRNA (miRNA or miR) binding site" refers to a sequence within a polynucleotide, e.g., within a DNA or within an RNA transcript, including in the 5′UTR and/or 3′UTR, that has sufficient complementarity to all or a region of a miRNA to interact with, associate with or bind to the miRNA. In some embodiments, a polynucleotide of the invention comprising an ORF encoding a polypeptide of interest and further comprises one or more miRNA binding site(s). In exemplary embodiments, a 5' UTR and/or 3' UTR of the polynucleotide (e.g., a ribonucleic acid (RNA), e.g., a messenger RNA (mRNA)) comprises the one or more miRNA binding site(s).

[0283] A miRNA binding site having sufficient complementarity to a miRNA refers to a degree of complementarity sufficient to facilitate miRNA-mediated regulation of a polynucleotide, e.g., miRNA-mediated translational repression or degradation of the polynucleotide. In exemplary aspects of the invention, a miRNA binding site having sufficient complementarity to the miRNA refers to a degree of complementarity sufficient to facilitate miRNA-mediated degradation of the polynucleotide, e.g., miRNA-guided RNA-induced silencing complex (RISC)-mediated cleavage of mRNA. The miRNA binding site can have complementarity to, for example, a 19-25 nucleotide long miRNA sequence, to a 19-23 nucleotide long miRNA sequence, or to a 22-nucleotide long miRNA sequence. A miRNA binding site can be complementary to only a portion of a miRNA, e.g., to a portion less than 1, 2, 3, or 4 nucleotides of the full length of a naturally-occurring miRNA sequence, or to a portion less than 1, 2, 3, or 4 nucleotides shorter than a naturally-occurring miRNA sequence. Full or complete complementarity (e.g., full complementarity or complete complementarity over all or a significant portion of the length of a naturally-occurring miRNA) is preferred when the desired regulation is mRNA degradation.

[0284] In some embodiments, a miRNA binding site includes a sequence that has complementarity (e.g., partial or complete complementarity) with a miRNA seed sequence. In some embodiments, the miRNA binding site includes a sequence that has complete complementarity with a miRNA seed sequence. In some embodiments, a miRNA binding site includes a sequence that has complementarity (e.g., partial or complete complementarity) with a miRNA sequence. In some embodiments, the miRNA binding site includes a sequence that has complete complementarity with a miRNA sequence. In some embodiments, a miRNA binding site has complete

complementarity with a miRNA sequence but for 1, 2, or 3 nucleotide substitutions, terminal additions, and/or truncations.

[0285] In some embodiments, the miRNA binding site is the same length as the corresponding miRNA. In other embodiments, the miRNA binding site is one, two, three, four, five, six, seven, eight, nine, ten, eleven or twelve nucleotide(s) shorter than the corresponding miRNA at the 5' terminus, the 3' terminus, or both. In still other embodiments, the microRNA binding site is two nucleotides shorter than the corresponding microRNA at the 5' terminus, the 3' terminus, or both. The miRNA binding sites that are shorter than the corresponding miRNAs are still capable of degrading the mRNA incorporating one or more of the miRNA binding sites or preventing the mRNA from translation.

[0286] In some embodiments, the miRNA binding site binds the corresponding mature miRNA that is part of an active RISC containing Dicer. In another embodiment, binding of the miRNA binding site to the corresponding miRNA in RISC degrades the mRNA containing the miRNA binding site or prevents the mRNA from being translated. In some embodiments, the miRNA binding site has sufficient complementarity to miRNA so that a RISC complex comprising the miRNA cleaves the polynucleotide comprising the miRNA binding site. In other embodiments, the miRNA binding site has imperfect complementarity so that a RISC complex comprising the miRNA induces instability in the polynucleotide comprising the miRNA binding site. In another embodiment, the miRNA binding site has imperfect complementarity so that a RISC complex comprising the miRNA represses transcription of the polynucleotide comprising the miRNA binding site.

[0287] In some embodiments, the miRNA binding site has one, two, three, four, five, six, seven, eight, nine, ten, eleven or twelve mismatch(es) from the corresponding miRNA.

[0288] In some embodiments, the miRNA binding site has at least about ten, at least about eleven, at least about twelve, at least about thirteen, at least about fourteen, at least about fifteen, at least about sixteen, at least about seventeen, at least about eighteen, at least about nineteen, at least about twenty, or at least about twenty-one contiguous nucleotides complementary to at least about ten, at least about eleven, at least about twelve, at least about thirteen, at least about fourteen, at least about fifteen, at least about sixteen, at least about seventeen, at least about eighteen, at least about

nineteen, at least about twenty, or at least about twenty-one, respectively, contiguous nucleotides of the corresponding miRNA.

[0289] By engineering one or more miRNA binding sites into a polynucleotide of the invention, the polynucleotide can be targeted for degradation or reduced translation, provided the miRNA in question is available. This can reduce off-target effects upon delivery of the polynucleotide. For example, if a polynucleotide of the invention is not intended to be delivered to a tissue or cell but ends up is said tissue or cell, then a miRNA abundant in the tissue or cell can inhibit the expression of the gene of interest if one or multiple binding sites of the miRNA are engineered into the 5′ UTR and/or 3′ UTR of the polynucleotide. Thus, in some embodiments, incorporation of one or more miRNA binding sites into an mRNA of the disclosure may reduce the hazard of off-target effects upon nucleic acid molecule delivery and/or enable tissue-specific regulation of expression of a polypeptide encoded by the mRNA. In yet other embodiments, incorporation of one or more miRNA binding sites into an mRNA of the disclosure can modulate immune responses upon nucleic acid delivery *in vivo*. In further embodiments, incorporation of one or more miRNA binding sites into an mRNA of the disclosure can modulate accelerated blood clearance (ABC) of lipid-comprising compounds and compositions described herein.

[0290] Conversely, miRNA binding sites can be removed from polynucleotide sequences in which they naturally occur to increase protein expression in specific tissues. For example, a binding site for a specific miRNA can be removed from a polynucleotide to improve protein expression in tissues or cells containing the miRNA.

[0291] Regulation of expression in multiple tissues can be accomplished through introduction or removal of one or more miRNA binding sites, e.g., one or more distinct miRNA binding sites. The decision whether to remove or insert a miRNA binding site can be made based on miRNA expression patterns and/or their profiling in tissues and/or cells in development and/or disease. Identification of miRNAs, miRNA binding sites, and their expression patterns and role in biology have been reported (e.g., Bonauer et al., Curr Drug Targets 2010 11:943-949; Anand and Cheresh Curr Opin Hematol 2011 18:171-176; Contreras and Rao Leukemia 2012 26:404-413 (2011 Dec 20. doi: 10.1038/leu.2011.356); Bartel Cell 2009 136:215-233; Landgraf et al, Cell, 2007 129:1401-1414; Gentner and Naldini, Tissue Antigens.

2012 80:393-403 and all references therein; each of which is incorporated herein by reference in its entirety).

[0292] Examples of tissues where miRNA are known to regulate mRNA, and thereby protein expression, include, but are not limited to, liver (miR-122), muscle (miR-133, miR-206, miR-208), endothelial cells (miR-17-92, miR-126), myeloid cells (miR-142-3p, miR-142-5p, miR-16, miR-21, miR-223, miR-24, miR-27), adipose tissue (let-7, miR-30c), heart (miR-1d, miR-149), kidney (miR-192, miR-194, miR-204), and lung epithelial cells (let-7, miR-133, miR-126).

[0293] Specifically, miRNAs are known to be differentially expressed in immune cells (also called hematopoietic cells), such as antigen presenting cells (APCs) (e.g., dendritic cells and macrophages), macrophages, monocytes, B lymphocytes, T lymphocytes, granulocytes, natural killer cells, etc. Immune cell specific miRNAs are involved in immunogenicity, autoimmunity, the immune-response to infection, inflammation, as well as unwanted immune response after gene therapy and tissue/organ transplantation. Immune cells specific miRNAs also regulate many aspects of development, proliferation, differentiation and apoptosis of hematopoietic cells (immune cells). For example, miR-142 and miR-146 are exclusively expressed in immune cells, particularly abundant in myeloid dendritic cells. It has been demonstrated that the immune response to a polynucleotide can be shut-off by adding miR-142 binding sites to the 3′-UTR of the polynucleotide, enabling more stable gene transfer in tissues and cells. miR-142 efficiently degrades exogenous polynucleotides in antigen presenting cells and suppresses cytotoxic elimination of transduced cells (e.g., Annoni A et al., blood, 2009, 114, 5152-5161; Brown BD, et al., Nat med. 2006, 12(5), 585-591; Brown BD, et al., blood, 2007, 110(13): 4144-4152, each of which is incorporated herein by reference in its entirety).

[0294] An antigen-mediated immune response can refer to an immune response triggered by foreign antigens, which, when entering an organism, are processed by the antigen presenting cells and displayed on the surface of the antigen presenting cells. T cells can recognize the presented antigen and induce a cytotoxic elimination of cells that express the antigen.

[0295] Introducing a miR-142 binding site into the 5' UTR and/or 3′UTR of a polynucleotide of the invention can selectively repress gene expression in antigen presenting cells through miR-142 mediated degradation, limiting antigen presentation in antigen presenting cells (e.g., dendritic cells) and thereby preventing antigen-

mediated immune response after the delivery of the polynucleotide. The polynucleotide is then stably expressed in target tissues or cells without triggering cytotoxic elimination.

[0296] In one embodiment, binding sites for miRNAs that are known to be expressed in immune cells, in particular, antigen presenting cells, can be engineered into a polynucleotide of the invention to suppress the expression of the polynucleotide in antigen presenting cells through miRNA mediated RNA degradation, subduing the antigen-mediated immune response. Expression of the polynucleotide is maintained in non-immune cells where the immune cell specific miRNAs are not expressed. For example, in some embodiments, to prevent an immunogenic reaction against a liver specific protein, any miR-122 binding site can be removed and a miR-142 (and/or mirR-146) binding site can be engineered into the 5' UTR and/or 3' UTR of a polynucleotide of the invention.

[0297] To further drive the selective degradation and suppression in APCs and macrophage, a polynucleotide of the invention can include a further negative regulatory element in the 5' UTR and/or 3' UTR, either alone or in combination with miR-142 and/or miR-146 binding sites. As a non-limiting example, the further negative regulatory element is a Constitutive Decay Element (CDE).

[0298] Immune cell specific miRNAs include, but are not limited to, hsa-let-7a-2-3p, hsa-let-7a-3p, hsa-7a-5p, hsa-let-7c, hsa-let-7e-3p, hsa-let-7e-5p, hsa-let-7g-3p, hsa-let-7g-5p, hsa-let-7i-3p, hsa-let-7i-5p, miR-10a-3p, miR-10a-5p, miR-1184, hsa-let-7f-1--3p, hsa-let-7f-2--5p, hsa-let-7f-5p, miR-125b-1-3p, miR-125b-2-3p, miR-125b-5p, miR-1279, miR-130a-3p, miR-130a-5p, miR-132-3p, miR-132-5p, miR-142-3p, miR-142-5p, miR-143-3p, miR-143-5p, miR-146a-3p, miR-146a-5p, miR-146b-3p, miR-146b-5p, miR-147a, miR-147b, miR-148a-5p, miR-148a-3p, miR-150-3p, miR-150-5p, miR-151b, miR-155-3p, miR-155-5p, miR-15a-3p, miR-15a-5p, miR-15b-5p, miR-15b-3p, miR-16-1-3p, miR-16-2-3p, miR-16-5p, miR-17-5p, miR-181a-3p, miR-181a-5p, miR-181a-2-3p, miR-182-3p, miR-182-5p, miR-197-3p, miR-197-5p, miR-21-5p, miR-21-3p, miR-214-3p, miR-214-5p, miR-223-3p, miR-223-5p, miR-221-3p, miR-221-5p, miR-23b-3p, miR-23b-5p, miR-24-1-5p,miR-24-2-5p, miR-24-3p, miR-26a-1-3p, miR-26a-2-3p, miR-26a-5p, miR-26b-3p, miR-26b-5p, miR-27a-3p, miR-27a-5p, miR-27b-3p,miR-27b-5p, miR-28-3p, miR-28-5p, miR-2909, miR-29a-3p, miR-29a-5p, miR-29b-1-5p, miR-29b-2-5p, miR-29c-3p, miR-29c-5p,, miR-30e-3p, miR-30e-5p, miR-331-5p, miR-339-3p, miR-339-5p, miR-345-3p, miR-345-5p, miR-

346, miR-34a-3p, miR-34a-5p, , miR-363-3p, miR-363-5p, miR-372, miR-377-3p, miR-377-5p, miR-493-3p, miR-493-5p, miR-542, miR-548b-5p, miR548c-5p, miR-548i, miR-548j, miR-548n, miR-574-3p, miR-598, miR-718, miR-935, miR-99a-3p, miR-99a-5p, miR-99b-3p, and miR-99b-5p. Furthermore, novel miRNAs can be identified in immune cell through micro-array hybridization and microtome analysis (e.g., Jima DD et al, Blood, 2010, 116:e118-e127; Vaz C et al., BMC Genomics, 2010, 11,288, the content of each of which is incorporated herein by reference in its entirety.)

[0299] miRNAs that are known to be expressed in the liver include, but are not limited to, miR-107, miR-122-3p, miR-122-5p, miR-1228-3p, miR-1228-5p, miR-1249, miR-129-5p, miR-1303, miR-151a-3p, miR-151a-5p, miR-152, miR-194-3p, miR-194-5p, miR-199a-3p, miR-199a-5p, miR-199b-3p, miR-199b-5p, miR-296-5p, miR-557, miR-581, miR-939-3p, and miR-939-5p. miRNA binding sites from any liver specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the liver. Liver specific miRNA binding sites can be engineered alone or further in combination with immune cell (e.g., APC) miRNA binding sites in a polynucleotide of the invention.

[0300] miRNAs that are known to be expressed in the lung include, but are not limited to, let-7a-2-3p, let-7a-3p, let-7a-5p, miR-126-3p, miR-126-5p, miR-127-3p, miR-127-5p, miR-130a-3p, miR-130a-5p, miR-130b-3p, miR-130b-5p, miR-133a, miR-133b, miR-134, miR-18a-3p, miR-18a-5p, miR-18b-3p, miR-18b-5p, miR-24-1-5p, miR-24-2-5p, miR-24-3p, miR-296-3p, miR-296-5p, miR-32-3p, miR-337-3p, miR-337-5p, miR-381-3p, and miR-381-5p. miRNA binding sites from any lung specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the lung. Lung specific miRNA binding sites can be engineered alone or further in combination with immune cell (e.g., APC) miRNA binding sites in a polynucleotide of the invention.

[0301] miRNAs that are known to be expressed in the heart include, but are not limited to, miR-1, miR-133a, miR-133b, miR-149-3p, miR-149-5p, miR-186-3p, miR-186-5p, miR-208a, miR-208b, miR-210, miR-296-3p, miR-320, miR-451a, miR-451b, miR-499a-3p, miR-499a-5p, miR-499b-3p, miR-499b-5p, miR-744-3p, miR-744-5p, miR-92b-3p, and miR-92b-5p. miRNA binding sites from any heart specific microRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the heart. Heart specific miRNA binding

sites can be engineered alone or further in combination with immune cell (e.g., APC) miRNA binding sites in a polynucleotide of the invention.

[0302] miRNAs that are known to be expressed in the nervous system include, but are not limited to, miR-124-5p, miR-125a-3p, miR-125a-5p, miR-125b-1-3p, miR-125b-2-3p, miR-125b-5p,miR-1271-3p, miR-1271-5p, miR-128, miR-132-5p, miR-135a-3p, miR-135a-5p, miR-135b-3p, miR-135b-5p, miR-137, miR-139-5p, miR-139-3p, miR-149-3p, miR-149-5p, miR-153, miR-181c-3p, miR-181c-5p, miR-183-3p, miR-183-5p, miR-190a, miR-190b, miR-212-3p, miR-212-5p, miR-219-1-3p, miR-219-2-3p, miR-23a-3p, miR-23a-5p,miR-30a-5p, miR-30b-3p, miR-30b-5p, miR-30c-1-3p, miR-30c-2-3p, miR-30c-5p, miR-30d-3p, miR-30d-5p, miR-329, miR-342-3p, miR-3665, miR-3666, miR-380-3p, miR-380-5p, miR-383, miR-410, miR-425-3p, miR-425-5p, miR-454-3p, miR-454-5p, miR-483, miR-510, miR-516a-3p, miR-548b-5p, miR-548c-5p, miR-571, miR-7-1-3p, miR-7-2-3p, miR-7-5p, miR-802, miR-922, miR-9-3p, and miR-9-5p. miRNAs enriched in the nervous system further include those specifically expressed in neurons, including, but not limited to, miR-132-3p, miR-132-3p, miR-148b-3p, miR-148b-5p, miR-151a-3p, miR-151a-5p, miR-212-3p, miR-212-5p, miR-320b, miR-320e, miR-323a-3p, miR-323a-5p, miR-324-5p, miR-325, miR-326, miR-328, miR-922 and those specifically expressed in glial cells, including, but not limited to, miR-1250, miR-219-1-3p, miR-219-2-3p, miR-219-5p, miR-23a-3p, miR-23a-5p, miR-3065-3p, miR-3065-5p, miR-30e-3p, miR-30e-5p, miR-32-5p, miR-338-5p, and miR-657. miRNA binding sites from any CNS specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the nervous system. Nervous system specific miRNA binding sites can be engineered alone or further in combination with immune cell (e.g., APC) miRNA binding sites in a polynucleotide of the invention.

[0303] miRNAs that are known to be expressed in the pancreas include, but are not limited to, miR-105-3p, miR-105-5p, miR-184, miR-195-3p, miR-195-5p, miR-196a-3p, miR-196a-5p, miR-214-3p, miR-214-5p, miR-216a-3p, miR-216a-5p, miR-30a-3p, miR-33a-3p, miR-33a-5p, miR-375, miR-7-1-3p, miR-7-2-3p, miR-493-3p, miR-493-5p, and miR-944. miRNA binding sites from any pancreas specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the pancreas. Pancreas specific miRNA binding sites can be engineered alone or further in combination with immune cell (e.g. APC) miRNA binding sites in a polynucleotide of the invention.

[0304] miRNAs that are known to be expressed in the kidney include, but are not limited to, miR-122-3p, miR-145-5p, miR-17-5p, miR-192-3p, miR-192-5p, miR-194-3p, miR-194-5p, miR-20a-3p, miR-20a-5p, miR-204-3p, miR-204-5p, miR-210, miR-216a-3p, miR-216a-5p, miR-296-3p, miR-30a-3p, miR-30a-5p, miR-30b-3p, miR-30b-5p, miR-30c-1-3p, miR-30c-2-3p, miR30c-5p, miR-324-3p, miR-335-3p, miR-335-5p, miR-363-3p, miR-363-5p, and miR-562. miRNA binding sites from any kidney specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the kidney. Kidney specific miRNA binding sites can be engineered alone or further in combination with immune cell (e.g., APC) miRNA binding sites in a polynucleotide of the invention.

[0305] miRNAs that are known to be expressed in the muscle include, but are not limited to, let-7g-3p, let-7g-5p, miR-1, miR-1286, miR-133a, miR-133b, miR-140-3p, miR-143-3p, miR-143-5p, miR-145-3p, miR-145-5p, miR-188-3p, miR-188-5p, miR-206, miR-208a, miR-208b, miR-25-3p, and miR-25-5p. MiRNA binding sites from any muscle specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the muscle. Muscle specific miRNA binding sites can be engineered alone or further in combination with immune cell (e.g., APC) miRNA binding sites in a polynucleotide of the invention.

[0306] miRNAs are also differentially expressed in different types of cells, such as, but not limited to, endothelial cells, epithelial cells, and adipocytes.

[0307] miRNAs that are known to be expressed in endothelial cells include, but are not limited to, let-7b-3p, let-7b-5p, miR-100-3p, miR-100-5p, miR-101-3p, miR-101-5p, miR-126-3p, miR-126-5p, miR-1236-3p, miR-1236-5p, miR-130a-3p, miR-130a-5p, miR-17-5p, miR-17-3p, miR-18a-3p, miR-18a-5p, miR-19a-3p, miR-19a-5p, miR-19b-1-5p, miR-19b-2-5p, miR-19b-3p, miR-20a-3p, miR-20a-5p, miR-217, miR-210, miR-21-3p, miR-21-5p, miR-221-3p, miR-221-5p, miR-222-3p, miR-222-5p, miR-23a-3p, miR-23a-5p, miR-296-5p, miR-361-3p, miR-361-5p, miR-421, miR-424-3p, miR-424-5p, miR-513a-5p, miR-92a-1-5p, miR-92a-2-5p, miR-92a-3p, miR-92b-3p, and miR-92b-5p. Many novel miRNAs are discovered in endothelial cells from deep-sequencing analysis (e.g., Voellenkle C et al., RNA, 2012, 18, 472-484, herein incorporated by reference in its entirety). miRNA binding sites from any endothelial cell specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the endothelial cells.

[0308] miRNAs that are known to be expressed in epithelial cells include, but are not limited to, let-7b-3p, let-7b-5p, miR-1246, miR-200a-3p, miR-200a-5p, miR-200b-3p, miR-200b-5p, miR-200c-3p, miR-200c-5p, miR-338-3p, miR-429, miR-451a, miR-451b, miR-494, miR-802 and miR-34a, miR-34b-5p, miR-34c-5p, miR-449a, miR-449b-3p, miR-449b-5p specific in respiratory ciliated epithelial cells, let-7 family, miR-133a, miR-133b, miR-126 specific in lung epithelial cells, miR-382-3p, miR-382-5p specific in renal epithelial cells, and miR-762 specific in corneal epithelial cells. miRNA binding sites from any epithelial cell specific miRNA can be introduced to or removed from a polynucleotide of the invention to regulate expression of the polynucleotide in the epithelial cells.

[0309] In addition, a large group of miRNAs are enriched in embryonic stem cells, controlling stem cell self-renewal as well as the development and/or differentiation of various cell lineages, such as neural cells, cardiac, hematopoietic cells, skin cells, osteogenic cells and muscle cells (e.g., Kuppusamy KT et al., Curr. Mol Med, 2013, 13(5), 757-764; Vidigal JA and Ventura A, Semin Cancer Biol. 2012, 22(5-6), 428-436; Goff LA et al., PLoS One, 2009, 4:e7192; Morin RD et al., Genome Res,2008,18, 610-621; Yoo JK et al., Stem Cells Dev. 2012, 21(11), 2049-2057, each of which is herein incorporated by reference in its entirety). miRNAs abundant in embryonic stem cells include, but are not limited to, let-7a-2-3p, let-a-3p, let-7a-5p, let7d-3p, let-7d-5p, miR-103a-2-3p, miR-103a-5p, miR-106b-3p, miR-106b-5p, miR-1246, miR-1275, miR-138-1-3p, miR-138-2-3p, miR-138-5p, miR-154-3p, miR-154-5p, miR-200c-3p, miR-200c-5p, miR-290, miR-301a-3p, miR-301a-5p, miR-302a-3p, miR-302a-5p, miR-302b-3p, miR-302b-5p, miR-302c-3p, miR-302c-5p, miR-302d-3p, miR-302d-5p, miR-302e, miR-367-3p, miR-367-5p, miR-369-3p, miR-369-5p, miR-370, miR-371, miR-373, miR-380-5p, miR-423-3p, miR-423-5p, miR-486-5p, miR-520c-3p, miR-548e, miR-548f, miR-548g-3p, miR-548g-5p, miR-548i, miR-548k, miR-548l, miR-548m, miR-548n, miR-548o-3p, miR-548o-5p, miR-548p, miR-664a-3p, miR-664a-5p, miR-664b-3p, miR-664b-5p, miR-766-3p, miR-766-5p, miR-885-3p, miR-885-5p,miR-93-3p, miR-93-5p, miR-941,miR-96-3p, miR-96-5p, miR-99b-3p and miR-99b-5p. Many predicted novel miRNAs are discovered by deep sequencing in human embryonic stem cells (e.g., Morin RD et al., Genome Res,2008,18, 610-621; Goff LA et al., PLoS One, 2009, 4:e7192; Bar M et al., Stem cells, 2008, 26, 2496-2505, the content of each of which is incorporated herein by reference in its entirety).

[0310] In some embodiments, miRNAs are selected based on expression and abundance in immune cells of the hematopoietic lineage, such as B cells, T cells, macrophages, dendritic cells, and cells that are known to express TLR7/ TLR8 and/or able to secrete cytokines such as endothelial cells and platelets. In some embodiments, the miRNA set thus includes miRs that may be responsible in part for the immunogenicity of these cells, and such that a corresponding miR-site incorporation in polynucleotides of the present invention (e.g., mRNAs) could lead to destabilization of the mRNA and/or suppression of translation from these mRNAs in the specific cell type. Non-limiting representative examples include miR-142, miR-144, miR-150, miR-155 and miR-223, which are specific for many of the hematopoietic cells; miR-142, miR150, miR-16 and miR-223, which are expressed in B cells; miR-223, miR-451, miR-26a, miR-16, which are expressed in progenitor hematopoietic cells; and miR-126, which is expressed in plasmacytoid dendritic cells, platelets and endothelial cells. For further discussion of tissue expression of miRs see e.g., Teruel-Montoya, R. et al. (2014) *PLoS One* 9:e102259; Landgraf, P. et al. (2007) *Cell* 129:1401-1414; Bissels, U. et al. (2009) *RNA* 15:2375-2384. Any one miR-site incorporation in the 3' UTR and/or 5' UTR may mediate such effects in multiple cell types of interest (e.g., miR-142 is abundant in both B cells and dendritic cells).

[0311] In some embodiments, it may be beneficial to target the same cell type with multiple miRs and to incorporate binding sites to each of the 3p and 5p arm if both are abundant (e.g., both miR-142-3p and miR142-5p are abundant in hematopoietic stem cells). Thus, in certain embodiments, polynucleotides of the invention contain two or more (e.g., two, three, four or more) miR bindings sites from: (i) the group consisting of miR-142, miR-144, miR-150, miR-155 and miR-223 (which are expressed in many hematopoietic cells); or (ii) the group consisting of miR-142, miR150, miR-16 and miR-223 (which are expressed in B cells); or the group consisting of miR-223, miR-451, miR-26a, miR-16 (which are expressed in progenitor hematopoietic cells).

[0312] In some embodiments, it may also be beneficial to combine various miRs such that multiple cell types of interest are targeted at the same time (e.g., miR-142 and miR-126 to target many cells of the hematopoietic lineage and endothelial cells). Thus, for example, in certain embodiments, polynucleotides of the invention comprise two or more (e.g., two, three, four or more) miRNA bindings sites, wherein: (i) at least one of the miRs targets cells of the hematopoietic lineage (e.g., miR-142, miR-144, miR-150, miR-155 or miR-223) and at least one of the miRs targets

plasmacytoid dendritic cells, platelets or endothelial cells (e.g., miR-126); or (ii) at least one of the miRs targets B cells (e.g., miR-142, miR150, miR-16 or miR-223) and at least one of the miRs targets plasmacytoid dendritic cells, platelets or endothelial cells (e.g., miR-126); or (iii) at least one of the miRs targets progenitor hematopoietic cells (e.g., miR-223, miR-451, miR-26a or miR-16) and at least one of the miRs targets plasmacytoid dendritic cells, platelets or endothelial cells (e.g., miR-126); or (iv) at least one of the miRs targets cells of the hematopoietic lineage (e.g., miR-142, miR-144, miR-150, miR-155 or miR-223), at least one of the miRs targets B cells (e.g., miR-142, miR150, miR-16 or miR-223) and at least one of the miRs targets plasmacytoid dendritic cells, platelets or endothelial cells (e.g., miR-126); or any other possible combination of the foregoing four classes of miR binding sites (i.e., those targeting the hematopoietic lineage, those targeting B cells, those targeting progenitor hematopoietic cells and/or those targeting plasmacytoid dendritic cells/platelets/endothelial cells).

[0313] In one embodiment, to modulate immune responses, polynucleotides of the present invention can comprise one or more miRNA binding sequences that bind to one or more miRs that are expressed in conventional immune cells or any cell that expresses TLR7 and/or TLR8 and secrete pro-inflammatory cytokines and/or chemokines (e.g., in immune cells of peripheral lymphoid organs and/or splenocytes and/or endothelial cells). It has now been discovered that incorporation into an mRNA of one or more miRs that are expressed in conventional immune cells or any cell that expresses TLR7 and/or TLR8 and secrete pro-inflammatory cytokines and/or chemokines (e.g., in immune cells of peripheral lymphoid organs and/or splenocytes and/or endothelial cells) reduces or inhibits immune cell activation (e.g., B cell activation, as measured by frequency of activated B cells) and/or cytokine production (e.g., production of IL-6, IFN-γ and/or TNFα). Furthermore, it has now been discovered that incorporation into an mRNA of one or more miRs that are expressed in conventional immune cells or any cell that expresses TLR7 and/or TLR8 and secrete pro-inflammatory cytokines and/or chemokines (e.g., in immune cells of peripheral lymphoid organs and/or splenocytes and/or endothelial cells) can reduce or inhibit an anti-drug antibody (ADA) response against a protein of interest encoded by the mRNA.

[0314] In another embodiment, to modulate accelerated blood clearance of a polynucleotide delivered in a lipid-comprising compound or composition, polynucleotides of the invention can comprise one or more miR binding sequences that bind to one or more miRNAs expressed in conventional immune cells or any cell that expresses TLR7 and/or TLR8 and secrete pro-inflammatory cytokines and/or chemokines (e.g., in immune cells of peripheral lymphoid organs and/or splenocytes and/or endothelial cells). It has now been discovered that incorporation into an mRNA of one or more miR binding sites reduces or inhibits accelerated blood clearance (ABC) of the lipid-comprising compound or composition for use in delivering the mRNA. Furthermore, it has now been discovered that incorporation of one or more miR binding sites into an mRNA reduces serum levels of anti-PEG anti-IgM (e.g., reduces or inhibits the acute production of IgMs that recognize polyethylene glycol (PEG) by B cells) and/or reduces or inhibits proliferation and/or activation of plasmacytoid dendritic cells following administration of a lipid-comprising compound or composition comprising the mRNA.

[0315] In some embodiments, miR sequences may correspond to any known microRNA expressed in immune cells, including but not limited to those taught in US Publication US2005/0261218 and US Publication US2005/0059005, the contents of which are incorporated herein by reference in their entirety. Non-limiting examples of miRs expressed in immune cells include those expressed in spleen cells, myeloid cells, dendritic cells, plasmacytoid dendritic cells, B cells, T cells and/or macrophages. For example, miR-142-3p, miR-142-5p, miR-16, miR-21, miR-223, miR-24 and miR-27 are expressed in myeloid cells, miR-155 is expressed in dendritic cells, B cells and T cells, miR-146 is upregulated in macrophages upon TLR stimulation and miR-126 is expressed in plasmacytoid dendritic cells. In certain embodiments, the miR(s) is expressed abundantly or preferentially in immune cells. For example, miR-142 (miR-142-3p and/or miR-142-5p), miR-126 (miR-126-3p and/or miR-126-5p), miR-146 (miR-146-3p and/or miR-146-5p) and miR-155 (miR-155-3p and/or miR155-5p) are expressed abundantly in immune cells. These microRNA sequences are known in the art and, thus, one of ordinary skill in the art can readily design binding sequences or target sequences to which these microRNAs will bind based upon Watson-Crick complementarity.

[0316] Accordingly, in various embodiments, polynucleotides of the present invention comprise at least one microRNA binding site for a miR selected from the

group consisting of miR-142, miR-146, miR-155, miR-126, miR-16, miR-21, miR-223, miR-24 and miR-27. In another embodiment, the mRNA comprises at least two miR binding sites for microRNAs expressed in immune cells. In various embodiments, the polynucleotide of the invention comprises 1-4, one, two, three or four miR binding sites for microRNAs expressed in immune cells. In another embodiment, the polynucleotide of the invention comprises three miR binding sites. These miR binding sites can be for microRNAs selected from the group consisting of miR-142, miR-146, miR-155, miR-126, miR-16, miR-21, miR-223, miR-24, miR-27, and combinations thereof. In one embodiment, the polynucleotide of the invention comprises two or more (e.g., two, three, four) copies of the same miR binding site expressed in immune cells, e.g., two or more copies of a miR binding site selected from the group of miRs consisting of miR-142, miR-146, miR-155, miR-126, miR-16, miR-21, miR-223, miR-24, miR-27.

[0317] In one embodiment, the polynucleotide of the invention comprises three copies of the same miRNA binding site. In certain embodiments, use of three copies of the same miR binding site can exhibit beneficial properties as compared to use of a single miRNA binding site. Non-limiting examples of sequences for 3' UTRs containing three miRNA bindings sites are shown in SEQ ID NO: 155 (three miR-142-3p binding sites) and SEQ ID NO: 157 (three miR-142-5p binding sites).

[0318] In another embodiment, the polynucleotide of the invention comprises two or more (e.g., two, three, four) copies of at least two different miR binding sites expressed in immune cells. Non-limiting examples of sequences of 3' UTRs containing two or more different miR binding sites are shown in SEQ ID NO:111 (one miR-142-3p binding site and one miR-126-3p binding site), SEQ ID NO: 158 (two miR-142-5p binding sites and one miR-142-3p binding sites), and SEQ ID NO: 161 (two miR-155-5p binding sites and one miR-142-3p binding sites).

[0319] In another embodiment, the polynucleotide of the invention comprises at least two miR binding sites for microRNAs expressed in immune cells, wherein one of the miR binding sites is for miR-142-3p. In various embodiments, the polynucleotide of the invention comprises binding sites for miR-142-3p and miR-155 (miR-155-3p or miR-155-5p), miR-142-3p and miR-146 (miR-146-3 or miR-146-5p), or miR-142-3p and miR-126 (miR-126-3p or miR-126-5p).

[0320] In another embodiment, the polynucleotide of the invention comprises at least two miR binding sites for microRNAs expressed in immune cells, wherein one of the

miR binding sites is for miR-126-3p. In various embodiments, the polynucleotide of the invention comprises binding sites for miR-126-3p and miR-155 (miR-155-3p or miR-155-5p), miR-126-3p and miR-146 (miR-146-3p or miR-146-5p), or miR-126-3p and miR-142 (miR-142-3p or miR-142-5p).

[0321] In another embodiment, the polynucleotide of the invention comprises at least two miR binding sites for microRNAs expressed in immune cells, wherein one of the miR binding sites is for miR-142-5p. In various embodiments, the polynucleotide of the invention comprises binding sites for miR-142-5p and miR-155 (miR-155-3p or miR-155-5p), miR-142-5p and miR-146 (miR-146-3 or miR-146-5p), or miR-142-5p and miR-126 (miR-126-3p or miR-126-5p).

[0322] In yet another embodiment, the polynucleotide of the invention comprises at least two miR binding sites for microRNAs expressed in immune cells, wherein one of the miR binding sites is for miR-155-5p. In various embodiments, the polynucleotide of the invention comprises binding sites for miR-155-5p and miR-142 (miR-142-3p or miR-142-5p), miR-155-5p and miR-146 (miR-146-3 or miR-146-5p), or miR-155-5p and miR-126 (miR-126-3p or miR-126-5p).

[0323] miRNA can also regulate complex biological processes such as angiogenesis (e.g., miR-132) (Anand and Cheresh Curr Opin Hematol 2011 18:171-176). In the polynucleotides of the invention, miRNA binding sites that are involved in such processes can be removed or introduced, to tailor the expression of the polynucleotides to biologically relevant cell types or relevant biological processes. In this context, the polynucleotides of the invention are defined as auxotrophic polynucleotides.

[0324] In some embodiments, a polynucleotide of the invention comprises a miRNA binding site, wherein the miRNA binding site comprises one or more nucleotide sequences selected from Table 3, including one or more copies of any one or more of the miRNA binding site sequences. In some embodiments, a polynucleotide of the invention further comprises at least one, two, three, four, five, six, seven, eight, nine, ten, or more of the same or different miRNA binding sites selected from Table 3, including any combination thereof.

[0325] In some embodiments, the miRNA binding site binds to miR-142 or is complementary to miR-142. In some embodiments, the miR-142 comprises SEQ ID NO:114. In some embodiments, the miRNA binding site binds to miR-142-3p or miR-142-5p. In some embodiments, the miR-142-3p binding site comprises SEQ ID

NO:116. In some embodiments, the miR-142-5p binding site comprises SEQ ID NO:118. In some embodiments, the miRNA binding site comprises a nucleotide sequence at least 80%, at least 85%, at least 90%, at least 95%, or 100% identical to SEQ ID NO:116 or SEQ ID NO:118.

[0326] In some embodiments, the miRNA binding site binds to miR-126 or is complementary to miR-126. In some embodiments, the miR-126 comprises SEQ ID NO: 119. In some embodiments, the miRNA binding site binds to miR-126-3p or miR-126-5p. In some embodiments, the miR-126-3p binding site comprises SEQ ID NO: 121. In some embodiments, the miR-126-5p binding site comprises SEQ ID NO: 123. In some embodiments, the miRNA binding site comprises a nucleotide sequence at least 80%, at least 85%, at least 90%, at least 95%, or 100% identical to SEQ ID NO: 121 or SEQ ID NO: 123.

[0327] In one embodiment, the 3' UTR comprises two miRNA binding sites, wherein a first miRNA binding site binds to miR-142 and a second miRNA binding site binds to miR-126. In a specific embodiment, the 3' UTR binding to miR-142 and miR-126 comprises, consists, or consists essentially of the sequence of SEQ ID NO: 163.

TABLE 3. miR-142, miR-126, and miR-142 and miR-126 binding sites

SEQ ID NO.	Description	Sequence
114	miR-142	GACAGUGCAGUCACCCAUAAAGUAGAAAGCACUACUAACAGCACUGGAGGGUGUAGUGUUUCCUACUUUAUGGAUGAGUGUACUGUG
115	miR-142-3p	UGUAGUGUUUCCUACUUUAUGGA
116	miR-142-3p binding site	UCCAUAAAGUAGGAAACACUACA
117	miR-142-5p	CAUAAAGUAGAAAGCACUACU
118	miR-142-5p binding site	AGUAGUGCUUUCUACUUUAUG
119	miR-126	CGCUGGCGACGGGACAUUAUUACUUUUGGUACGCGCUGUGACACUUCAAACUCGUACCGUGAGUAAUAAUGCGCCGUCCACGGCA
120	miR-126-3p	UCGUACCGUGAGUAAUAAUGCG
121	miR-126-3p binding site	CGCAUUAUUACUCACGGUACGA
122	miR-126-5p	CAUUAUUACUUUUGGUACGCG
123	miR-126-5p binding site	CGCGUACCAAAAGUAAUAAUG

[0328] In some embodiments, a miRNA binding site is inserted in the polynucleotide of the invention in any position of the polynucleotide (e.g., the 5' UTR and/or 3' UTR). In some embodiments, the 5' UTR comprises a miRNA binding site. In some embodiments, the 3' UTR comprises a miRNA binding site. In some embodiments, the 5' UTR and the 3' UTR comprise a miRNA binding site. The insertion site in the

polynucleotide can be anywhere in the polynucleotide as long as the insertion of the miRNA binding site in the polynucleotide does not interfere with the translation of a functional polypeptide in the absence of the corresponding miRNA; and in the presence of the miRNA, the insertion of the miRNA binding site in the polynucleotide and the binding of the miRNA binding site to the corresponding miRNA are capable of degrading the polynucleotide or preventing the translation of the polynucleotide.

[0329] In some embodiments, a miRNA binding site is inserted in at least about 30 nucleotides downstream from the stop codon of an ORF in a polynucleotide of the invention comprising the ORF. In some embodiments, a miRNA binding site is inserted in at least about 10 nucleotides, at least about 15 nucleotides, at least about 20 nucleotides, at least about 25 nucleotides, at least about 30 nucleotides, at least about 35 nucleotides, at least about 40 nucleotides, at least about 45 nucleotides, at least about 50 nucleotides, at least about 55 nucleotides, at least about 60 nucleotides, at least about 65 nucleotides, at least about 70 nucleotides, at least about 75 nucleotides, at least about 80 nucleotides, at least about 85 nucleotides, at least about 90 nucleotides, at least about 95 nucleotides, or at least about 100 nucleotides downstream from the stop codon of an ORF in a polynucleotide of the invention. In some embodiments, a miRNA binding site is inserted in about 10 nucleotides to about 100 nucleotides, about 20 nucleotides to about 90 nucleotides, about 30 nucleotides to about 80 nucleotides, about 40 nucleotides to about 70 nucleotides, about 50 nucleotides to about 60 nucleotides, about 45 nucleotides to about 65 nucleotides downstream from the stop codon of an ORF in a polynucleotide of the invention.

[0330] In some embodiments, a miRNA binding site is inserted within the 3' UTR immediately following the stop codon of the coding region within the polynucleotide of the invention, e.g., mRNA. In some embodiments, if there are multiple copies of a stop codon in the construct, a miRNA binding site is inserted immediately following the final stop codon. In some embodiments, a miRNA binding site is inserted further downstream of the stop codon, in which case there are 3' UTR bases between the stop codon and the miR binding site(s). In some embodiments, three non-limiting examples of possible insertion sites for a miR in a 3' UTR are shown in SEQ ID NOs: 162, 163, and 164, which show a 3' UTR sequence with a miR-142-3p site inserted in one of three different possible insertion sites, respectively, within the 3' UTR.

[0331] In some embodiments, one or more miRNA binding sites can be positioned within the 5' UTR at one or more possible insertion sites. For example, three non-

limiting examples of possible insertion sites for a miR in a 5' UTR are shown in SEQ ID NOs: 165, 166, or 167, which show a 5' UTR sequence with a miR-142-3p site inserted into one of three different possible insertion sites, respectively, within the 5' UTR.

[0332] In one embodiment, a codon optimized open reading frame encoding a polypeptide of interest comprises a stop codon and the at least one microRNA binding site is located within the 3' UTR 1-100 nucleotides after the stop codon. In one embodiment, the codon optimized open reading frame encoding the polypeptide of interest comprises a stop codon and the at least one microRNA binding site for a miR expressed in immune cells is located within the 3' UTR 30-50 nucleotides after the stop codon. In another embodiment, the codon optimized open reading frame encoding the polypeptide of interest comprises a stop codon and the at least one microRNA binding site for a miR expressed in immune cells is located within the 3' UTR at least 50 nucleotides after the stop codon. In other embodiments, the codon optimized open reading frame encoding the polypeptide of interest comprises a stop codon and the at least one microRNA binding site for a miR expressed in immune cells is located within the 3' UTR immediately after the stop codon, or within the 3' UTR 15-20 nucleotides after the stop codon or within the 3' UTR 70-80 nucleotides after the stop codon. In other embodiments, the 3' UTR comprises more than one miRNA binding site (e.g., 2-4 miRNA binding sites), wherein there can be a spacer region (e.g., of 10-100, 20-70 or 30-50 nucleotides in length) between each miRNA binding site. In another embodiment, the 3' UTR comprises a spacer region between the end of the miRNA binding site(s) and the poly A tail nucleotides. For example, a spacer region of 10-100, 20-70 or 30-50 nucleotides in length can be situated between the end of the miRNA binding site(s) and the beginning of the poly A tail.

[0333] In one embodiment, a codon optimized open reading frame encoding a polypeptide of interest comprises a start codon and the at least one microRNA binding site is located within the 5' UTR 1-100 nucleotides before (upstream of) the start codon. In one embodiment, the codon optimized open reading frame encoding the polypeptide of interest comprises a start codon and the at least one microRNA binding site for a miR expressed in immune cells is located within the 5' UTR 10-50 nucleotides before (upstream of) the start codon. In another embodiment, the codon optimized open reading frame encoding the polypeptide of interest comprises a start codon and the at least one microRNA binding site for a miR expressed in immune

cells is located within the 5' UTR at least 25 nucleotides before (upstream of) the start codon. In other embodiments, the codon optimized open reading frame encoding the polypeptide of interest comprises a start codon and the at least one microRNA binding site for a miR expressed in immune cells is located within the 5' UTR immediately before the start codon, or within the 5' UTR 15-20 nucleotides before the start codon or within the 5' UTR 70-80 nucleotides before the start codon. In other embodiments, the 5' UTR comprises more than one miRNA binding site (e.g., 2-4 miRNA binding sites), wherein there can be a spacer region (e.g., of 10-100, 20-70 or 30-50 nucleotides in length) between each miRNA binding site.

[0334] In one embodiment, the 3' UTR comprises more than one stop codon, wherein at least one miRNA binding site is positioned downstream of the stop codons. For example, a 3' UTR can comprise 1, 2 or 3 stop codons. Non-limiting examples of triple stop codons that can be used include: UGAUAAUAG (SEQ ID NO:124), UGAUAGUAA (SEQ ID NO:125), UAAUGAUAG (SEQ ID NO:126), UGAUAAUAA (SEQ ID NO:127), UGAUAGUAG (SEQ ID NO:128), UAAUGAUGA (SEQ ID NO:129), UAAUAGUAG (SEQ ID NO:130), UGAUGAUGA (SEQ ID NO:131), UAAUAAUAA (SEQ ID NO:132), and UAGUAGUAG (SEQ ID NO:133). Within a 3' UTR, for example, 1, 2, 3 or 4 miRNA binding sites, e.g., miR-142-3p binding sites, can be positioned immediately adjacent to the stop codon(s) or at any number of nucleotides downstream of the final stop codon. When the 3' UTR comprises multiple miRNA binding sites, these binding sites can be positioned directly next to each other in the construct (i.e., one after the other) or, alternatively, spacer nucleotides can be positioned between each binding site.

[0335] In one embodiment, the 3' UTR comprises three stop codons with a single miR-142-3p binding site located downstream of the 3rd stop codon. Non-limiting examples of sequences of 3' UTR having three stop codons and a single miR-142-3p binding site located at different positions downstream of the final stop codon are shown in SEQ ID NOs: 151, 162, 163, and 164.

TABLE 4A. 5' UTRs, 3'UTRs, miR sequences, and miR binding sites

SEQ ID NO:	Sequence
134	GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCC UCCUCCCCUUCCUGCACCCGUACCCCCCUCCAUAAAGUAGGAAACACUACAGU GGUCUUUGAAUAAAGUCUGAGUGGGCGGC

	(3' UTR with miR 142-3p binding site)
116	UCCAUAAAGUAGGAAACACUACA (miR 142-3p binding site)
115	UGUAGUGUUUCCUACUUUAUGGA (miR 142-3p sequence)
117	CAUAAAGUAGAAAGCACUACU (miR 142-5p sequence)
135	CCUCUGAAAUUCAGUUCUUCAG (miR 146-3p sequence)
136	UGAGAACUGAAUUCCAUGGGUU (miR 146-5p sequence)
137	CUCCUACAUAUUAGCAUUAACA (miR 155-3p sequence)
138	UUAAUGCUAAUCGUGAUAGGGGU (miR 155-5p sequence)
120	UCGUACCGUGAGUAAUAAUGCG (miR 126-3p sequence)
122	CAUUAUUACUUUUGGUACGCG (miR 126-5p sequence)
139	CCAGUAUUAACUGUGCUGCUGA (miR 16-3p sequence)
140	UAGCAGCACGUAAAUAUUGGCG (miR 16-5p sequence)
141	CAACACCAGUCGAUGGGCUGU (miR 21-3p sequence)
142	UAGCUUAUCAGACUGAUGUUGA (miR 21-5p sequence)
143	UGUCAGUUUGUCAAAUACCCCA (miR 223-3p sequence)
144	CGUGUAUUUGACAAGCUGAGUU (miR 223-5p sequence)
145	UGGCUCAGUUCAGCAGGAACAG (miR 24-3p sequence)
146	UGCCUACUGAGCUGAUAUCAGU (miR 24-5p sequence)
147	UUCACAGUGGCUAAGUUCCGC (miR 27-3p sequence)
148	AGGGCUUAGCUGCUUGUGAGCA (miR 27-5p sequence)
121	CGCAUUAUUACUCACGGUACGA (miR 126-3p binding site)
149	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCC<u>CGCAUUAUUACUCACG</u> <u>GUACGA</u>GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 126-3p binding site)
150	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUAAA GUCUGAGUGGGCGGC (3' UTR, no miR binding sites)

151	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGUAGGAAA CACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 142-3p binding site)
111	**UGAUAAUAG**UCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCAU GCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCCGCAUUAUUACUCACGGUACGAGUGGUCUUUGAAUAAAGUCUGAG UGGGCGGC (3' UTR with miR 142-3p and miR 126-3p binding sites variant 1)
153	UUAAUGCUAAUUGUGAUAGGGGU (miR 155-5p sequence)
154	ACCCCUAUCACAAUUAGCAUUAA (miR 155-5p binding site)
155	**UGAUAAUAG**UCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCAU GCUUCUUGCCCCUUGGGCCUCCAUAAAGUAGGAAACACUACAUCCCCCCAGC CCCUCCUCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGUAGGAAACACUAC AGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 3 miR 142-3p binding sites)
156	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCC*AGUAGUGCUUUCUACU UUAUG*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 142-5p binding site)
157	**UGAUAAUAG***AGUAGUGCUUUCUACUUUAUG*GCUGGAGCCUCGGUGGCCAUGC UUCUUGCCCCUUGGGCC*AGUAGUGCUUUCUACUUUAUG*UCCCCCCAGCCCCU CCUCCCCUUCCUGCACCCGUACCCCC*AGUAGUGCUUUCUACUUUAUG*GUGGU CUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 3 miR 142-5p binding sites)
158	**UGAUAAUAG***AGUAGUGCUUUCUACUUUAUG*GCUGGAGCCUCGGUGGCCAUGC UUCUUGCCCCUUGGGCCUCCAUAAAGUAGGAAACACUACAUCCCCCCAGCCC CUCCUCCCCUUCCUGCACCCGUACCCCC*AGUAGUGCUUUCUACUUUAUG*GUG GUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 2 miR 142-5p binding sites and 1 miR 142-3p binding site)
159	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCC*ACCCCUAUCACAAUUA GCAUUAA*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 155-5p binding site)
160	**UGAUAAUAG***ACCCCUAUCACAAUUAGCAUUAA*GCUGGAGCCUCGGUGGCCAU GCUUCUUGCCCCUUGGGCC*ACCCCUAUCACAAUUAGCAUUAA*UCCCCCCAGC CCCUCCUCCCCUUCCUGCACCCGUACCCCC*ACCCCUAUCACAAUUAGCAUUA A*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 3 miR 155-5p binding sites)
161	**UGAUAAUAG***ACCCCUAUCACAAUUAGCAUUAA*GCUGGAGCCUCGGUGGCCAU GCUUCUUGCCCCUUGGGCCUCCAUAAAGUAGGAAACACUACAUCCCCCCAGC CCCUCCUCCCCUUCCUGCACCCGUACCCCC*ACCCCUAUCACAAUUAGCAUUA A*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 2 miR 155-5p binding sites and 1 miR 142-3p binding site)
162	**UGAUAAUAG**UCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCAU GCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 142-3p binding site, P1 insertion)

163	**UGAUAAUAG**GCUGGAGCCUCGGUGGCUCCAUAAAGUAGGAAACACUACACAU GCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 142-3p binding site, P2 insertion)
164	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCA UAAAGUAGGAAACACUACAUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 142-3p binding site, P3 insertion)
118	AGUAGUGCUUUCUACUUUAUG (miR-142-5p binding site)
114	GACAGUGCAGUCACCCAUAAAGUAGAAAGCACUACUAACAGCACUGGAGGGU GUAGUGUUUCCUACUUUAUGGAUGAGUGUACUGUG (miR-142)
3	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC (5' UTR)
165	GGGAAAUAAGAGUCCAUAAAGUAGGAAACACUACAAGAAAAGAAGAGUAAGA AGAAAUAUAAGAGCCACC (5' UTR with miR142-3p binding site at position p1)
166	GGGAAAUAAGAGAGAAAAGAAGAGUAAUCCAUAAAGUAGGAAACACUACAGA AGAAAUAUAAGAGCCACC (5' UTR with miR142-3p binding site at position p2)
167	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAUCCAUAAAGUAGG AAACACUACAGAGCCACC (5' UTR with miR142-3p binding site at position p3)
168	ACCCCUAUCACAAUUAGCAUUAA (miR 155-5p binding site)
169	**UGAUAAUAG***AGUAGUGCUUUCUACUUUAUG*GCUGGAGCCUCGGUGGCCAUGC UUCUUGCCCCUUGGGCC*AGUAGUGCUUUCUACUUUAUG*UCCCCCCAGCCCCU CUCCCCUUCCUGCACCCGUACCCCC*AGUAGUGCUUUCUACUUUAUG*GUGGUC UUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 3 miR 142-5p binding sites)
170	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUCCAUAAAGU AGGAAACACUACAUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR including miR142-3p binding site)
171	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGUCCAUAAAGUAGGAAACACUACACCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR including miR142-3p binding site)
172	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCUUCUCCAUAAAGUAGGAAACACUACACUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR including miR142-3p binding site)
173	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUAAA GUUCCAUAAAGUAGGAAACACUACACUGAGUGGGCGGC (3'UTR including miR142-3p binding site)
174	**UGAUAAUAG**UCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCUA GCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG

	UACCCCCCGCAUUAUUACUCACGGUACGAGUGGUCUUUGAAUAAAGUCUGAG UGGGCGGC (3' UTR with miR 142-3p and miR 126-3p binding sites variant 2)
175	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUAAA GUCUGAGUGGGCGGC (3' UTR, no miR binding sites variant 2)
4	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGUAGGAAA CACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 142-3p binding site variant 3)
177	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCCCGCAUUAUUACUCACG GUACGAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with miR 126-3p binding site variant 3)
178	**UGAUAAUAG**UCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCUA GCUUCUUGCCCCUUGGGCCUCCAUAAAGUAGGAAACACUACAUCCCCCCAGC CCCUCCUCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGUAGGAAACACUAC AGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 3 miR 142-3p binding sites variant 2)
179	**UGAUAAUAG**UCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCUA GCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR with miR 142-3p binding site, P1 insertion variant 2)
180	**UGAUAAUAG**GCUGGAGCCUCGGUGGCUCCAUAAAGUAGGAAACACUACACUA GCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR with miR 142-3p binding site, P2 insertion variant 2)
181	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCA UAAAGUAGGAAACACUACAUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCG UACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR with miR 142-3p binding site, P3 insertion variant 2)
182	**UGAUAAUAG**GCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCC CCCAGCCCCUCCUCCCCCUUCCUGCACCCGUACCCCC*ACCCCUAUCACAAUUA GCAUUAA*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR with miR 155-5p binding site variant 2)
183	**UGAUAAUAG***ACCCCUAUCACAAUUAGCAUUAA*GCUGGAGCCUCGGUGGCCUA GCUUCUUGCCCCUUGGGCC*ACCCCUAUCACAAUUAGCAUUAA*UCCCCCCCAGC CCCUCCUCCCCUUCCUGCACCCGUACCCCC*ACCCCUAUCACAAUUAGCAUUA A*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3' UTR with 3 miR 155-5p binding sites variant 2)
184	**UGAUAAUAG***ACCCCUAUCACAAUUAGCAUUAA*GCUGGAGCCUCGGUGGCCUA GCUUCUUGCCCCUUGGGCCUCCAUAAAGUAGGAAACACUACAUCCCCCCAGC CCCUCCUCCCCUUCCUGCACCCGUACCCCC*ACCCCUAUCACAAUUAGCAUUA A*GUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (3'UTR with 2 miR 155-5p binding sites and 1 miR 142-3p binding site variant 2)

Stop codon = bold

miR 142-3p binding site = underline

miR 126-3p binding site = bold underline

miR 155-5p binding site = italicized

miR 142-5p binding site = italicized and bold underline

TABLE 4B. Exemplary Preferred UTRs

SEQ ID NO:	**Sequence**
5' UTR (v1) (SEQ ID NO:3)	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC
5'UTR (v1 A) (SEQ ID NO:210)	AGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC
5' UTR (v1.1) (SEQ ID NO:39)	GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGA**CCCCGGC GCC**GCCACC
5' UTR (v1.1 A) (SEQ ID NO:211)	AGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGA**CCCCGGC GCC**GCCACC
3' UTR (v1) (SEQ ID NO:150)	UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCC UCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCGUGGUCUU UGAAUAAAGUCUGAGUGGGCGGC
3' UTR (v1.1) (SEQ ID NO:175)	UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCC UCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCGUGGUCUU UGAAUAAAGUCUGAGUGGGCGGC
3' UTR (miR122) (SEQ ID NO:212)	UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCC UCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCCAAACACC AUUGUCACACUCCAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC
3' UTR (v1.1 miR122) (SEQ ID NO:213)	UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCC UCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCCAAACACC AUUGUCACACUCCAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC
3' UTR (v1.1 mir142-3p) (SEQ ID NO:4)	UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCC UCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCUCCAUAAA GUAGGAAACACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC
3' UTR (v1.1 mir 126-3p) (SEQ ID NO:177)	UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCC UCCCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCCGCAUUAU UACUCACGGUACGAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC
3' UTR (mir-126, miR-142-3p) (SEQ ID NO:111)	UGAUAAUAGUCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGG CCAUGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUCCUCCCCUUCC UGCACCCGUACCCCCCGCAUUAUUACUCACGGUACGAGUGGUCUUUGA AUAAAGUCUGAGUGGGCGGC
3' UTR (v.1.1 3x miR142-3p) (SEQ ID NO:178)	UGAUAAUAGUCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGG CCUAGCUUCUUGCCCCUUGGGCCUCCAUAAAGUAGGAAACACUACAUC CCCCCAGCCCCUCCUCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGU AGGAAACACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC

[0336] In one embodiment, the polynucleotide of the invention comprises a 5' UTR, a codon optimized open reading frame encoding a polypeptide of interest, a 3' UTR comprising the at least one miRNA binding site for a miR expressed in immune cells, and a 3' tailing region of linked nucleosides. In various embodiments, the 3' UTR comprises 1-4, at least two, one, two, three or four miRNA binding sites for miRs expressed in immune cells, preferably abundantly or preferentially expressed in immune cells.

[0337] In one embodiment, the at least one miRNA expressed in immune cells is a miR-142-3p microRNA binding site. In one embodiment, the miR-142-3p microRNA binding site comprises the sequence shown in SEQ ID NO: 116. In one embodiment, the 3' UTR of the mRNA comprising the miR-142-3p microRNA binding site comprises the sequence shown in SEQ ID NO: 134.

[0338] In one embodiment, the at least one miRNA expressed in immune cells is a miR-126 microRNA binding site. In one embodiment, the miR-126 binding site is a miR-126-3p binding site. In one embodiment, the miR-126-3p microRNA binding site comprises the sequence shown in SEQ ID NO: 121. In one embodiment, the 3' UTR of the mRNA of the invention comprising the miR-126-3p microRNA binding site comprises the sequence shown in SEQ ID NO: 149.

[0339] Non-limiting exemplary sequences for miRs to which a microRNA binding site(s) of the disclosure can bind include the following: miR-142-3p (SEQ ID NO: 115), miR-142-5p (SEQ ID NO: 117), miR-146-3p (SEQ ID NO: 135), miR-146-5p (SEQ ID NO: 136), miR-155-3p (SEQ ID NO: 137), miR-155-5p (SEQ ID NO: 138), miR-126-3p (SEQ ID NO: 120), miR-126-5p (SEQ ID NO: 122), miR-16-3p (SEQ ID NO: 139), miR-16-5p (SEQ ID NO: 140), miR-21-3p (SEQ ID NO: 141), miR-21-5p (SEQ ID NO: 142), miR-223-3p (SEQ ID NO: 143), miR-223-5p (SEQ ID NO: 144), miR-24-3p (SEQ ID NO: 145), miR-24-5p (SEQ ID NO: 146), miR-27-3p (SEQ ID NO: 147) and miR-27-5p (SEQ ID NO: 148). Other suitable miR sequences expressed in immune cells (e.g., abundantly or preferentially expressed in immune cells) are known and available in the art, for example at the University of Manchester's microRNA database, miRBase. Sites that bind any of the aforementioned miRs can be designed based on Watson-Crick complementarity to the miR, typically 100% complementarity to the miR, and inserted into an mRNA construct of the disclosure as described herein.

[0340] In another embodiment, a polynucleotide of the present invention (e.g., and mRNA, e.g., the 3' UTR thereof) can comprise at least one miRNA binding site to thereby reduce or inhibit accelerated blood clearance, for example by reducing or inhibiting production of IgMs, e.g., against PEG, by B cells and/or reducing or inhibiting proliferation and/or activation of pDCs, and can comprise at least one miRNA binding site for modulating tissue expression of an encoded protein of interest.

[0341] miRNA gene regulation can be influenced by the sequence surrounding the miRNA such as, but not limited to, the species of the surrounding sequence, the type of sequence (e.g., heterologous, homologous, exogenous, endogenous, or artificial), regulatory elements in the surrounding sequence and/or structural elements in the surrounding sequence. The miRNA can be influenced by the 5′UTR and/or 3′UTR. As a non-limiting example, a non-human 3′UTR can increase the regulatory effect of the miRNA sequence on the expression of a polypeptide of interest compared to a human 3′ UTR of the same sequence type.

[0342] In one embodiment, other regulatory elements and/or structural elements of the 5′ UTR can influence miRNA mediated gene regulation. One example of a regulatory element and/or structural element is a structured IRES (Internal Ribosome Entry Site) in the 5′ UTR, which is necessary for the binding of translational elongation factors to initiate protein translation. EIF4A2 binding to this secondarily structured element in the 5′-UTR is necessary for miRNA mediated gene expression (Meijer HA et al., Science, 2013, 340, 82-85, herein incorporated by reference in its entirety). The polynucleotides of the invention can further include this structured 5′ UTR to enhance microRNA mediated gene regulation.

[0343] At least one miRNA binding site can be engineered into the 3′ UTR of a polynucleotide of the invention. In this context, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, at least ten, or more miRNA binding sites can be engineered into a 3' UTR of a polynucleotide of the invention. For example, 1 to 10, 1 to 9, 1 to 8, 1 to 7, 1 to 6, 1 to 5, 1 to 4, 1 to 3, 2, or 1 miRNA binding sites can be engineered into the 3′UTR of a polynucleotide of the invention. In one embodiment, miRNA binding sites incorporated into a polynucleotide of the invention can be the same or can be different miRNA sites. A combination of different miRNA binding sites incorporated into a polynucleotide of the invention can include combinations in which more than one copy of any of the

different miRNA sites are incorporated. In another embodiment, miRNA binding sites incorporated into a polynucleotide of the invention can target the same or different tissues in the body. As a non-limiting example, through the introduction of tissue-, cell-type-, or disease-specific miRNA binding sites in the 3′-UTR of a polynucleotide of the invention, the degree of expression in specific cell types (e.g., myeloid cells, endothelial cells, etc.) can be reduced.

[0344] In one embodiment, a miRNA binding site can be engineered near the 5′ terminus of the 3′UTR, about halfway between the 5′ terminus and 3′ terminus of the 3′UTR and/or near the 3′ terminus of the 3′ UTR in a polynucleotide of the invention. As a non-limiting example, a miRNA binding site can be engineered near the 5′ terminus of the 3′UTR and about halfway between the 5′ terminus and 3′ terminus of the 3′UTR. As another non-limiting example, a miRNA binding site can be engineered near the 3′ terminus of the 3′UTR and about halfway between the 5′ terminus and 3′ terminus of the 3′ UTR. In another non-limiting example, a miRNA binding site can be engineered near the 5′ terminus of the 3′ UTR and near the 3′ terminus of the 3′ UTR.

[0345] In another embodiment, a 3′UTR can comprise 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 miRNA binding sites. The miRNA binding sites can be complementary to a miRNA, miRNA seed sequence, and/or miRNA sequences flanking the seed sequence.

[0346] In some embodiments, the expression of a polynucleotide of the invention can be controlled by incorporating at least one sensor sequence in the polynucleotide and formulating the polynucleotide for administration. As a non-limiting example, a polynucleotide of the invention can be targeted to a tissue or cell by incorporating a miRNA binding site and formulating the polynucleotide in a lipid nanoparticle comprising a ionizable lipid, including any of the lipids described herein.

[0347] A polynucleotide of the invention can be engineered for more targeted expression in specific tissues, cell types, or biological conditions based on the expression patterns of miRNAs in the different tissues, cell types, or biological conditions. Through introduction of tissue-specific miRNA binding sites, a polynucleotide of the invention can be designed for optimal protein expression in a tissue or cell, or in the context of a biological condition.

[0348] In some embodiments, a polynucleotide of the invention can be designed to incorporate miRNA binding sites that either have 100% identity to known miRNA seed sequences or have less than 100% identity to miRNA seed sequences. In some

embodiments, a polynucleotide of the invention can be designed to incorporate miRNA binding sites that have at least: 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, or 99% identity to known miRNA seed sequences. The miRNA seed sequence can be partially mutated to decrease miRNA binding affinity and as such result in reduced downmodulation of the polynucleotide. In essence, the degree of match or mis-match between the miRNA binding site and the miRNA seed can act as a rheostat to more finely tune the ability of the miRNA to modulate protein expression. In addition, mutation in the non-seed region of a miRNA binding site can also impact the ability of a miRNA to modulate protein expression.

[0349] In one embodiment, a miRNA sequence can be incorporated into the loop of a stem loop.

[0350] In another embodiment, a miRNA seed sequence can be incorporated in the loop of a stem loop and a miRNA binding site can be incorporated into the 5′ or 3′ stem of the stem loop.

[0351] In one embodiment the miRNA sequence in the 5′ UTR can be used to stabilize a polynucleotide of the invention described herein.

[0352] In another embodiment, a miRNA sequence in the 5′ UTR of a polynucleotide of the invention can be used to decrease the accessibility of the site of translation initiation such as, but not limited to a start codon. See, e.g., Matsuda et al., PLoS One. 2010 11(5):e15057; incorporated herein by reference in its entirety, which used antisense locked nucleic acid (LNA) oligonucleotides and exon-junction complexes (EJCs) around a start codon (-4 to +37 where the A of the AUG codons is +1) to decrease the accessibility to the first start codon (AUG). Matsuda showed that altering the sequence around the start codon with an LNA or EJC affected the efficiency, length and structural stability of a polynucleotide. A polynucleotide of the invention can comprise a miRNA sequence, instead of the LNA or EJC sequence described by Matsuda et al, near the site of translation initiation to decrease the accessibility to the site of translation initiation. The site of translation initiation can be prior to, after or within the miRNA sequence. As a non-limiting example, the site of translation initiation can be located within a miRNA sequence such as a seed sequence or binding site.

[0353] In some embodiments, a polynucleotide of the invention can include at least one miRNA to dampen the antigen presentation by antigen presenting cells. The miRNA can be the complete miRNA sequence, the miRNA seed sequence, the

miRNA sequence without the seed, or a combination thereof. As a non-limiting example, a miRNA incorporated into a polynucleotide of the invention can be specific to the hematopoietic system. As another non-limiting example, a miRNA incorporated into a polynucleotide of the invention to dampen antigen presentation is miR-142-3p.

[0354] In some embodiments, a polynucleotide of the invention can include at least one miRNA to dampen expression of the encoded polypeptide in a tissue or cell of interest. As a non-limiting example, a polynucleotide of the invention can include at least one miR-142-3p binding site, miR-142-3p seed sequence, miR-142-3p binding site without the seed, miR-142-5p binding site, miR-142-5p seed sequence, miR-142-5p binding site without the seed, miR-146 binding site, miR-146 seed sequence and/or miR-146 binding site without the seed sequence.

[0355] In some embodiments, a polynucleotide of the invention can comprise at least one miRNA binding site in the 3′UTR to selectively degrade mRNA therapeutics in the immune cells to subdue unwanted immunogenic reactions caused by therapeutic delivery. As a non-limiting example, the miRNA binding site can make a polynucleotide of the invention more unstable in antigen presenting cells. Non-limiting examples of these miRNAs include miR-142-5p, miR-142-3p, miR-146a-5p, and miR-146-3p.

[0356] In one embodiment, a polynucleotide of the invention comprises at least one miRNA sequence in a region of the polynucleotide that can interact with a RNA binding protein.

[0357] In some embodiments, the polynucleotide of the invention (e.g., a RNA, e.g., an mRNA) comprising (i) a sequence-optimized nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof) and (ii) a miRNA binding site (e.g., a miRNA binding site that binds to miR-142) and/or a miRNA binding site that binds to miR-126.

12. 3′ UTRs

[0358] In certain embodiments, a polynucleotide of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide of the invention) further comprises a 3' UTR.

[0359] 3'-UTR is the section of mRNA that immediately follows the translation termination codon and often contains regulatory regions that post-transcriptionally

influence gene expression. Regulatory regions within the 3'-UTR can influence polyadenylation, translation efficiency, localization, and stability of the mRNA. In one embodiment, the 3'-UTR useful for the invention comprises a binding site for regulatory proteins or microRNAs.

[0360] In certain embodiments, the 3' UTR useful for the polynucleotides of the invention comprises a 3' UTR selected from the group consisting of SEQ ID NO: 4, 104 to 113, 150, and 175, or any combination thereof. In some embodiments, the 3' UTR comprises a nucleic acid sequence selected from the group consisting of SEQ ID NOs: 111, 112, or 113 or any combination thereof. In some embodiments, the 3' UTR comprises a nucleic acid sequence of SEQ ID NO: 111. In some embodiments, the 3' UTR comprises a nucleic acid sequence of SEQ ID NO: 112. In some embodiments, the 3' UTR comprises a nucleic acid sequences of SEQ ID NO: 113. In some embodiments, the 3' UTR comprises a nucleic acid sequences of SEQ ID NO: 150. In some embodiments, the 3' UTR comprises a nucleic acid sequences of SEQ ID NO: 175. In some embodiments, the 3' UTR comprises a nucleic acid sequences of SEQ ID NO:212. In some embodiments, the 3' UTR comprises a nucleic acid sequences of SEQ ID NO:213.

[0361] In certain embodiments, the 3' UTR sequence useful for the invention comprises a nucleotide sequence at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or about 100% identical to a sequence selected from the group consisting of 3' UTR sequences selected from the group consisting of SEQ ID NOs: 4, 104 to 113, 150, and 175, or any combination thereof.

[0362] In certain embodiments, the 3' UTR sequence useful for the invention comprises a nucleotide sequence at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or about 100% identical to a sequence selected from the group consisting of 3' UTR sequences selected from the group consisting of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213, or any combination thereof.

13. Regions having a 5′ Cap

[0363] The disclosure also includes a polynucleotide that comprises both a 5′ Cap and a polynucleotide of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide).

[0364] The 5′ cap structure of a natural mRNA is involved in nuclear export, increasing mRNA stability and binds the mRNA Cap Binding Protein (CBP), which is responsible for mRNA stability in the cell and translation competency through the association of CBP with poly(A) binding protein to form the mature cyclic mRNA species. The cap further assists the removal of 5′ proximal introns during mRNA splicing.

[0365] Endogenous mRNA molecules can be 5′-end capped generating a 5′-ppp-5′-triphosphate linkage between a terminal guanosine cap residue and the 5′-terminal transcribed sense nucleotide of the mRNA molecule. This 5′-guanylate cap can then be methylated to generate an N7-methyl-guanylate residue. The ribose sugars of the terminal and/or ante-terminal transcribed nucleotides of the 5′ end of the mRNA can optionally also be 2′-O-methylated. 5′-decapping through hydrolysis and cleavage of the guanylate cap structure can target a nucleic acid molecule, such as an mRNA molecule, for degradation.

[0366] In some embodiments, the polynucleotides of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) incorporate a cap moiety.

[0367] In some embodiments, polynucleotides of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) comprise a non-hydrolyzable cap structure preventing decapping and thus increasing mRNA half-life. Because cap structure hydrolysis requires cleavage of 5′-ppp-5′ phosphorodiester linkages, modified nucleotides can be used during the capping reaction. For example, a Vaccinia Capping Enzyme from New England Biolabs (Ipswich, MA) can be used with α-thio-guanosine nucleotides according to the manufacturer's instructions to create a phosphorothioate linkage in the 5′-ppp-5′ cap. Additional modified guanosine nucleotides can be used such as α-methyl-phosphonate and seleno-phosphate nucleotides.

[0368] Additional modifications include, but are not limited to, 2′-O-methylation of the ribose sugars of 5′-terminal and/or 5′-anteterminal nucleotides of the

polynucleotide (as mentioned above) on the 2′-hydroxyl group of the sugar ring. Multiple distinct 5′-cap structures can be used to generate the 5′-cap of a nucleic acid molecule, such as a polynucleotide that functions as an mRNA molecule. Cap analogs, which herein are also referred to as synthetic cap analogs, chemical caps, chemical cap analogs, or structural or functional cap analogs, differ from natural (*i.e.*, endogenous, wild-type or physiological) 5′-caps in their chemical structure, while retaining cap function. Cap analogs can be chemically (*i.e.*, non-enzymatically) or enzymatically synthesized and/or linked to the polynucleotides of the invention.

[0369] For example, the Anti-Reverse Cap Analog (ARCA) cap contains two guanines linked by a 5′-5′-triphosphate group, wherein one guanine contains an N7 methyl group as well as a 3′-O-methyl group (*i.e.*, N7,3′-O-dimethyl-guanosine-5′-triphosphate-5′-guanosine (m^7G-3′mppp-G; which can equivalently be designated 3′ O-Me-m7G(5')ppp(5')G). The 3′-O atom of the other, unmodified, guanine becomes linked to the 5′-terminal nucleotide of the capped polynucleotide. The N7- and 3′-O-methlyated guanine provides the terminal moiety of the capped polynucleotide.

[0370] Another exemplary cap is mCAP, which is similar to ARCA but has a 2′-O-methyl group on guanosine (*i.e.*, N7,2′-O-dimethyl-guanosine-5′-triphosphate-5′-guanosine, m^7Gm-ppp-G).

[0371] In some embodiments, the cap is a dinucleotide cap analog. As a non-limiting example, the dinucleotide cap analog can be modified at different phosphate positions with a boranophosphate group or a phosphoroselenoate group such as the dinucleotide cap analogs described in U.S. Patent No. US 8,519,110, the contents of which are herein incorporated by reference in its entirety.

[0372] In another embodiment, the cap is a cap analog is a N7-(4-chlorophenoxyethyl) substituted dinucleotide form of a cap analog known in the art and/or described herein. Non-limiting examples of a N7-(4-chlorophenoxyethyl) substituted dinucleotide form of a cap analog include a N7-(4-chlorophenoxyethyl)-G(5')ppp(5')G and a N7-(4-chlorophenoxyethyl)-$m^{3'\text{-}O}$G(5')ppp(5')G cap analog (*See, e.g.,* the various cap analogs and the methods of synthesizing cap analogs described in Kore et al. Bioorganic & Medicinal Chemistry 2013 21:4570-4574; the contents of which are herein incorporated by reference in its entirety). In another embodiment, a cap analog of the present invention is a 4-chloro/bromophenoxyethyl analog.

[0373] While cap analogs allow for the concomitant capping of a polynucleotide or a region thereof, in an *in vitro* transcription reaction, up to 20% of transcripts can

remain uncapped. This, as well as the structural differences of a cap analog from an endogenous 5′-cap structures of nucleic acids produced by the endogenous, cellular transcription machinery, can lead to reduced translational competency and reduced cellular stability.

[0374] Polynucleotides of the invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) can also be capped post-manufacture (whether IVT or chemical synthesis), using enzymes, to generate more authentic 5′-cap structures. As used herein, the phrase "more authentic" refers to a feature that closely mirrors or mimics, either structurally or functionally, an endogenous or wild type feature. That is, a "more authentic" feature is better representative of an endogenous, wild-type, natural or physiological cellular function and/or structure as compared to synthetic features or analogs, etc., of the prior art, or which outperforms the corresponding endogenous, wild-type, natural or physiological feature in one or more respects. Non-limiting examples of more authentic 5′cap structures of the present invention are those that, among other things, have enhanced binding of cap binding proteins, increased half-life, reduced susceptibility to 5′ endonucleases and/or reduced 5′decapping, as compared to synthetic 5′cap structures known in the art (or to a wild-type, natural or physiological 5′cap structure). For example, recombinant Vaccinia Virus Capping Enzyme and recombinant 2′-O-methyltransferase enzyme can create a canonical 5′-5′-triphosphate linkage between the 5′-terminal nucleotide of a polynucleotide and a guanine cap nucleotide wherein the cap guanine contains an N7 methylation and the 5′-terminal nucleotide of the mRNA contains a 2′-O-methyl. Such a structure is termed the Cap1 structure. This cap results in a higher translational-competency and cellular stability and a reduced activation of cellular pro-inflammatory cytokines, as compared, *e.g.,* to other 5′cap analog structures known in the art. Cap structures include, but are not limited to, 7mG(5')ppp(5')N,pN2p (cap 0), 7mG(5')ppp(5')NlmpNp (cap 1), and 7mG(5')-ppp(5')NlmpN2mp (cap 2).

[0375] As a non-limiting example, capping chimeric polynucleotides post-manufacture can be more efficient as nearly 100% of the chimeric polynucleotides can be capped. This is in contrast to ~80% efficiency when a cap analog is linked to a chimeric polynucleotide during an *in vitro* transcription reaction.

[0376] According to the present invention, 5′ terminal caps can include endogenous caps or cap analogs. According to the present invention, a 5′ terminal cap can

comprise a guanine analog. Useful guanine analogs include, but are not limited to, inosine, N1-methyl-guanosine, 2′fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, and 2-azido-guanosine.

14. Poly-A Tails

[0377] In some embodiments, the polynucleotides of the present disclosure (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) further comprise a poly-A tail. In further embodiments, terminal groups on the poly-A tail can be incorporated for stabilization. In other embodiments, a poly-A tail comprises des-3' hydroxyl tails.

[0378] During RNA processing, a long chain of adenine nucleotides (poly-A tail) can be added to a polynucleotide such as an mRNA molecule to increase stability. Immediately after transcription, the 3' end of the transcript can be cleaved to free a 3' hydroxyl. Then poly-A polymerase adds a chain of adenine nucleotides to the RNA. The process, called polyadenylation, adds a poly-A tail that can be between, for example, approximately 80 to approximately 250 residues long, including approximately 80, 90, 100, 110, 120, 130, 140, 150, 160, 170, 180, 190, 200, 210, 220, 230, 240 or 250 residues long. In one embodiment, the poly-A tail is 100 nucleotides in length (SEQ ID NO:223).

[0379] PolyA tails can also be added after the construct is exported from the nucleus.

[0380] According to the present invention, terminal groups on the poly A tail can be incorporated for stabilization. Polynucleotides of the present invention can include des-3' hydroxyl tails. They can also include structural moieties or 2'-Omethyl modifications as taught by Junjie Li, et al. (Current Biology, Vol. 15, 1501–1507, August 23, 2005, the contents of which are incorporated herein by reference in its entirety).

[0381] The polynucleotides of the present invention can be designed to encode transcripts with alternative polyA tail structures including histone mRNA. According to Norbury, "Terminal uridylation has also been detected on human replication-dependent histone mRNAs. The turnover of these mRNAs is thought to be important for the prevention of potentially toxic histone accumulation following the completion or inhibition of chromosomal DNA replication. These mRNAs are distinguished by their lack of a 3′ poly(A) tail, the function of which is instead assumed by a stable stem–loop structure and its cognate stem–loop binding protein (SLBP); the latter

carries out the same functions as those of PABP on polyadenylated mRNAs" (Norbury, "Cytoplasmic RNA: a case of the tail wagging the dog," Nature Reviews Molecular Cell Biology; AOP, published online 29 August 2013; doi:10.1038/nrm3645) the contents of which are incorporated herein by reference in its entirety.

[0382] Unique poly-A tail lengths provide certain advantages to the polynucleotides of the present invention. Generally, the length of a poly-A tail, when present, is greater than 30 nucleotides in length. In another embodiment, the poly-A tail is greater than 35 nucleotides in length (*e.g.*, at least or greater than about 35, 40, 45, 50, 55, 60, 70, 80, 90, 100, 120, 140, 160, 180, 200, 250, 300, 350, 400, 450, 500, 600, 700, 800, 900, 1,000, 1,100, 1,200, 1,300, 1,400, 1,500, 1,600, 1,700, 1,800, 1,900, 2,000, 2,500, and 3,000 nucleotides).

[0383] In some embodiments, the polynucleotide or region thereof includes from about 30 to about 3,000 nucleotides (*e.g.*, from 30 to 50, from 30 to 100, from 30 to 250, from 30 to 500, from 30 to 750, from 30 to 1,000, from 30 to 1,500, from 30 to 2,000, from 30 to 2,500, from 50 to 100, from 50 to 250, from 50 to 500, from 50 to 750, from 50 to 1,000, from 50 to 1,500, from 50 to 2,000, from 50 to 2,500, from 50 to 3,000, from 100 to 500, from 100 to 750, from 100 to 1,000, from 100 to 1,500, from 100 to 2,000, from 100 to 2,500, from 100 to 3,000, from 500 to 750, from 500 to 1,000, from 500 to 1,500, from 500 to 2,000, from 500 to 2,500, from 500 to 3,000, from 1,000 to 1,500, from 1,000 to 2,000, from 1,000 to 2,500, from 1,000 to 3,000, from 1,500 to 2,000, from 1,500 to 2,500, from 1,500 to 3,000, from 2,000 to 3,000, from 2,000 to 2,500, and from 2,500 to 3,000).

[0384] In some embodiments, the poly-A tail is designed relative to the length of the overall polynucleotide or the length of a particular region of the polynucleotide. This design can be based on the length of a coding region, the length of a particular feature or region or based on the length of the ultimate product expressed from the polynucleotides.

[0385] In this context, the poly-A tail can be 10, 20, 30, 40, 50, 60, 70, 80, 90, or 100% greater in length than the polynucleotide or feature thereof. The poly-A tail can also be designed as a fraction of the polynucleotides to which it belongs. In this context, the poly-A tail can be 10, 20, 30, 40, 50, 60, 70, 80, or 90% or more of the total length of the construct, a construct region or the total length of the construct

minus the poly-A tail. Further, engineered binding sites and conjugation of polynucleotides for Poly-A binding protein can enhance expression.

[0386] Additionally, multiple distinct polynucleotides can be linked together via the PABP (Poly-A binding protein) through the 3′-end using modified nucleotides at the 3′-terminus of the poly-A tail. Transfection experiments can be conducted in relevant cell lines at and protein production can be assayed by ELISA at 12hr, 24hr, 48hr, 72hr and day 7 post-transfection.

[0387] In some embodiments, the polynucleotides of the present invention are designed to include a polyA-G Quartet region. The G-quartet is a cyclic hydrogen bonded array of four guanine nucleotides that can be formed by G-rich sequences in both DNA and RNA. In this embodiment, the G-quartet is incorporated at the end of the poly-A tail. The resultant polynucleotide is assayed for stability, protein production and other parameters including half-life at various time points. It has been discovered that the polyA-G quartet results in protein production from an mRNA equivalent to at least 75% of that seen using a poly-A tail of 120 nucleotides alone (SEQ ID NO:233).

15. Start codon region

[0388] The invention also includes a polynucleotide that comprises both a start codon region and the polynucleotide described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide). In some embodiments, the polynucleotides of the present invention can have regions that are analogous to or function like a start codon region.

[0389] In some embodiments, the translation of a polynucleotide can initiate on a codon that is not the start codon AUG. Translation of the polynucleotide can initiate on an alternative start codon such as, but not limited to, ACG, AGG, AAG, CTG/CUG, GTG/GUG, ATA/AUA, ATT/AUU, TTG/UUG (see Touriol et al. Biology of the Cell 95 (2003) 169-178 and Matsuda and Mauro PLoS ONE, 2010 5:11; the contents of each of which are herein incorporated by reference in its entirety).

[0390] As a non-limiting example, the translation of a polynucleotide begins on the alternative start codon ACG. As another non-limiting example, polynucleotide translation begins on the alternative start codon CTG or CUG. As another non-

limiting example, the translation of a polynucleotide begins on the alternative start codon GTG or GUG.

[0391] Nucleotides flanking a codon that initiates translation such as, but not limited to, a start codon or an alternative start codon, are known to affect the translation efficiency, the length and/or the structure of the polynucleotide. (*See, e.g.,* Matsuda and Mauro PLoS ONE, 2010 5:11; the contents of which are herein incorporated by reference in its entirety). Masking any of the nucleotides flanking a codon that initiates translation can be used to alter the position of translation initiation, translation efficiency, length and/or structure of a polynucleotide.

[0392] In some embodiments, a masking agent can be used near the start codon or alternative start codon to mask or hide the codon to reduce the probability of translation initiation at the masked start codon or alternative start codon. Non-limiting examples of masking agents include antisense locked nucleic acids (LNA) polynucleotides and exon-junction complexes (EJCs) (*See, e.g.,* Matsuda and Mauro describing masking agents LNA polynucleotides and EJCs (PLoS ONE, 2010 5:11); the contents of which are herein incorporated by reference in its entirety).

[0393] In another embodiment, a masking agent can be used to mask a start codon of a polynucleotide to increase the likelihood that translation will initiate on an alternative start codon. In some embodiments, a masking agent can be used to mask a first start codon or alternative start codon to increase the chance that translation will initiate on a start codon or alternative start codon downstream to the masked start codon or alternative start codon.

[0394] In some embodiments, a start codon or alternative start codon can be located within a perfect complement for a miRNA binding site. The perfect complement of a miRNA binding site can help control the translation, length and/or structure of the polynucleotide similar to a masking agent. As a non-limiting example, the start codon or alternative start codon can be located in the middle of a perfect complement for a miRNA binding site. The start codon or alternative start codon can be located after the first nucleotide, second nucleotide, third nucleotide, fourth nucleotide, fifth nucleotide, sixth nucleotide, seventh nucleotide, eighth nucleotide, ninth nucleotide, tenth nucleotide, eleventh nucleotide, twelfth nucleotide, thirteenth nucleotide, fourteenth nucleotide, fifteenth nucleotide, sixteenth nucleotide, seventeenth nucleotide, eighteenth nucleotide, nineteenth nucleotide, twentieth nucleotide or twenty-first nucleotide.

[0395] In another embodiment, the start codon of a polynucleotide can be removed from the polynucleotide sequence to have the translation of the polynucleotide begin on a codon that is not the start codon. Translation of the polynucleotide can begin on the codon following the removed start codon or on a downstream start codon or an alternative start codon. In a non-limiting example, the start codon ATG or AUG is removed as the first 3 nucleotides of the polynucleotide sequence to have translation initiate on a downstream start codon or alternative start codon. The polynucleotide sequence where the start codon was removed can further comprise at least one masking agent for the downstream start codon and/or alternative start codons to control or attempt to control the initiation of translation, the length of the polynucleotide and/or the structure of the polynucleotide.

16. Stop Codon Region

[0396] The invention also includes a polynucleotide that comprises both a stop codon region and the polynucleotide described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide). In some embodiments, the polynucleotides of the present invention can include at least two stop codons before the 3' untranslated region (UTR). The stop codon can be selected from TGA, TAA and TAG in the case of DNA, or from UGA, UAA and UAG in the case of RNA. In some embodiments, the polynucleotides of the present invention include the stop codon TGA in the case or DNA, or the stop codon UGA in the case of RNA, and one additional stop codon. In a further embodiment the addition stop codon can be TAA or UAA. In another embodiment, the polynucleotides of the present invention include three consecutive stop codons, four stop codons, or more.

17. Polynucleotide Comprising an mRNA Encoding a G6PC Polypeptide

[0397] In certain embodiments, a polynucleotide of the present disclosure, for example a polynucleotide comprising an mRNA nucleotide sequence encoding a G6PC polypeptide, comprises from 5’ to 3’ end:

(i) a 5' cap provided above;

(ii) a 5' UTR, such as the sequences provided above;

(iii) an open reading frame encoding a G6PC polypeptide, e.g., a sequence optimized nucleic acid sequence encoding a G6PC disclosed herein;

(iv) at least one stop codon;

(v) a 3' UTR, such as the sequences provided above; and

(vi) a poly-A tail provided above.

[0398] In some embodiments, the polynucleotide further comprises a miRNA binding site, e.g., a miRNA binding site that binds to miRNA-142. In some embodiments, the 5' UTR comprises the miRNA binding site. In some embodiments, the 3' UTR comprises the miRNA binding site.

[0399] In some embodiments, a polynucleotide of the present disclosure comprises a nucleotide sequence encoding a polypeptide sequence at least 70%, at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96% , at least 97%, at least 98%, at least 99%, or 100% identical to the protein sequence of a wild type human G6PC (SEQ ID NO:199) or a variant thereof (e.g., SEQ ID NOs:194-198, 200-206, 208, and 209).

[0400] In some embodiments, a polynucleotide of the present disclosure, for example a polynucleotide comprising an mRNA nucleotide sequence encoding a polypeptide, comprises (1) a 5' cap provided above, for example, CAP1, (2) a 5' UTR, (3) a nucleotide sequence ORF selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207, (3) a stop codon, (4) a 3'UTR, and (5) a poly-A tail provided above, for example, a poly-A tail of about 100 residues.

[0401] Exemplary G6PC nucleotide constructs are described below:

[0402] SEQ ID NO: 45 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 2, and 3' UTR of SEQ ID NO: 150.

[0403] SEQ ID NO: 46 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 5, and 3' UTR of SEQ ID NO: 150.

[0404] SEQ ID NO: 47 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 6, and 3' UTR of SEQ ID NO: 150.

[0405] SEQ ID NO: 48 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 7, and 3' UTR of SEQ ID NO: 150.

[0406] SEQ ID NO: 49 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 7, and 3' UTR of SEQ ID NO: 150.

[0407] SEQ ID NO: 50 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 9, and 3' UTR of SEQ ID NO: 150.

[0408] SEQ ID NO: 51 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 10, and 3' UTR of SEQ ID NO: 150.

[0409] SEQ ID NO: 52 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 7, and 3' UTR of SEQ ID NO: 175.

[0410] SEQ ID NO: 53 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 12, and 3' UTR of SEQ ID NO: 175.

[0411] SEQ ID NO: 54 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 13, and 3' UTR of SEQ ID NO: 175.

[0412] SEQ ID NO: 55 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 14, and 3' UTR of SEQ ID NO: 175.

[0413] SEQ ID NO: 56 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 15, and 3' UTR of SEQ ID NO: 175.

[0414] SEQ ID NO: 57 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 16, and 3' UTR of SEQ ID NO: 175.

[0415] SEQ ID NO: 58 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 17, and 3' UTR of SEQ ID NO: 175.

[0416] SEQ ID NO: 59 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 18, and 3' UTR of SEQ ID NO: 175.

[0417] SEQ ID NO: 60 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 19, and 3' UTR of SEQ ID NO: 175.

[0418] SEQ ID NO: 61 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 20, and 3' UTR of SEQ ID NO: 175.

[0419] SEQ ID NO: 62 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 11, and 3' UTR of SEQ ID NO: 175.

[0420] SEQ ID NO: 63 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 8, and 3' UTR of SEQ ID NO: 175.

[0421] SEQ ID NO: 64 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 207, and 3' UTR of SEQ ID NO: 175.

[0422] SEQ ID NO: 65 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 22, and 3' UTR of SEQ ID NO: 175.

[0423] SEQ ID NO: 66 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 23, and 3' UTR of SEQ ID NO: 175.

[0424] SEQ ID NO: 67 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 24, and 3' UTR of SEQ ID NO: 175.

[0425] SEQ ID NO: 68 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 25, and 3' UTR of SEQ ID NO: 175.

[0426] SEQ ID NO: 69 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 26, and 3' UTR of SEQ ID NO: 175.

[0427] SEQ ID NO: 70 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 27, and 3' UTR of SEQ ID NO: 175.

[0428] SEQ ID NO: 71 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 28, and 3' UTR of SEQ ID NO: 175.

[0429] SEQ ID NO: 72 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 29, and 3' UTR of SEQ ID NO: 150.

[0430] SEQ ID NO: 73 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 30, and 3' UTR of SEQ ID NO: 150.

[0431] SEQ ID NO: 74 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 31, and 3' UTR of SEQ ID NO: 150.

[0432] SEQ ID NO: 75 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 32, and 3' UTR of SEQ ID NO: 150.

[0433] SEQ ID NO: 76 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 33, and 3' UTR of SEQ ID NO: 150.

[0434] SEQ ID NO: 77 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 34, and 3' UTR of SEQ ID NO: 150.

[0435] SEQ ID NO: 78 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 35, and 3' UTR of SEQ ID NO: 150.

[0436] SEQ ID NO: 79 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 36, and 3' UTR of SEQ ID NO: 150.

[0437] SEQ ID NO: 80 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 37, and 3' UTR of SEQ ID NO: 150.

[0438] SEQ ID NO: 81 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 38, and 3' UTR of SEQ ID NO: 150.

[0439] SEQ ID NO: 214 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 210, G6PC nucleotide ORF of SEQ ID NO: 23, and 3' UTR of SEQ ID NO: 175.

[0440] SEQ ID NO: 215 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 210, G6PC nucleotide ORF of SEQ ID NO: 23, and 3' UTR of SEQ ID NO: 178.

[0441] SEQ ID NO: 217 consists from 5' to 3' end: 5' UTR of SEQ ID NO: 3, G6PC nucleotide ORF of SEQ ID NO: 23, and 3' UTR of SEQ ID NO: 178.

[0442] In certain embodiments, in constructs with SEQ ID NO:53-81, 214, 215, or 217, all uracils therein are replaced by N1-methylpseudouracil. In certain

embodiments, in constructs with SEQ ID NO:53-81, 214, 215, or 217, all uracils therein are replaced by 5-methoxyuracil.

[0443] In some embodiments, a polynucleotide of the present disclosure, for example a polynucleotide comprising an mRNA nucleotide sequence encoding a G6PC polypeptide, comprises (1) a 5' cap provided above, for example, CAP1, (2) a nucleotide sequence selected from the group consisting of SEQ ID NO:53-81, 214, 215, and 217, and (3) a poly-A tail provided above, for example, a poly A tail of ~100 residues. In certain embodiments, in constructs with SEQ ID NO:53-81, 214, 215, or 217, all uracils therein are replaced by N1-methylpseudouracil. In certain embodiments, in constructs with SEQ ID NO:53-81, 214, 215, or 217, all uracils therein are replaced by 5-methoxyuracil.

TABLE 5 – Modified mRNA constructs including ORFs encoding human G6PC or a variant thereof (each of constructs #1 to #39 comprises a Cap1 5' terminal cap and a 3' terminal PolyA region)

G6PC mRNA construct	5'UTR SEQ ID NO	G6PC ORF		3' UTR SEQ ID NO:	Protein SEQ ID NO
		Name (Chemistry)	SEQ ID NO		
#1 (SEQ ID NO: 45)	3	hG6PC-FLAG (G5)	2	150	1
#2 (SEQ ID NO: 46)	3	mG6PC (G5)	5	150	21
#3 (SEQ ID NO: 47)	3	mG6PC-FLAG (G5)	6	150	82
#4 (SEQ ID NO: 48)	3	hG6PC (G5)	7	150	199
#5 (SEQ ID NO: 49)	3	hG6PC WT1 (G5)	7	150	199
#6 (SEQ ID NO: 50)	3	hG6PC V1 (G5)	9	150	83
#7 (SEQ ID NO: 51)	3	HG6PC V2 (G5)	10	150	84
#8 (SEQ ID NO: 52)	3	hG6PC WT2 (G5)	7	175	199
#9 (SEQ ID NO: 53)	3	hG6PC Q139R (G5)	12	175	195
#10 (SEQ ID NO: 54)	3	hG6PC I142K (G5)	13	175	197
#11 (SEQ ID NO: 55)	3	hG6PC Q247R (G6)	14	175	196

#12 (SEQ ID NO: 56)	3	hG6PC S297G (G5)	15	175	198
#13 (SEQ ID NO: 57)	3	hG6PC S298C (G5)	16	175	194
#14 (SEQ ID NO: 58)	3	hG6PC V308L (G5)	17	175	200
#15 (SEQ ID NO: 59)	3	hG6PC V321I (G5)	18	175	201
#16 (SEQ ID NO: 60)	3	hG6PC V321L (G5)	19	175	202
#17 (SEQ ID NO: 61)	3	hG6PC V340L(G5)	20	175	203
#18 (SEQ ID NO: 62)	3	hG6PC V332A (G5)	11	175	204
#19 (SEQ ID NO: 63)	3	hG6PCQ139R/Q247R/S298C (G5)	8	175	205
#20 (SEQ ID NO: 64)	3	hG6PC Q247R/S298C (G5)	207	175	206
#21(SEQ ID NO: 65)	3	hG6PC S298A (G5)	22	175	208
#22 (SEQ ID NO: 66)	3	hG6PC S298C_CoV1 (G5)	23	175	194
#23(SEQ ID NO: 67)	3	hG6PC Q139R/Q247R/S298C_CoV1 (G5)	24	175	205
#24 (SEQ ID NO: 68)	3	hG6PC M8-CoV1 (G5)	25	175	209
#25 (SEQ ID NO: 69)	3	hG6PC S298C_CoV2 (G5)	26	175	194
#26 (SEQ ID NO: 70)	3	hG6PC Q139R/Q247R/S298C_CoV2 (G5)	27	175	205
#27 (SEQ ID NO: 71)	3	hG6PC M8-CoV2 (G5)	28	175	209
#28 (SEQ ID NO: 72)	3	hG6PC V1_Co-1 (G5)	29	150	83
#29 (SEQ ID NO: 73)	3	hG6PC V1_Co-2 (G5)	30	150	83
#30 (SEQ ID NO: 74)	3	hG6PC V1-Co-3 (G5)	31	150	83
#31 (SEQ ID NO: 75)	3	hG6PC V1_Co-4 (G5)	32	150	83
#32 (SEQ ID NO: 76)	3	hG6PC V1_Co-5 (G5)	33	150	83
#33 (SEQ ID NO: 77)	3	hG6PC V2_Co-1 (G5)	34	150	84
#34 (SEQ ID NO: 78)	3	hG6PC V2_Co-2 (G5)	35	150	84

#35 (SEQ ID NO: 79)	3	hG6Pc V2_Co-3 (G5)	36	150	84
#36 (SEQ ID NO: 80)	3	hG6PC V2_Co-4 (G5)	37	150	84
#37 (SEQ ID NO: 81)	3	hG6PC V2-Co-5 (G5)	38	150	84
#38 (SEQ ID NO: 214)	210	G6PC_S298C_Astart (G5)	23	175	194
#39 (SEQ ID NO: 215)	210	G6PC_S298C_Astart_3X miR142 (G5)	23	178	194
#40 (SEQ ID NO: 217)	3	G6PC_S298C_Gstart_3X miR142	23	178	194

18. Methods of Making Polynucleotides

[0444] The present disclosure also provides methods for making a polynucleotide of the invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) or a complement thereof.

[0445] In some aspects, a polynucleotide (e.g., a RNA, e.g., an mRNA) disclosed herein, and encoding a G6PC polypeptide, can be constructed using *in vitro* transcription (IVT). In other aspects, a polynucleotide (e.g., a RNA, e.g., an mRNA) disclosed herein, and encoding a G6PC polypeptide, can be constructed by chemical synthesis using an oligonucleotide synthesizer.

[0446] In other aspects, a polynucleotide (e.g., a RNA, e.g., an mRNA) disclosed herein, and encoding a G6PC polypeptide is made by using a host cell. In certain aspects, a polynucleotide (e.g., a RNA, e.g., an mRNA) disclosed herein, and encoding a G6PC polypeptide is made by one or more combination of the IVT, chemical synthesis, host cell expression, or any other methods known in the art.

[0447] Naturally occurring nucleosides, non-naturally occurring nucleosides, or combinations thereof, can totally or partially naturally replace occurring nucleosides present in the candidate nucleotide sequence and can be incorporated into a sequence-optimized nucleotide sequence (e.g., a RNA, e.g., an mRNA) encoding a G6PC polypeptide. The resultant polynucleotides, e.g., mRNAs, can then be examined for their ability to produce protein and/or produce a therapeutic outcome.

a. In Vitro Transcription / Enzymatic Synthesis

[0448] The polynucleotides of the present invention disclosed herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) can

be transcribed using an *in vitro* transcription (IVT) system. The system typically comprises a transcription buffer, nucleotide triphosphates (NTPs), an RNase inhibitor and a polymerase. The NTPs can be selected from, but are not limited to, those described herein including natural and unnatural (modified) NTPs. The polymerase can be selected from, but is not limited to, T7 RNA polymerase, T3 RNA polymerase and mutant polymerases such as, but not limited to, polymerases able to incorporate polynucleotides disclosed herein. See U.S. Publ. No. US20130259923, which is herein incorporated by reference in its entirety.

[0449] Any number of RNA polymerases or variants can be used in the synthesis of the polynucleotides of the present invention. RNA polymerases can be modified by inserting or deleting amino acids of the RNA polymerase sequence. As a non-limiting example, the RNA polymerase can be modified to exhibit an increased ability to incorporate a 2′-modified nucleotide triphosphate compared to an unmodified RNA polymerase (see International Publication WO2008078180 and U.S. Patent 8,101,385; herein incorporated by reference in their entireties).

[0450] Variants can be obtained by evolving an RNA polymerase, optimizing the RNA polymerase amino acid and/or nucleic acid sequence and/or by using other methods known in the art. As a non-limiting example, T7 RNA polymerase variants can be evolved using the continuous directed evolution system set out by Esvelt et al. (Nature 472:499-503 (2011); herein incorporated by reference in its entirety) where clones of T7 RNA polymerase can encode at least one mutation such as, but not limited to, lysine at position 93 substituted for threonine (K93T), I4M, A7T, E63V, V64D, A65E, D66Y, T76N, C125R, S128R, A136T, N165S, G175R, H176L, Y178H, F182L, L196F, G198V, D208Y, E222K, S228A, Q239R, T243N, G259D, M267I, G280C, H300R, D351A, A354S, E356D, L360P, A383V, Y385C, D388Y, S397R, M401T, N410S, K450R, P451T, G452V, E484A, H523L, H524N, G542V, E565K, K577E, K577M, N601S, S684Y, L699I, K713E, N748D, Q754R, E775K, A827V, D851N or L864F. As another non-limiting example, T7 RNA polymerase variants can encode at least mutation as described in U.S. Pub. Nos. 20100120024 and 20070117112; herein incorporated by reference in their entireties. Variants of RNA polymerase can also include, but are not limited to, substitutional variants, conservative amino acid substitution, insertional variants, and/or deletional variants.

[0451] In one aspect, the polynucleotide can be designed to be recognized by the wild type or variant RNA polymerases. In doing so, the polynucleotide can be modified to

contain sites or regions of sequence changes from the wild type or parent chimeric polynucleotide.

[0452] Polynucleotide or nucleic acid synthesis reactions can be carried out by enzymatic methods utilizing polymerases. Polymerases catalyze the creation of phosphodiester bonds between nucleotides in a polynucleotide or nucleic acid chain. Currently known DNA polymerases can be divided into different families based on amino acid sequence comparison and crystal structure analysis. DNA polymerase I (pol I) or A polymerase family, including the Klenow fragments of *E. coli*, *Bacillus* DNA polymerase I, *Thermus aquaticus* (Taq) DNA polymerases, and the T7 RNA and DNA polymerases, is among the best studied of these families. Another large family is DNA polymerase α (pol α) or B polymerase family, including all eukaryotic replicating DNA polymerases and polymerases from phages T4 and RB69. Although they employ similar catalytic mechanism, these families of polymerases differ in substrate specificity, substrate analog-incorporating efficiency, degree and rate for primer extension, mode of DNA synthesis, exonuclease activity, and sensitivity against inhibitors.

[0453] DNA polymerases are also selected based on the optimum reaction conditions they require, such as reaction temperature, pH, and template and primer concentrations. Sometimes a combination of more than one DNA polymerases is employed to achieve the desired DNA fragment size and synthesis efficiency. For example, Cheng et al. increase pH, add glycerol and dimethyl sulfoxide, decrease denaturation times, increase extension times, and utilize a secondary thermostable DNA polymerase that possesses a 3′ to 5′ exonuclease activity to effectively amplify long targets from cloned inserts and human genomic DNA. (Cheng et al., PNAS 91:5695-5699 (1994), the contents of which are incorporated herein by reference in their entirety). RNA polymerases from bacteriophage T3, T7, and SP6 have been widely used to prepare RNAs for biochemical and biophysical studies. RNA polymerases, capping enzymes, and poly-A polymerases are disclosed in the co-pending International Publication No. WO2014/028429, the contents of which are incorporated herein by reference in their entirety.

[0454] In one aspect, the RNA polymerase which can be used in the synthesis of the polynucleotides of the present invention is a Syn5 RNA polymerase. (see Zhu et al. Nucleic Acids Research 2013, doi:10.1093/nar/gkt1193, which is herein incorporated by reference in its entirety). The Syn5 RNA polymerase was recently characterized

from marine cyanophage Syn5 by Zhu et al. where they also identified the promoter sequence (see Zhu et al. Nucleic Acids Research 2013, the contents of which is herein incorporated by reference in its entirety). Zhu et al. found that Syn5 RNA polymerase catalyzed RNA synthesis over a wider range of temperatures and salinity as compared to T7 RNA polymerase. Additionally, the requirement for the initiating nucleotide at the promoter was found to be less stringent for Syn5 RNA polymerase as compared to the T7 RNA polymerase making Syn5 RNA polymerase promising for RNA synthesis.

[0455] In one aspect, a Syn5 RNA polymerase can be used in the synthesis of the polynucleotides described herein. As a non-limiting example, a Syn5 RNA polymerase can be used in the synthesis of the polynucleotide requiring a precise 3′-terminus.

[0456] In one aspect, a Syn5 promoter can be used in the synthesis of the polynucleotides. As a non-limiting example, the Syn5 promoter can be 5′-ATTGGGCACCCGTAAGGG-3′ (SEQ ID NO: 185 as described by Zhu et al. (Nucleic Acids Research 2013).

[0457] In one aspect, a Syn5 RNA polymerase can be used in the synthesis of polynucleotides comprising at least one chemical modification described herein and/or known in the art (see e.g., the incorporation of pseudo-UTP and 5Me-CTP described in Zhu et al. Nucleic Acids Research 2013).

[0458] In one aspect, the polynucleotides described herein can be synthesized using a Syn5 RNA polymerase which has been purified using modified and improved purification procedure described by Zhu et al. (Nucleic Acids Research 2013).

[0459] Various tools in genetic engineering are based on the enzymatic amplification of a target gene which acts as a template. For the study of sequences of individual genes or specific regions of interest and other research needs, it is necessary to generate multiple copies of a target gene from a small sample of polynucleotides or nucleic acids. Such methods can be applied in the manufacture of the polynucleotides of the invention.

For example, polymerase chain reaction (PCR), strand displacement amplification (SDA), nucleic acid sequence-based amplification (NASBA), also called transcription mediated amplification (TMA), and/or rolling-circle amplification (RCA) can be utilized in the manufacture of one or more regions of the polynucleotides of the present invention. Assembling polynucleotides or nucleic acids by a ligase is also widely used.

b. Chemical synthesis

[0460] Standard methods can be applied to synthesize an isolated polynucleotide sequence encoding an isolated polypeptide of interest, such as a polynucleotide of the invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide). For example, a single DNA or RNA oligomer containing a codon-optimized nucleotide sequence coding for the particular isolated polypeptide can be synthesized. In other aspects, several small oligonucleotides coding for portions of the desired polypeptide can be synthesized and then ligated. In some aspects, the individual oligonucleotides typically contain 5' or 3' overhangs for complementary assembly.

[0461] A polynucleotide disclosed herein (e.g., a RNA, e.g., an mRNA) can be chemically synthesized using chemical synthesis methods and potential nucleobase substitutions known in the art. See, for example, International Publication Nos. WO2014093924, WO2013052523; WO2013039857, WO2012135805, WO2013151671; U.S. Publ. No. US20130115272; or U.S. Pat. Nos. US8999380 or US8710200, all of which are herein incorporated by reference in their entireties.

c. Purification of Polynucleotides Encoding G6PC

[0462] Purification of the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) can include, but is not limited to, polynucleotide clean-up, quality assurance and quality control. Clean-up can be performed by methods known in the arts such as, but not limited to, AGENCOURT® beads (Beckman Coulter Genomics, Danvers, MA), poly-T beads, LNA™ oligo-T capture probes (EXIQON® Inc., Vedbaek, Denmark) or HPLC based purification methods such as, but not limited to, strong anion exchange HPLC, weak anion exchange HPLC, reverse phase HPLC (RP-HPLC), and hydrophobic interaction HPLC (HIC-HPLC).

[0463] The term "purified" when used in relation to a polynucleotide such as a "purified polynucleotide" refers to one that is separated from at least one contaminant. As used herein, a "contaminant" is any substance that makes another unfit, impure or inferior. Thus, a purified polynucleotide (*e.g.*, DNA and RNA) is present in a form or setting different from that in which it is found in nature, or a form or setting different from that which existed prior to subjecting it to a treatment or purification method.

[0464] In some embodiments, purification of a polynucleotide of the invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) removes impurities that can reduce or remove an unwanted immune response, *e.g.*, reducing cytokine activity.

[0465] In some embodiments, the polynucleotide of the invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) is purified prior to administration using column chromatography (*e.g.*, strong anion exchange HPLC, weak anion exchange HPLC, reverse phase HPLC (RP-HPLC), and hydrophobic interaction HPLC (HIC-HPLC), or (LCMS)).

[0466] In some embodiments, the polynucleotide of the invention (e.g., a polynucleotide comprising a nucleotide sequence a G6PC polypeptide) purified using column chromatography (*e.g.*, strong anion exchange HPLC, weak anion exchange HPLC, reverse phase HPLC (RP-HPLC, hydrophobic interaction HPLC (HIC-HPLC), or (LCMS)) presents increased expression of the encoded G6PC protein compared to the expression level obtained with the same polynucleotide of the present disclosure purified by a different purification method.

[0467] In some embodiments, a column chromatography (*e.g.*, strong anion exchange HPLC, weak anion exchange HPLC, reverse phase HPLC (RP-HPLC), hydrophobic interaction HPLC (HIC-HPLC), or (LCMS)) purified polynucleotide comprises a nucleotide sequence encoding a G6PC polypeptide comprising one or more of the point mutations known in the art.

[0468] In some embodiments, the use of RP-HPLC purified polynucleotide increases G6PC protein expression levels in cells when introduced into those cells, *e.g.*, by 10-100%, i.e., at least about 10%, at least about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 90%, at least about 95%, or at least about 100% with respect to the expression levels of G6PC protein in the cells before the RP-HPLC purified polynucleotide was introduced in the cells, or after a non-RP-HPLC purified polynucleotide was introduced in the cells.

[0469] In some embodiments, the use of RP-HPLC purified polynucleotide increases functional G6PC protein expression levels in cells when introduced into those cells, *e.g.*, by 10-100%, i.e., at least about 10%, at least about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least

about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 90%, at least about 95%, or at least about 100% with respect to the functional expression levels of G6PC protein in the cells before the RP-HPLC purified polynucleotide was introduced in the cells, or after a non-RP-HPLC purified polynucleotide was introduced in the cells.

[0470] In some embodiments, the use of RP-HPLC purified polynucleotide increases detectable G6PC activity in cells when introduced into those cells, *e.g.,* by 10-100%, i.e., at least about 10%, at least about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 90%, at least about 95%, or at least about 100% with respect to the activity levels of functional G6PC in the cells before the RP-HPLC purified polynucleotide was introduced in the cells, or after a non-RP-HPLC purified polynucleotide was introduced in the cells.

[0471] In some embodiments, the purified polynucleotide is at least about 80% pure, at least about 85% pure, at least about 90% pure, at least about 95% pure, at least about 96% pure, at least about 97% pure, at least about 98% pure, at least about 99% pure, or about 100% pure.

[0472] A quality assurance and/or quality control check can be conducted using methods such as, but not limited to, gel electrophoresis, UV absorbance, or analytical HPLC. In another embodiment, the polynucleotide can be sequenced by methods including, but not limited to reverse-transcriptase-PCR.

d. Quantification of Expressed Polynucleotides Encoding G6PC

[0473] In some embodiments, the polynucleotides of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide), their expression products, as well as degradation products and metabolites can be quantified according to methods known in the art.

[0474] In some embodiments, the polynucleotides of the present invention can be quantified in exosomes or when derived from one or more bodily fluid. As used herein "bodily fluids" include peripheral blood, serum, plasma, ascites, urine, cerebrospinal fluid (CSF), sputum, saliva, bone marrow, synovial fluid, aqueous humor, amniotic fluid, cerumen, breast milk, broncheoalveolar lavage fluid, semen, prostatic fluid, cowper's fluid or pre-ejaculatory fluid, sweat, fecal matter, hair, tears,

cyst fluid, pleural and peritoneal fluid, pericardial fluid, lymph, chyme, chyle, bile, interstitial fluid, menses, pus, sebum, vomit, vaginal secretions, mucosal secretion, stool water, pancreatic juice, lavage fluids from sinus cavities, bronchopulmonary aspirates, blastocyl cavity fluid, and umbilical cord blood. Alternatively, exosomes can be retrieved from an organ selected from the group consisting of lung, heart, pancreas, stomach, intestine, bladder, kidney, ovary, testis, skin, colon, breast, prostate, brain, esophagus, liver, and placenta.

[0475] In the exosome quantification method, a sample of not more than 2mL is obtained from the subject and the exosomes isolated by size exclusion chromatography, density gradient centrifugation, differential centrifugation, nanomembrane ultrafiltration, immunoabsorbent capture, affinity purification, microfluidic separation, or combinations thereof. In the analysis, the level or concentration of a polynucleotide can be an expression level, presence, absence, truncation or alteration of the administered construct. It is advantageous to correlate the level with one or more clinical phenotypes or with an assay for a human disease biomarker.

[0476] The assay can be performed using construct specific probes, cytometry, qRT-PCR, real-time PCR, PCR, flow cytometry, electrophoresis, mass spectrometry, or combinations thereof while the exosomes can be isolated using immunohistochemical methods such as enzyme linked immunosorbent assay (ELISA) methods. Exosomes can also be isolated by size exclusion chromatography, density gradient centrifugation, differential centrifugation, nanomembrane ultrafiltration, immunoabsorbent capture, affinity purification, microfluidic separation, or combinations thereof.

[0477] These methods afford the investigator the ability to monitor, in real time, the level of polynucleotides remaining or delivered. This is possible because the polynucleotides of the present invention differ from the endogenous forms due to the structural or chemical modifications.

[0478] In some embodiments, the polynucleotide can be quantified using methods such as, but not limited to, ultraviolet visible spectroscopy (UV/Vis). A non-limiting example of a UV/Vis spectrometer is a NANODROP® spectrometer (ThermoFisher, Waltham, MA). The quantified polynucleotide can be analyzed to determine if the polynucleotide can be of proper size, check that no degradation of the polynucleotide has occurred. Degradation of the polynucleotide can be checked by methods such as,

but not limited to, agarose gel electrophoresis, HPLC based purification methods such as, but not limited to, strong anion exchange HPLC, weak anion exchange HPLC, reverse phase HPLC (RP-HPLC), and hydrophobic interaction HPLC (HIC-HPLC), liquid chromatography-mass spectrometry (LCMS), capillary electrophoresis (CE) and capillary gel electrophoresis (CGE).

19. Pharmaceutical Compositions and Formulations

[0479] The present invention provides pharmaceutical compositions and formulations that comprise any of the polynucleotides described above. In some embodiments, the composition or formulation further comprises a delivery agent.

[0480] In some embodiments, the composition or formulation can contain a polynucleotide comprising a sequence optimized nucleic acid sequence disclosed herein which encodes a G6PC polypeptide. In some embodiments, the composition or formulation can contain a polynucleotide (e.g., a RNA, e.g., an mRNA) comprising a polynucleotide (e.g., an ORF) having significant sequence identity to a sequence optimized nucleic acid sequence disclosed herein which encodes a G6PC polypeptide. In some embodiments, the polynucleotide further comprises a miRNA binding site, e.g., a miRNA binding site that binds miR-126, miR-142, miR-144, miR-146, miR-150, miR-155, miR-16, miR-21, miR-223, miR-24, miR-27 and miR-26a.

[0481] Pharmaceutical compositions or formulation can optionally comprise one or more additional active substances, *e.g.*, therapeutically and/or prophylactically active substances. Pharmaceutical compositions or formulation of the present invention can be sterile and/or pyrogen-free. General considerations in the formulation and/or manufacture of pharmaceutical agents can be found, for example, in *Remington: The Science and Practice of Pharmacy* 21st ed., Lippincott Williams & Wilkins, 2005 (incorporated herein by reference in its entirety). In some embodiments, compositions are administered to humans, human patients or subjects. For the purposes of the present disclosure, the phrase "active ingredient" generally refers to polynucleotides to be delivered as described herein.

[0482] Formulations and pharmaceutical compositions described herein can be prepared by any method known or hereafter developed in the art of pharmacology. In general, such preparatory methods include the step of associating the active ingredient with an excipient and/or one or more other accessory ingredients, and then, if

necessary and/or desirable, dividing, shaping and/or packaging the product into a desired single- or multi-dose unit.

[0483] A pharmaceutical composition or formulation in accordance with the present disclosure can be prepared, packaged, and/or sold in bulk, as a single unit dose, and/or as a plurality of single unit doses. As used herein, a "unit dose" refers to a discrete amount of the pharmaceutical composition comprising a predetermined amount of the active ingredient. The amount of the active ingredient is generally equal to the dosage of the active ingredient that would be administered to a subject and/or a convenient fraction of such a dosage such as, for example, one-half or one-third of such a dosage.

[0484] Relative amounts of the active ingredient, the pharmaceutically acceptable excipient, and/or any additional ingredients in a pharmaceutical composition in accordance with the present disclosure can vary, depending upon the identity, size, and/or condition of the subject being treated and further depending upon the route by which the composition is to be administered.

[0485] In some embodiments, the compositions and formulations described herein can contain at least one polynucleotide of the invention. As a non-limiting example, the composition or formulation can contain 1, 2, 3, 4 or 5 polynucleotides of the invention. In some embodiments, the compositions or formulations described herein can comprise more than one type of polynucleotide. In some embodiments, the composition or formulation can comprise a polynucleotide in linear and circular form. In another embodiment, the composition or formulation can comprise a circular polynucleotide and an in vitro transcribed (IVT) polynucleotide. In yet another embodiment, the composition or formulation can comprise an IVT polynucleotide, a chimeric polynucleotide and a circular polynucleotide.

[0486] Although the descriptions of pharmaceutical compositions and formulations provided herein are principally directed to pharmaceutical compositions and formulations that are suitable for administration to humans, it will be understood by the skilled artisan that such compositions are generally suitable for administration to any other animal, *e.g.,* to non-human animals, *e.g.* non-human mammals.

[0487] The present invention provides pharmaceutical formulations that comprise a polynucleotide described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide). The polynucleotides described herein can be formulated using one or more excipients to: (1) increase stability; (2) increase cell transfection; (3) permit the sustained or delayed release (*e.g.,* from a depot

formulation of the polynucleotide); (4) alter the biodistribution (*e.g.*, target the polynucleotide to specific tissues or cell types); (5) increase the translation of encoded protein *in vivo*; and/or (6) alter the release profile of encoded protein *in vivo*. In some embodiments, the pharmaceutical formulation further comprises a delivery agent comprising, e.g., a compound having the Formula (I), e.g., any of Compounds 1-232, e.g., Compound II; a compound having the Formula (III), (IV), (V), or (VI), e.g., any of Compounds 233-342, e.g., Compound VI; or a compound having the Formula (VIII), e.g., any of Compounds 419-428, e.g., Compound I, or any combination thereof. In some embodiments, the delivery agent comprises Compound II, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 50:10:38.5:1.5. In some embodiments, the delivery agent comprises Compound II, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0. In some embodiments, the delivery agent comprises Compound VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 50:10:38.5:1.5. In some embodiments, the delivery agent comprises Compound VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0.

[0488] A pharmaceutically acceptable excipient, as used herein, includes, but are not limited to, any and all solvents, dispersion media, or other liquid vehicles, dispersion or suspension aids, diluents, granulating and/or dispersing agents, surface active agents, isotonic agents, thickening or emulsifying agents, preservatives, binders, lubricants or oil, coloring, sweetening or flavoring agents, stabilizers, antioxidants, antimicrobial or antifungal agents, osmolality adjusting agents, pH adjusting agents, buffers, chelants, cytoprotectants, and/or bulking agents, as suited to the particular dosage form desired. Various excipients for formulating pharmaceutical compositions and techniques for preparing the composition are known in the art (see Remington: The Science and Practice of Pharmacy, 21st Edition, A. R. Gennaro (Lippincott, Williams & Wilkins, Baltimore, MD, 2006; incorporated herein by reference in its entirety).

[0489] Exemplary diluents include, but are not limited to, calcium or sodium carbonate, calcium phosphate, calcium hydrogen phosphate, sodium phosphate, lactose, sucrose, cellulose, microcrystalline cellulose, kaolin, mannitol, sorbitol, etc., and/or combinations thereof.

[0490] Exemplary granulating and/or dispersing agents include, but are not limited to, starches, pregelatinized starches, or microcrystalline starch, alginic acid, guar gum, agar, poly(vinyl-pyrrolidone), (providone), cross-linked poly(vinyl-pyrrolidone) (crospovidone), cellulose, methylcellulose, carboxymethyl cellulose, cross-linked sodium carboxymethyl cellulose (croscarmellose), magnesium aluminum silicate (VEEGUM®), sodium lauryl sulfate, etc., and/or combinations thereof.

[0491] Exemplary surface active agents and/or emulsifiers include, but are not limited to, natural emulsifiers (e.g., acacia, agar, alginic acid, sodium alginate, tragacanth, chondrux, cholesterol, xanthan, pectin, gelatin, egg yolk, casein, wool fat, cholesterol, wax, and lecithin), sorbitan fatty acid esters (e.g., polyoxyethylene sorbitan monooleate [TWEEN®80], sorbitan monopalmitate [SPAN®40], glyceryl monooleate, polyoxyethylene esters, polyethylene glycol fatty acid esters (e.g., CREMOPHOR®), polyoxyethylene ethers (e.g., polyoxyethylene lauryl ether [BRIJ®30]), PLUORINC®F 68, POLOXAMER®188, etc. and/or combinations thereof.

[0492] Exemplary binding agents include, but are not limited to, starch, gelatin, sugars (e.g., sucrose, glucose, dextrose, dextrin, molasses, lactose, lactitol, mannitol), amino acids (e.g., glycine), natural and synthetic gums (e.g., acacia, sodium alginate), ethylcellulose, hydroxyethylcellulose, hydroxypropyl methylcellulose, etc., and combinations thereof.

[0493] Oxidation is a potential degradation pathway for mRNA, especially for liquid mRNA formulations. To prevent oxidation, antioxidants can be added to the formulations. Exemplary antioxidants include, but are not limited to, alpha tocopherol, ascorbic acid, ascorbyl palmitate, benzyl alcohol, butylated hydroxyanisole, m-cresol, methionine, butylated hydroxytoluene, monothioglycerol, sodium or potassium metabisulfite, propionic acid, propyl gallate, sodium ascorbate, etc., and combinations thereof.

[0494] Exemplary chelating agents include, but are not limited to, ethylenediaminetetraacetic acid (EDTA), citric acid monohydrate, disodium edetate, fumaric acid, malic acid, phosphoric acid, sodium edetate, tartaric acid, trisodium edetate, etc., and combinations thereof.

[0495] Exemplary antimicrobial or antifungal agents include, but are not limited to, benzalkonium chloride, benzethonium chloride, methyl paraben, ethyl paraben, propyl paraben, butyl paraben, benzoic acid, hydroxybenzoic acid, potassium or sodium

benzoate, potassium or sodium sorbate, sodium propionate, sorbic acid, etc., and combinations thereof.

[0496] Exemplary preservatives include, but are not limited to, vitamin A, vitamin C, vitamin E, beta-carotene, citric acid, ascorbic acid, butylated hydroxyanisol, ethylenediamine, sodium lauryl sulfate (SLS), sodium lauryl ether sulfate (SLES), etc., and combinations thereof.

[0497] In some embodiments, the pH of polynucleotide solutions is maintained between pH 5 and pH 8 to improve stability. Exemplary buffers to control pH can include, but are not limited to sodium phosphate, sodium citrate, sodium succinate, histidine (or histidine-HCl), sodium malate, sodium carbonate, etc., and/or combinations thereof.

[0498] Exemplary lubricating agents include, but are not limited to, magnesium stearate, calcium stearate, stearic acid, silica, talc, malt, hydrogenated vegetable oils, polyethylene glycol, sodium benzoate, sodium or magnesium lauryl sulfate, etc., and combinations thereof.

[0499] The pharmaceutical composition or formulation described here can contain a cryoprotectant to stabilize a polynucleotide described herein during freezing. Exemplary cryoprotectants include, but are not limited to mannitol, sucrose, trehalose, lactose, glycerol, dextrose, etc., and combinations thereof.

[0500] The pharmaceutical composition or formulation described here can contain a bulking agent in lyophilized polynucleotide formulations to yield a "pharmaceutically elegant" cake, stabilize the lyophilized polynucleotides during long term (e.g., 36 month) storage. Exemplary bulking agents of the present invention can include, but are not limited to sucrose, trehalose, mannitol, glycine, lactose, raffinose, and combinations thereof.

[0501] In some embodiments, the pharmaceutical composition or formulation further comprises a delivery agent. The delivery agent of the present disclosure can include, without limitation, liposomes, lipid nanoparticles, lipidoids, polymers, lipoplexes, microvesicles, exosomes, peptides, proteins, cells transfected with polynucleotides, hyaluronidase, nanoparticle mimics, nanotubes, conjugates, and combinations thereof.

20. Delivery Agents

a. Lipid Compound

[0502] The present disclosure provides pharmaceutical compositions with advantageous properties. The lipid compositions described herein may be advantageously used in lipid nanoparticle compositions for the delivery of therapeutic and/or prophylactic agents, e.g., mRNAs, to mammalian cells or organs. For example, the lipids described herein have little or no immunogenicity. For example, the lipid compounds disclosed herein have a lower immunogenicity as compared to a reference lipid (*e.g.*, MC3, KC2, or DLinDMA). For example, a formulation comprising a lipid disclosed herein and a therapeutic or prophylactic agent, e.g., mRNA, has an increased therapeutic index as compared to a corresponding formulation which comprises a reference lipid (*e.g.*, MC3, KC2, or DLinDMA) and the same therapeutic or prophylactic agent.

[0503] In certain embodiments, the present application provides pharmaceutical compositions comprising:

(a) a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide; and

(b) a delivery agent.

Lipid Nanoparticle Formulations

[0504] In some embodiments, nucleic acids of the invention (e.g. G6PC mRNA) are formulated in a lipid nanoparticle (LNP). Lipid nanoparticles typically comprise ionizable cationic lipid, non-cationic lipid, sterol and PEG lipid components along with the nucleic acid cargo of interest. The lipid nanoparticles of the invention can be generated using components, compositions, and methods as are generally known in the art, see for example PCT/US2016/052352; PCT/US2016/068300; PCT/US2017/037551; PCT/US2015/027400; PCT/US2016/047406; PCT/US2016000129; PCT/US2016/014280; PCT/US2016/014280; PCT/US2017/038426; PCT/US2014/027077; PCT/US2014/055394; PCT/US2016/52117; PCT/US2012/069610; PCT/US2017/027492; PCT/US2016/059575 and PCT/US2016/069491 all of which are incorporated by reference herein in their entirety.

[0505] Nucleic acids of the present disclosure (e.g. G6PC mRNA) are typically formulated in lipid nanoparticle. In some embodiments, the lipid nanoparticle comprises at least one ionizable cationic lipid, at least one non-cationic lipid, at least one sterol, and/or at least one polyethylene glycol (PEG)-modified lipid.

[0506] In some embodiments, the lipid nanoparticle comprises a molar ratio of 20-60% ionizable cationic lipid. For example, the lipid nanoparticle may comprise a molar ratio of 20-50%, 20-40%, 20-30%, 30-60%, 30-50%, 30-40%, 40-60%, 40-50%, or 50-60% ionizable cationic lipid. In some embodiments, the lipid nanoparticle comprises a molar ratio of 20%, 30%, 40%, 50, or 60% ionizable cationic lipid.

[0507] In some embodiments, the lipid nanoparticle comprises a molar ratio of 5-25% non-cationic lipid. For example, the lipid nanoparticle may comprise a molar ratio of 5-20%, 5-15%, 5-10%, 10-25%, 10-20%, 10-25%, 15-25%, 15-20%, or 20-25% non-cationic lipid. In some embodiments, the lipid nanoparticle comprises a molar ratio of 5%, 10%, 15%, 20%, or 25% non-cationic lipid.

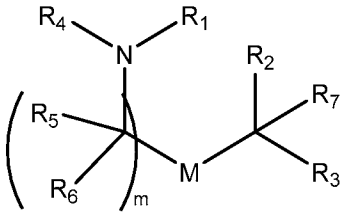
[0508] In some embodiments, the lipid nanoparticle comprises a molar ratio of 25-55% sterol. For example, the lipid nanoparticle may comprise a molar ratio of 25-50%, 25-45%, 25-40%, 25-35%, 25-30%, 30-55%, 30-50%, 30-45%, 30-40%, 30-35%, 35-55%, 35-50%, 35-45%, 35-40%, 40-55%, 40-50%, 40-45%, 45-55%, 45-50%, or 50-55% sterol. In some embodiments, the lipid nanoparticle comprises a molar ratio of 25%, 30%, 35%, 40%, 45%, 50%, or 55% sterol.

[0509] In some embodiments, the lipid nanoparticle comprises a molar ratio of 0.5-15% PEG-modified lipid. For example, the lipid nanoparticle may comprise a molar ratio of 0.5-10%, 0.5-5%, 1-15%, 1-10%, 1-5%, 2-15%, 2-10%, 2-5%, 5-15%, 5-10%, or 10-15%. In some embodiments, the lipid nanoparticle comprises a molar ratio of 0.5%, 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 11%, 12%, 13%, 14%, or 15% PEG-modified lipid.

[0510] In some embodiments, the lipid nanoparticle comprises a molar ratio of 20-60% ionizable cationic lipid, 5-25% non-cationic lipid, 25-55% sterol, and 0.5-15% PEG-modified lipid.

Ionizable Lipids

[0511] In some aspects, the ionizable lipids of the present disclosure may be one or more of compounds of Formula (I):



(I),

or their N-oxides, or salts or isomers thereof, wherein:

R_1 is selected from the group consisting of C_{5-30} alkyl, C_{5-20} alkenyl, -R*YR", -YR", and -R"M'R';

R_2 and R_3 are independently selected from the group consisting of H, C_{1-14} alkyl, C_{2-14} alkenyl, -R*YR", -YR", and -R*OR", or R_2 and R_3, together with the atom to which they are attached, form a heterocycle or carbocycle;

R_4 is selected from the group consisting of hydrogen, a C_{3-6} carbocycle, $-(CH_2)_nQ$, $-(CH_2)_nCHQR$, -CHQR, $-CQ(R)_2$, and unsubstituted C_{1-6} alkyl, where Q is selected from a carbocycle, heterocycle, -OR, $-O(CH_2)_nN(R)_2$, -C(O)OR, -OC(O)R, $-CX_3$, $-CX_2H$, $-CXH_2$, -CN, $-N(R)_2$, $-C(O)N(R)_2$, $-N(R)C(O)R$, $-N(R)S(O)_2R$, $-N(R)C(O)N(R)_2$, $-N(R)C(S)N(R)_2$, $-N(R)R_8$, $-N(R)S(O)_2R_8$, $-O(CH_2)_nOR$, $-N(R)C(=NR_9)N(R)_2$, $-N(R)C(=CHR_9)N(R)_2$, $-OC(O)N(R)_2$, -N(R)C(O)OR, -N(OR)C(O)R, $-N(OR)S(O)_2R$, -N(OR)C(O)OR, $-N(OR)C(O)N(R)_2$, $-N(OR)C(S)N(R)_2$, $-N(OR)C(=NR_9)N(R)_2$, $-N(OR)C(=CHR_9)N(R)_2$, $-C(=NR_9)N(R)_2$, $-C(=NR_9)R$, -C(O)N(R)OR, and $-C(R)N(R)_2C(O)OR$, and each n is independently selected from 1, 2, 3, 4, and 5;

each R_5 is independently selected from the group consisting of C_{1-3} alkyl, C_{2-3} alkenyl, and H;

each R_6 is independently selected from the group consisting of C_{1-3} alkyl, C_{2-3} alkenyl, and H;

M and M' are independently selected from -C(O)O-, -OC(O)-, -OC(O)-M"-C(O)O-, -C(O)N(R')-, -N(R')C(O)-, -C(O)-, -C(S)-, -C(S)S-, -SC(S)-, -CH(OH)-, -P(O)(OR')O-, $-S(O)_2-$, -S-S-, an aryl group, and a heteroaryl group, in which M" is a bond, C_{1-13} alkyl or C_{2-13} alkenyl;

R_7 is selected from the group consisting of C_{1-3} alkyl, C_{2-3} alkenyl, and H;

R_8 is selected from the group consisting of C_{3-6} carbocycle and heterocycle;

R_9 is selected from the group consisting of H, CN, NO_2, C_{1-6} alkyl, -OR, $-S(O)_2R$, $-S(O)_2N(R)_2$, C_{2-6} alkenyl, C_{3-6} carbocycle and heterocycle;

each R is independently selected from the group consisting of C_{1-3} alkyl, C_{2-3} alkenyl, and H;

each R' is independently selected from the group consisting of C_{1-18} alkyl, C_{2-18} alkenyl, -R*YR", -YR", and H;

each R" is independently selected from the group consisting of C_{3-15} alkyl and C_{3-15} alkenyl;

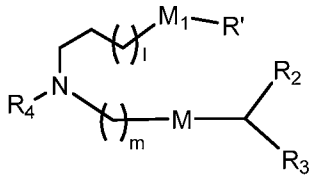
each R* is independently selected from the group consisting of C_{1-12} alkyl and C_{2-12} alkenyl;

each Y is independently a C_{3-6} carbocycle;

each X is independently selected from the group consisting of F, Cl, Br, and I; and

m is selected from 5, 6, 7, 8, 9, 10, 11, 12, and 13; and wherein when R_4 is $-(CH_2)_nQ$, $-(CH_2)_nCHQR$, –CHQR, or $-CQ(R)_2$, then (i) Q is not $-N(R)_2$ when n is 1, 2, 3, 4 or 5, or (ii) Q is not 5, 6, or 7-membered heterocycloalkyl when n is 1 or 2.

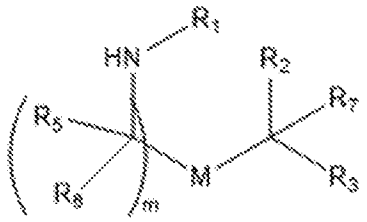
[0512] In certain embodiments, a subset of compounds of Formula (I) includes those of Formula (IA):



(IA),

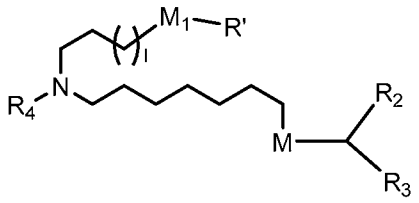
or its N-oxide, or a salt or isomer thereof, wherein l is selected from 1, 2, 3, 4, and 5; m is selected from 5, 6, 7, 8, and 9; M_1 is a bond or M'; R_4 is hydrogen, unsubstituted C_{1-3} alkyl, or $-(CH_2)_nQ$, in which Q is OH, $-NHC(S)N(R)_2$, $-NHC(O)N(R)_2$, $-N(R)C(O)R$, $-N(R)S(O)_2R$, $-N(R)R_8$, $-NHC(=NR_9)N(R)_2$, $-NHC(=CHR_9)N(R)_2$, $-OC(O)N(R)_2$, $-N(R)C(O)OR$, heteroaryl or heterocycloalkyl; M and M' are independently selected from -C(O)O-, -OC(O)-, -OC(O)-M"-C(O)O-, -C(O)N(R')-, -P(O)(OR')O-, -S-S-, an aryl group, and a heteroaryl group,; and R_2 and R_3 are independently selected from the group consisting of H, C_{1-14} alkyl, and C_{2-14} alkenyl. For example, m is 5, 7, or 9. For example, Q is OH, $-NHC(S)N(R)_2$, or $-NHC(O)N(R)_2$. For example, Q is $-N(R)C(O)R$, or $-N(R)S(O)_2R$.

[0513] In certain embodiments, a subset of compounds of Formula (I) includes those of Formula (IB):



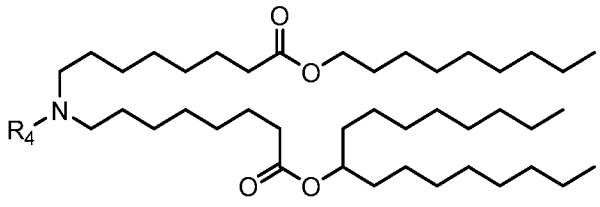
(IB), or its N-oxide, or a salt or isomer thereof in which all variables are as defined herein. For example, m is selected from 5, 6, 7, 8, and 9; R_4 is hydrogen, unsubstituted C_{1-3} alkyl, or $-(CH_2)_nQ$, in which Q is OH, $-NHC(S)N(R)_2$, $-NHC(O)N(R)_2$, $-N(R)C(O)R$, $-N(R)S(O)_2R$, $-N(R)R_8$, $-NHC(=NR_9)N(R)_2$, $-NHC(=CHR_9)N(R)_2$, $-OC(O)N(R)_2$, $-N(R)C(O)OR$, heteroaryl or heterocycloalkyl; M and M' are independently selected from $-C(O)O-$, $-OC(O)-$, $-OC(O)-M''-C(O)O-$, $-C(O)N(R')-$, $-P(O)(OR')O-$, -S-S-, an aryl group, and a heteroaryl group; and R_2 and R_3 are independently selected from the group consisting of H, C_{1-14} alkyl, and C_{2-14} alkenyl. For example, m is 5, 7, or 9. For example, Q is OH, $-NHC(S)N(R)_2$, or $-NHC(O)N(R)_2$. For example, Q is $-N(R)C(O)R$, or $-N(R)S(O)_2R$.

[0514] In certain embodiments, a subset of compounds of Formula (I) includes those of Formula (II):



(II), or its N-oxide, or a salt or isomer thereof, wherein l is selected from 1, 2, 3, 4, and 5; M_1 is a bond or M'; R_4 is hydrogen, unsubstituted C_{1-3} alkyl, or $-(CH_2)_nQ$, in which n is 2, 3, or 4, and Q is OH, $-NHC(S)N(R)_2$, $-NHC(O)N(R)_2$, $-N(R)C(O)R$, $-N(R)S(O)_2R$, $-N(R)R_8$, $-NHC(=NR_9)N(R)_2$, $-NHC(=CHR_9)N(R)_2$, $-OC(O)N(R)_2$, $-N(R)C(O)OR$, heteroaryl or heterocycloalkyl; M and M' are independently selected from $-C(O)O-$, $-OC(O)-$, $-OC(O)-M''-C(O)O-$, $-C(O)N(R')-$, $-P(O)(OR')O-$, -S-S-, an aryl group, and a heteroaryl group; and R_2 and R_3 are independently selected from the group consisting of H, C_{1-14} alkyl, and C_{2-14} alkenyl.

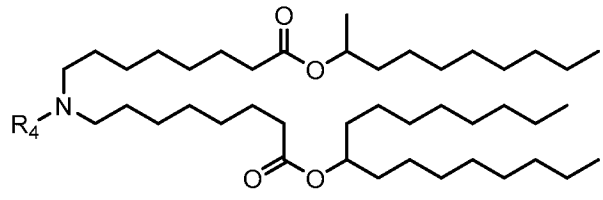
[0515] In one embodiment, the compounds of Formula (I) are of Formula (IIa),



(IIa),

or their N-oxides, or salts or isomers thereof, wherein R_4 is as described herein.

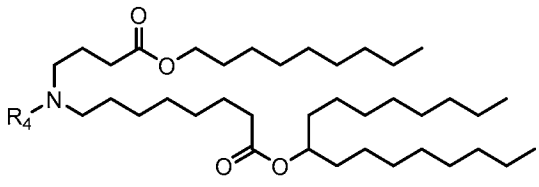
[0516] In another embodiment, the compounds of Formula (I) are of Formula (IIb),



(IIb),

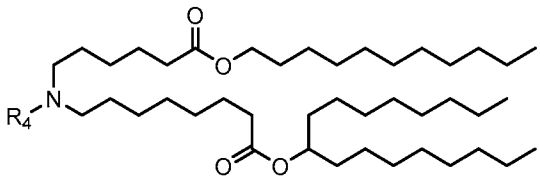
or their N-oxides, or salts or isomers thereof, wherein R_4 is as described herein.

[0517] In another embodiment, the compounds of Formula (I) are of Formula (IIc) or (IIe):



(IIc)

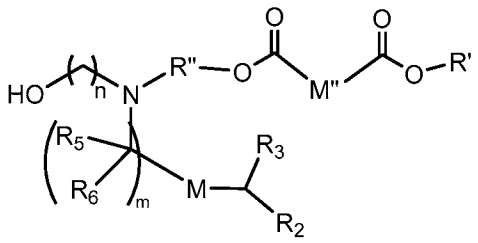
or



(IIe),

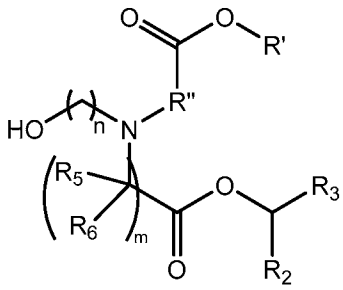
or their N-oxides, or salts or isomers thereof, wherein R_4 is as described herein.

[0518] In another embodiment, the compounds of Formula (I) are of Formula (IIf):



(IIf) or their N-oxides, or salts or isomers thereof, wherein M is -C(O)O- or –OC(O)-, M" is C_{1-6} alkyl or C_{2-6} alkenyl, R_2 and R_3 are independently selected from the group consisting of C_{5-14} alkyl and C_{5-14} alkenyl, and n is selected from 2, 3, and 4.

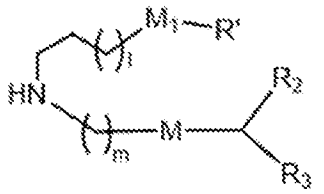
[0519] In a further embodiment, the compounds of Formula (I) are of Formula (IId),



(IId),

or their N-oxides, or salts or isomers thereof, wherein n is 2, 3, or 4; and m, R', R", and R_2 through R_6 are as described herein. For example, each of R_2 and R_3 may be independently selected from the group consisting of C_{5-14} alkyl and C_{5-14} alkenyl.

[0520] In a further embodiment, the compounds of Formula (I) are of Formula (IIg),

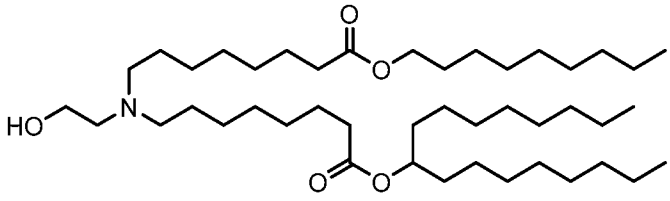


(IIg), or their N-oxides, or salts or isomers thereof, wherein l is selected from 1, 2, 3, 4, and 5; m is selected from 5, 6, 7, 8, and 9; M_1 is a bond or M'; M and M' are independently selected from -C(O)O-, -OC(O)-, -OC(O)-M"-C(O)O-, -C(O)N(R')-, -P(O)(OR')O-, -S-S-, an aryl group, and a heteroaryl group; and R_2 and R_3 are independently selected from the group consisting of H, C_{1-14} alkyl, and C_{2-14} alkenyl. For example, M" is C_{1-6} alkyl (e.g., C_{1-4} alkyl) or C_{2-6} alkenyl (e.g. C_{2-4} alkenyl). For example, R_2 and R_3 are independently selected from the group consisting of C_{5-14} alkyl and C_{5-14} alkenyl.

[0521] In some embodiments, the ionizable lipids are one or more of the compounds described in U.S. Application Nos. 62/220,091, 62/252,316, 62/253,433, 62/266,460, 62/333,557, 62/382,740, 62/393,940, 62/471,937, 62/471,949, 62/475,140, and 62/475,166, and PCT Application No. PCT/US2016/052352.

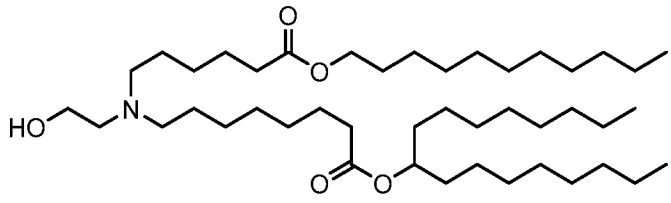
[0522] In some embodiments, the ionizable lipids are selected from Compounds 1-280 described in U.S. Application No. 62/475,166.

[0523] In some embodiments, the ionizable lipid is



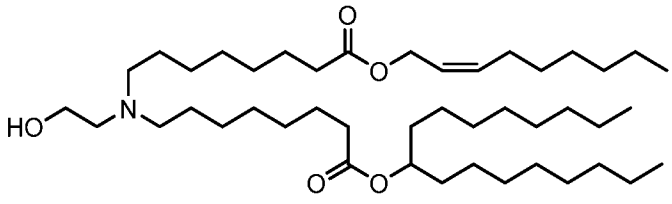
(Compound II), or a salt thereof.

[0524] In some embodiments, the ionizable lipid is



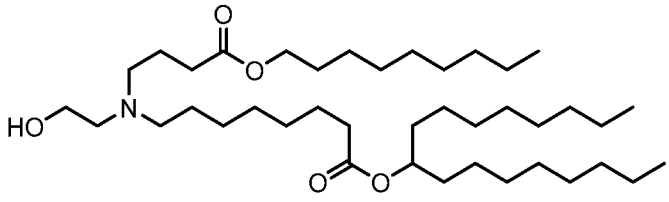
(Compound III), or a salt thereof.

[0525] In some embodiments, the ionizable lipid is



(Compound IV), or a salt thereof.

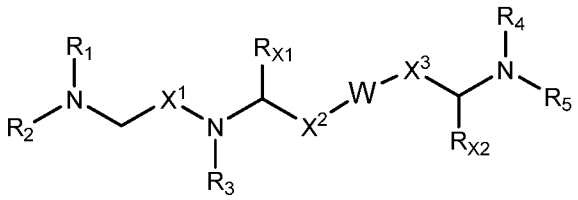
[0526] In some embodiments, the ionizable lipid is



(Compound V), or a salt thereof.

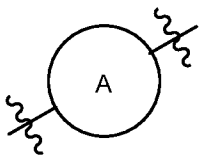
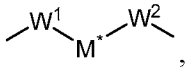
[0527] The central amine moiety of a lipid according to Formula (I), (IA), (IB), (II), (IIa), (IIb), (IIc), (IId), (IIe), (IIf), or (IIg) may be protonated at a physiological pH. Thus, a lipid may have a positive or partial positive charge at physiological pH. Such lipids may be referred to as cationic or ionizable (amino)lipids. Lipids may also be zwitterionic, i.e., neutral molecules having both a positive and a negative charge.

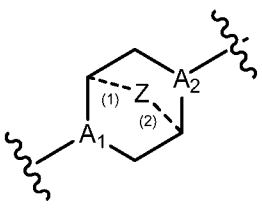
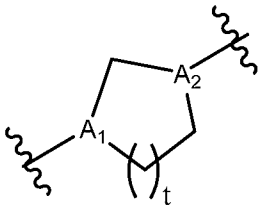
[0528] In some aspects, the ionizable lipids of the present disclosure may be one or more of compounds of formula (III),



(III),

or salts or isomers thereof, wherein

W is  or  ,

ring A is  or  ;

t is 1 or 2;

A_1 and A_2 are each independently selected from CH or N;

Z is CH_2 or absent wherein when Z is CH_2, the dashed lines (1) and (2) each represent a single bond; and when Z is absent, the dashed lines (1) and (2) are both absent;

R_1, R_2, R_3, R_4, and R_5 are independently selected from the group consisting of C_{5-20} alkyl, C_{5-20} alkenyl, -R"MR', -R*YR", -YR", and -R*OR";

R_{X1} and R_{X2} are each independently H or C_{1-3} alkyl;

each M is independently selected from the group consisting of -C(O)O-, -OC(O)-, -OC(O)O-, -C(O)N(R')-, -N(R')C(O)-, -C(O)-, -C(S)-, -C(S)S-, -SC(S)-,
-CH(OH)-, -P(O)(OR')O-, -$S(O)_2$-, -C(O)S-, -SC(O)-, an aryl group, and a heteroaryl group;

M* is C_1-C_6 alkyl,

W^1 and W^2 are each independently selected from the group consisting of -O- and -N(R_6)-;

each R_6 is independently selected from the group consisting of H and C_{1-5} alkyl;

X^1, X^2, and X^3 are independently selected from the group consisting of a bond, -CH_2-, -$(CH_2)_2$-, -CHR-, -CHY-, -C(O)-, -C(O)O-, -OC(O)-, -$(CH_2)_n$-C(O)-, -C(O)-$(CH_2)_n$-, -$(CH_2)_n$-C(O)O-, -OC(O)-$(CH_2)_n$-, -$(CH_2)_n$-OC(O)-, -C(O)O-$(CH_2)_n$-, -CH(OH)-, -C(S)-, and -CH(SH)-;

each Y is independently a C_{3-6} carbocycle;

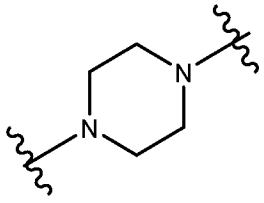
each R* is independently selected from the group consisting of C_{1-12} alkyl and C_{2-12} alkenyl;

each R is independently selected from the group consisting of C_{1-3} alkyl and a C_{3-6} carbocycle;

each R' is independently selected from the group consisting of C_{1-12} alkyl, C_{2-12} alkenyl, and H;

each R" is independently selected from the group consisting of C_{3-12} alkyl, C_{3-12} alkenyl and -R*MR'; and

n is an integer from 1-6;

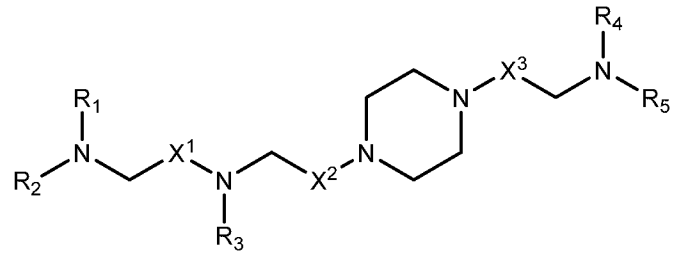
when ring A is 

, then

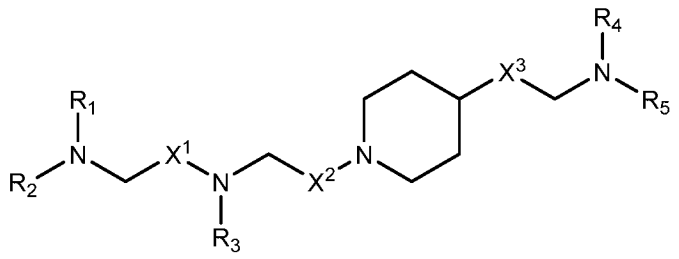
i) at least one of X^1, X^2, and X^3 is not -CH_2-; and/or

ii) at least one of R_1, R_2, R_3, R_4, and R_5 is -R"MR'.

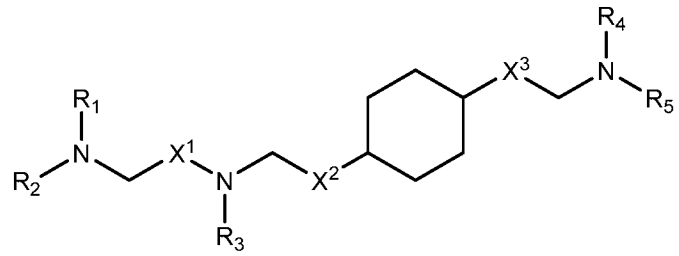
[0529] In some embodiments, the compound is of any of formulae (IIIa1)-(IIIa8):



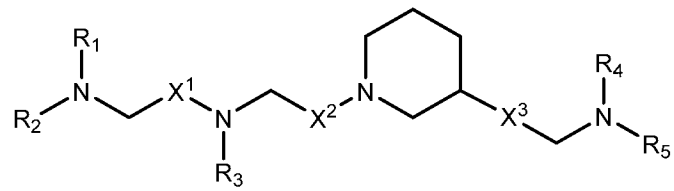
(IIIa1),



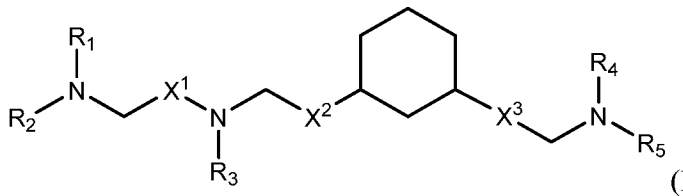
(IIIa2),



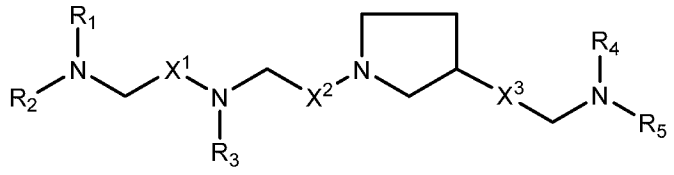
(IIIa3),



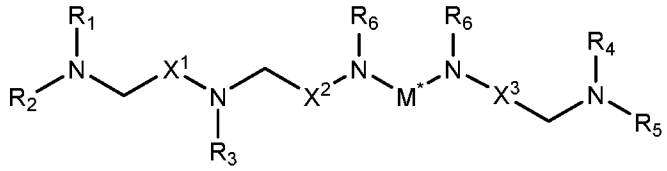
(IIIa4),



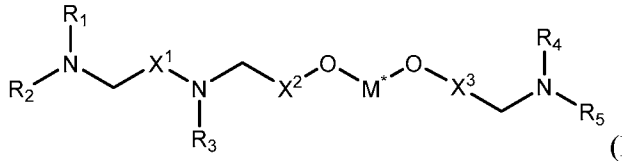
(IIIa5'),



(IIIa6),



(IIIa7), or



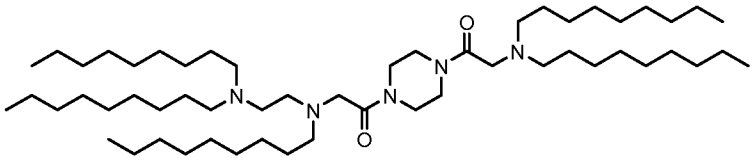
(IIIa8).

[0530] In some embodiments, the ionizable lipids are one or more of the compounds described in U.S. Application Nos. 62/271,146, 62/338,474, 62/413,345, and 62/519,826, and PCT Application No. PCT/US2016/068300.

[0531] In some embodiments, the ionizable lipids are selected from Compounds 1-156 described in U.S. Application No. 62/519,826.

[0532] In some embodiments, the ionizable lipids are selected from Compounds 1-16, 42-66, 68-76, and 78-156 described in U.S. Application No. 62/519,826.

[0533] In some embodiments, the ionizable lipid is



(Compound VI), or a salt thereof.

[0534] In some embodiments, the ionizable lipid is

(Compound VII), or a salt thereof.

[0535] The central amine moiety of a lipid according to Formula (III), (IIIa1), (IIIa2), (IIIa3), (IIIa4), (IIIa5), (IIIa6), (IIIa7), or (IIIa8) may be protonated at a physiological pH. Thus, a lipid may have a positive or partial positive charge at physiological pH. Such lipids may be referred to as cationic or ionizable (amino)lipids. Lipids may also be zwitterionic, i.e., neutral molecules having both a positive and a negative charge.

Phospholipids

[0536] The lipid composition of the lipid nanoparticle composition disclosed herein can comprise one or more phospholipids, for example, one or more saturated or (poly)unsaturated phospholipids or a combination thereof. In general, phospholipids comprise a phospholipid moiety and one or more fatty acid moieties.

[0537] A phospholipid moiety can be selected, for example, from the non-limiting group consisting of phosphatidyl choline, phosphatidyl ethanolamine, phosphatidyl glycerol, phosphatidyl serine, phosphatidic acid, 2-lysophosphatidyl choline, and a sphingomyelin.

[0538] A fatty acid moiety can be selected, for example, from the non-limiting group consisting of lauric acid, myristic acid, myristoleic acid, palmitic acid, palmitoleic acid, stearic acid, oleic acid, linoleic acid, alpha-linolenic acid, erucic acid, phytanoic acid, arachidic acid, arachidonic acid, eicosapentaenoic acid, behenic acid, docosapentaenoic acid, and docosahexaenoic acid.

[0539] Particular phospholipids can facilitate fusion to a membrane. For example, a cationic phospholipid can interact with one or more negatively charged phospholipids of a membrane (e.g., a cellular or intracellular membrane). Fusion of a phospholipid to a membrane can allow one or more elements (e.g., a therapeutic agent) of a lipid-containing composition (e.g., LNPs) to pass through the membrane permitting, e.g., delivery of the one or more elements to a target tissue.

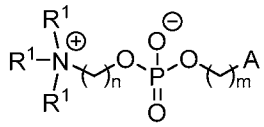
[0540] Non-natural phospholipid species including natural species with modifications and substitutions including branching, oxidation, cyclization, and alkynes are also contemplated. For example, a phospholipid can be functionalized with or cross-linked to one or more alkynes (e.g., an alkenyl group in which one or more double bonds is replaced with a triple bond). Under appropriate reaction conditions, an alkyne group can undergo a copper-catalyzed cycloaddition upon exposure to an azide. Such reactions can be useful in functionalizing a lipid bilayer of a nanoparticle composition to facilitate membrane permeation or cellular recognition or in conjugating a nanoparticle composition to a useful component such as a targeting or imaging moiety (e.g., a dye).

[0541] Phospholipids include, but are not limited to, glycerophospholipids such as phosphatidylcholines, phosphatidylethanolamines, phosphatidylserines, phosphatidylinositols, phosphatidy glycerols, and phosphatidic acids. Phospholipids also include phosphosphingolipid, such as sphingomyelin.

[0542] In some embodiments, a phospholipid of the invention comprises 1,2-distearoyl-sn-glycero-3-phosphocholine (DSPC), 1,2-dioleoyl-sn-glycero-3-phosphoethanolamine (DOPE), 1,2-dilinoleoyl-sn-glycero-3-phosphocholine (DLPC), 1,2-dimyristoyl-sn-gly cero-phosphocholine (DMPC), 1,2-dioleoyl-sn-glycero-3-phosphocholine (DOPC), l,2-dipalmitoyl-sn-glycero-3-phosphocholine (DPPC), 1,2-

diundecanoyl-sn-glycero-phosphocholine (DUPC), 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC), 1,2-di-O-octadecenyl-sn-glycero-3-phosphocholine (18:0 Diether PC), 1-oleoyl-2 cholesterylhemisuccinoyl-sn-glycero-3-phosphocholine (OChemsPC), 1-hexadecyl-sn-glycero-3-phosphocholine (C16 Lyso PC), 1,2-dilinolenoyl-sn-glycero-3-phosphocholine,1,2-diarachidonoyl-sn-glycero-3-phosphocholine, 1,2-didocosahexaenoyl-sn-glycero-3-phosphocholine, 1,2-diphytanoyl-sn-glycero-3-phosphoethanolamine (ME 16.0 PE), 1,2-distearoyl-sn-glycero-3-phosphoethanolamine, 1,2-dilinoleoyl-sn-glycero-3-phosphoethanolamine, 1,2-dilinolenoyl-sn-glycero-3-phosphoethanolamine, 1,2-diarachidonoyl-sn-glycero-3-phosphoethanolamine, 1,2-didocosahexaenoyl-sn-glycero-3-phosphoethanolamine, 1,2-dioleoyl-sn-glycero-3-phospho-rac-(1-glycerol) sodium salt (DOPG), sphingomyelin, and mixtures thereof.

[0543] In certain embodiments, a phospholipid useful or potentially useful in the present invention is an analog or variant of DSPC. In certain embodiments, a phospholipid useful or potentially useful in the present invention is a compound of Formula (**IV**):



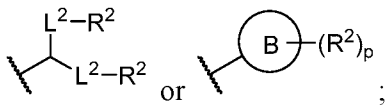
(**IV**),

or a salt thereof, wherein:

each R^1 is independently optionally substituted alkyl; or optionally two R^1 are joined together with the intervening atoms to form optionally substituted monocyclic carbocyclyl or optionally substituted monocyclic heterocyclyl; or optionally three R^1 are joined together with the intervening atoms to form optionally substituted bicyclic carbocyclyl or optionally substitute bicyclic heterocyclyl;

n is 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10;

m is 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10;

A is of the formula:  or ;

each instance of L^2 is independently a bond or optionally substituted C_{1-6} alkylene, wherein one methylene unit of the optionally substituted C_{1-6} alkylene is optionally replaced

with O, $N(R^N)$, S, C(O), $C(O)N(R^N)$, $NR^NC(O)$, C(O)O, OC(O), OC(O)O, $OC(O)N(R^N)$, $-NR^NC(O)O$, or $NR^NC(O)N(R^N)$;

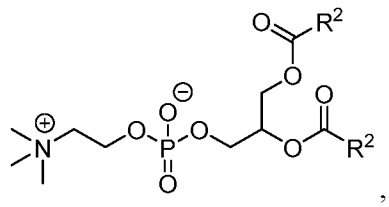
each instance of R^2 is independently optionally substituted C_{1-30} alkyl, optionally substituted C_{1-30} alkenyl, or optionally substituted C_{1-30} alkynyl; optionally wherein one or more methylene units of R^2 are independently replaced with optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted arylene, optionally substituted heteroarylene, $N(R^N)$, O, S, C(O), $C(O)N(R^N)$, $NR^NC(O)$, $-NR^NC(O)N(R^N)$, C(O)O, OC(O), OC(O)O, $OC(O)N(R^N)$, $NR^NC(O)O$, C(O)S, SC(O), $-C(=NR^N)$, $C(=NR^N)N(R^N)$, $NR^NC(=NR^N)$, $NR^NC(=NR^N)N(R^N)$, C(S), $C(S)N(R^N)$, $NR^NC(S)$, $NR^NC(S)N(R^N)$, S(O), OS(O), S(O)O, OS(O)O, $OS(O)_2$, $S(O)_2O$, $OS(O)_2O$, $N(R^N)S(O)$, $-S(O)N(R^N)$, $N(R^N)S(O)N(R^N)$, $OS(O)N(R^N)$, $N(R^N)S(O)O$, $S(O)_2$, $N(R^N)S(O)_2$, $S(O)_2N(R^N)$, $N(R^N)S(O)_2N(R^N)$, $OS(O)_2N(R^N)$, or $N(R^N)S(O)_2O$;

each instance of R^N is independently hydrogen, optionally substituted alkyl, or a nitrogen protecting group;

Ring B is optionally substituted carbocyclyl, optionally substituted heterocyclyl, optionally substituted aryl, or optionally substituted heteroaryl; and

p is 1 or 2;

provided that the compound is not of the formula:



,

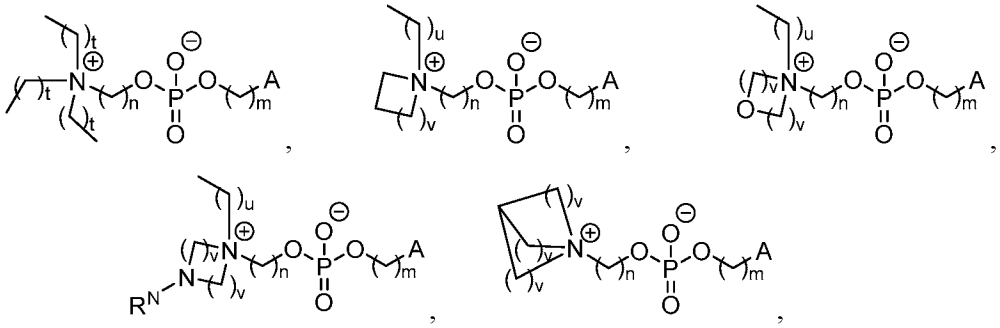
wherein each instance of R^2 is independently unsubstituted alkyl, unsubstituted alkenyl, or unsubstituted alkynyl.

[0544] In some embodiments, the phospholipids may be one or more of the phospholipids described in U.S. Application No. 62/520,530.

i) Phospholipid Head Modifications

[0545] In certain embodiments, a phospholipid useful or potentially useful in the present invention comprises a modified phospholipid head (*e.g.*, a modified choline group). In certain embodiments, a phospholipid with a modified head is DSPC, or analog thereof, with a modified quaternary amine. For example, in embodiments of Formula (**IV**), at least one of R^1 is not methyl. In certain embodiments, at least one of

R^1 is not hydrogen or methyl. In certain embodiments, the compound of Formula (**IV**) is of one of the following formulae:



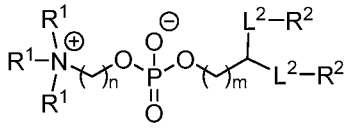
or a salt thereof, wherein:

each t is independently 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10;

each u is independently 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10; and

each v is independently 1, 2, or 3.

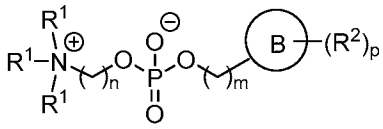
In certain embodiments, a compound of Formula (**IV**) is of Formula (**IV-a**):



(**IV-a**),

or a salt thereof.

[0546] In certain embodiments, a phospholipid useful or potentially useful in the present invention comprises a cyclic moiety in place of the glyceride moiety. In certain embodiments, a phospholipid useful in the present invention is DSPC, or analog thereof, with a cyclic moiety in place of the glyceride moiety. In certain embodiments, the compound of Formula (**IV**) is of Formula (**IV-b**):



(**IV-b**),

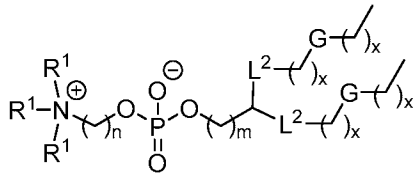
or a salt thereof.

(ii) Phospholipid Tail Modifications

[0547] In certain embodiments, a phospholipid useful or potentially useful in the present invention comprises a modified tail. In certain embodiments, a phospholipid useful or potentially useful in the present invention is DSPC, or analog thereof, with a

modified tail. As described herein, a "modified tail" may be a tail with shorter or longer aliphatic chains, aliphatic chains with branching introduced, aliphatic chains with substituents introduced, aliphatic chains wherein one or more methylenes are replaced by cyclic or heteroatom groups, or any combination thereof. For example, in certain embodiments, the compound of (**IV**) is of Formula (**IV-a**), or a salt thereof, wherein at least one instance of R^2 is each instance of R^2 is optionally substituted C_{1-30} alkyl, wherein one or more methylene units of R^2 are independently replaced with optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted arylene, optionally substituted heteroarylene, $N(R^N)$, O, S, -C(O), $C(O)N(R^N)$, $NR^NC(O)$, $NR^NC(O)N(R^N)$, C(O)O, OC(O), OC(O)O, -$OC(O)N(R^N)$, $NR^NC(O)O$, C(O)S, SC(O), $C(=NR^N)$, $C(=NR^N)N(R^N)$, $NR^NC(=NR^N)$, $NR^NC(=NR^N)N(R^N)$, C(S), $C(S)N(R^N)$, $NR^NC(S)$, $NR^NC(S)N(R^N)$, S(O), OS(O), -S(O)O, OS(O)O, $OS(O)_2$, $S(O)_2O$, $OS(O)_2O$, $N(R^N)S(O)$, $S(O)N(R^N)$, -$N(R^N)S(O)N(R^N)$, $OS(O)N(R^N)$, $N(R^N)S(O)O$, $S(O)_2$, $N(R^N)S(O)_2$, $S(O)_2N(R^N)$, -$N(R^N)S(O)_2N(R^N)$, $OS(O)_2N(R^N)$, or $N(R^N)S(O)_2O$.

[0548] In certain embodiments, the compound of Formula (**IV**) is of Formula (**IV-c**):



(**IV-c**),

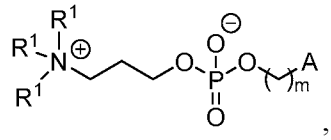
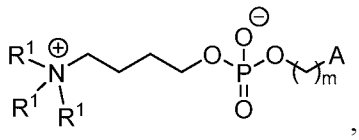
or a salt thereof, wherein:

each x is independently an integer between 0-30, inclusive; and

each instance is G is independently selected from the group consisting of optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted arylene, optionally substituted heteroarylene, $N(R^N)$, O, S, C(O), $C(O)N(R^N)$, $NR^NC(O)$, -$NR^NC(O)N(R^N)$, C(O)O, OC(O), OC(O)O, $OC(O)N(R^N)$, $NR^NC(O)O$, C(O)S, SC(O), -$C(=NR^N)$, $C(=NR^N)N(R^N)$, $NR^NC(=NR^N)$, $NR^NC(=NR^N)N(R^N)$, C(S), $C(S)N(R^N)$, $NR^NC(S)$, $NR^NC(S)N(R^N)$, S(O), OS(O), S(O)O, OS(O)O, $OS(O)_2$, $S(O)_2O$, $OS(O)_2O$, $N(R^N)S(O)$, -$S(O)N(R^N)$, $N(R^N)S(O)N(R^N)$, $OS(O)N(R^N)$, $N(R^N)S(O)O$, $S(O)_2$, $N(R^N)S(O)_2$, $S(O)_2N(R^N)$, $N(R^N)S(O)_2N(R^N)$, $OS(O)_2N(R^N)$, or $N(R^N)S(O)_2O$. Each possibility represents a separate embodiment of the present invention.

[0549] In certain embodiments, a phospholipid useful or potentially useful in the present invention comprises a modified phosphocholine moiety, wherein the alkyl chain linking the quaternary amine to the phosphoryl group is not ethylene (*e.g.*, n is

not 2). Therefore, in certain embodiments, a phospholipid useful or potentially useful in the present invention is a compound of Formula (**IV**), wherein n is 1, 3, 4, 5, 6, 7, 8, 9, or 10. For example, in certain embodiments, a compound of Formula (**IV**) is of one of the following formulae:


,

,

or a salt thereof.

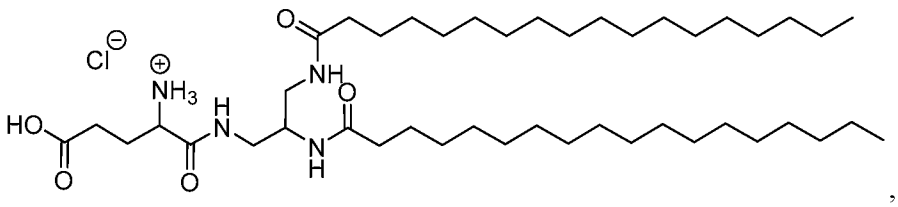
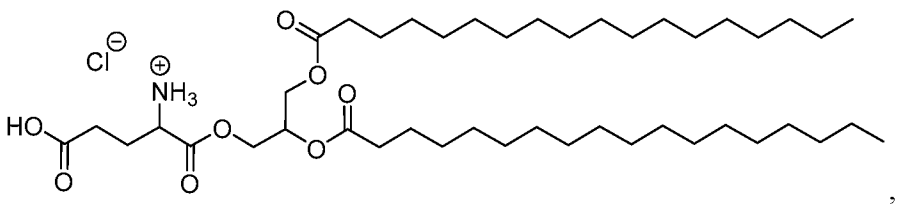
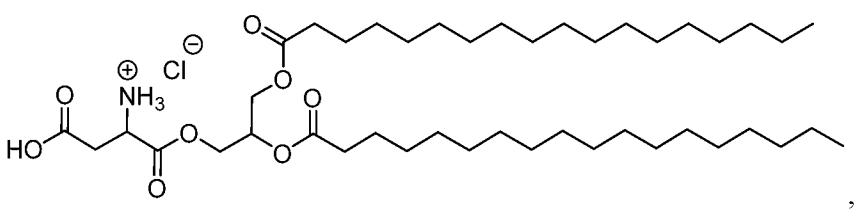
Alternative Lipids

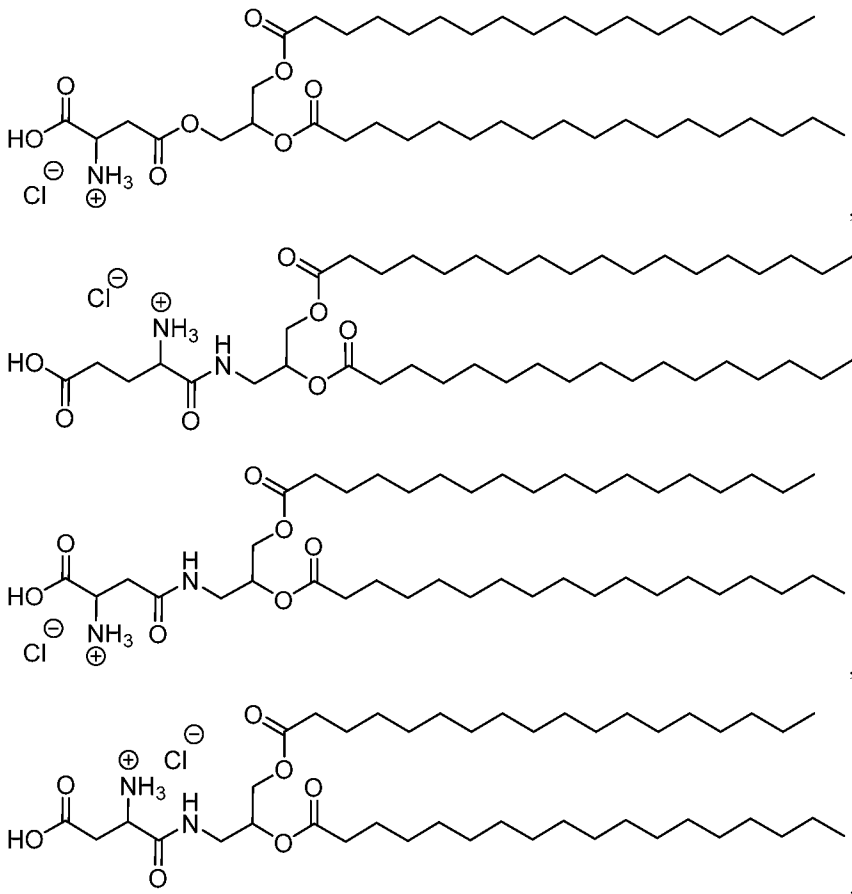
[0550] In certain embodiments, a phospholipid useful or potentially useful in the present invention comprises a modified phosphocholine moiety, wherein the alkyl chain linking the quaternary amine to the phosphoryl group is not ethylene (*e.g.*, n is not 2). Therefore, in certain embodiments, a phospholipid useful.

[0551] In certain embodiments, an alternative lipid is used in place of a phospholipid of the present disclosure.

[0552] In certain embodiments, an alternative lipid of the invention is oleic acid.

[0553] In certain embodiments, the alternative lipid is one of the following:


,

,

,



, and

Structural Lipids

[0554] The lipid composition of a pharmaceutical composition disclosed herein can comprise one or more structural lipids. As used herein, the term "structural lipid" refers to sterols and to lipids containing sterol moieties.

[0555] Incorporation of structural lipids in the lipid nanoparticle may help mitigate aggregation of other lipids in the particle. Structural lipids can be selected from the group including but not limited to, cholesterol, fecosterol, sitosterol, ergosterol, campesterol, stigmasterol, brassicasterol, tomatidine, tomatine, ursolic acid, alpha-tocopherol, hopanoids, phytosterols, steroids, and mixtures thereof. In some embodiments, the structural lipid is a sterol. As defined herein, "sterols" are a subgroup of steroids consisting of steroid alcohols. In certain embodiments, the structural lipid is a steroid. In certain embodiments, the structural lipid is cholesterol. In certain embodiments, the structural lipid is an analog of cholesterol. In certain embodiments, the structural lipid is alpha-tocopherol.

[0556] In some embodiments, the structural lipids may be one or more of the structural lipids described in U.S. Application No. 62 /520,530.

Polyethylene Glycol (PEG)-Lipids

[0557] The lipid composition of a pharmaceutical composition disclosed herein can comprise one or more a polyethylene glycol (PEG) lipid.

[0558] As used herein, the term "PEG-lipid" refers to polyethylene glycol (PEG)-modified lipids. Non-limiting examples of PEG-lipids include PEG-modified phosphatidylethanolamine and phosphatidic acid, PEG-ceramide conjugates (e.g., PEG-CerC14 or PEG-CerC20), PEG-modified dialkylamines and PEG-modified 1,2-diacyloxypropan-3-amines. Such lipids are also referred to as PEGylated lipids. For example, a PEG lipid can be PEG-c-DOMG, PEG-DMG, PEG-DLPE, PEG-DMPE, PEG-DPPC, or a PEG-DSPE lipid.

[0559] In some embodiments, the PEG-lipid includes, but not limited to 1,2-dimyristoyl-sn-glycerol methoxypolyethylene glycol (PEG-DMG), 1,2-distearoyl-sn-glycero-3-phosphoethanolamine-N-[amino(polyethylene glycol)] (PEG-DSPE), PEG-disteryl glycerol (PEG-DSG), PEG-dipalmetoleyl, PEG-dioleyl, PEG-distearyl, PEG-diacylglycamide (PEG-DAG), PEG-dipalmitoyl phosphatidylethanolamine (PEG-DPPE), or PEG-l,2-dimyristyloxlpropyl-3-amine (PEG-c-DMA).

[0560] In one embodiment, the PEG-lipid is selected from the group consisting of a PEG-modified phosphatidylethanolamine, a PEG-modified phosphatidic acid, a PEG-modified ceramide, a PEG-modified dialkylamine, a PEG-modified diacylglycerol, a PEG-modified dialkylglycerol, and mixtures thereof.

[0561] In some embodiments, the lipid moiety of the PEG-lipids includes those having lengths of from about C_{14} to about C_{22}, preferably from about C_{14} to about C_{16}. In some embodiments, a PEG moiety, for example an mPEG-NH_2, has a size of about 1000, 2000, 5000, 10,000, 15,000 or 20,000 daltons. In one embodiment, the PEG-lipid is PEG_{2k}-DMG.

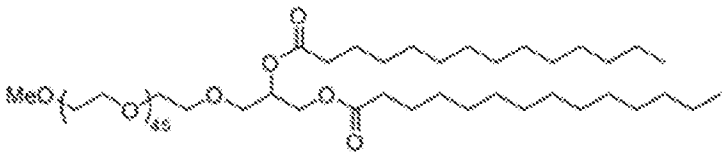
[0562] In one embodiment, the lipid nanoparticles described herein can comprise a PEG lipid which is a non-diffusible PEG. Non-limiting examples of non-diffusible PEGs include PEG-DSG and PEG-DSPE.

[0563] PEG-lipids are known in the art, such as those described in U.S. Patent No. 8158601 and International Publ. No. WO 2015/130584 A2, which are incorporated herein by reference in their entirety.

[0564] In general, some of the other lipid components (e.g., PEG lipids) of various formulae, described herein may be synthesized as described International Patent Application No. PCT/US2016/000129, filed December 10, 2016, entitled "Compositions and Methods for Delivery of Therapeutic Agents," which is incorporated by reference in its entirety.

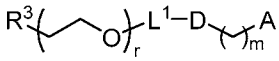
[0565] The lipid component of a lipid nanoparticle composition may include one or more molecules comprising polyethylene glycol, such as PEG or PEG-modified lipids. Such species may be alternately referred to as PEGylated lipids. A PEG lipid is a lipid modified with polyethylene glycol. A PEG lipid may be selected from the non-limiting group including PEG-modified phosphatidylethanolamines, PEG-modified phosphatidic acids, PEG-modified ceramides, PEG-modified dialkylamines, PEG-modified diacylglycerols, PEG-modified dialkylglycerols, and mixtures thereof. For example, a PEG lipid may be PEG-c-DOMG, PEG-DMG, PEG-DLPE, PEG-DMPE, PEG-DPPC, or a PEG-DSPE lipid.

[0566] In some embodiments the PEG-modified lipids are a modified form of PEG DMG. PEG-DMG has the following structure:



[0567] In one embodiment, PEG lipids useful in the present invention can be PEGylated lipids described in International Publication No. WO2012099755, the contents of which is herein incorporated by reference in its entirety. Any of these exemplary PEG lipids described herein may be modified to comprise a hydroxyl group on the PEG chain. In certain embodiments, the PEG lipid is a PEG-OH lipid. As generally defined herein, a "PEG-OH lipid" (also referred to herein as "hydroxy-PEGylated lipid") is a PEGylated lipid having one or more hydroxyl (–OH) groups on the lipid. In certain embodiments, the PEG-OH lipid includes one or more hydroxyl groups on the PEG chain. In certain embodiments, a PEG-OH or hydroxy-PEGylated lipid comprises an –OH group at the terminus of the PEG chain. Each possibility represents a separate embodiment of the present invention.

[0568] In certain embodiments, a PEG lipid useful in the present invention is a compound of Formula (**V**). Provided herein are compounds of Formula (**V**):



(V),

or salts thereof, wherein:

R^3 is $-OR^O$;

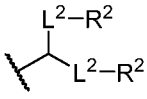
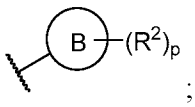
R^O is hydrogen, optionally substituted alkyl, or an oxygen protecting group;

r is an integer between 1 and 100, inclusive;

L^1 is optionally substituted C_{1-10} alkylene, wherein at least one methylene of the optionally substituted C_{1-10} alkylene is independently replaced with optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted arylene, optionally substituted heteroarylene, O, $N(R^N)$, S, C(O), $C(O)N(R^N)$, $NR^NC(O)$, C(O)O, -OC(O), OC(O)O, $OC(O)N(R^N)$, $NR^NC(O)O$, or $NR^NC(O)N(R^N)$;

D is a moiety obtained by click chemistry or a moiety cleavable under physiological conditions;

m is 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10;

A is of the formula:  or  ;

each instance of L^2 is independently a bond or optionally substituted C_{1-6} alkylene, wherein one methylene unit of the optionally substituted C_{1-6} alkylene is optionally replaced with O, $N(R^N)$, S, C(O), $C(O)N(R^N)$, $NR^NC(O)$, C(O)O, OC(O), OC(O)O, $OC(O)N(R^N)$, -$NR^NC(O)O$, or $NR^NC(O)N(R^N)$;

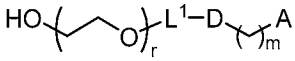
each instance of R^2 is independently optionally substituted C_{1-30} alkyl, optionally substituted C_{1-30} alkenyl, or optionally substituted C_{1-30} alkynyl; optionally wherein one or more methylene units of R^2 are independently replaced with optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted arylene, optionally substituted heteroarylene, $N(R^N)$, O, S, C(O), $C(O)N(R^N)$, $NR^NC(O)$, -$NR^NC(O)N(R^N)$, C(O)O, OC(O), OC(O)O, $OC(O)N(R^N)$, $NR^NC(O)O$, C(O)S, SC(O), -$C(=NR^N)$, $C(=NR^N)N(R^N)$, $NR^NC(=NR^N)$, $NR^NC(=NR^N)N(R^N)$, C(S), $C(S)N(R^N)$, $NR^NC(S)$, $NR^NC(S)N(R^N)$, S(O) , OS(O), S(O)O, OS(O)O, $OS(O)_2$, $S(O)_2O$, $OS(O)_2O$, $N(R^N)S(O)$, -$S(O)N(R^N)$, $N(R^N)S(O)N(R^N)$, $OS(O)N(R^N)$, $N(R^N)S(O)O$, $S(O)_2$, $N(R^N)S(O)_2$, $S(O)_2N(R^N)$, $N(R^N)S(O)_2N(R^N)$, $OS(O)_2N(R^N)$, or $N(R^N)S(O)_2O$;

each instance of R^N is independently hydrogen, optionally substituted alkyl, or a nitrogen protecting group;

Ring B is optionally substituted carbocyclyl, optionally substituted heterocyclyl, optionally substituted aryl, or optionally substituted heteroaryl; and

p is 1 or 2.

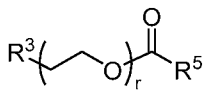
[0569] In certain embodiments, the compound of Fomula (**V**) is a PEG-OH lipid (*i.e.*, R^3 is $-OR^O$, and R^O is hydrogen). In certain embodiments, the compound of Formula (**V**) is of Formula (**V-OH**):



(V-OH),

or a salt thereof.

[0570] In certain embodiments, a PEG lipid useful in the present invention is a PEGylated fatty acid. In certain embodiments, a PEG lipid useful in the present invention is a compound of Formula (**VI**). Provided herein are compounds of Formula (**VI**):



(**VI**),

or a salt thereof, wherein:

R^3 is $-OR^O$;

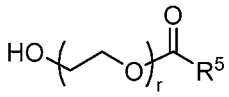
R^O is hydrogen, optionally substituted alkyl or an oxygen protecting group;

r is an integer between 1 and 100, inclusive;

R^5 is optionally substituted C_{10-40} alkyl, optionally substituted C_{10-40} alkenyl, or optionally substituted C_{10-40} alkynyl; and optionally one or more methylene groups of R^5 are replaced with optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted arylene, optionally substituted heteroarylene, $N(R^N)$, O, S, C(O), $-C(O)N(R^N)$, $NR^NC(O)$, $NR^NC(O)N(R^N)$, C(O)O, OC(O), OC(O)O, $OC(O)N(R^N)$, $-NR^NC(O)O$, C(O)S, SC(O), $C(=NR^N)$, $C(=NR^N)N(R^N)$, $NR^NC(=NR^N)$, $NR^NC(=NR^N)N(R^N)$, C(S), $C(S)N(R^N)$, $NR^NC(S)$, $NR^NC(S)N(R^N)$, S(O), OS(O), S(O)O, OS(O)O, $OS(O)_2$, $-S(O)_2O$, $OS(O)_2O$, $N(R^N)S(O)$, $S(O)N(R^N)$, $N(R^N)S(O)N(R^N)$, $OS(O)N(R^N)$, $N(R^N)S(O)O$, $S(O)_2$, $N(R^N)S(O)_2$, $S(O)_2N(R^N)$, $N(R^N)S(O)_2N(R^N)$, $OS(O)_2N(R^N)$, or $N(R^N)S(O)_2O$; and

each instance of R^N is independently hydrogen, optionally substituted alkyl, or a nitrogen protecting group.

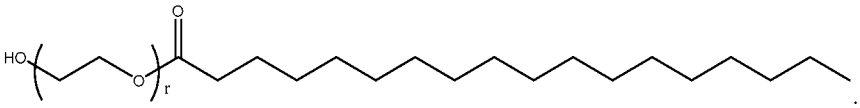
[0571] In certain embodiments, the compound of Formula (**VI**) is of Formula (**VI-OH**):



(VI-OH),

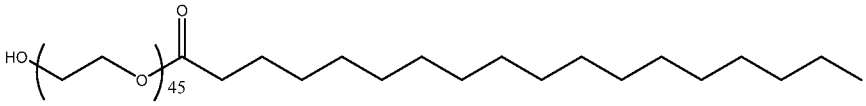
or a salt thereof. In one embodiment, r is an integer between 1 and 100, inclusive. In some embodiments, r is 45.

[0572] In yet other embodiments the compound of Formula (VI) is:



or a salt thereof.

[0573] In one embodiment, the compound of Formula (VI) is



(Compound I).

[0574] In some aspects, the lipid composition of the pharmaceutical compositions disclosed herein does not comprise a PEG-lipid.

[0575] In some embodiments, the PEG-lipids may be one or more of the PEG lipids described in U.S. Application No. 62/520,530.

[0576] In some embodiments, a PEG lipid of the invention comprises a PEG-modified phosphatidylethanolamine, a PEG-modified phosphatidic acid, a PEG-modified ceramide, a PEG-modified dialkylamine, a PEG-modified diacylglycerol, a PEG-modified dialkylglycerol, and mixtures thereof. In some embodiments, the PEG-modified lipid is PEG-DMG, PEG-c-DOMG (also referred to as PEG-DOMG), PEG-DSG and/or PEG-DPG.

[0577] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of any of Formula I, II or III, a phospholipid comprising DSPC, a structural lipid, and a PEG lipid comprising PEG-DMG.

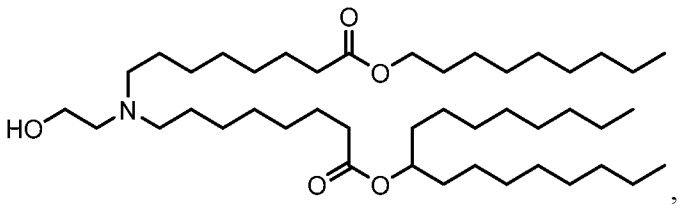
[0578] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of any of Formula I, II or III, a phospholipid comprising DSPC, a structural lipid, and a PEG lipid comprising a compound having Formula VI.

[0579] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of Formula I, II or III, a phospholipid comprising a compound having Formula IV, a structural lipid, and the PEG lipid comprising a compound having Formula V or VI.

[0580] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of Formula I, II or III, a phospholipid comprising a compound having Formula IV, a structural lipid, and the PEG lipid comprising a compound having Formula V or VI.

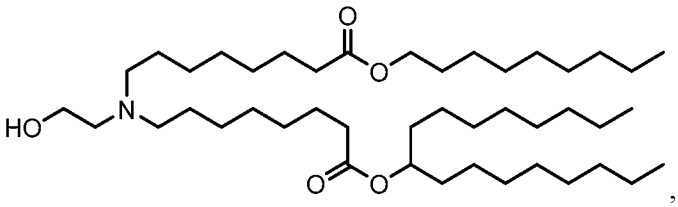
[0581] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of Formula I, II or III, a phospholipid having Formula IV, a structural lipid, and a PEG lipid comprising a compound having Formula VI.

[0582] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of



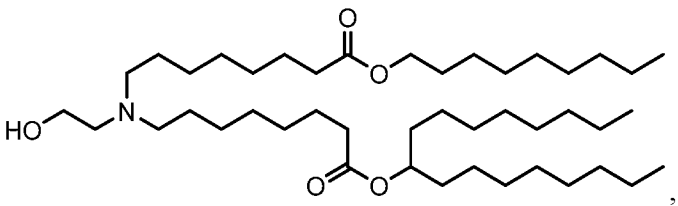
and a PEG lipid comprising Formula VI.

[0583] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of



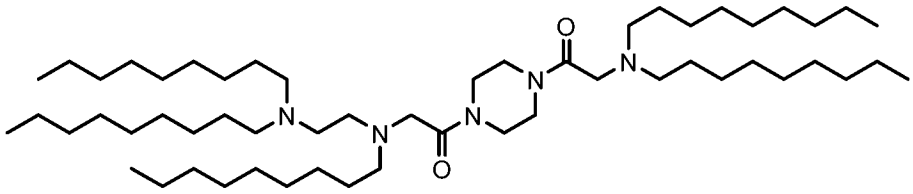
and an alternative lipid comprising oleic acid.

[0584] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of



an alternative lipid comprising oleic acid, a structural lipid comprising cholesterol, and a PEG lipid comprising a compound having Formula VI.

[0585] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of



a phospholipid comprising DOPE, a structural lipid comprising cholesterol, and a PEG lipid comprising a compound having Formula VI.

[0586] In some embodiments, a LNP of the invention comprises an ionizable cationic lipid of

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a phospholipid comprising DOPE, a structural lipid comprising cholesterol, and a PEG lipid comprising a compound having Formula VII.

[0587] In some embodiments, a LNP of the invention comprises an N:P ratio of from about 2:1 to about 30:1.

[0588] In some embodiments, a LNP of the invention comprises an N:P ratio of about 6:1.

[0589] In some embodiments, a LNP of the invention comprises an N:P ratio of about 3:1.

[0590] In some embodiments, a LNP of the invention comprises a wt/wt ratio of the ionizable cationic lipid component to the RNA of from about 10:1 to about 100:1.

[0591] In some embodiments, a LNP of the invention comprises a wt/wt ratio of the ionizable cationic lipid component to the RNA of about 20:1.

[0592] In some embodiments, a LNP of the invention comprises a wt/wt ratio of the ionizable cationic lipid component to the RNA of about 10:1.

[0593] In some embodiments, a LNP of the invention has a mean diameter from about 50nm to about 150nm.

[0594] In some embodiments, a LNP of the invention has a mean diameter from about 70nm to about 120nm.

[0595] As used herein, the term "alkyl", "alkyl group", or "alkylene" means a linear or branched, saturated hydrocarbon including one or more carbon atoms (e.g., one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen, fifteen, sixteen, seventeen, eighteen, nineteen, twenty, or more carbon atoms), which is

optionally substituted. The notation "C1-14 alkyl" means an optionally substituted linear or branched, saturated hydrocarbon including 1 14 carbon atoms. Unless otherwise specified, an alkyl group described herein refers to both unsubstituted and substituted alkyl groups.

[0596] As used herein, the term "alkenyl", "alkenyl group", or "alkenylene" means a linear or branched hydrocarbon including two or more carbon atoms (e.g., two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen, fifteen, sixteen, seventeen, eighteen, nineteen, twenty, or more carbon atoms) and at least one double bond, which is optionally substituted. The notation "C2-14 alkenyl" means an optionally substituted linear or branched hydrocarbon including 2 14 carbon atoms and at least one carbon-carbon double bond. An alkenyl group may include one, two, three, four, or more carbon-carbon double bonds. For example, C18 alkenyl may include one or more double bonds. A C18 alkenyl group including two double bonds may be a linoleyl group. Unless otherwise specified, an alkenyl group described herein refers to both unsubstituted and substituted alkenyl groups.

[0597] As used herein, the term "alkynyl", "alkynyl group", or "alkynylene" means a linear or branched hydrocarbon including two or more carbon atoms (e.g., two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen, fifteen, sixteen, seventeen, eighteen, nineteen, twenty, or more carbon atoms) and at least one carbon-carbon triple bond, which is optionally substituted. The notation "C2-14 alkynyl" means an optionally substituted linear or branched hydrocarbon including 2 14 carbon atoms and at least one carbon-carbon triple bond. An alkynyl group may include one, two, three, four, or more carbon-carbon triple bonds. For example, C18 alkynyl may include one or more carbon-carbon triple bonds. Unless otherwise specified, an alkynyl group described herein refers to both unsubstituted and substituted alkynyl groups.

[0598] As used herein, the term "carbocycle" or "carbocyclic group" means an optionally substituted mono- or multi-cyclic system including one or more rings of carbon atoms. Rings may be three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen, fifteen, sixteen, seventeen, eighteen, nineteen, or twenty membered rings. The notation "C3-6 carbocycle" means a carbocycle including a single ring having 3-6 carbon atoms. Carbocycles may include one or more carbon-carbon double or triple bonds and may be non-aromatic or aromatic (e.g., cycloalkyl or aryl groups). Examples of carbocycles include cyclopropyl, cyclopentyl,

cyclohexyl, phenyl, naphthyl, and 1,2 dihydronaphthyl groups. The term "cycloalkyl" as used herein means a non-aromatic carbocycle and may or may not include any double or triple bond. Unless otherwise specified, carbocycles described herein refers to both unsubstituted and substituted carbocycle groups, i.e., optionally substituted carbocycles.

[0599] As used herein, the term "heterocycle" or "heterocyclic group" means an optionally substituted mono- or multi-cyclic system including one or more rings, where at least one ring includes at least one heteroatom. Heteroatoms may be, for example, nitrogen, oxygen, or sulfur atoms. Rings may be three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, or fourteen membered rings. Heterocycles may include one or more double or triple bonds and may be non-aromatic or aromatic (e.g., heterocycloalkyl or heteroaryl groups). Examples of heterocycles include imidazolyl, imidazolidinyl, oxazolyl, oxazolidinyl, thiazolyl, thiazolidinyl, pyrazolidinyl, pyrazolyl, isoxazolidinyl, isoxazolyl, isothiazolidinyl, isothiazolyl, morpholinyl, pyrrolyl, pyrrolidinyl, furyl, tetrahydrofuryl, thiophenyl, pyridinyl, piperidinyl, quinolyl, and isoquinolyl groups. The term "heterocycloalkyl" as used herein means a non-aromatic heterocycle and may or may not include any double or triple bond. Unless otherwise specified, heterocycles described herein refers to both unsubstituted and substituted heterocycle groups, i.e., optionally substituted heterocycles.

[0600] As used herein, the term "heteroalkyl", "heteroalkenyl", or "heteroalkynyl", refers respectively to an alkyl, alkenyl, alkynyl group, as defined herein, which further comprises one or more (e.g., 1, 2, 3, or 4) heteroatoms (e.g., oxygen, sulfur, nitrogen, boron, silicon, phosphorus) wherein the one or more heteroatoms is inserted between adjacent carbon atoms within the parent carbon chain and/or one or more heteroatoms is inserted between a carbon atom and the parent molecule, i.e., between the point of attachment. Unless otherwise specified, heteroalkyls, heteroalkenyls, or heteroalkynyls described herein refers to both unsubstituted and substituted heteroalkyls, heteroalkenyls, or heteroalkynyls, i.e., optionally substituted heteroalkyls, heteroalkenyls, or heteroalkynyls.

[0601] As used herein, a "biodegradable group" is a group that may facilitate faster metabolism of a lipid in a mammalian entity. A biodegradable group may be selected from the group consisting of, but is not limited to, -C(O)O-, -OC(O)-, -C(O)N(R')-, -N(R')C(O)-, -C(O)-, -C(S)-, -C(S)S-, -SC(S)-, -CH(OH)-, -P(O)(OR')O-, -S(O)2-,

an aryl group, and a heteroaryl group. As used herein, an "aryl group" is an optionally substituted carbocyclic group including one or more aromatic rings. Examples of aryl groups include phenyl and naphthyl groups. As used herein, a "heteroaryl group" is an optionally substituted heterocyclic group including one or more aromatic rings. Examples of heteroaryl groups include pyrrolyl, furyl, thiophenyl, imidazolyl, oxazolyl, and thiazolyl. Both aryl and heteroaryl groups may be optionally substituted. For example, M and M' can be selected from the non-limiting group consisting of optionally substituted phenyl, oxazole, and thiazole. In the formulas herein, M and M' can be independently selected from the list of biodegradable groups above. Unless otherwise specified, aryl or heteroaryl groups described herein refers to both unsubstituted and substituted groups, i.e., optionally substituted aryl or heteroaryl groups.

[0602] Alkyl, alkenyl, and cyclyl (e.g., carbocyclyl and heterocyclyl) groups may be optionally substituted unless otherwise specified. Optional substituents may be selected from the group consisting of, but are not limited to, a halogen atom (e.g., a chloride, bromide, fluoride, or iodide group), a carboxylic acid (e.g., C(O)OH), an alcohol (e.g., a hydroxyl, OH), an ester (e.g., C(O)OR OC(O)R), an aldehyde (e.g., C(O)H), a carbonyl (e.g., C(O)R, alternatively represented by C=O), an acyl halide (e.g., C(O)X, in which X is a halide selected from bromide, fluoride, chloride, and iodide), a carbonate (e.g., OC(O)OR), an alkoxy (e.g., OR), an acetal (e.g., C(OR)2R"", in which each OR are alkoxy groups that can be the same or different and R"" is an alkyl or alkenyl group), a phosphate (e.g., P(O)43-), a thiol (e.g., SH), a sulfoxide (e.g., S(O)R), a sulfinic acid (e.g., S(O)OH), a sulfonic acid (e.g., S(O)2OH), a thial (e.g., C(S)H), a sulfate (e.g., S(O)42-), a sulfonyl (e.g., S(O)2), an amide (e.g., C(O)NR2, or N(R)C(O)R), an azido (e.g., N3), a nitro (e.g., NO2), a cyano (e.g., CN), an isocyano (e.g., NC), an acyloxy (e.g., OC(O)R), an amino (e.g., NR2, NRH, or NH2), a carbamoyl (e.g., OC(O)NR2, OC(O)NRH, or OC(O)NH2), a sulfonamide (e.g., S(O)2NR2, S(O)2NRH, S(O)2NH2, N(R)S(O)2R, N(H)S(O)2R, N(R)S(O)2H, or N(H)S(O)2H), an alkyl group, an alkenyl group, and a cyclyl (e.g., carbocyclyl or heterocyclyl) group. In any of the preceding, R is an alkyl or alkenyl group, as defined herein. In some embodiments, the substituent groups themselves may be further substituted with, for example, one, two, three, four, five, or six substituents as defined herein. For example, a C1 6 alkyl

group may be further substituted with one, two, three, four, five, or six substituents as described herein.

[0603] Compounds of the disclosure that contain nitrogens can be converted to N-oxides by treatment with an oxidizing agent (e.g., 3-chloroperoxybenzoic acid (mCPBA) and/or hydrogen peroxides) to afford other compounds of the disclosure. Thus, all shown and claimed nitrogen-containing compounds are considered, when allowed by valency and structure, to include both the compound as shown and its N-oxide derivative (which can be designated as N□O or N+-O-). Furthermore, in other instances, the nitrogens in the compounds of the disclosure can be converted to N-hydroxy or N-alkoxy compounds. For example, N-hydroxy compounds can be prepared by oxidation of the parent amine by an oxidizing agent such as m CPBA. All shown and claimed nitrogen-containing compounds are also considered, when allowed by valency and structure, to cover both the compound as shown and its N-hydroxy (i.e., N-OH) and N-alkoxy (i.e., N-OR, wherein R is substituted or unsubstituted C1-C 6 alkyl, C1-C6 alkenyl, C1-C6 alkynyl, 3-14-membered carbocycle or 3-14-membered heterocycle) derivatives.

(vi) Other Lipid Composition Components

[0604] The lipid composition of a pharmaceutical composition disclosed herein can include one or more components in addition to those described above. For example, the lipid composition can include one or more permeability enhancer molecules, carbohydrates, polymers, surface altering agents (e.g., surfactants), or other components. For example, a permeability enhancer molecule can be a molecule described by U.S. Patent Application Publication No. 2005/0222064. Carbohydrates can include simple sugars (e.g., glucose) and polysaccharides (e.g., glycogen and derivatives and analogs thereof).

[0605] A polymer can be included in and/or used to encapsulate or partially encapsulate a pharmaceutical composition disclosed herein (e.g., a pharmaceutical composition in lipid nanoparticle form). A polymer can be biodegradable and/or biocompatible. A polymer can be selected from, but is not limited to, polyamines, polyethers, polyamides, polyesters, polycarbamates, polyureas, polycarbonates, polystyrenes, polyimides, polysulfones, polyurethanes, polyacetylenes, polyethylenes, polyethyleneimines, polyisocyanates, polyacrylates, polymethacrylates, polyacrylonitriles, and polyarylates.

[0606] The ratio between the lipid composition and the polynucleotide range can be from about 10:1 to about 60:1 (wt/wt).

[0607] In some embodiments, the ratio between the lipid composition and the polynucleotide can be about 10:1, 11:1, 12:1, 13:1, 14:1, 15:1, 16:1, 17:1, 18:1, 19:1, 20:1, 21:1, 22:1, 23:1, 24:1, 25:1, 26:1, 27:1, 28:1, 29:1, 30:1, 31:1, 32:1, 33:1, 34:1, 35:1, 36:1, 37:1, 38:1, 39:1, 40:1, 41:1, 42:1, 43:1, 44:1, 45:1, 46:1, 47:1, 48:1, 49:1, 50:1, 51:1, 52:1, 53:1, 54:1, 55:1, 56:1, 57:1, 58:1, 59:1 or 60:1 (wt/wt). In some embodiments, the wt/wt ratio of the lipid composition to the polynucleotide encoding a therapeutic agent is about 20:1 or about 15:1.

[0608] In some embodiments, the pharmaceutical composition disclosed herein can contain more than one polypeptides. For example, a pharmaceutical composition disclosed herein can contain two or more polynucleotides (e.g., RNA, e.g., mRNA).

[0609] In one embodiment, the lipid nanoparticles described herein can comprise polynucleotides (e.g., mRNA) in a lipid:polynucleotide weight ratio of 5:1, 10:1, 15:1, 20:1, 25:1, 30:1, 35:1, 40:1, 45:1, 50:1, 55:1, 60:1 or 70:1, or a range or any of these ratios such as, but not limited to, 5:1 to about 10:1, from about 5:1 to about 15:1, from about 5:1 to about 20:1, from about 5:1 to about 25:1, from about 5:1 to about 30:1, from about 5:1 to about 35:1, from about 5:1 to about 40:1, from about 5:1 to about 45:1, from about 5:1 to about 50:1, from about 5:1 to about 55:1, from about 5:1 to about 60:1, from about 5:1 to about 70:1, from about 10:1 to about 15:1, from about 10:1 to about 20:1, from about 10:1 to about 25:1, from about 10:1 to about 30:1, from about 10:1 to about 35:1, from about 10:1 to about 40:1, from about 10:1 to about 45:1, from about 10:1 to about 50:1, from about 10:1 to about 55:1, from about 10:1 to about 60:1, from about 10:1 to about 70:1, from about 15:1 to about 20:1, from about 15:1 to about 25:1,from about 15:1 to about 30:1, from about 15:1 to about 35:1, from about 15:1 to about 40:1, from about 15:1 to about 45:1, from about 15:1 to about 50:1, from about 15:1 to about 55:1, from about 15:1 to about 60:1 or from about 15:1 to about 70:1.

[0610] In one embodiment, the lipid nanoparticles described herein can comprise the polynucleotide in a concentration from approximately 0.1 mg/ml to 2 mg/ml such as, but not limited to, 0.1 mg/ml, 0.2 mg/ml, 0.3 mg/ml, 0.4 mg/ml, 0.5 mg/ml, 0.6 mg/ml, 0.7 mg/ml, 0.8 mg/ml, 0.9 mg/ml, 1.0 mg/ml, 1.1 mg/ml, 1.2 mg/ml, 1.3 mg/ml, 1.4 mg/ml, 1.5 mg/ml, 1.6 mg/ml, 1.7 mg/ml, 1.8 mg/ml, 1.9 mg/ml, 2.0 mg/ml or greater than 2.0 mg/ml.

(vii) Nanoparticle Compositions

[0611] In some embodiments, the pharmaceutical compositions disclosed herein are formulated as lipid nanoparticles (LNP). Accordingly, the present disclosure also provides nanoparticle compositions comprising (i) a lipid composition comprising a delivery agent such as compound as described herein, and (ii) a polynucleotide encoding a G6PC polypeptide. In such nanoparticle composition, the lipid composition disclosed herein can encapsulate the polynucleotide encoding a G6PC polypeptide.

[0612] Nanoparticle compositions are typically sized on the order of micrometers or smaller and can include a lipid bilayer. Nanoparticle compositions encompass lipid nanoparticles (LNPs), liposomes (e.g., lipid vesicles), and lipoplexes. For example, a nanoparticle composition can be a liposome having a lipid bilayer with a diameter of 500 nm or less.

[0613] Nanoparticle compositions include, for example, lipid nanoparticles (LNPs), liposomes, and lipoplexes. In some embodiments, nanoparticle compositions are vesicles including one or more lipid bilayers. In certain embodiments, a nanoparticle composition includes two or more concentric bilayers separated by aqueous compartments. Lipid bilayers can be functionalized and/or crosslinked to one another. Lipid bilayers can include one or more ligands, proteins, or channels.

[0614] In one embodiment, a lipid nanoparticle comprises an ionizable lipid, a structural lipid, a phospholipid, and mRNA. In some embodiments, the LNP comprises an ionizable lipid, a PEG-modified lipid, a sterol and a structural lipid. In some embodiments, the LNP has a molar ratio of about 20-60% ionizable lipid: about 5-25% structural lipid: about 25-55% sterol; and about 0.5-15% PEG-modified lipid.

[0615] In some embodiments, the LNP has a polydispersity value of less than 0.4. In some embodiments, the LNP has a net neutral charge at a neutral pH. In some embodiments, the LNP has a mean diameter of 50-150 nm. In some embodiments, the LNP has a mean diameter of 80-100 nm.

[0616] As generally defined herein, the term "lipid" refers to a small molecule that has hydrophobic or amphiphilic properties. Lipids may be naturally occurring or synthetic. Examples of classes of lipids include, but are not limited to, fats, waxes, sterol-containing metabolites, vitamins, fatty acids, glycerolipids, glycerophospholipids, sphingolipids, saccharolipids, and polyketides, and prenol

lipids. In some instances, the amphiphilic properties of some lipids lead them to form liposomes, vesicles, or membranes in aqueous media.

[0617] In some embodiments, a lipid nanoparticle (LNP) may comprise an ionizable lipid. As used herein, the term "ionizable lipid" has its ordinary meaning in the art and may refer to a lipid comprising one or more charged moieties. In some embodiments, an ionizable lipid may be positively charged or negatively charged. An ionizable lipid may be positively charged, in which case it can be referred to as "cationic lipid". In certain embodiments, an ionizable lipid molecule may comprise an amine group, and can be referred to as an ionizable amino lipid. As used herein, a "charged moiety" is a chemical moiety that carries a formal electronic charge, e.g., monovalent (+1, or -1), divalent (+2, or -2), trivalent (+3, or -3), etc. The charged moiety may be anionic (i.e., negatively charged) or cationic (i.e., positively charged). Examples of positively-charged moieties include amine groups (e.g., primary, secondary, and/or tertiary amines), ammonium groups, pyridinium group, guanidine groups, and imidizolium groups. In a particular embodiment, the charged moieties comprise amine groups. Examples of negatively- charged groups or precursors thereof, include carboxylate groups, sulfonate groups, sulfate groups, phosphonate groups, phosphate groups, hydroxyl groups, and the like. The charge of the charged moiety may vary, in some cases, with the environmental conditions, for example, changes in pH may alter the charge of the moiety, and/or cause the moiety to become charged or uncharged. In general, the charge density of the molecule may be selected as desired.

[0618] It should be understood that the terms "charged" or "charged moiety" does not refer to a "partial negative charge" or "partial positive charge" on a molecule. The terms "partial negative charge" and "partial positive charge" are given its ordinary meaning in the art. A "partial negative charge" may result when a functional group comprises a bond that becomes polarized such that electron density is pulled toward one atom of the bond, creating a partial negative charge on the atom. Those of ordinary skill in the art will, in general, recognize bonds that can become polarized in this way.

[0619] In some embodiments, the ionizable lipid is an ionizable amino lipid, sometimes referred to in the art as an "ionizable cationic lipid". In one embodiment, the ionizable amino lipid may have a positively charged hydrophilic head and a hydrophobic tail that are connected via a linker structure.

[0620] In addition to these, an ionizable lipid may also be a lipid including a cyclic amine group.

[0621] In one embodiment, the ionizable lipid may be selected from, but not limited to, a ionizable lipid described in International Publication Nos. WO2013086354 and WO2013116126; the contents of each of which are herein incorporated by reference in their entirety.

[0622] In yet another embodiment, the ionizable lipid may be selected from, but not limited to, formula CLI-CLXXXXII of US Patent No. 7,404,969; each of which is herein incorporated by reference in their entirety.

[0623] In one embodiment, the lipid may be a cleavable lipid such as those described in International Publication No. WO2012170889, herein incorporated by reference in its entirety. In one embodiment, the lipid may be synthesized by methods known in the art and/or as described in International Publication Nos. WO2013086354; the contents of each of which are herein incorporated by reference in their entirety.

[0624] Nanoparticle compositions can be characterized by a variety of methods. For example, microscopy (e.g., transmission electron microscopy or scanning electron microscopy) can be used to examine the morphology and size distribution of a nanoparticle composition. Dynamic light scattering or potentiometry (e.g., potentiometric titrations) can be used to measure zeta potentials. Dynamic light scattering can also be utilized to determine particle sizes. Instruments such as the Zetasizer Nano ZS (Malvern Instruments Ltd, Malvern, Worcestershire, UK) can also be used to measure multiple characteristics of a nanoparticle composition, such as particle size, polydispersity index, and zeta potential.

[0625] The size of the nanoparticles can help counter biological reactions such as, but not limited to, inflammation, or can increase the biological effect of the polynucleotide.

[0626] As used herein, "size" or "mean size" in the context of nanoparticle compositions refers to the mean diameter of a nanoparticle composition.

[0627] In one embodiment, the polynucleotide encoding a G6PC polypeptide are formulated in lipid nanoparticles having a diameter from about 10 to about 100 nm such as, but not limited to, about 10 to about 20 nm, about 10 to about 30 nm, about 10 to about 40 nm, about 10 to about 50 nm, about 10 to about 60 nm, about 10 to about 70 nm, about 10 to about 80 nm, about 10 to about 90 nm, about 20 to about 30 nm, about 20 to about 40 nm, about 20 to about 50 nm, about 20 to about 60 nm,

about 20 to about 70 nm, about 20 to about 80 nm, about 20 to about 90 nm, about 20 to about 100 nm, about 30 to about 40 nm, about 30 to about 50 nm, about 30 to about 60 nm, about 30 to about 70 nm, about 30 to about 80 nm, about 30 to about 90 nm, about 30 to about 100 nm, about 40 to about 50 nm, about 40 to about 60 nm, about 40 to about 70 nm, about 40 to about 80 nm, about 40 to about 90 nm, about 40 to about 100 nm, about 50 to about 60 nm, about 50 to about 70 nm, about 50 to about 80 nm, about 50 to about 90 nm, about 50 to about 100 nm, about 60 to about 70 nm, about 60 to about 80 nm, about 60 to about 90 nm, about 60 to about 100 nm, about 70 to about 80 nm, about 70 to about 90 nm, about 70 to about 100 nm, about 80 to about 90 nm, about 80 to about 100 nm and/or about 90 to about 100 nm.

[0628] In one embodiment, the nanoparticles have a diameter from about 10 to 500 nm. In one embodiment, the nanoparticle has a diameter greater than 100 nm, greater than 150 nm, greater than 200 nm, greater than 250 nm, greater than 300 nm, greater than 350 nm, greater than 400 nm, greater than 450 nm, greater than 500 nm, greater than 550 nm, greater than 600 nm, greater than 650 nm, greater than 700 nm, greater than 750 nm, greater than 800 nm, greater than 850 nm, greater than 900 nm, greater than 950 nm or greater than 1000 nm.

[0629] In some embodiments, the largest dimension of a nanoparticle composition is 1 μm or shorter (e.g., 1 μm, 900 nm, 800 nm, 700 nm, 600 nm, 500 nm, 400 nm, 300 nm, 200 nm, 175 nm, 150 nm, 125 nm, 100 nm, 75 nm, 50 nm, or shorter).

[0630] A nanoparticle composition can be relatively homogenous. A polydispersity index can be used to indicate the homogeneity of a nanoparticle composition, e.g., the particle size distribution of the nanoparticle composition. A small (e.g., less than 0.3) polydispersity index generally indicates a narrow particle size distribution. A nanoparticle composition can have a polydispersity index from about 0 to about 0.25, such as 0.01, 0.02, 0.03, 0.04, 0.05, 0.06, 0.07, 0.08, 0.09, 0.10, 0.11, 0.12, 0.13, 0.14, 0.15, 0.16, 0.17, 0.18, 0.19, 0.20, 0.21, 0.22, 0.23, 0.24, or 0.25. In some embodiments, the polydispersity index of a nanoparticle composition disclosed herein can be from about 0.10 to about 0.20.

[0631] The zeta potential of a nanoparticle composition can be used to indicate the electrokinetic potential of the composition. For example, the zeta potential can describe the surface charge of a nanoparticle composition. Nanoparticle compositions with relatively low charges, positive or negative, are generally desirable, as more highly charged species can interact undesirably with cells, tissues, and other elements

in the body. In some embodiments, the zeta potential of a nanoparticle composition disclosed herein can be from about -10 mV to about +20 mV, from about -10 mV to about +15 mV, from about 10 mV to about +10 mV, from about -10 mV to about +5 mV, from about -10 mV to about 0 mV, from about -10 mV to about -5 mV, from about -5 mV to about +20 mV, from about -5 mV to about +15 mV, from about -5 mV to about +10 mV, from about -5 mV to about +5 mV, from about -5 mV to about 0 mV, from about 0 mV to about +20 mV, from about 0 mV to about +15 mV, from about 0 mV to about +10 mV, from about 0 mV to about +5 mV, from about +5 mV to about +20 mV, from about +5 mV to about +15 mV, or from about +5 mV to about +10 mV.

[0632] In some embodiments, the zeta potential of the lipid nanoparticles can be from about 0 mV to about 100 mV, from about 0 mV to about 90 mV, from about 0 mV to about 80 mV, from about 0 mV to about 70 mV, from about 0 mV to about 60 mV, from about 0 mV to about 50 mV, from about 0 mV to about 40 mV, from about 0 mV to about 30 mV, from about 0 mV to about 20 mV, from about 0 mV to about 10 mV, from about 10 mV to about 100 mV, from about 10 mV to about 90 mV, from about 10 mV to about 80 mV, from about 10 mV to about 70 mV, from about 10 mV to about 60 mV, from about 10 mV to about 50 mV, from about 10 mV to about 40 mV, from about 10 mV to about 30 mV, from about 10 mV to about 20 mV, from about 20 mV to about 100 mV, from about 20 mV to about 90 mV, from about 20 mV to about 80 mV, from about 20 mV to about 70 mV, from about 20 mV to about 60 mV, from about 20 mV to about 50 mV, from about 20 mV to about 40 mV, from about 20 mV to about 30 mV, from about 30 mV to about 100 mV, from about 30 mV to about 90 mV, from about 30 mV to about 80 mV, from about 30 mV to about 70 mV, from about 30 mV to about 60 mV, from about 30 mV to about 50 mV, from about 30 mV to about 40 mV, from about 40 mV to about 100 mV, from about 40 mV to about 90 mV, from about 40 mV to about 80 mV, from about 40 mV to about 70 mV, from about 40 mV to about 60 mV, and from about 40 mV to about 50 mV. In some embodiments, the zeta potential of the lipid nanoparticles can be from about 10 mV to about 50 mV, from about 15 mV to about 45 mV, from about 20 mV to about 40 mV, and from about 25 mV to about 35 mV. In some embodiments, the zeta potential of the lipid nanoparticles can be about 10 mV, about 20 mV, about 30 mV, about 40 mV, about 50 mV, about 60 mV, about 70 mV, about 80 mV, about 90 mV, and about 100 mV.

[0633] The term "encapsulation efficiency" of a polynucleotide describes the amount of the polynucleotide that is encapsulated by or otherwise associated with a nanoparticle composition after preparation, relative to the initial amount provided. As used herein, "encapsulation" can refer to complete, substantial, or partial enclosure, confinement, surrounding, or encasement.

[0634] Encapsulation efficiency is desirably high (e.g., close to 100%). The encapsulation efficiency can be measured, for example, by comparing the amount of the polynucleotide in a solution containing the nanoparticle composition before and after breaking up the nanoparticle composition with one or more organic solvents or detergents.

[0635] Fluorescence can be used to measure the amount of free polynucleotide in a solution. For the nanoparticle compositions described herein, the encapsulation efficiency of a polynucleotide can be at least 50%, for example 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100%. In some embodiments, the encapsulation efficiency can be at least 80%. In certain embodiments, the encapsulation efficiency can be at least 90%.

[0636] The amount of a polynucleotide present in a pharmaceutical composition disclosed herein can depend on multiple factors such as the size of the polynucleotide, desired target and/or application, or other properties of the nanoparticle composition as well as on the properties of the polynucleotide.

[0637] For example, the amount of an mRNA useful in a nanoparticle composition can depend on the size (expressed as length, or molecular mass), sequence, and other characteristics of the mRNA. The relative amounts of a polynucleotide in a nanoparticle composition can also vary.

[0638] The relative amounts of the lipid composition and the polynucleotide present in a lipid nanoparticle composition of the present disclosure can be optimized according to considerations of efficacy and tolerability. For compositions including an mRNA as a polynucleotide, the N:P ratio can serve as a useful metric.

[0639] As the N:P ratio of a nanoparticle composition controls both expression and tolerability, nanoparticle compositions with low N:P ratios and strong expression are desirable. N:P ratios vary according to the ratio of lipids to RNA in a nanoparticle composition.

[0640] In general, a lower N:P ratio is preferred. The one or more RNA, lipids, and amounts thereof can be selected to provide an N:P ratio from about 2:1 to about 30:1,

such as 2:1, 3:1, 4:1, 5:1, 6:1, 7:1, 8:1, 9:1, 10:1, 12:1, 14:1, 16:1, 18:1, 20:1, 22:1, 24:1, 26:1, 28:1, or 30:1. In certain embodiments, the N:P ratio can be from about 2:1 to about 8:1. In other embodiments, the N:P ratio is from about 5:1 to about 8:1. In certain embodiments, the N:P ratio is between 5:1 and 6:1. In one specific aspect, the N:P ratio is about is about 5.67:1.

[0641] In addition to providing nanoparticle compositions, the present disclosure also provides methods of producing lipid nanoparticles comprising encapsulating a polynucleotide. Such method comprises using any of the pharmaceutical compositions disclosed herein and producing lipid nanoparticles in accordance with methods of production of lipid nanoparticles known in the art. *See*, e.g., Wang et al. (2015) "Delivery of oligonucleotides with lipid nanoparticles" Adv. Drug Deliv. Rev. 87:68-80; Silva et al. (2015) "Delivery Systems for Biopharmaceuticals. Part I: Nanoparticles and Microparticles" Curr. Pharm. Technol. 16: 940-954; Naseri et al. (2015) "Solid Lipid Nanoparticles and Nanostructured Lipid Carriers: Structure, Preparation and Application" Adv. Pharm. Bull. 5:305-13; Silva et al. (2015) "Lipid nanoparticles for the delivery of biopharmaceuticals" Curr. Pharm. Biotechnol. 16:291-302, and references cited therein.

21. Other Delivery Agents

a. Liposomes, Lipoplexes, and Lipid Nanoparticles

[0642] In some embodiments, the compositions or formulations of the present disclosure comprise a delivery agent, e.g., a liposome, a lipolexes, a lipid nanoparticle, or any combination thereof. The polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) can be formulated using one or more liposomes, lipoplexes, or lipid nanoparticles. Liposomes, lipoplexes, or lipid nanoparticles can be used to improve the efficacy of the polynucleotides directed protein production as these formulations can increase cell transfection by the polynucleotide; and/or increase the translation of encoded protein. The liposomes, lipoplexes, or lipid nanoparticles can also be used to increase the stability of the polynucleotides.

[0643] Liposomes are artificially-prepared vesicles that can primarily be composed of a lipid bilayer and can be used as a delivery vehicle for the administration of

pharmaceutical formulations. Liposomes can be of different sizes. A multilamellar vesicle (MLV) can be hundreds of nanometers in diameter, and can contain a series of concentric bilayers separated by narrow aqueous compartments. A small unicellular vesicle (SUV) can be smaller than 50 nm in diameter, and a large unilamellar vesicle (LUV) can be between 50 and 500 nm in diameter. Liposome design can include, but is not limited to, opsonins or ligands to improve the attachment of liposomes to unhealthy tissue or to activate events such as, but not limited to, endocytosis. Liposomes can contain a low or a high pH value to improve the delivery of the pharmaceutical formulations.

[0644] The formation of liposomes can depend on the pharmaceutical formulation entrapped and the liposomal ingredients, the nature of the medium in which the lipid vesicles are dispersed, the effective concentration of the entrapped substance and its potential toxicity, any additional processes involved during the application and/or delivery of the vesicles, the optimal size, polydispersity and the shelf-life of the vesicles for the intended application, and the batch-to-batch reproducibility and scale up production of safe and efficient liposomal products, etc.

[0645] As a non-limiting example, liposomes such as synthetic membrane vesicles can be prepared by the methods, apparatus and devices described in U.S. Pub. Nos. US20130177638, US20130177637, US20130177636, US20130177635, US20130177634, US20130177633, US20130183375, US20130183373, and US20130183372. In some embodiments, the polynucleotides described herein can be encapsulated by the liposome and/or it can be contained in an aqueous core that can then be encapsulated by the liposome as described in, e.g., Intl. Pub. Nos. WO2012031046, WO2012031043, WO2012030901, WO2012006378, and WO2013086526; and U.S. Pub.Nos. US20130189351, US20130195969 and US20130202684. Each of the references in herein incorporated by reference in its entirety.

[0646] In some embodiments, the polynucleotides described herein can be formulated in a cationic oil-in-water emulsion where the emulsion particle comprises an oil core and a cationic lipid that can interact with the polynucleotide anchoring the molecule to the emulsion particle. In some embodiments, the polynucleotides described herein can be formulated in a water-in-oil emulsion comprising a continuous hydrophobic phase in which the hydrophilic phase is dispersed. Exemplary emulsions can be made

by the methods described in Intl. Pub. Nos. WO2012006380 and WO201087791, each of which is herein incorporated by reference in its entirety.

[0647] In some embodiments, the polynucleotides described herein can be formulated in a lipid-polycation complex. The formation of the lipid-polycation complex can be accomplished by methods as described in, e.g., U.S. Pub. No. US20120178702. As a non-limiting example, the polycation can include a cationic peptide or a polypeptide such as, but not limited to, polylysine, polyornithine and/or polyarginine and the cationic peptides described in Intl. Pub. No. WO2012013326 or U.S. Pub. No. US20130142818. Each of the references is herein incorporated by reference in its entirety.

[0648] In some embodiments, the polynucleotides described herein can be formulated in a lipid nanoparticle (LNP) such as those described in Intl. Pub. Nos. WO2013123523, WO2012170930, WO2011127255 and WO2008103276; and U.S. Pub. No. US20130171646, each of which is herein incorporated by reference in its entirety.

[0649] Lipid nanoparticle formulations typically comprise one or more lipids. In some embodiments, the lipid is an ionizable lipid (*e.g.*, an ionizable amino lipid), sometimes referred to in the art as an "ionizable cationic lipid". In some embodiments, lipid nanoparticle formulations further comprise other components, including a phospholipid, a structural lipid, and a molecule capable of reducing particle aggregation, for example a PEG or PEG-modified lipid.

[0650] Exemplary ionizable lipids include, but not limited to, any one of Compounds 1-342 disclosed herein, DLin-MC3-DMA (MC3), DLin-DMA, DLenDMA, DLin-D-DMA, DLin-K-DMA, DLin-M-C2-DMA, DLin-K-DMA, DLin-KC2-DMA, DLin-KC3-DMA, DLin-KC4-DMA, DLin-C2K-DMA, DLin-MP-DMA, DODMA, 98N12-5, C12-200, DLin-C-DAP, DLin-DAC, DLinDAP, DLinAP, DLin-EG-DMA, DLin-2-DMAP, KL10, KL22, KL25, Octyl-CLinDMA, Octyl-CLinDMA (2R), Octyl-CLinDMA (2S), and any combination thereof. Other exemplary ionizable lipids include, (13Z,16Z)-N,N-dimethyl-3-nonyldocosa-13,16-dien-1-amine (L608), (20Z,23Z)-N,N-dimethylnonacosa-20,23-dien-10-amine, (17Z,20Z)-N,N-dimemylhexacosa-17,20-dien-9-amine, (16Z,19Z)-N5N-dimethylpentacosa-16,19-dien-8-amine, (13Z,16Z)-N,N-dimethyldocosa-13,16-dien-5-amine, (12Z,15Z)-N,N-dimethylhenicosa-12,15-dien-4-amine, (14Z,17Z)-N,N-dimethyltricosa-14,17-dien-6-amine, (15Z,18Z)-N,N-dimethyltetracosa-15,18-dien-7-amine, (18Z,21Z)-N,N-

dimethylheptacosa-18,21-dien-10-amine, (15Z,18Z)-N,N-dimethyltetracosa-15,18-dien-5-amine, (14Z,17Z)-N,N-dimethyltricosa-14,17-dien-4-amine, (19Z,22Z)-N,N-dimeihyloctacosa-19,22-dien-9-amine, (18Z,21Z)-N,N-dimethylheptacosa-18,21-dien-8-amine, (17Z,20Z)-N,N-dimethylhexacosa-17,20-dien-7-amine, (16Z,19Z)-N,N-dimethylpentacosa-16,19-dien-6-amine, (22Z,25Z)-N,N-dimethylhentriaconta-22,25-dien-10-amine, (21Z,24Z)-N,N-dimethyltriaconta-21,24-dien-9-amine, (18Z)-N,N-dimetylheptacos-18-en-10-amine, (17Z)-N,N-dimethylhexacos-17-en-9-amine, (19Z,22Z)-N,N-dimethyloctacosa-19,22-dien-7-amine, N,N-dimethylheptacosan-10-amine, (20Z,23Z)-N-ethyl-N-methylnonacosa-20,23-dien-10-amine, 1-[(11Z,14Z)-l-nonylicosa-11,14-dien-l-yl]pyrrolidine, (20Z)-N,N-dimethylheptacos-20-en-10-amine, (15Z)-N,N-dimethyl eptacos-15-en-10-amine, (14Z)-N,N-dimethylnonacos-14-en-10-amine, (17Z)-N,N-dimethylnonacos-17-en-10-amine, (24Z)-N,N-dimethyltritriacont-24-en-10-amine, (20Z)-N,N-dimethylnonacos-20-en-10-amine, (22Z)-N,N-dimethylhentriacont-22-en-10-amine, (16Z)-N,N-dimethylpentacos-16-en-8-amine, (12Z,15Z)-N,N-dimethyl-2-nonylhenicosa-12,15-dien-1-amine, N,N-dimethyl-l-[(lS,2R)-2-octylcyclopropyl] eptadecan-8-amine, l-[(1S,2R)-2-hexylcyclopropyl]-N,N-dimethylnonadecan-10-amine, N,N-dimethyl-1-[(1S,2R)-2-octylcyclopropyl]nonadecan-10-amine, N,N-dimethyl-21-[(1S,2R)-2-octylcyclopropyl]henicosan-10-amine, N,N-dimethyl-l-[(lS,2S)-2-{[(1R,2R)-2-pentylcycIopropyl]methyl}cyclopropyl]nonadecan-10-amine, N,N-dimethyl-l-[(1S,2R)-2-octylcyclopropyl]hexadecan-8-amine, N,N-dimethyl-[(lR,2S)-2-undecyIcyclopropyl]tetradecan-5-amine, N,N-dimethyl-3-{7-[(lS,2R)-2-octylcyclopropyl]heptyl}dodecan-1-amine, 1-[(1R,2S)-2-heptylcyclopropyl]-N,N-dimethyloctadecan-9-amine, 1-[(1S,2R)-2-decylcyclopropyl]-N,N-dimethylpentadecan-6-amine, N,N-dimethyl-l-[(lS,2R)-2-octylcyclopropyl]pentadecan-8-amine, R-N,N-dimethyl-1-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]-3-(octyloxy)propan-2-amine, S-N,N-dimethyl-1-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]-3-(octyloxy)propan-2-amine, 1-{2-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]-1-[(octyloxy)methyl]ethyl}pyrrolidine, (2S)-N,N-dimethyl-1-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]-3-[(5Z)-oct-5-en-1-yloxy]propan-2-amine, 1-{2-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]-1-[(octyloxy)methyl]ethyl}azetidine, (2S)-1-(hexyloxy)-N,N-dimethyl-3-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]propan-2-amine, (2S)-1-(heptyloxy)-N,N-dimethyl-3-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]propan-2-amine, N,N-dimethyl-1-(nonyloxy)-3-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]propan-

2-amine, N,N-dimethyl-1-[(9Z)-octadec-9-en-l-yloxy]-3-(octyloxy)propan-2-amine; (2S)-N,N-dimethyl-l-[(6Z,9Z,12Z)-octadeca-6,9,12-trien-l-yloxy]-3-(octyloxy)propan-2-amine, (2S)-1-[(11Z,14Z)-icosa-11,14-dien-1-yloxy]-N,N-dimethyl-3-(pentyloxy)propan-2-amine, (2S)-1-(hexyloxy)-3-[(11Z,14Z)-icosa-11,14-dien-l-yloxy]-N,N-dimethylpropan-2-amine, 1-[(11Z,14Z)-icosa-11,14-dien-l-yloxy]-N,N-dimethyl-3-(octyloxy)propan-2-amine, 1-[(13Z,16Z)-docosa-13,16-dien-l-yloxy]-N,N-dimethyl-3-(octyloxy)propan-2-amine, (2S)-1-[(13Z, 16Z)-docosa-13,16-dien-1-yloxy]-3-(hexyloxy)-N,N-dimethylpropan-2-amine, (2S)-1-[(13Z)-docos-13-en-1-yloxy]-3-(hexyloxy)-N,N-dimethylpropan-2-amine, 1-[(13Z)-docos-13-en-1-yloxy]-N,N-dimethyl-3-(octyloxy)propan-2-amine, 1-[(9Z)-hexadec-9-en-1-yloxy]-N,N-dimethyl-3-(octyloxy)propan-2-amine, (2R)-N,N-dimethyl-H(1-metoyloctyl)oxy]-3-[(9Z,12Z)-octadeca-9,12-dien-1-yloxy]propan-2-amine, (2R)-1-[(3,7-dimethyloctyl)oxy]-N,N-dimethyl-3-[(9Z,12Z)-octadeca-9,12-dien-l-yloxy]propan-2-amine, N,N-dimethyl-1-(octyloxy)-3-({8-[(1S,2S)-2-{[(1R,2R)-2-pentylcyclopropyl]methyl}cyclopropyl]octyl}oxy)propan-2-amine, N,N-dimethyl-1-{[8-(2-oclylcyclopropyl)octyl]oxy}-3-(octyloxy)propan-2-amine, and (11E,20Z,23Z)-N,N-dimethylnonacosa-11,20,2-trien-10-amine, and any combination thereof.

[0651] Phospholipids include, but are not limited to, glycerophospholipids such as phosphatidylcholines, phosphatidylethanolamines, phosphatidylserines, phosphatidylinositols, phosphatidy glycerols, and phosphatidic acids. Phospholipids also include phosphosphingolipid, such as sphingomyelin. In some embodiments, the phospholipids are DLPC, DMPC, DOPC, DPPC, DSPC, DUPC, 18:0 Diether PC, DLnPC, DAPC, DHAPC, DOPE, 4ME 16:0 PE, DSPE, DLPE, DLnPE, DAPE, DHAPE, DOPG, and any combination thereof. In some embodiments, the phospholipids are MPPC, MSPC, PMPC, PSPC, SMPC, SPPC, DHAPE, DOPG, and any combination thereof. In some embodiments, the amount of phospholipids (e.g., DSPC) in the lipid composition ranges from about 1 mol% to about 20 mol%.

[0652] The structural lipids include sterols and lipids containing sterol moieties. In some embodiments, the structural lipids include cholesterol, fecosterol, sitosterol, ergosterol, campesterol, stigmasterol, brassicasterol, tomatidine, tomatine, ursolic acid, alpha-tocopherol, and mixtures thereof. In some embodiments, the structural lipid is cholesterol. In some embodiments, the amount of the structural lipids (e.g., cholesterol) in the lipid composition ranges from about 20 mol% to about 60 mol%.

[0653] The PEG-modified lipids include PEG-modified phosphatidylethanolamine and phosphatidic acid, PEG-ceramide conjugates (e.g., PEG-CerC14 or PEG-CerC20), PEG-modified dialkylamines and PEG-modified 1,2-diacyloxypropan-3-amines. Such lipids are also referred to as PEGylated lipids. For example, a PEG lipid can be PEG-c-DOMG, PEG-DMG, PEG-DLPE, PEG DMPE, PEG-DPPC, or a PEG-DSPE lipid. In some embodiments, the PEG-lipid are 1,2-dimyristoyl-sn-glycerol methoxypolyethylene glycol (PEG-DMG), 1,2-distearoyl-sn-glycero-3-phosphoethanolamine-N-[amino(polyethylene glycol)] (PEG-DSPE), PEG-disteryl glycerol (PEG-DSG), PEG-dipalmetoleyl, PEG-dioleyl, PEG-distearyl, PEG-diacylglycamide (PEG-DAG), PEG-dipalmitoyl phosphatidylethanolamine (PEG-DPPE), or PEG-l,2-dimyristyloxlpropyl-3-amine (PEG-c-DMA). In some embodiments, the PEG moiety has a size of about 1000, 2000, 5000, 10,000, 15,000 or 20,000 daltons. In some embodiments, the amount of PEG-lipid in the lipid composition ranges from about 0 mol% to about 5 mol%.

[0654] In some embodiments, the LNP formulations described herein can additionally comprise a permeability enhancer molecule. Non-limiting permeability enhancer molecules are described in U.S. Pub. No. US20050222064, herein incorporated by reference in its entirety.

[0655] The LNP formulations can further contain a phosphate conjugate. The phosphate conjugate can increase in vivo circulation times and/or increase the targeted delivery of the nanoparticle. Phosphate conjugates can be made by the methods described in, e.g., Intl. Pub. No. WO2013033438 or U.S. Pub. No. US20130196948. The LNP formulation can also contain a polymer conjugate (e.g., a water-soluble conjugate) as described in, e.g., U.S. Pub. Nos. US20130059360, US20130196948, and US20130072709. Each of the references is herein incorporated by reference in its entirety.

[0656] The LNP formulations can comprise a conjugate to enhance the delivery of nanoparticles of the present invention in a subject. Further, the conjugate can inhibit phagocytic clearance of the nanoparticles in a subject. In some embodiments, the conjugate can be a "self" peptide designed from the human membrane protein CD47 (e.g., the "self" particles described by Rodriguez et al, Science 2013 339, 971-975, herein incorporated by reference in its entirety). As shown by Rodriguez et al. the self-peptides delayed macrophage-mediated clearance of nanoparticles which enhanced delivery of the nanoparticles.

[0657] The LNP formulations can comprise a carbohydrate carrier. As a non-limiting example, the carbohydrate carrier can include, but is not limited to, an anhydride-modified phytoglycogen or glycogen-type material, phytoglycogen octenyl succinate, phytoglycogen beta-dextrin, anhydride-modified phytoglycogen beta-dextrin (e.g., Intl. Pub. No. WO2012109121, herein incorporated by reference in its entirety).

[0658] The LNP formulations can be coated with a surfactant or polymer to improve the delivery of the particle. In some embodiments, the LNP can be coated with a hydrophilic coating such as, but not limited to, PEG coatings and/or coatings that have a neutral surface charge as described in U.S. Pub. No. US20130183244, herein incorporated by reference in its entirety.

[0659] The LNP formulations can be engineered to alter the surface properties of particles so that the lipid nanoparticles can penetrate the mucosal barrier as described in U.S. Pat. No. 8,241,670 or Intl. Pub. No. WO2013110028, each of which is herein incorporated by reference in its entirety.

[0660] The LNP engineered to penetrate mucus can comprise a polymeric material (i.e., a polymeric core) and/or a polymer-vitamin conjugate and/or a tri-block co-polymer. The polymeric material can include, but is not limited to, polyamines, polyethers, polyamides, polyesters, polycarbamates, polyureas, polycarbonates, poly(styrenes), polyimides, polysulfones, polyurethanes, polyacetylenes, polyethylenes, polyethyeneimines, polyisocyanates, polyacrylates, polymethacrylates, polyacrylonitriles, and polyarylates.

[0661] LNP engineered to penetrate mucus can also include surface altering agents such as, but not limited to, polynucleotides, anionic proteins (e.g., bovine serum albumin), surfactants (e.g., cationic surfactants such as for example dimethyldioctadecyl-ammonium bromide), sugars or sugar derivatives (e.g., cyclodextrin), nucleic acids, polymers (e.g., heparin, polyethylene glycol and poloxamer), mucolytic agents (e.g., N-acetylcysteine, mugwort, bromelain, papain, clerodendrum, acetylcysteine, bromhexine, carbocisteine, eprazinone, mesna, ambroxol, sobrerol, domiodol, letosteine, stepronin, tiopronin, gelsolin, thymosin β4 dornase alfa, neltenexine, erdosteine) and various DNases including rhDNase.

[0662] In some embodiments, the mucus penetrating LNP can be a hypotonic formulation comprising a mucosal penetration enhancing coating. The formulation can be hypotonic for the epithelium to which it is being delivered. Non-limiting

examples of hypotonic formulations can be found in, e.g., Intl. Pub. No. WO2013110028, herein incorporated by reference in its entirety.

[0663] In some embodiments, the polynucleotide described herein is formulated as a lipoplex, such as, without limitation, the ATUPLEXTM system, the DACC system, the DBTC system and other siRNA-lipoplex technology from Silence Therapeutics (London, United Kingdom), STEMFECTTM from STEMGENT® (Cambridge, MA), and polyethylenimine (PEI) or protamine-based targeted and non-targeted delivery of nucleic acids (Aleku et al. Cancer Res. 2008 68:9788-9798; Strumberg et al. Int J Clin Pharmacol Ther 2012 50:76-78; Santel et al., Gene Ther 2006 13:1222-1234; Santel et al., Gene Ther 2006 13:1360-1370; Gutbier et al., Pulm Pharmacol. Ther. 2010 23:334-344; Kaufmann et al. Microvasc Res 2010 80:286-293Weide et al. J Immunother. 2009 32:498-507; Weide et al. J Immunother. 2008 31:180-188; Pascolo Expert Opin. Biol. Ther. 4:1285-1294; Fotin-Mleczek et al., 2011 J. Immunother. 34:1-15; Song et al., Nature Biotechnol. 2005, 23:709-717; Peer et al., Proc Natl Acad Sci U S A. 2007 6;104:4095-4100; deFougerolles Hum Gene Ther. 2008 19:125-132; all of which are incorporated herein by reference in its entirety).

[0664] In some embodiments, the polynucleotides described herein are formulated as a solid lipid nanoparticle (SLN), which can be spherical with an average diameter between 10 to 1000 nm. SLN possess a solid lipid core matrix that can solubilize lipophilic molecules and can be stabilized with surfactants and/or emulsifiers. Exemplary SLN can be those as described in Intl. Pub. No. WO2013105101, herein incorporated by reference in its entirety.

[0665] In some embodiments, the polynucleotides described herein can be formulated for controlled release and/or targeted delivery. As used herein, "controlled release" refers to a pharmaceutical composition or compound release profile that conforms to a particular pattern of release to effect a therapeutic outcome. In one embodiment, the polynucleotides can be encapsulated into a delivery agent described herein and/or known in the art for controlled release and/or targeted delivery. As used herein, the term "encapsulate" means to enclose, surround or encase. As it relates to the formulation of the compounds of the invention, encapsulation can be substantial, complete or partial. The term "substantially encapsulated" means that at least greater than 50, 60, 70, 80, 85, 90, 95, 96, 97, 98, 99, or greater than 99% of the pharmaceutical composition or compound of the invention can be enclosed, surrounded or encased within the delivery agent. "Partially encapsulation" means that

less than 10, 10, 20, 30, 40 50 or less of the pharmaceutical composition or compound of the invention can be enclosed, surrounded or encased within the delivery agent.

[0666] Advantageously, encapsulation can be determined by measuring the escape or the activity of the pharmaceutical composition or compound of the invention using fluorescence and/or electron micrograph. For example, at least 1, 5, 10, 20, 30, 40, 50, 60, 70, 80, 85, 90, 95, 96, 97, 98, 99, 99.9, or greater than 99% of the pharmaceutical composition or compound of the invention are encapsulated in the delivery agent.

[0667] In some embodiments, the polynucleotides described herein can be encapsulated in a therapeutic nanoparticle, referred to herein as "therapeutic nanoparticle polynucleotides." Therapeutic nanoparticles can be formulated by methods described in, e.g., Intl. Pub. Nos. WO2010005740, WO2010030763, WO2010005721, WO2010005723, and WO2012054923; and U.S. Pub. Nos. US20110262491, US20100104645, US20100087337, US20100068285, US20110274759, US20100068286, US20120288541, US20120140790, US20130123351 and US20130230567; and U.S. Pat. Nos. 8,206,747, 8,293,276, 8,318,208 and 8,318,211, each of which is herein incorporated by reference in its entirety.

[0668] In some embodiments, the therapeutic nanoparticle polynucleotide can be formulated for sustained release. As used herein, "sustained release" refers to a pharmaceutical composition or compound that conforms to a release rate over a specific period of time. The period of time can include, but is not limited to, hours, days, weeks, months and years. As a non-limiting example, the sustained release nanoparticle of the polynucleotides described herein can be formulated as disclosed in Intl. Pub. No. WO2010075072 and U.S. Pub. Nos. US20100216804, US20110217377, US20120201859 and US20130150295, each of which is herein incorporated by reference in their entirety.

[0669] In some embodiments, the therapeutic nanoparticle polynucleotide can be formulated to be target specific, such as those described in Intl. Pub. Nos. WO2008121949, WO2010005726, WO2010005725, WO2011084521 and WO2011084518; and U.S. Pub. Nos. US20100069426, US20120004293 and US20100104655, each of which is herein incorporated by reference in its entirety.

[0670] The LNPs can be prepared using microfluidic mixers or micromixers. Exemplary microfluidic mixers can include, but are not limited to, a slit interdigital micromixer including, but not limited to those manufactured by Microinnova

(Allerheiligen bei Wildon, Austria) and/or a staggered herringbone micromixer (SHM) (see Zhigaltsevet al., "Bottom-up design and synthesis of limit size lipid nanoparticle systems with aqueous and triglyceride cores using millisecond microfluidic mixing," Langmuir 28:3633-40 (2012); Belliveau et al., "Microfluidic synthesis of highly potent limit-size lipid nanoparticles for in vivo delivery of siRNA," Molecular Therapy-Nucleic Acids. 1:e37 (2012); Chen et al., "Rapid discovery of potent siRNA-containing lipid nanoparticles enabled by controlled microfluidic formulation," J. Am. Chem. Soc. 134(16):6948-51 (2012); each of which is herein incorporated by reference in its entirety). Exemplary micromixers include Slit Interdigital Microstructured Mixer (SIMM-V2) or a Standard Slit Interdigital Micro Mixer (SSIMM) or Caterpillar (CPMM) or Impinging-jet (IJMM,) from the Institut für Mikrotechnik Mainz GmbH, Mainz Germany. In some embodiments, methods of making LNP using SHM further comprise mixing at least two input streams wherein mixing occurs by microstructure-induced chaotic advection (MICA). According to this method, fluid streams flow through channels present in a herringbone pattern causing rotational flow and folding the fluids around each other. This method can also comprise a surface for fluid mixing wherein the surface changes orientations during fluid cycling. Methods of generating LNPs using SHM include those disclosed in U.S. Pub. Nos. US20040262223 and US20120276209, each of which is incorporated herein by reference in their entirety.

[0671] In some embodiments, the polynucleotides described herein can be formulated in lipid nanoparticles using microfluidic technology (see Whitesides, George M., "The Origins and the Future of Microfluidics," Nature 442: 368-373 (2006); and Abraham et al., "Chaotic Mixer for Microchannels," Science 295: 647-651 (2002); each of which is herein incorporated by reference in its entirety). In some embodiments, the polynucleotides can be formulated in lipid nanoparticles using a micromixer chip such as, but not limited to, those from Harvard Apparatus (Holliston, MA) or Dolomite Microfluidics (Royston, UK). A micromixer chip can be used for rapid mixing of two or more fluid streams with a split and recombine mechanism.

[0672] In some embodiments, the polynucleotides described herein can be formulated in lipid nanoparticles having a diameter from about 1 nm to about 100 nm such as, but not limited to, about 1 nm to about 20 nm, from about 1 nm to about 30 nm, from about 1 nm to about 40 nm, from about 1 nm to about 50 nm, from about 1 nm to about 60 nm, from about 1 nm to about 70 nm, from about 1 nm to about 80 nm, from

about 1 nm to about 90 nm, from about 5 nm to about from 100 nm, from about 5 nm to about 10 nm, about 5 nm to about 20 nm, from about 5 nm to about 30 nm, from about 5 nm to about 40 nm, from about 5 nm to about 50 nm, from about 5 nm to about 60 nm, from about 5 nm to about 70 nm, from about 5 nm to about 80 nm, from about 5 nm to about 90 nm, about 10 to about 20 nm, about 10 to about 30 nm, about 10 to about 40 nm, about 10 to about 50 nm, about 10 to about 60 nm, about 10 to about 70 nm, about 10 to about 80 nm, about 10 to about 90 nm, about 20 to about 30 nm, about 20 to about 40 nm, about 20 to about 50 nm, about 20 to about 60 nm, about 20 to about 70 nm, about 20 to about 80 nm, about 20 to about 90 nm, about 20 to about 100 nm, about 30 to about 40 nm, about 30 to about 50 nm, about 30 to about 60 nm, about 30 to about 70 nm, about 30 to about 80 nm, about 30 to about 90 nm, about 30 to about 100 nm, about 40 to about 50 nm, about 40 to about 60 nm, about 40 to about 70 nm, about 40 to about 80 nm, about 40 to about 90 nm, about 40 to about 100 nm, about 50 to about 60 nm, about 50 to about 70 nm about 50 to about 80 nm, about 50 to about 90 nm, about 50 to about 100 nm, about 60 to about 70 nm, about 60 to about 80 nm, about 60 to about 90 nm, about 60 to about 100 nm, about 70 to about 80 nm, about 70 to about 90 nm, about 70 to about 100 nm, about 80 to about 90 nm, about 80 to about 100 nm and/or about 90 to about 100 nm.

[0673] In some embodiments, the lipid nanoparticles can have a diameter from about 10 to 500 nm. In one embodiment, the lipid nanoparticle can have a diameter greater than 100 nm, greater than 150 nm, greater than 200 nm, greater than 250 nm, greater than 300 nm, greater than 350 nm, greater than 400 nm, greater than 450 nm, greater than 500 nm, greater than 550 nm, greater than 600 nm, greater than 650 nm, greater than 700 nm, greater than 750 nm, greater than 800 nm, greater than 850 nm, greater than 900 nm, greater than 950 nm or greater than 1000 nm.

[0674] In some embodiments, the polynucleotides can be delivered using smaller LNPs. Such particles can comprise a diameter from below 0.1 μm up to 100 nm such as, but not limited to, less than 0.1 μm, less than 1.0 μm, less than 5μm, less than 10 μm, less than 15 um, less than 20 um, less than 25 um, less than 30 um, less than 35 um, less than 40 um, less than 50 um, less than 55 um, less than 60 um, less than 65 um, less than 70 um, less than 75 um, less than 80 um, less than 85 um, less than 90 um, less than 95 um, less than 100 um, less than 125 um, less than 150 um, less than 175 um, less than 200 um, less than 225 um, less than 250 um, less than 275 um, less than 300 um, less than 325 um, less than 350 um, less than 375 um, less than 400 um,

less than 425 um, less than 450 um, less than 475 um, less than 500 um, less than 525 um, less than 550 um, less than 575 um, less than 600 um, less than 625 um, less than 650 um, less than 675 um, less than 700 um, less than 725 um, less than 750 um, less than 775 um, less than 800 um, less than 825 um, less than 850 um, less than 875 um, less than 900 um, less than 925 um, less than 950 um, or less than 975 um.

[0675] The nanoparticles and microparticles described herein can be geometrically engineered to modulate macrophage and/or the immune response. The geometrically engineered particles can have varied shapes, sizes and/or surface charges to incorporate the polynucleotides described herein for targeted delivery such as, but not limited to, pulmonary delivery (see, e.g., Intl. Pub. No. WO2013082111, herein incorporated by reference in its entirety). Other physical features the geometrically engineering particles can include, but are not limited to, fenestrations, angled arms, asymmetry and surface roughness, charge that can alter the interactions with cells and tissues.

[0676] In some embodiment, the nanoparticles described herein are stealth nanoparticles or target-specific stealth nanoparticles such as, but not limited to, those described in U.S. Pub. No. US20130172406, herein incorporated by reference in its entirety. The stealth or target-specific stealth nanoparticles can comprise a polymeric matrix, which can comprise two or more polymers such as, but not limited to, polyethylenes, polycarbonates, polyanhydrides, polyhydroxyacids, polypropylfumerates, polycaprolactones, polyamides, polyacetals, polyethers, polyesters, poly(orthoesters), polycyanoacrylates, polyvinyl alcohols, polyurethanes, polyphosphazenes, polyacrylates, polymethacrylates, polycyanoacrylates, polyureas, polystyrenes, polyamines, polyesters, polyanhydrides, polyethers, polyurethanes, polymethacrylates, polyacrylates, polycyanoacrylates, or combinations thereof.

b. Lipidoids

[0677] In some embodiments, the compositions or formulations of the present disclosure comprise a delivery agent, e.g., a lipidoid. The polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) can be formulated with lipidoids. Complexes, micelles, liposomes or particles can be prepared containing these lipidoids and therefore to achieve an effective delivery of the polynucleotide, as judged by the production of an encoded protein, following the injection of a lipidoid formulation via localized and/or systemic

routes of administration. Lipidoid complexes of polynucleotides can be administered by various means including, but not limited to, intravenous, intramuscular, or subcutaneous routes.

[0678] The synthesis of lipidoids is described in literature (see Mahon et al., Bioconjug. Chem. 2010 21:1448-1454; Schroeder et al., J Intern Med. 2010 267:9-21; Akinc et al., Nat Biotechnol. 2008 26:561-569; Love et al., Proc Natl Acad Sci U S A. 2010 107:1864-1869; Siegwart et al., Proc Natl Acad Sci U S A. 2011 108:12996-3001; all of which are incorporated herein in their entireties).

[0679] Formulations with the different lipidoids, including, but not limited to penta[3-(1-laurylaminopropionyl)]-triethylenetetramine hydrochloride (TETA-5LAP; also known as 98N12-5, see Murugaiah et al., Analytical Biochemistry, 401:61 (2010)), C12-200 (including derivatives and variants), and MD1, can be tested for in vivo activity. The lipidoid "98N12-5" is disclosed by Akinc et al., Mol Ther. 2009 17:872-879. The lipidoid "C12-200" is disclosed by Love et al., Proc Natl Acad Sci U S A. 2010 107:1864-1869 and Liu and Huang, Molecular Therapy. 2010 669-670. Each of the references is herein incorporated by reference in its entirety.

[0680] In one embodiment, the polynucleotides described herein can be formulated in an aminoalcohol lipidoid. Aminoalcohol lipidoids can be prepared by the methods described in U.S. Patent No. 8,450,298 (herein incorporated by reference in its entirety).

[0681] The lipidoid formulations can include particles comprising either 3 or 4 or more components in addition to polynucleotides. Lipidoids and polynucleotide formulations comprising lipidoids are described in Intl. Pub. No. WO 2015051214 (herein incorporated by reference in its entirety.

c. Hyaluronidase

[0682] In some embodiments, the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) and hyaluronidase for injection (e.g., intramuscular or subcutaneous injection). Hyaluronidase catalyzes the hydrolysis of hyaluronan, which is a constituent of the interstitial barrier. Hyaluronidase lowers the viscosity of hyaluronan, thereby increases tissue permeability (Frost, Expert Opin. Drug Deliv. (2007) 4:427-440). Alternatively, the hyaluronidase can be used to increase the number of cells exposed to the polynucleotides administered intramuscularly, or subcutaneously.

d. Nanoparticle Mimics

[0683] In some embodiments, the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) is encapsulated within and/or absorbed to a nanoparticle mimic. A nanoparticle mimic can mimic the delivery function organisms or particles such as, but not limited to, pathogens, viruses, bacteria, fungus, parasites, prions and cells. As a non-limiting example, the polynucleotides described herein can be encapsulated in a non-viron particle that can mimic the delivery function of a virus (see e.g., Intl. Pub. No. WO2012006376 and U.S. Pub. Nos. US20130171241 and US20130195968, each of which is herein incorporated by reference in its entirety).

e. Self-Assembled Nanoparticles, or Self-Assembled Macromolecules

[0684] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) in self-assembled nanoparticles, or amphiphilic macromolecules (AMs) for delivery. AMs comprise biocompatible amphiphilic polymers that have an alkylated sugar backbone covalently linked to poly(ethylene glycol). In aqueous solution, the AMs self-assemble to form micelles. Nucleic acid self-assembled nanoparticles are described in Intl. Appl. No. PCT/US2014/027077, and AMs and methods of forming AMs are described in U.S. Pub. No. US20130217753, each of which is herein incorporated by reference in its entirety.

f. Cations and Anions

[0685] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) and a cation or anion, such as Zn2+, Ca2+, Cu2+, Mg2+ and combinations thereof. Exemplary formulations can include polymers and a polynucleotide complexed with a metal cation as described in, e.g., U.S. Pat. Nos. 6,265,389 and 6,555,525, each of which is herein incorporated by reference in its entirety. In some embodiments, cationic nanoparticles can contain a combination of divalent and monovalent cations. The delivery of polynucleotides in cationic nanoparticles or in one or more depot

comprising cationic nanoparticles can improve polynucleotide bioavailability by acting as a long-acting depot and/or reducing the rate of degradation by nucleases.

g. Amino Acid Lipids

[0686] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) that is formulation with an amino acid lipid. Amino acid lipids are lipophilic compounds comprising an amino acid residue and one or more lipophilic tails. Non-limiting examples of amino acid lipids and methods of making amino acid lipids are described in U.S. Pat. No. 8,501,824. The amino acid lipid formulations can deliver a polynucleotide in releasable form that comprises an amino acid lipid that binds and releases the polynucleotides. As a non-limiting example, the release of the polynucleotides described herein can be provided by an acid-labile linker as described in, e.g., U.S. Pat. Nos. 7,098,032, 6,897,196, 6,426,086, 7,138,382, 5,563,250, and 5,505,931, each of which is herein incorporated by reference in its entirety.

h. Interpolyelectrolyte Complexes

[0687] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) in an interpolyelectrolyte complex. Interpolyelectrolyte complexes are formed when charge-dynamic polymers are complexed with one or more anionic molecules. Non-limiting examples of charge-dynamic polymers and interpolyelectrolyte complexes and methods of making interpolyelectrolyte complexes are described in U.S. Pat. No. 8,524,368, herein incorporated by reference in its entirety.

i. Crystalline Polymeric Systems

[0688] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) in crystalline polymeric systems. Crystalline polymeric systems are polymers with crystalline moieties and/or terminal units comprising crystalline moieties. Exemplary polymers

are described in U.S. Pat. No. 8,524,259 (herein incorporated by reference in its entirety).

j. Polymers, Biodegradable Nanoparticles, and Core-Shell Nanoparticles

[0689] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) and a natural and/or synthetic polymer. The polymers include, but not limited to, polyethenes, polyethylene glycol (PEG), poly(l-lysine)(PLL), PEG grafted to PLL, cationic lipopolymer, biodegradable cationic lipopolymer, polyethyleneimine (PEI), cross-linked branched poly(alkylene imines), a polyamine derivative, a modified poloxamer, elastic biodegradable polymer, biodegradable copolymer, biodegradable polyester copolymer, biodegradable polyester copolymer, multiblock copolymers, poly[α-(4-aminobutyl)-L-glycolic acid) (PAGA), biodegradable cross-linked cationic multi-block copolymers, polycarbonates, polyanhydrides, polyhydroxyacids, polypropylfumerates, polycaprolactones, polyamides, polyacetals, polyethers, polyesters, poly(orthoesters), polycyanoacrylates, polyvinyl alcohols, polyurethanes, polyphosphazenes, polyureas, polystyrenes, polyamines, polylysine, poly(ethylene imine), poly(serine ester), poly(L-lactide-co-L-lysine), poly(4-hydroxy-L-proline ester), amine-containing polymers, dextran polymers, dextran polymer derivatives or combinations thereof.

[0690] Exemplary polymers include, DYNAMIC POLYCONJUGATE® (Arrowhead Research Corp., Pasadena, CA) formulations from MIRUS® Bio (Madison, WI) and Roche Madison (Madison, WI), PHASERXTM polymer formulations such as, without limitation, SMARTT POLYMER TECHNOLOGY™ (PHASERX®, Seattle, WA), DMRI/DOPE, poloxamer, VAXFECTIN® adjuvant from Vical (San Diego, CA), chitosan, cyclodextrin from Calando Pharmaceuticals (Pasadena, CA), dendrimers and poly(lactic-co-glycolic acid) (PLGA) polymers. RONDELTM (RNAi/Oligonucleotide Nanoparticle Delivery) polymers (Arrowhead Research Corporation, Pasadena, CA) and pH responsive co-block polymers such as PHASERX® (Seattle, WA).

[0691] The polymer formulations allow a sustained or delayed release of the polynucleotide (e.g., following intramuscular or subcutaneous injection). The altered

release profile for the polynucleotide can result in, for example, translation of an encoded protein over an extended period of time. The polymer formulation can also be used to increase the stability of the polynucleotide. Sustained release formulations can include, but are not limited to, PLGA microspheres, ethylene vinyl acetate (EVAc), poloxamer, GELSITE® (Nanotherapeutics, Inc. Alachua, FL), HYLENEX® (Halozyme Therapeutics, San Diego CA), surgical sealants such as fibrinogen polymers (Ethicon Inc. Cornelia, GA), TISSELL® (Baxter International, Inc. Deerfield, IL), PEG-based sealants, and COSEAL® (Baxter International, Inc. Deerfield, IL).

[0692] As a non-limiting example modified mRNA can be formulated in PLGA microspheres by preparing the PLGA microspheres with tunable release rates (e.g., days and weeks) and encapsulating the modified mRNA in the PLGA microspheres while maintaining the integrity of the modified mRNA during the encapsulation process. EVAc are non-biodegradable, biocompatible polymers that are used extensively in pre-clinical sustained release implant applications (e.g., extended release products Ocusert a pilocarpine ophthalmic insert for glaucoma or progestasert a sustained release progesterone intrauterine device; transdermal delivery systems Testoderm, Duragesic and Selegiline; catheters). Poloxamer F-407 NF is a hydrophilic, non-ionic surfactant triblock copolymer of polyoxyethylene-polyoxypropylene-polyoxyethylene having a low viscosity at temperatures less than 5ºC and forms a solid gel at temperatures greater than 15ºC.

[0693] As a non-limiting example, the polynucleotides described herein can be formulated with the polymeric compound of PEG grafted with PLL as described in U.S. Pat. No. 6,177,274. As another non-limiting example, the polynucleotides described herein can be formulated with a block copolymer such as a PLGA-PEG block copolymer (see e.g., U.S. Pub. No. US20120004293 and U.S. Pat. Nos. 8,236,330 and 8,246,968), or a PLGA-PEG-PLGA block copolymer (see e.g., U.S. Pat. No. 6,004,573). Each of the references is herein incorporated by reference in its entirety.

[0694] In some embodiments, the polynucleotides described herein can be formulated with at least one amine-containing polymer such as, but not limited to polylysine, polyethylene imine, poly(amidoamine) dendrimers, poly(amine-co-esters) or combinations thereof. Exemplary polyamine polymers and their use as delivery

agents are described in, e.g., U.S. Pat. Nos. 8,460,696, 8,236,280, each of which is herein incorporated by reference in its entirety.

[0695] In some embodiments, the polynucleotides described herein can be formulated in a biodegradable cationic lipopolymer, a biodegradable polymer, or a biodegradable copolymer, a biodegradable polyester copolymer, a biodegradable polyester polymer, a linear biodegradable copolymer, PAGA, a biodegradable cross-linked cationic multi-block copolymer or combinations thereof as described in, e.g., U.S. Pat. Nos. 6,696,038, 6,517,869, 6,267,987, 6,217,912, 6,652,886, 8,057,821, and 8,444,992; U.S. Pub. Nos. US20030073619, US20040142474, US20100004315, US2012009145 and US20130195920; and Intl Pub. Nos. WO2006063249 and WO2013086322, each of which is herein incorporated by reference in its entirety.

[0696] In some embodiments, the polynucleotides described herein can be formulated in or with at least one cyclodextrin polymer as described in U.S. Pub. No. US20130184453. In some embodiments, the polynucleotides described herein can be formulated in or with at least one crosslinked cation-binding polymers as described in Intl. Pub. Nos. WO2013106072, WO2013106073 and WO2013106086. In some embodiments, the polynucleotides described herein can be formulated in or with at least PEGylated albumin polymer as described in U.S. Pub. No. US20130231287. Each of the references is herein incorporated by reference in its entirety.

[0697] In some embodiments, the polynucleotides disclosed herein can be formulated as a nanoparticle using a combination of polymers, lipids, and/or other biodegradable agents, such as, but not limited to, calcium phosphate. Components can be combined in a core-shell, hybrid, and/or layer-by-layer architecture, to allow for fine-tuning of the nanoparticle for delivery (Wang et al., Nat Mater. 2006 5:791-796; Fuller et al., Biomaterials. 2008 29:1526-1532; DeKoker et al., Adv Drug Deliv Rev. 2011 63:748-761; Endres et al., Biomaterials. 2011 32:7721-7731; Su et al., Mol Pharm. 2011 Jun 6;8(3):774-87; herein incorporated by reference in their entireties). As a non-limiting example, the nanoparticle can comprise a plurality of polymers such as, but not limited to hydrophilic-hydrophobic polymers (e.g., PEG-PLGA), hydrophobic polymers (e.g., PEG) and/or hydrophilic polymers (Intl. Pub. No. WO20120225129, herein incorporated by reference in its entirety).

[0698] The use of core-shell nanoparticles has additionally focused on a high-throughput approach to synthesize cationic cross-linked nanogel cores and various shells (Siegwart et al., Proc Natl Acad Sci U S A. 2011 108:12996-13001; herein

incorporated by reference in its entirety). The complexation, delivery, and internalization of the polymeric nanoparticles can be precisely controlled by altering the chemical composition in both the core and shell components of the nanoparticle. For example, the core-shell nanoparticles can efficiently deliver siRNA to mouse hepatocytes after they covalently attach cholesterol to the nanoparticle.

[0699] In some embodiments, a hollow lipid core comprising a middle PLGA layer and an outer neutral lipid layer containing PEG can be used to delivery of the polynucleotides as described herein. In some embodiments, the lipid nanoparticles can comprise a core of the polynucleotides disclosed herein and a polymer shell, which is used to protect the polynucleotides in the core. The polymer shell can be any of the polymers described herein and are known in the art. The polymer shell can be used to protect the polynucleotides in the core.

[0700] Core–shell nanoparticles for use with the polynucleotides described herein are described in U.S. Pat. No. 8,313,777 or Intl. Pub. No. WO2013124867, each of which is herein incorporated by reference in their entirety.

k. Peptides and Proteins

[0701] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) that is formulated with peptides and/or proteins to increase transfection of cells by the polynucleotide, and/or to alter the biodistribution of the polynucleotide (e.g., by targeting specific tissues or cell types), and/or increase the translation of encoded protein (e.g., Intl. Pub. Nos. WO2012110636 and WO2013123298. In some embodiments, the peptides can be those described in U.S. Pub. Nos. US20130129726, US20130137644 and US20130164219. Each of the references is herein incorporated by reference in its entirety.

l. Conjugates

[0702] In some embodiments, the compositions or formulations of the present disclosure comprise the polynucleotides described herein (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide) that is covalently linked to a carrier or targeting group, or including two encoding regions that together produce a fusion protein (e.g., bearing a targeting group and therapeutic protein or

peptide) as a conjugate. The conjugate can be a peptide that selectively directs the nanoparticle to neurons in a tissue or organism, or assists in crossing the blood-brain barrier.

[0703] The conjugates include a naturally occurring substance, such as a protein (e.g., human serum albumin (HSA), low-density lipoprotein (LDL), high-density lipoprotein (HDL), or globulin); a carbohydrate (e.g., a dextran, pullulan, chitin, chitosan, inulin, cyclodextrin or hyaluronic acid); or a lipid. The ligand can also be a recombinant or synthetic molecule, such as a synthetic polymer, e.g., a synthetic polyamino acid, an oligonucleotide (e.g., an aptamer). Examples of polyamino acids include polyamino acid is a polylysine (PLL), poly L-aspartic acid, poly L-glutamic acid, styrene-maleic acid anhydride copolymer, poly(L-lactide-co-glycolied) copolymer, divinyl ether-maleic anhydride copolymer, N-(2-hydroxypropyl)methacrylamide copolymer (HMPA), polyethylene glycol (PEG), polyvinyl alcohol (PVA), polyurethane, poly(2-ethylacryllic acid), N-isopropylacrylamide polymers, or polyphosphazine. Example of polyamines include: polyethylenimine, polylysine (PLL), spermine, spermidine, polyamine, pseudopeptide-polyamine, peptidomimetic polyamine, dendrimer polyamine, arginine, amidine, protamine, cationic lipid, cationic porphyrin, quaternary salt of a polyamine, or an alpha helical peptide.

[0704] In some embodiments, the conjugate can function as a carrier for the polynucleotide disclosed herein. The conjugate can comprise a cationic polymer such as, but not limited to, polyamine, polylysine, polyalkylenimine, and polyethylenimine that can be grafted to with poly(ethylene glycol). Exemplary conjugates and their preparations are described in U.S. Pat. No. 6,586,524 and U.S. Pub. No. US20130211249, each of which herein is incorporated by reference in its entirety.

[0705] The conjugates can also include targeting groups, e.g., a cell or tissue targeting agent, e.g., a lectin, glycoprotein, lipid or protein, e.g., an antibody, that binds to a specified cell type such as a kidney cell. A targeting group can be a thyrotropin, melanotropin, lectin, glycoprotein, surfactant protein A, Mucin carbohydrate, multivalent lactose, multivalent galactose, N-acetyl-galactosamine, N-acetyl-glucosamine multivalent mannose, multivalent fucose, glycosylated polyaminoacids, multivalent galactose, transferrin, bisphosphonate, polyglutamate, polyaspartate, a lipid, cholesterol, a steroid, bile acid, folate, vitamin B12, biotin, an RGD peptide, an RGD peptide mimetic or an aptamer.

[0706] Targeting groups can be proteins, e.g., glycoproteins, or peptides, e.g., molecules having a specific affinity for a co-ligand, or antibodies e.g., an antibody, that binds to a specified cell type such as an endothelial cell or bone cell. Targeting groups can also include hormones and hormone receptors. They can also include non-peptidic species, such as lipids, lectins, carbohydrates, vitamins, cofactors, multivalent lactose, multivalent galactose, N-acetyl-galactosamine, N-acetyl-glucosamine multivalent mannose, multivalent frucose, or aptamers. The ligand can be, for example, a lipopolysaccharide, or an activator of p38 MAP kinase.

[0707] The targeting group can be any ligand that is capable of targeting a specific receptor. Examples include, without limitation, folate, GalNAc, galactose, mannose, mannose-6P, aptamers, integrin receptor ligands, chemokine receptor ligands, transferrin, biotin, serotonin receptor ligands, PSMA, endothelin, GCPII, somatostatin, LDL, and HDL ligands. In particular embodiments, the targeting group is an aptamer. The aptamer can be unmodified or have any combination of modifications disclosed herein. As a non-limiting example, the targeting group can be a glutathione receptor (GR)-binding conjugate for targeted delivery across the blood-central nervous system barrier as described in, e.g., U.S. Pub. No. US2013021661012 (herein incorporated by reference in its entirety).

[0708] In some embodiments, the conjugate can be a synergistic biomolecule-polymer conjugate, which comprises a long-acting continuous-release system to provide a greater therapeutic efficacy. The synergistic biomolecule-polymer conjugate can be those described in U.S. Pub. No. US20130195799. In some embodiments, the conjugate can be an aptamer conjugate as described in Intl. Pat. Pub. No. WO2012040524. In some embodiments, the conjugate can be an amine containing polymer conjugate as described in U.S. Pat. No. 8,507,653. Each of the references is herein incorporated by reference in its entirety. In some embodiments, the polynucleotides can be conjugated to SMARTT POLYMER TECHNOLOGY® (PHASERX®, Inc. Seattle, WA).

[0709] In some embodiments, the polynucleotides described herein are covalently conjugated to a cell penetrating polypeptide, which can also include a signal sequence or a targeting sequence. The conjugates can be designed to have increased stability, and/or increased cell transfection; and/or altered the biodistribution (e.g., targeted to specific tissues or cell types).

[0710] In some embodiments, the polynucleotides described herein can be conjugated to an agent to enhance delivery. In some embodiments, the agent can be a monomer or polymer such as a targeting monomer or a polymer having targeting blocks as described in Intl. Pub. No. WO2011062965. In some embodiments, the agent can be a transport agent covalently coupled to a polynucleotide as described in, e.g., U.S. Pat. Nos. 6,835.393 and 7,374,778. In some embodiments, the agent can be a membrane barrier transport enhancing agent such as those described in U.S. Pat. Nos. 7,737,108 and 8,003,129. Each of the references is herein incorporated by reference in its entirety.

22. Accelerated Blood Clearance

[0711] The disclosure provides compounds, compositions and methods of use thereof for reducing the effect of ABC on a repeatedly administered active agent such as a biologically active agent. As will be readily apparent, reducing or eliminating altogether the effect of ABC on an administered active agent effectively increases its half-life and thus its efficacy.

[0712] In some embodiments the term reducing ABC refers to any reduction in ABC in comparison to a positive reference control ABC inducing LNP such as an MC3 LNP. ABC inducing LNPs cause a reduction in circulating levels of an active agent upon a second or subsequent administration within a given time frame. Thus, a reduction in ABC refers to less clearance of circulating agent upon a second or subsequent dose of agent, relative to a standard LNP. The reduction may be, for instance, at least 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 98%, or 100%. In some embodiments the reduction is 10-100%, 10-50%, 20-100%, 20-50%, 30-100%, 30-50%, 40%-100%, 40-80%, 50-90%, or 50-100%. Alternatively, the reduction in ABC may be characterized as at least a detectable level of circulating agent following a second or subsequent administration or at least a 2-fold, 3-fold, 4-fold, 5-fold increase in circulating agent relative to circulating agent following administration of a standard LNP. In some embodiments the reduction is a 2-100 fold, 2-50 fold, 3-100 fold, 3-50 fold, 3-20 fold, 4-100 fold, 4-50 fold, 4-40 fold, 4-30 fold, 4-25 fold, 4-20 fold, 4-15 fold, 4-10 fold, 4-5 fold, 5 -100 fold, 5-50 fold, 5-40 fold, 5-30 fold, 5-25 fold, 5-20 fold, 5-15 fold, 5-10 fold, 6-100 fold, 6-50 fold, 6-40 fold, 6-30 fold, 6-25 fold, 6-20 fold, 6-15 fold, 6-10 fold, 8-100 fold, 8-50 fold, 8-40 fold, 8-30 fold, 8-25 fold, 8-20 fold, 8-15 fold,

8-10 fold, 10-100 fold, 10-50 fold, 10-40 fold, 10-30 fold, 10-25 fold, 10-20 fold, 10-15 fold, 20-100 fold, 20-50 fold, 20-40 fold, 20-30 fold, or 20-25 fold.

[0713] The disclosure provides lipid-comprising compounds and compositions that are less susceptible to clearance and thus have a longer half-life in vivo. This is particularly the case where the compositions are intended for repeated including chronic administration, and even more particularly where such repeated administration occurs within days or weeks.

[0714] Significantly, these compositions are less susceptible or altogether circumvent the observed phenomenon of accelerated blood clearance (ABC). ABC is a phenomenon in which certain exogenously administered agents are rapidly cleared from the blood upon second and subsequent administrations. This phenomenon has been observed, in part, for a variety of lipid-containing compositions including but not limited to lipidated agents, liposomes or other lipid-based delivery vehicles, and lipid-encapsulated agents. Heretofore, the basis of ABC has been poorly understood and in some cases attributed to a humoral immune response and accordingly strategies for limiting its impact in vivo particularly in a clinical setting have remained elusive.

[0715] This disclosure provides compounds and compositions that are less susceptible, if at all susceptible, to ABC. In some important aspects, such compounds and compositions are lipid-comprising compounds or compositions. The lipid-containing compounds or compositions of this disclosure, surprisingly, do not experience ABC upon second and subsequent administration in vivo. This resistance to ABC renders these compounds and compositions particularly suitable for repeated use in vivo, including for repeated use within short periods of time, including days or 1-2 weeks. This enhanced stability and/or half-life is due, in part, to the inability of these compositions to activate B1a and/or B1b cells and/or conventional B cells, pDCs and/or platelets.

[0716] This disclosure therefore provides an elucidation of the mechanism underlying accelerated blood clearance (ABC). It has been found, in accordance with this disclosure and the inventions provided herein, that the ABC phenomenon at least as it relates to lipids and lipid nanoparticles is mediated, at least in part an innate immune response involving B1a and/or B1b cells, pDC and/or platelets. B1a cells are normally responsible for secreting natural antibody, in the form of circulating IgM. This IgM is poly-reactive, meaning that it can bind to a variety of antigens, albeit with a relatively low affinity for each.

[0717] It has been found in accordance with the invention that some lipidated agents or lipid-comprising formulations such as lipid nanoparticles administered in vivo trigger and are subject to ABC. It has now been found in accordance with the invention that upon administration of a first dose of the LNP, one or more cells involved in generating an innate immune response (referred to herein as sensors) bind such agent, are activated, and then initiate a cascade of immune factors (referred to herein as effectors) that promote ABC and toxicity. For instance, B1a and B1b cells may bind to LNP, become activated (alone or in the presence of other sensors such as pDC and/or effectors such as IL6) and secrete natural IgM that binds to the LNP. Pre-existing natural IgM in the subject may also recognize and bind to the LNP, thereby triggering complement fixation. After administration of the first dose, the production of natural IgM begins within 1-2 hours of administration of the LNP. Typically, by about 2-3 weeks the natural IgM is cleared from the system due to the natural half-life of IgM. Natural IgG is produced beginning around 96 hours after administration of the LNP. The agent, when administered in a naïve setting, can exert its biological effects relatively unencumbered by the natural IgM produced post-activation of the B1a cells or B1b cells or natural IgG. The natural IgM and natural IgG are non-specific and thus are distinct from anti-PEG IgM and anti-PEG IgG.

[0718] Although Applicant is not bound by mechanism, it is proposed that LNPs trigger ABC and/or toxicity through the following mechanisms. It is believed that when an LNP is administered to a subject the LNP is rapidly transported through the blood to the spleen. The LNPs may encounter immune cells in the blood and/or the spleen. A rapid innate immune response is triggered in response to the presence of the LNP within the blood and/or spleen. Applicant has shown herein that within hours of administration of an LNP several immune sensors have reacted to the presence of the LNP. These sensors include but are not limited to immune cells involved in generating an immune response, such as B cells, pDC, and platelets. The sensors may be present in the spleen, such as in the marginal zone of the spleen and/or in the blood. The LNP may physically interact with one or more sensors, which may interact with other sensors. In such a case the LNP is directly or indirectly interacting with the sensors. The sensors may interact directly with one another in response to recognition of the LNP. For instance, many sensors are located in the spleen and can easily interact with one another. Alternatively, one or more of the sensors may interact with LNP in the blood and become activated. The activated sensor may then interact

directly with other sensors or indirectly (e.g., through the stimulation or production of a messenger such as a cytokine e.g., IL6).

[0719] In some embodiments the LNP may interact directly with and activate each of the following sensors: pDC, B1a cells, B1b cells, and platelets. These cells may then interact directly or indirectly with one another to initiate the production of effectors which ultimately lead to the ABC and/or toxicity associated with repeated doses of LNP. For instance, Applicant has shown that LNP administration leads to pDC activation, platelet aggregation and activation and B cell activation. In response to LNP platelets also aggregate and are activated and aggregate with B cells. pDC cells are activated. LNP has been found to interact with the surface of platelets and B cells relatively quickly. Blocking the activation of any one or combination of these sensors in response to LNP is useful for dampening the immune response that would ordinarily occur. This dampening of the immune response results in the avoidance of ABC and/or toxicity.

[0720] The sensors once activated produce effectors. An effector, as used herein, is an immune molecule produced by an immune cell, such as a B cell. Effectors include but are not limited to immunoglobulin such as natural IgM and natural IgG and cytokines such as IL6. B1a and B1b cells stimulate the production of natural IgMs within 2-6 hours following administration of an LNP. Natural IgG can be detected within 96 hours. IL6 levels are increased within several hours. The natural IgM and IgG circulate in the body for several days to several weeks. During this time the circulating effectors can interact with newly administered LNPs, triggering those LNPs for clearance by the body. For instance, an effector may recognize and bind to an LNP. The Fc region of the effector may be recognized by and trigger uptake of the decorated LNP by macrophage. The macrophage is then transported to the spleen. The production of effectors by immune sensors is a transient response that correlates with the timing observed for ABC.

[0721] If the administered dose is the second or subsequent administered dose, and if such second or subsequent dose is administered before the previously induced natural IgM and/or IgG is cleared from the system (e.g., before the 2-3 window time period), then such second or subsequent dose is targeted by the circulating natural IgM and/or natural IgG or Fc which trigger alternative complement pathway activation and is itself rapidly cleared. When LNP are administered after the effectors have cleared from the body or are reduced in number, ABC is not observed.

[0722] Thus, it is useful according to aspects of the invention to inhibit the interaction between LNP and one or more sensors, to inhibit the activation of one or more sensors by LNP (direct or indirect), to inhibit the production of one or more effectors, and/or to inhibit the activity of one or more effectors. In some embodiments the LNP is designed to limit or block interaction of the LNP with a sensor. For instance, the LNP may have an altered PC and/or PEG to prevent interactions with sensors. Alternatively or additionally, an agent that inhibits immune responses induced by LNPs may be used to achieve any one or more of these effects.

[0723] It has also been determined that conventional B cells are also implicated in ABC. Specifically, upon first administration of an agent, conventional B cells, referred to herein as CD19(+), bind to and react against the agent. Unlike B1a and B1b cells though, conventional B cells can mount first an IgM response (beginning around 96 hours after administration of the LNPs) followed by an IgG response (beginning around 14 days after administration of the LNPs) concomitant with a memory response. Thus, conventional B cells react against the administered agent and contribute to IgM (and eventually IgG) that mediates ABC. The IgM and IgG are typically anti-PEG IgM and anti-PEG IgG.

[0724] It is contemplated that in some instances, much of the ABC response is mediated through B1a cells and B1a-mediated immune responses. It is further contemplated that in some instances, the ABC response is mediated by both IgM and IgG, with both conventional B cells and B1a cells mediating such effects. In yet still other instances, the ABC response is mediated by natural IgM molecules, some of which are capable of binding to natural IgM, which may be produced by activated B1a cells. The natural IgMs may bind to one or more components of the LNPs, e.g., binding to a phospholipid component of the LNPs (such as binding to the PC moiety of the phospholipid) and/or binding to a PEG-lipid component of the LNPs (such as binding to PEG-DMG, in particular, binding to the PEG moiety of PEG-DMG). Since B1a expresses CD36, to which phosphatidylcholine is a ligand, it is contemplated that the CD36 receptor may mediate the activation of B1a cells and thus production of natural IgM. In yet still other instances, the ABC response is mediated primarily by conventional B cells.

[0725] It has been found in accordance with the invention that the ABC phenomenon can be reduced or abrogated, at least in part, through the use of compounds and compositions (such as agents, delivery vehicles, and formulations) that do not activate

B1a cells. Compounds and compositions that do not activate B1a cells may be referred to herein as B1a inert compounds and compositions. It has been further found in accordance with the invention that the ABC phenomenon can be reduced or abrogated, at least in part, through the use of compounds and compositions that do not activate conventional B cells. Compounds and compositions that do not activate conventional B cells may in some embodiments be referred to herein as CD19-inert compounds and compositions. Thus, in some embodiments provided herein, the compounds and compositions do not activate B1a cells and they do not activate conventional B cells. Compounds and compositions that do not activate B1a cells and conventional B cells may in some embodiments be referred to herein as B1a/CD19-inert compounds and compositions.

[0726] These underlying mechanisms were not heretofore understood, and the role of B1a and B1b cells and their interplay with conventional B cells in this phenomenon was also not appreciated.

[0727] Accordingly, this disclosure provides compounds and compositions that do not promote ABC. These may be further characterized as not capable of activating B1a and/or B1b cells, platelets and/or pDC, and optionally conventional B cells also. These compounds (e.g., agents, including biologically active agents such as prophylactic agents, therapeutic agents and diagnostic agents, delivery vehicles, including liposomes, lipid nanoparticles, and other lipid-based encapsulating structures, etc.) and compositions (e.g., formulations, etc.) are particularly desirable for applications requiring repeated administration, and in particular repeated administrations that occur within with short periods of time (e.g., within 1-2 weeks). This is the case, for example, if the agent is a nucleic acid based therapeutic that is provided to a subject at regular, closely-spaced intervals. The findings provided herein may be applied to these and other agents that are similarly administered and/or that are subject to ABC.

[0728] Of particular interest are lipid-comprising compounds, lipid-comprising particles, and lipid-comprising compositions as these are known to be susceptible to ABC. Such lipid-comprising compounds particles, and compositions have been used extensively as biologically active agents or as delivery vehicles for such agents. Thus, the ability to improve their efficacy of such agents, whether by reducing the effect of ABC on the agent itself or on its delivery vehicle, is beneficial for a wide variety of active agents.

[0729] Also provided herein are compositions that do not stimulate or boost an acute phase response (ARP) associated with repeat dose administration of one or more biologically active agents.

[0730] The composition, in some instances, may not bind to IgM, including but not limited to natural IgM.

[0731] The composition, in some instances, may not bind to an acute phase protein such as but not limited to C-reactive protein.

[0732] The composition, in some instances, may not trigger a CD5(+) mediated immune response. As used herein, a CD5(+) mediated immune response is an immune response that is mediated by B1a and/or B1b cells. Such a response may include an ABC response, an acute phase response, induction of natural IgM and/or IgG, and the like.

[0733] The composition, in some instances, may not trigger a CD19(+) mediated immune response. As used herein, a CD19(+) mediated immune response is an immune response that is mediated by conventional CD19(+), CD5(-) B cells. Such a response may include induction of IgM, induction of IgG, induction of memory B cells, an ABC response, an anti-drug antibody (ADA) response including an anti-protein response where the protein may be encapsulated within an LNP, and the like.

[0734] B1a cells are a subset of B cells involved in innate immunity. These cells are the source of circulating IgM, referred to as natural antibody or natural serum antibody. Natural IgM antibodies are characterized as having weak affinity for several antigens, and therefore they are referred to as "poly-specific" or "poly-reactive", indicating their ability to bind to more than one antigen. B1a cells are not able to produce IgG. Additionally, they do not develop into memory cells and thus do not contribute to an adaptive immune response. However, they can secrete IgM upon activation. The secreted IgM is typically cleared within about 2-3 weeks, at which point the immune system is rendered relatively naïve to the previously administered antigen. If the same antigen is presented after this time period (e.g., at about 3 weeks after the initial exposure), the antigen is not rapidly cleared. However, significantly, if the antigen is presented within that time period (e.g., within 2 weeks, including within 1 week, or within days), then the antigen is rapidly cleared. This delay between consecutive doses has rendered certain lipid-containing therapeutic or diagnostic agents unsuitable for use.

[0735] In humans, B1a cells are CD19(+), CD20(+), CD27(+), CD43(+), CD70(-) and CD5(+). In mice, B1a cells are CD19(+), CD5(+), and CD45 B cell isoform B220(+). It is the expression of CD5 which typically distinguishes B1a cells from other convention B cells. B1a cells may express high levels of CD5, and on this basis, may be distinguished from other B-1 cells such as B-1b cells which express low or undetectable levels of CD5. CD5 is a pan-T cell surface glycoprotein. B1a cells also express CD36, also known as fatty acid translocase. CD36 is a member of the class B scavenger receptor family. CD36 can bind many ligands, including oxidized low-density lipoproteins, native lipoproteins, oxidized phospholipids, and long-chain fatty acids.

[0736] B1b cells are another subset of B cells involved in innate immunity. These cells are another source of circulating natural IgM. Several antigens, including PS, are capable of inducing T cell independent immunity through B1b activation. CD27 is typically upregulated on B1b cells in response to antigen activation. Similar to B1a cells, the B1b cells are typically located in specific body locations such as the spleen and peritoneal cavity and are in very low abundance in the blood. The B1b secreted natural IgM is typically cleared within about 2-3 weeks, at which point the immune system is rendered relatively naïve to the previously administered antigen. If the same antigen is presented after this time period (e.g., at about 3 weeks after the initial exposure), the antigen is not rapidly cleared. However, significantly, if the antigen is presented within that time period (e.g., within 2 weeks, including within 1 week, or within days), then the antigen is rapidly cleared. This delay between consecutive doses has rendered certain lipid-containing therapeutic or diagnostic agents unsuitable for use.

[0737] In some embodiments it is desirable to block B1a and/or B1b cell activation. One strategy for blocking B1a and/or B1b cell activation involves determining which components of a lipid nanoparticle promote B cell activation and neutralizing those components. It has been discovered herein that at least PEG and phosphatidylcholine (PC) contribute to B1a and B1b cell interaction with other cells and/or activation. PEG may play a role in promoting aggregation between B1 cells and platelets, which may lead to activation. PC (a helper lipid in LNPs) is also involved in activating the B1 cells, likely through interaction with the CD36 receptor on the B cell surface. Numerous particles have PEG-lipid alternatives, PEG-less, and/or PC replacement lipids (e.g. oleic acid or analogs thereof) have been designed and tested. Applicant

has established that replacement of one or more of these components within an LNP that otherwise would promote ABC upon repeat administration, is useful in preventing ABC by reducing the production of natural IgM and/or B cell activation. Thus, the invention encompasses LNPs that have reduced ABC as a result of a design which eliminates the inclusion of B cell triggers.

[0738] Another strategy for blocking B1a and/or B1b cell activation involves using an agent that inhibits immune responses induced by LNPs. These types of agents are discussed in more detail below. In some embodiments these agents block the interaction between B1a/B1b cells and the LNP or platelets or pDC. For instance, the agent may be an antibody or other binding agent that physically blocks the interaction. An example of this is an antibody that binds to CD36 or CD6. The agent may also be a compound that prevents or disables the B1a/B1b cell from signaling once activated or prior to activation. For instance, it is possible to block one or more components in the B1a/B1b signaling cascade the results from B cell interaction with LNP or other immune cells. In other embodiments the agent may act one or more effectors produced by the B1a/B1b cells following activation. These effectors include for instance, natural IgM and cytokines.

[0739] It has been demonstrated according to aspects of the invention that when activation of pDC cells is blocked, B cell activation in response to LNP is decreased. Thus, to avoid ABC and/or toxicity, it may be desirable to prevent pDC activation. Similar to the strategies discussed above, pDC cell activation may be blocked by agents that interfere with the interaction between pDC and LNP and/or B cells/platelets. Alternatively, agents that act on the pDC to block its ability to get activated or on its effectors can be used together with the LNP to avoid ABC.

[0740] Platelets may also play an important role in ABC and toxicity. Very quickly after a first dose of LNP is administered to a subject, platelets associate with the LNP, aggregate and are activated. In some embodiments it is desirable to block platelet aggregation and/or activation. One strategy for blocking platelet aggregation and/or activation involves determining which components of a lipid nanoparticle promote platelet aggregation and/or activation and neutralizing those components. It has been discovered herein that at least PEG contribute to platelet aggregation, activation and/or interaction with other cells. Numerous particles have PEG-lipid alternatives and PEG-less have been designed and tested. Applicant has established that replacement of one or more of these components within an LNP that otherwise would

promote ABC upon repeat administration, is useful in preventing ABC by reducing the production of natural IgM and/or platelet aggregation. Thus, the invention encompasses LNPs that have reduced ABC as a result of a design which eliminates the inclusion of platelet triggers. Alternatively, agents that act on the platelets to block its activity once it is activated or on its effectors can be used together with the LNP to avoid ABC.

(i) Measuring ABC Activity and related activities

[0741] Various compounds and compositions provided herein, including LNPs, do not promote ABC activity upon administration in vivo. These LNPs may be characterized and/or identified through any of several assays, such as but not limited to those described below, as well as any of the assays disclosed in the Examples section, include the methods subsection of the Examples.

[0742] In some embodiments the methods involve administering an LNP without producing an immune response that promotes ABC. An immune response that promotes ABC involves activation of one or more sensors, such as B1 cells, pDC, or platelets, and one or more effectors, such as natural IgM, natural IgG or cytokines such as IL6. Thu,s administration of an LNP without producing an immune response that promotes ABC, at a minimum involves administration of an LNP without significant activation of one or more sensors and significant production of one or more effectors. Significant used in this context refers to an amount that would lead to the physiological consequence of accelerated blood clearance of all or part of a second dose with respect to the level of blood clearance expected for a second dose of an ABC triggering LNP. For instance, the immune response should be dampened such that the ABC observed after the second dose is lower than would have been expected for an ABC triggering LNP.

(ii) B1a or B1b activation assay

[0743] Certain compositions provided in this disclosure do not activate B cells, such as B1a or B1b cells (CD19+ CD5+) and/or conventional B cells (CD19+ CD5-). Activation of B1a cells, B1b cells, or conventional B cells may be determined in several ways, some of which are provided below. B cell population may be provided as fractionated B cell populations or unfractionated populations of splenocytes or peripheral blood mononuclear cells (PBMC). If the latter, the cell population may be

incubated with the LNP of choice for a period of time, and then harvested for further analysis. Alternatively, the supernatant may be harvested and analyzed.

(iii) Upregulation of activation marker cell surface expression

[0744] Activation of B1a cells, B1b cells, or conventional B cells may be demonstrated as increased expression of B cell activation markers including late activation markers such as CD86. In an exemplary non-limiting assay, unfractionated B cells are provided as a splenocyte population or as a PBMC population, incubated with an LNP of choice for a particular period of time, and then stained for a standard B cell marker such as CD19 and for an activation marker such as CD86, and analyzed using for example flow cytometry. A suitable negative control involves incubating the same population with medium, and then performing the same staining and visualization steps. An increase in CD86 expression in the test population compared to the negative control indicates B cell activation.

(iv) Pro-inflammatory cytokine release

[0745] B cell activation may also be assessed by cytokine release assay. For example, activation may be assessed through the production and/or secretion of cytokines such as IL-6 and/or TNF-alpha upon exposure with LNPs of interest.

[0746] Such assays may be performed using routine cytokine secretion assays well known in the art. An increase in cytokine secretion is indicative of B cell activation.

(v) LNP binding/association to and/or uptake by B cells

[0747] LNP association or binding to B cells may also be used to assess an LNP of interest and to further characterize such LNP. Association/binding and/or uptake/internalization may be assessed using a detectably labeled, such as fluorescently labeled, LNP and tracking the location of such LNP in or on B cells following various periods of incubation.

[0748] The invention further contemplates that the compositions provided herein may be capable of evading recognition or detection and optionally binding by downstream mediators of ABC such as circulating IgM and/or acute phase response mediators such as acute phase proteins (e.g., C-reactive protein (CRP).

(vi) Methods of use for reducing ABC

[0749] Also provided herein are methods for delivering LNPs, which may encapsulate an agent such as a therapeutic agent, to a subject without promoting ABC.

[0750] In some embodiments, the method comprises administering any of the LNPs described herein, which do not promote ABC, for example, do not induce production of natural IgM binding to the LNPs, do not activate B1a and/or B1b cells. As used herein, an LNP that "does not promote ABC" refers to an LNP that induces no immune responses that would lead to substantial ABC or a substantially low level of immune responses that is not sufficient to lead to substantial ABC. An LNP that does not induce the production of natural IgMs binding to the LNP refers to LNPs that induce either no natural IgM binding to the LNPs or a substantially low level of the natural IgM molecules, which is insufficient to lead to substantial ABC. An LNP that does not activate B1a and/or B1b cells refer to LNPs that induce no response of B1a and/or B1b cells to produce natural IgM binding to the LNPs or a substantially low level of B1a and/or B1b responses, which is insufficient to lead to substantial ABC.

[0751] In some embodiments the terms do not activate and do not induce production are a relative reduction to a reference value or condition. In some embodiments the reference value or condition is the amount of activation or induction of production of a molecule such as IgM by a standard LNP such as an MC3 LNP. In some embodiments the relative reduction is a reduction of at least 30%, for example at least 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100%. In other embodiments the terms do not activate cells such as B cells and do not induce production of a protein such as IgM may refer to an undetectable amount of the active cells or the specific protein.

(vii) Platelet effects and toxicity

[0752] The invention is further premised in part on the elucidation of the mechanism underlying dose-limiting toxicity associated with LNP administration. Such toxicity may involve coagulopathy, disseminated intravascular coagulation (DIC, also referred to as consumptive coagulopathy), whether acute or chronic, and/or vascular thrombosis. In some instances, the dose-limiting toxicity associated with LNPs is

acute phase response (APR) or complement activation-related pseudoallergy (CARPA).

[0753] As used herein, coagulopathy refers to increased coagulation (blood clotting) in vivo. The findings reported in this disclosure are consistent with such increased coagulation and significantly provide insight on the underlying mechanism. Coagulation is a process that involves several different factors and cell types, and heretofore the relationship between and interaction of LNPs and platelets has not been understood in this regard. This disclosure provides evidence of such interaction and provides compounds and compositions that are modified to have reduced platelet effect, including reduced platelet association, reduced platelet aggregation, and/or reduced platelet aggregation. The ability to modulate, including preferably down-modulate, such platelet effects can reduce the incidence and/or severity of coagulopathy post-LNP administration. This in turn will reduce toxicity relating to such LNP, thereby allowing higher doses of LNPs and importantly their cargo to be administered to patients in need thereof.

[0754] CARPA is a class of acute immune toxicity manifested in hypersensitivity reactions (HSRs), which may be triggered by nanomedicines and biologicals. Unlike allergic reactions, CARPA typically does not involve IgE but arises as a consequence of activation of the complement system, which is part of the innate immune system that enhances the body's abilities to clear pathogens. One or more of the following pathways, the classical complement pathway (CP), the alternative pathway (AP), and the lectin pathway (LP), may be involved in CARPA. Szebeni, Molecular Immunology, 61:163-173 (2014).

[0755] The classical pathway is triggered by activation of the C1-complex, which contains. C1q, C1r, C1s, or C1qr2s2. Activation of the C1-complex occurs when C1q binds to IgM or IgG complexed with antigens, or when C1q binds directly to the surface of the pathogen. Such binding leads to conformational changes in the C1q molecule, which leads to the activation of C1r, which in turn, cleave C1s. The C1r2s2 component now splits C4 and then C2, producing C4a, C4b, C2a, and C2b. C4b and C2b bind to form the classical pathway C3-convertase (C4b2b complex), which promotes cleavage of C3 into C3a and C3b. C3b then binds the C3 convertase to from the C5 convertase (C4b2b3b complex). The alternative pathway is continuously activated as a result of spontaneous C3 hydrolysis. Factor P (properdin) is a positive regulator of the alternative pathway. Oligomerization of properdin stabilizes the C3

convertase, which can then cleave much more C3. The C3 molecules can bind to surfaces and recruit more B, D, and P activity, leading to amplification of the complement activation.

[0756] Acute phase response (APR) is a complex, systemic innate immune response for preventing infection and clearing potential pathogens. Numerous proteins are involved in APR and C-reactive protein is a well-characterized one.

[0757] It has been found, in accordance with the invention, that certain LNP are able to associate physically with platelets almost immediately after administration in vivo, while other LNP do not associate with platelets at all or only at background levels. Significantly, those LNPs that associate with platelets also apparently stabilize the platelet aggregates that are formed thereafter. Physical contact of the platelets with certain LNPs correlates with the ability of such platelets to remain aggregated or to form aggregates continuously for an extended period of time after administration. Such aggregates comprise activated platelets and innate immune cells such as macrophages and B cells.

23. Methods of Use

[0758] The polynucleotides, pharmaceutical compositions and formulations described herein are used in the preparation, manufacture and therapeutic use of to treat and/or prevent G6PC-related diseases, disorders or conditions, e.g., glycogen storage disease type 1a (GSD-Ia). In some embodiments, the polynucleotides, compositions and formulations of the invention are used to treat and/or prevent resulting from defective G6PC or deficient G6PC. In some embodiments, the polynucleotides, compositions and formulations of the invention are used to improve glycemia or improve blood glucose homeostasis resulting from defective G6PC or deficient G6PC. In some embodiments, the polynucleotides, compositions and formulations of the invention are used to treat and/or prevent the excessive accumulation of glycogen resulting from defective or deficient G6PC. In some embodiments, the polynucleotides, compositions and formulations of the invention are used to treat and/or prevent hepatomegaly, nephromegaly, hypercholesterolemia, hypertriglyceridemia, hyperuricemia, and/or lactic academia in a subject with defective or deficient G6PC, e.g., a subject with GSD-Ia. In some embodiments, the polynucleotides, compositions and formulations of the invention are used to treat and/or prevent hepatocellular adenoma and/or hepatocellular carcinoma.

[0759] In some embodiments, the polynucleotides, pharmaceutical compositions and formulations of the invention are used in methods for reducing the levels of glycogen in a subject in need thereof, e.g., a subject having GSD-Ia that is fasting, e.g., a subject who has not had a meal. In some embodiments, the polynucleotides, pharmaceutical compositions and formulations of the invention are used in methods for increasing the levels of glucose (e.g., increasing blood glucose levels) in a subject in need thereof, e.g., a subject having GSD-Ia that is fasting, e.g., a subject who has not had a meal. For instance, one aspect of the invention provides a method of alleviating the symptoms of GSD-Ia in a subject (e.g., a subject with GSD-Ia that is fasting) comprising the administration of a composition or formulation comprising a polynucleotide encoding G6PC to that subject (e.g., an mRNA encoding a G6PC polypeptide). In some embodiments, the subject with GSD-Ia has been fasting for 1, 1.5, 2, 2.5, 3, 3.5, 4, 4.5, 5, 5.5, 6, 6.5, 7, 7.5, 8, 8.5, 9, 9.5, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, or 24 or more hours. In some embodiments, the subject with GSD-Ia has been fasting for 1 to 24 hours, 1 to 12 hours, 2 to 24 hours, 2 to 12 hours, 3 to 12 hours, 4 to 10 hours, 6 to 24 hours, 6 to 12 hours, or 5 to 18 hours. In some embodiments, the subject has been fasting for more than one day, e.g., 1, 2, or 3 days.

[0760] In some embodiments, the polynucleotides, pharmaceutical compositions and formulations of the invention are used to reduce the level of glycogen, the method comprising administering to the subject an effective amount of a polynucleotide encoding a G6PC polypeptide. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in a meaningful reduction in the level of glycogen (e.g., a statistically significant reduction in glycogen relative to the level of glycogen in the subject prior to administration), within a short period of time (e.g., within about 1 hour, within about 2 hours, within about 4 hours, within about 6 hours, within about 8 hours, within about 12 hours, within about 16 hours, within about 20 hours, or within about 24 hours) after administration of the polynucleotide, pharmaceutical composition or formulation of the invention.

[0761] In some embodiments, the polynucleotides, pharmaceutical compositions and formulations of the invention are used to reduce the level of glucose-6-phosphate (G6P), the method comprising administering to the subject an effective amount of a polynucleotide encoding a G6PC polypeptide. In some embodiments, the G6PC

encoded by the polynucleotide hydrolyzes the G6P to glucose and phosphate. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in a meaningful reduction in the level of glucose-6-phosphate (e.g., a statistically significant reduction in glucose-6-phosphate relative to the level of glucose-6-phosphate in the subject prior to administration), within a short period of time (e.g., within about 1 hour, within about 2 hours, within about 4 hours, within about 6 hours, within about 8 hours, within about 12 hours, within about 16 hours, within about 20 hours, or within about 24 hours) after administration of the polynucleotide, pharmaceutical composition or formulation of the invention.

[0762] In some embodiments, the polynucleotides, pharmaceutical compositions and formulations of the invention are used to increase the level of glucose, the method comprising administering to the subject an effective amount of a polynucleotide encoding a G6PC polypeptide. In some embodiments, the polynucleotide encoding a G6PC polypeptide increases or improves blood glucose homeostasis in the subject, e.g., when the subject is fasted or has not eaten. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in an increase in the level of blood glucose to at least 70 mg/dl (e.g., at least 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117, 118, 119, 120, 125, 130, 135, 140, 145, or 150 mg/dl), within a short period of time (e.g., within about 1 hour, within about 2 hours, within about 4 hours, within about 6 hours, within about 8 hours, within about 12 hours, within about 16 hours, within about 20 hours, or within about 24 hours) after administration of the polynucleotide, pharmaceutical composition or formulation of the invention. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in an increase in the level of glucose to at least 3.5 mmol/l (e.g., at least 3.5, 3.6, 3.7, 3.8, 3.9, 4.0, 4.1, 4.2, 4.3, 4.4, 4.5, 4.6, 4.7, 4.8, 4.9, 5.0, 5.1, 5.2, 5.3, 5.4, 5.5, 5.6, 5.7, 5.8, 5.9, 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.7, 6.8, 6.9, 7.0, 7.1, 7.2, 7.3, 7.4, 7.5, 7.6, 7.7, 7.8, 7.9, 8.0, or 8.5 mmol/l), within a short period of time (e.g., within about 1 hour, within about 2 hours, within about 4 hours, within about 6 hours, within about 8 hours, within about 12 hours, within about 16 hours, within about 20 hours, or within about 24 hours) after administration of the polynucleotide, pharmaceutical composition or formulation

of the invention. For instance, one aspect of the invention provides a method of increasing the blood glucose levels in a subject with GSD-Ia that is fasting, comprising the administration of a composition or formulation comprising a polynucleotide encoding G6PC to that subject (e.g., an mRNA encoding a G6PC polypeptide). In some embodiments, administering the composition or formulation alleviates a symptom of GSD-Ia. In some embodiments, administering the composition or formulation alleviates a symptom of GSD-Ia, e.g., improves glycemia or blood glucose homeostasis in the subject.

[0763] In some embodiments, the administration of an effective amount of a polynucleotide, pharmaceutical composition or formulation of the invention reduces the levels of a biomarker of GSD-Ia. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in reduction in the level of one or more biomarkers of GSD-Ia, within a short period of time (e.g., within about 1 hour, within about 2 hours, within about 4 hours, within 6 hours, within about 8 hours, within about 12 hours, within about 16 hours, within about 20 hours, or within about 24 hours) after administration of the polynucleotide, pharmaceutical composition or formulation of the invention.

[0764] In some embodiments, the administration of an effective amount of a polynucleotide, pharmaceutical composition or formulation of the invention increases the levels of a biomarker of GSD-Ia. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in an increase in the level of one or more biomarkers of GSD-Ia, within a short period of time (e.g., within about 1 hour, within about 2 hours, within about 4 hours, within 6 hours, within about 8 hours, within about 12 hours, within about 16 hours, within about 20 hours, or within about 24 hours) after administration of the polynucleotide, pharmaceutical composition or formulation of the invention.

[0765] Replacement therapy is a potential treatment for GSD-Ia. Thus, in certain aspects of the invention, the polynucleotides, e.g., mRNA, disclosed herein comprise one or more sequences encoding a G6PC polypeptide that is suitable for use in gene replacement therapy for GSD-Ia. In some embodiments, the present disclosure treats a lack of G6PC or G6PC activity, or decreased or abnormal G6PC activity in a subject by providing a polynucleotide, e.g., mRNA, that encodes a G6PC polypeptide to the subject. In some embodiments, the polynucleotide is sequence-optimized. In some embodiments, the polynucleotide (e.g., an mRNA) comprises a nucleic acid sequence

(e.g., an ORF) encoding a G6PC polypeptide, wherein the nucleic acid is sequence-optimized, e.g., by modifying its G/C, uridine, or thymidine content, and/or the polynucleotide comprises at least one chemically modified nucleoside. In some embodiments, the polynucleotide comprises a miRNA binding site, e.g., a miRNA binding site that binds miRNA-142 and/or a miRNA binding site that binds miRNA-126. In some embodiments, the polypeptide does not comprise a miRNA binding site.

[0766] In some embodiments, the administration of a composition or formulation comprising polynucleotide, pharmaceutical composition or formulation of the invention

[0767] to a subject results in a decrease in glycogen to a level at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or to 100% lower than the level observed prior to the administration of the composition or formulation. In some embodiments, the administration of a composition or formulation comprising polynucleotide, pharmaceutical composition or formulation of the invention to a subject results in a decrease in glucose-6-phosphate (G6P) to a level at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or to 100% lower than the level observed prior to the administration of the composition or formulation. In some embodiments, the decrease in glycogen and/or G6P is in hepatic cells or tissue. In some embodiments, the decrease in glycogen and/or G6P is in blood, plasma or serum.

[0768] In some embodiments, the administration of a composition or formulation comprising polynucleotide, pharmaceutical composition or formulation of the invention to a subject results in an increase in glucose to a level at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or to 100% lower than the level observed prior to the administration of the composition or formulation. In some embodiments, the increase in glucose is in hepatic cells or tissue. In some embodiments, the increase in glucose is in blood, plasma or serum.

[0769] In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in expression of G6PC in cells of the subject. In some embodiments, administering the polynucleotide, pharmaceutical composition or formulation of the invention results in an increase of G6PCexpression and/or enzymatic activity in the subject. For example, in some embodiments, the polynucleotides of the present invention are used in methods of administering a composition or formulation comprising an mRNA encoding a G6PC polypeptide to a subject, wherein the method results in an increase of G6PC expression and/or enzymatic activity in at least some cells of a subject.

[0770] In some embodiments, the administration of a composition or formulation comprising an mRNA encoding a G6PC polypeptide to a subject results in an increase of G6PC expression and/or enzymatic activity in cells subject to a level at least 1%, at least 2%, at least 5%, at least 10%, at least 15%, at least 20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or to 100% or more of the expression and/or activity level expected in a normal subject, *e.g.*, a human not suffering from GSD-Ia. In some embodiments, the administration of a composition or formulation comprising an mRNA encoding a G6PC polypeptide to a subject results in an increase of 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 505%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, or 95% or more of G6PC expression and/or enzymatic activity in cells of a subject, e.g., a human subject with GSD-Ia.

[0771] In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in expression of G6PC protein in at least some of the cells of a subject that persists for a period of time sufficient to allow significant glucose production to occur. In some embodiments, the administration of the polynucleotide, pharmaceutical composition or formulation of the invention results in expression of G6PC protein in at least some of the cells of a subject that persists for a period of time sufficient to allow significant amounts of hydrolysis of glucose-6-phosphate (G6P) to glucose and phosphate to occur.

[0772] In some embodiments, the expression of the encoded polypeptide is increased. In some embodiments, the polynucleotide increases G6PC expression and/or enzymatic activity levels in cells when introduced into those cells, *e.g.,* by at least 1%, at least 2%, at least 3%, at least 4%, at least 5%, at least 10%, at least 15%, at least

20%, at least 25%, at least 30%, at least 35%, at least 40%, at least 45%, at least 50%, at least 55%, at least 60%, at least 65%, at least 70%, at least 75%, at least 80%, at least 85%, at least 90%, at least 95%, or to 100% with respect to the G6PC expression and/or enzymatic activity level in the cells before the polypeptide is introduced in the cells.

[0773] In some embodiments, the method or use comprises administering a polynucleotide, *e.g.,* mRNA, comprising a nucleotide sequence having sequence similarity to a polynucleotide selected from the group of SEQ ID NOs:8, 11-20, 22-38, and 207, wherein the polynucleotide encodes an G6PC polypeptide.

[0774] Other aspects of the present disclosure relate to transplantation of cells containing polynucleotides to a mammalian subject. Administration of cells to mammalian subjects is known to those of ordinary skill in the art, and includes, but is not limited to, local implantation (*e.g.,* topical or subcutaneous administration), organ delivery or systemic injection (*e.g.,* intravenous injection or inhalation), and the formulation of cells in pharmaceutically acceptable carriers.

[0775] In some embodiments, the polynucleotides (e.g., mRNA), pharmaceutical compositions and formulations used in the methods of the invention comprise a uracil-modified sequence encoding a G6PC polypeptide disclosed herein and a miRNA binding site disclosed herein, e.g., a miRNA binding site that binds to miR-142 and/or a miRNA binding site that binds to miR-126. In some embodiments, the uracil-modified sequence encoding a G6PC polypeptide comprises at least one chemically modified nucleobase, e.g., N1-methylpseudouracil or 5-methoxyuracil. In some embodiments, at least 95% of a type of nucleobase (e.g., uracil) in a uracil-modified sequence encoding a G6PC polypeptide of the invention are modified nucleobases. In some embodiments, at least 95% of uracil in a uracil-modified sequence encoding a G6PC polypeptide is 1-N-methylpseudouridine or 5-methoxyuridine. In some embodiments, the polynucleotide (e.g., a RNA, e.g., a mRNA) disclosed herein is formulated with a delivery agent comprising, e.g., a compound having the Formula (I), e.g., any of Compounds 1-232, e.g., Compound II; a compound having the Formula (III), (IV), (V), or (VI), e.g., any of Compounds 233-342, e.g., Compound VI; or a compound having the Formula (VIII), e.g., any of Compounds 419-428, e.g., Compound I, or any combination thereof. In some embodiments, the delivery agent comprises Compound II, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0 or

about 50:10:38.5:1.5. In some embodiments, the delivery agent comprises Compound VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio in the range of about 30 to about 60 mol% Compound II or VI (or related suitable amino lipid) (e.g., 30-40, 40-45, 45-50, 50-55 or 55-60 mol% Compound II or VI (or related suitable amino lipid)), about 5 to about 20 mol% phospholipid (or related suitable phospholipid or "helper lipid") (e.g., 5-10, 10-15, or 15-20 mol% phospholipid (or related suitable phospholipid or "helper lipid")), about 20 to about 50 mol% cholesterol (or related sterol or "non-cationic" lipid) (e.g., about 20-30, 30-35, 35-40, 40-45, or 45-50 mol% cholesterol (or related sterol or "non-cationic" lipid)) and about 0.05 to about 10 mol% PEG lipid (or other suitable PEG lipid) (e.g., 0.05-1, 1-2, 2-3, 3-4, 4-5, 5-7, or 7-10 mol% PEG lipid (or other suitable PEG lipid)). An exemplary delivery agent can comprise mole ratios of, for example, 47.5:10.5:39.0:3.0 or 50:10:38.5:1.5. In certain instances, an exemplary delivery agent can comprise mole ratios of, for example, 47.5:10.5:39.0:3; 47.5:10:39.5:3; 47.5:11:39.5:2; 47.5:10.5:39.5:2.5; 47.5:11:39:2.5; 48.5:10:38.5:3; 48.5:10.5:39:2; 48.5:10.5:38.5:2.5; 48.5:10.5:39.5:1.5; 48.5:10.5:38.0:3; 47:10.5:39.5:3; 47:10:40.5:2.5; 47:11:40:2; 47:10.5:39.5:3; 48:10.5:38.5:3; 48:10:39.5:2.5; 48:11:39:2; or 48:10.5:38.5:3. In some embodiments, the delivery agent comprises Compound II or VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0. In some embodiments, the delivery agent comprises Compound II or VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0 or about 50:10:38.5:1.5.

[0776] The skilled artisan will appreciate that the therapeutic effectiveness of a drug or a treatment of the instant invention can be characterized or determined by measuring the level of expression of an encoded protein (e.g., enzyme) in a sample or in samples taken from a subject (e.g., from a preclinical test subject (rodent, primate, etc.) or from a clinical subject (human). Likewise, the therapeutic effectiveness of a drug or a treatment of the instant invention can be characterized or determined by measuring the level of activity of an encoded protein (e.g., enzyme) in a sample or in samples taken from a subject (e.g., from a preclinical test subject (rodent, primate, etc.) or from a clinical subject (human). Furthermore, the therapeutic effectiveness of a drug or a treatment of the instant invention can be characterized or determined by measuring the level of an appropriate biomarker in sample(s) taken from a subject. Levels of protein and/or biomarkers can be determined post-administration with a

single dose of an mRNA therapeutic of the invention or can be determined and/or monitored at several time points following administration with a single dose or can be determined and/or monitored throughout a course of treatment, e.g., a multi-dose treatment.

[0777] GSD-Ia is associated with the impaired ability of G6PC to hydrolyze glucose-6-phosphate (G6P) to glucose and phosphate. Accordingly, subjects with GSD-Ia commonly show high levels of glycogen or and/or G6P in their cells and tissues, and low levels of glucose in their blood/plasma.

[0778] GSD-Ia is an autosomal recessive metabolic disease characterized by an impaired ability to maintain glucose homeostasis. GSD-Ia patients typically present with fasting hypoglycemia, hepatomegaly, nephromegaly, hyperlipidemia, hyperuricemia, lactic academia, and/or growth retardation. GSD-Ia patients can be asymptomatic carriers of the disorder or suffer from the various symptoms associated with the disease. Subjects with GSD-Ia commonly show higher levels of glycogen and/or glucose-6-phosphate (G6P) in cells and tissues (e.g., in hepatic cells, in liver, in kidneys, in the intestine) when fasted. Subjects with GSD-Ia also often show higher levels of triglycerides and/or cholesterol in their blood, plasma, or serum, and/or higher levels of lactate, when fasted. Subjects with GSD-Ia also commonly present with lower levels of glucose, e.g., in blood, plasma or serum glucose, when fasted. Unless otherwise specified, the methods of treating GSD-Ia patients or human subjects disclosed herein include treatment of both asymptomatic carriers and those individuals with abnormal levels of biomarkers.

G6PC Protein Expression Levels

[0779] Certain aspects disclosed herein feature measurement, determination and/or monitoring of the expression level or levels of glucose-6-phosphatase (G6Pase or G6PC) protein in a subject, for example, in an animal (e.g., rodents, primates, and the like) or in a human subject. Animals include normal, healthy or wildtype animals, as well as animal models for use in understanding GSD-Ia and treatments thereof. Exemplary animal models include rodent models, for example, G6PC deficient mice also referred to as G6PC mice, e.g., global *G6pc* knockout mice (G6pc(-/-)) or liver-specific knockout mice (L-G6pc(-/-)) (see Lei et al., Nat. Genet., 1996, 13:203-209; Mutel et al., J. Hepatol., 2011, 54:529-537).

[0780] G6PC protein expression levels can be measured or determined by any art-recognized method for determining protein levels in biological samples, e.g., from blood or tissue samples or a needle biopsy. The term "level" or "level of a protein" as used herein, preferably means the weight, mass or concentration of the protein within a sample or a subject. It will be understood by the skilled artisan that in certain embodiments the sample may be subjected, e.g., to any of the following: purification, precipitation, separation, e.g. centrifugation and/or HPLC, and subsequently subjected to determining the level of the protein, e.g., using mass and/or spectrometric analysis. In exemplary embodiments, enzyme-linked immunosorbent assay (ELISA) can be used to determine protein expression levels. In other exemplary embodiments, protein purification, separation and LC-MS can be used as a means for determining the level of a protein according to the invention. In some embodiments, an mRNA therapy disclosed herein (e.g., a single intravenous dose) results in increased G6PC protein expression levels in the liver tissue of the subject (e.g., 1.5-fold, 2-fold, 3-fold, 4-fold, 5-fold, 6-fold, 7-fold, 8-fold, 9-fold, 10-fold, 20-fold, 30-fold, 40-fold, 50-fold increase and/or increased to at least 1%, 2%, 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, at least 60%, at least 70%, at least 75%, 80%, at least 85%, at least 90%, at least 95%, or at least 100% of normal levels) for at least 6 hours, at least 12 hours, at least 24 hours, at least 36 hours, at least 48 hours, at least 60 hours, at least 72 hours, at least 84 hours, at least 96 hours, at least 108 hours, at least 120 hours, at least 132 hours, at least 144 hours, at least 156 hours, at least 168 hours, at least 192 hours, at least 216 hours, at least 240 hours, at least 264 hours, at least 288 hours, at least 312 hours, or at least 336 hours after administration of a single dose of the mRNA therapy. In some embodiments, an mRNA therapy of the invention (e.g., a single intravenous dose) results in increased glucose levels in the blood, plasma, or liver tissue of the subject (e.g., at least 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117, 118, 119, 120, 125, 130, 135, 140, 145, or 150 mg/dl) for at least 6 hours, at least 12 hours, at least 24 hours, at least 36 hours, at least 48 hours, at least 60 hours, at least 72 hours, at least 84 hours, at least 96 hours, at least 108 hours, at least 120 hours, at least 132 hours, at least 144 hours, at least 156 hours, at least 168 hours, at least 192 hours, at least 216 hours, at least 240 hours, at least 264 hours, at least 288 hours, at least 312 hours, or at least 336 hours after administration of a single dose of the mRNA

therapy. In some embodiments, an mRNA therapy of the invention (e.g., a single intravenous dose) results in increased glucose levels in the blood, plasma, or liver tissue of the subject (e.g., at least 3.5, 3.6, 3.7, 3.8, 3.9, 4.0, 4.1, 4.2, 4.3, 4.4, 4.5, 4.6, 4.7, 4.8, 4.9, 5.0, 5.1, 5.2, 5.3, 5.4, 5.5, 5.6, 5.7, 5.8, 5.9, 6.0, 6.1, 6.2, 6.3, 6.4, 6.5, 6.7, 6.8, 6.9, 7.0, 7.1, 7.2, 7.3, 7.4, 7.5, 7.6, 7.7, 7.8, 7.9, 8.0, or 8.5 mmol/l) for at least 6 hours, at least 12 hours, at least 24 hours, at least 36 hours, at least 48 hours, at least 60 hours, at least 72 hours, at least 84 hours, at least 96 hours, at least 108 hours, at least 1202 hours, at least 132 hours, at least 144 hours, at least 156 hours, at least 168 hours, at least 192 hours, at least 216 hours, at least 240 hours, at least 264 hours, at least 288 hours, at least 312 hours, or at least 336 hours after administration of a single dose of the mRNA therapy. In some embodiments, an mRNA therapy of the invention (e.g., a single intravenous dose) results in meaningfully decreased glycogen and/or G6P levels (e.g., statistically significant decrease relative to prior to administration of the mRNA therapy) in the blood, plasma, or liver tissue of the subject (e.g., 100, 125, 150, 175, 200, 225, 250, 275, 300, 325, 350, 375, 400, 425, 450, 475, 500, 525, 550, 575, 600, 625, 650, 675, 700, 725, 750, 775, 800, 825, 850, 875, 900, 925, 950, 975, 1000, 1025, 1050, 1075, 1100, 1125, 1150, 1175 or 1,200 μM or lower) for at least 6 hours, at least 12 hours, at least 24 hours, at least 36 hours, at least 48 hours, at least 60 hours, at least 72 hours, at least 84 hours, at least 96 hours, at least 108 hours, at least 1202 hours, at least 132 hours, at least 144 hours, at least 156 hours, at least 168 hours, at least 192 hours, at least 216 hours, at least 240 hours, at least 264 hours, at least 288 hours, at least 312 hours, or at least 336 hours after administration of a single dose of the mRNA therapy.

G6PC Protein Activity

[0781] In subjects with GSD-Ia, G6PC enzymatic activity is reduced compared to a normal physiological activity level. Further aspects disclosed herein feature measurement, determination and/or monitoring of the activity level(s) (*i.e.*, enzymatic activity level(s)) of G6PC protein in a subject, for example, in an animal (e.g., rodent, primate, and the like) or in a human subject. Activity levels can be measured or determined by any art-recognized method for determining enzymatic activity levels in biological samples. The term "activity level" or "enzymatic activity level" as used herein, preferably means the activity of the enzyme per volume, mass or weight of sample or total protein within a sample. In exemplary embodiments, the "activity

level" or "enzymatic activity level" is described in terms of units per milliliter of fluid (e.g., bodily fluid, e.g., serum, plasma, urine and the like) or is described in terms of units per weight of tissue or per weight of protein (*e.g.*, total protein) within a sample. Units ("U") of enzyme activity can be described in terms of weight or mass of substrate hydrolyzed per unit time. In certain embodiments of the invention feature G6PC activity described in terms of U/ml plasma or U/mg protein (tissue), where units ("U") are described in terms of nmol substrate hydrolyzed per hour (or nmol/hr).

[0782] In certain embodiments, an mRNA therapy of the invention features a pharmaceutical composition comprising a dose of mRNA effective to result in at least 5 U/mg, at least 10 U/mg, at least 20 U/mg, at least 30 U/mg, at least 40 U/mg, at least 50 U/mg, at least 60 U/mg, at least 70 U/mg, at least 80 U/mg, at least 90 U/mg, at least 100 U/mg, or at least 150 U/mg of G6PC activity in tissue (e.g., liver) between 2 and 6 hours, between 6 and 12 hours, or between 12 and 24, between 24 and 48, or between 48 and 72 hours post administration (e.g., at 48 or at 72 hours post administration).

[0783] In exemplary embodiments, an mRNA therapy described herein features a pharmaceutical composition comprising a single intravenous dose of mRNA that results in the above-described levels of activity. In another embodiment, an mRNA therapy of the invention features a pharmaceutical composition which can be administered in multiple single unit intravenous doses of mRNA that maintain the above-described levels of activity.

GSD-Ia Biomarkers

[0784] Further aspects of the invention feature determining the level (or levels) of a biomarker determined in a sample as compared to a level (e.g., a reference level) of the same or another biomarker in another sample, e.g., from the same subject, from another subject, from a control and/or from the same or different time points, and/or a physiologic level, and/or an elevated level, and/or a supraphysiologic level, and/or a level of a control. The skilled artisan will be familiar with physiologic levels of biomarkers, for example, levels in normal or wildtype animals, normal or healthy subjects, and the like, in particular, the level or levels characteristic of subjects who are healthy and/or normal functioning. As used herein, the phrase "elevated level" means amounts greater than normally found in a normal or wildtype preclinical animal or in a normal or healthy subject, e.g. a human subject. As used herein, the

term “supraphysiologic” means amounts greater than normally found in a normal or wildtype preclinical animal or in a normal or healthy subject, e.g. a human subject, optionally producing a significantly enhanced physiologic response. As used herein, the term "comparing" or "compared to" preferably means the mathematical comparison of the two or more values, e.g., of the levels of the biomarker(s). It will thus be readily apparent to the skilled artisan whether one of the values is higher, lower or identical to another value or group of values if at least two of such values are compared with each other. Comparing or comparison to can be in the context, for example, of comparing to a control value, e.g., as compared to a reference blood, serum, plasma, and/or tissue (e.g., liver) biomarker level, in said subject prior to administration (e.g., in a person suffering from GSD-Ia) or in a normal or healthy subject. Comparing or comparison to can also be in the context, for example, of comparing to a control value, e.g., as compared to a reference blood, serum, plasma and/or tissue (e.g., liver) biomarker level in said subject prior to administration (e.g., in a person suffering from GSD-Ia) or in a normal or healthy subject. As used herein, a “control” is preferably a sample from a subject wherein the GSD-Ia status of said subject is known. In one embodiment, a control is a sample of a healthy patient. In another embodiment, the control is a sample from at least one subject having a known GSD-Ia status, for example, a severe, mild, or healthy GSD-Ia status, e.g. a control patient. In another embodiment, the control is a sample from a subject not being treated for GSD-Ia. In a still further embodiment, the control is a sample from a single subject or a pool of samples from different subjects and/or samples taken from the subject(s) at different time points.

[0785] The term "level" or "level of a biomarker" as used herein, preferably means the mass, weight or concentration of a biomarker of the invention within a sample or a subject. It will be understood by the skilled artisan that in certain embodiments the sample may be subjected to, e.g., one or more of the following: substance purification, precipitation, separation, e.g. centrifugation and/or HPLC and subsequently subjected to determining the level of the biomarker, e.g. using mass spectrometric analysis. In certain embodiments, LC-MS can be used as a means for determining the level of a biomarker according to the invention.

[0786] The term "determining the level" of a biomarker as used herein can mean methods which include quantifying an amount of at least one substance in a sample

from a subject, for example, in a bodily fluid from the subject (e.g., serum, plasma, urine, lymph, etc.) or in a tissue of the subject (e.g., liver, etc.).

[0787] The term "reference level" as used herein can refer to levels (e.g., of a biomarker) in a subject prior to administration of an mRNA therapy of the invention (e.g., in a person suffering from GSD-Ia) or in a normal or healthy subject.

[0788] As used herein, the term "normal subject" or "healthy subject" refers to a subject not suffering from symptoms associated with GSD-Ia. Moreover, a subject will be considered to be normal (or healthy) if it has no mutation of the functional portions or domains of the G6PC gene and/or no mutation of the G6PC gene resulting in a reduction of or deficiency of the enzyme G6PC or the activity thereof, resulting in symptoms associated with GSD-Ia. Said mutations will be detected if a sample from the subject is subjected to a genetic testing for such G6PC mutations. In certain embodiments of the present invention, a sample from a healthy subject is used as a control sample, or the known or standardized value for the level of biomarker from samples of healthy or normal subjects is used as a control.

[0789] Since G6PC hydrolyzes G6P into glucose and phosphate during gluconeogenesis and glycogenolysis, low levels of glucose and/or phosphate, and high levels of glycogen and/or G6P, during fasting conditions is indicative that a subject has GSD-Ia or should be treated for GSD-Ia. Thus, glucose, phosphate, glycogen, and G6P can be used as biomarkers for GSD-Ia.

[0790] In some embodiments, comparing the level of the biomarker in a sample from a subject in need of treatment for GSD-Ia or in a subject being treated for GSD-Ia to a control level of the biomarker comprises comparing the level of the biomarker in the sample from the subject (in need of treatment or being treated for GSD-Ia) to a baseline or reference level, wherein if a level of the biomarker in the sample from the subject (in need of treatment or being treated for GSD-Ia) is elevated, increased or higher compared to the baseline or reference level, this is indicative that the subject is suffering from GSD-Ia and/or is in need of treatment; and/or wherein if a level of the biomarker in the sample from the subject (in need of treatment or being treated for GSD-Ia) is decreased or lower or the same compared to the baseline level this is indicative that the subject is not suffering from, is successfully being treated for GSD-Ia, or is not in need of treatment for GSD-Ia. The stronger the reduction (e.g., at least 1.5-fold, at least 2-fold, at least 3-fold, at least 4-fold, at least 5-fold, at least 6-fold, at least 7-fold, at least 8-fold, at least 10-fold, at least 20-fold, at least-30 fold, at least

40-fold, at least 50-fold reduction and/or at least 2%, at least 5%, at least 10%, at least 20%, at least 30% at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% reduction) of the level of a biomarker, within a certain time period, e.g., within 2 hours, within 4 hours, within 6 hours, within 12 hours, 24 hours, 36 hours, 48 hours, 60 hours, or 72 hours, and/or for a certain duration of time, e.g., 48 hours, 72 hours, 96 hours, 120 hours, 144 hours, 1 week, 2 weeks, 3 weeks, 4 weeks, 1 month, 2 months, 3 months, 4 months, 5 months, 6 months, 7 months, 8 months, 9 months, 10 months, 11 months, 12 months, 18 months, 24 months, etc. the more successful is a therapy, such as for example an mRNA therapy of the invention (e.g., a single dose or a multiple regimen). In some embodiments, the biomarker that is elevated, increased or higher compared to the baseline or reference level, and is indicative of GSD-Ia, is glycogen and/or glucose-6-phosphate (G6P). In some embodiments, the biomarker that is elevated, increased or higher compared to the baseline or reference level, and is indicative of GSD-Ia, is lactate. In some embodiments, the biomarker that is elevated, increased or higher compared to the baseline or reference level, and is indicative of GSD-Ia, is triglycerides and/or cholesterol, e.g., in blood, plasma, or serum.

[0791] A reduction of at least about 5%, at least about 10%, at least about 20%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least 100% or more of the level of biomarker (e.g., glycogen and/or G6P), in particular, in bodily fluid (e.g., plasma, serum, urine, e.g., urinary sediment) or in tissue(s) in a subject (e.g., liver), within 12 hours, 1, 2, 3, 4, 5, 6 or more days following administration is indicative of a dose suitable for successful treatment GSD-Ia, wherein reduction as used herein, preferably means that the level of biomarker determined at the end of a specified time period (e.g., post-administration, for example, of a single intravenous dose) is compared to the level of the same biomarker determined at the beginning of said time period (e.g., pre-administration of said dose). Exemplary time periods include 6, 12, 24, 48, 72, 96, 120 or 144 hours post administration, in particular 24, 48, 72 or 96 hours post administration.

[0792] A sustained reduction in some substrate levels (e.g., biomarkers, such as glycogen and/or G6P) is particularly indicative of mRNA therapeutic dosing and/or administration regimens successful for treatment of GSD-Ia. Such sustained reduction can be referred to herein as "duration" of effect. In exemplary

embodiments, a reduction of at least about 5%, at least about 10%, at least about 20%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, or at least about 95% at least about 96%, at least about 97%, at least about 98%, at least about 99%, at least about 100% or more of the level of biomarker, in particular, in a bodily fluid (e.g., plasma, serum, urine, e.g., urinary sediment) or in tissue(s) in a subject (e.g., liver), within 12 hours, 1, 2, 3, 4, 5, 6, 7, 8 or more days following administration is indicative of a successful therapeutic approach. In exemplary embodiments, sustained reduction in substrate (e.g., biomarker) levels in one or more samples (e.g., fluids and/or tissues) is preferred. For example, mRNA therapies resulting in sustained reduction in a biomarker, optionally in combination with sustained reduction of said biomarker in at least one tissue, preferably two, three, four, five or more tissues, is indicative of successful treatment.

[0793] In other embodiments, comparing the level of the biomarker in a sample from a subject in need of treatment for GSD-Ia or in a subject being treated for GSD-Ia to a control level of the biomarker comprises comparing the level of the biomarker in the sample from the subject (in need of treatment or being treated for GSD-Ia) to a baseline or reference level, wherein if a level of the biomarker in the sample from the subject (in need of treatment or being treated for GSD-Ia) is reduced, decreased or lower compared to the baseline or reference level, this is indicative that the subject is suffering from GSD-Ia and/or is in need of treatment; and/or wherein if a level of the biomarker in the sample from the subject (in need of treatment or being treated for GSD-Ia) is increased or higher or the same compared to the baseline level this is indicative that the subject is not suffering from, is successfully being treated for GSD-Ia, or is not in need of treatment for GSD-Ia. The stronger the increase (e.g., at least 1.5-fold, at least 2-fold, at least 3-fold, at least 4-fold, at least 5-fold, at least 6-fold, at least 7-fold, at least 8-fold, at least 10-fold, at least 20-fold, at least-30 fold, at least 40-fold, at least 50-fold increase and/or at least 2%, at least 5%, at least 10%, at least 20%, at least 30% at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% increase) of the level of a biomarker, within a certain time period, e.g., within 2 hours, within 4 hours, within 6 hours, within 12 hours, 24 hours, 36 hours, 48 hours, 60 hours, or 72 hours, and/or for a certain duration of time, e.g., 48 hours, 72 hours, 96 hours, 120 hours, 144 hours, 1 week, 2 weeks, 3 weeks, 4 weeks, 1 month, 2 months, 3 months, 4 months, 5 months, 6 months, 7 months, 8

months, 9 months, 10 months, 11 months, 12 months, 18 months, 24 months, etc. the more successful is a therapy, such as for example an mRNA therapy of the invention (e.g., a single dose or a multiple regimen). In some embodiments, the biomarker that is reduced, decreased or lower compared to the baseline or reference level, and is indicative of GSD-Ia, is glucose, e.g., glucose in blood, plasma or serum. In some embodiments, the biomarker that is reduced, decreased or lower compared to the baseline or reference level, and is indicative of GSD-Ia, is phosphate.

[0794] An increase of at least about 5%, at least about 10%, at least about 20%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least 100% or more of the level of biomarker (e.g., glucose), in particular, in bodily fluid (e.g., plasma, serum, urine, e.g., urinary sediment) or in tissue(s) in a subject (e.g., liver), within 12 hours, 1, 2, 3, 4, 5, 6 or more days following administration is indicative of a dose suitable for successful treatment GSD-Ia, wherein increase as used herein, preferably means that the level of biomarker determined at the end of a specified time period (e.g., post-administration, for example, of a single intravenous dose) is compared to the level of the same biomarker determined at the beginning of said time period (e.g., pre-administration of said dose). Exemplary time periods include 6, 12, 24, 48, 72, 96, 120 or 144 hours post administration, in particular 24, 48, 72 or 96 hours post administration.

[0795] A sustained increase in some substrate levels (e.g., biomarkers, e.g., glucose) is particularly indicative of mRNA therapeutic dosing and/or administration regimens successful for treatment of GSD-Ia. Such sustained increase can be referred to herein as "duration" of effect. In exemplary embodiments, an increase of at least about 5%, at least about 10%, at least about 20%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, or at least about 95% at least about 96%, at least about 97%, at least about 98%, at least about 99%, at least about 100% or more of the level of biomarker, in particular, in a bodily fluid (e.g., plasma, serum, urine, e.g., urinary sediment) or in tissue(s) in a subject (e.g., liver), within 12 hours, 1, 2, 3, 4, 5, 6, 7, 8 or more days following administration is indicative of a successful therapeutic approach. In exemplary embodiments, sustained increase in substrate (e.g., biomarker) levels in one or more samples (e.g., fluids and/or tissues) is preferred. For example, mRNA therapies resulting in sustained

reduction in a biomarker, optionally in combination with sustained reduction of said biomarker in at least one tissue, preferably two, three, four, five or more tissues, is indicative of successful treatment.

[0796] In some embodiments, a single dose of an mRNA therapy described herein is about 0.2 to about 0.8 mpk. about 0.3 to about 0.7 mpk, about 0.4 to about 0.8 mpk, or about 0.5 mpk. In another embodiment, a single dose of an mRNA therapy of the invention is less than 1.5 mpk, less than 1.25 mpk, less than 1 mpk, or less than 0.75 mpk.

[0797] In some embodiments, a subject with GSD-Ia, at risk of developing GSD-Ia, or in need of treatment for GSD-Ia, has a larger liver compared to the size of a liver in a normal or healthy subject, i.e., a subject without GSD-Ia. In some embodiments, a larger liver in a subject indicates that the subject has GSD-Ia, is at risk of having GSD-Ia, or should be treated for GSD-Ia. In some embodiments, a subject having a liver that is 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, or 50% larger than the liver in a normal or healthy subject, or to a reference weight for a normal liver, indicates that the subject has GSD-Ia or should be treated for GSD-Ia.

[0798] In some embodiments, the biomarker or biomarkers are measured in a sample taken from a subject in need of treatment for GSD-Ia or in a subject being treated for GSD-Ia that has been fasting (e.g., a subject that has not eaten for some time, e.g., for 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 or more hours). In some embodiments, the control sample is taken from a healthy or normal subject that has also been fasting, or the reference level of the biomarker or biomarkers are the levels in a healthy or normal subject that was fasting.

24. Compositions and Formulations for Use

[0799] Certain aspects of the invention are directed to compositions or formulations comprising any of the polynucleotides disclosed above.

[0800] In some embodiments, the composition or formulation comprises:

(i) a polynucleotide (e.g., a RNA, e.g., an mRNA) comprising a sequence-optimized nucleotide sequence (e.g., an ORF) encoding a G6PC polypeptide (e.g., the wild-type sequence, functional fragment, or variant thereof), wherein the polynucleotide comprises at least one chemically modified nucleobase, e.g., N1-methylpseudouracil or 5-methoxyuracil (e.g., wherein at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about

70%, at least about 80%, at least about 90%, at least about 95%, at least about 99%, or 100% of the uracils are N1-methylpseudouracils or 5-methoxyuracils), and wherein the polynucleotide further comprises a miRNA binding site, e.g., a miRNA binding site that binds to miR-142 (e.g., a miR-142-3p or miR-142-5p binding site) and/or a miRNA binding site that binds to miR-126 (e.g., a miR-126-3p or miR-126-5p binding site); and

(ii) a delivery agent comprising, e.g., a compound having the Formula (I), e.g., any of Compounds 1-232, e.g., Compound II; a compound having the Formula (III), (IV), (V), or (VI), e.g., any of Compounds 233-342, e.g., Compound VI; or a compound having the Formula (VIII), e.g., any of Compounds 419-428, e.g., Compound I, or any combination thereof. In some embodiments, the delivery agent is a lipid nanoparticle comprising Compound II, Compound VI, a salt or a stereoisomer thereof, or any combination thereof. In some embodiments, the delivery agent comprises Compound II, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 50:10:38.5:1.5. In some embodiments, the delivery agent comprises Compound II, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0. In some embodiments, the delivery agent comprises Compound VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 50:10:38.5:1.5. In some embodiments, the delivery agent comprises Compound VI, DSPC, Cholesterol, and Compound I or PEG-DMG, e.g., with a mole ratio of about 47.5:10.5:39.0:3.0.

[0801] In some embodiments, the uracil or thymine content of the ORF relative to the theoretical minimum uracil or thymine content of a nucleotide sequence encoding the G6PC polypeptide (%U_{TM} or %T_{TM}), is between about 100% and about 150%.

[0802] In some embodiments, the polynucleotides, compositions or formulations above are used to treat and/or prevent G6PC -related diseases, disorders or conditions, e.g., GSD-Ia.

25. Forms of Administration

[0803] The polynucleotides, pharmaceutical compositions and formulations of the invention described above can be administered by any route that results in a therapeutically effective outcome. These include, but are not limited to enteral (into the intestine), gastroenteral, epidural (into the dura matter), oral (by way of the mouth), transdermal, peridural, intracerebral (into the cerebrum),

intracerebroventricular (into the cerebral ventricles), epicutaneous (application onto the skin), intradermal, (into the skin itself), subcutaneous (under the skin), nasal administration (through the nose), intravenous (into a vein), intravenous bolus, intravenous drip, intraarterial (into an artery), intramuscular (into a muscle), intracardiac (into the heart), intraosseous infusion (into the bone marrow), intrathecal (into the spinal canal), intraperitoneal, (infusion or injection into the peritoneum), intravesical infusion, intravitreal, (through the eye), intracavernous injection (into a pathologic cavity) intracavitary (into the base of the penis), intravaginal administration, intrauterine, extra-amniotic administration, transdermal (diffusion through the intact skin for systemic distribution), transmucosal (diffusion through a mucous membrane), transvaginal, insufflation (snorting), sublingual, sublabial, enema, eye drops (onto the conjunctiva), in ear drops, auricular (in or by way of the ear), buccal (directed toward the cheek), conjunctival, cutaneous, dental (to a tooth or teeth), electro-osmosis, endocervical, endosinusial, endotracheal, extracorporeal, hemodialysis, infiltration, interstitial, intra-abdominal, intra-amniotic, intra-articular, intrabiliary, intrabronchial, intrabursal, intracartilaginous (within a cartilage), intracaudal (within the cauda equine), intracisternal (within the cisterna magna cerebellomedularis), intracorneal (within the cornea), dental intracornal, intracoronary (within the coronary arteries), intracorporus cavernosum (within the dilatable spaces of the corporus cavernosa of the penis), intradiscal (within a disc), intraductal (within a duct of a gland), intraduodenal (within the duodenum), intradural (within or beneath the dura), intraepidermal (to the epidermis), intraesophageal (to the esophagus), intragastric (within the stomach), intragingival (within the gingivae), intraileal (within the distal portion of the small intestine), intralesional (within or introduced directly to a localized lesion), intraluminal (within a lumen of a tube), intralymphatic (within the lymph), intramedullary (within the marrow cavity of a bone), intrameningeal (within the meninges), intraocular (within the eye), intraovarian (within the ovary), intrapericardial (within the pericardium), intrapleural (within the pleura), intraprostatic (within the prostate gland), intrapulmonary (within the lungs or its bronchi), intrasinal (within the nasal or periorbital sinuses), intraspinal (within the vertebral column), intrasynovial (within the synovial cavity of a joint), intratendinous (within a tendon), intratesticular (within the testicle), intrathecal (within the cerebrospinal fluid at any level of the cerebrospinal axis), intrathoracic (within the thorax), intratubular (within the tubules of an organ), intratympanic (within the aurus

media), intravascular (within a vessel or vessels), intraventricular (within a ventricle), iontophoresis (by means of electric current where ions of soluble salts migrate into the tissues of the body), irrigation (to bathe or flush open wounds or body cavities), laryngeal (directly upon the larynx), nasogastric (through the nose and into the stomach), occlusive dressing technique (topical route administration that is then covered by a dressing that occludes the area), ophthalmic (to the external eye), oropharyngeal (directly to the mouth and pharynx), parenteral, percutaneous, periarticular, peridural, perineural, periodontal, rectal, respiratory (within the respiratory tract by inhaling orally or nasally for local or systemic effect), retrobulbar (behind the pons or behind the eyeball), intramyocardial (entering the myocardium), soft tissue, subarachnoid, subconjunctival, submucosal, topical, transplacental (through or across the placenta), transtracheal (through the wall of the trachea), transtympanic (across or through the tympanic cavity), ureteral (to the ureter), urethral (to the urethra), vaginal, caudal block, diagnostic, nerve block, biliary perfusion, cardiac perfusion, photopheresis or spinal. In specific embodiments, compositions can be administered in a way that allows them cross the blood-brain barrier, vascular barrier, or other epithelial barrier. In some embodiments, a formulation for a route of administration can include at least one inactive ingredient.

[0804] The polynucleotides of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide or a functional fragment or variant thereof) can be delivered to a cell naked. As used herein in, "naked" refers to delivering polynucleotides free from agents that promote transfection. The naked polynucleotides can be delivered to the cell using routes of administration known in the art and described herein.

[0805] The polynucleotides of the present invention (e.g., a polynucleotide comprising a nucleotide sequence encoding a G6PC polypeptide or a functional fragment or variant thereof) can be formulated, using the methods described herein. The formulations can contain polynucleotides that can be modified and/or unmodified. The formulations can further include, but are not limited to, cell penetration agents, a pharmaceutically acceptable carrier, a delivery agent, a bioerodible or biocompatible polymer, a solvent, and a sustained-release delivery depot. The formulated polynucleotides can be delivered to the cell using routes of administration known in the art and described herein.

[0806] A pharmaceutical composition for parenteral administration can comprise at least one inactive ingredient. Any or none of the inactive ingredients used can have been approved by the US Food and Drug Administration (FDA). A non-exhaustive list of inactive ingredients for use in pharmaceutical compositions for parenteral administration includes hydrochloric acid, mannitol, nitrogen, sodium acetate, sodium chloride and sodium hydroxide.

[0807] Injectable preparations, for example, sterile injectable aqueous or oleaginous suspensions can be formulated according to the known art using suitable dispersing agents, wetting agents, and/or suspending agents. Sterile injectable preparations can be sterile injectable solutions, suspensions, and/or emulsions in nontoxic parenterally acceptable diluents and/or solvents, for example, as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that can be employed are water, Ringer's solution, U.S.P., and isotonic sodium chloride solution. Sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil can be employed including synthetic mono- or diglycerides. Fatty acids such as oleic acid can be used in the preparation of injectables. The sterile formulation can also comprise adjuvants such as local anesthetics, preservatives and buffering agents.

[0808] Injectable formulations can be sterilized, for example, by filtration through a bacterial-retaining filter, and/or by incorporating sterilizing agents in the form of sterile solid compositions that can be dissolved or dispersed in sterile water or other sterile injectable medium prior to use.

[0809] Injectable formulations can be for direct injection into a region of a tissue, organ and/or subject. As a non-limiting example, a tissue, organ and/or subject can be directly injected a formulation by intramyocardial injection into the ischemic region. (*See, e.g.,* Zangi et al. Nature Biotechnology 2013; the contents of which are herein incorporated by reference in its entirety).

[0810] To prolong the effect of an active ingredient, it is often desirable to slow the absorption of the active ingredient from subcutaneous or intramuscular injection. This can be accomplished by the use of a liquid suspension of crystalline or amorphous material with poor water solubility. The rate of absorption of the drug then depends upon its rate of dissolution which, in turn, can depend upon crystal size and crystalline form. Alternatively, delayed absorption of a parenterally administered drug form is accomplished by dissolving or suspending the drug in an oil vehicle. Injectable depot

forms are made by forming microencapsule matrices of the drug in biodegradable polymers such as polylactide-polyglycolide. Depending upon the ratio of drug to polymer and the nature of the particular polymer employed, the rate of drug release can be controlled. Examples of other biodegradable polymers include poly(orthoesters) and poly(anhydrides). Depot injectable formulations are prepared by entrapping the drug in liposomes or microemulsions that are compatible with body tissues.

26. Kits and Devices

a. Kits

[0811] The invention provides a variety of kits for conveniently and/or effectively using the claimed nucleotides of the present invention. Typically, kits will comprise sufficient amounts and/or numbers of components to allow a user to perform multiple treatments of a subject(s) and/or to perform multiple experiments.

[0812] In one aspect, the present invention provides kits comprising the molecules (polynucleotides) of the invention.

[0813] The kits can be for protein production (e.g., protein production in a subject following administration), comprising a first polynucleotide comprising a translatable region. The kit can further comprise packaging and instructions and/or a delivery agent to form a formulation composition. The delivery agent can comprise a saline, a buffered solution, a lipidoid or any delivery agent disclosed herein.

[0814] In some embodiments, the buffer solution can include sodium chloride, calcium chloride, phosphate and/or EDTA. In another embodiment, the buffer solution can include, but is not limited to, saline, saline with 2mM calcium, 5% sucrose, 5% sucrose with 2mM calcium, 5% Mannitol, 5% Mannitol with 2mM calcium, Ringer's lactate, sodium chloride, sodium chloride with 2mM calcium and mannose (*See, e.g.*, U.S. Pub. No. 20120258046; herein incorporated by reference in its entirety). In a further embodiment, the buffer solutions can be precipitated or it can be lyophilized. The amount of each component can be varied to enable consistent, reproducible higher concentration saline or simple buffer formulations. The components can also be varied to increase the stability of modified RNA in the buffer solution over a period of time and/or under a variety of conditions. In one aspect, the present invention provides kits for protein production, comprising: a polynucleotide

comprising a translatable region, provided in an amount effective to produce a desired amount of a protein encoded by the translatable region when introduced into a target cell; a second polynucleotide comprising an inhibitory nucleic acid, provided in an amount effective to substantially inhibit the innate immune response of the cell; and packaging and instructions.

[0815] In one aspect, the present invention provides kits for protein production, comprising a polynucleotide comprising a translatable region, wherein the polynucleotide exhibits reduced degradation by a cellular nuclease, and packaging and instructions.

[0816] In one aspect, the present invention provides kits for protein production, comprising a polynucleotide comprising a translatable region, wherein the polynucleotide exhibits reduced degradation by a cellular nuclease, and a mammalian cell suitable for translation of the translatable region of the first nucleic acid.

b. Devices

[0817] The present invention provides for devices that can incorporate polynucleotides that encode polypeptides of interest. These devices contain in a stable formulation the reagents to synthesize a polynucleotide in a formulation available to be immediately delivered to a subject in need thereof, such as a human patient

[0818] Devices for administration can be employed to deliver the polynucleotides of the present invention according to single, multi- or split-dosing regimens taught herein. Such devices are taught in, for example, International Application Publ. No. WO2013151666, the contents of which are incorporated herein by reference in their entirety.

[0819] Method and devices known in the art for multi-administration to cells, organs and tissues are contemplated for use in conjunction with the methods and compositions disclosed herein as embodiments of the present invention. These include, for example, those methods and devices having multiple needles, hybrid devices employing for example lumens or catheters as well as devices utilizing heat, electric current or radiation driven mechanisms.

[0820] According to the present invention, these multi-administration devices can be utilized to deliver the single, multi- or split doses contemplated herein. Such devices are taught for example in, International Application Publ. No. WO2013151666, the contents of which are incorporated herein by reference in their entirety.

[0821] In some embodiments, the polynucleotide is administered subcutaneously or intramuscularly via at least 3 needles to three different, optionally adjacent, sites simultaneously, or within a 60 minutes period (*e.g.*, administration to 4, 5, 6, 7, 8, 9, or 10 sites simultaneously or within a 60-minute period).

c. Methods and Devices utilizing catheters and/or lumens

[0822] Methods and devices using catheters and lumens can be employed to administer the polynucleotides of the present invention on a single, multi- or split dosing schedule. Such methods and devices are described in International Application Publication No. WO2013151666, the contents of which are incorporated herein by reference in their entirety.

d. Methods and Devices utilizing electrical current

[0823] Methods and devices utilizing electric current can be employed to deliver the polynucleotides of the present invention according to the single, multi- or split dosing regimens taught herein. Such methods and devices are described in International Application Publication No. WO2013151666, the contents of which are incorporated herein by reference in their entirety.

27. Definitions

[0824] In order that the present disclosure can be more readily understood, certain terms are first defined. As used in this application, except as otherwise expressly provided herein, each of the following terms shall have the meaning set forth below. Additional definitions are set forth throughout the application.

[0825] The invention includes embodiments in which exactly one member of the group is present in, employed in, or otherwise relevant to a given product or process. The invention includes embodiments in which more than one, or all of the group members are present in, employed in, or otherwise relevant to a given product or process.

[0826] In this specification and the appended claims, the singular forms "a", "an" and "the" include plural referents unless the context clearly dictates otherwise. The terms "a" (or "an"), as well as the terms "one or more," and "at least one" can be used interchangeably herein. In certain aspects, the term "a" or "an" means "single." In other aspects, the term "a" or "an" includes "two or more" or "multiple."

[0827] Furthermore, "and/or" where used herein is to be taken as specific disclosure of each of the two specified features or components with or without the other. Thus, the term "and/or" as used in a phrase such as "A and/or B" herein is intended to include "A and B," "A or B," "A" (alone), and "B" (alone). Likewise, the term "and/or" as used in a phrase such as "A, B, and/or C" is intended to encompass each of the following aspects: A, B, and C; A, B, or C; A or C; A or B; B or C; A and C; A and B; B and C; A (alone); B (alone); and C (alone).

[0828] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this disclosure is related. For example, the Concise Dictionary of Biomedicine and Molecular Biology, Juo, Pei-Show, 2nd ed., 2002, CRC Press; The Dictionary of Cell and Molecular Biology, 3rd ed., 1999, Academic Press; and the Oxford Dictionary Of Biochemistry And Molecular Biology, Revised, 2000, Oxford University Press, provide one of skill with a general dictionary of many of the terms used in this disclosure.

[0829] Wherever aspects are described herein with the language "comprising," otherwise analogous aspects described in terms of "consisting of" and/or "consisting essentially of" are also provided.

[0830] Units, prefixes, and symbols are denoted in their Système International de Unites (SI) accepted form. Numeric ranges are inclusive of the numbers defining the range. Where a range of values is recited, it is to be understood that each intervening integer value, and each fraction thereof, between the recited upper and lower limits of that range is also specifically disclosed, along with each subrange between such values. The upper and lower limits of any range can independently be included in or excluded from the range, and each range where either, neither or both limits are included is also encompassed within the invention. Where a value is explicitly recited, it is to be understood that values which are about the same quantity or amount as the recited value are also within the scope of the invention. Where a combination is disclosed, each subcombination of the elements of that combination is also specifically disclosed and is within the scope of the invention. Conversely, where different elements or groups of elements are individually disclosed, combinations thereof are also disclosed. Where any element of an invention is disclosed as having a plurality of alternatives, examples of that invention in which each alternative is excluded singly or in any combination with the other alternatives are also hereby

disclosed; more than one element of an invention can have such exclusions, and all combinations of elements having such exclusions are hereby disclosed.

[0831] Nucleotides are referred to by their commonly accepted single-letter codes. Unless otherwise indicated, nucleic acids are written left to right in 5′ to 3′ orientation. Nucleobases are referred to herein by their commonly known one-letter symbols recommended by the IUPAC-IUB Biochemical Nomenclature Commission. Accordingly, A represents adenine, C represents cytosine, G represents guanine, T represents thymine, U represents uracil.

[0832] Amino acids are referred to herein by either their commonly known three letter symbols or by the one-letter symbols recommended by the IUPAC-IUB Biochemical Nomenclature Commission. Unless otherwise indicated, amino acid sequences are written left to right in amino to carboxy orientation.

[0833] *About:* The term "about" as used in connection with a numerical value throughout the specification and the claims denotes an interval of accuracy, familiar and acceptable to a person skilled in the art, such interval of accuracy is ± 10 %.

[0834] Where ranges are given, endpoints are included. Furthermore, unless otherwise indicated or otherwise evident from the context and understanding of one of ordinary skill in the art, values that are expressed as ranges can assume any specific value or subrange within the stated ranges in different embodiments of the invention, to the tenth of the unit of the lower limit of the range, unless the context clearly dictates otherwise.

[0835] *Administered in combination:* As used herein, the term "administered in combination" or "combined administration" means that two or more agents are administered to a subject at the same time or within an interval such that there can be an overlap of an effect of each agent on the patient. In some embodiments, they are administered within about 60, 30, 15, 10, 5, or 1 minute of one another. In some embodiments, the administrations of the agents are spaced sufficiently closely together such that a combinatorial (*e.g.*, a synergistic) effect is achieved.

[0836] *Amino acid substitution:* The term "amino acid substitution" refers to replacing an amino acid residue present in a parent or reference sequence (*e.g.*, a wild type G6PC sequence) with another amino acid residue. An amino acid can be substituted in a parent or reference sequence (e.g., a wild type G6PC polypeptide sequence), for example, via chemical peptide synthesis or through recombinant methods known in the art. Accordingly, a reference to a "substitution at position X"

refers to the substitution of an amino acid present at position X with an alternative amino acid residue. In some aspects, substitution patterns can be described according to the schema AnY, wherein A is the single letter code corresponding to the amino acid naturally or originally present at position n, and Y is the substituting amino acid residue. In other aspects, substitution patterns can be described according to the schema An(YZ), wherein A is the single letter code corresponding to the amino acid residue substituting the amino acid naturally or originally present at position X, and Y and Z are alternative substituting amino acid residue.

[0837] In the context of the present disclosure, substitutions (even when they referred to as amino acid substitution) are conducted at the nucleic acid level, *i.e.,* substituting an amino acid residue with an alternative amino acid residue is conducted by substituting the codon encoding the first amino acid with a codon encoding the second amino acid.

[0838] *Animal:* As used herein, the term "animal" refers to any member of the animal kingdom. In some embodiments, "animal" refers to humans at any stage of development. In some embodiments, "animal" refers to non-human animals at any stage of development. In certain embodiments, the non-human animal is a mammal (*e.g.*, a rodent, a mouse, a rat, a rabbit, a monkey, a dog, a cat, a sheep, cattle, a primate, or a pig). In some embodiments, animals include, but are not limited to, mammals, birds, reptiles, amphibians, fish, and worms. In some embodiments, the animal is a transgenic animal, genetically-engineered animal, or a clone.

[0839] *Approximately:* As used herein, the term "approximately," as applied to one or more values of interest, refers to a value that is similar to a stated reference value. In certain embodiments, the term "approximately" refers to a range of values that fall within 25%, 20%, 19%, 18%, 17%, 16%, 15%, 14%, 13%, 12%, 11%, 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, 1%, or less in either direction (greater than or less than) of the stated reference value unless otherwise stated or otherwise evident from the context (except where such number would exceed 100% of a possible value).

[0840] *Associated with:* As used herein with respect to a disease, the term "associated with" means that the symptom, measurement, characteristic, or status in question is linked to the diagnosis, development, presence, or progression of that disease. As association can, but need not, be causatively linked to the disease. For example, symptoms, sequelae, or any effects causing a decrease in the quality of life of a patient of GSD-IA are considered associated with GSD-Ia and in some embodiments

of the present invention can be treated, ameliorated, or prevented by administering the polynucleotides of the present invention to a subject in need thereof.

[0841] When used with respect to two or more moieties, the terms "associated with," "conjugated," "linked," "attached," and "tethered," when used with respect to two or more moieties, means that the moieties are physically associated or connected with one another, either directly or via one or more additional moieties that serves as a linking agent, to form a structure that is sufficiently stable so that the moieties remain physically associated under the conditions in which the structure is used, *e.g.*, physiological conditions. An "association" need not be strictly through direct covalent chemical bonding. It can also suggest ionic or hydrogen bonding or a hybridization based connectivity sufficiently stable such that the "associated" entities remain physically associated.

[0842] *Bifunctional:* As used herein, the term "bifunctional" refers to any substance, molecule or moiety that is capable of or maintains at least two functions. The functions can affect the same outcome or a different outcome. The structure that produces the function can be the same or different. For example, bifunctional modified RNAs of the present invention can encode a G6PC peptide (a first function) while those nucleosides that comprise the encoding RNA are, in and of themselves, capable of extending the half-life of the RNA (second function). In this example, delivery of the bifunctional modified RNA to a subject suffering from a protein deficiency would produce not only a peptide or protein molecule that can ameliorate or treat a disease or conditions, but would also maintain a population modified RNA present in the subject for a prolonged period of time. In other aspects, a bifunction modified mRNA can be a chimeric or quimeric molecule comprising, for example, an RNA encoding a G6PC peptide (a first function) and a second protein either fused to first protein or co-expressed with the first protein.

[0843] *Biocompatible*: As used herein, the term "biocompatible" means compatible with living cells, tissues, organs or systems posing little to no risk of injury, toxicity or rejection by the immune system.

[0844] *Biodegradable*: As used herein, the term "biodegradable" means capable of being broken down into innocuous products by the action of living things.

[0845] *Biologically active*: As used herein, the phrase "biologically active" refers to a characteristic of any substance that has activity in a biological system and/or organism. For instance, a substance that, when administered to an organism, has a

biological effect on that organism, is considered to be biologically active. In particular embodiments, a polynucleotide of the present invention can be considered biologically active if even a portion of the polynucleotide is biologically active or mimics an activity considered biologically relevant.

[0846] *Chimera*: As used herein, "chimera" is an entity having two or more incongruous or heterogeneous parts or regions. For example, a chimeric molecule can comprise a first part comprising a G6PC polypeptide, and a second part (e.g., genetically fused to the first part) comprising a second therapeutic protein (e.g., a protein with a distinct enzymatic activity, an antigen binding moiety, or a moiety capable of extending the plasma half-life of G6PC, for example, an Fc region of an antibody).

[0847] *Sequence Optimization*: The term "sequence optimization" refers to a process or series of processes by which nucleobases in a reference nucleic acid sequence are replaced with alternative nucleobases, resulting in a nucleic acid sequence with improved properties, e.g., improved protein expression or decreased immunogenicity.

[0848] In general, the goal in sequence optimization is to produce a synonymous nucleotide sequence than encodes the same polypeptide sequence encoded by the reference nucleotide sequence. Thus, there are no amino acid substitutions (as a result of codon optimization) in the polypeptide encoded by the codon optimized nucleotide sequence with respect to the polypeptide encoded by the reference nucleotide sequence.

[0849] *Codon substitution*: The terms "codon substitution" or "codon replacement" in the context of sequence optimization refer to replacing a codon present in a reference nucleic acid sequence with another codon. A codon can be substituted in a reference nucleic acid sequence, for example, via chemical peptide synthesis or through recombinant methods known in the art. Accordingly, references to a "substitution" or "replacement" at a certain location in a nucleic acid sequence (e.g., an mRNA) or within a certain region or subsequence of a nucleic acid sequence (e.g., an mRNA) refer to the substitution of a codon at such location or region with an alternative codon.

[0850] As used herein, the terms "coding region" and "region encoding" and grammatical variants thereof, refer to an Open Reading Frame (ORF) in a polynucleotide that upon expression yields a polypeptide or protein.

[0851] *Compound:* As used herein, the term "compound," is meant to include all stereoisomers and isotopes of the structure depicted. As used herein, the term "stereoisomer" means any geometric isomer (e.g., cis- and trans- isomer), enantiomer, or diastereomer of a compound. The present disclosure encompasses any and all stereoisomers of the compounds described herein, including stereomerically pure forms (e.g., geometrically pure, enantiomerically pure, or diastereomerically pure) and enantiomeric and stereoisomeric mixtures, e.g., racemates. Enantiomeric and stereomeric mixtures of compounds and means of resolving them into their component enantiomers or stereoisomers are well-known. "Isotopes" refers to atoms having the same atomic number but different mass numbers resulting from a different number of neutrons in the nuclei. For example, isotopes of hydrogen include tritium and deuterium. Further, a compound, salt, or complex of the present disclosure can be prepared in combination with solvent or water molecules to form solvates and hydrates by routine methods.

[0852] *Contacting*: As used herein, the term "contacting" means establishing a physical connection between two or more entities. For example, contacting a mammalian cell with a nanoparticle composition means that the mammalian cell and a nanoparticle are made to share a physical connection. Methods of contacting cells with external entities both in vivo and ex vivo are well known in the biological arts. For example, contacting a nanoparticle composition and a mammalian cell disposed within a mammal can be performed by varied routes of administration (*e.g.*, intravenous, intramuscular, intradermal, and subcutaneous) and can involve varied amounts of nanoparticle compositions. Moreover, more than one mammalian cell can be contacted by a nanoparticle composition.

[0853] *Conservative amino acid substitution:* A "conservative amino acid substitution" is one in which the amino acid residue in a protein sequence is replaced with an amino acid residue having a similar side chain. Families of amino acid residues having similar side chains have been defined in the art, including basic side chains (*e.g.*, lysine, arginine, or histidine), acidic side chains (*e.g.*, aspartic acid or glutamic acid), uncharged polar side chains (*e.g.*, glycine, asparagine, glutamine, serine, threonine, tyrosine, or cysteine), nonpolar side chains (*e.g.*, alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, or tryptophan), beta-branched side chains (*e.g.*, threonine, valine, isoleucine) and aromatic side chains (*e.g.*, tyrosine, phenylalanine, tryptophan, or histidine). Thus, if an amino acid in a

polypeptide is replaced with another amino acid from the same side chain family, the amino acid substitution is considered to be conservative. In another aspect, a string of amino acids can be conservatively replaced with a structurally similar string that differs in order and/or composition of side chain family members.

[0854] *Non-conservative amino acid substitution*: Non-conservative amino acid substitutions include those in which (i) a residue having an electropositive side chain (*e.g.,* Arg, His or Lys) is substituted for, or by, an electronegative residue (*e.g.,* Glu or Asp), (ii) a hydrophilic residue (*e.g.,* Ser or Thr) is substituted for, or by, a hydrophobic residue (*e.g.,* Ala, Leu, Ile, Phe or Val), (iii) a cysteine or proline is substituted for, or by, any other residue, or (iv) a residue having a bulky hydrophobic or aromatic side chain (*e.g.,* Val, His, Ile or Trp) is substituted for, or by, one having a smaller side chain (*e.g.,* Ala or Ser) or no side chain (*e.g.,* Gly).

[0855] Other amino acid substitutions can be readily identified by workers of ordinary skill. For example, for the amino acid alanine, a substitution can be taken from any one of D-alanine, glycine, beta-alanine, L-cysteine and D-cysteine. For lysine, a replacement can be any one of D-lysine, arginine, D-arginine, homo-arginine, methionine, D-methionine, ornithine, or D- ornithine. Generally, substitutions in functionally important regions that can be expected to induce changes in the properties of isolated polypeptides are those in which (i) a polar residue, *e.g.,* serine or threonine, is substituted for (or by) a hydrophobic residue, *e.g.,* leucine, isoleucine, phenylalanine, or alanine; (ii) a cysteine residue is substituted for (or by) any other residue; (iii) a residue having an electropositive side chain, *e.g.,* lysine, arginine or histidine, is substituted for (or by) a residue having an electronegative side chain, *e.g.,* glutamic acid or aspartic acid; or (iv) a residue having a bulky side chain, *e.g.,* phenylalanine, is substituted for (or by) one not having such a side chain, *e.g.,* glycine. The likelihood that one of the foregoing non-conservative substitutions can alter functional properties of the protein is also correlated to the position of the substitution with respect to functionally important regions of the protein: some non-conservative substitutions can accordingly have little or no effect on biological properties.

[0856] *Conserved*: As used herein, the term "conserved" refers to nucleotides or amino acid residues of a polynucleotide sequence or polypeptide sequence, respectively, that are those that occur unaltered in the same position of two or more sequences being compared. Nucleotides or amino acids that are relatively conserved

are those that are conserved amongst more related sequences than nucleotides or amino acids appearing elsewhere in the sequences.

[0857] In some embodiments, two or more sequences are said to be "completely conserved" if they are 100% identical to one another. In some embodiments, two or more sequences are said to be "highly conserved" if they are at least 70% identical, at least 80% identical, at least 90% identical, or at least 95% identical to one another. In some embodiments, two or more sequences are said to be "highly conserved" if they are about 70% identical, about 80% identical, about 90% identical, about 95%, about 98%, or about 99% identical to one another. In some embodiments, two or more sequences are said to be "conserved" if they are at least 30% identical, at least 40% identical, at least 50% identical, at least 60% identical, at least 70% identical, at least 80% identical, at least 90% identical, or at least 95% identical to one another. In some embodiments, two or more sequences are said to be "conserved" if they are about 30% identical, about 40% identical, about 50% identical, about 60% identical, about 70% identical, about 80% identical, about 90% identical, about 95% identical, about 98% identical, or about 99% identical to one another. Conservation of sequence can apply to the entire length of a polynucleotide or polypeptide or can apply to a portion, region or feature thereof.

[0858] *Controlled Release:* As used herein, the term "controlled release" refers to a pharmaceutical composition or compound release profile that conforms to a particular pattern of release to effect a therapeutic outcome.

[0859] *Cyclic or Cyclized:* As used herein, the term "cyclic" refers to the presence of a continuous loop. Cyclic molecules need not be circular, only joined to form an unbroken chain of subunits. Cyclic molecules such as the engineered RNA or mRNA of the present invention can be single units or multimers or comprise one or more components of a complex or higher order structure.

[0860] *Cytotoxic*: As used herein, "cytotoxic" refers to killing or causing injurious, toxic, or deadly effect on a cell (*e.g.*, a mammalian cell (*e.g.*, a human cell)), bacterium, virus, fungus, protozoan, parasite, prion, or a combination thereof.

[0861] *Delivering*: As used herein, the term “delivering” means providing an entity to a destination. For example, delivering a polynucleotide to a subject can involve administering a nanoparticle composition including the polynucleotide to the subject (*e.g.*, by an intravenous, intramuscular, intradermal, or subcutaneous route).

Administration of a nanoparticle composition to a mammal or mammalian cell can involve contacting one or more cells with the nanoparticle composition.

[0862] *Delivery Agent*: As used herein, "delivery agent" refers to any substance that facilitates, at least in part, the *in vivo, in vitro, or ex vivo* delivery of a polynucleotide to targeted cells.

[0863] *Destabilized:* As used herein, the term "destable," "destabilize," or "destabilizing region" means a region or molecule that is less stable than a starting, wild-type or native form of the same region or molecule.

[0864] *Diastereomer:* As used herein, the term "diastereomer," means stereoisomers that are not mirror images of one another and are non-superimposable on one another.

[0865] *Digest*: As used herein, the term "digest" means to break apart into smaller pieces or components. When referring to polypeptides or proteins, digestion results in the production of peptides.

[0866] *Distal:* As used herein, the term "distal" means situated away from the center or away from a point or region of interest.

[0867] *Domain*: As used herein, when referring to polypeptides, the term "domain" refers to a motif of a polypeptide having one or more identifiable structural or functional characteristics or properties (e.g., binding capacity, serving as a site for protein-protein interactions).

[0868] *Dosing regimen*: As used herein, a "dosing regimen" or a "dosing regimen" is a schedule of administration or physician determined regimen of treatment, prophylaxis, or palliative care.

[0869] *Effective Amount:* As used herein, the term "effective amount" of an agent is that amount sufficient to effect beneficial or desired results, for example, clinical results, and, as such, an "effective amount" depends upon the context in which it is being applied. For example, in the context of administering an agent that treats a protein deficiency (e.g., a G6PC deficiency), an effective amount of an agent is, for example, an amount of mRNA expressing sufficient G6PC to ameliorate, reduce, eliminate, or prevent the symptoms associated with the G6PC deficiency, as compared to the severity of the symptom observed without administration of the agent. The term "effective amount" can be used interchangeably with "effective dose," "therapeutically effective amount," or "therapeutically effective dose."

[0870] *Enantiomer:* As used herein, the term "enantiomer" means each individual optically active form of a compound of the invention, having an optical purity or

enantiomeric excess (as determined by methods standard in the art) of at least 80% (*i.e.*, at least 90% of one enantiomer and at most 10% of the other enantiomer), at least 90%, or at least 98%.

[0871] *Encapsulate:* As used herein, the term "encapsulate" means to enclose, surround or encase.

[0872] *Encapsulation Efficiency*: As used herein, "encapsulation efficiency" refers to the amount of a polynucleotide that becomes part of a nanoparticle composition, relative to the initial total amount of polynucleotide used in the preparation of a nanoparticle composition. For example, if 97 mg of polynucleotide are encapsulated in a nanoparticle composition out of a total 100 mg of polynucleotide initially provided to the composition, the encapsulation efficiency can be given as 97%. As used herein, "encapsulation" can refer to complete, substantial, or partial enclosure, confinement, surrounding, or encasement.

[0873] *Encoded protein cleavage signal*: As used herein, "encoded protein cleavage signal" refers to the nucleotide sequence that encodes a protein cleavage signal.

[0874] *Engineered:* As used herein, embodiments of the invention are "engineered" when they are designed to have a feature or property, whether structural or chemical, that varies from a starting point, wild type or native molecule.

[0875] *Enhanced Delivery*: As used herein, the term "enhanced delivery" means delivery of more (e.g., at least 1.5 fold more, at least 2-fold more, at least 3-fold more, at least 4-fold more, at least 5-fold more, at least 6-fold more, at least 7-fold more, at least 8-fold more, at least 9-fold more, at least 10-fold more) of a polynucleotide by a nanoparticle to a target tissue of interest (e.g., mammalian liver) compared to the level of delivery of a polynucleotide by a control nanoparticle to a target tissue of interest (e.g., MC3, KC2, or DLinDMA). The level of delivery of a nanoparticle to a particular tissue can be measured by comparing the amount of protein produced in a tissue to the weight of said tissue, comparing the amount of polynucleotide in a tissue to the weight of said tissue, comparing the amount of protein produced in a tissue to the amount of total protein in said tissue, or comparing the amount of polynucleotide in a tissue to the amount of total polynucleotide in said tissue. It will be understood that the enhanced delivery of a nanoparticle to a target tissue need not be determined in a subject being treated, it can be determined in a surrogate such as an animal model (e.g., a rat model).

[0876] *Exosome:* As used herein, "exosome" is a vesicle secreted by mammalian cells or a complex involved in RNA degradation.

[0877] *Expression*: As used herein, "expression" of a nucleic acid sequence refers to one or more of the following events: (1) production of an mRNA template from a DNA sequence (*e.g.*, by transcription); (2) processing of an mRNA transcript (*e.g.*, by splicing, editing, 5′ cap formation, and/or 3′ end processing); (3) translation of an mRNA into a polypeptide or protein; and (4) post-translational modification of a polypeptide or protein.

[0878] *Ex Vivo*: As used herein, the term "ex vivo" refers to events that occur outside of an organism (*e.g.*, animal, plant, or microbe or cell or tissue thereof). Ex vivo events can take place in an environment minimally altered from a natural (*e.g.*, in vivo) environment.

[0879] *Feature:* As used herein, a "feature" refers to a characteristic, a property, or a distinctive element. When referring to polypeptides, "features" are defined as distinct amino acid sequence-based components of a molecule. Features of the polypeptides encoded by the polynucleotides of the present invention include surface manifestations, local conformational shape, folds, loops, half-loops, domains, half-domains, sites, termini or any combination thereof.

[0880] *Formulation*: As used herein, a "formulation" includes at least a polynucleotide and one or more of a carrier, an excipient, and a delivery agent.

[0881] *Fragment:* A "fragment," as used herein, refers to a portion. For example, fragments of proteins can comprise polypeptides obtained by digesting full-length protein isolated from cultured cells. In some embodiments, a fragment is a subsequence of a full-length protein (e.g., G6PC) wherein N-terminal, and/or C-terminal, and/or internal subsequences have been deleted. In some preferred aspects of the present invention, the fragments of a protein of the present invention are functional fragments.

[0882] *Functional*: As used herein, a "functional" biological molecule is a biological molecule in a form in which it exhibits a property and/or activity by which it is characterized. Thus, a functional fragment of a polynucleotide of the present invention is a polynucleotide capable of expressing a functional G6PC fragment. As used herein, a functional fragment of G6PC refers to a fragment of wild type G6PC (i.e., a fragment of any of its naturally occurring isoforms), or a mutant or variant thereof, wherein the fragment retains a least about 10%, at least about 15%, at least

about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, or at least about 95% of the biological activity of the corresponding full length protein.

[0883] *G6PC Associated Disease:* As use herein the terms " G6PC -associated disease" or " G6PC -associated disorder" refer to diseases or disorders, respectively, which result from aberrant G6PC activity (e.g., decreased activity or increased activity). As a non-limiting example, GSD-IA is a G6PC associated disease.

[0884] The terms " G6PC enzymatic activity" and " G6PC activity," are used interchangeably in the present disclosure and refer to G6PC 's ability to convert phenylalanine to tyrosine. Accordingly, a fragment or variant retaining or having G6PC enzymatic activity or G6PC activity refers to a fragment or variant that has measurable enzymatic activity in catalyzing the conversion of phenylalanine to tyrosine.

[0885] *Helper Lipid*: As used herein, the term "helper lipid" refers to a compound or molecule that includes a lipidic moiety (for insertion into a lipid layer, e.g., lipid bilayer) and a polar moiety (for interaction with physiologic solution at the surface of the lipid layer). Typically, the helper lipid is a phospholipid. A function of the helper lipid is to "complement" the amino lipid and increase the fusogenicity of the bilayer and/or to help facilitate endosomal escape, e.g., of nucleic acid delivered to cells. Helper lipids are also believed to be a key structural component to the surface of the LNP.

[0886] *Homology*: As used herein, the term "homology" refers to the overall relatedness between polymeric molecules, *e.g.* between nucleic acid molecules (*e.g.* DNA molecules and/or RNA molecules) and/or between polypeptide molecules. Generally, the term "homology" implies an evolutionary relationship between two molecules. Thus, two molecules that are homologous will have a common evolutionary ancestor. In the context of the present invention, the term homology encompasses both to identity and similarity.

[0887] In some embodiments, polymeric molecules are considered to be "homologous" to one another if at least 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or 99% of the monomers in the molecule are identical (exactly the same monomer) or are similar (conservative substitutions). The

term "homologous" necessarily refers to a comparison between at least two sequences (polynucleotide or polypeptide sequences).

[0888] *Identity*: As used herein, the term "identity" refers to the overall monomer conservation between polymeric molecules, *e.g.*, between polynucleotide molecules (*e.g.* DNA molecules and/or RNA molecules) and/or between polypeptide molecules. Calculation of the percent identity of two polynucleotide sequences, for example, can be performed by aligning the two sequences for optimal comparison purposes (*e.g.*, gaps can be introduced in one or both of a first and a second nucleic acid sequences for optimal alignment and non-identical sequences can be disregarded for comparison purposes). In certain embodiments, the length of a sequence aligned for comparison purposes is at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, at least 95%, or 100% of the length of the reference sequence. The nucleotides at corresponding nucleotide positions are then compared. When a position in the first sequence is occupied by the same nucleotide as the corresponding position in the second sequence, then the molecules are identical at that position. The percent identity between the two sequences is a function of the number of identical positions shared by the sequences, taking into account the number of gaps, and the length of each gap, which needs to be introduced for optimal alignment of the two sequences. The comparison of sequences and determination of percent identity between two sequences can be accomplished using a mathematical algorithm. When comparing DNA and RNA, thymine (T) and uracil (U) can be considered equivalent.

[0889] Suitable software programs are available from various sources, and for alignment of both protein and nucleotide sequences. One suitable program to determine percent sequence identity is bl2seq, part of the BLAST suite of program available from the U.S. government's National Center for Biotechnology Information BLAST web site (blast.ncbi.nlm.nih.gov). Bl2seq performs a comparison between two sequences using either the BLASTN or BLASTP algorithm. BLASTN is used to compare nucleic acid sequences, while BLASTP is used to compare amino acid sequences. Other suitable programs are, e.g., Needle, Stretcher, Water, or Matcher, part of the EMBOSS suite of bioinformatics programs and also available from the European Bioinformatics Institute (EBI) at www.ebi.ac.uk/Tools/psa.

[0890] Sequence alignments can be conducted using methods known in the art such as MAFFT, Clustal (ClustalW, Clustal X or Clustal Omega), MUSCLE, etc.

[0891] Different regions within a single polynucleotide or polypeptide target sequence that aligns with a polynucleotide or polypeptide reference sequence can each have their own percent sequence identity. It is noted that the percent sequence identity value is rounded to the nearest tenth. For example, 80.11, 80.12, 80.13, and 80.14 are rounded down to 80.1, while 80.15, 80.16, 80.17, 80.18, and 80.19 are rounded up to 80.2. It also is noted that the length value will always be an integer.

[0892] In certain aspects, the percentage identity "%ID" of a first amino acid sequence (or nucleic acid sequence) to a second amino acid sequence (or nucleic acid sequence) is calculated as %ID = 100 x (Y/Z), where Y is the number of amino acid residues (or nucleobases) scored as identical matches in the alignment of the first and second sequences (as aligned by visual inspection or a particular sequence alignment program) and Z is the total number of residues in the second sequence. If the length of a first sequence is longer than the second sequence, the percent identity of the first sequence to the second sequence will be higher than the percent identity of the second sequence to the first sequence.

[0893] One skilled in the art will appreciate that the generation of a sequence alignment for the calculation of a percent sequence identity is not limited to binary sequence-sequence comparisons exclusively driven by primary sequence data. It will also be appreciated that sequence alignments can be generated by integrating sequence data with data from heterogeneous sources such as structural data (e.g., crystallographic protein structures), functional data (e.g., location of mutations), or phylogenetic data. A suitable program that integrates heterogeneous data to generate a multiple sequence alignment is T-Coffee, available at www.tcoffee.org, and alternatively available, e.g., from the EBI. It will also be appreciated that the final alignment used to calculate percent sequence identity can be curated either automatically or manually.

[0894] *Immune response:* The term “immune response” refers to the action of, for example, lymphocytes, antigen presenting cells, phagocytic cells, granulocytes, and soluble macromolecules produced by the above cells or the liver (including antibodies, cytokines, and complement) that results in selective damage to, destruction of, or elimination from the human body of invading pathogens, cells or tissues infected with pathogens, cancerous cells, or, in cases of autoimmunity or pathological inflammation, normal human cells or tissues. In some cases, the administration of a nanoparticle comprising a lipid component and an encapsulated

therapeutic agent can trigger an immune response, which can be caused by (i) the encapsulated therapeutic agent (e.g., an mRNA), (ii) the expression product of such encapsulated therapeutic agent (e.g., a polypeptide encoded by the mRNA), (iii) the lipid component of the nanoparticle, or (iv) a combination thereof.

[0895] *Inflammatory response*: "Inflammatory response" refers to immune responses involving specific and non-specific defense systems. A specific defense system reaction is a specific immune system reaction to an antigen. Examples of specific defense system reactions include antibody responses. A non-specific defense system reaction is an inflammatory response mediated by leukocytes generally incapable of immunological memory, e.g., macrophages, eosinophils and neutrophils. In some aspects, an immune response includes the secretion of inflammatory cytokines, resulting in elevated inflammatory cytokine levels.

[0896] *Inflammatory cytokines*: The term "inflammatory cytokine" refers to cytokines that are elevated in an inflammatory response. Examples of inflammatory cytokines include interleukin-6 (IL-6), CXCL1 (chemokine (C-X-C motif) ligand 1; also known as GROα, interferon-γ (IFNγ), tumor necrosis factor α (TNFα), interferon γ-induced protein 10 (IP-10), or granulocyte-colony stimulating factor (G-CSF). The term inflammatory cytokines includes also other cytokines associated with inflammatory responses known in the art, e.g., interleukin-1 (IL-1), interleukin-8 (IL-8), interleukin-12 (IL-12), interleukin-13 (Il-13), interferon α (IFN-α), etc.

[0897] *In Vitro*: As used herein, the term "in vitro" refers to events that occur in an artificial environment, e.g., in a test tube or reaction vessel, in cell culture, in a Petri dish, etc., rather than within an organism (e.g., animal, plant, or microbe).

[0898] *In Vivo*: As used herein, the term "in vivo" refers to events that occur within an organism (e.g., animal, plant, or microbe or cell or tissue thereof).

[0899] *Insertional and deletional variants:* "Insertional variants" when referring to polypeptides are those with one or more amino acids inserted immediately adjacent to an amino acid at a particular position in a native or starting sequence. "Immediately adjacent" to an amino acid means connected to either the alpha-carboxy or alpha-amino functional group of the amino acid. "Deletional variants" when referring to polypeptides are those with one or more amino acids in the native or starting amino acid sequence removed. Ordinarily, deletional variants will have one or more amino acids deleted in a particular region of the molecule.

[0900] *Intact:* As used herein, in the context of a polypeptide, the term "intact" means retaining an amino acid corresponding to the wild type protein, *e.g.,* not mutating or substituting the wild type amino acid. Conversely, in the context of a nucleic acid, the term "intact" means retaining a nucleobase corresponding to the wild type nucleic acid, e.g., not mutating or substituting the wild type nucleobase.

[0901] *Ionizable amino lipid*: The term "ionizable amino lipid" includes those lipids having one, two, three, or more fatty acid or fatty alkyl chains and a pH-titratable amino head group (e.g., an alkylamino or dialkylamino head group). An ionizable amino lipid is typically protonated (i.e., positively charged) at a pH below the pKa of the amino head group and is substantially not charged at a pH above the pKa. Such ionizable amino lipids include, but are not limited to DLin-MC3-DMA (MC3) and (13Z,165Z)-N,N-dimethyl-3-nonydocosa-13-16-dien-1-amine (L608).

[0902] *Isolated*: As used herein, the term "isolated" refers to a substance or entity that has been separated from at least some of the components with which it was associated (whether in nature or in an experimental setting). Isolated substances (*e.g.*, polynucleotides or polypeptides) can have varying levels of purity in reference to the substances from which they have been isolated. Isolated substances and/or entities can be separated from at least about 10%, about 20%, about 30%, about 40%, about 50%, about 60%, about 70%, about 80%, about 90%, or more of the other components with which they were initially associated. In some embodiments, isolated substances are more than about 80%, about 85%, about 90%, about 91%, about 92%, about 93%, about 94%, about 95%, about 96%, about 97%, about 98%, about 99%, or more than about 99% pure. As used herein, a substance is "pure" if it is substantially free of other components.

[0903] *Substantially isolated*: By "substantially isolated" is meant that the compound is substantially separated from the environment in which it was formed or detected. Partial separation can include, for example, a composition enriched in the compound of the present disclosure. Substantial separation can include compositions containing at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 97%, or at least about 99% by weight of the compound of the present disclosure, or salt thereof.

[0904] A polynucleotide, vector, polypeptide, cell, or any composition disclosed herein which is "isolated" is a polynucleotide, vector, polypeptide, cell, or composition which is in a form not found in nature. Isolated polynucleotides, vectors,

polypeptides, or compositions include those which have been purified to a degree that they are no longer in a form in which they are found in nature. In some aspects, a polynucleotide, vector, polypeptide, or composition which is isolated is substantially pure.

[0905] *Isomer:* As used herein, the term "isomer" means any tautomer, stereoisomer, enantiomer, or diastereomer of any compound of the invention. It is recognized that the compounds of the invention can have one or more chiral centers and/or double bonds and, therefore, exist as stereoisomers, such as double-bond isomers (*i.e.*, geometric E/Z isomers) or diastereomers (*e.g.*, enantiomers (*i.e.*, (+) or (-)) or cis/trans isomers). According to the invention, the chemical structures depicted herein, and therefore the compounds of the invention, encompass all of the corresponding stereoisomers, that is, both the stereomerically pure form (*e.g.*, geometrically pure, enantiomerically pure, or diastereomerically pure) and enantiomeric and stereoisomeric mixtures, *e.g.*, racemates. Enantiomeric and stereoisomeric mixtures of compounds of the invention can typically be resolved into their component enantiomers or stereoisomers by well-known methods, such as chiral-phase gas chromatography, chiral-phase high performance liquid chromatography, crystallizing the compound as a chiral salt complex, or crystallizing the compound in a chiral solvent. Enantiomers and stereoisomers can also be obtained from stereomerically or enantiomerically pure intermediates, reagents, and catalysts by well-known asymmetric synthetic methods.

[0906] *Linker*: As used herein, a "linker" refers to a group of atoms, e.g., 10-1,000 atoms, and can be comprised of the atoms or groups such as, but not limited to, carbon, amino, alkylamino, oxygen, sulfur, sulfoxide, sulfonyl, carbonyl, and imine. The linker can be attached to a modified nucleoside or nucleotide on the nucleobase or sugar moiety at a first end, and to a payload, e.g., a detectable or therapeutic agent, at a second end. The linker can be of sufficient length as to not interfere with incorporation into a nucleic acid sequence. The linker can be used for any useful purpose, such as to form polynucleotide multimers (e.g., through linkage of two or more chimeric polynucleotides molecules or IVT polynucleotides) or polynucleotides conjugates, as well as to administer a payload, as described herein. Examples of chemical groups that can be incorporated into the linker include, but are not limited to, alkyl, alkenyl, alkynyl, amido, amino, ether, thioether, ester, alkylene, heteroalkylene, aryl, or heterocyclyl, each of which can be optionally substituted, as

described herein. Examples of linkers include, but are not limited to, unsaturated alkanes, polyethylene glycols (e.g., ethylene or propylene glycol monomeric units, e.g., diethylene glycol, dipropylene glycol, triethylene glycol, tripropylene glycol, tetraethylene glycol, or tetraethylene glycol), and dextran polymers and derivatives thereof., Other examples include, but are not limited to, cleavable moieties within the linker, such as, for example, a disulfide bond (-S-S-) or an azo bond (-N=N-), which can be cleaved using a reducing agent or photolysis. Non-limiting examples of a selectively cleavable bond include an amido bond can be cleaved for example by the use of tris(2-carboxyethyl)phosphine (TCEP), or other reducing agents, and/or photolysis, as well as an ester bond can be cleaved for example by acidic or basic hydrolysis.

[0907] *Methods of Administration*: As used herein, "methods of administration" can include intravenous, intramuscular, intradermal, subcutaneous, or other methods of delivering a composition to a subject. A method of administration can be selected to target delivery (*e.g.*, to specifically deliver) to a specific region or system of a body.

[0908] *Modified:* As used herein "modified" refers to a changed state or structure of a molecule of the invention. Molecules can be modified in many ways including chemically, structurally, and functionally. In some embodiments, the mRNA molecules of the present invention are modified by the introduction of non-natural nucleosides and/or nucleotides, *e.g.,* as it relates to the natural ribonucleotides A, U, G, and C. Noncanonical nucleotides such as the cap structures are not considered "modified" although they differ from the chemical structure of the A, C, G, U ribonucleotides.

[0909] *Mucus*: As used herein, "mucus" refers to the natural substance that is viscous and comprises mucin glycoproteins.

[0910] *Nanoparticle Composition*: As used herein, a "nanoparticle composition" is a composition comprising one or more lipids. Nanoparticle compositions are typically sized on the order of micrometers or smaller and can include a lipid bilayer. Nanoparticle compositions encompass lipid nanoparticles (LNPs), liposomes (*e.g.*, lipid vesicles), and lipoplexes. For example, a nanoparticle composition can be a liposome having a lipid bilayer with a diameter of 500 nm or less.

[0911] *Naturally occurring:* As used herein, "naturally occurring" means existing in nature without artificial aid.

[0912] *Non-human vertebrate:* As used herein, a "non-human vertebrate" includes all vertebrates except *Homo sapiens*, including wild and domesticated species. Examples of non-human vertebrates include, but are not limited to, mammals, such as alpaca, banteng, bison, camel, cat, cattle, deer, dog, donkey, gayal, goat, guinea pig, horse, llama, mule, pig, rabbit, reindeer, sheep water buffalo, and yak.

[0913] *Nucleic acid sequence:* The terms "nucleic acid sequence," "nucleotide sequence," or "polynucleotide sequence" are used interchangeably and refer to a contiguous nucleic acid sequence. The sequence can be either single stranded or double stranded DNA or RNA, *e.g.,* an mRNA.

[0914] The term "nucleic acid," in its broadest sense, includes any compound and/or substance that comprises a polymer of nucleotides. These polymers are often referred to as polynucleotides. Exemplary nucleic acids or polynucleotides of the invention include, but are not limited to, ribonucleic acids (RNAs), deoxyribonucleic acids (DNAs), threose nucleic acids (TNAs), glycol nucleic acids (GNAs), peptide nucleic acids (PNAs), locked nucleic acids (LNAs, including LNA having a β- D-ribo configuration, α-LNA having an α-L-ribo configuration (a diastereomer of LNA), 2′-amino-LNA having a 2′-amino functionalization, and 2′-amino- α-LNA having a 2′-amino functionalization), ethylene nucleic acids (ENA), cyclohexenyl nucleic acids (CeNA) or hybrids or combinations thereof.

[0915] The phrase "nucleotide sequence encoding" refers to the nucleic acid (*e.g.,* an mRNA or DNA molecule) coding sequence which encodes a polypeptide. The coding sequence can further include initiation and termination signals operably linked to regulatory elements including a promoter and polyadenylation signal capable of directing expression in the cells of an individual or mammal to which the nucleic acid is administered. The coding sequence can further include sequences that encode signal peptides.

[0916] *Off-target:* As used herein, "off target" refers to any unintended effect on any one or more target, gene, or cellular transcript.

[0917] *Open reading frame:* As used herein, "open reading frame" or "ORF" refers to a sequence which does not contain a stop codon in a given reading frame.

[0918] *Operably linked:* As used herein, the phrase "operably linked" refers to a functional connection between two or more molecules, constructs, transcripts, entities, moieties or the like.

[0919] *Optionally substituted:* Herein a phrase of the form "optionally substituted X" (*e.g.*, optionally substituted alkyl) is intended to be equivalent to "X, wherein X is optionally substituted" (*e.g.*, "alkyl, wherein said alkyl is optionally substituted"). It is not intended to mean that the feature "X" (*e.g.*, alkyl) *per se* is optional.

[0920] *Part*: As used herein, a "part" or "region" of a polynucleotide is defined as any portion of the polynucleotide that is less than the entire length of the polynucleotide.

[0921] *Patient:* As used herein, "patient" refers to a subject who can seek or be in need of treatment, requires treatment, is receiving treatment, will receive treatment, or a subject who is under care by a trained professional for a particular disease or condition. In some embodiments, the treatment is needed, required, or received to prevent or decrease the risk of developing acute disease, i.e., it is a prophylactic treatment.

[0922] *Pharmaceutically acceptable*: The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms that are, within the scope of sound medical judgment, suitable for use in contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, or other problem or complication, commensurate with a reasonable benefit/risk ratio.

[0923] *Pharmaceutically acceptable excipients:* The phrase "pharmaceutically acceptable excipient," as used herein, refers any ingredient other than the compounds described herein (for example, a vehicle capable of suspending or dissolving the active compound) and having the properties of being substantially nontoxic and non-inflammatory in a patient. Excipients can include, for example: antiadherents, antioxidants, binders, coatings, compression aids, disintegrants, dyes (colors), emollients, emulsifiers, fillers (diluents), film formers or coatings, flavors, fragrances, glidants (flow enhancers), lubricants, preservatives, printing inks, sorbents, suspensing or dispersing agents, sweeteners, and waters of hydration. Exemplary excipients include, but are not limited to: butylated hydroxytoluene (BHT), calcium carbonate, calcium phosphate (dibasic), calcium stearate, croscarmellose, crosslinked polyvinyl pyrrolidone, citric acid, crospovidone, cysteine, ethylcellulose, gelatin, hydroxypropyl cellulose, hydroxypropyl methylcellulose, lactose, magnesium stearate, maltitol, mannitol, methionine, methylcellulose, methyl paraben, microcrystalline cellulose, polyethylene glycol, polyvinyl pyrrolidone, povidone, pregelatinized starch, propyl paraben, retinyl palmitate, shellac, silicon dioxide,

sodium carboxymethyl cellulose, sodium citrate, sodium starch glycolate, sorbitol, starch (corn), stearic acid, sucrose, talc, titanium dioxide, vitamin A, vitamin E, vitamin C, and xylitol.

[0924] *Pharmaceutically acceptable salts*: The present disclosure also includes pharmaceutically acceptable salts of the compounds described herein. As used herein, "pharmaceutically acceptable salts" refers to derivatives of the disclosed compounds wherein the parent compound is modified by converting an existing acid or base moiety to its salt form (*e.g.*, by reacting the free base group with a suitable organic acid). Examples of pharmaceutically acceptable salts include, but are not limited to, mineral or organic acid salts of basic residues such as amines; alkali or organic salts of acidic residues such as carboxylic acids; and the like. Representative acid addition salts include acetate, acetic acid, adipate, alginate, ascorbate, aspartate, benzenesulfonate, benzene sulfonic acid, benzoate, bisulfate, borate, butyrate, camphorate, camphorsulfonate, citrate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, glucoheptonate, glycerophosphate, hemisulfate, heptonate, hexanoate, hydrobromide, hydrochloride, hydroiodide, 2-hydroxy-ethanesulfonate, lactobionate, lactate, laurate, lauryl sulfate, malate, maleate, malonate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, nitrate, oleate, oxalate, palmitate, pamoate, pectinate, persulfate, 3-phenylpropionate, phosphate, picrate, pivalate, propionate, stearate, succinate, sulfate, tartrate, thiocyanate, toluenesulfonate, undecanoate, valerate salts, and the like. Representative alkali or alkaline earth metal salts include sodium, lithium, potassium, calcium, magnesium, and the like, as well as nontoxic ammonium, quaternary ammonium, and amine cations, including, but not limited to ammonium, tetramethylammonium, tetraethylammonium, methylamine, dimethylamine, trimethylamine, triethylamine, ethylamine, and the like. The pharmaceutically acceptable salts of the present disclosure include the conventional non-toxic salts of the parent compound formed, for example, from non-toxic inorganic or organic acids. The pharmaceutically acceptable salts of the present disclosure can be synthesized from the parent compound that contains a basic or acidic moiety by conventional chemical methods. Generally, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, nonaqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are used. Lists of suitable salts are

found in *Remington's Pharmaceutical Sciences*, 17th ed., Mack Publishing Company, Easton, Pa., 1985, p. 1418, *Pharmaceutical Salts: Properties, Selection, and Use*, P.H. Stahl and C.G. Wermuth (eds.), Wiley-VCH, 2008, and Berge et al., *Journal of Pharmaceutical Science*, 66, 1-19 (1977), each of which is incorporated herein by reference in its entirety.

[0925] *Pharmaceutically acceptable solvate*: The term "pharmaceutically acceptable solvate," as used herein, means a compound of the invention wherein molecules of a suitable solvent are incorporated in the crystal lattice. A suitable solvent is physiologically tolerable at the dosage administered. For example, solvates can be prepared by crystallization, recrystallization, or precipitation from a solution that includes organic solvents, water, or a mixture thereof. Examples of suitable solvents are ethanol, water (for example, mono-, di-, and tri-hydrates), *N*-methylpyrrolidinone (NMP), dimethyl sulfoxide (DMSO), *N*,*N*′-dimethylformamide (DMF), *N*,*N*′-dimethylacetamide (DMAC), 1,3-dimethyl-2-imidazolidinone (DMEU), 1,3-dimethyl-3,4,5,6-tetrahydro-2-(1H)-pyrimidinone (DMPU), acetonitrile (ACN), propylene glycol, ethyl acetate, benzyl alcohol, 2-pyrrolidone, benzyl benzoate, and the like. When water is the solvent, the solvate is referred to as a "hydrate."

[0926] *Pharmacokinetic:* As used herein, "pharmacokinetic" refers to any one or more properties of a molecule or compound as it relates to the determination of the fate of substances administered to a living organism. Pharmacokinetics is divided into several areas including the extent and rate of absorption, distribution, metabolism and excretion. This is commonly referred to as ADME where: (A) Absorption is the process of a substance entering the blood circulation; (D) Distribution is the dispersion or dissemination of substances throughout the fluids and tissues of the body; (M) Metabolism (or Biotransformation) is the irreversible transformation of parent compounds into daughter metabolites; and (E) Excretion (or Elimination) refers to the elimination of the substances from the body. In rare cases, some drugs irreversibly accumulate in body tissue.

[0927] *Physicochemical:* As used herein, "physicochemical" means of or relating to a physical and/or chemical property.

[0928] *Polynucleotide:* The term "polynucleotide" as used herein refers to polymers of nucleotides of any length, including ribonucleotides, deoxyribonucleotides, analogs thereof, or mixtures thereof. This term refers to the primary structure of the molecule. Thus, the term includes triple-, double- and single-stranded deoxyribonucleic acid

("DNA"), as well as triple-, double- and single-stranded ribonucleic acid ("RNA"). It also includes modified, for example by alkylation, and/or by capping, and unmodified forms of the polynucleotide. More particularly, the term "polynucleotide" includes polydeoxyribonucleotides (containing 2-deoxy-D-ribose), polyribonucleotides (containing D-ribose), including tRNA, rRNA, hRNA, siRNA and mRNA, whether spliced or unspliced, any other type of polynucleotide which is an N- or C-glycoside of a purine or pyrimidine base, and other polymers containing normucleotidic backbones, for example, polyamide (*e.g.,* peptide nucleic acids "PNAs") and polymorpholino polymers, and other synthetic sequence-specific nucleic acid polymers providing that the polymers contain nucleobases in a configuration which allows for base pairing and base stacking, such as is found in DNA and RNA. In particular aspects, the polynucleotide comprises an mRNA. In other aspect, the mRNA is a synthetic mRNA. In some aspects, the synthetic mRNA comprises at least one unnatural nucleobase. In some aspects, all nucleobases of a certain class have been replaced with unnatural nucleobases (*e.g.,* all uridines in a polynucleotide disclosed herein can be replaced with an unnatural nucleobase, *e.g.,* 5-methoxyuridine). In some aspects, the polynucleotide (*e.g.,* a synthetic RNA or a synthetic DNA) comprises only natural nucleobases, *i.e.,* A (adenosine), G (guanosine), C (cytidine), and T (thymidine) in the case of a synthetic DNA, or A, C, G, and U (uridine) in the case of a synthetic RNA.

[0929] The skilled artisan will appreciate that the T bases in the codon maps disclosed herein are present in DNA, whereas the T bases would be replaced by U bases in corresponding RNAs. For example, a codon-nucleotide sequence disclosed herein in DNA form, *e.g.,* a vector or an in-vitro translation (IVT) template, would have its T bases transcribed as U based in its corresponding transcribed mRNA. In this respect, both codon-optimized DNA sequences (comprising T) and their corresponding mRNA sequences (comprising U) are considered codon-optimized nucleotide sequence of the present invention. A skilled artisan would also understand that equivalent codon-maps can be generated by replaced one or more bases with non-natural bases. Thus, *e.g.,* a TTC codon (DNA map) would correspond to a UUC codon (RNA map), which in turn would correspond to a ΨΨC codon (RNA map in which U has been replaced with pseudouridine).

[0930] Standard A-T and G-C base pairs form under conditions which allow the formation of hydrogen bonds between the N3-H and C4-oxy of thymidine and the N1

and C6-NH2, respectively, of adenosine and between the C2-oxy, N3 and C4-NH2, of cytidine and the C2-NH2, N′—H and C6-oxy, respectively, of guanosine. Thus, for example, guanosine (2-amino-6-oxy-9-β-D-ribofuranosyl-purine) can be modified to form isoguanosine (2-oxy-6-amino-9-β-D-ribofuranosyl-purine). Such modification results in a nucleoside base which will no longer effectively form a standard base pair with cytosine. However, modification of cytosine (1-β-D-ribofuranosyl-2-oxy-4-amino-pyrimidine) to form isocytosine (1-β-D-ribofuranosyl-2-amino-4-oxy-pyrimidine-) results in a modified nucleotide which will not effectively base pair with guanosine but will form a base pair with isoguanosine (U.S. Pat. No. 5,681,702 to Collins et al.). Isocytosine is available from Sigma Chemical Co. (St. Louis, Mo.); isocytidine can be prepared by the method described by Switzer et al. (1993) Biochemistry 32:10489-10496 and references cited therein; 2′-deoxy-5-methyl-isocytidine can be prepared by the method of Tor et al., 1993, J. Am. Chem. Soc. 115:4461-4467 and references cited therein; and isoguanine nucleotides can be prepared using the method described by Switzer et al., 1993, supra, and Mantsch et al., 1993, Biochem. 14:5593-5601, or by the method described in U.S. Pat. No. 5,780,610 to Collins et al. Other nonnatural base pairs can be synthesized by the method described in Piccirilli et al., 1990, Nature 343:33-37, for the synthesis of 2,6-diaminopyrimidine and its complement (1-methylpyrazolo-[4,3]pyrimidine-5,7-(4H,6H)-dione. Other such modified nucleotide units which form unique base pairs are known, such as those described in Leach et al. (1992) J. Am. Chem. Soc. 114:3675-3683 and Switzer et al., supra.

[0931] *Polypeptide:* The terms "polypeptide," "peptide," and "protein" are used interchangeably herein to refer to polymers of amino acids of any length. The polymer can comprise modified amino acids. The terms also encompass an amino acid polymer that has been modified naturally or by intervention; for example, disulfide bond formation, glycosylation, lipidation, acetylation, phosphorylation, or any other manipulation or modification, such as conjugation with a labeling component. Also included within the definition are, for example, polypeptides containing one or more analogs of an amino acid (including, for example, unnatural amino acids such as homocysteine, ornithine, p-acetylphenylalanine, D-amino acids, and creatine), as well as other modifications known in the art.

[0932] The term, as used herein, refers to proteins, polypeptides, and peptides of any size, structure, or function. Polypeptides include encoded polynucleotide products,

naturally occurring polypeptides, synthetic polypeptides, homologs, orthologs, paralogs, fragments and other equivalents, variants, and analogs of the foregoing. A polypeptide can be a monomer or can be a multi-molecular complex such as a dimer, trimer or tetramer. They can also comprise single chain or multichain polypeptides. Most commonly disulfide linkages are found in multichain polypeptides. The term polypeptide can also apply to amino acid polymers in which one or more amino acid residues are an artificial chemical analogue of a corresponding naturally occurring amino acid. In some embodiments, a "peptide" can be less than or equal to 50 amino acids long, *e.g.,* about 5, 10, 15, 20, 25, 30, 35, 40, 45, or 50 amino acids long.

[0933] *Polypeptide variant*: As used herein, the term "polypeptide variant" refers to molecules that differ in their amino acid sequence from a native or reference sequence. The amino acid sequence variants can possess substitutions, deletions, and/or insertions at certain positions within the amino acid sequence, as compared to a native or reference sequence. Ordinarily, variants will possess at least about 50% identity, at least about 60% identity, at least about 70% identity, at least about 80% identity, at least about 90% identity, at least about 95% identity, at least about 99% identity to a native or reference sequence. In some embodiments, they will be at least about 80%, or at least about 90% identical to a native or reference sequence.

[0934] *Polypeptide per unit drug (PUD)*: As used herein, a PUD or product per unit drug, is defined as a subdivided portion of total daily dose, usually 1 mg, pg, kg, etc., of a product (such as a polypeptide) as measured in body fluid or tissue, usually defined in concentration such as pmol/mL, mmol/mL, etc. divided by the measure in the body fluid.

[0935] *Preventing*: As used herein, the term "preventing" refers to partially or completely delaying onset of an infection, disease, disorder and/or condition; partially or completely delaying onset of one or more symptoms, features, or clinical manifestations of a particular infection, disease, disorder, and/or condition; partially or completely delaying onset of one or more symptoms, features, or manifestations of a particular infection, disease, disorder, and/or condition; partially or completely delaying progression from an infection, a particular disease, disorder and/or condition; and/or decreasing the risk of developing pathology associated with the infection, the disease, disorder, and/or condition.

[0936] *Proliferate:* As used herein, the term "proliferate" means to grow, expand or increase or cause to grow, expand or increase rapidly. "Proliferative" means having

the ability to proliferate. "Anti-proliferative" means having properties counter to or inapposite to proliferative properties.

[0937] *Prophylactic*: As used herein, "prophylactic" refers to a therapeutic or course of action used to prevent the spread of disease.

[0938] *Prophylaxis:* As used herein, a "prophylaxis" refers to a measure taken to maintain health and prevent the spread of disease. An "immune prophylaxis" refers to a measure to produce active or passive immunity to prevent the spread of disease.

[0939] *Protein cleavage site*: As used herein, "protein cleavage site" refers to a site where controlled cleavage of the amino acid chain can be accomplished by chemical, enzymatic or photochemical means.

[0940] *Protein cleavage signal:* As used herein "protein cleavage signal" refers to at least one amino acid that flags or marks a polypeptide for cleavage.

[0941] *Protein of interest:* As used herein, the terms "proteins of interest" or "desired proteins" include those provided herein and fragments, mutants, variants, and alterations thereof.

[0942] *Proximal:* As used herein, the term "proximal" means situated nearer to the center or to a point or region of interest.

[0943] *Pseudouridine:* As used herein, pseudouridine (ψ) refers to the C-glycoside isomer of the nucleoside uridine. A "pseudouridine analog" is any modification, variant, isoform or derivative of pseudouridine. For example, pseudouridine analogs include but are not limited to 1-carboxymethyl-pseudouridine, 1-propynyl-pseudouridine, 1-taurinomethyl-pseudouridine, 1-taurinomethyl-4-thio-pseudouridine, 1-methylpseudouridine ($m^1\psi$) (also known as N1-methyl-pseudouridine), 1-methyl-4-thio-pseudouridine ($m^1s^4\psi$), 4-thio-1-methyl-pseudouridine, 3-methyl-pseudouridine ($m^3\psi$), 2-thio-1-methyl-pseudouridine, 1-methyl-1-deaza-pseudouridine, 2-thio-1-methyl-1-deaza-pseudouridine, dihydropseudouridine, 2-thio-dihydropseudouridine, 2-methoxyuridine, 2-methoxy-4-thio-uridine, 4-methoxy-pseudouridine, 4-methoxy-2-thio-pseudouridine, 1-methyl-3-(3-amino-3-carboxypropyl)pseudouridine ($acp^3\ \psi$), and 2′-O-methyl-pseudouridine (ψm).

[0944] *Purified:* As used herein, "purify," "purified," "purification" means to make substantially pure or clear from unwanted components, material defilement, admixture or imperfection.

[0945] *Reference Nucleic Acid Sequence*: The term "reference nucleic acid sequence" or “reference nucleic acid” or “reference nucleotide sequence” or “reference

sequence" refers to a starting nucleic acid sequence (e.g., a RNA, e.g., an mRNA sequence) that can be sequence optimized. In some embodiments, the reference nucleic acid sequence is a wild type nucleic acid sequence, a fragment or a variant thereof. In some embodiments, the reference nucleic acid sequence is a previously sequence optimized nucleic acid sequence.

[0946] *Salts*: In some aspects, the pharmaceutical composition for delivery disclosed herein and comprises salts of some of their lipid constituents. The term "salt" includes any anionic and cationic complex. Non-limiting examples of anions include inorganic and organic anions, e.g., fluoride, chloride, bromide, iodide, oxalate (e.g., hemioxalate), phosphate, phosphonate, hydrogen phosphate, dihydrogen phosphate, oxide, carbonate, bicarbonate, nitrate, nitrite, nitride, bisulfite, sulfide, sulfite, bisulfate, sulfate, thiosulfate, hydrogen sulfate, borate, formate, acetate, benzoate, citrate, tartrate, lactate, acrylate, polyacrylate, fumarate, maleate, itaconate, glycolate, gluconate, malate, mandelate, tiglate, ascorbate, salicylate, polymethacrylate, perchlorate, chlorate, chlorite, hypochlorite, bromate, hypobromite, iodate, an alkylsulfonate, an arylsulfonate, arsenate, arsenite, chromate, dichromate, cyanide, cyanate, thiocyanate, hydroxide, peroxide, permanganate, and mixtures thereof.

[0947] *Sample:* As used herein, the term "sample" or "biological sample" refers to a subset of its tissues, cells or component parts (*e.g.,* body fluids, including but not limited to blood, mucus, lymphatic fluid, synovial fluid, cerebrospinal fluid, saliva, amniotic fluid, amniotic cord blood, urine, vaginal fluid and semen). A sample further can include a homogenate, lysate or extract prepared from a whole organism or a subset of its tissues, cells or component parts, or a fraction or portion thereof, including but not limited to, for example, plasma, serum, spinal fluid, lymph fluid, the external sections of the skin, respiratory, intestinal, and genitourinary tracts, tears, saliva, milk, blood cells, tumors, organs. A sample further refers to a medium, such as a nutrient broth or gel, which can contain cellular components, such as proteins or nucleic acid molecule.

[0948] *Signal Sequence:* As used herein, the phrases "signal sequence," "signal peptide," and "transit peptide" are used interchangeably and refer to a sequence that can direct the transport or localization of a protein to a certain organelle, cell compartment, or extracellular export. The term encompasses both the signal sequence polypeptide and the nucleic acid sequence encoding the signal sequence. Thus,

references to a signal sequence in the context of a nucleic acid refer in fact to the nucleic acid sequence encoding the signal sequence polypeptide.

[0949] *Signal transduction pathway*: A "signal transduction pathway" refers to the biochemical relationship between a variety of signal transduction molecules that play a role in the transmission of a signal from one portion of a cell to another portion of a cell. As used herein, the phrase "cell surface receptor" includes, for example, molecules and complexes of molecules capable of receiving a signal and the transmission of such a signal across the plasma membrane of a cell.

[0950] *Similarity*: As used herein, the term "similarity" refers to the overall relatedness between polymeric molecules, *e.g.* between polynucleotide molecules (*e.g.* DNA molecules and/or RNA molecules) and/or between polypeptide molecules. Calculation of percent similarity of polymeric molecules to one another can be performed in the same manner as a calculation of percent identity, except that calculation of percent similarity takes into account conservative substitutions as is understood in the art.

[0951] *Single unit dose*: As used herein, a "single unit dose" is a dose of any therapeutic administered in one dose/at one time/single route/single point of contact, *i.e.*, single administration event.

[0952] *Split dose*: As used herein, a "split dose" is the division of single unit dose or total daily dose into two or more doses.

[0953] *Specific delivery*: As used herein, the term “specific delivery,” “specifically deliver,” or “specifically delivering” means delivery of more (*e.g.*, at least 1.5 fold more, at least 2-fold more, at least 3-fold more, at least 4-fold more, at least 5-fold more, at least 6-fold more, at least 7-fold more, at least 8-fold more, at least 9-fold more, at least 10-fold more) of a polynucleotide by a nanoparticle to a target tissue of interest (*e.g.*, mammalian liver) compared to an off-target tissue (*e.g.*, mammalian spleen). The level of delivery of a nanoparticle to a particular tissue can be measured by comparing the amount of protein produced in a tissue to the weight of said tissue, comparing the amount of polynucleotide in a tissue to the weight of said tissue, comparing the amount of protein produced in a tissue to the amount of total protein in said tissue, or comparing the amount of polynucleotide in a tissue to the amount of total polynucleotide in said tissue. For example, for renovascular targeting, a polynucleotide is specifically provided to a mammalian kidney as compared to the liver and spleen if 1.5, 2-fold, 3-fold, 5-fold, 10-fold, 15-fold, or 20-fold more

polynucleotide per 1 g of tissue is delivered to a kidney compared to that delivered to the liver or spleen following systemic administration of the polynucleotide. It will be understood that the ability of a nanoparticle to specifically deliver to a target tissue need not be determined in a subject being treated, it can be determined in a surrogate such as an animal model (*e.g.*, a rat model).

[0954] *Stable:* As used herein "stable" refers to a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and in some cases capable of formulation into an efficacious therapeutic agent.

[0955] *Stabilized:* As used herein, the term "stabilize," "stabilized," "stabilized region" means to make or become stable.

[0956] *Stereoisomer:* As used herein, the term "stereoisomer" refers to all possible different isomeric as well as conformational forms that a compound can possess (*e.g.*, a compound of any formula described herein), in particular all possible stereochemically and conformationally isomeric forms, all diastereomers, enantiomers and/or conformers of the basic molecular structure. Some compounds of the present invention can exist in different tautomeric forms, all of the latter being included within the scope of the present invention.

[0957] *Subject:* By "subject" or "individual" or "animal" or "patient" or "mammal," is meant any subject, particularly a mammalian subject, for whom diagnosis, prognosis, or therapy is desired. Mammalian subjects include, but are not limited to, humans, domestic animals, farm animals, zoo animals, sport animals, pet animals such as dogs, cats, guinea pigs, rabbits, rats, mice, horses, cattle, cows; primates such as apes, monkeys, orangutans, and chimpanzees; canids such as dogs and wolves; felids such as cats, lions, and tigers; equids such as horses, donkeys, and zebras; bears, food animals such as cows, pigs, and sheep; ungulates such as deer and giraffes; rodents such as mice, rats, hamsters and guinea pigs; and so on. In certain embodiments, the mammal is a human subject. In other embodiments, a subject is a human patient. In a particular embodiment, a subject is a human patient in need of treatment.

[0958] *Substantially*: As used herein, the term "substantially" refers to the qualitative condition of exhibiting total or near-total extent or degree of a characteristic or property of interest. One of ordinary skill in the biological arts will understand that biological and chemical characteristics rarely, if ever, go to completion and/or proceed to completeness or achieve or avoid an absolute result. The term

"substantially" is therefore used herein to capture the potential lack of completeness inherent in many biological and chemical characteristics.

[0959] *Substantially equal*: As used herein as it relates to time differences between doses, the term means plus/minus 2%.

[0960] *Substantially simultaneous*: As used herein and as it relates to plurality of doses, the term means within 2 seconds.

[0961] *Suffering from*: An individual who is "suffering from" a disease, disorder, and/or condition has been diagnosed with or displays one or more symptoms of the disease, disorder, and/or condition.

[0962] *Susceptible to*: An individual who is "susceptible to" a disease, disorder, and/or condition has not been diagnosed with and/or cannot exhibit symptoms of the disease, disorder, and/or condition but harbors a propensity to develop a disease or its symptoms. In some embodiments, an individual who is susceptible to a disease, disorder, and/or condition (for example, GSD-Ia) can be characterized by one or more of the following: (1) a genetic mutation associated with development of the disease, disorder, and/or condition; (2) a genetic polymorphism associated with development of the disease, disorder, and/or condition; (3) increased and/or decreased expression and/or activity of a protein and/or nucleic acid associated with the disease, disorder, and/or condition; (4) habits and/or lifestyles associated with development of the disease, disorder, and/or condition; (5) a family history of the disease, disorder, and/or condition; and (6) exposure to and/or infection with a microbe associated with development of the disease, disorder, and/or condition. In some embodiments, an individual who is susceptible to a disease, disorder, and/or condition will develop the disease, disorder, and/or condition. In some embodiments, an individual who is susceptible to a disease, disorder, and/or condition will not develop the disease, disorder, and/or condition.

[0963] *Sustained release:* As used herein, the term "sustained release" refers to a pharmaceutical composition or compound release profile that conforms to a release rate over a specific period of time.

[0964] *Synthetic*: The term "synthetic" means produced, prepared, and/or manufactured by the hand of man. Synthesis of polynucleotides or other molecules of the present invention can be chemical or enzymatic.

[0965] *Targeted Cells:* As used herein, "targeted cells" refers to any one or more cells of interest. The cells can be found *in vitro*, *in vivo*, *in situ* or in the tissue or organ of

an organism. The organism can be an animal, for example a mammal, a human, a subject or a patient.

[0966] *Target tissue*: As used herein "target tissue" refers to any one or more tissue types of interest in which the delivery of a polynucleotide would result in a desired biological and/or pharmacological effect. Examples of target tissues of interest include specific tissues, organs, and systems or groups thereof. In particular applications, a target tissue can be a liver, a kidney, a lung, a spleen, or a vascular endothelium in vessels (e.g., intra-coronary or intra-femoral). An "off-target tissue" refers to any one or more tissue types in which the expression of the encoded protein does not result in a desired biological and/or pharmacological effect.

[0967] The presence of a therapeutic agent in an off-target issue can be the result of: (i) leakage of a polynucleotide from the administration site to peripheral tissue or distant off-target tissue via diffusion or through the bloodstream (e.g., a polynucleotide intended to express a polypeptide in a certain tissue would reach the off-target tissue and the polypeptide would be expressed in the off-target tissue); or (ii) leakage of an polypeptide after administration of a polynucleotide encoding such polypeptide to peripheral tissue or distant off-target tissue via diffusion or through the bloodstream (e.g., a polynucleotide would expressed a polypeptide in the target tissue, and the polypeptide would diffuse to peripheral tissue).

[0968] *Targeting sequence*: As used herein, the phrase "targeting sequence" refers to a sequence that can direct the transport or localization of a protein or polypeptide.

[0969] *Terminus*: As used herein the terms "termini" or "terminus," when referring to polypeptides, refers to an extremity of a peptide or polypeptide. Such extremity is not limited only to the first or final site of the peptide or polypeptide but can include additional amino acids in the terminal regions. The polypeptide based molecules of the invention can be characterized as having both an N-terminus (terminated by an amino acid with a free amino group (NH_2)) and a C-terminus (terminated by an amino acid with a free carboxyl group (COOH)). Proteins of the invention are in some cases made up of multiple polypeptide chains brought together by disulfide bonds or by non-covalent forces (multimers, oligomers). These sorts of proteins will have multiple N- and C-termini. Alternatively, the termini of the polypeptides can be modified such that they begin or end, as the case can be, with a non-polypeptide based moiety such as an organic conjugate.

[0970] *Therapeutic Agent:* The term "therapeutic agent" refers to an agent that, when administered to a subject, has a therapeutic, diagnostic, and/or prophylactic effect and/or elicits a desired biological and/or pharmacological effect. For example, in some embodiments, an mRNA encoding a G6PC polypeptide can be a therapeutic agent.

[0971] *Therapeutically effective amount:* As used herein, the term "therapeutically effective amount" means an amount of an agent to be delivered (*e.g.*, nucleic acid, drug, therapeutic agent, diagnostic agent, prophylactic agent, etc.) that is sufficient, when administered to a subject suffering from or susceptible to an infection, disease, disorder, and/or condition, to treat, improve symptoms of, diagnose, prevent, and/or delay the onset of the infection, disease, disorder, and/or condition.

[0972] *Therapeutically effective outcome*: As used herein, the term "therapeutically effective outcome" means an outcome that is sufficient in a subject suffering from or susceptible to an infection, disease, disorder, and/or condition, to treat, improve symptoms of, diagnose, prevent, and/or delay the onset of the infection, disease, disorder, and/or condition.

[0973] *Total daily dose:* As used herein, a "total daily dose" is an amount given or prescribed in 24 hr. period. The total daily dose can be administered as a single unit dose or a split dose.

[0974] *Transcription factor:* As used herein, the term "transcription factor" refers to a DNA-binding protein that regulates transcription of DNA into RNA, for example, by activation or repression of transcription. Some transcription factors effect regulation of transcription alone, while others act in concert with other proteins. Some transcription factor can both activate and repress transcription under certain conditions. In general, transcription factors bind a specific target sequence or sequences highly similar to a specific consensus sequence in a regulatory region of a target gene. Transcription factors can regulate transcription of a target gene alone or in a complex with other molecules.

[0975] *Transcription*: As used herein, the term "transcription" refers to methods to produce mRNA (*e.g.*, an mRNA sequence or template) from DNA (*e.g.*, a DNA template or sequence)

[0976] *Transfection*: As used herein, "transfection" refers to the introduction of a polynucleotide (e.g., exogenous nucleic acids) into a cell wherein a polypeptide encoded by the polynucleotide is expressed (e.g., mRNA) or the polypeptide

modulates a cellular function (e.g., siRNA, miRNA). As used herein, "expression" of a nucleic acid sequence refers to translation of a polynucleotide (e.g., an mRNA) into a polypeptide or protein and/or post-translational modification of a polypeptide or protein. Methods of transfection include, but are not limited to, chemical methods, physical treatments and cationic lipids or mixtures.

[0977] *Treating, treatment, therapy*: As used herein, the term "treating" or "treatment" or "therapy" refers to partially or completely alleviating, ameliorating, improving, relieving, delaying onset of, inhibiting progression of, reducing severity of, and/or reducing incidence of one or more symptoms or features of a disease, *e.g.*, hypoglycemia, GSD-Ia. For example, "treating" GSD-Ia can refer to diminishing symptoms associate with the disease, prolong the lifespan (increase the survival rate) of patients, reducing the severity of the disease, preventing or delaying the onset of the disease, etc. Treatment can be administered to a subject who does not exhibit signs of a disease, disorder, and/or condition and/or to a subject who exhibits only early signs of a disease, disorder, and/or condition for the purpose of decreasing the risk of developing pathology associated with the disease, disorder, and/or condition.

[0978] *Unmodified*: As used herein, "unmodified" refers to any substance, compound or molecule prior to being changed in some way. Unmodified can, but does not always, refer to the wild type or native form of a biomolecule. Molecules can undergo a series of modifications whereby each modified molecule can serve as the "unmodified" starting molecule for a subsequent modification.

[0979] *Uracil*: Uracil is one of the four nucleobases in the nucleic acid of RNA, and it is represented by the letter U. Uracil can be attached to a ribose ring, or more specifically, a ribofuranose via a β-N_1-glycosidic bond to yield the nucleoside uridine. The nucleoside uridine is also commonly abbreviated according to the one letter code of its nucleobase, i.e., U. Thus, in the context of the present disclosure, when a monomer in a polynucleotide sequence is U, such U is designated interchangeably as a "uracil" or a "uridine."

[0980] *Uridine Content*: The terms "uridine content" or "uracil content" are interchangeable and refer to the amount of uracil or uridine present in a certain nucleic acid sequence. Uridine content or uracil content can be expressed as an absolute value (total number of uridine or uracil in the sequence) or relative (uridine

or uracil percentage respect to the total number of nucleobases in the nucleic acid sequence).

[0981] *Uridine-Modified Sequence*: The terms "uridine-modified sequence" refers to a sequence optimized nucleic acid (e.g., a synthetic mRNA sequence) with a different overall or local uridine content (higher or lower uridine content) or with different uridine patterns (e.g., gradient distribution or clustering) with respect to the uridine content and/or uridine patterns of a candidate nucleic acid sequence. In the content of the present disclosure, the terms "uridine-modified sequence" and "uracil-modified sequence" are considered equivalent and interchangeable.

[0982] A "high uridine codon" is defined as a codon comprising two or three uridines, a "low uridine codon" is defined as a codon comprising one uridine, and a "no uridine codon" is a codon without any uridines. In some embodiments, a uridine-modified sequence comprises substitutions of high uridine codons with low uridine codons, substitutions of high uridine codons with no uridine codons, substitutions of low uridine codons with high uridine codons, substitutions of low uridine codons with no uridine codons, substitution of no uridine codons with low uridine codons, substitutions of no uridine codons with high uridine codons, and combinations thereof. In some embodiments, a high uridine codon can be replaced with another high uridine codon. In some embodiments, a low uridine codon can be replaced with another low uridine codon. In some embodiments, a no uridine codon can be replaced with another no uridine codon. A uridine-modified sequence can be uridine enriched or uridine rarefied.

[0983] *Uridine Enriched*: As used herein, the terms "uridine enriched" and grammatical variants refer to the increase in uridine content (expressed in absolute value or as a percentage value) in a sequence optimized nucleic acid (e.g., a synthetic mRNA sequence) with respect to the uridine content of the corresponding candidate nucleic acid sequence. Uridine enrichment can be implemented by substituting codons in the candidate nucleic acid sequence with synonymous codons containing less uridine nucleobases. Uridine enrichment can be global (i.e., relative to the entire length of a candidate nucleic acid sequence) or local (i.e., relative to a subsequence or region of a candidate nucleic acid sequence).

[0984] *Uridine Rarefied*: As used herein, the terms "uridine rarefied" and grammatical variants refer to a decrease in uridine content (expressed in absolute value or as a percentage value) in a sequence optimized nucleic acid (e.g., a synthetic

mRNA sequence) with respect to the uridine content of the corresponding candidate nucleic acid sequence. Uridine rarefication can be implemented by substituting codons in the candidate nucleic acid sequence with synonymous codons containing less uridine nucleobases. Uridine rarefication can be global (i.e., relative to the entire length of a candidate nucleic acid sequence) or local (i.e., relative to a subsequence or region of a candidate nucleic acid sequence).

[0985] *Variant*: The term variant as used in present disclosure refers to both natural variants (e.g., polymorphisms, isoforms, etc.) and artificial variants in which at least one amino acid residue in a native or starting sequence (e.g., a wild type sequence) has been removed and a different amino acid inserted in its place at the same position. These variants can be described as "substitutional variants." The substitutions can be single, where only one amino acid in the molecule has been substituted, or they can be multiple, where two or more amino acids have been substituted in the same molecule. If amino acids are inserted or deleted, the resulting variant would be an "insertional variant" or a "deletional variant" respectively.

[0986] *Initiation Codon*: As used herein, the term “initiation codon”, used interchangeably with the term “start codon”, refers to the first codon of an open reading frame that is translated by the ribosome and is comprised of a triplet of linked adenine-uracil-guanine nucleobases. The initiation codon is depicted by the first letter codes of adenine (A), uracil (U), and guanine (G) and is often written simply as “AUG”. Although natural mRNAs may use codons other than AUG as the initiation codon, which are referred to herein as “alternative initiation codons”, the initiation codons of polynucleotides described herein use the AUG codon. During the process of translation initiation, the sequence comprising the initiation codon is recognized via complementary base-pairing to the anticodon of an initiator tRNA (Met-tRNA$_i^{Met}$) bound by the ribosome. Open reading frames may contain more than one AUG initiation codon, which are referred to herein as “alternate initiation codons”.

[0987] The initiation codon plays a critical role in translation initiation. The initiation codon is the first codon of an open reading frame that is translated by the ribosome. Typically, the initiation codon comprises the nucleotide triplet AUG, however, in some instances translation initiation can occur at other codons comprised of distinct nucleotides. The initiation of translation in eukaryotes is a multistep biochemical process that involves numerous protein-protein, protein-RNA, and RNA-RNA interactions between messenger RNA molecules (mRNAs), the 40S ribosomal

subunit, other components of the translation machinery (e.g., eukaryotic initiation factors; eIFs). The current model of mRNA translation initiation postulates that the pre-initiation complex (alternatively "43S pre-initiation complex"; abbreviated as "PIC") translocates from the site of recruitment on the mRNA (typically the 5' cap) to the initiation codon by scanning nucleotides in a 5′ to 3′ direction until the first AUG codon that resides within a specific translation-promotive nucleotide context (the Kozak sequence) is encountered (Kozak (1989) J Cell Biol 108:229-241). Scanning by the PIC ends upon complementary base-pairing between nucleotides comprising the anticodon of the initiator Met-$tRNA_i^{Met}$ transfer RNA and nucleotides comprising the initiation codon of the mRNA. Productive base-pairing between the AUG codon and the Met-$tRNA_i^{Met}$ anticodon elicits a series of structural and biochemical events that culminate in the joining of the large 60S ribosomal subunit to the PIC to form an active ribosome that is competent for translation elongation.

[0988] *Kozak Sequence:* The term "Kozak sequence" (also referred to as "Kozak consensus sequence") refers to a translation initiation enhancer element to enhance expression of a gene or open reading frame, and which in eukaryotes, is located in the 5' UTR. The Kozak consensus sequence was originally defined as the sequence GCCRCC (SEQ ID NO:41), where R = a purine, following an analysis of the effects of single mutations surrounding the initiation codon (AUG) on translation of the preproinsulin gene (Kozak (1986) Cell 44:283-292). Polynucleotides disclosed herein comprise a Kozak consensus sequence, or a derivative or modification thereof. (Examples of translational enhancer compositions and methods of use thereof, see U.S. Pat. No. 5,807,707 to Andrews et al., incorporated herein by reference in its entirety; U.S. Pat. No. 5,723,332 to Chernajovsky, incorporated herein by reference in its entirety; U.S. Pat. No. 5,891,665 to Wilson, incorporated herein by reference in its entirety.)

[0989] *Modified*: As used herein "modified" or "modification" refers to a changed state or a change in composition or structure of a polynucleotide (e.g., mRNA). Polynucleotides may be modified in various ways including chemically, structurally, and/or functionally. For example, polynucleotides may be structurally modified by the incorporation of one or more RNA elements, wherein the RNA element comprises a sequence and/or an RNA secondary structure(s) that provides one or more functions (e.g., translational regulatory activity). Accordingly, polynucleotides of the disclosure

may be comprised of one or more modifications (e.g., may include one or more chemical, structural, or functional modifications, including any combination thereof).

[0990] *Nucleobase*: As used herein, the term "nucleobase" (alternatively "nucleotide base" or "nitrogenous base") refers to a purine or pyrimidine heterocyclic compound found in nucleic acids, including any derivatives or analogs of the naturally occurring purines and pyrimidines that confer improved properties (e.g., binding affinity, nuclease resistance, chemical stability) to a nucleic acid or a portion or segment thereof. Adenine, cytosine, guanine, thymine, and uracil are the nucleobases predominately found in natural nucleic acids. Other natural, non-natural, and/or synthetic nucleobases, as known in the art and/or described herein, can be incorporated into nucleic acids.

[0991] *Nucleoside/Nucleotide*: As used herein, the term "nucleoside" refers to a compound containing a sugar molecule (e.g., a ribose in RNA or a deoxyribose in DNA), or derivative or analog thereof, covalently linked to a nucleobase (e.g., a purine or pyrimidine), or a derivative or analog thereof (also referred to herein as "nucleobase"), but lacking an internucleoside linking group (e.g., a phosphate group). As used herein, the term "nucleotide" refers to a nucleoside covalently bonded to an internucleoside linking group (e.g., a phosphate group), or any derivative, analog, or modification thereof that confers improved chemical and/or functional properties (e.g., binding affinity, nuclease resistance, chemical stability) to a nucleic acid or a portion or segment thereof.

[0992] *Nucleic acid:* As used herein, the term "nucleic acid" is used in its broadest sense and encompasses any compound and/or substance that includes a polymer of nucleotides, or derivatives or analogs thereof. These polymers are often referred to as "polynucleotides". Accordingly, as used herein the terms "nucleic acid" and "polynucleotide" are equivalent and are used interchangeably. Exemplary nucleic acids or polynucleotides of the disclosure include, but are not limited to, ribonucleic acids (RNAs), deoxyribonucleic acids (DNAs), DNA-RNA hybrids, RNAi-inducing agents, RNAi agents, siRNAs, shRNAs, mRNAs, modified mRNAs, miRNAs, antisense RNAs, ribozymes, catalytic DNA, RNAs that induce triple helix formation, threose nucleic acids (TNAs), glycol nucleic acids (GNAs), peptide nucleic acids (PNAs), locked nucleic acids (LNAs, including LNA having a β-D-ribo configuration, α-LNA having an α-L-ribo configuration (a diastereomer of LNA), 2'-amino-LNA

having a 2'-amino functionalization, and 2'-amino-α-LNA having a 2'-amino functionalization) or hybrids thereof.

[0993] *Nucleic Acid Structure*: As used herein, the term "nucleic acid structure" (used interchangeably with "polynucleotide structure") refers to the arrangement or organization of atoms, chemical constituents, elements, motifs, and/or sequence of linked nucleotides, or derivatives or analogs thereof, that comprise a nucleic acid (e.g., an mRNA). The term also refers to the two-dimensional or three-dimensional state of a nucleic acid. Accordingly, the term "RNA structure" refers to the arrangement or organization of atoms, chemical constituents, elements, motifs, and/or sequence of linked nucleotides, or derivatives or analogs thereof, comprising an RNA molecule (e.g., an mRNA) and/or refers to a two-dimensional and/or three-dimensional state of an RNA molecule. Nucleic acid structure can be further demarcated into four organizational categories referred to herein as "molecular structure", "primary structure", "secondary structure", and "tertiary structure" based on increasing organizational complexity.

[0994] *Open Reading Frame*: As used herein, the term "open reading frame", abbreviated as "ORF", refers to a segment or region of an mRNA molecule that encodes a polypeptide. The ORF comprises a continuous stretch of non-overlapping, in-frame codons, beginning with the initiation codon and ending with a stop codon, and is translated by the ribosome.

[0995] *Pre-Initiation Complex (PIC)*: As used herein, the term "pre-initiation complex" (alternatively "43S pre-initiation complex"; abbreviated as "PIC") refers to a ribonucleoprotein complex comprising a 40S ribosomal subunit, eukaryotic initiation factors (eIF1, eIF1A, eIF3, eIF5), and the eIF2-GTP-Met-tRNA$_i^{Met}$ ternary complex, that is intrinsically capable of attachment to the 5' cap of an mRNA molecule and, after attachment, of performing ribosome scanning of the 5' UTR.

[0996] *RNA element*: As used herein, the term "RNA element" refers to a portion, fragment, or segment of an RNA molecule that provides a biological function and/or has biological activity (e.g., translational regulatory activity). Modification of a polynucleotide by the incorporation of one or more RNA elements, such as those described herein, provides one or more desirable functional properties to the modified polynucleotide. RNA elements, as described herein, can be naturally-occurring, non-naturally occurring, synthetic, engineered, or any combination thereof. For example, naturally-occurring RNA elements that provide a regulatory activity include elements

found throughout the transcriptomes of viruses, prokaryotic and eukaryotic organisms (e.g., humans). RNA elements in particular eukaryotic mRNAs and translated viral RNAs have been shown to be involved in mediating many functions in cells. Exemplary natural RNA elements include, but are not limited to, translation initiation elements (e.g., internal ribosome entry site (IRES), see Kieft et al., (2001) RNA 7(2):194-206), translation enhancer elements (e.g., the APP mRNA translation enhancer element, see Rogers et al., (1999) J Biol Chem 274(10):6421-6431), mRNA stability elements (e.g., AU-rich elements (AREs), see Garneau et al., (2007) Nat Rev Mol Cell Biol 8(2):113-126), translational repression element (see e.g., Blumer et al., (2002) Mech Dev 110(1-2):97-112), protein-binding RNA elements (e.g., iron-responsive element, see Selezneva et al., (2013) J Mol Biol 425(18):3301-3310), cytoplasmic polyadenylation elements (Villalba et al., (2011) Curr Opin Genet Dev 21(4):452-457), and catalytic RNA elements (e.g., ribozymes, see Scott et al., (2009) Biochim Biophys Acta 1789(9-10):634-641).

[0997] *Residence time*: As used herein, the term "residence time" refers to the time of occupancy of a pre-initiation complex (PIC) or a ribosome at a discrete position or location along an mRNA molecule.

[0998] *Translational Regulatory Activity*: As used herein, the term "translational regulatory activity" (used interchangeably with "translational regulatory function") refers to a biological function, mechanism, or process that modulates (e.g., regulates, influences, controls, varies) the activity of the translational apparatus, including the activity of the PIC and/or ribosome. In some aspects, the desired translation regulatory activity promotes and/or enhances the translational fidelity of mRNA translation. In some aspects, the desired translational regulatory activity reduces and/or inhibits leaky scanning.

28. Equivalents and Scope

[0999] Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments in accordance with the invention described herein. The scope of the present invention is not intended to be limited to the above Description, but rather is as set forth in the appended claims.

[1000] In the claims, articles such as "a," "an," and "the" can mean one or more than one unless indicated to the contrary or otherwise evident from the context. Claims or

descriptions that include "or" between one or more members of a group are considered satisfied if one, more than one, or all of the group members are present in, employed in, or otherwise relevant to a given product or process unless indicated to the contrary or otherwise evident from the context. The invention includes embodiments in which exactly one member of the group is present in, employed in, or otherwise relevant to a given product or process. The invention includes embodiments in which more than one, or all of the group members are present in, employed in, or otherwise relevant to a given product or process.

[1001] It is also noted that the term "comprising" is intended to be open and permits but does not require the inclusion of additional elements or steps. When the term "comprising" is used herein, the term "consisting of" is thus also encompassed and disclosed.

[1002] Where ranges are given, endpoints are included. Furthermore, it is to be understood that unless otherwise indicated or otherwise evident from the context and understanding of one of ordinary skill in the art, values that are expressed as ranges can assume any specific value or subrange within the stated ranges in different embodiments of the invention, to the tenth of the unit of the lower limit of the range, unless the context clearly dictates otherwise.

[1003] In addition, it is to be understood that any particular embodiment of the present invention that falls within the prior art can be explicitly excluded from any one or more of the claims. Since such embodiments are deemed to be known to one of ordinary skill in the art, they can be excluded even if the exclusion is not set forth explicitly herein. Any particular embodiment of the compositions of the invention (*e.g.*, any nucleic acid or protein encoded thereby; any method of production; any method of use; etc.) can be excluded from any one or more claims, for any reason, whether or not related to the existence of prior art.

[1004] All cited sources, for example, references, publications, databases, database entries, and art cited herein, are incorporated into this application by reference, even if not expressly stated in the citation. In case of conflicting statements of a cited source and the instant application, the statement in the instant application shall control.

[1005] Section and table headings are not intended to be limiting.

CONSTRUCT SEQUENCES

By "**G5**" is meant that all uracils (U) in the mRNA are replaced by N1-methylpseudouracils.

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
SEQ ID NO:		1	2	3	150	45
	hG6PC-FLAG G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSLDY KDDDDK	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 45 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:2, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUGGACUA CAAGGACGACGAUG ACAAG			
SEQ ID NO:		**21**	**5**	**3**	**150**	**46**
	mG6PC G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNILHDFGIQS TRYLQVNYQDSQDW FILVSVIADLRNAFYV LFPIWFHLKETVGINL LWVAVVGDWFNLVF KWILFGQRPYWWVL DTDYYSNSSVPIIKQF PVTCETGPGSPSGHA MGAAGVYYVMVTST LAIFRGKKKPTYGFR CLNVILWLGFWAVQ LNVCLSRIYLAAHFP HQVVAGVLSGIAVAE TFSHIRGIYNASLRKY CLITIFLFGFALGFYL LLKGLGVDLLWTLE KAKRWCERPEWVHL DTTPFASLFKNLGTL LGLGLALNSSMYRKS	AUGGAGGAAGGAA UGAACAUUCUCCAU GACUUUGGGAUCCA GUCGACUCGCUAUC UCCAAGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCCUUG UGUCUGUGAUUGCU GACCUGAGGAACGC CUUCUAUGUCCUCU UUCCCAUCUGGUUC CAUCUUAAAGAGAC UGUGGGCAUCAAUC UCCUCUGGGUGGCA GUGGUCGGAGACUG GUUCAACCUCGUCU UCAAGUGGAUUCUG UUUGGACAACGCCC GUAUUGGUGGGUCC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU	SEQ ID NO: 46 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:5, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		CKGELSKLLPFRFACI VASLVLLHLFDSLKP PSQVELIFYILSFCKS ATVPFASVSLIPYCLA RILGQTHKKSL	UGGACACCGACUAC UACAGCAACAGCUC CGUGCCUAUAAUAA AGCAGUUCCCUGUC ACCUGUGAGACCGG ACCAGGAAGUCCCU CUGGCCAUGCCAUG GGCGCAGCAGGUGU AUACUAUGUUAUGG UCACUUCUACUCUU GCUAUCUUUCGAGG AAAGAAAAAGCCAA CGUAUGGAUUCCGG UGUUUGAACGUCAU CUUGUGGUUGGGAU UCUGGGCUGUGCAG CUGAACGUCUGUCU GUCCCGGAUCUACC UUGCUGCUCACUUU CCCCACCAGGUCGU GGCUGGAGUCUUGU CAGGCAUUGCUGUG GCUGAAACUUUCAG CCACAUCCGGGGCA UCUACAAUGCCAGC CUCCGGAAGUAUUG UCUCAUCACCAUCU UCUUGUUUGGUUUC GCGCUUGGAUUCUA CCUGCUACUAAAAG GGCUAGGGGUGGAC CUCCUGUGGACUUU GGAGAAAGCUAAGA GAUGGUGUGAGCGG CCAGAAUGGGUCCA CCUUGACACUACAC CCUUUGCCAGCCUC UUCAAAAACCUGGG AACCCUCUUGGGGU UGGGGCUGGCCCUC AACUCCAGCAUGUA CCGGAAGAGCUGCA AGGGAGAACUCAGC AAGUUGCUCCCAUU CCGCUUCGCCUGCA UUGUGGCUUCCUUG GUCCUCCUGCAUCU CUUUGACUCUCUGA AGCCCCCAUCCCAG GUUGAGUUGAUCUU CUACAUCUUGUCUU UCUGCAAGAGCGCA ACAGUUCCCUUUGC		GAGUGG GCGGC	

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AUCUGUCAGUCUUA UCCCAUACUGCCUA GCCCGGAUCCUGGG ACAGACACACAAGA AGUCUUUG			
SEQ ID NO:		**82**	**6**	**3**	**150**	**47**
	mG6PC-FLAG G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNILHDFGIQS TRYLQVNYQDSQDW FILVSVIADLRNAFYV LFPIWFHLKETVGINL LWVAVVGDWFNLVF KWILFGQRPYWWVL DTDYYSNSSVPIIKQF PVTCETGPGSPSGHA MGAAGVYYVMVTST LAIFRGKKKPTYGFR CLNVILWLGFWAVQ LNVCLSRIYLAAHFP HQVVAGVLSGIAVAE TFSHIRGIYNASLRKY CLITIFLFGFALGFYL LLKGLGVDLLWTLE KAKRWCERPEWVHL DTTPFASLFKNLGTL LGLGLALNSSMYRKS CKGELSKLLPFRFACI VASLVLLHLFDSLKP PSQVELIFYILSFCKS ATVPFASVSLIPYCLA RILGQTHKKSLDYKD DDDK	AUGGAGGAAGGAA UGAACAUUCUCCAU GACUUUGGGAUCCA GUCGACUCGCUAUC UCCAAGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCCUUG UGUCUGUGAUUGCU GACCUGAGGAACGC CUUCUAUGUCCUCU UUCCCAUCUGGUUC CAUCUUAAAGAGAC UGUGGGCAUCAAUC UCCUCUGGGUGGCA GUGGUCGGAGACUG GUUCAACCUCGUCU UCAAGUGGAUUCUG UUUGGACAACGCCC GUAUUGGUGGGUCC UGGACACCGACUAC UACAGCAACAGCUC CGUGCCUAUAAUAA AGCAGUUCCCUGUC ACCUGUGAGACCGG ACCAGGAAGUCCCU CUGGCCAUGCCAUG GGCGCAGCAGGUGU AUACUAUGUUAUGG UCACUUCUACUCUU GCUAUCUUUCGAGG AAAGAAAAAGCCAA CGUAUGGAUUCCGG UGUUUGAACGUCAU CUUGUGGUUGGGAU UCUGGGCUGUGCAG CUGAACGUCUGUCU GUCCCGGAUCUACC UUGCUGCUCACUUU CCCCACCAGGUCGU GGCUGGAGUCUUGU CAGGCAUUGCUGUG GCUGAAACUUUCAG CCACAUCCGGGGCA UCUACAAUGCCAGC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 47 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:6, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CUCCGGAAGUAUUG UCUCAUCACCAUCU UCUUGUUUGGUUUC GCGCUUGGAUUCUA CCUGCUACUAAAAG GGCUAGGGGUGGAC CUCCUGUGGACUUU GGAGAAAGCUAAGA GAUGGUGUGAGCGG CCAGAAUGGGUCCA CCUUGACACUACAC CCUUUGCCAGCCUC UUCAAAAACCUGGG AACCCUCUUGGGGU UGGGGCUGGCCCUC AACUCCAGCAUGUA CCGGAAGAGCUGCA AGGGAGAACUCAGC AAGUUGCUCCCAUU CCGCUUCGCCUGCA UUGUGGCUUCCUUG GUCCUCCUGCAUCU CUUUGACUCUCUGA AGCCCCCAUCCCAG GUUGAGUUGAUCUU CUACAUCUUGUCUU UCUGCAAGAGCGCA ACAGUUCCCUUUGC AUCUGUCAGUCUUA UCCCAUACUGCCUA GCCCGGAUCCUGGG ACAGACACACAAGA AGUCUUUGGACUAC AAGGACGACGAUGA CAAG			
SEQ ID NO:		**199**	**7**	**3**	**150**	**48**
	hG6PC G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC	SEQ ID NO: 48 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:7, and 3' UTR of

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG		CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**199**	**7**	**3**	**150**	**49**
	hG6PC WT1 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 49 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:7, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**83**	**9**	**3**	**150**	**50**
	hG6PC V1 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC	SEQ ID NO: 50 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCGGCUGU		CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	NO:9, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AUUGUAGCCUCCCU CGUCCUCCUGCACC UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGAUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUCUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**84**	**10**	**3**	**150**	**51**
	hG6PC V2 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUAGGGG AAAGAAGAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 51 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:10, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGAG GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUGU AUUGUAGCCUCCCU CGUCCUCCUGCACC UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**199**	**7**	**3**	**175**	**52**
	hG6PC WT2 G5 Cap: C1	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC	SEQ ID NO: 52 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3,

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	PolyA tail: 100nt (SEQ ID NO:223)	GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC	UAAGAG CCACC	UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	ORF Sequence of SEQ ID NO:7, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**195**	**12**	**3**	**175**	**53**
	hG6PC Q139R G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUAGGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 53 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:12, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**197**	**13**	**3**	**175**	**54**
	hG6PC I142K G5 Cap: C1	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG	SEQ ID NO: 54 consists from 5' to 3' end: 5'

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	PolyA tail: 100nt (SEQ ID NO:223)	VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAAGAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC	GAAAUA UAAGAG CCACC	CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:13, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**196**	**14**	**3**	**175**	**55**
	hG6PC Q247R G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 46 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:14, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGAG GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**198**	**15**	**3**	**175**	**56**
	hG6PC S297G	MEEGMNVLHDFGIQ STHYLQVNYQDSQD	AUGGAGGAAGGAA UGAAUGUUCUCCAU	GGGAAA UAAGAG	UGAUAA UAGGCU	SEQ ID NO: 56

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA	AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:15, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCGGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**194**	**16**	**3**	**175**	**57**
	hG6PC S298C G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 57 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:16, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUGU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
SEQ ID NO:		200	17	3	175	58
	hG6PC V308L G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 58 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:17, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACC UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**201**	**18**	**3**	**175**	**59**
	hG6PC V321I G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 59 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:18, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		PSQVELIFYVLSFCKS AVVPLASVSVIPYCL AQVLGQPHKKSL	CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGAUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU			

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**202**	**19**	**3**	**175**	**60**
	hG6PC V321L G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELLFYVLSFCKS AVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 60 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:19, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGCUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**203**	**20**	**3**	**175**	**61**
	hG6PC V340L G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU	SEQ ID NO: 61 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:20, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSLIPYCL AQVLGQPHKKSL	AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC		GAGUGG GCGGC	

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			GGUAGUGCCCCUGG CAUCCGUCAGUCUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**204**	**11**	**3**	**175**	**62**
	hG6PC V332A G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSS IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAAVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 62 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:11, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUCU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGCAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**205**	**8**	**3**	**175**	**63**
	hG6PC Q139R/Q247R/S 298C G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU	SEQ ID NO: 63 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:8, and 3' UTR of

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUAGGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGAG GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUGU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA		GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**206**	**207**	**3**	**175**	**64**
	hG6PC Q247R/S298C G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 64 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:207, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGAG GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCUGU AUUGUAGCCUCCCU CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**208**	**22**	**3**	**175**	**65**
	hG6PC S298A G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH	AUGGAGGAAGGAA UGAAUGUUCUCCAU GACUUUGGGAUCCA GUCAACACAUUACC UCCAGGUGAAUUAC CAAGACUCCCAGGA CUGGUUCAUCUUGG UGUCCGUGAUCGCA GACCUCAGGAAUGC CUUCUACGUCCUCU UCCCCAUCUGGUUC CAUCUUCAGGAAGC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC	SEQ ID NO: 65 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:22,

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSA IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	UGUGGGCAUUAAAC UCCUUUGGGUAGCU GUGAUUGGAGACUG GCUCAACCUCGUCU UUAAGUGGAUUCUC UUUGGACAGCGUCC AUACUGGUGGGUUU UGGAUACUGACUAC UACAGCAACACUUC CGUGCCCCUGAUAA AGCAGUUCCCUGUA ACCUGUGAGACUGG ACCAGGGAGCCCCU CUGGCCAUGCCAUG GGCACAGCAGGUGU AUACUACGUGAUGG UCACAUCUACUCUU UCCAUCUUUCAGGG AAAGAUAAAGCCGA CCUACAGAUUUCGG UGCUUGAAUGUCAU UUUGUGGUUGGGA UUCUGGGCUGUGCA GCUGAAUGUCUGUC UGUCACGAAUCUAC CUUGCUGCUCAUUU UCCUCAUCAAGUUG UUGCUGGAGUCCUG UCAGGCAUUGCUGU UGCAGAAACUUUCA GCCACAUCCACAGC AUCUAUAAUGCCAG CCUCAAGAAAUAUU UUCUCAUUACCUUC UUCCUGUUCAGCUU CGCCAUCGGAUUUU AUCUGCUGCUCAAG GGACUGGGUGUAGA CCUCCUGUGGACUC UGGAGAAAGCCCAG AGGUGGUGCGAGCA GCCAGAAUGGGUCC ACAUUGACACCACA CCCUUUGCCAGCCU CCUCAAGAACCUGG GCACGCUCUUUGGC CUGGGGCUGGCUCU CAACUCCAGCAUGU ACAGGGAGAGCUGC AAGGGGAAACUCAG CAAGUGGCUCCCAU UCCGCCUCAGCGCU AUUGUAGCCUCCCU		CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CGUCCUCCUGCACG UCUUUGACUCCUUG AAACCCCCAUCCCA AGUCGAGCUGGUCU UCUACGUCUUGUCC UUCUGCAAGAGUGC GGUAGUGCCCCUGG CAUCCGUCAGUGUC AUCCCCUACUGCCU CGCCCAGGUCCUGG GCCAGCCGCACAAG AAGUCGUUG			
SEQ ID NO:		**194**	**23**	**3**	**175**	**66**
	hG6PC S298C_CoV1 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 66 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:23, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACAAC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA AGUGGCUCCCUUUC AGACUUAGCUGCAU CGUGGCGUCUCUCG UGCUGCUGCACGUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGGUGUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCGUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			
SEQ ID NO:		**205**	**24**	**3**	**175**	**67**
	hG6PC Q139R/Q247R/S 298C_CoV1 G5 Cap: C1	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC	SEQ ID NO: 67 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	PolyA tail: 100nt (SEQ ID NO:223)	VTSTLSIFRGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCAGGGGC AAGAUCAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACGGC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA		CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	Sequence of SEQ ID NO:24, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AGUGGCUCCCUUUC AGACUUAGCUGCAU CGUGGCGUCUCUCG UGCUGCUGCACGUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGGUGUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCGUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			
SEQ ID NO:		**209**	**25**	**3**	**175**	**68**
	hG6PC M8-CoV1 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCAGGGGC AAGAAGAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 68 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:25, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACGGC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA AGUGGCUCCCUUUC AGACUUGGAUGCAU CGUGGCGUCUCUCG UGCUGCUGCACCUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGAUUUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCCUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			
SEQ ID NO:		**194**	**26**	**3**	**175**	**69**
	hG6PC S298C_CoV2 G5 Cap: C1	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW	AUGGAGGAGGGCAU GAACGUACUUCACG ACUUCGGCAUCCAG AGCACCCACUACUU GCAGGUGAACUACC AGGACUCCCAGGAU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU	SEQ ID NO: 69 consists from 5' to 3' end: 5' UTR of

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	PolyA tail: 100nt (SEQ ID NO:223)	VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	UGGUUCAUUCUCGU GAGCGUGAUUGCCG AUCUCCGGAACGCU UUCUACGUGCUGUU CCCUAUUUGGUUCC ACCUGCAGGAGGCU GUGGGCAUCAAGUU GCUGUGGGUCGCCG UGAUAGGCGACUGG CUCAACUUGGUGUU CAAGUGGAUUCUGU UCGGCCAGCGUCCA UACUGGUGGGUGCU GGACACAGACUAUU AUUCUAAUACAAGC GUACCUCUGAUCAA GCAGUUCCCAGUGA CCUGCGAAACGGGA CCUGGAUCACCAAG CGGCCACGCCAUGG GAACGGCUGGCGUG UACUACGUGAUGGU GACCAGCACCCUCA GCAUCUUCCAGGGA AAGAUCAAGCCGAC UUACAGAUUCCGUU GCCUGAACGUGAUC CUGUGGCUCGGAUU CUGGGCCGUGCAGC UCAACGUGUGCCUG AGUAGAAUCUACCU GGCCGCUCACUUCC CUCACCAGGUAGUG GCCGGCGUGCUGAG CGGCAUUGCCGUGG CCGAGACAUUCAGU CACAUCCACAGCAU CUAUAACGCUAGCC UGAAGAAGUAUUUC CUGAUUACAUUCUU CCUGUUCUCUUUCG CUAUCGGCUUCUAC CUCCUCCUGAAGGG CCUGGGCGUGGACC UGUUGUGGACCCUG GAGAAGGCUCAGAG GUGGUGCGAGCAAC CUGAGUGGGUCCAU AUCGAUACCACCCC AUUCGCAAGUCUGC UGAAGAACCUGGGC ACACUCUUCGGCCU GGGCCUGGCCCUGA	UAAGAG CCACC	GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO:3, ORF Sequence of SEQ ID NO:26, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			ACAGCUCGAUGUAC AGAGAGUCUUGCAA GGGAAAGCUUAGCA AGUGGCUCCCUUUC AGACUGUCUUGCAU CGUGGCCAGCCUGG UGCUGCUGCAUGUG UUCGACUCUCUGAA GCCUCCUUCACAGG UCGAGCUGGUUUUC UAUGUGCUGAGUUU CUGCAAGUCCGCUG UGGUGCCUCUGGCU AGCGUGAGCGUGAU CCCUUACUGCCUGG CCCAGGUGCUGGGC CAGCCUCACAAGAA GUCCCUA			
SEQ ID NO:		**205**	**27**	**3**	**175**	**70**
	hG6PC Q139R/Q247R/S298C_CoV2 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUACUUCACG ACUUCGGCAUCCAG AGCACCCACUACUU GCAGGUGAACUACC AGGACUCCCAGGAU UGGUUCAUUCUCGU GAGCGUGAUUGCCG AUCUCCGGAACGCU UUCUACGUGCUGUU CCCUAUUUGGUUCC ACCUGCAGGAGGCU GUGGGCAUCAAGUU GCUGUGGGUCGCCG UGAUAGGCGACUGG CUCAACUUGGUGUU CAAGUGGAUUCUGU UCGGCCAGCGUCCA UACUGGUGGGUGCU GGACACAGACUAUU AUUCUAAUACAAGC GUACCUCUGAUCAA GCAGUUCCCAGUGA CCUGCGAAACGGGA CCUGGAUCACCAAG CGGCCACGCCAUGG GAACGGCUGGCGUG UACUACGUGAUGGU GACCAGCACCCUCA GCAUCUUCCGGGGA AAGAUCAAGCCGAC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 70 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:27, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UUACAGAUUCCGUU GCCUGAACGUGAUC CUGUGGCUCGGAUU CUGGGCCGUGCAGC UCAACGUGUGCCUG AGUAGAAUCUACCU GGCCGCUCACUUCC CUCACCAGGUAGUG GCCGGCGUGCUGAG CGGCAUUGCCGUGG CCGAGACAUUCAGU CACAUCCACAGCAU CUAUAACGCUAGCC UGAAGAAGUAUUUC CUGAUUACAUUCUU CCUGUUCUCUUUCG CUAUCGGCUUCUAC CUCCUCCUGAAGGG CCUGGGCGUGGACC UGUUGUGGACCCUG GAGAAGGCUCAGAG GUGGUGCGAGAGGC CUGAGUGGGUCCAU AUCGAUACCACCCC AUUCGCAAGUCUGC UGAAGAACCUGGGC ACACUCUUCGGCCU GGGCCUGGCCCUGA ACAGCUCGAUGUAC AGAGAGUCUUGCAA GGGAAAGCUUAGCA AGUGGCUCCCUUUC AGACUGUCUUGCAU CGUGGCCAGCCUGG UGCUGCUGCAUGUG UUCGACUCUCUGAA GCCUCCUUCACAGG UCGAGCUGGUUUUC UAUGUGCUGAGUUU CUGCAAGUCCGCUG UGGUGCCUCUGGCU AGCGUGAGCGUGAU CCCUUACUGCCUGG CCCAGGUGCUGGGC CAGCCUCACAAGAA GUCCCUA			
SEQ ID NO:		**209**	**28**	**3**	**175**	**71**
	hG6PC M8_CoV2	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF	AUGGAGGAGGGCAU GAACGUACUUCACG ACUUCGGCAUCCAG	GGGAAA UAAGAG AGAAAA	UGAUAA UAGGCU GGAGCC	SEQ ID NO: 71 consists

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	AGCACCCACUACUU GCAGGUGAACUACC AGGACUCCCAGGAU UGGUUCAUUCUCGU GAGCGUGAUUGCCG AUCUCCGGAACGCU UUCUACGUGCUGUU CCCUAUUUGGUUCC ACCUGCAGGAGGCU GUGGGCAUCAAGUU GCUGUGGGUCGCCG UGAUAGGCGACUGG CUCAACUUGGUGUU CAAGUGGAUUCUGU UCGGCCAGCGUCCA UACUGGUGGGUGCU GGACACAGACUAUU AUUCUAAUACAAGC GUACCUCUGAUCAA GCAGUUCCCAGUGA CCUGCGAAACGGGA CCUGGAUCACCAAG CGGCCACGCCAUGG GAACGGCUGGCGUG UACUACGUGAUGGU GACCAGCACCCUCA GCAUCUUCCGGGGA AAGAAGAAGCCGAC UUACAGAUUCCGUU GCCUGAACGUGAUC CUGUGGCUCGGAUU CUGGGCCGUGCAGC UCAACGUGUGCCUG AGUAGAAUCUACCU GGCCGCUCACUUCC CUCACCAGGUAGUG GCCGGCGUGCUGAG CGGCAUUGCCGUGG CCGAGACAUUCAGU CACAUCCACAGCAU CUAUAACGCUAGCC UGAAGAAGUAUUUC CUGAUUACAUUCUU CCUGUUCUCUUUCG CUAUCGGCUUCUAC CUCCUCCUGAAGGG CCUGGGCGUGGACC UGUUGUGGACCCUG GAGAAGGCUCAGAG GUGGUGCGAGAGGC CUGAGUGGGUCCAU AUCGAUACCACCCC AUUCGCAAGUCUGC	GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:28, and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UGAAGAACCUGGGC ACACUCUUCGGCCU GGGCCUGGCCCUGA ACAGCUCGAUGUAC AGAGAGUCUUGCAA GGGAAAGCUUAGCA AGUGGCUCCCUUUC AGACUGGGAUGCAU CGUGGCCAGCCUGG UGCUGCUGCAUCUG UUCGACUCUCUGAA GCCUCCUUCACAGG UCGAGCUGAUUUUC UAUGUGCUGAGUUU CUGCAAGUCCGCUG UGGUGCCUCUGGCU AGCGUGAGCCUGAU CCCUUACUGCCUGG CCCAGGUGCUGGGC CAGCCUCACAAGAA GUCCCUA			
SEQ ID NO:		**83**	**29**	**3**	**150**	**72**
	hG6PC V1_Co-1 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	AUGGAGGAAGGAA UGAACGUGCUUCAC GAUUUCGGCAUUCA AAGCACCCAUUACC UCCAGGUGAACUAC CAGGACAGCCAGGA CUGGUUCAUUUUGG UCAGCGUAAUUGCC GAUCUCAGGAACGC AUUCUACGUGCUGU UCCCUAUCUGGUUC CACCUGCAGGAAGC CGUGGGCAUUAAGC UGCUUUGGGUGGCC GUGAUCGGCGACUG GCUGAACCUGGUGU UCAAGUGGAUCCUG UUCGGCCAGCGGCC AUACUGGUGGGUGC UGGAUACCGACUAC UACAGCAACACUUC UGUCCCGCUGAUAA AGCAGUUCCCGGUA ACGUGCGAAACGGG CCCUGGCAGCCCUA GCGGACACGCCAUG GGCACCGCCGGCGU GUACUACGUGAUGG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 72 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:29, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UGACUUCGACCCUC AGCAUUUUCCAGGG AAAGAUCAAGCCAA CCUACAGAUUCAGG UGUCUCAAUGUGAU UCUGUGGCUGGGCU UCUGGGCUGUGCAG UUGAACGUGUGCCU CAGCAGAAUUUACC UGGCCGCCCACUUC CCUCACCAGGUGGU GGCCGGUGUGCUGA GCGGCAUUGCUGUG GCCGAGACGUUCUC GCAUAUCCACAGCA UCUACAACGCCAGC UUAAAGAAGUACUU CCUGAUCACCUUCU UCCUCUUCUCCUUC GCUAUCGGAUUCUA UCUGCUGCUGAAGG GACUGGGCGUCGAC CUGCUGUGGACGCU GGAGAAGGCCCAGC GGUGGUGCGAGCAG CCAGAGUGGGUGCA UAUCGAUACUACCC CAUUCGCCUCUCUG UUAAAGAACCUGGG CACACUGUUCGGUC UGGGCCUGGCCCUC AACUCCAGCAUGUA CCGGGAGAGCUGCA AGGGAAAGCUGUCC AAGUGGCUGCCUUU CCGCCUGGGCUGCA UCGUGGCUAGCCUU GUCCUGCUGCAUCU CUUCGACAGCCUCA AGCCUCCAAGCCAG GUGGAACUGAUUUU CUACGUGCUGAGCU UCUGCAAGUCAGCU GUGGUGCCACUGGC UAGCGUGUCACUGA UUCCUUACUGCCUC GCCCAGGUCUUAGG CCAGCCUCAUAAGA AGUCGCUG			
SEQ ID NO:		**83**	**30**	**3**	**150**	**73**

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	hG6PC V1_Co-2 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	AUGGAGGAGGGAA UGAACGUCUUACAC GACUUCGGCAUCCA GUCCACCCACUACC UACAGGUGAAUUAC CAGGACAGCCAAGA UUGGUUCAUCCUCG UGAGCGUUAUAGCC GACUUGAGAAACGC CUUCUACGUGUUGU UCCCAAUCUGGUUC CACCUCCAGGAGGC AGUUGGAAUUAAGC UGCUGUGGGUGGCA GUGAUCGGCGACUG GCUGAACCUGGUCU UCAAGUGGAUUUUG UUCGGCCAGAGGCC UUACUGGUGGGUGC UGGAUACAGACUAC UACUCCAACACAAG CGUGCCGCUGAUUA AGCAGUUCCCUGUG ACCUGUGAGACAGG CCCUGGCUCCCCAA GCGGCCACGCCAUG GGCACAGCCGGUGU GUACUACGUUAUGG UGACUAGCACGCUG UCGAUUUUCCAGGG CAAGAUCAAGCCAA CCUACAGAUUCAGG UGCCUGAAUGUGAU CCUGUGGCUCGGCU UCUGGGCCGUCCAG CUGAACGUGUGCCU GAGCCGCAUCUAUC UGGCGGCUCACUUC CCACACCAAGUGGU UGCCGGAGUGCUCA GCGGCAUCGCAGUC GCCGAAACCUUCAG CCACAUUCACAGCA UUUAUAAUGCUAGC CUGAAGAAGUAUUU CCUGAUCACCUUCU UCCUCUUCAGCUUC GCCAUUGGCUUCUA UCUGCUGCUCAAGG GCCUGGGCGUGGAC CUUCUGUGGACACU GGAGAAGGCUCAGC GCUGGUGCGAACAG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 73 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:30, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CCAGAGUGGGUGCA CAUCGACACCACCC CGUUCGCCUCCCUG CUGAAGAAUCUGGG CACCCUGUUCGGCC UCGGCCUGGCCCUG AACAGUAGCAUGUA CAGGGAAAGCUGUA AGGGCAAGCUGAGC AAGUGGCUCCCAUU CCGGCUGGGCUGCA UCGUUGCCAGCCUG GUGCUGCUGCACCU UUUCGACUCCCUGA AGCCACCUUCACAG GUGGAGCUCAUCUU CUACGUGCUCUCCU UCUGCAAGUCCGCU GUGGUGCCGCUGGC GAGUGUGAGCCUGA UCCCAUACUGCCUU GCCCAGGUGCUGGG UCAGCCUCAUAAGA AGUCCUUG			
SEQ ID NO:		**83**	**31**	**3**	**150**	**74**
	hG6PC V1-Co-3 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 74 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:31, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACAAC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA AGUGGCUCCCUUUC AGACUUGGAUGCAU CGUGGCGUCUCUCG UGCUGCUGCACCUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGAUUUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCCUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
SEQ ID NO:		83	32	3	150	75
	hG6PC V1_Co-4 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUGCUCCACG ACUUCGGCAUCCAG AGCACCCACUACCU CCAGGUGAACUACC AGGACAGCCAGGAC UGGUUCAUCCUCGU GAGCGUGAUCGCCG ACUUGCGGAACGCC UUCUACGUGUUGUU CCCCAUCUGGUUCC ACUUGCAGGAGGCC GUGGGCAUCAAGCU GCUGUGGGUGGCCG UGAUCGGCGACUGG CUGAACCUGGUGUU CAAGUGGAUCCUGU UCGGCCAGCGGCCC UACUGGUGGGUGCU GGACACCGACUACU ACAGCAACACCAGC GUGCCCCUGAUCAA GCAGUUCCCCGUGA CCUGCGAGACGGGC CCCGGCAGCCCUUC UGGUCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACCAGCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCCAC CUACCGGUUCCGGU GCCUGAACGUGAUC CUGUGGCUGGGCUU CUGGGCCGUGCAGC UGAAUGUGUGCCUG AGCCGGAUCUACCU GGCCGCCCACUUCC CACACCAGGUGGUG GCCGGCGUCCUGAG CGGCAUCGCCGUGG CCGAGACGUUCAGC CACAUCCACAGCAU CUACAACGCCAGCC UGAAGAAGUACUUC CUGAUCACCUUCUU CCUGUUCAGCUUCG CCAUCGGCUUCUAC CUGCUGCUGAAGGG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 75 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:32, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CCUGGGCGUGGACC UUCUGUGGACCCUG GAGAAGGCCCAGCG GUGGUGCGAGCAGC CCGAGUGGGUGCAC AUCGACACCACCCC AUUCGCAAGCCUGC UAAAGAACCUGGGC ACACUCUUCGGCCU GGGACUGGCCCUGA ACAGCAGCAUGUAC CGGGAGAGCUGCAA GGGCAAGCUGAGCA AGUGGCUGCCCUUC AGGCUUGGCUGCAU CGUGGCUAGCCUGG UGCUGCUGCACCUG UUCGACUCCCUGAA GCCUCCCAGCCAGG UGGAGCUGAUCUUC UAUGUCCUCAGCUU CUGCAAGAGCGCCG UGGUGCCACUCGCC AGCGUGAGCCUGAU CCCCUACUGCCUGG CCCAGGUGCUGGGC CAGCCCCACAAGAA GUCCCUG			
SEQ ID NO:		**83**	**33**	**3**	**150**	**76**
	hG6PC V1_Co-5 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLGC IVASLVLLHLFDSLKP	AUGGAGGAGGGCAU GAACGUGCUCCACG ACUUCGGCAUCCAG AGCACCCACUACCU CCAGGUGAACUACC AGGACAGCCAGGAC UGGUUCAUCUUGGU GAGCGUGAUCGCCG ACUUGCGGAACGCC UUCUACGUGCUGUU CCCCAUCUGGUUCC ACCUGCAAGAGGCC GUGGGCAUCAAGCU GCUGUGGGUGGCCG UGAUCGGCGACUGG CUGAACCUGGUGUU CAAGUGGAUCCUGU UCGGCCAGCGGCCC UACUGGUGGGUGCU GGACACCGACUACU ACAGCAACACCAGC	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 76 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:33, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		PSQVELIFYVLSFCKS AVVPLASVSLIPYCL AQVLGQPHKKSL	GUGCCCCUGAUCAA GCAGUUCCCCGUGA CCUGCGAGACUGGC CCCGGCAGCCCUAG CGGACACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACCAGCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCCAC CUACCGGUUCCGGU GCCUGAACGUGAUC CUGUGGCUGGGCUU CUGGGCCGUGCAGC UCAAUGUCUGCCUG AGCCGGAUCUACCU GGCCGCCCACUUCC CUCACCAGGUGGUG GCCGGAGUUCUGAG CGGCAUCGCCGUGG CCGAGACUUUCAGC CACAUCCACAGCAU CUACAACGCCAGCC UGAAGAAGUACUUC CUGAUCACCUUCUU CCUGUUCAGCUUCG CCAUCGGCUUCUAC CUGCUGCUGAAGGG CCUGGGCGUGGACC UUCUGUGGACCCUG GAGAAGGCCCAGCG GUGGUGCGAGCAGC CCGAGUGGGUGCAC AUCGACACCACACC UUUCGCCUCACUGC UCAAGAACCUGGGC ACGCUGUUCGGACU GGGCCUGGCCCUGA ACAGCAGCAUGUAC CGGGAGAGCUGCAA GGGCAAGCUGAGCA AGUGGCUGCCCUUC AGACUGGGAUGCAU CGUGGCGAGCUUGG UGCUGCUGCACCUG UUCGAUUCCCUGAA GCCACCCAGCCAGG UGGAGCUGAUCUUC UAUGUCCUCAGCUU CUGUAAGAGCGCCG UGGUGCCGCUGGCC AGCGUGAGCCUGAU CCCCUACUGUCUGG			

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CCCAGGUGCUGGGC CAGCCCCACAAGAA GUCCCUG			
SEQ ID NO:		**84**	**34**	**3**	**150**	77
	hG6PC V2_Co-1 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUACUUCACG ACUUCGGCAUCCAG AGCACCCACUACUU GCAGGUGAACUACC AGGACUCCCAGGAU UGGUUCAUUCUCGU GAGCGUGAUUGCCG AUCUCCGGAACGCU UUCUACGUGCUGUU CCCUAUUUGGUUCC ACCUGCAGGAGGCU GUGGGCAUCAAGUU GCUGUGGGUCGCCG UGAUAGGCGACUGG CUCAACUUGGUGUU CAAGUGGAUUCUGU UCGGCCAGCGUCCA UACUGGUGGGUGCU GGACACAGACUAUU AUUCUAAUACAAGC GUACCUCUGAUCAA GCAGUUCCCAGUGA CCUGCGAAACGGGA CCUGGAUCACCAAG CGGCCACGCCAUGG GAACGGCUGGCGUG UACUACGUGAUGGU GACCAGCACCCUCA GCAUCUUCCGGGGA AAGAAGAAGCCGAC UUACAGAUUCCGUU GCCUGAACGUGAUC CUGUGGCUCGGAUU CUGGGCCGUGCAGC UCAACGUGUGCCUG AGUAGAAUCUACCU GGCCGCUCACUUCC CUCACCAGGUAGUG GCCGGCGUGCUGAG CGGCAUUGCCGUGG CCGAGACAUUCAGU CACAUCCACAGCAU CUAUAACGCUAGCC UGAAGAAGUAUUUC CUGAUUACAUUCUU	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 77 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:34, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CCUGUUCUCUUUCG CUAUCGGCUUCUAC CUCCUCCUGAAGGG CCUGGGCGUGGACC UGUUGUGGACCCUG GAGAAGGCUCAGAG GUGGUGCGAGAGGC CUGAGUGGGUCCAU AUCGAUACCACCCC AUUCGCAAGUCUGC UGAAGAACCUGGGC ACACUCUUCGGCCU GGGCCUGGCCCUGA ACAGCUCGAUGUAC AGAGAGUCUUGCAA GGGAAAGCUUAGCA AGUGGCUCCCUUUC AGACUGUCUUGCAU CGUGGCCAGCCUGG UGCUGCUGCAUCUG UUCGACUCUCUGAA GCCUCCUUCACAGG UCGAGCUGGUUUUC UAUGUGCUGAGUUU CUGCAAGUCCGCUG UGGUGCCUCUGGCU AGCGUGAGCGUGAU CCCUUACUGCCUGG CCCAGGUGCUGGGC CAGCCUCACAAGAA GUCCCUA			
SEQ ID NO:		**84**	**35**	**3**	**150**	**78**
	hG6PC V2_Co-2G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL	AUGGAGGAGGGCAU GAACGUCCUUCACG ACUUCGGUAUCCAG UCGACCCACUACCU CCAAGUGAACUAUC AGGAUAGCCAGGAC UGGUUCAUUUUGGU GAGCGUAAUAGCCG ACCUCAGAAACGCC UUCUACGUGUUAUU CCCUAUCUGGUUCC ACCUGCAGGAGGCC GUGGGAAUAAAGCU UCUUUGGGUGGCUG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUCCUGU UCGGCCAGAGGCCA	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU	SEQ ID NO: 78 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:35, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	UACUGGUGGGUCCU GGAUACCGACUACU ACAGCAACACAAGC GUGCCACUGAUUAA GCAGUUCCCGGUGA CAUGCGAAACAGGC CCUGGCUCCCCUAG CGGCCAUGCUAUGG GCACCGCCGGCGUG UAUUACGUGAUGGU GACCAGCACCUUGU CUAUCUUCCGAGGC AAGAAGAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUGAUC CUCUGGCUGGGCUU CUGGGCCGUGCAGC UAAACGUUUGCCUC UCCAGAAUCUACCU CGCCGCCCAUUUCC CACACCAGGUGGUG GCCGGCGUUCUGAG UGGCAUCGCCGUGG CUGAGACAUUCAGC CACAUCCAUAGCAU UUACAACGCCAGCU UGAAGAAGUAUUUC CUGAUUACCUUCUU CCUGUUCAGCUUCG CCAUAGGCUUCUAU CUUCUUCUGAAGGG CCUGGGCGUGGACC UGCUGUGGACCCUG GAGAAGGCCCAGCG GUGGUGCGAGAGGC CUGAAUGGGUGCAU AUCGAUACAACCCC UUUCGCCUCCCUAC UCAAGAACCUCGGC ACUCUGUUCGGCCU CGGCCUCGCCCUGA ACAGCUCCAUGUAC AGAGAGAGCUGCAA GGGCAAGCUCAGCA AGUGGCUGCCAUUC CGGCUGAGCUGCAU UGUUGCCAGCCUCG UGCUCCUGCACCUG UUCGACAGUCUCAA GCCUCCUAGUCAGG UGGAGCUGGUGUUC UAUGUCCUGUCAUU CUGCAAGUCCGCCG		GAGUGG GCGGC	

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UGGUGCCUCUGGCU AGCGUGAGCGUCAU CCCAUACUGCCUGG CCCAAGUGCUGGGA CAGCCACACAAGAA GUCGUUA			
SEQ ID NO:		**84**	**36**	**3**	**150**	**79**
	hG6Pc V2_Co-3 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAGGGUA UGAACGUCCUACAC GACUUCGGCAUACA GUCUACCCACUAUC UUCAGGUUAACUAC CAGGACAGCCAGGA CUGGUUCAUCCUUG UGAGCGUUAUCGCC GACCUUCGCAACGC GUUCUACGUGCUUU UCCCUAUCUGGUUC CACCUGCAGGAGGC UGUAGGCAUUAAGC UGCUGUGGGUCGCC GUGAUCGGCGACUG GCUGAAUCUGGUGU UCAAGUGGAUCCUG UUCGGCCAACGCCC UUAUUGGUGGGUCC UGGACACAGAUUAC UAUUCAAAUACCAG CGUGCCAUUGAUUA AGCAGUUCCCUGUU ACCUGCGAAACCGG CCCUGGCUCCCCUU CCGGCCAUGCAAUG GGAACCGCCGGCGU GUACUAUGUCAUGG UGACCAGCACCCUU UCAAUUUUCCGGGG CAAGAAGAAGCCUA CUUACAGGUUCAGA UGCCUGAACGUGAU CCUGUGGCUGGGCU UCUGGGCCGUGCAG CUCAACGUGUGCCU GAGCAGAAUCUAUC UCGCCGCUCACUUC CCUCAUCAGGUGGU CGCCGGAGUGCUCU CAGGCAUCGCCGUG GCGGAGACGUUCUC CCACAUACACAGCA	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 79 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:36, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UUUACAACGCAAGC CUGAAGAAGUACUU CCUGAUCACCUUCU UCCUGUUCAGCUUC GCCAUCGGAUUCUA CCUGCUGCUGAAGG GCCUCGGCGUUGAC CUGCUGUGGACCCU GGAGAAGGCCCAGA GAUGGUGUGAGCGU CCUGAGUGGGUGCA CAUUGACACCACUC CGUUCGCUAGCCUG CUGAAGAACCUGGG CACCCUGUUCGGCC UGGGACUGGCUCUG AACAGCAGCAUGUA UAGGGAGUCCUGCA AGGGCAAGCUAUCA AAGUGGCUGCCUUU CCGGCUGAGCUGCA UUGUGGCCAGCCUG GUGCUGCUGCACCU GUUCGACUCCCUGA AGCCGCCUUCCCAG GUGGAGCUGGUCUU CUACGUGCUUAGCU UCUGUAAGAGCGCC GUGGUGCCACUCGC UAGCGUGUCUGUUA UCCCUUACUGCCUG GCCCAGGUGCUGGG CCAGCCUCAUAAGA AGUCUCUC			
SEQ ID NO:		**84**	**37**	**3**	**150**	**80**
	hG6PC V2_Co-4 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY	AUGGAGGAGGGCAU GAACGUGCUUCACG AUUUCGGAAUUCAG AGCACCCACUACUU GCAGGUGAACUACC AGGACUCUCAAGAC UGGUUCAUCCUUGU GAGCGUGAUCGCCG AUCUCAGGAACGCC UUCUACGUGUUGUU CCCGAUUUGGUUCC ACCUUCAGGAGGCC GUCGGCAUCAAGCU GCUGUGGGUGGCCG UGAUCGGCGACUGG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU	SEQ ID NO: 80 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:37, and 3' UTR of

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	CUGAACUUGGUGUU CAAGUGGAUUCUGU UCGGACAGAGGCCA UACUGGUGGGUGCU GGACACUGACUAUU AUAGUAACACUAGC GUCCCACUGAUAAA GCAAUUCCCUGUGA CCUGCGAGACAGGC CCUGGCAGCCCUUC CGGCCACGCCAUGG GCACUGCCGGAGUA UACUAUGUGAUGGU GACCAGCACCCUGA GCAUCUUCAGGGGA AAGAAGAAGCCUAC GUACAGAUUCAGAU GCCUGAACGUGAUC CUGUGGCUGGGCUU CUGGGCCGUGCAAC UGAAUGUGUGCCUC UCCAGGAUUUACCU GGCAGCGCACUUCC CUCACCAGGUCGUG GCUGGAGUGCUGUC UGGAAUCGCCGUGG CCGAAACAUUCAGC CACAUUCACUCAAU UUACAAUGCCAGCC UGAAGAAGUACUUC CUGAUCACCUUCUU CCUGUUCAGUUUCG CCAUAGGUUUCUAC CUGCUGCUCAAGGG CCUAGGCGUUGACC UGCUGUGGACAUUG GAGAAGGCACAGCG CUGGUGCGAGAGGC CAGAAUGGGUGCAC AUUGACACAACCCC UUUCGCCAGCCUCC UGAAGAAUCUCGGC ACCCUGUUCGGCCU GGGCCUGGCCCUGA ACAGCUCAAUGUAU AGAGAGAGCUGCAA GGGUAAGCUGAGCA AGUGGCUGCCUUUC AGGCUGAGCUGCAU CGUUGCCAGCCUCG UACUGCUCCACCUC UUCGACUCUCUGAA GCCACCAAGUCAGG		GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UGGAGCUGGUGUUC UAUGUGCUUAGCUU CUGCAAGAGCGCCG UGGUGCCUCUGGCC AGCGUGUCCGUGAU CCCUUACUGCCUGG CCCAGGUCCUGGGC CAGCCGCAUAAGAA GUCUCUG			
SEQ ID NO:		**84**	**38**	**3**	**150**	**81**
	hG6PC V2-Co-5 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFRGKKKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCERPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHLFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUGUUACACG ACUUCGGCAUCCAG AGCACCCACUACCU CCAGGUGAACUACC AGGACAGCCAGGAC UGGUUCAUCCUAGU GAGCGUGAUCGCCG ACUUACGGAACGCC UUCUACGUGCUUUU CCCCAUCUGGUUCC ACCUUCAGGAGGCC GUGGGCAUCAAGCU GCUGUGGGUGGCCG UGAUCGGCGACUGG CUGAACCUGGUGUU CAAGUGGAUCCUGU UCGGCCAGCGGCCC UACUGGUGGGUGCU GGACACCGACUACU ACAGCAACACCAGC GUGCCCCUGAUCAA GCAGUUCCCCGUGA CCUGCGAGACAGGC CCCGGCAGCCCGAG CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACCAGCACCCUGA GCAUCUUCCGGGGC AAGAAGAAGCCCAC CUACCGGUUCCGGU GCCUGAACGUGAUC CUGUGGCUGGGCUU CUGGGCCGUGCAGC UUAACGUCUGCCUG AGCCGGAUCUACCU GGCCGCCCACUUCC CUCACCAGGUGGUG GCUGGAGUGCUGAG	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCAUG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	SEQ ID NO: 81 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF Sequence of SEQ ID NO:38, and 3' UTR of SEQ ID NO:150

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CGGCAUCGCCGUGG CCGAGACAUUCAGC CACAUCCACAGCAU CUACAACGCCAGCC UGAAGAAGUACUUC CUGAUCACCUUCUU CCUGUUCAGCUUCG CCAUCGGCUUCUAC CUGCUGCUGAAGGG CCUGGGCGUGGACC UGCUCUGGACCCUG GAGAAGGCCCAGCG GUGGUGCGAAAGAC CCGAGUGGGUGCAC AUCGACACCACCCC UUUCGCCUCUCUGC UCAAGAACCUGGGC ACGCUUUUCGGUCU GGGCCUGGCCCUGA ACAGCAGCAUGUAC CGGGAGAGCUGCAA GGGCAAGCUGAGCA AGUGGCUGCCCUUC CGGCUGAGCUGCAU CGUGGCUAGCCUGG UGCUGCUGCACCUG UUCGAUUCCCUGAA GCCUCCCAGCCAGG UGGAGCUCGUGUUC UAUGUGCUGUCAUU CUGCAAGAGCGCCG UGGUGCCACUGGCC AGCGUGUCAGUGAU CCCCUACUGUCUGG CACAGGUGCUGGGC CAGCCCCACAAGAA GUCUCUG			
SEQ ID NO:		**194**	**23**	**210**	**175**	**214**
	G6PC_S298C_A start G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU	AGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGGCU GGAGCC UCGGUG GCCUAG CUUCUU GCCCCU UGGGCC UCCCCC CAGCCC CUCCUC CCCUUC	SEQ ID NO:214 consists from 5' to 3' end: 5' UTR of SEQ ID NO:210, ORF Sequence of SEQ ID NO:23,

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
		FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACAAC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA AGUGGCUCCCUUUC AGACUUAGCUGCAU CGUGGCGUCUCUCG		CUGCAC CCGUAC CCCCGU GGUCUU UGAAUA AAGUCU GAGUGG GCGGC	and 3' UTR of SEQ ID NO:175

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			UGCUGCUGCACGUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGGUGUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCGUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			
SEQ ID NO:		**194**	**23**	**210**	**178**	**215**
	G6PC_S298C_A start_3XmiR142 G5 Cap: C1 PolyA tail: 100nt (SEQ ID NO:223)	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU	AGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGUCC AUAAAG UAGGAA ACACUA CAGCUG GAGCCU CGGUGG CCUAGC UUCUUG CCCCUU GGGCCU CCAUAA AGUAGG AAACAC UACAUC CCCCCA GCCCCU CCUCCC CUUCCU GCACCC GUACCC CCUCCA UAAAGU AGGAAA CACUAC AGUGGU CUUUGA AUAAAG UCUGAG UGGGCG GC	SEQ ID NO:215 consists from 5' to 3' end: 5' UTR of SEQ ID NO:210, ORF Sequence of SEQ ID NO:23, and 3' UTR of SEQ ID NO:178

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACAAC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA AGUGGCUCCCUUUC AGACUUAGCUGCAU CGUGGCGUCUCUCG UGCUGCUGCACGUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGGUGUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCGUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			
SEQ ID NO:		**194**	**23**	**3**	**178**	**217**
	G6PC_S298C_G start_3XmiR142 G5 Cap: C1	MEEGMNVLHDFGIQ STHYLQVNYQDSQD WFILVSVIADLRNAF YVLFPIWFHLQEAVG IKLLWVAVIGDWLNL VFKWILFGQRPYWW VLDTDYYSNTSVPLI KQFPVTCETGPGSPS GHAMGTAGVYYVM	AUGGAGGAGGGCAU GAACGUGCUUCACG ACUUCGGCAUCCAG UCAACCCAUUACCU ACAGGUGAACUAUC AGGACUCCCAGGAU UGGUUCAUCCUCGU GAGCGUGAUCGCCG AUCUAAGAAACGCA	GGGAAA UAAGAG AGAAAA GAAGAG UAAGAA GAAAUA UAAGAG CCACC	UGAUAA UAGUCC AUAAAG UAGGAA ACACUA CAGCUG GAGCCU CGGUGG CCUAGC	SEQ ID NO:217 consists from 5' to 3' end: 5' UTR of SEQ ID NO:3, ORF

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
	PolyA tail: 100nt (SEQ ID NO:223)	VTSTLSIFQGKIKPTY RFRCLNVILWLGFWA VQLNVCLSRIYLAAH FPHQVVAGVLSGIAV AETFSHIHSIYNASLK KYFLITFFLFSFAIGFY LLLKGLGVDLLWTLE KAQRWCEQPEWVHI DTTPFASLLKNLGTL FGLGLALNSSMYRES CKGKLSKWLPFRLSC IVASLVLLHVFDSLKP PSQVELVFYVLSFCK SAVVPLASVSVIPYCL AQVLGQPHKKSL	UUCUACGUGUUGUU CCCAAUCUGGUUCC ACCUGCAGGAAGCU GUGGGCAUCAAGCU CCUUUGGGUGGCCG UGAUUGGCGAUUGG CUGAACCUGGUGUU CAAGUGGAUUCUCU UCGGUCAGCGGCCU UACUGGUGGGUGUU GGAUACCGACUAUU ACAGUAACACCUCU GUUCCGCUGAUCAA GCAGUUCCCUGUCA CCUGCGAAACCGGA CCUGGCUCCCCUUC CGGCCACGCCAUGG GCACCGCCGGCGUG UACUACGUGAUGGU GACAUCCACCCUGA GCAUCUUCCAGGGC AAGAUCAAGCCUAC CUACAGAUUCAGGU GCCUGAACGUUAUC UUGUGGCUGGGAUU CUGGGCAGUGCAGC UCAACGUGUGCCUG AGCAGGAUCUAUCU CGCAGCCCACUUCC CACAUCAGGUCGUG GCCGGCGUACUGAG CGGAAUCGCCGUGG CUGAAACCUUCAGC CAUAUCCACAGCAU CUACAAUGCCUCCC UGAAGAAGUACUUC CUUAUUACAUUCUU CCUGUUCUCUUUCG CCAUCGGCUUCUAU CUGCUGCUAAAGGG CCUGGGCGUCGACC UCCUGUGGACCCUG GAGAAGGCCCAGAG AUGGUGCGAACAAC CUGAGUGGGUGCAC AUCGACACCACUCC GUUCGCUAGCCUGC UCAAGAACCUGGGC ACUCUUUUCGGCCU GGGCCUGGCCCUCA AUAGCAGCAUGUAC CGGGAGAGCUGUAA GGGAAAGCUGUCCA		UUCUUG CCCCUU GGGCCU CCAUAA AGUAGG AAACAC UACAUC CCCCCA GCCCCU CCUCCC CUUCCU GCACCC GUACCC CCUCCA UAAAGU AGGAAA CACUAC AGUGGU CUUUGA AUAAAG UCUGAG UGGGCG GC	Sequence of SEQ ID NO:23, and 3' UTR of SEQ ID NO:178

	mRNA Name	ORF Sequence (Amino Acid)	ORF Sequence (Nucleotide)	5' UTR Sequence	3' UTR Sequence	Construct Sequence
			AGUGGCUCCCUUUC AGACUUAGCUGCAU CGUGGCGUCUCUCG UGCUGCUGCACGUG UUCGACAGCCUGAA GCCUCCUUCGCAGG UGGAGCUGGUGUUC UACGUGCUGAGCUU CUGCAAGAGCGCGG UGGUGCCACUGGCA AGUGUCAGCGUGAU CCCUUAUUGCCUGG CCCAGGUGCUGGGC CAGCCUCAUAAGAA GUCCCUG			

EXAMPLES

EXAMPLE 1

Chimeric Polynucleotide Synthesis

A. Triphosphate route

[1006] Two regions or parts of a chimeric polynucleotide can be joined or ligated using triphosphate chemistry. According to this method, a first region or part of 100 nucleotides or less can be chemically synthesized with a 5' monophosphate and terminal 3'desOH or blocked OH. If the region is longer than 80 nucleotides, it can be synthesized as two strands for ligation.

[1007] If the first region or part is synthesized as a non-positionally modified region or part using *in vitro* transcription (IVT), conversion the 5'monophosphate with subsequent capping of the 3' terminus can follow. Monophosphate protecting groups can be selected from any of those known in the art.

[1008] The second region or part of the chimeric polynucleotide can be synthesized using either chemical synthesis or IVT methods. IVT methods can include an RNA polymerase that can utilize a primer with a modified cap. Alternatively, a cap of up to 80 nucleotides can be chemically synthesized and coupled to the IVT region or part.

[1009] It is noted that for ligation methods, ligation with DNA T4 ligase, followed by treatment with DNAse should readily avoid concatenation.

[1010] The entire chimeric polynucleotide need not be manufactured with a phosphate-sugar backbone. If one of the regions or parts encodes a polypeptide, then such region or part can comprise a phosphate-sugar backbone.

[1011] Ligation can then be performed using any known click chemistry, orthoclick chemistry, solulink, or other bioconjugate chemistries known to those in the art.

B. Synthetic route

[1012] The chimeric polynucleotide can be made using a series of starting segments. Such segments include:

(a) Capped and protected 5' segment comprising a normal 3'OH (SEG. 1)

(b) 5' triphosphate segment which can include the coding region of a polypeptide and comprising a normal 3'OH (SEG. 2)

(c) 5' monophosphate segment for the 3' end of the chimeric polynucleotide (*e.g.*, the tail) comprising cordycepin or no 3'OH (SEG. 3)

[1013] After synthesis (chemical or IVT), segment 3 (SEG. 3) can be treated with cordycepin and then with pyrophosphatase to create the 5'monophosphate.

[1014] Segment 2 (SEG. 2) can then be ligated to SEG. 3 using RNA ligase. The ligated polynucleotide can then be purified and treated with pyrophosphatase to cleave the diphosphate. The treated SEG.2-SEG. 3 construct is then purified and SEG. 1 is ligated to the 5' terminus. A further purification step of the chimeric polynucleotide can be performed.

[1015] Where the chimeric polynucleotide encodes a polypeptide, the ligated or joined segments can be represented as: 5' UTR (SEG. 1), open reading frame or ORF (SEG. 2) and 3' UTR+PolyA (SEG. 3).

[1016] The yields of each step can be as much as 90-95%.

EXAMPLE 2

PCR for cDNA Production

[1017] PCR procedures for the preparation of cDNA can be performed using 2x KAPA HIFI™ HotStart ReadyMix by Kapa Biosystems (Woburn, MA). This system includes 2x KAPA ReadyMix12.5 µl; Forward Primer (10 µM) 0.75 µl; Reverse

Primer (10 µM) 0.75 µl; Template cDNA -100 ng; and dH_20 diluted to 25.0 µl. The PCR reaction conditions can be: at 95° C for 5 min. and 25 cycles of 98° C for 20 sec, then 58° C for 15 sec, then 72° C for 45 sec, then 72° C for 5 min. then 4° C to termination.

[1018] The reverse primer of the instant invention can incorporate a poly-T_{120} (SEQ ID NO:234) for a poly-A_{120} (SEQ ID NO:233) in the mRNA. Other reverse primers with longer or shorter poly(T) tracts can be used to adjust the length of the poly(A) tail in the polynucleotide mRNA.

[1019] The reaction can be cleaned up using Invitrogen's PURELINK™ PCR Micro Kit (Carlsbad, CA) per manufacturer's instructions (up to 5 µg). Larger reactions will require a cleanup using a product with a larger capacity. Following the cleanup, the cDNA can be quantified using the NANODROP™ and analyzed by agarose gel electrophoresis to confirm the cDNA is the expected size. The cDNA can then be submitted for sequencing analysis before proceeding to the *in vitro* transcription reaction.

EXAMPLE 3

In vitro Transcription (IVT)

[1020] The *in vitro* transcription reactions can generate polynucleotides containing uniformly modified polynucleotides. Such uniformly modified polynucleotides can comprise a region or part of the polynucleotides of the invention. The input nucleotide triphosphate (NTP) mix can be made using natural and un-natural NTPs.

[1021] A typical *in vitro* transcription reaction can include the following:

1 Template cDNA—1.0 µg

2 10x transcription buffer (400 mM Tris-HCl pH 8.0, 190 mM $MgCl_2$, 50 mM DTT, 10 mM Spermidine)—2.0 µl

3 Custom NTPs (25mM each) —7.2 µl

4 RNase Inhibitor—20 U

5 T7 RNA polymerase —3000 U

6 dH_20—Up to 20.0 µl. and

7 Incubation at 37° C for 3 hr-5 hours.

[1022] The crude IVT mix can be stored at 4° C overnight for cleanup the next day. 1 U of RNase-free DNase can then be used to digest the original template. After 15

minutes of incubation at 37° C, the mRNA can be purified using Ambion's MEGACLEAR™ Kit (Austin, TX) following the manufacturer's instructions. This kit can purify up to 500 µg of RNA. Following the cleanup, the RNA can be quantified using the NanoDrop and analyzed by agarose gel electrophoresis to confirm the RNA is the proper size and that no degradation of the RNA has occurred.

EXAMPLE 4

Enzymatic Capping

[1023] Capping of a polynucleotide can be performed with a mixture includes: IVT RNA 60 µg-180µg and dH_20 up to 72 µl. The mixture can be incubated at 65° C for 5 minutes to denature RNA, and then can be transferred immediately to ice.

[1024] The protocol can then involve the mixing of 10x Capping Buffer (0.5 M Tris-HCl (pH 8.0), 60 mM KCl, 12.5 mM $MgCl_2$) (10.0 µl); 20 mM GTP (5.0 µl); 20 mM S-Adenosyl Methionine (2.5 µl); RNase Inhibitor (100 U); 2′-O-Methyltransferase (400U); Vaccinia capping enzyme (Guanylyl transferase) (40 U); dH_20 (Up to 28 µl); and incubation at 37° C for 30 minutes for 60 µg RNA or up to 2 hours for 180 µg of RNA.

[1025] The polynucleotide can then be purified using Ambion's MEGACLEAR™ Kit (Austin, TX) following the manufacturer's instructions. Following the cleanup, the RNA can be quantified using the NANODROP™ (ThermoFisher, Waltham, MA) and analyzed by agarose gel electrophoresis to confirm the RNA is the proper size and that no degradation of the RNA has occurred. The RNA product can also be sequenced by running a reverse-transcription-PCR to generate the cDNA for sequencing.

EXAMPLE 5

PolyA Tailing Reaction

[1026] Without a poly-T in the cDNA, a poly-A tailing reaction must be performed before cleaning the final product. This can be done by mixing Capped IVT RNA (100 µl); RNase Inhibitor (20 U); 10x Tailing Buffer (0.5 M Tris-HCl (pH 8.0), 2.5 M NaCl, 100 mM $MgCl_2$) (12.0 µl); 20 mM ATP (6.0 µl); Poly-A Polymerase (20 U); dH_20 up to 123.5 µl and incubating at 37° C for 30 min. If the poly-A tail is already in the transcript, then the tailing reaction can be skipped and proceed directly to cleanup

with Ambion's MEGACLEAR™ kit (Austin, TX) (up to 500 µg). Poly-A Polymerase is, in some cases, a recombinant enzyme expressed in yeast.

[1027] It should be understood that the processivity or integrity of the polyA tailing reaction does not always result in an exact size polyA tail. Hence polyA tails of approximately between 40-200 nucleotides, *e.g.*, about 40, 50, 60, 70, 80, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 150-165, 155, 156, 157, 158, 159, 160, 161, 162, 163, 164 or 165 are within the scope of the invention.

EXAMPLE 6

Natural 5′ Caps and 5′ Cap Analogues

[1028] 5′-capping of polynucleotides can be completed concomitantly during the *in vitro*-transcription reaction using the following chemical RNA cap analogs to generate the 5′-guanosine cap structure according to manufacturer protocols: 3′-O-Me-m7G(5')ppp(5') G [the ARCA cap];G(5')ppp(5')A; G(5')ppp(5')G; m7G(5')ppp(5')A; m7G(5')ppp(5')G (New England BioLabs, Ipswich, MA). 5′-capping of modified RNA can be completed post-transcriptionally using a Vaccinia Virus Capping Enzyme to generate the "Cap 0" structure: m7G(5')ppp(5')G (New England BioLabs, Ipswich, MA). Cap 1 structure can be generated using both Vaccinia Virus Capping Enzyme and a 2′-O methyl-transferase to generate: m7G(5')ppp(5')G-2′-O-methyl. Cap 2 structure can be generated from the Cap 1 structure followed by the 2′-O-methylation of the 5′-antepenultimate nucleotide using a 2′-O methyl-transferase. Cap 3 structure can be generated from the Cap 2 structure followed by the 2′-O-methylation of the 5′-preantepenultimate nucleotide using a 2′-O methyl-transferase. Enzymes can be derived from a recombinant source.

[1029] When transfected into mammalian cells, the modified mRNAs can have a stability of between 12-18 hours or more than 18 hours, *e.g.*, 24, 36, 48, 60, 72 or greater than 72 hours.

EXAMPLE 7

Capping Assays

A. Protein Expression Assay

[1030] Polynucleotides encoding a polypeptide, containing any of the caps taught herein, can be transfected into cells at equal concentrations. After 6, 12, 24 and 36 hours post-transfection, the amount of protein secreted into the culture medium can be assayed by ELISA. Synthetic polynucleotides that secrete higher levels of protein into the medium would correspond to a synthetic polynucleotide with a higher translationally-competent Cap structure.

B. Purity Analysis Synthesis

[1031] Polynucleotides encoding a polypeptide, containing any of the caps taught herein, can be compared for purity using denaturing Agarose-Urea gel electrophoresis or HPLC analysis. Polynucleotides with a single, consolidated band by electrophoresis correspond to the higher purity product compared to polynucleotides with multiple bands or streaking bands. Synthetic polynucleotides with a single HPLC peak would also correspond to a higher purity product. The capping reaction with a higher efficiency would provide a more pure polynucleotide population.

C. Cytokine Analysis

[1032] Polynucleotides encoding a polypeptide, containing any of the caps taught herein, can be transfected into cells at multiple concentrations. After 6, 12, 24 and 36 hours post-transfection the amount of pro-inflammatory cytokines such as TNF-alpha and IFN-beta secreted into the culture medium can be assayed by ELISA. Polynucleotides resulting in the secretion of higher levels of pro-inflammatory cytokines into the medium would correspond to polynucleotides containing an immune-activating cap structure.

D. Capping Reaction Efficiency

[1033] Polynucleotides encoding a polypeptide, containing any of the caps taught herein, can be analyzed for capping reaction efficiency by LC-MS after nuclease treatment. Nuclease treatment of capped polynucleotides would yield a mixture of free nucleotides and the capped 5′-5-triphosphate cap structure detectable by LC-MS.

The amount of capped product on the LC-MS spectra can be expressed as a percent of total polynucleotide from the reaction and would correspond to capping reaction efficiency. The cap structure with higher capping reaction efficiency would have a higher amount of capped product by LC-MS.

EXAMPLE 8

Agarose Gel Electrophoresis of Modified RNA or RT PCR Products

[1034] Individual polynucleotides (200-400 ng in a 20 µl volume) or reverse transcribed PCR products (200-400 ng) can be loaded into a well on a non-denaturing 1.2% Agarose E-Gel (Invitrogen, Carlsbad, CA) and run for 12-15 minutes according to the manufacturer protocol.

EXAMPLE 9

Nanodrop Modified RNA Quantification and UV Spectral Data

[1035] Modified polynucleotides in TE buffer (1 µl) can be used for Nanodrop UV absorbance readings to quantitate the yield of each polynucleotide from a chemical synthesis or *in vitro* transcription reaction.

EXAMPLE 10

Formulation of Modified mRNA Using Lipidoids

[1036] Polynucleotides can be formulated for *in vitro* experiments by mixing the polynucleotides with the lipidoid at a set ratio prior to addition to cells. *In vivo* formulation can require the addition of extra ingredients to facilitate circulation throughout the body. To test the ability of these lipidoids to form particles suitable for *in vivo* work, a standard formulation process used for siRNA-lipidoid formulations can be used as a starting point. After formation of the particle, polynucleotide can be added and allowed to integrate with the complex. The encapsulation efficiency can be determined using a standard dye exclusion assays.

EXAMPLE 11

Method of Screening for Protein Expression

A. Electrospray Ionization

[1037] A biological sample that can contain proteins encoded by a polynucleotide administered to the subject can be prepared and analyzed according to the manufacturer protocol for electrospray ionization (ESI) using 1, 2, 3 or 4 mass analyzers. A biologic sample can also be analyzed using a tandem ESI mass spectrometry system.

[1038] Patterns of protein fragments, or whole proteins, can be compared to known controls for a given protein and identity can be determined by comparison.

B. Matrix-Assisted Laser Desorption/Ionization

[1039] A biological sample that can contain proteins encoded by one or more polynucleotides administered to the subject can be prepared and analyzed according to the manufacturer protocol for matrix-assisted laser desorption/ionization (MALDI).

[1040] Patterns of protein fragments, or whole proteins, can be compared to known controls for a given protein and identity can be determined by comparison.

C. Liquid Chromatography-Mass spectrometry-Mass spectrometry

[1041] A biological sample, which can contain proteins encoded by one or more polynucleotides, can be treated with a trypsin enzyme to digest the proteins contained within. The resulting peptides can be analyzed by liquid chromatography-mass spectrometry-mass spectrometry (LC/MS/MS). The peptides can be fragmented in the mass spectrometer to yield diagnostic patterns that can be matched to protein sequence databases via computer algorithms. The digested sample can be diluted to achieve 1 ng or less starting material for a given protein. Biological samples containing a simple buffer background (*e.g.,* water or volatile salts) are amenable to direct in-solution digest; more complex backgrounds (*e.g.,* detergent, non-volatile salts, glycerol) require an additional clean-up step to facilitate the sample analysis.

[1042] Patterns of protein fragments, or whole proteins, can be compared to known controls for a given protein and identity can be determined by comparison.

EXAMPLE 12

Synthesis of mRNA Encoding G6PC

[1043] Sequence optimized polynucleotides encoding G6PC polypeptides are synthesized and characterized as described in Examples 1 to 11.

[1044] An mRNA encoding human G6PC or a mutated variant thereof can be constructed, e.g., by using the ORF sequence (amino acid) provided in SEQ ID NO: 194, 199 or 205, respectively. The mRNA sequence includes both 5' and 3' UTR regions flanking the ORF sequence (nucleotide). In an exemplary construct, the 5' UTR and 3' UTR sequences are SEQ ID NO: 3 and SEQ ID NO: 150, respectively.

5'UTR:

GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC(SEQ ID NO:3)

3'UTR:

UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUC
CUCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (SEQ
ID NO:150)

In another exemplary construct, the 5' UTR and 3' UTR sequences are SEQ ID NOs.: 3 and 175, respectively (see below):

5'UTR:

GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC(SEQ ID NO:3)

3'UTR:

UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUC
CUCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (SEQ
ID NO:175)

In another exemplary construct, the 5' UTR and 3' UTR sequences are SEQ ID NOs.: 3 and 4, respectively (see below):

5'UTR:

GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC (SEQ ID NO:3)

3'UTR:

UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUC
CUCCCCUUCCUGCACCCGUACCCCCUCCAUAAAGUAGGAAACACUACAGUGGUCUUUGAAUAA
AGUCUGAGUGGGCGGC (SEQ ID NO:4)

In another exemplary construct, the 5' UTR and 3' UTR sequences are SEQ ID NOs.: 3 and 177, respectively (see below):

5'UTR:

GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC(SEQ ID NO:3)

3'UTR:

UGAUAAUAGGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCCCUUGGGCCUCCCCCCAGCCCCUC
CUCCCCUUCCUGCACCCGUACCCCCCGCAUUAUUACUCACGGUACGAGUGGUCUUUGAAUAAA
GUCUGAGUGGGCGGC (SEQ ID NO: 177)

In another exemplary construct, the 5' UTR and 3' UTR sequences are SEQ ID NOs.: 3 and 178, respectively (see below):

5′UTR:

GGGAAAUAAGAGAGAAAAGAAGAGUAAGAAGAAAUAUAAGAGCCACC(SEQ ID NO:3)

3′UTR:

UGAUAAUAGUCCAUAAAGUAGGAAACACUACAGCUGGAGCCUCGGUGGCCUAGCUUCUUGCCC
CUUGGGCCUCCAUAAAGUAGGAAACACUACAUCCCCCCAGCCCCUCCUCCCCUUCCUGCACCC
GUACCCCCUCCAUAAAGUAGGAAACACUACAGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC
(SEQ ID NO:178)

In another exemplary construct, the 5' UTR and 3' UTR sequences are SEQ ID NOs.: 193 and 150, respectively (see below):

5′UTR:

UCAAGCUUUUGGACCCUCGUACAGAAGCUAAUACGACUCACUAUAGGGAAAUAAGAGAGAAAA
GAAGAGUAAGAAGAAAUAUAAGAGCCACC (SEQ ID NO:193)

3′UTR:

UGAUAAUAGGCUGGAGCCUCGGUGGCCAUGCUUCUUGCCCCUUGGGCCUCCCCCCCAGCCCCUC
CUCCCCUUCCUGCACCCGUACCCCCGUGGUCUUUGAAUAAAGUCUGAGUGGGCGGC (SEQ
ID NO:150)

[1045] The G6PC mRNA sequence is prepared as modified mRNA. Specifically, during in vitro transcription, modified mRNA can be generated using N1-methylpseudouridine-5'-triphosphate to ensure that the mRNAs contain 100% N1-methylpseudouridine instead of uridine. Further, G6PC-mRNA can be synthesized with a primer that introduces a polyA-tail, and a Cap 1 structure is generated on both mRNAs using Vaccinia Virus Capping Enzyme and a 2′-O methyl-transferase to generate: m7G(5')ppp(5')G-2′-O-methyl.

EXAMPLE 13

In Vitro Expression of mRNA Encoding G6Pase

[1046] The *in vitro* expression of 1-methyl-pseudouridine modified mRNA constructs encoding human G6PC (hG6PC; SEQ ID NO:48), a FLAG-tagged hG6PC (SEQ ID NO:45), mouse G6PC (mG6PC; SEQ ID NO:46), and a FLAG-tagged mG6PC (SEQ ID NO:47) was measured in a human embryonic kidney cell line. HEK 293 cells (ATCC® CRL1573™), were seeded on 6-well plates (BD Biosciences, San Jose, USA) one day prior to transfection. Modified mRNA formulations encoding each of the human or mouse G6PC sequences, or a GFP control, were transfected using 2 μg

mRNA and 6 μL Lipofectamine MessengerMAX in 250 μL OPTI-MEM per well and incubated.

[1047] After 24 hours, the cells were lysed using a consistent amount of lysis buffer. Protein concentrations of each were determined. To analyze G6PC expression, equal loads of each lysate containing 20 μg of total protein were prepared in a loading buffer and subjected to capillary electrophoresis (CE) analysis. Beta-actin was used as a control. For detection of G6PC, commercial antibodies were used (Atlas HPA052324 anti-G6PC antibody) according to the manufacturer's instructions. The results were compared to G6PC expression levels in mock transfected cells.

[1048] **FIG. 1A** shows that G6PC was expressed from each of the mRNA constructs, but was not detected in cells transfected with eGFP mRNA or in mock transfected cells. **FIG. 1B** shows the relative expression levels of each of the constructs derived from a quantitative analysis of the CE signals.

[1049] Transfection of primary patient fibroblasts with an mRNA encoding human G6PC (SEQ ID NO:48) resulted in G6PC that localized to the endoplasmic reticulum (ER) (data not shown). The G6PC protein co-localized with calnexin (detected with calnexin-specific antibody, Abcam 22595), an ER-specific marker (data not shown).

EXAMPLE 14

In Vitro Activity of mRNA-Encoded G6Pase

[1050] An *in vitro* G6PC activity assay was performed to determine whether G6PC exogenously-expressed after introduction of mRNA comprising a G6PC sequence is active.

A. Expression Assay

[1051] HEK 293 cells were transfected with mRNA formulations comprising human G6PC (hG6PC; SEQ ID NO:48), a FLAG-tagged hG6PC (SEQ ID NO:45), mouse G6PC (mG6PC; SEQ ID NO:46), or a FLAG-tagged mG6PC (SEQ ID NO:47), or a GFP control. Cells were transfected with Lipofectamin 2000 and lysed as described in Example 13 above.

B. Activity Assay

[1052] To assess whether mRNA-expressed G6PC can function, an *in vitro* activity assay was performed using transfected HEK 293 cell lysates as the source of enzymatic activity. G6PC catalyzes the hydrolysis of glucose-6-P into glucose and

inorganic phosphate (Pi). A Taussky-Shorr Color Reagent assay was used to determine the levels of inorganic phosphate in transfected cells, thereby indicating the activity level of G6PC (see Taussky and Shorr, J. Biol. Chem. 1953, 202(2):675-685, herein incorporated by reference in its entirety, and **FIG. 2A**). **FIG. 2B** shows that HEK 293 cells transfected with each of the mRNAs encoding human G6PC (hG6PC; SEQ ID NO:48), a FLAG-tagged hG6PC (SEQ ID NO:45), mouse G6PC (mG6PC; SEQ ID NO:46), or a FLAG-tagged mG6PC (SEQ ID NO:47) expressed G6PC that exhibited G6Pase activity, as exhibited by increases in Pi levels relative to eGFP control or mock transfected cells.

EXAMPLE 15

Production of nanoparticle compositions

A. Production of nanoparticle compositions

[1053] Nanoparticles can be made with mixing processes such as microfluidics and T-junction mixing of two fluid streams, one of which contains the polynucleotide and the other has the lipid components.

[1054] Lipid compositions are prepared by combining an ionizable amino lipid disclosed herein, e.g., a lipid according to Formula (I) such as Compound II or a lipid according to Formula (III) such as Compound VI, a phospholipid (such as DOPE or DSPC, obtainable from Avanti Polar Lipids, Alabaster, AL), a PEG lipid (such as 1,2-dimyristoyl-*sn*-glycerol methoxypolyethylene glycol, also known as PEG-DMG, obtainable from Avanti Polar Lipids, Alabaster, AL), and a structural lipid (such as cholesterol, obtainable from Sigma-Aldrich, Taufkirchen, Germany, or a corticosteroid (such as prednisolone, dexamethasone, prednisone, and hydrocortisone), or a combination thereof) at concentrations of about 50 mM in ethanol. Solutions should be refrigerated for storage at, for example, -20° C. Lipids are combined to yield desired molar ratios and diluted with water and ethanol to a final lipid concentration of between about 5.5 mM and about 25 mM.

[1055] Nanoparticle compositions including a polynucleotide and a lipid composition are prepared by combining the lipid solution with a solution including the a polynucleotide at lipid composition to polynucleotide wt:wt ratios between about 5:1 and about 50:1. The lipid solution is rapidly injected using a NanoAssemblr microfluidic based system at flow rates between about 10 ml/min and about 18

ml/min into the polynucleotide solution to produce a suspension with a water to ethanol ratio between about 1:1 and about 4:1.

[1056] For nanoparticle compositions including an RNA, solutions of the RNA at concentrations of 0.1 mg/ml in deionized water are diluted in 50 mM sodium citrate buffer at a pH between 3 and 4 to form a stock solution.

[1057] Nanoparticle compositions can be processed by dialysis to remove ethanol and achieve buffer exchange. Formulations are dialyzed twice against phosphate buffered saline (PBS), pH 7.4, at volumes 200 times that of the primary product using Slide-A-Lyzer cassettes (Thermo Fisher Scientific Inc., Rockford, IL) with a molecular weight cutoff of 10 kD. The first dialysis is carried out at room temperature for 3 hours. The formulations are then dialyzed overnight at 4° C. The resulting nanoparticle suspension is filtered through 0.2 μm sterile filters (Sarstedt, Nümbrecht, Germany) into glass vials and sealed with crimp closures. Nanoparticle composition solutions of 0.01 mg/ml to 0.10 mg/ml are generally obtained.

[1058] The method described above induces nano-precipitation and particle formation. Alternative processes including, but not limited to, T-junction and direct injection, can be used to achieve the same nano-precipitation.

B. Characterization of nanoparticle compositions

[1059] A Zetasizer Nano ZS (Malvern Instruments Ltd, Malvern, Worcestershire, UK) can be used to determine the particle size, the polydispersity index (PDI) and the zeta potential of the nanoparticle compositions in 1×PBS in determining particle size and 15 mM PBS in determining zeta potential.

[1060] Ultraviolet-visible spectroscopy can be used to determine the concentration of a polynucleotide (*e.g.*, RNA) in nanoparticle compositions. 100 μL of the diluted formulation in 1×PBS is added to 900 μL of a 4:1 (v/v) mixture of methanol and chloroform. After mixing, the absorbance spectrum of the solution is recorded, for example, between 230 nm and 330 nm on a DU 800 spectrophotometer (Beckman Coulter, Beckman Coulter, Inc., Brea, CA). The concentration of polynucleotide in the nanoparticle composition can be calculated based on the extinction coefficient of the polynucleotide used in the composition and on the difference between the absorbance at a wavelength of, for example, 260 nm and the baseline value at a wavelength of, for example, 330 nm.

[1061] For nanoparticle compositions including an RNA, a QUANT-IT™ RIBOGREEN® RNA assay (Invitrogen Corporation Carlsbad, CA) can be used to evaluate the encapsulation of an RNA by the nanoparticle composition. The samples are diluted to a concentration of approximately 5 μg/mL in a TE buffer solution (10 mM Tris-HCl, 1 mM EDTA, pH 7.5). 50 μL of the diluted samples are transferred to a polystyrene 96 well plate and either 50 μL of TE buffer or 50 μL of a 2% Triton X-100 solution is added to the wells. The plate is incubated at a temperature of 37° C for 15 minutes. The RIBOGREEN® reagent is diluted 1:100 in TE buffer, and 100 μL of this solution is added to each well. The fluorescence intensity can be measured using a fluorescence plate reader (Wallac Victor 1420 Multilablel Counter; Perkin Elmer, Waltham, MA) at an excitation wavelength of, for example, about 480 nm and an emission wavelength of, for example, about 520 nm. The fluorescence values of the reagent blank are subtracted from that of each of the samples and the percentage of free RNA is determined by dividing the fluorescence intensity of the intact sample (without addition of Triton X-100) by the fluorescence value of the disrupted sample (caused by the addition of Triton X-100).

[1062] Exemplary formulations of the nanoparticle compositions are presented in the Table 6 below. The term "Compound" refers to an ionizable lipid such as MC3, Compound II, or Compound VI. "Phospholipid" can be DSPC or DOPE. "PEG-lipid" can be PEG-DMG or Compound I.

Table 6. Exemplary Formulations of Nanoparticles

Composition (mol %)	Components
40:20:38.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
45:15:38.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
50:10:38.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
55:5:38.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
60:5:33.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
45:20:33.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
50:20:28.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
55:20:23.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
60:20:18.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
40:15:43.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
50:15:33.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
55:15:28.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
60:15:23.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
40:10:48.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
45:10:43.5:1.5	Compound:Phospholipid:Chol:PEG-lipid

Composition (mol %)	Components
55:10:33.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
60:10:28.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
40:5:53.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
45:5:48.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
50:5:43.5:1.5	Compound:Phospholipid:Chol:PEG-lipid
40:20:40:0	Compound:Phospholipid:Chol:PEG-lipid
45:20:35:0	Compound:Phospholipid:Chol:PEG-lipid
50:20:30:0	Compound:Phospholipid:Chol:PEG-lipid
55:20:25:0	Compound:Phospholipid:Chol:PEG-lipid
60:20:20:0	Compound:Phospholipid:Chol:PEG-lipid
40:15:45:0	Compound:Phospholipid:Chol:PEG-lipid
45:15:40:0	Compound:Phospholipid:Chol:PEG-lipid
50:15:35:0	Compound:Phospholipid:Chol:PEG-lipid
55:15:30:0	Compound:Phospholipid:Chol:PEG-lipid
60:15:25:0	Compound:Phospholipid:Chol:PEG-lipid
40:10:50:0	Compound:Phospholipid:Chol:PEG-lipid
45:10:45:0	Compound:Phospholipid:Chol:PEG-lipid
50:10:40:0	Compound:Phospholipid:Chol:PEG-lipid
55:10:35:0	Compound:Phospholipid:Chol:PEG-lipid
60:10:30:0	Compound:Phospholipid:Chol:PEG-lipid

EXAMPLE 16

Liver-Specific G6PC(-/-) Knockout Mouse Model for GSD-Ia

[1063] To assess the ability of 1-methyl-pseudouridine modified mRNA constructs encoding G6PC to express G6PC *in vivo*, mRNA encoding wild-type human G6PC or a mutated G6PC, was introduced into an animal model of GSD-Ia.

[1064] A genetic mouse model of GSD-Ia is the liver-specific G6PC knock-out mouse (L-G6PC(-/-)) mouse model (Mutel et al., J. Hepatol., 2011, 54(3):529-37, herein incorporated by reference in its entirety). A mouse model for GSD-Ia has been generated that has a G6PC knocked-out completely, but these mice usually do not live longer than 3 months, making it impossible to evaluate long-term pathogenesis. By contrast, L-G6PC(-/-) mice were generated by tamoxifen-induced liver deletion of G6PC in adult mice. These mice are viable and exhibit normoglycemia when in the fed state. However, L-G6PC(-/-) mice cannot mobilize glycogen like wild-type mice, as they exhibit significantly elevated glycogen levels and much lower blood glucose levels (and are hypoglycemic) after a 6-hour fast. As time progresses, blood glucose levels gradually increase in these mice, possibly due to compensation from

extrahepatic tissues. L-G6PC(-/-) mice develop enlarged livers and hepatic steatosis as a result of glycogen and triglycerides accumulation.

[1065] As a result, mRNA efficacy was evaluated by intravenously injecting L-G6pc(-/-) mice with either G6PC mRNA or control mRNA encoding GFP. The mRNA was formulated in lipid nanoparticles for delivery into mice, e.g., in MC3 or other lipid formulations. Mice were fasted for 2.5 to 24 hours after mRNA injection. Glucose levels and body weight were measured at the time of injection (time 0) and at numerous time points after injection (e.g., at 2.5 hours, 6 hours, 9 hours, and 24 hours after injection). Under some circumstances, at 24 hours after injection, mice were sacrificed, and mRNA levels, G6PC activity, glycogen levels, and G6P levels were measured in liver; tissue weight was measured (liver weight); and triglycerides, cholesterol, uric acid, lactic acid, ALT were measured in serum.

EXAMPLE 17

Engineered G6PC Variants to Increase Expression and Activity

[1066] Mutations were introduced into wild-type human G6PC to improve the potency and/or stability of G6PC. Mutants were expressed in mice to determine whether they exhibited increased expression levels, increased G6Pase activity, and longer duration of action relative to wild-type G6PC. Two G6PC protein variants had been identified that exhibited increased protein expression levels and increased hepatic G6Pase activity relative to the wild-type protein when expressed in wild-type CD-1 mice. One mutant G6PC construct (Variant 1 (V1), SEQ ID NO:50) contains the S297G, S298C, V308L, V321I, and V340L substitutions. The second mutant G6PC construct (Variant 2 (V2), SEQ ID NO:51) contains the Q139R, I142K, Q247R, S298C, and V308L substitutions. Both variants (V1 and V2) showed higher protein expression and G6Pase activity than the wild-type G6PC (WT), while the Variant 2 (V2) exhibited higher protein expression and G6Pase activity than Variant 1 (see **FIGs. 3A** and **3B**).

[1067] Bioinformatics tools were used to identify individual amino acid substitutions that exhibit enhanced expression and/or activity level in comparison to wild-type human G6PC sequence. The goal was to identify G6PC mutants likely to exhibit expression and/or activity that are the same or better relative to the Variant 2 G6PC mutant, but carry fewer mutations than the Variant 2 mutant. **FIG. 4** shows the results of this bioinformatics analysis. Greater than 100 mammalian G6PC protein

sequences were aligned and the consensus amino acids used in each position was compared with that used in human G6PC. Accordingly, a consensus score and reference score in each position was calculated (**FIG. 4**). Positions with larger difference in the consensus and reference scores were identified, and substitution of those individual amino acid in the human G6PC with the consensus amino acid (or more frequently used by non-human mammalian species) were made and tested to determine whether the substitution improves G6PC expression and/or activity.

[1068] The G6Pase activity of a subset of these individual G6PC substitutions (with greatest difference between the reference and consensus score, boxed in **FIG.4**) was tested in Hela cells. Hela cells were grown in 6-well plates, and transfected with 1 μg of 1-methyl-pseudouridine modified mRNA encoding G6PC with a single amino acid substitution. G6Pase activity was tested 48 hours later in cell lysates, as described in Example 15. Specifically, cells were transfected with mRNA encoding human G6PC with a single substitution selected from Q139R (SEQ ID NO:53), I142K (SEQ ID NO:54), Q247R (SEQ ID NO:55), S297G (SEQ ID NO:56), S298C (SEQ ID NO:57), V308L (SEQ ID NO:58), V321I (SEQ ID NO:59), V321L (SEQ ID NO:60), V340L (SEQ ID NO:61), V332A (SEQ ID NO:62), and S298A (SEQ ID NO:65), or mRNA encoding human G6PC with multiple substitutions selected from Q139R/Q247R/S298C (SEQ ID NO:63) and Q247R/S298C (SEQ ID NO:64). G6Pase activity was compared to control cells transfected with eGFP, mRNA encoding wild-type human G6PC, or mRNA encoding Variant 2 G6PC. **FIG. 5A** shows that the S298C single substitution confers enhanced G6PC activity at a level that is similar to the Variant 2 G6PC mutant, which has five amino acid substitutions (Q139R, I142K, Q247R, S298C, and V308L). Thus, the S298C substitution is likely responsible for most of the enhanced activity observed for the Variant 2 G6PC mutant.

[1069] Additional mutations were made to G6PC to confirm that the S298C substitution is largely responsible for the enhanced activity and expression of the G6PC Variant 2 mutant. 1-methyl-pseudouridine modified mRNAs encoding G6PC with an S298A substitution, an S298T substitution, Q247R and S298C substitutions, Q139R and Q247R substitutions were transfected into Hela cells as described above, and G6Pase activity was tested 48 hours later in cell lysates, as described in Example 15. G6Pase activity was compared to control cells transfected with eGFP, mRNA encoding wild-type human G6PC, or mRNA encoding Variant 1 G6PC. **FIG. 5B**

confirms that the S298C substitution is most responsible for the enhanced activity of the Variant 2 G6PC mutant. Mutation of serine to cysteine is necessary for the enhanced stability and activity of G6PC, as the S298A and S298T substitutions exhibited low activity. In addition, combining S298C with Q139R or Q247R, other substitutions that exhibited higher G6Pase activity (see **FIG. 5A**), had moderate effect on the elevated G6Pase activity of the S298C substitution. Western blotting analysis was conducted, using Erp72 as a control to normalize the relative expression of human G6PC. **FIG. 5C** shows that mRNA encoding G6PC with the S298C substitution exhibited a high expression level and/or enhanced stability, near the expression level and/or enhanced stability of the Variant 2 mutant. The mRNA encoding G6PC with the Q139R and Q247R substitutions also exhibited a high expression level and/or stability, near the level observed for the Variant 2 mutant.

[1070] The activity and expression levels of the individual Q139R and Q247R G6PC substitutions were also examined as described above. **FIG. 5D** shows that G6PC with a Q139R or Q247R substitution exhibits greater G6Pase activity than wild-type G6PC, but less G6Pase activity than G6PC having an S298C substitution. **FIG. 5E** shows the specificity of this effect, as mutation of glutamine to alanine, rather than arginine, at amino acid position 139 or 247 of G6PC eliminated the enhanced G6PAse activity. **FIG. 5F** shows that G6PC with a Q139R or Q247R substitution exhibits greater expression than wild-type G6PC, but less expression than G6PC having an S298C substitution. **FIG. 5G** shows the specificity of this effect, as mutation of glutamine to alanine, rather than arginine, at amino acid position 139 or 247 of G6PC eliminated the enhanced expression. Therefore, in addition to S198C, the Q139R and Q247R substitution in human G6PC also appears to be beneficial in enhancing the protein expression and activity of human G6PC mRNA in mammalian cells.

EXAMPLE 18

Codon Optimization of mRNA Constructs

[1071] The mRNAs encoding engineered G6PC variants were codon optimized using an algorithm to further improve the performance of the mRNAs to increase protein expression and activity.

[1072] The Variant 1 and Variant 2 mutants were each codon optimized. Five codon-optimized mRNA constructs were created for each of the Variable 1 and Variable 2 G6PC mutants (hG6PC V1 Co-1 (SEQ ID NO:72), hG6PC V1 Co-2 (SEQ ID

NO:73), hG6PC V1 Co-3 (SEQ ID NO:74), hG6PC V1 Co-4 (SEQ ID NO:75), hG6PC V1 Co-5 (SEQ ID NO:76), hG6PC V2 Co-1 (SEQ ID NO:77), hG6PC V2 Co-2 (SEQ ID NO:78), hG6PC V2 Co-3 (SEQ ID NO:79), and hG6PC V2 Co-4 (SEQ ID NO:80), hG6PC V2 Co-5 (SEQ ID NO:81)). The codon optimized Variant 1 mRNA (hG6PC V1_Co-3; SEQ ID NO:74), codon optimized Variant 2 mRNA (hG6PC V2_Co-1; SEQ ID NO:77), or Variant 2 mRNA that was not codon optimized (SEQ ID NO:51) was injected in a single 1.0 mg/kg dose into wild-type mice via a single injection, and liver G6PC activity was measured at 48 or 96 hours post-injection, as described in Example 15 but using liver microsomes. Control mice were injected with mRNA encoding eGFP. The mRNAs were formulated in either MC3 lipid nanoparticles or in Compound II lipid nanoparticles for delivery into mice. **FIGs. 6A** and **6B** show that G6PC expressed from both the codon optimized Variant 1 and Variant 2 mRNAs exhibited higher G6Pase activity compared to G6PC expressed from non-codon-optimized Variant 2 (V2) mRNA at both 48 and 96 hours post-injection, regardless of with lipid nanoparticles were used. * $P<0.05$ vs Variant 2 (V2) one-way ANOVA followed by Dunnett's test, within each time point.

[1073] In addition, codon optimized Variant 1 mRNA (hG6PC V1 Co-3 (SEQ ID NO:74), hG6PC V1 Co-4 (SEQ ID NO:75), hG6PC V1 Co-5 (SEQ ID NO:76)), and codon optimized Variant 2 mRNA (hG6PC V2 Co-1 (SEQ ID NO:77), hG6PC V2 Co-3 (SEQ ID NO:79), and hG6PC V2 Co-5 (SEQ ID NO:81)), or Variant 1 and Variant 2 mRNA that was not codon optimized (SEQ ID NO:50 and SEQ ID NO: 51) was injected in a single 1.0 mg/kg dose into wild-type mice via a single injection (n=6/per construct), and liver G6PC activity was measured at 96 hours post-injection, as described in Example 15 but using liver microsomes. Control mice were injected with mRNA encoding eGFP. mRNAs were formulated in Compound II lipid nanoparticles for delivery into mice. **FIG. 6C** shows that injection of codon optimized mRNA encoding Variant 1 mutant proteins (hG6PC V2 CoV2-1) (Variant 2 or CoV2 as shown below) resulted in higher G6Pase activity generally relative to injection of non-codon optimized mRNA encoding the Variant 1 and Variant 2 mutants, although the hG6PC V2 CoV2-5 mRNA also exhibited higher G6PAse activity. **FIG. 6D** shows that injection of mRNA encoding the codon optimized hG6PC V1 CoV2-3 mRNA construct (Variant 1 or CoV1 as shown below) resulted in the highest G6PC protein expression. **$P<0.01$, ***$P<0.001$, ****$P<0.0001$ vs. hG6PC V2 One-Way ANOVA followed by Dunnett's test. Therefore, CoV2-1

(Variant 1) and CoV2-3 (Variant 2) were selected as templates for further improvement.

[1074] The mRNA encoding G6PC with an S298C substitution was codon optimized based on the codon-optimized Variant 1 mRNA sequence (the substitution was introduced into SEQ ID NO:74, producing S298C_CoV1, SEQ ID NO:66) or on the codon-optimized Variant 2 mRNA sequence (the substitution was introduced into SEQ ID NO:77, producing S298C_CoV2, SEQ ID NO:69). The mRNA encoding G6PC with Q139R, Q247R, and S298C substitutions was codon optimized based on the codon-optimized Variant 1 mRNA sequence (the substitution was introduced into SEQ ID NO:74, producing Q139R/Q247R/S298C_CoV1, SEQ ID NO:67) or on the codon-optimized Variant 2 mRNA sequence (the substitution was introduced into SEQ ID NO:77, producing Q139R/Q247R/S298C/CoV2, SEQ ID NO:70). The mRNA encoding G6PC with Q139R, I142K, Q247R, S297G, S298C, V308L, V321I, and V340L substitutions was codon optimized based on the codon-optimized Variant 1 mRNA sequence (the substitution was introduced into SEQ ID NO:74, producing M8_CoV1, SEQ ID NO:68) or on the codon-optimized Variant 2 mRNA sequence (the substitution was introduced into SEQ ID NO:77, producing M8 _CoV2, SEQ ID NO:71).

EXAMPLE 19

In vivo Expression and Activity of an mRNA-Encoded G6PC Variants in Wild-Type Mice

[1075] To test the ability of codon-optimized and non-codon-optimized 1-methyl-pseudouridine modified mRNA constructs to express G6PC *in vivo*, and to test the activity of the G6PC, a single 1.0 mg/kg dose of each mRNA construct was IV administered to wild-type CD-1 mice via tail vein injection. Non-codon-optimized mRNA constructs encoding G6PC with an S298C substitution (SEQ ID NO:57), and S298A mutation (SEQ ID NO:65), Q139R/Q247R/S298C mutations (SEQ ID NO:63), or Q247R/S298C mutations (SEQ ID NO:64); or codon-optimized mRNA constructs, including S298C_CoV1 (SEQ ID NO:66), S298C_CoV2 (SEQ ID NO:69), Q139R/Q247R/S298C_CoV1 (SEQ ID NO:67), Q139R/Q247R/S298C_CoV2 (SEQ ID NO:70), M8_CoV1 (SEQ ID NO:68), and M8_CoV2 (SEQ ID NO:71). The mRNA was formulated in lipid nanoparticles (Compound II) for delivery into mice. Control mice were administered eGFP or mRNA encoding wild-type human G6PC. Four mice were injected per mRNA

construct. Animals were sacrificed 96 hours after injection, and 0.5 g of liver was removed from each animal and homogenized in 5 ml of saline to prepare liver microsomes. G6PC protein expression was determined by capillary electrophoresis or Western blot, and G6Pase activity was determined, using the liver microsomes.

[1076] **FIG. 7A** shows that the mRNA constructs that were codon optimized based on the codon-optimized Variant 1 G6PC mutant exhibited higher G6Pase activity levels compared to those constructs that were codon optimized based on the Variant 2 G6PC mutant or those constructs that were not codon optimized. The G6PC proteins with the S298C, Q139R/Q247R/S298C, or the M8 substitutions exhibited similar G6Pase levels. **FIG. 7B** shows that the mRNA constructs that were codon optimized based on the codon-optimized Variant 1 G6PC mutant exhibited higher G6PC expression levels compared to those constructs that were codon optimized based on the Variant 2 G6PC mutant or those constructs that were not codon optimized. Based on this data, the S298C-CoV1 was selected for further pharmacology study in Gsd1a disease models in mice.

EXAMPLE 20

Assessing Duration of Action of mRNA-Encoded G6PC in a Liver-Specific Knockout Mouse Model – Single Dose Studies

[1077] To test the efficacy and duration of action of 1-methyl-pseudouridine modified mRNA encoding a codon-optimized human G6PC S298C variant (S298C_CoV1; SEQ ID NO:66) *in vivo*, a single 0.5 mg/kg dose or 1.0 mg/kg dose of mRNA was IV administered to L-G6PC(-/-) mice (n = 8-10 mice per dose) via tail vein injection. The mRNA was formulated in lipid nanoparticles (Compound II/ PEG-DMG) for delivery into mice. As controls, mRNA encoding eGFP (at 1 mg/kg) was injected into L-G6PC(-/-) mice (n = 9 mice), and PBS was injected into wild-type mice (n = 8 mice). Blood glucose levels were monitored at 2.5 hours and 6 hours fasting at 3 days prior to mRNA injection (-3 days), and on each of 1, 3, 5, 8, 11, and 15 days post-injection.

[1078] **FIGs. 8A-8F** show that L-G6PC(-/-) mice that were administered mRNA encoding G6PC with the S298C substitution exhibited higher blood glucose levels than mice injected with the eGFP control after being fasted for 2.5 hours and for 6 hours for up to 11 days post-injection. Thus, injection of G6PC mRNA resulted in fasting blood glucose levels in L-G6PC(-/-) mice that were higher than eGFP control

and closer to normal levels, although the levels were not as high as the glucose levels observed in wild-type mice. This effect was observed when the mRNA was injected at 0.5 mg/kg or at 1 mg/kg dose. By day 15 (see **FIG. 8F**), mRNA injected mice no longer exhibited higher blood glucose levels relative to eGFP control mice. As expected, control wild-type mice injected with PBS maintained high blood glucose levels when fasted, and all mice (wild-type control and L-G6PC(-/-)) had higher blood glucose levels when fed. **P<0.01 vs. GFP one-way ANOVA followed by multiple comparison. ****P<0.0001 vs. GFP one-way ANOVA followed by multiple comparison.

[1079] **FIGs. 8G-8H** show the time course effect of administration of the 0.5 mg/kg dose (**FIG. 8G**) and the 1 mg/kg dose (**FIG. 8H**). In both figures, the therapeutic threshold of 70 mg/dL is indicated by a solid bar.

[1080] **FIGs. 9A-9C** show the effects of injecting mRNA encoding G6PC with the S298C substitution on fasting glucose levels in L-G6PC(-/-) mice, as compared to fasting glucose levels prior to mRNA injection. **FIG. 9A** shows that fasting glucose were elevated significantly through 11 days post-injection when mRNA was administered at 1 mg/kg. **FIG. 9B** shows that fasting glucose were elevated significantly through 8 days post-injection when mRNA was administered at 0.5 mg/kg. **FIG. 9C** shows that administering eGFP to L-G6PC(-/-) mice did not result in elevated fasting glucose levels, as expected for this control. **P<0.01 vs. pre-treatment Paired t-test. ***P<0.001 vs. pre-treatment Paired t-test. ****P<0.0001 vs. pre-treatment Paired t-test.

EXAMPLE 21

Assessing Dose Response of mRNA Encoding G6PC in a Liver-Specific Knockout Mouse Model

[1081] To test the dose response of 1-methyl-pseudouridine modified mRNA encoding a codon-optimized human G6PC S298C variant (S298C_CoV1; SEQ ID NO:66) *in vivo*, a single 0.2 mg/kg dose, 0.5 mg/kg dose or 1.0 mg/kg dose of mRNA was IV administered to L-G6PC(-/-) mice via tail vein injection. Seven to eight mice were injected per mRNA construct. The mRNA was formulated in lipid nanoparticles (Compound II/PEG-DMG) for delivery into mice. As controls, mRNA encoding eGFP (at 1 mg/kg) was injected into L-G6PC(-/-) mice (n = 8 mice), and PBS was injected into wild-type mice (n = 8 mice). Fasting glucose levels were measured in

mice fasted for 2.5 hours, 6 hours, and 24 hours, and compared to glucose levels in mice that were fed. At 24 hours post-injection, liver samples were collected, and liver weight and appearance was recorded, and the levels of glycogen, G-6-phosphate, triglyceride, G6PC protein expression, G6PC activity, hG6PC mRNA, and IHC were measured in the samples. At 24 hours post-injection, plasma was collected from the mice, and the liver enzyme ALT, the lipids TG and cholesterol, lactate, and hG6PC mRNA levels were measured in the plasma.

[1082] **FIG. 10A** shows that G6PC mRNA (SEQ ID NO:66) injected at each of the 0.2 mg/kg, 0.5 mg/kg, and 1.0 mg/kg doses increased glucose levels in L-G6PC(-/-) mice at 2.5 hours, 6 hours, and 24 hours of fasting relative to the glucose levels in control mice injected with eGFP mRNA. At 2.5 hours fasting, a single mRNA dose of 1 mg/kg resulted in glucose levels that were near the glucose levels observed in wild-type control mice injected with PBS. However, single doses of mRNA at 0.2 mg/kg, 0.5 mg/kg, and 1.0 mg/kg resulted in similar elevated levels of glucose at 6 hours and 24 hours fasting that were near the levels see in fasted wild-type control mice. *P<0.05, **P<0.01, ***P<0.001, ****P<0.0001 vs. GFP Two-way ANOVA followed by multiple comparisons. **FIG. 10B** shows that injection of 0.2 mg/kg, 0.5 mg/kg, or 1.0 mg/kg of the G6PC mRNA resulted in fasting blood glucose levels in L-G6PC(-/-) that were above the therapeutic threshold of 70 mg/dL (indicated by a solid bar in **FIG. 10B**).

[1083] **FIG. 10C** shows the effects of injecting single 0.2 mg/kg, 0.5 mg/kg, and 1 mg/kg doses of mRNA encoding G6PC with the S298C substitution on fasting glucose levels in L-G6PC(-/-) mice, as compared to fasting glucose levels prior to mRNA injection. Injecting each dose results in an increase in blood glucose levels post-injection, and injection of mRNA at a 1 mg/kg dose results in the largest increase in glucose. By contrast, injecting PBS in control wild-type mice, and injecting mRNA encoding GFP in L-G6PC(-/-) mice did not increase blood glucose levels. **P<0.01, ****P<0.0001 vs. pre-treatment Paired t-test.

[1084] **FIGs. 11A-11B** show the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on liver weight in L-G6PC(-/-) mice, both in the total weight (in grams) of the livers (**Fig. 11A**) and as the weight of the liver as a percentage of total body weight (**FIG. 11B**). As expected, control L-G6PC(-/-) mice injected with mRNA encoding GFP had substantially larger livers compared to wild-type control mice injected with PBS. Injection of any single 0.2 mg/kg, 0.5 mg/kg or

1 mg/kg dose of mRNA encoding the human G6PC protein resulted in a significant decrease in liver weight relative to the L-G6PC(-/-) mice treated with the control eGFP mRNA. Injecting the 1 mg/kg dose of mRNA resulted in the largest decrease in liver size, and the livers of these mice were similar in weight to those of the wild-type control mice. ****P<0.0001 vs. GFP One-way ANOVA followed by multiple comparisons.

[1085] **FIG. 12** shows representative liver images from wild-type mice or L-G6PC(-/-) mice injected with mRNA encoding G6PC with the S298C substitution at 0.2, 0.5, or 1.0 mg/kg, or with mRNA encoding GFP. As expected, the liver from L-G6PC(-/-) mice were enlarged and had a whitish and pale appearance compared to the liver in wild-type mice. Injection with mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) resulted in significant normalization of the liver appearance in L-G6PC(-/-) mice.

[1086] **FIG. 13A** shows the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on hepatic glucose-6 phosphate (G6P), the endogenous substrate of G6Pase. As expected, the hepatic G6P content from L-G6PC(-/-) mice is significantly elevated in comparison with wild-type mice (compare L-G6PC(-/-) mice injected with eGFP mRNA and wild-type mice injected with PBS). Injection of mRNA encoding G6PC with the S298C substitution led to a significant and dose-dependent decrease in hepatic G6P content. **P<0.01, ***P<0.001, ****P<0.0001 vs. eGFP, one-way ANOVA followed by multiple comparisons.

[1087] **FIG. 13B** shows the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on hepatic glycogen levels. The hepatic glycogen content from L-G6PC(-/-) mice is significantly elevated in comparison with wild-type mice (compare L-G6PC(-/-) mice injected with eGFP mRNA and wild-type mice injected with PBS). Injection of mRNA encoding G6PC with the S298C substitution led to a significant decrease in hepatic glycogen content.

[1088] **FIG. 13C** shows the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on serum triglyceride levels. The serum triglyceride level from L-G6PC(-/-) mice is significantly elevated in comparison with wild-type mice (compare L-G6PC(-/-) mice injected with eGFP mRNA and wild-type mice injected with PBS). Injection of mRNA encoding G6PC with the S298C substitution led to a significant decrease in serum triglyceride level.

[1089] **FIG. 13D** shows the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on serum ALT (liver enzyme) levels. The serum ALT level from L-G6PC(-/-) mice is significantly elevated in comparison with wild-type mice (compare L-G6PC(-/-) mice injected with eGFP mRNA and wild-type mice injected with PBS). Injection of mRNA encoding G6PC with the S298C substitution led to a significant decrease in serum ALT level.

[1090] **FIG. 13E** shows the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on serum lactate levels in 3-4 month old mice. **FIG. 13F** shows the effects of injecting mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) on liver lactate levels in 3-4 month old mice.

[1091] **FIGs. 14A-14C** show hepatic G6Pase activity (**FIG. 14A**) and G6PC protein expression (**FIGs. 14B-C**) in L-G6PC(-/-) mice injected with mRNA encoding G6PC with the S298C substitution (SEQ ID NO:66) at 0.2, 0.5, or 1.0 mg/kg, or with mRNA encoding GFP. As expected, the L-G6PC(-/-) mice injected with mRNA encoding GFP had a very low hepatic G-6-phosphatase activity in comparison with the wild-type mice (injected with PBS). Injection with mRNA encoding G6PC with the S298C substitution resulted in a dose-dependent increase in hepatic G-6-Phosphatase activity (**FIG. 14A**). *P<0.05, **P<0.01, ****P<0.0001 vs. GFP, one-way ANOVA followed by multiple comparisons. Injection with mRNA encoding the G6PC with the S298C substitution also resulted in a dose-dependent increase in hepatic G6PC protein expression (**FIGs. 14B-C**; as analyzed by western blot and quantitated using the imaging instrument Odyssey CLx and accompanying applications (Li-Cor, Lincoln, NE)). **P<0.01, ***P<0.001, ****P<0.0001 vs. eGFP, one-way ANOVA followed by multiple comparisons.

EXAMPLE 22

Pharmacokinetic (PK) Analysis of mRNA-Encoded G6PC Variants in Wild-Type Mice

[1092] A pharmacokinetic analysis of 1-methyl-pseudouridine modified mRNAs encoding human G6PC variants was conducted in wild-type CD-1 mice. Mice were administered a single 1 mg/kg dose of mRNA encoding G6PC with the S298C substitution (S298C_CoV1; SEQ ID NO:66) or mRNA encoding G6PC with S298C/Q139R/Q247R substitutions (Q139R/Q247R/S298C_CoV1; SEQ ID NO:67) via tail vein injection. The mRNA was formulated in lipid nanoparticles (Compound II/PEG-DMG) for delivery into mice. Control mice were administered eGFP. Liver

samples were isolated at 6 hours, 1 day, 3 days, 7 days, and 14 days following injection, as described. mRNA levels were analyzed in the samples using RT-PCR and normalized by ActB (β-actin) mRNA levels. G6PC protein expression levels and G6PC activity levels were also assayed in the liver samples at each time point, as described above.

[1093] **FIG. 15A** shows the amount of mRNA (pg of G6PC mRNA per 100 ng of total RNA) in the livers of injected animals over time. The amount of each of the mRNAs encoding the G6PC variants (S298C_CoV1 and Q139R/Q247R/S298C_CoV1) reduced rapidly over 24 hours, and were no longer detectable by 72 hours post-injection. **FIG. 15B** and **15C** show the protein expression levels of the mRNA-expressed G6PC S298C and G6PC Q139R/Q247R/S298C proteins over time, and **FIG. 15D** compares the time courses of protein expression for these G6PC variants. G6PC S298C protein levels decrease to lower levels at an earlier time (e.g., day 7) following mRNA injection than G6PC Q139R/Q247R/S298C protein levels, although both proteins have similar low levels of expression by 14 days (336 hours) post-injection. **FIG. 15E** shows that the G6PC S298C protein exhibited a significantly lower area under the curve (AUC) value based on protein expression than the G6PC Q139R/Q247R/S298C protein. **P<0.01. **FIG. 15F** shows that the G6Pase activity of mRNA-expressed G6PC S298C protein was slightly but consistently lower than the G6Pase activity of mRNA-expressed G6PC Q139R/Q247R/S298C protein over the course of 14 days following mRNA injection. **FIG. 15G** shows that the G6PC S298C protein exhibited a significantly lower area under the curve (AUC) value based on protein activity than the G6PC Q139R/Q247R/S298C protein. **P<0.01.

[1094] Mice were administered a single 1 mg/kg dose of mRNA encoding G6PC with the S298C substitution (hG6PC S298C_CoV1 (SEQ ID NO:66) or G6PC_S298C_Astart_3XmiR142 (SEQ ID NO:215)) via tail vein injection. The mRNA was formulated in lipid nanoparticles (Compound II/PEG-DMG) for delivery into the mice. Control mice were administered eGFP. Liver samples were isolated 24 hours following injection. G6PC protein expression levels and G6PC activity levels were assayed in the liver samples, as described above. **FIGS. 15H-15I** show the G6PC protein expression levels (**FIG. 15H**) and G6Pase activity (**FIG. 15I**) following administration of each of the mRNAs.

[1095]

EXAMPLE 23

Assessing Sustained Efficacy of mRNA-Encoded G6PC in a Liver-Specific Knockout Mouse Model – Repeat Dose Studies

[1096] L-G6PC(-/-) mice received five consecutive intravenous injections of 1-methyl-pseudouridine modified mRNA encoding G6PC with the S298C substitution (S298C_CoV1; SEQ ID NO:66), 1-methyl-pseudouridine modified mRNA encoding G6PC with S298C/Q139R/Q247R substitutions (Q139R/Q247R/S298C_CoV1; SEQ ID NO:67), or eGFP mRNA, each at doses of 0.5 mg/kg or 0.25 mg/kg and at an interval of 11 to 14 days over the course of 51 days. The mRNA was formulated in lipid nanoparticles (Compound II/PEG-DMG) for delivery into the mice. Wild-type C57BL/6J mice receiving PBS were used as control. The fed and fasting glucose levels (2.5 hours post-fasting) were measured every 3-4 days after each injection. The baseline fed and fasting glucose levels (2.5 hours post-fasting) were also measured 5 days before injecting the mRNAs. After the last (5th) injection, the mice were fasted and the fasting glucose levels were monitored at 2.5, 6, and 24 hours post-fasting. At the end of the in-life study, the mice were sacrificed by cervical dislocation for blood and tissue collections. The livers were harvested, weighed, and photographed. Liver pieces were either snap-frozen in liquid nitrogen for further analysis or fixed in formalin and embedded in paraffin for immunohistochemistry. The plasma samples were also collected and snap-frozen in liquid nitrogen for further analysis.

[1097] **FIG. 16** shows that L-G6PC(-/-) mice that were administered multiple doses of 0.25 mg/kg of mRNA encoding G6PC with the S298C substitution ("single") or mRNA encoding G6PC with the S298C/Q139R/Q247R substitutions ("triple") exhibited higher blood glucose levels than mice injected with the eGFP control after being fasted for 2.5 hours. Injection of each of five doses of both of the G6PC mRNAs resulted in blood glucose levels in L-G6PC(-/-) mice that were above the therapeutic threshold of 70 mg/dL (indicated by a solid bar in **FIG. 16**). As expected, control wild-type mice injected with PBS had high blood glucose levels when fasted.

[1098] **Fig. 17** shows that injection of mRNA encoding G6PC with the S298C substitution ("single") or mRNA encoding G6PC with the S298C/Q139R/Q247R substitutions ("triple") resulted in a significant decrease in liver weight relative to the L-G6PC(-/-) control mice. Liver weight was measured in mice receiving mRNA at 0.5 mg/kg, after the 5th dose.

EXAMPLE 24

Assessing Action of mRNA-Encoded G6PC in Different Lipid Nanoparticle Formulations

[1099] To test the efficacy of different lipid nanoparticle formulations, 1-methyl-pseudouridine modified mRNA encoding the G6PC S298C variant (G6PC_S298C_Astart; SEQ ID NO:214) was IV administered to L-G6PC(-/-) mice (n = 8 or 9 mice per group) in a single 1.0 mg/kg dose of mRNA via tail vein injection. The mRNA was formulated in lipid nanoparticles (Compound II/PEG-DMG or Compound II/ Compound I) for delivery into the mice. As controls, mRNA encoding eGFP (at 1 mg/kg) was injected into L-G6PC(-/-) mice (Compound II/ Compound I), and Tris-Sucrose buffer was injected into wild-type mice. Blood glucose levels were monitored at 2.5 hours fasting on each of days 0, 1, 3, 7, 10, and 14 post-injection.

[1100] **FIG. 18** shows that L-G6PC(-/-) mice that were administered mRNA encoding G6PC with the S298C substitution in either Compound II/PEG-DMG or Compound II/ Compound I lipid nanoparticles exhibited higher blood glucose levels than mice injected with the eGFP control after being fasted for 2.5 hours. Injection of the G6PC mRNA in both lipid nanoparticles resulted in blood glucose levels in L-G6PC(-/-) mice that were above the therapeutic threshold of 70 mg/dL (indicated by a solid bar in **FIG. 18**).

EXAMPLE 25

In Vivo Expression of miRNA-Targeted mRNA-Encoded G6Pase in Wild-Type Mice

[1101] The impact of miRNA-142 target sites in the 3′ UTR of 1-methyl-pseudouridine-modified mRNA encoding a human G6PC variant was tested in wild-type CD-1 mice. Mice were administered a single 1 mg/kg dose of mRNA encoding G6PC with the S298C substitution and lacking miR-142 target sites in the 3′ UTR (hG6PC S298C_CoV1; SEQ ID NO:66) or mRNA encoding G6PC with the S298C substitution and having three miR-142 target sites in the 3′ UTR (G6PC_S298C_Astart_3XmiR142; SEQ ID NO:215). The mRNA was formulated in lipid nanoparticles (Compound II/PEG-DMG) for delivery into mice. Control mice were administered eGFP. Liver and spleen samples were isolated 24 hours following injection, as described. G6PC protein expression levels were assayed in the liver and spleen samples as described above.

[1102] **FIG. 19A** shows the liver protein expression levels of the mRNA-expressed G6PC S298C with (miR142; SEQ ID NO:215) and without (no miR; SEQ ID NO:66) miR-142 target sites in the 3′ UTR. mRNA encoding G6PC S298C with miR-142 target sites in the 3′ UTR (SEQ ID NO:215) was as effective as mRNA encoding G6PC S298C without miR-142 target sites in the 3′ UTR (SEQ ID NO:66) in hepatic expression of G6Pase protein. **FIG. 19B** shows the spleen protein expression levels of the mRNA-expressed G6PC S298C with (miR142; SEQ ID NO:215) and without (no miR; SEQ ID NO:66) miR-142 target sites in the 3′ UTR. Neither mRNA encoding G6PC S298C resulted in G6Pase protein expression in the spleen.

EXAMPLE 26

In Vitro Expression of mRNA-Encoded G6Pase in Kidney Proximal Tubule Segment (TKPTS) Cells

[1103] To test the delivery to kidney cells of LNP-formulated 1-methyl-pseudouridine-modified mRNA encoding a hG6PC variant, various transfection methods were tested in TKPTS cells. Mouse TKPTS cells were mock-transfected (no mRNA, negative control) or transfected with mRNA encoding hG6PC having an S298C substitution (hG6PC S298C_CoV1; SEQ ID NO:66). mRNAs were formulated in Compound II/PEG-DMG or Compound II/Compound I LNPs, or were transfected using Lipofectamine MessengerMAX (ThermoFisher Scientific). At 4 and 24 hours post-transfection, cells were monitored for LNP uptake, hG6PC mRNA levels, and hG6Pase expression levels. LNPs were labeled with rhodamine to track accumulation in the cells. G6PC expression levels were assayed as described above.

[1104] **FIG. 20A** shows the LNP uptake in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C substitution (SEQ ID NO:66) in the presence of 10% fetal bovine serum (FBS; left) or 5% human serum (right); negative control represents non-transfected cells. Compound II/PEG-DMG and Compound II/Compound I LNPs displayed superior LNP uptake compared to MessengerMAX uptake. **FIG. 20B** shows the eGFP expression in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C substitution (SEQ ID NO:66) in the presence of 10% FBS (left) or 5% human serum (right). **FIG. 20C** shows hG6PC expression in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C substitution (SEQ ID NO:66) in the presence of 10% FBS (left) or 5%

human serum (right). **FIG. 20D** shows the percentage of cells expressing transfected mRNA in TKPTS cells at the indicated timepoints post-transfection of the cells with mRNA encoding hG6PC with a S298C substitution in the presence of 10% FBS (left) or 5% human serum (right).

EXAMPLE 27

In Vivo Expression of mRNA-Encoded G6Pase in Kidneys

[1105] The delivery to kidneys of LNP-formulated 1-methyl-pseudouridine-modified hG6PC mRNA was tested in K.G6PC$^{-/-}$ mice. K.G6PC$^{-/-}$ mice were administered 1 mg/kg of Compound II/PEG-DMG LNPs comprising mRNA encoding eGFP or hG6PC having a S298C substitution (hG6PC S298C_CoV1; SEQ ID NO:66) via the intravenous route. As a control, wild-type mice were administered PBS intravenously. Six or 24 hours post-administration, mice were sacrificed and protein expression and enzymatic activity were assayed as described above.

[1106] **FIG. 21A** shows G6Pase expression, as a percentage of the level in wild-type mice, in wild-type mice administered PBS or K.G6PC$^{-/-}$ mice administered eGFP mRNA or hG6PC mRNA (SEQ ID NO:66). **FIG. 21B** shows G6Pase activity in wild-type mice administered PBS or K.G6PC -/- mice administered eGFP mRNA or hG6PC mRNA. G6Pase expression was determined by western blot. K.G6PC$^{-/-}$ mice had ~50% G6Pase expression and activity in kidneys compared to that of wild type mice.

[1107] It is to be appreciated that the Detailed Description section, and not the Summary and Abstract sections, is intended to be used to interpret the claims. The Summary and Abstract sections can set forth one or more but not all exemplary embodiments of the present invention as contemplated by the inventor(s), and thus, are not intended to limit the present invention and the appended claims in any way.

[1108] The present invention has been described above with the aid of functional building blocks illustrating the implementation of specified functions and relationships thereof. The boundaries of these functional building blocks have been arbitrarily defined herein for the convenience of the description. Alternate boundaries can be defined so long as the specified functions and relationships thereof are appropriately performed.

[1109] The foregoing description of the specific embodiments will so fully reveal the general nature of the invention that others can, by applying knowledge within the skill of the art, readily modify and/or adapt for various applications such specific embodiments, without undue experimentation, without departing from the general concept of the present invention. Therefore, such adaptations and modifications are intended to be within the meaning and range of equivalents of the disclosed embodiments, based on the teaching and guidance presented herein. It is to be understood that the phraseology or terminology herein is for the purpose of description and not of limitation, such that the terminology or phraseology of the present specification is to be interpreted by the skilled artisan in light of the teachings and guidance.

[1110] The breadth and scope of the present invention should not be limited by any of the above-described exemplary embodiments, but should be defined only in accordance with the following claims and their equivalents.

[1111] All publications, patent applications, patents, and other references mentioned herein are incorporated by reference in their entirety. In case of conflict, the present specification, including definitions, will control.

WHAT IS CLAIMED IS:

1. A pharmaceutical composition comprising an mRNA comprising an open reading frame (ORF) encoding a glucose-6-phosphatase (G6PC) polypeptide, wherein the composition when administered as a single intravenous dose to a human subject in need thereof is sufficient to:

(i) increase the level of G6PC activity in liver tissue to within at least 2%, at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% of normal G6PC activity level for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(ii) increase the level of G6PC activity in liver tissue at least 1.5-fold, at least 2-fold, at least 3-fold, at least 4-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold compared to the human subject's baseline G6PC activity level or a reference G6PC activity level in a human subject having glycogen storage disease type 1a (GSD-Ia) for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(iii) reduce liver levels of glycogen at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(iv) reduce liver levels of glucose-6-phosphate (G6P) at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(v) reduce liver levels of glycogen at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold when the human subject is fasting as compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(vi) reduce liver levels of G6P at least 1.5-fold, at least 2-fold at least 5-fold, at least 10-fold, at least 20-fold or at least 50-fold as compared to the human subject's baseline liver G6P levels while fasting or a reference liver G6P level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(vii) reduce serum levels of triglycerides at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(viii) reduce serum levels of triglyceride at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold when the human subject is fasting as compared to the human subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(ix) increase blood, plasma, and/or serum levels of glucose at least 2%, at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting compared to the human subject's baseline blood, plasma, or serum glucose level while fasting or a reference blood, plasma, or serum glucose level in a fasting human subject having GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration;

(x) increase blood, plasma, and/or serum levels of glucose at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold when the human subject is fasting as compared to the human subject's baseline blood, plasma, and/or serum glucose level while fasting or a reference blood, plasma, and/or serum glucose level in a fasting patient with GSD-Ia for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration; and/or

(xi) increase blood, plasma, or serum levels of glucose to at least 70 mg/dl, at least 75 mg/dl , at least 80 mg/dl , at least 85 mg/dl , at least 90 mg/dl , at least 95 mg/dl , at least 100 mg/dl, at least 105 mg/dl, at least 110 mg/dl, at least 115 mg/dl, at least 120 mg/dl, at least 125 mg/dl, at least 130 mg/dl, at least 135 mg/dl, at least 140 mg/dl, at least 145 mg/dl, or at least 150 mg/dl in a fasting patient with GSD-Ia for at least 6 hours, at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration.

2. The pharmaceutical composition of claim 1, further comprising a delivery agent.

3. The pharmaceutical composition of claim 2, wherein the delivery agent comprises a lipid nanoparticle comprising:

(i) Compound II, (ii) Cholesterol, and (iii) PEG-DMG or Compound I;

(i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) PEG-DMG or Compound I;

(i) Compound II, (ii) Cholesterol, and (iii) Compound I; or

(i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) Compound I.

4. The pharmaceutical composition of any one of the preceding claims, wherein the G6PC polypeptide consists of the amino acid sequence set forth in SEQ ID NO:83, SEQ ID NO:84, SEQ ID NO:194-206, 208, or 209.

5. The pharmaceutical composition of any one of the preceding claims, wherein the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to a nucleic acid sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207.

6. The pharmaceutical composition of any one of the preceding claims, wherein the mRNA comprises a microRNA (miR) binding site.

7. The pharmaceutical composition of claim 6, wherein the microRNA is expressed in an immune cell of hematopoietic lineage or a cell that expresses TLR7 and/or TLR8 and secretes pro-inflammatory cytokines and/or chemokines.

8. The pharmaceutical composition of claim 6, wherein the microRNA binding site is for a microRNA selected from the group consisting of miR-126, miR-142, miR-144, miR-146, miR-150, miR-155, miR-16, miR-21, miR-223, miR-24, miR-27, miR-26a, or any combination thereof.

9. The pharmaceutical composition of claim 6, wherein the microRNA binding site is for a microRNA selected from the group consisting of miR126-3p, miR-142-3p, miR-142-5p, miR-155, or any combination thereof.

10. The pharmaceutical composition of claim 6, wherein the microRNA binding site is a miR-142-3p binding site.

11. The pharmaceutical composition of any one of claims 6 to 10, wherein the microRNA binding site is located in the 3' UTR of the mRNA.

12. The pharmaceutical composition of any one of the preceding claims, wherein the mRNA comprises a 3' UTR comprising a nucleic acid sequence at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 3' UTR sequence of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213.

13. The pharmaceutical composition of any one of the preceding claims, wherein the mRNA comprises a 5' UTR comprising a nucleic acid sequence at least 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 5' UTR sequence of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211.

14. The pharmaceutical composition of any one of the preceding claims, wherein the mRNA comprises a 5' terminal cap.

15. The pharmaceutical composition of claim 14, wherein the 5' terminal cap comprises a Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5' methylG cap, or an analog thereof.

16. The pharmaceutical composition of any one of the preceding claims, wherein the mRNA comprises a poly-A region.

17. The pharmaceutical composition of claim 16, wherein the poly-A region is at least about 10, at least about 20, at least about 30, at least about 40, at least about 50, at least about 60, at least about 70, at least about 80, at least about 90 nucleotides in length, or at least about 100 nucleotides in length.

18. The pharmaceutical composition of claim 16, wherein the poly-A region has about 10 to about 200, about 20 to about 180, about 50 to about 160, about 70 to about 140, or about 80 to about 120 nucleotides in length.

19. The pharmaceutical composition of any one of the preceding claims, wherein the mRNA comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof.

20. The pharmaceutical composition of claim 19, wherein the at least one chemically modified nucleobase is selected from the group consisting of pseudouracil (ψ), N1-methylpseudouracil (m1ψ), 1-ethylpseudouracil, 2-thiouracil (s2U), 4'-thiouracil, 5-methylcytosine, 5-methyluracil, 5-methoxyuracil, and any combination thereof.

21. The pharmaceutical composition of claim 19 or 20, wherein at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, at least about 90%, at least about 95%, at least about 99%, or 100% of the uracils are chemically modified to N1-methylpseudouracils.

22. The pharmaceutical composition of any one of claims 1-21, wherein the human subject has GSD-Ia.

23. The pharmaceutical composition of any one of claims 1-22, wherein the human subject has been fasting.

24. The pharmaceutical composition of claim 23, wherein the human subject has been fasting for at least 2 hours.

25. A polynucleotide comprising an mRNA comprising:

(i) a 5' UTR;

(ii) an open reading frame (ORF) encoding a human G6PC polypeptide, wherein the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to a nucleic acid sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207;

(iii) a stop codon; and

(iv) a 3' UTR.

26. The polynucleotide of claim 25, wherein:

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:8 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:205;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:11 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:204;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:12 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:195;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:13 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:197;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:14 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:196;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:15 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:198;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:16 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:194;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:17 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:200;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:18 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:201;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:19 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:202;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:20 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:203;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at

least 99%, or 100% sequence identity to SEQ ID NO:22 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:208;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:23 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:194;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:24 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:205;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:25 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:209;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:26 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:194;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:27 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:205;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:28 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:209;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:29 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:30 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:31 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:32 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:33 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:83;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:34 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:35 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:36 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:37 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84;

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or 100% sequence identity to SEQ ID NO:38 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:84; or

the ORF has at least 79%, at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at

least 99%, or 100% sequence identity to SEQ ID NO:207 and the G6PC polypeptide consists of the amino acid sequence of SEQ ID NO:206.

27. The polynucleotide of claim 25 or 26, wherein the mRNA comprises a microRNA (miR) binding site.

28. The polynucleotide of claim 27, wherein the microRNA is expressed in an immune cell of hematopoietic lineage or a cell that expresses TLR7 and/or TLR8 and secretes pro-inflammatory cytokines and/or chemokines.

29. The polynucleotide of claim 27, wherein the microRNA binding site is for a microRNA selected from the group consisting of miR-126, miR-142, miR-144, miR-146, miR-150, miR-155, miR-16, miR-21, miR-223, miR-24, miR-27, miR-26a, or any combination thereof.

30. The polynucleotide of claim 27, wherein the microRNA binding site is for a microRNA selected from the group consisting of miR126-3p, miR-142-3p, miR-142-5p, miR-155, or any combination thereof.

31. The polynucleotide of claim 27, wherein the microRNA binding site is a miR-142-3p binding site.

32. The polynucleotide of any one of claims 25 to 31, wherein the microRNA binding site is located in the 3' UTR of the mRNA.

33. The polynucleotide of any one of claims 25 to 32, wherein the 3' UTR comprises a nucleic acid sequence at least about 90%, at least about 95%, at least about 96%, at least about 97%, at least about 98%, at least about 99%, or 100% identical to a 3' UTR of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213.

34. The polynucleotide of any one of claims 25 to 33, wherein the 5' UTR comprises a nucleic acid sequence at least 90%, at least about 95%, at least about 96%, at least about

97%, at least about 98%, at least about 99%, or 100% identical to a 5' UTR sequence of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211.

35. The polynucleotide of any one of claims 25 to 34, wherein the mRNA comprises a 5' terminal cap.

36. The polynucleotide of claim 35, wherein the 5' terminal cap comprises a Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5' methylG cap, or an analog thereof.

37. The polynucleotide of any one of claims 25 to 36, wherein the mRNA comprises a poly-A region.

38. The polynucleotide of claim 37, wherein the poly-A region is at least about 10, at least about 20, at least about 30, at least about 40, at least about 50, at least about 60, at least about 70, at least about 80, at least about 90 nucleotides in length, or at least about 100 nucleotides in length.

39. The polynucleotide of claim 37, wherein the poly-A region has about 10 to about 200, about 20 to about 180, about 50 to about 160, about 70 to about 140, or about 80 to about 120 nucleotides in length.

40. The polynucleotide of any one of claims 25 to 39, wherein the mRNA comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof.

41. The polynucleotide of claim 40, wherein the at least one chemically modified nucleobase is selected from the group consisting of pseudouracil (ψ), N1-methylpseudouracil (m1ψ), 1-ethylpseudouracil, 2-thiouracil (s2U), 4’-thiouracil, 5-methylcytosine, 5-methyluracil, 5-methoxyuracil, and any combination thereof.

42. The polynucleotide of claim 40 or 41, wherein at least about 25%, at least about 30%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least

about 80%, at least about 90%, at least about 95%, at least about 99%, or 100% of the uracils are chemically modified to N1-methylpseudouracils.

43. The polynucleotide of claim 25, comprising a nucleic acid sequence selected from the group consisting of SEQ ID NO:53-81, 214, 215, and 217.

44. A polynucleotide comprising an mRNA comprising:

(i) a 5'-terminal cap;

(ii) a 5' UTR comprising the nucleic acid sequence of SEQ ID NO:3, SEQ ID NO:39, SEQ ID NO:210, or SEQ ID NO:211;

(iii) an open reading frame (ORF) encoding a G6PC polypeptide, wherein the ORF comprises a sequence selected from the group consisting of SEQ ID NOs:8, 11-20, 22-38, and 207;

(iv) a 3' UTR comprising the nucleic acid sequence of SEQ ID NO:4, SEQ ID NO:111, SEQ ID NO:150, SEQ ID NO:175, SEQ ID NO:177, SEQ ID NO:178, SEQ ID NO:212, or SEQ ID NO:213; and

(vi) a poly-A-region.

45. The polynucleotide of claim 44, wherein the 5' terminal cap comprises a Cap0, Cap1, ARCA, inosine, N1-methyl-guanosine, 2′-fluoro-guanosine, 7-deaza-guanosine, 8-oxo-guanosine, 2-amino-guanosine, LNA-guanosine, 2-azidoguanosine, Cap2, Cap4, 5' methylG cap, or an analog thereof.

46. The polynucleotide of claim 44 or 45, wherein the poly-A region is at least about 10, at least about 20, at least about 30, at least about 40, at least about 50, at least about 60, at least about 70, at least about 80, at least about 90 nucleotides in length, or at least about 100 nucleotides in length.

47. The polynucleotide of claim 44 or 45, wherein the poly-A region has about 10 to about 200, about 20 to about 180, about 50 to about 160, about 70 to about 140, or about 80 to about 120 nucleotides in length.

48. The polynucleotide of any one of claims 44 to 47, wherein the mRNA comprises at least one chemically modified nucleobase, sugar, backbone, or any combination thereof.

49. The polynucleotide of claim 48, wherein the at least one chemically modified nucleobase is selected from the group consisting of pseudouracil (ψ), N1-methylpseudouracil (m1ψ), 1-ethylpseudouracil, 2-thiouracil (s2U), 4'-thiouracil, 5-methylcytosine, 5-methyluracil, 5-methoxyuracil, and any combination thereof.

50. The polynucleotide of claim 44, comprising a nucleic acid sequence selected from the group consisting of SEQ ID NO:53-81, 214, 215, and 217.

51. The polynucleotide of claim 50, wherein the 5' terminal cap comprises Cap1 and all of the uracils of the polynucleotide are N1-methylpseudouracils.

52. The polynucleotide of claim 51, wherein the poly-A-region is 100 nucleotides in length.

53. A pharmaceutical composition comprising the polynucleotide of any one of claims 25 to 52, and a delivery agent.

54. The pharmaceutical composition of claim 53, wherein the delivery agent comprises a lipid nanoparticle comprising:

(i) Compound II, (ii) Cholesterol, and (iii) PEG-DMG or Compound I;

(i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) PEG-DMG or Compound I;

(i) Compound II, (ii) Cholesterol, and (iii) Compound I; or

(i) Compound II, (ii) DSPC or DOPE, (iii) Cholesterol, and (iv) Compound I.

55. A method of expressing a glucose-6-phosphatase (G6PC) polypeptide in a human subject in need thereof, comprising administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

56. A method of treating, preventing, or delaying the onset and/or progression of glycogen storage disease type 1a (GSD-Ia) in a human subject in need thereof, comprising

administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

57. A method of increasing glucose blood, plasma or serum level in a human subject in need thereof, comprising administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

58. A method of reducing liver glycogen level in a human subject in need thereof, comprising administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

59. A method of reducing liver glucose-6-phosphate (G6P) level in a human subject in need thereof, comprising administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

60. A method of reducing serum triglyceride level in a human subject in need thereof, comprising administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

61. The method of any one of claims 55 to 60, wherein:

(i) the liver glycogen level is reduced at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting as compared to the subject's baseline liver glycogen level while fasting or a reference liver glycogen level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(ii) the liver G6P level is reduced at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% as compared to the subject's baseline liver G6P level while

fasting or a reference liver G6P level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(iii) the liver glycogen level is reduced to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting as compared to a normal liver glycogen level while fasting within at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(iv) the liver G6P level is reduced to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting, as compared to a normal liver G6P level while fasting, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(v) the serum triglyceride level is reduced at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% when the human subject is fasting as compared to the subject's baseline serum triglyceride level while fasting or a reference liver glycogen level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(vi) the serum triglyceride level is reduced to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human subject is fasting as compared to a normal serum triglyceride level while fasting within at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(vii) the blood, plasma or serum glucose liver is increased at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, or at least 100% as compared to the subject's baseline glucose blood, plasma or serum level while fasting or a reference glucose blood, plasma or serum level in a patient with GSD-Ia, for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration;

(viii) the glucose blood, plasma or serum level is increased to at least within 10-fold, at least within 5-fold, at least within 2-fold, or at least within 1.5-fold when the human

subject is fasting as compared to a normal glucose blood, plasma or serum level while fasting within at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; and/or

(ix) the glucose blood, plasma or serum level is increased to at least 70 mg/dl, at least 75 mg/dl , at least 80 mg/dl , at least 85 mg/dl , at least 90 mg/dl , at least 95 mg/dl , at least 100 mg/dl, at least 105 mg/dl, at least 110 mg/dl, at least 115 mg/dl, at least 120 mg/dl, at least 125 mg/dl, at least 130 mg/dl, at least 135 mg/dl, at least 140 mg/dl, at least 145 mg/dl, or at least 150 mg/dl in the subject when fasting for at least 12 hours, at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks post-administration.

62. A method of increasing G6PC activity in a human subject in need thereof, comprising administering to the subject an effective amount of the pharmaceutical composition of any one of claims 1-24, 53, or 54 or the polynucleotide of any one of claims 25-52.

63. The method of any one of claims 55 to 62, wherein:

(i) the level of G6PC activity in the subject is increased at least 1.5-fold, at least 2-fold, at least 5-fold, at least 10-fold, at least 20-fold, or at least 50-fold as compared to a reference G6PC activity level in a subject having GSD-Ia for at least 24 hours, at least 48 hours, at least 72 hours, at least 96 hours, at least 120 hours, 6 days, 1 week, 8 days, 9 days, 10 days, 11 days, 12 days, 13 days, or 2 weeks after a single administration; and/or

(ii) 12 hours after a single administration of the pharmaceutical composition or polynucleotide is administered to the subject, the G6PC activity in the subject is increased at least 2%, at least 5%, at least 10%, at least 20%, at least 30%, at least 40%, at least 50%, at least 60%, at least 70%, at least 80%, at least 90%, at least 100%, at least 150%, at least 200%, at least 300%, at least 400%, at least 500%, or at least 600% compared to the subject's baseline G6PC activity.

64. The method of claim 62 or 63, wherein the G6PC activity is increased in the liver or blood of the subject.

65. The method of any one of claims 55-64, wherein the human subject has been fasting.

66. The method of claim 65, wherein the human subject has been fasting for at least 2 hours.

67. The method of any one of claims 55 to 66, comprising multiple administrations of the pharmaceutical composition or the polynucleotide to the human subject.

68. The method of claim 67, wherein the multiple administrations to the subject are about once a week, about once every two weeks, or about once a month.

69. The method of any one of claims 55 to 68, wherein the pharmaceutical composition or polynucleotide is administered intravenously.

FIGURE 1A

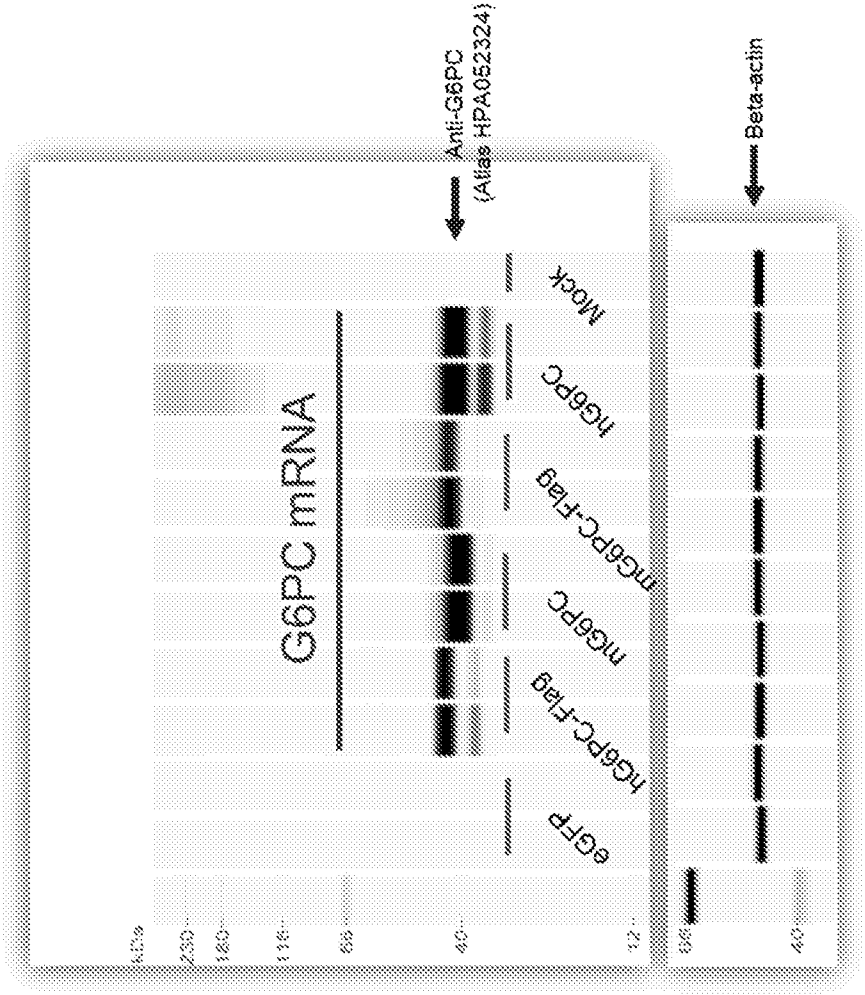


FIGURE 1B

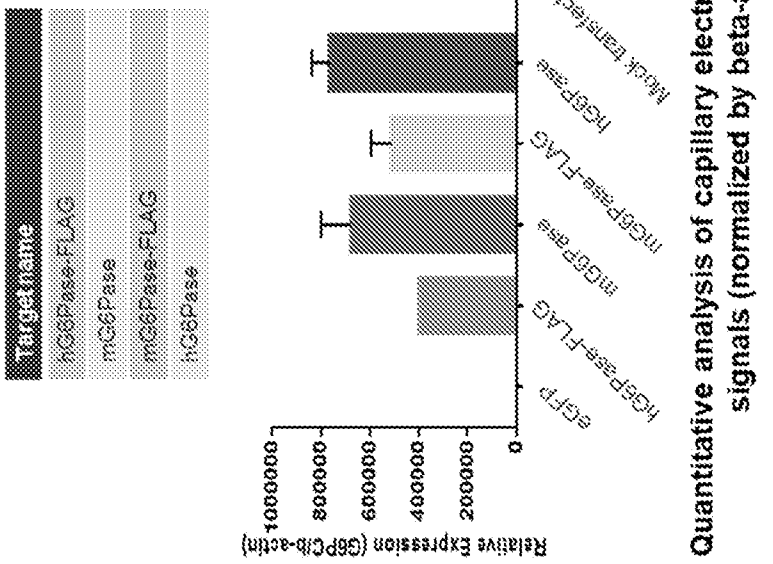


FIGURE 2A

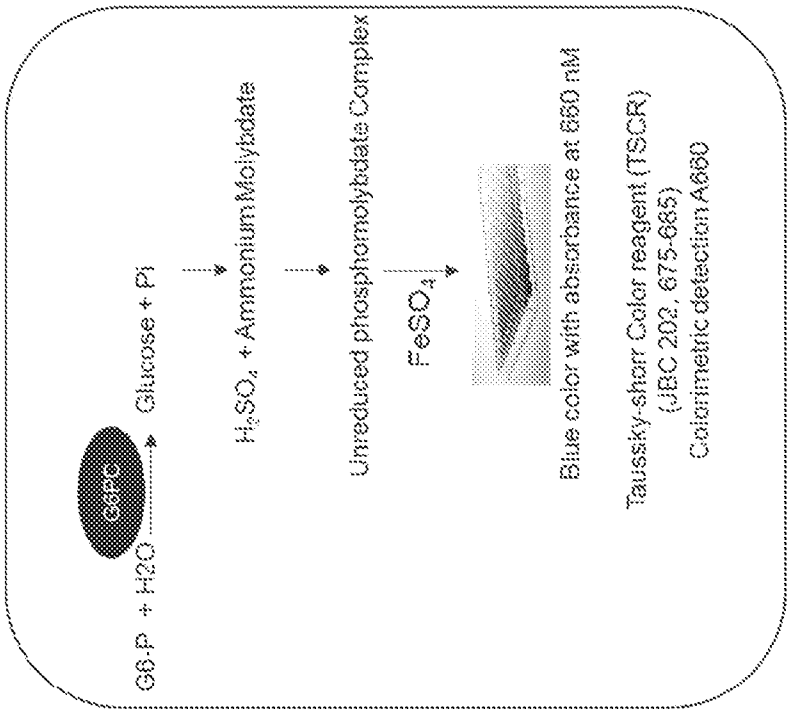


FIGURE 2B

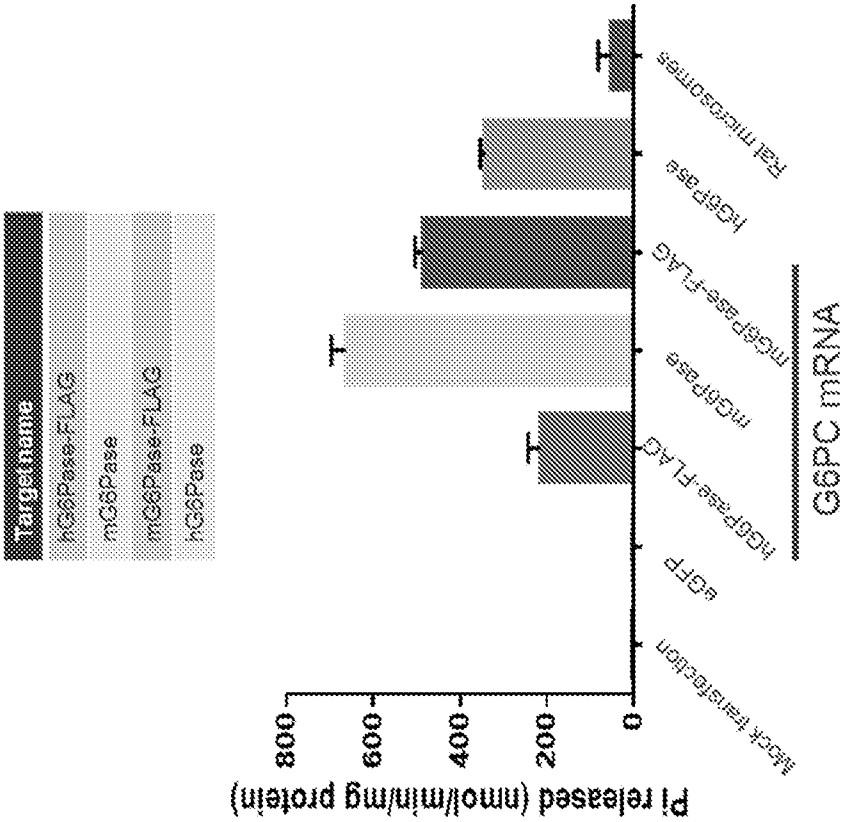


FIGURE 3B

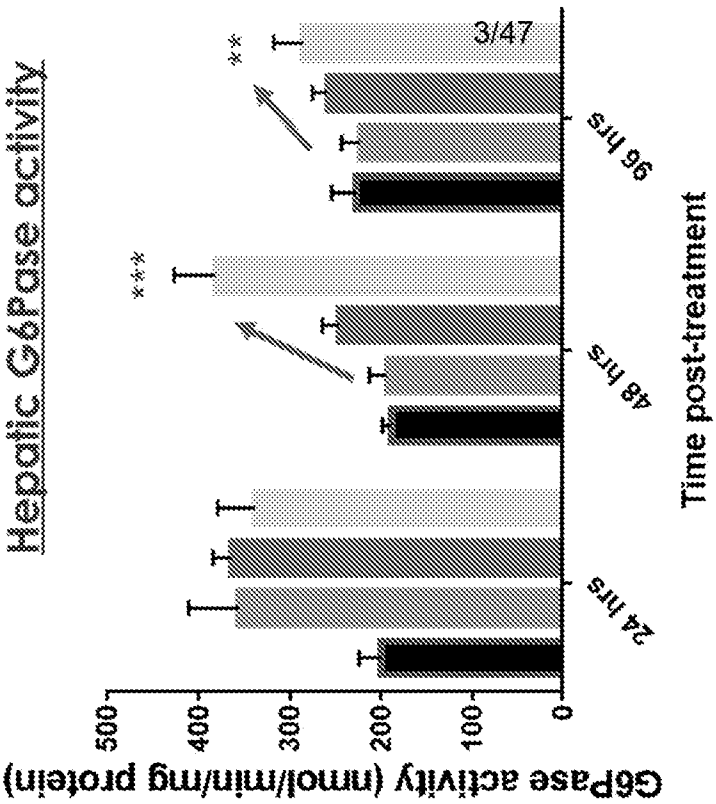


FIGURE 3A

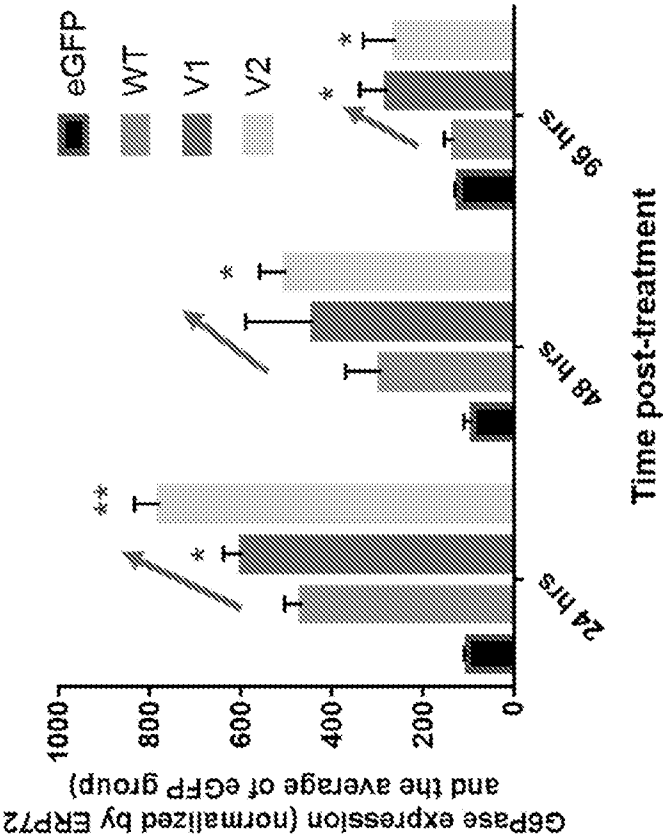


FIGURE 4

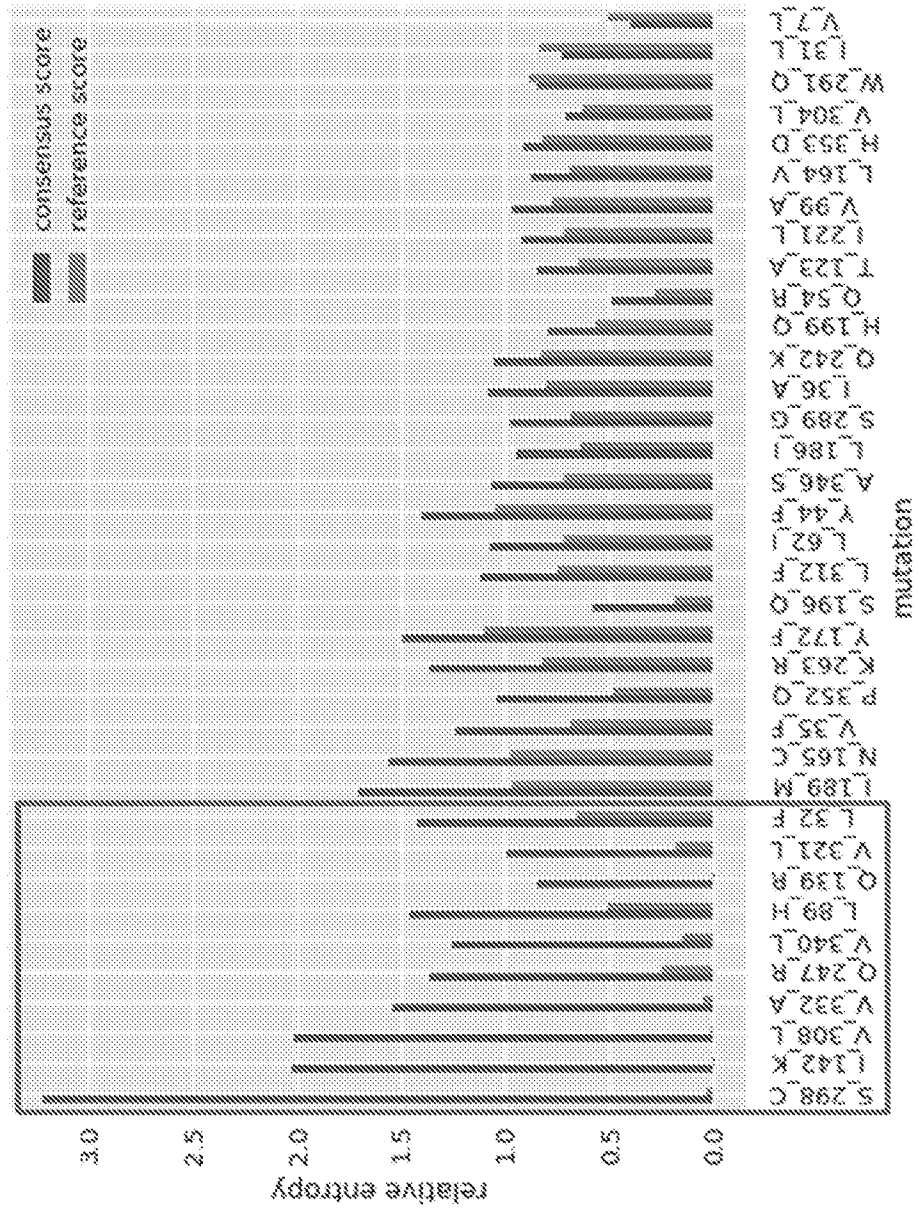

consensus score
reference score
relative entropy
3.0
2.5
2.0
1.5
1.0
0.5
0.0
mutation
S_298_C
L_142_K
V_308_L
V_332_A
Q_247_R
V_340_L
L_89_H
Q_139_R
V_321_L
L_32_F
L_189_M
N_165_C
V_35_F
P_352_Q
K_263_R
V_172_F
S_196_Q
L_312_F
L_62_I
Y_44_F
A_346_S
L_186_I
S_289_G
I_36_A
Q_242_K
H_199_Q
Q_54_R
T_123_A
I_221_L
Y_99_A
L_164_V
H_353_Q
V_304_L
W_291_Q
L_31_L
V_7_L

FIGURE 5A

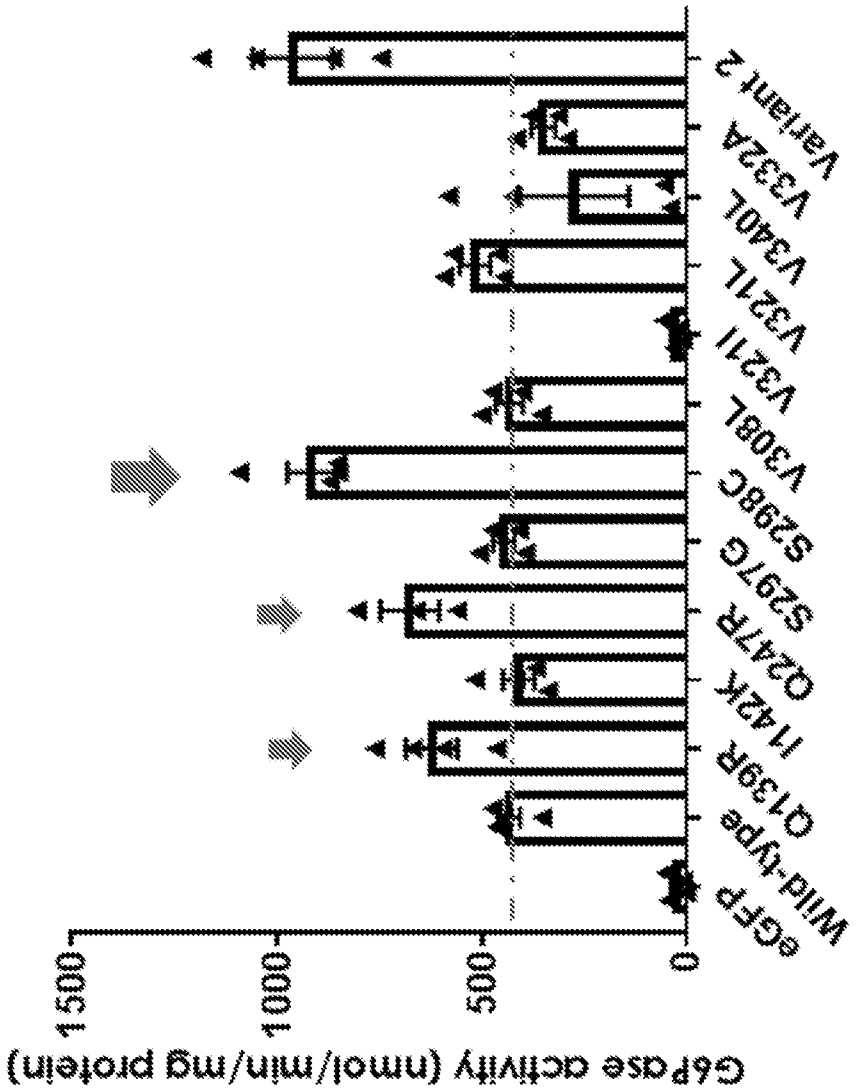

G6Pase activity (nmol/min/mg protein)
1500
1000
500
0
eGFP
Wild-type
Q139R
I142K
Q247R
S297G
S298C
V308L
V321I
V321L
V340L
V332A
Variant 2

FIGURE 5C

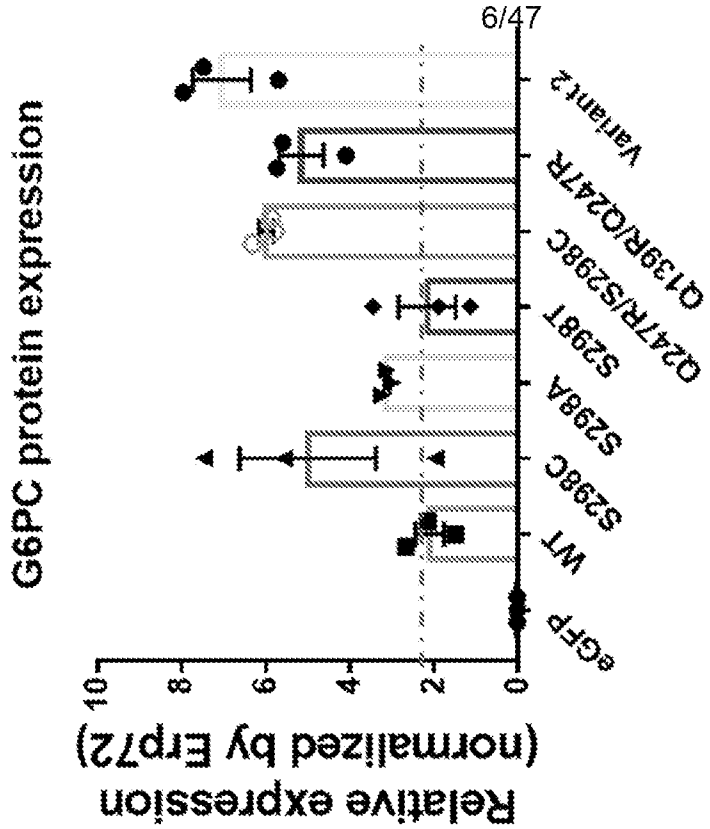


FIGURE 5B

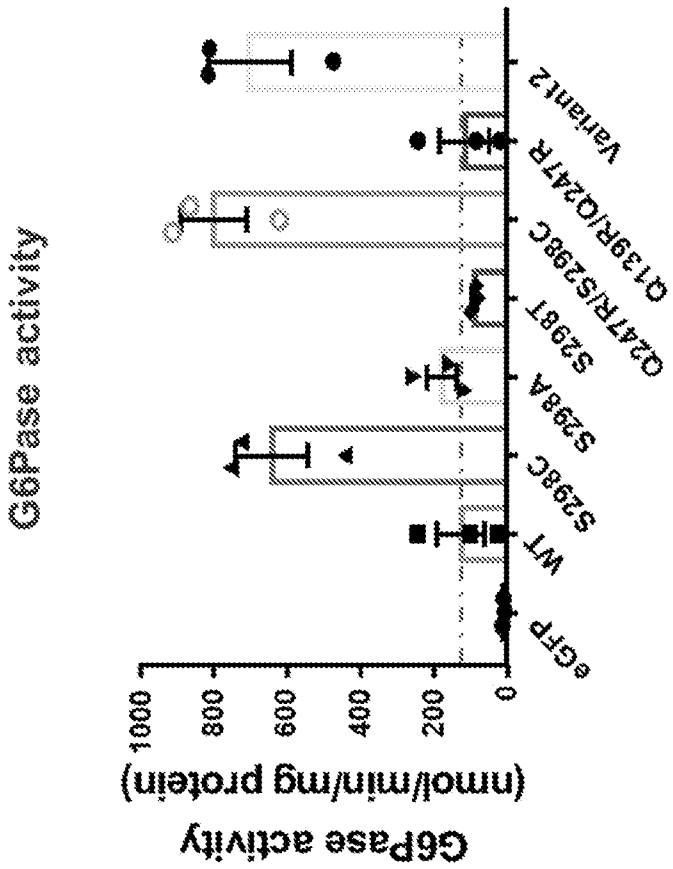


FIGURE 5E

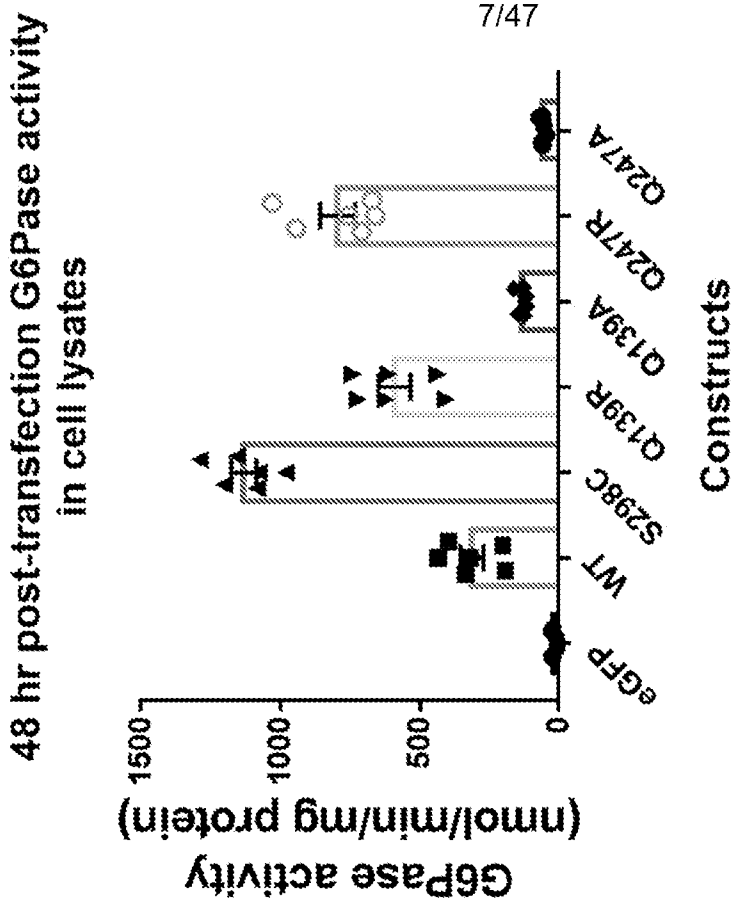


FIGURE 5D

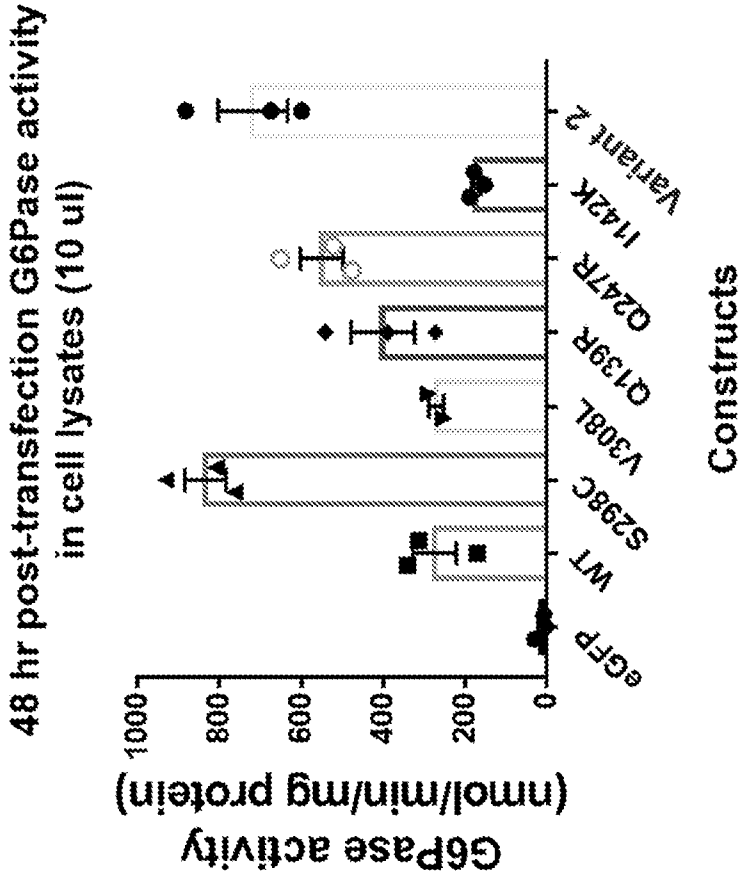


FIGURE 5G

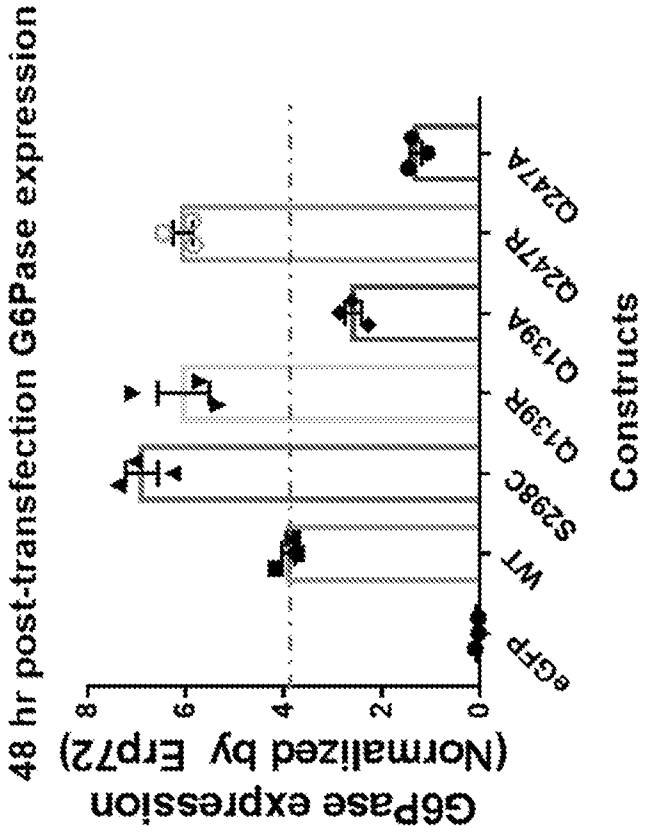


FIGURE 5F

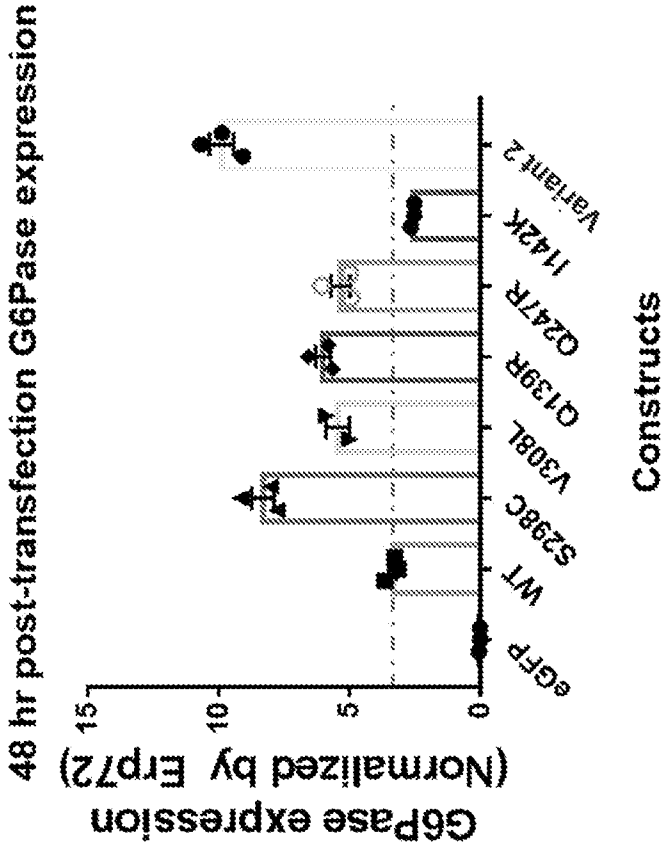


FIGURE 6A

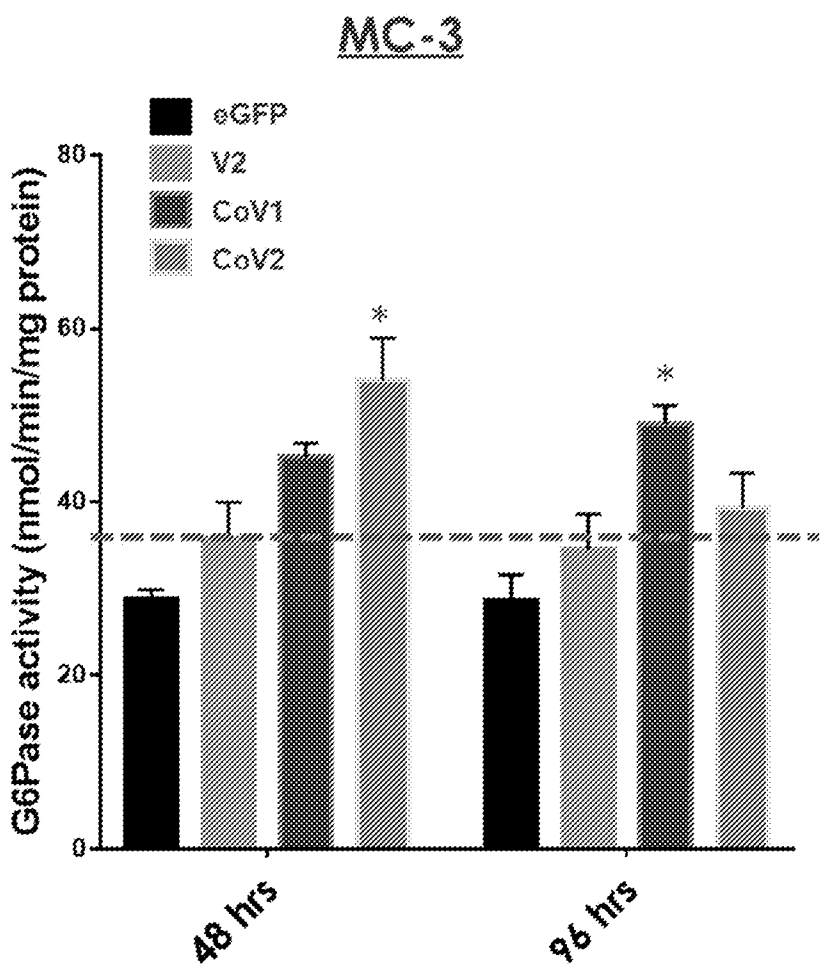
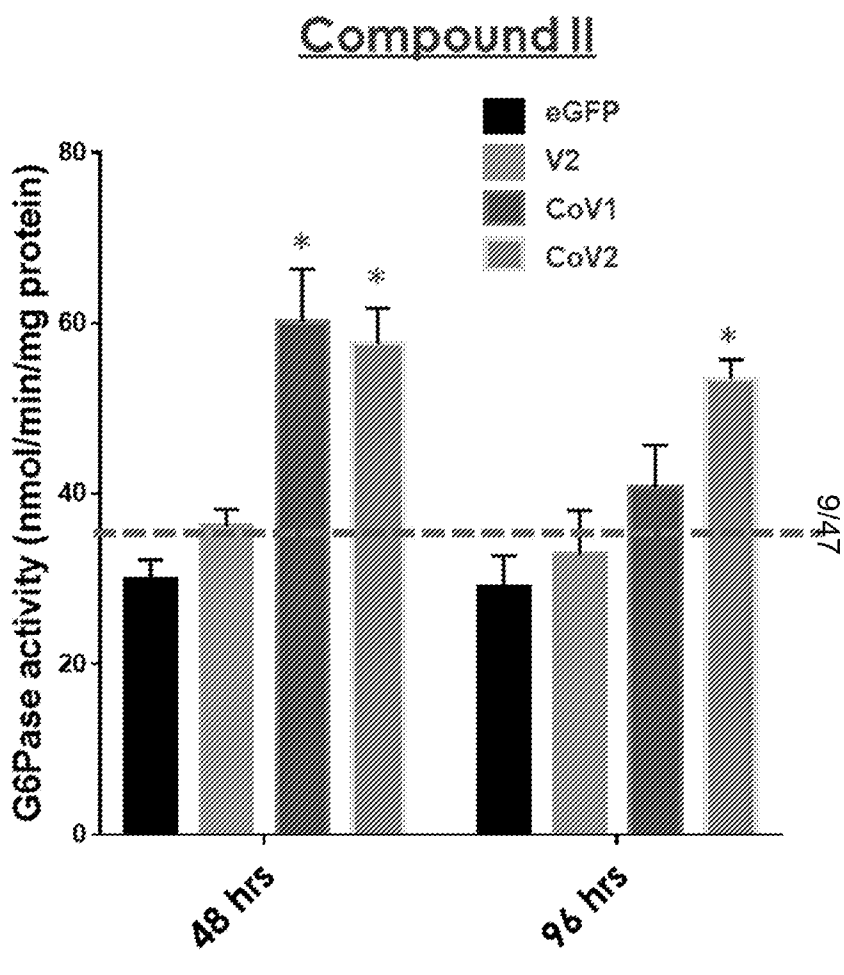


FIGURE 6B



* P<0.05 vs Variant 2 one-way ANOVA followed by Dunnett's test, within each time point

FIGURE 6D

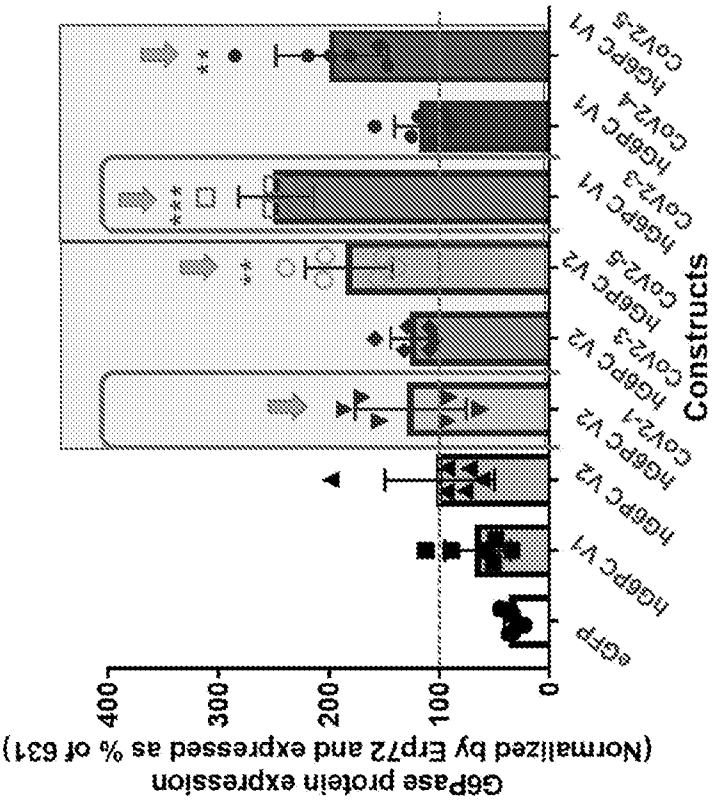


FIGURE 6C

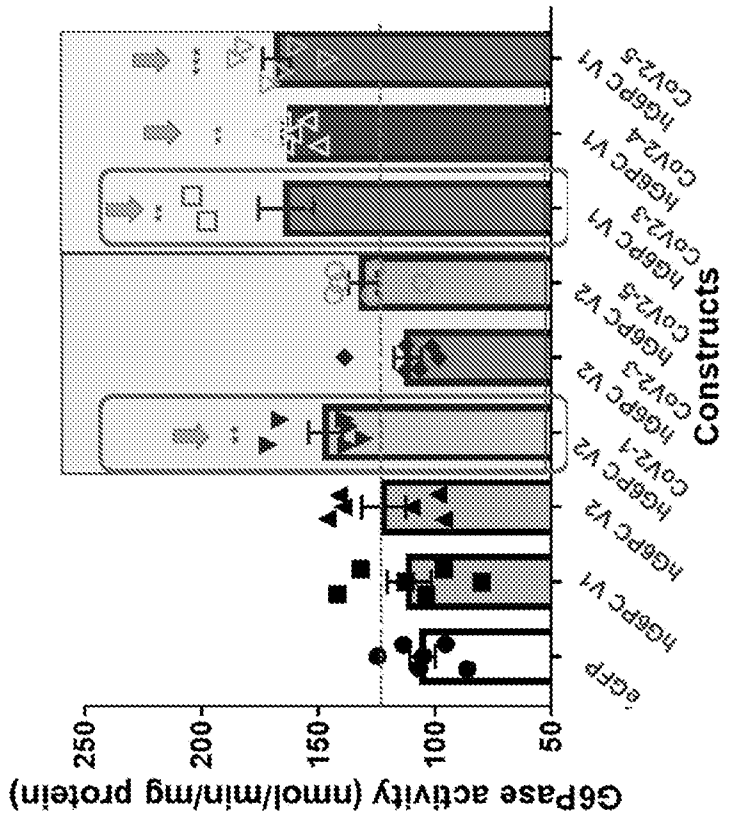


FIGURE 7B

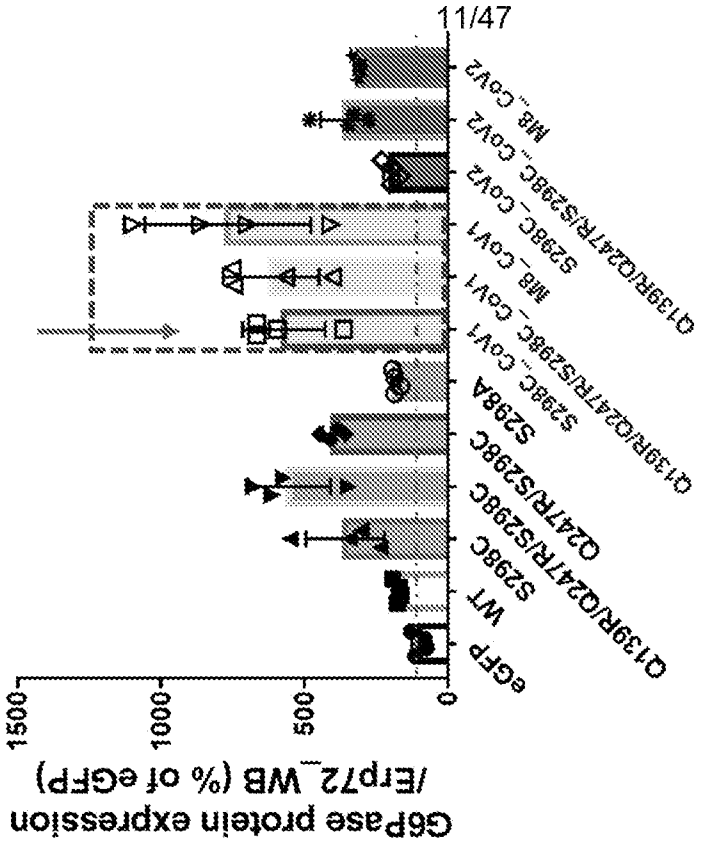


FIGURE 7A

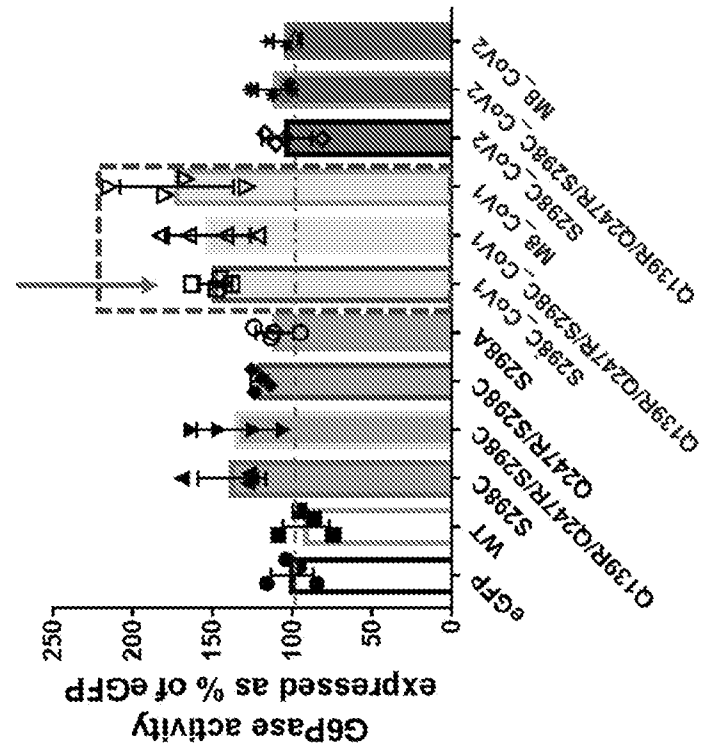


FIGURE 8A

Day 1

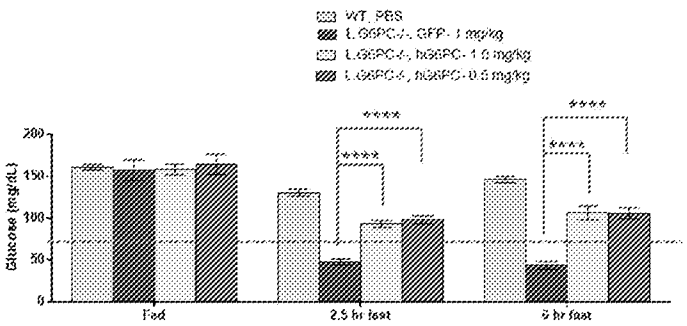


FIGURE 8B

Day 3

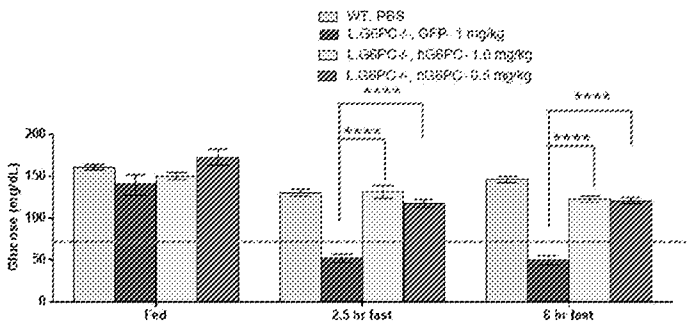


FIGURE 8C

Day 5

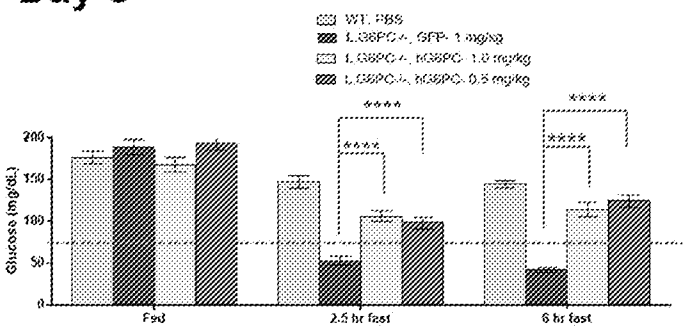
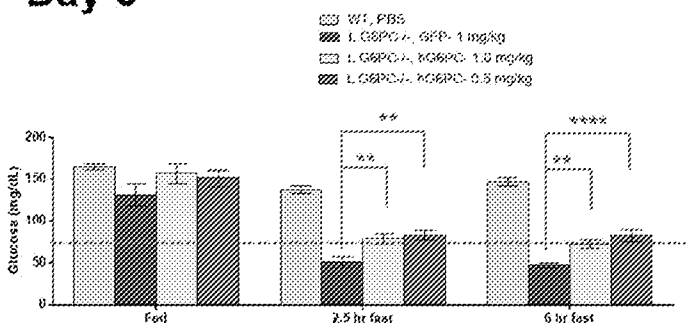


FIGURE 8D

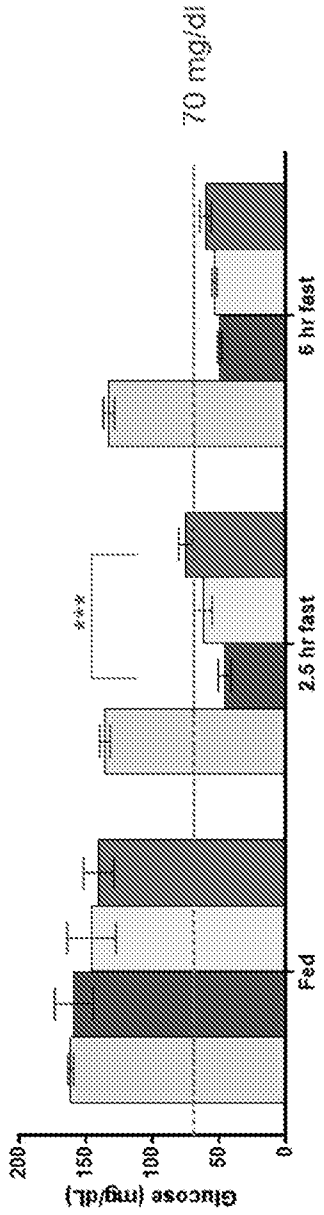
Day 8



**P<0.01
****P<0.0001
vs. GFP
one-way ANOVA
followed by multiple
comparison

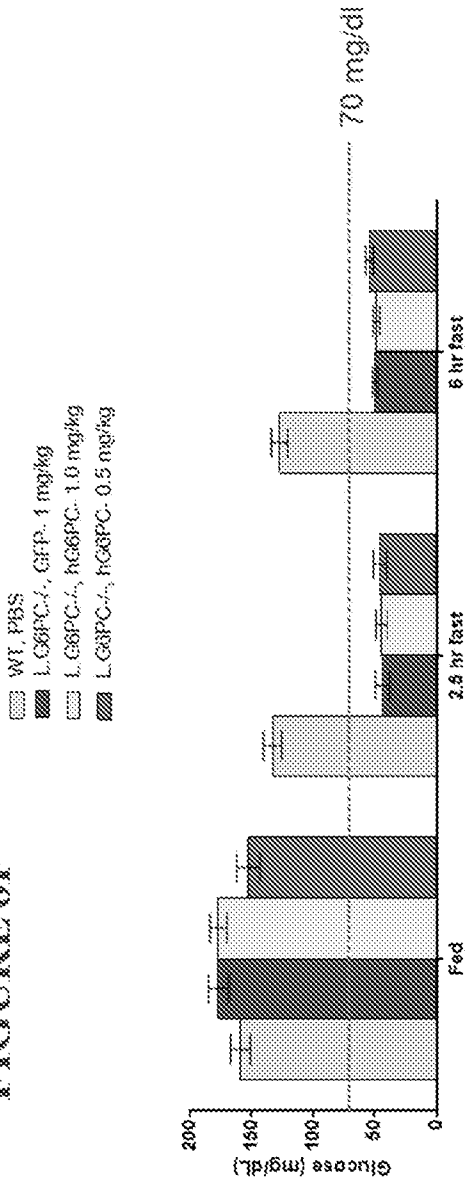
70 mg/dl

FIGURE 8E



Day 11

FIGURE 8F



Day 15

***P<0.001
vs. GFP
one-Way ANOVA
followed by multiple
comparisons

FIGURE 8G

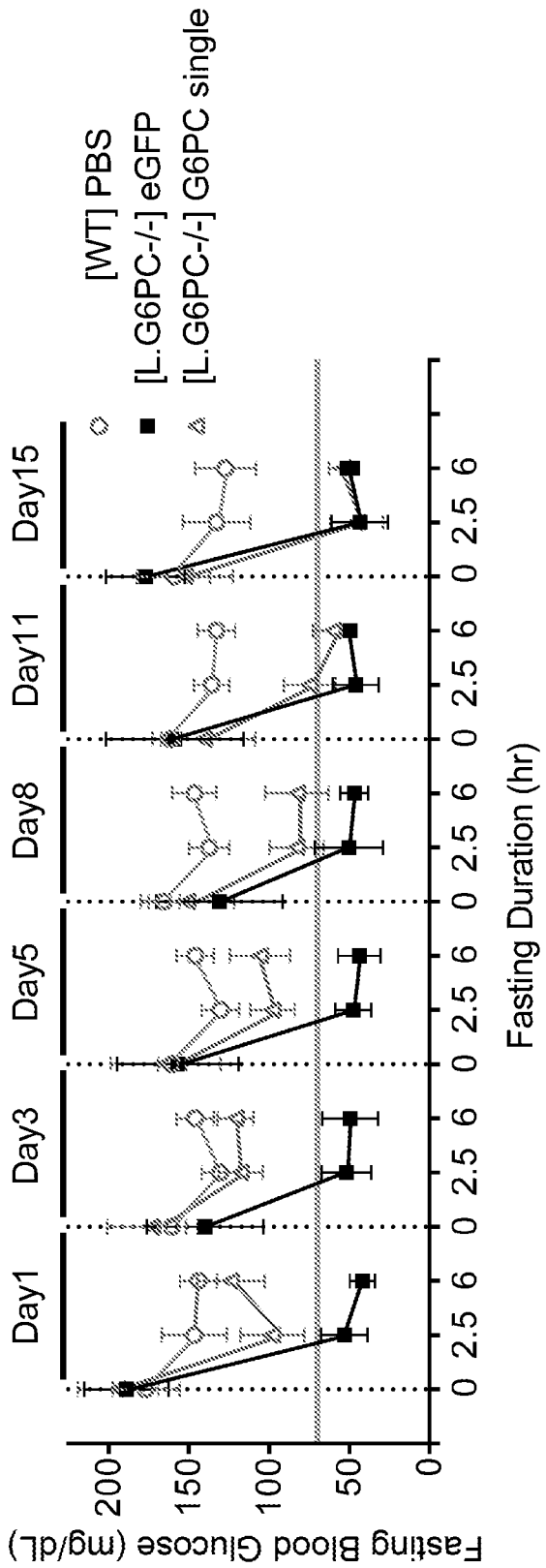

[WT] PBS
[L.G6PC-/-] eGFP
[L.G6PC-/-] G6PC single
Day1
Day3
Day5
Day8
Day11
Day15
0 2.5 6
Fasting Duration (hr)
Fasting Blood Glucose (mg/dL)
200
150
100
50
0

FIGURE 8H

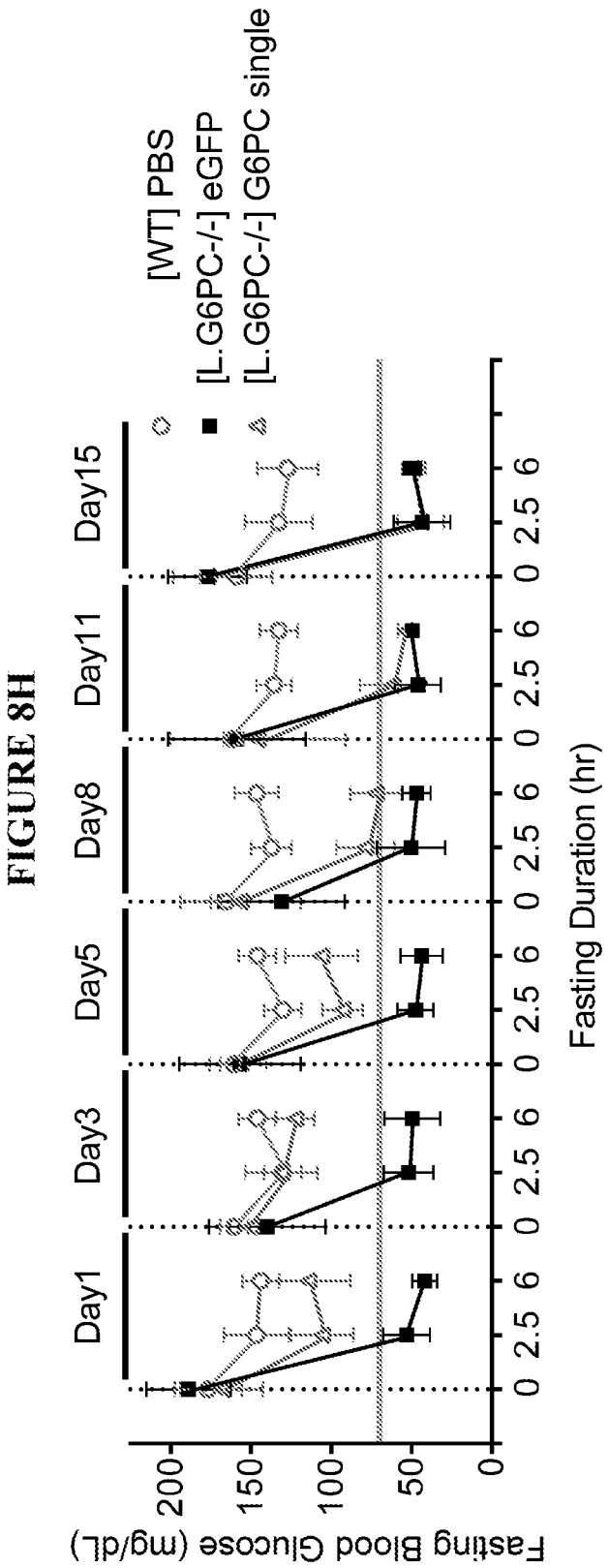

[WT] PBS
[L.G6PC-/-] eGFP
[L.G6PC-/-] G6PC single
Day1
Day3
Day5
Day8
Day11
Day15
0 2.5 6
Fasting Duration (hr)
Fasting Blood Glucose (mg/dL)
200
150
100
50
0

FIGURE 9A

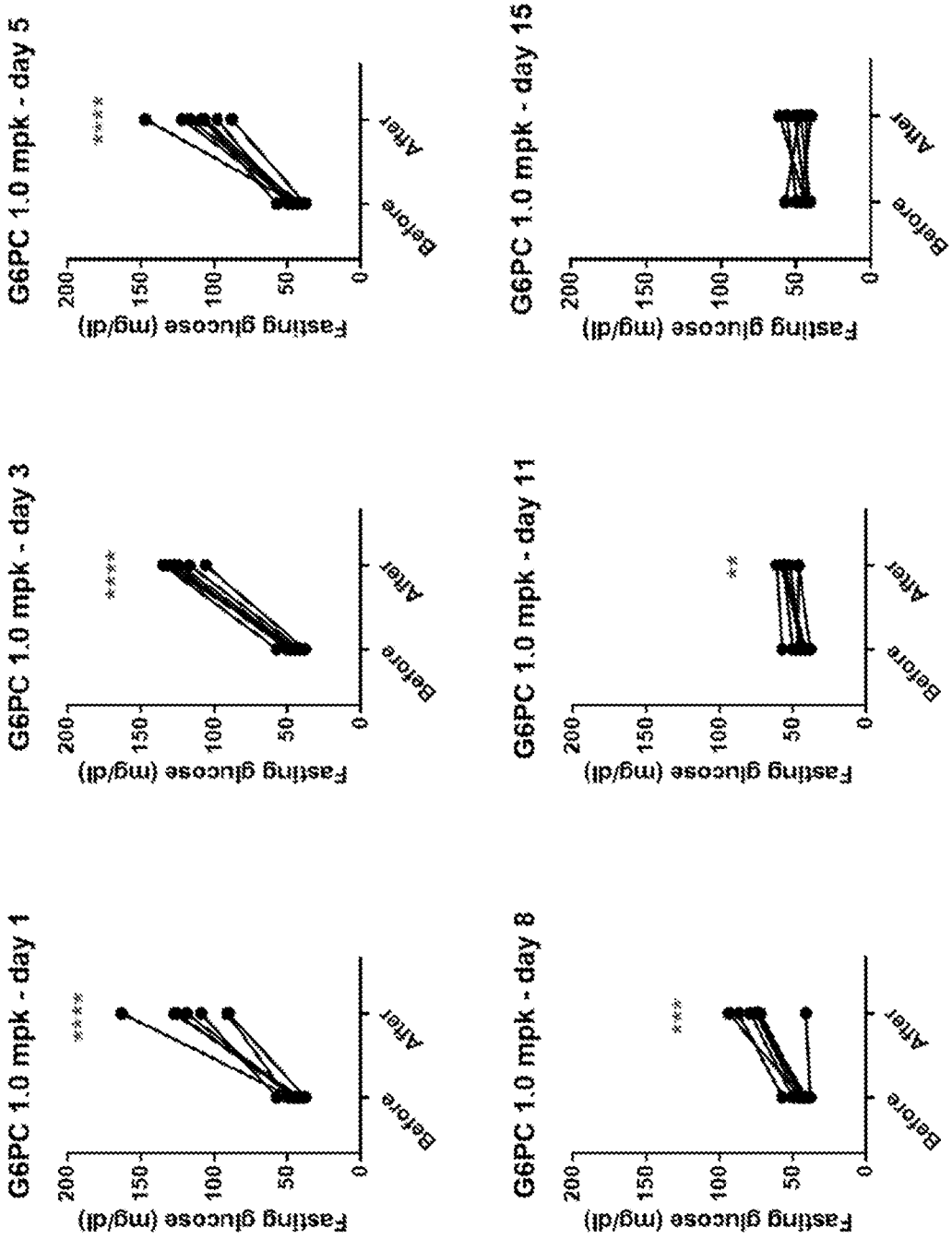

G6PC 1.0 mpk - day 1
G6PC 1.0 mpk - day 3
G6PC 1.0 mpk - day 5
G6PC 1.0 mpk - day 8
G6PC 1.0 mpk - day 11
G6PC 1.0 mpk - day 15
Fasting glucose (mg/dl)
200
150
100
50
0
Before
After
****
***
**

FIGURE 9B

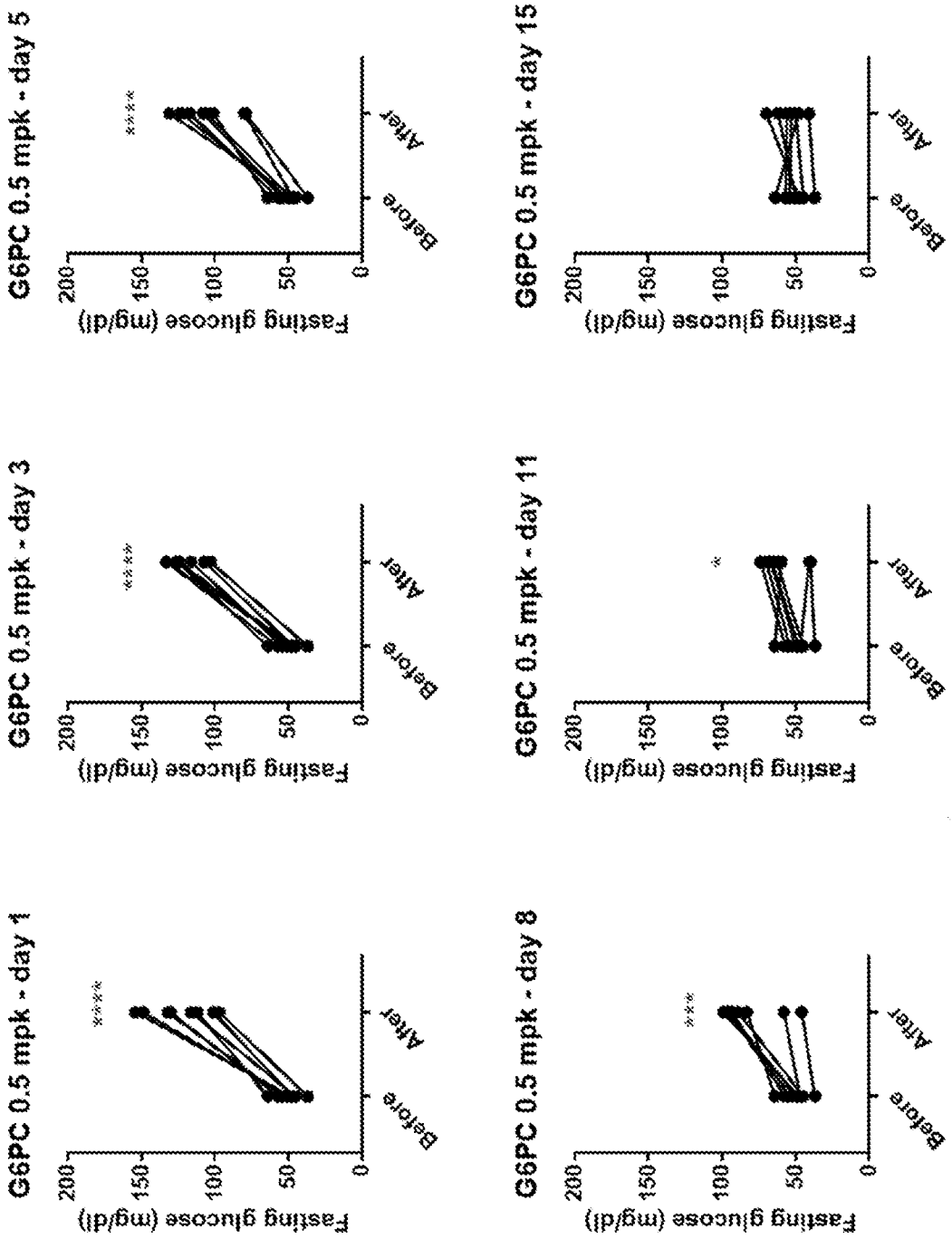

G6PC 0.5 mpk - day 1
G6PC 0.5 mpk - day 3
G6PC 0.5 mpk - day 5
G6PC 0.5 mpk - day 8
G6PC 0.5 mpk - day 11
G6PC 0.5 mpk - day 15
Fasting glucose (mg/dl)
200
150
100
50
0
Before
After
****
***
*

FIGURE 9C

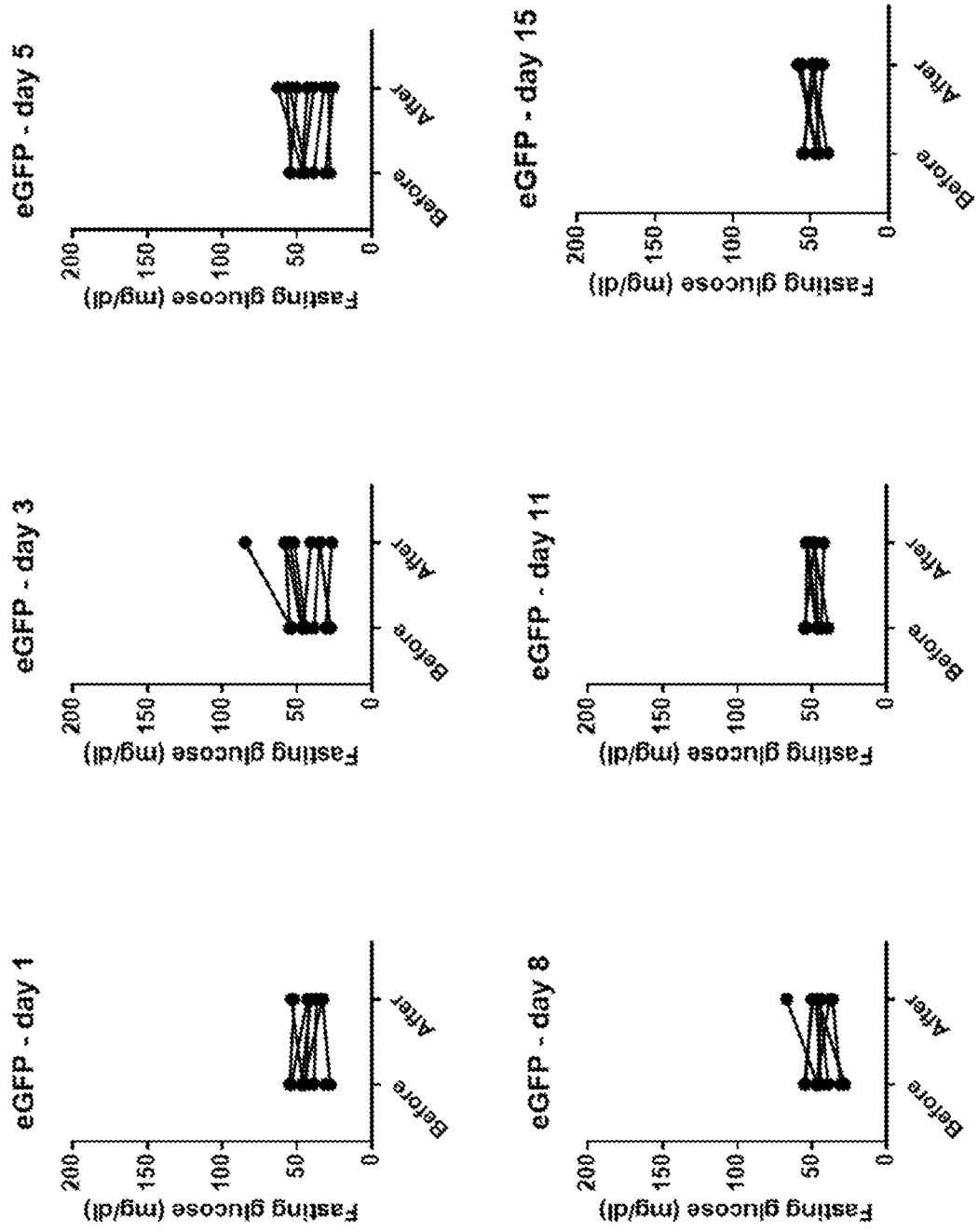

eGFP - day 1
eGFP - day 3
eGFP - day 5
eGFP - day 8
eGFP - day 11
eGFP - day 15
Fasting glucose (mg/dl)
0
50
100
150
200
Before
After

FIGURE 10A

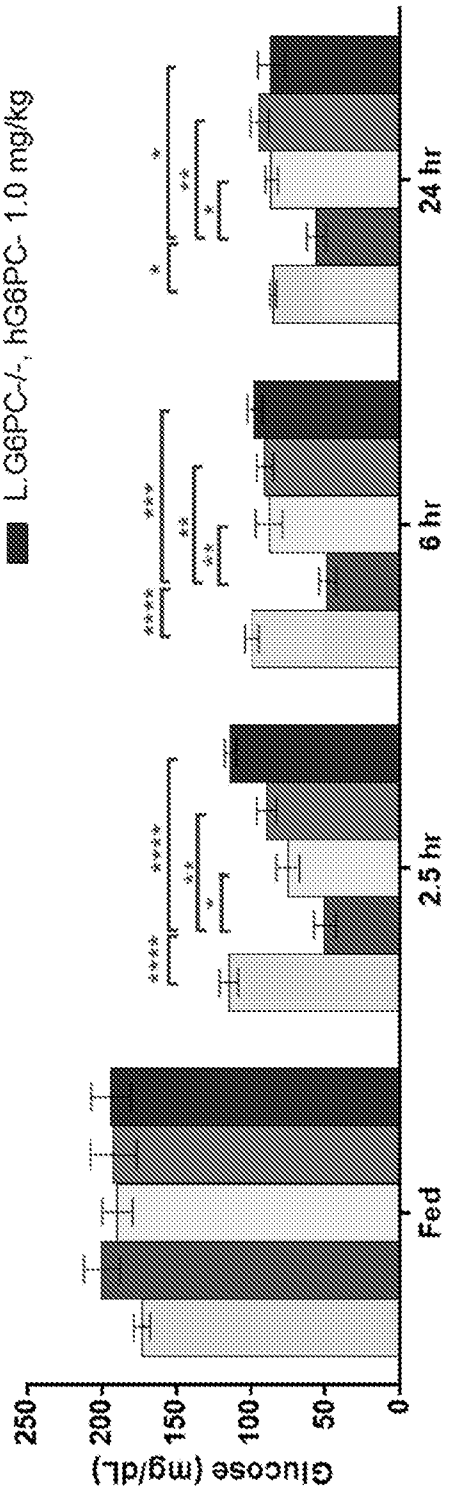

WT, PBS
L.G6PC-/-, GFP- 1 mg/kg
L.G6PC-/-, hG6PC- 0.2 mg/kg
L.G6PC-/-, hG6PC- 0.5 mg/kg
L.G6PC-/-, hG6PC- 1.0 mg/kg
Glucose (mg/dL)
250
200
150
100
50
0
Fed
2.5 hr
6 hr
24 hr

FIGURE 10B

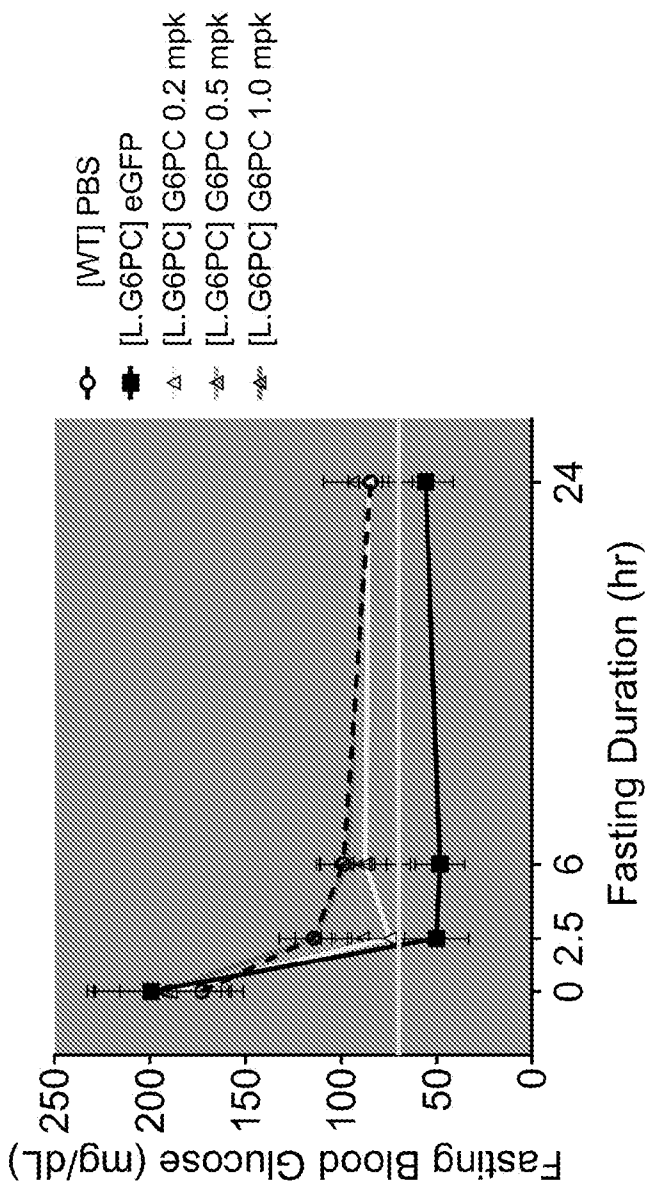

[WT] PBS
[L.G6PC] eGFP
[L.G6PC] G6PC 0.2 mpk
[L.G6PC] G6PC 0.5 mpk
[L.G6PC] G6PC 1.0 mpk
Fasting Blood Glucose (mg/dL)
250
200
150
100
50
0
0 2.5 6 24
Fasting Duration (hr)

FIGURE 10C

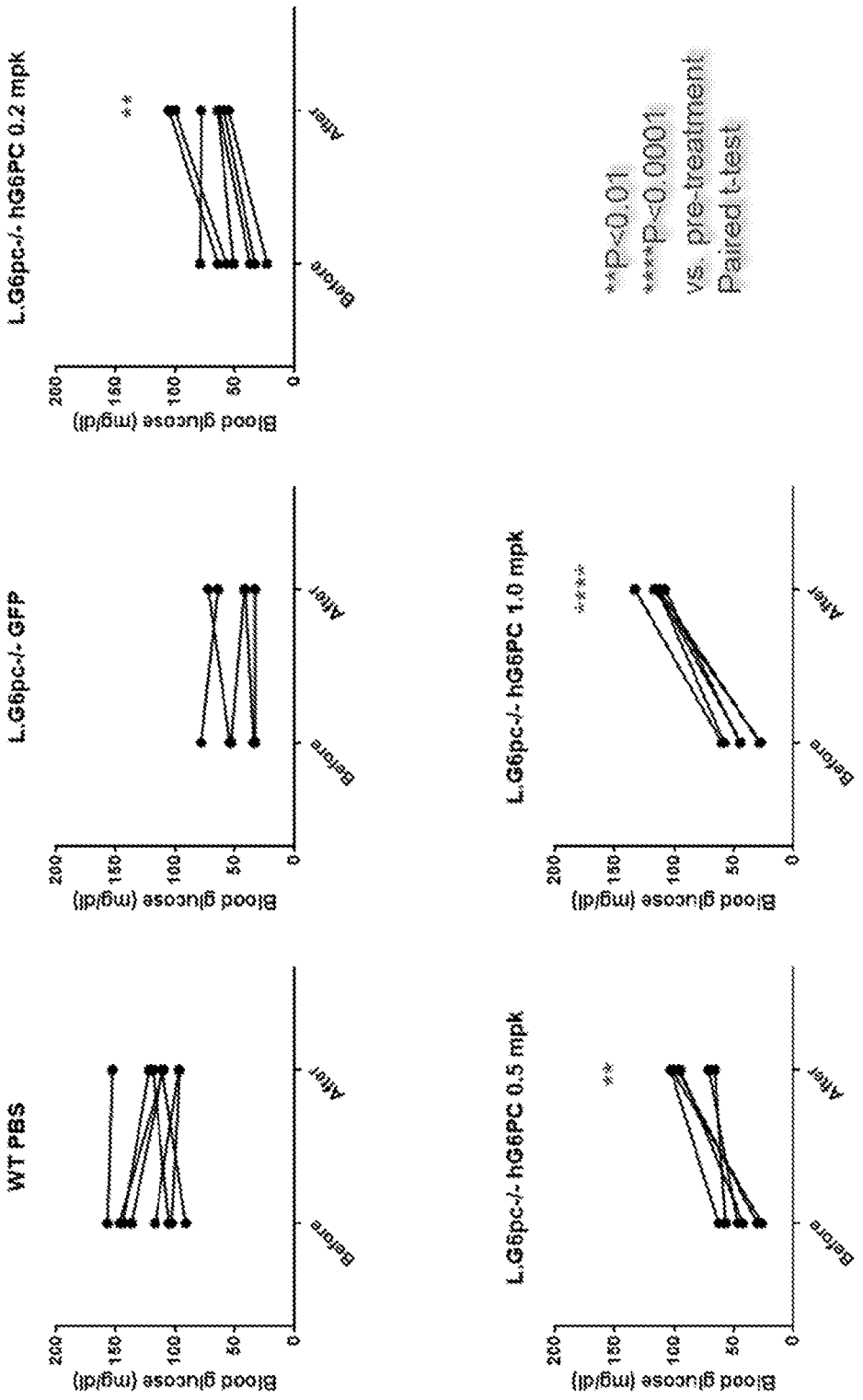

WT PBS
L.G6pc-/- GFP
L.G6pc-/- hG6PC 0.2 mpk
L.G6pc-/- hG6PC 0.5 mpk
L.G6pc-/- hG6PC 1.0 mpk
Blood glucose (mg/dl)
200
150
100
50
0
Before
After
**
***
****
**P<0.01
***P<0.0001
vs. pre-treatment
Paired t-test

FIGURE 11A

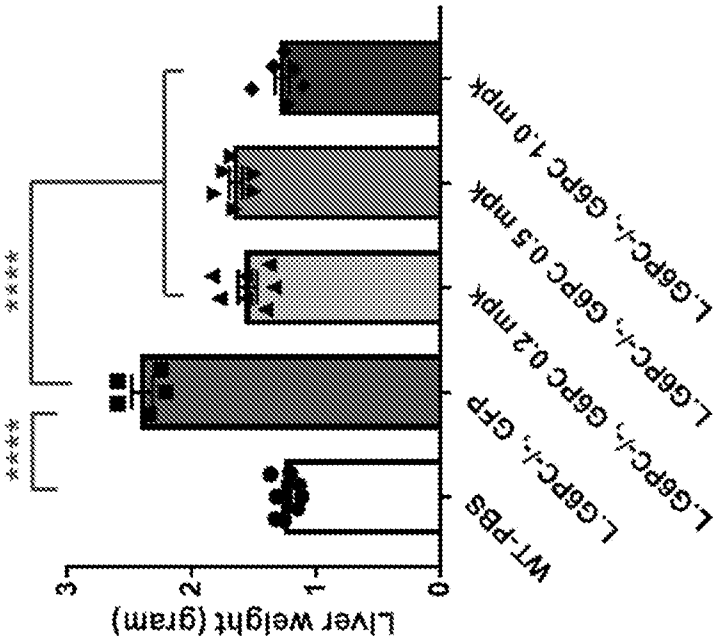
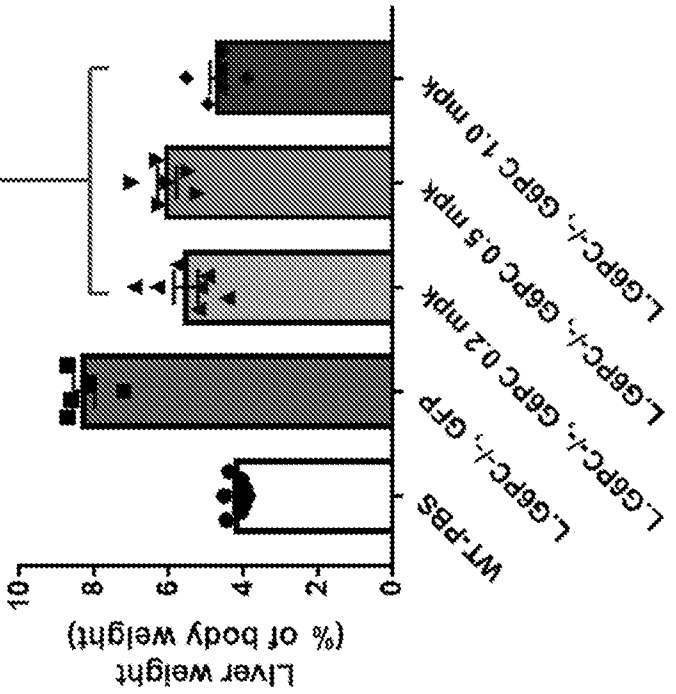


FIGURE 11B



****$p<0.0001$ vs. GFP
One-way ANOVA followed by multiple comparisons

FIGURE 12

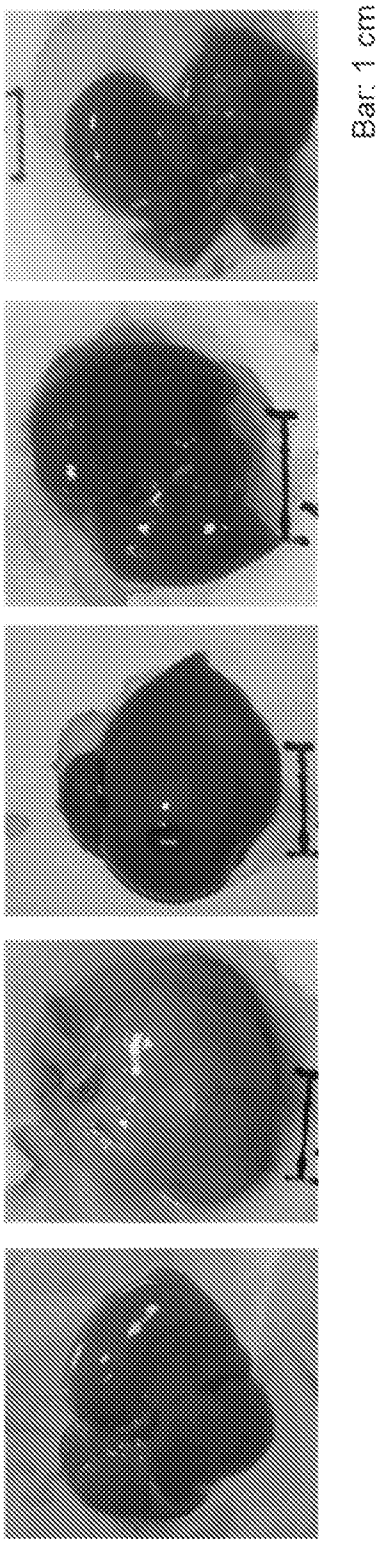

Wild-Type
PBS
L-G6pc-/-
GFP-mRNA 1.0 mpk
G6PC-mRNA 0.2 mpk
G6PC-mRNA 0.5 mpk
G6PC-mRNA 1.0 mpk
Bar: 1 cm

FIGURE 13A

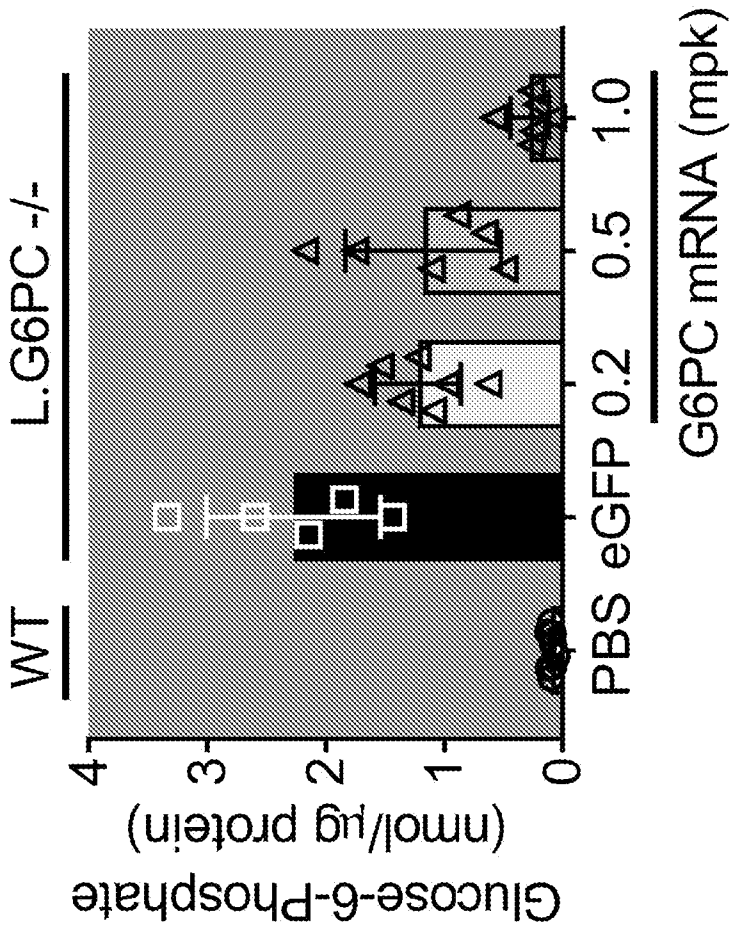

WT
L.G6PC -/-
Glucose-6-Phosphate
(nmol/µg protein)
0
1
2
3
4
PBS
eGFP 0.2
0.5
1.0
G6PC mRNA (mpk)

FIGURE 13B

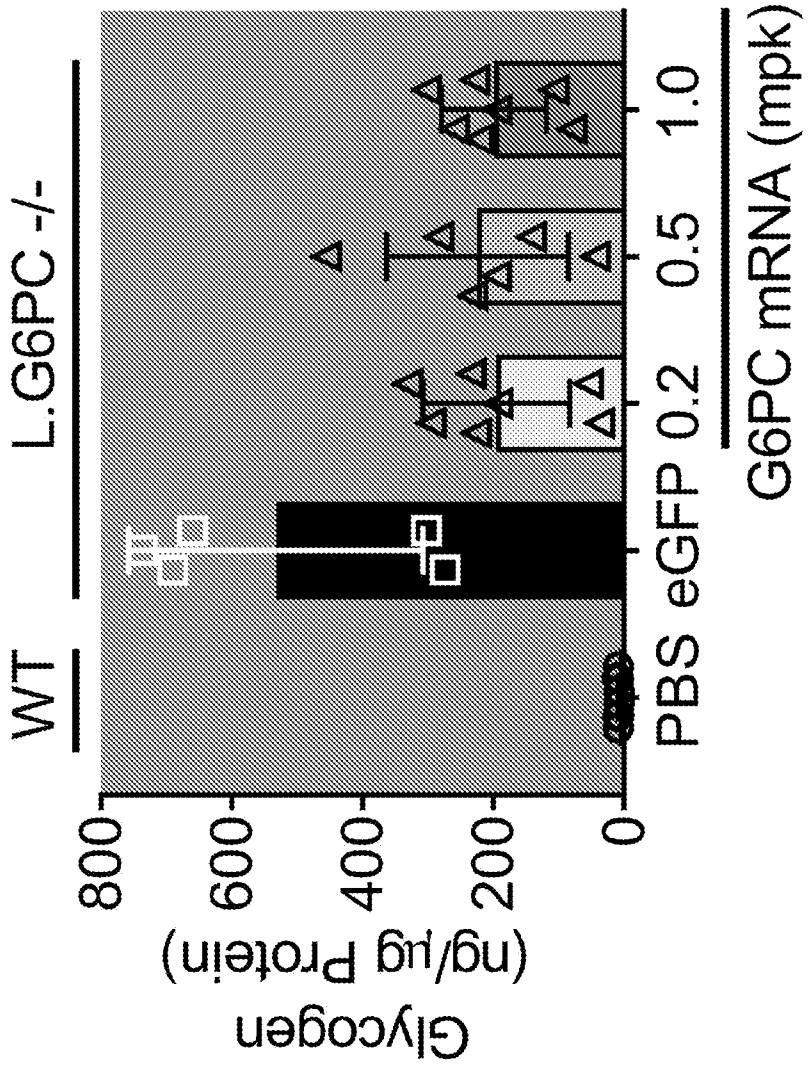

WT
L.G6PC -/-
Glycogen
(ng/μg Protein)
800
600
400
200
0
PBS
eGFP
0.2
0.5
1.0
G6PC mRNA (mpk)

FIGURE 13C

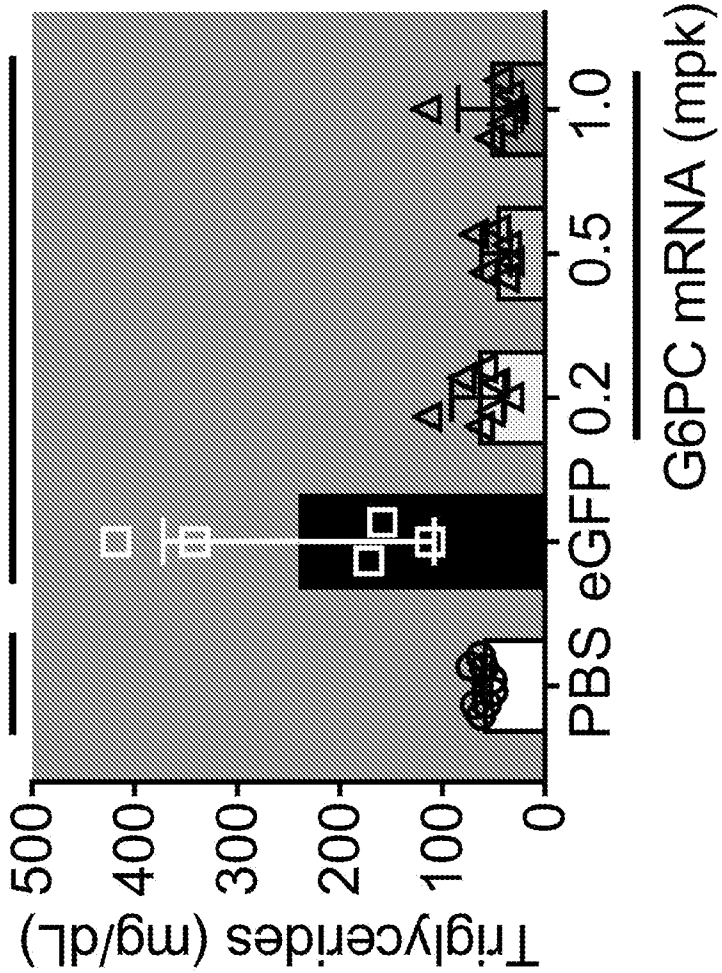

WT
L.G6PC -/-
Triglycerides (mg/dL)
500
400
300
200
100
0
PBS
eGFP
0.2
0.5
1.0
G6PC mRNA (mpk)

FIGURE 13D

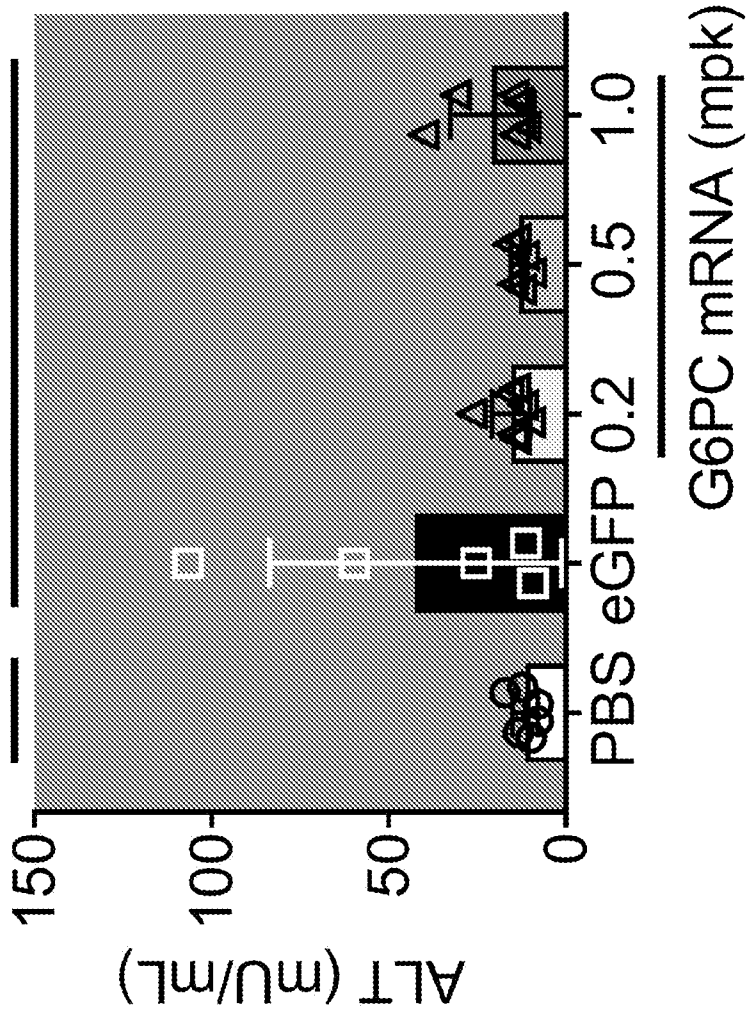


FIGURE 13E

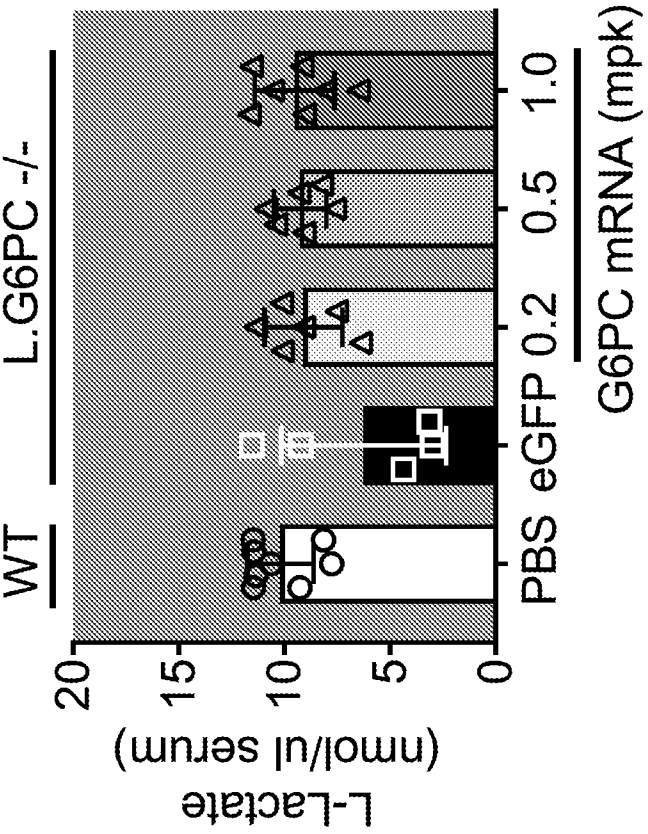

WT
L.G6PC -/-
L-Lactate
(nmol/ul serum)
20
15
10
5
0
PBS
eGFP 0.2
0.5
1.0
G6PC mRNA (mpk)

FIGURE 13F

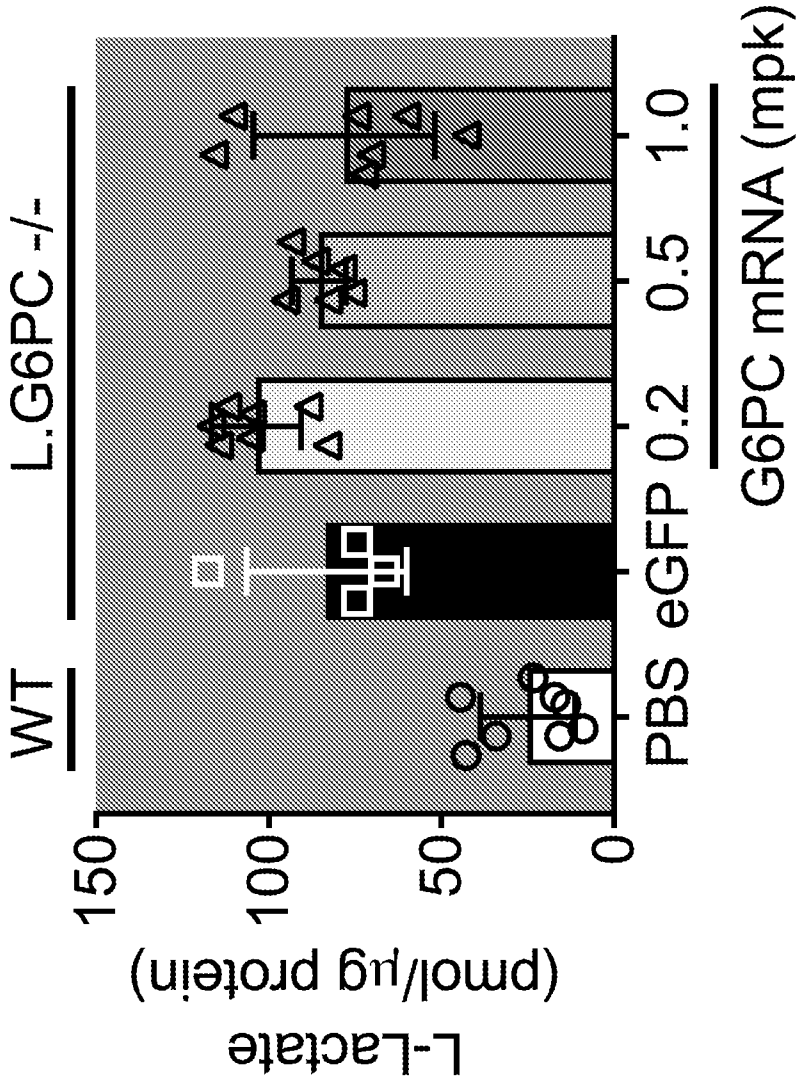

WT
L.G6PC -/-
L-Lactate
(pmol/μg protein)
150
100
50
0
PBS
eGFP
0.2
0.5
1.0
G6PC mRNA (mpk)

FIGURE 14A

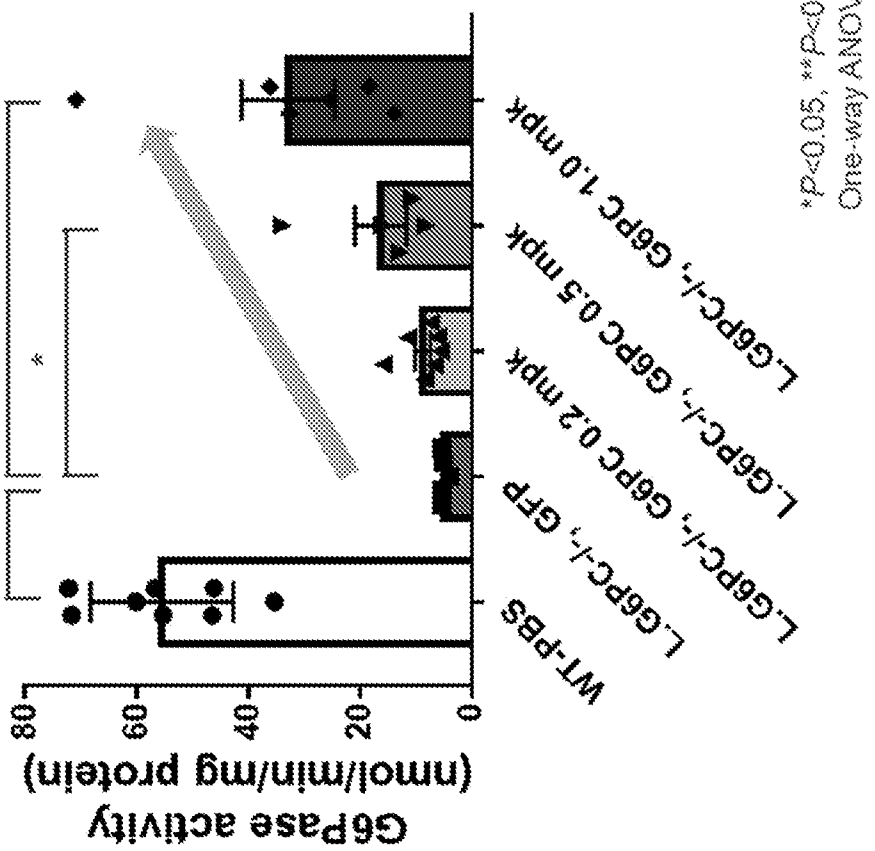

G6Pase activity
(nmol/min/mg protein)
0
20
40
60
80
****
**
*
WT-PBS
L-G6PC-/- GFP
L-G6PC-/- G6PC 0.2 mpk
L-G6PC-/- G6PC 0.5 mpk
L-G6PC-/- G6PC 1.0 mpk
*P<0.05, **P<0.01, ****P<0.0001 vs. GFP
One-way ANOVA followed by multiple comparisons

FIGURE 14B

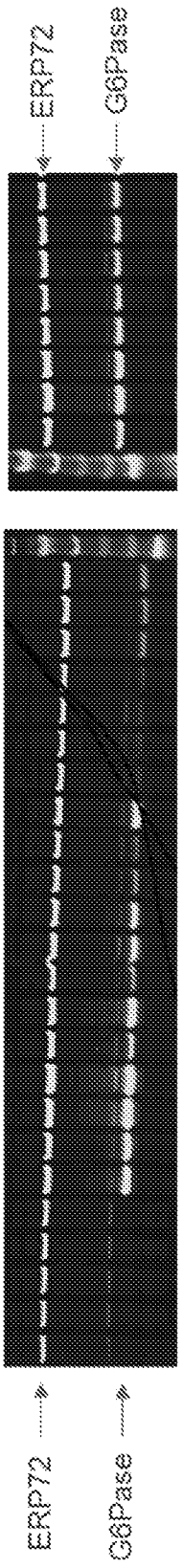

L-G6pc-/-
GFP-mRNA 1.0 mpk
G6PC-mRNA 1.0 mpk
G6PC-mRNA 0.5 mpk
G6PC-mRNA 0.2 mpk
ERP72
G6Pase
Wild-Type
PBS
ERP72
G6Pase

FIGURE 14C

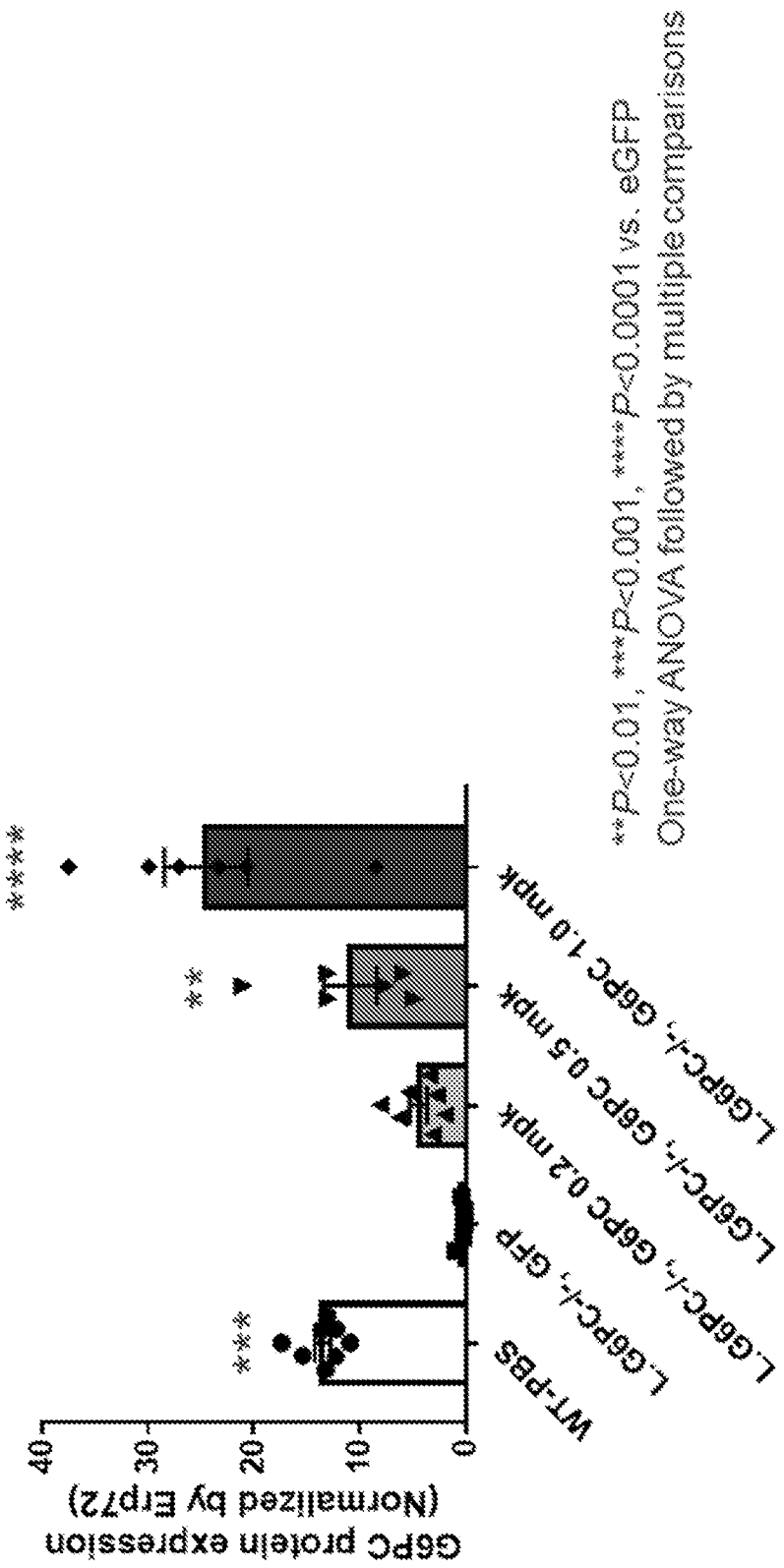

G6PC protein expression
(Normalized by Erp72)
0
10
20
30
40
WT-PBS
L-G6PC-/-, GFP
L-G6PC-/-, G6PC 0.2 mpk
L-G6PC-/-, G6PC 0.5 mpk
L-G6PC-/-, G6PC 1.0 mpk
***
**
****
**P<0.01, ***P<0.001, ****P<0.0001 vs. eGFP
One-way ANOVA followed by multiple comparisons

FIGURE 15A

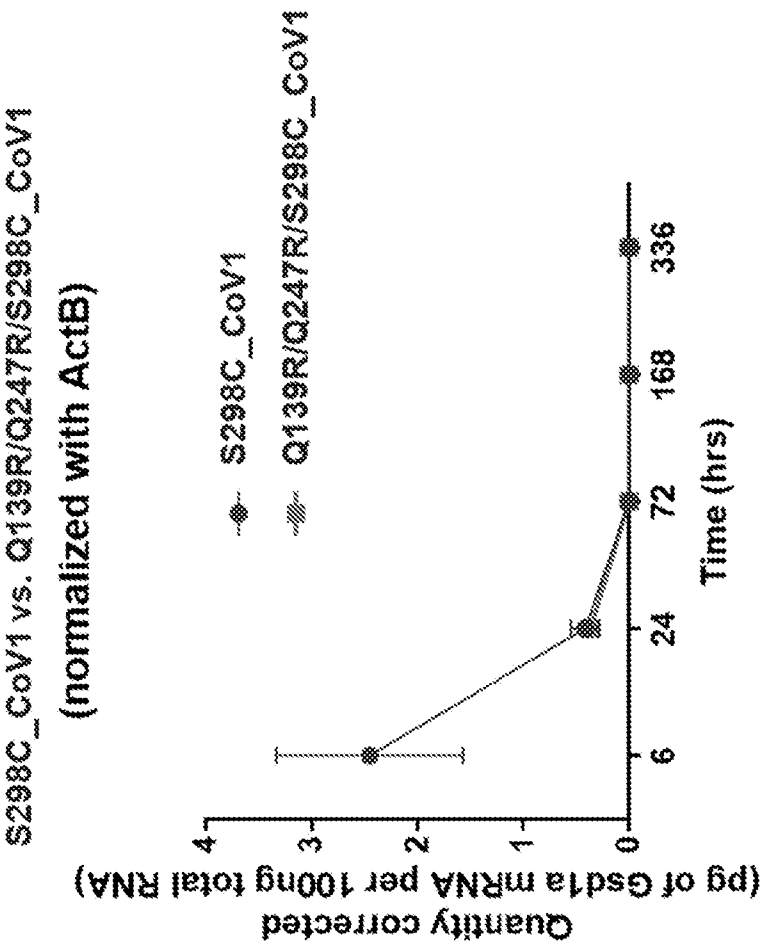

S298C_CoV1 vs. Q139R/Q247R/S298C_CoV1
(normalized with ActB)
S298C_CoV1
Q139R/Q247R/S298C_CoV1
Quantity corrected
(pg of Gsdl4a mRNA per 100ng total RNA)
0
1
2
3
4
6
24
72
168
336
Time (hrs)

FIGURE 15B

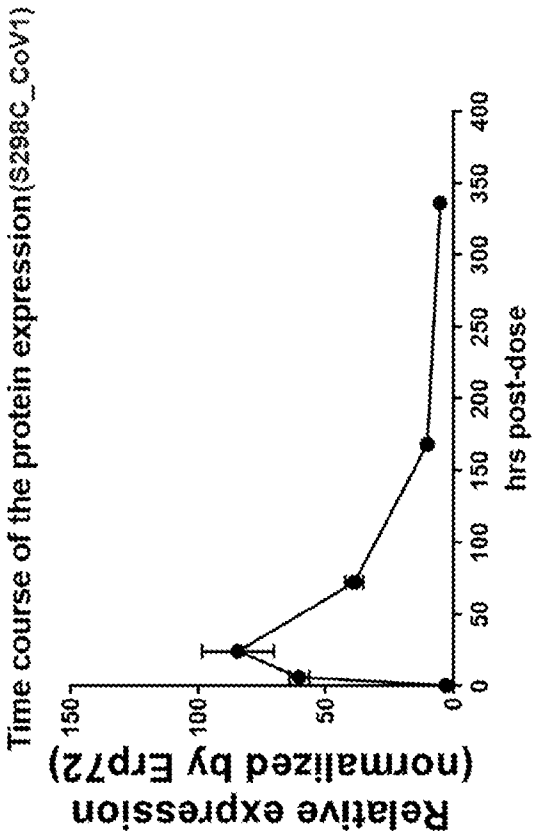


FIGURE 15C

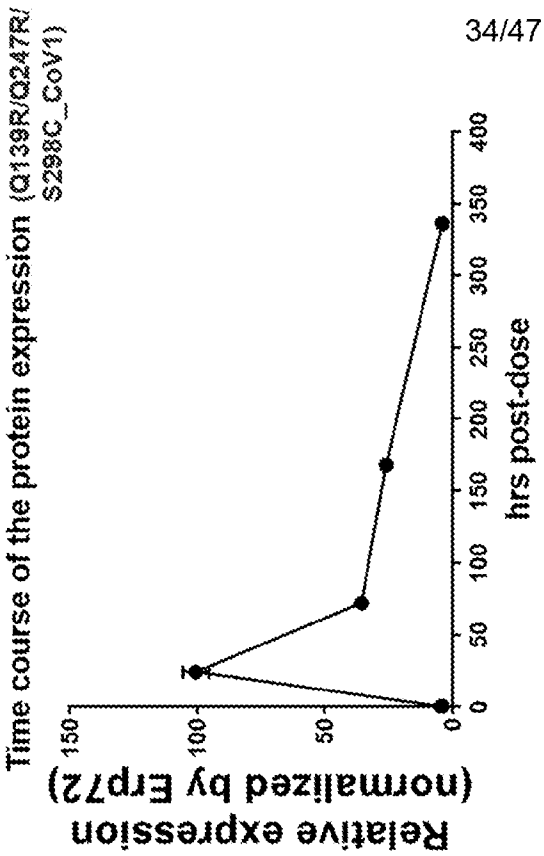


FIGURE 15E

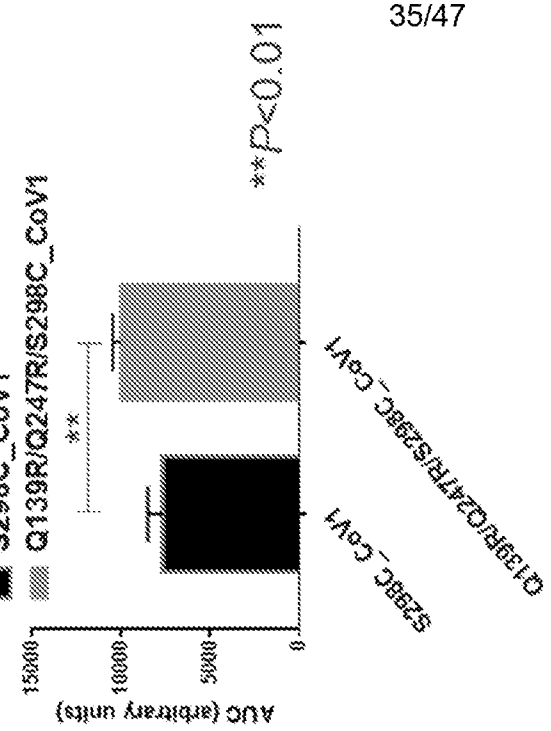


FIGURE 15D

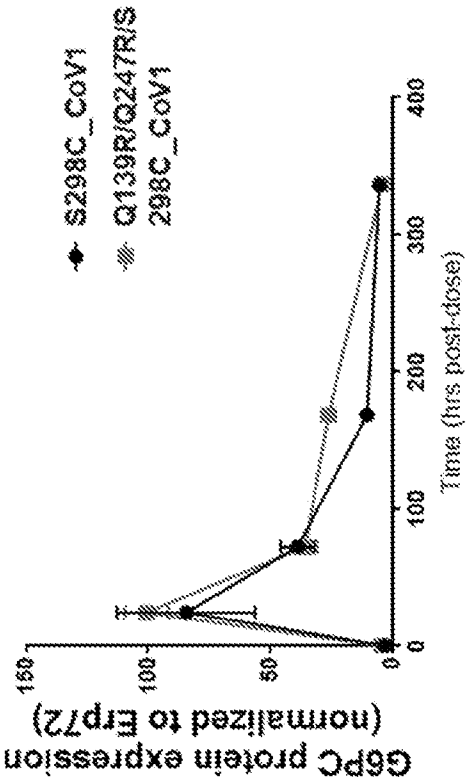


FIGURE 15G

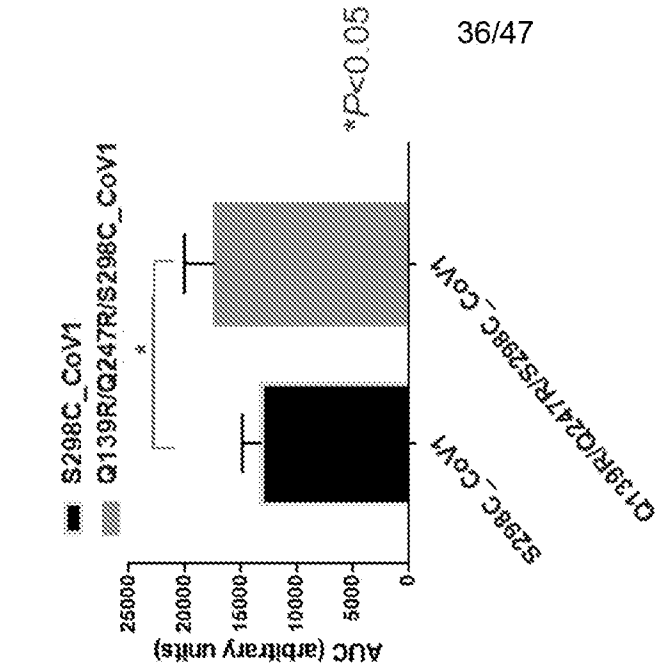


FIGURE 15F

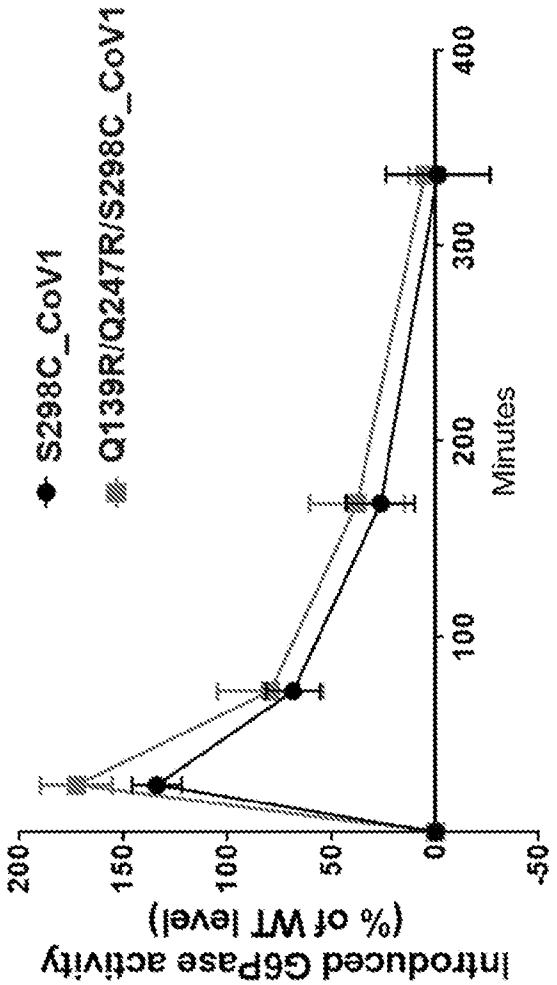


FIGURE 15H

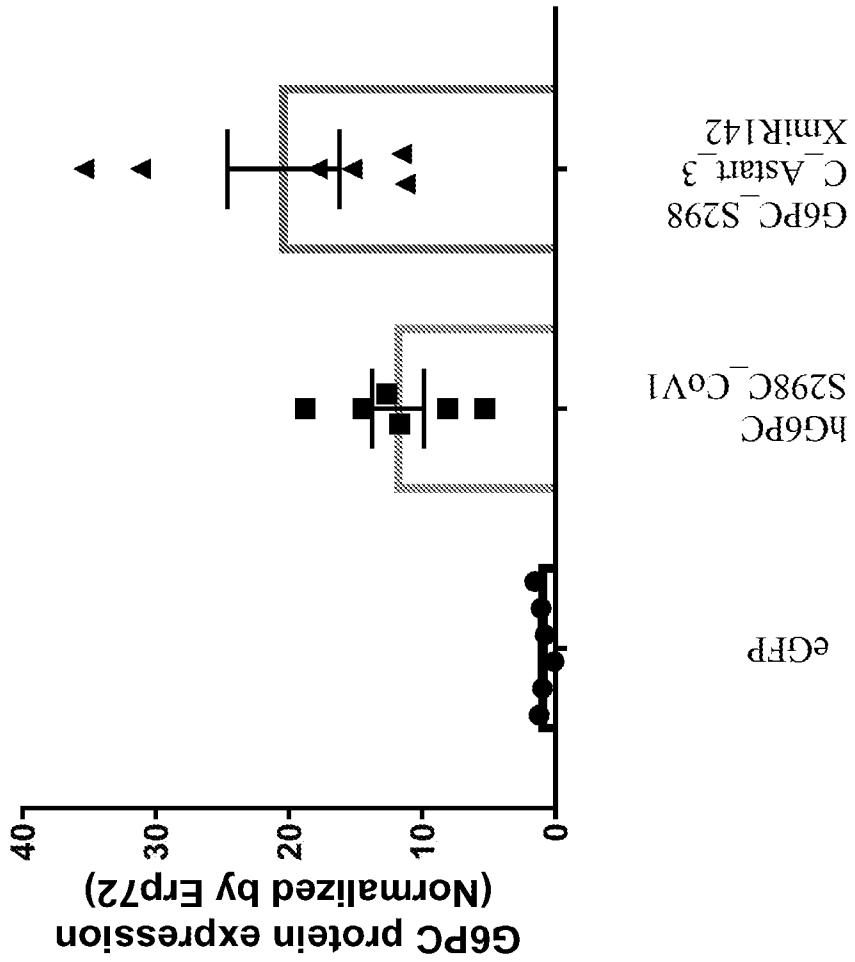

G6PC protein expression
(Normalized by Erp72)
0
10
20
30
40
eGFP
hG6PC
S298C_CoV1
G6PC_S298
C_Astart_3
XmiR142

FIGURE 15I

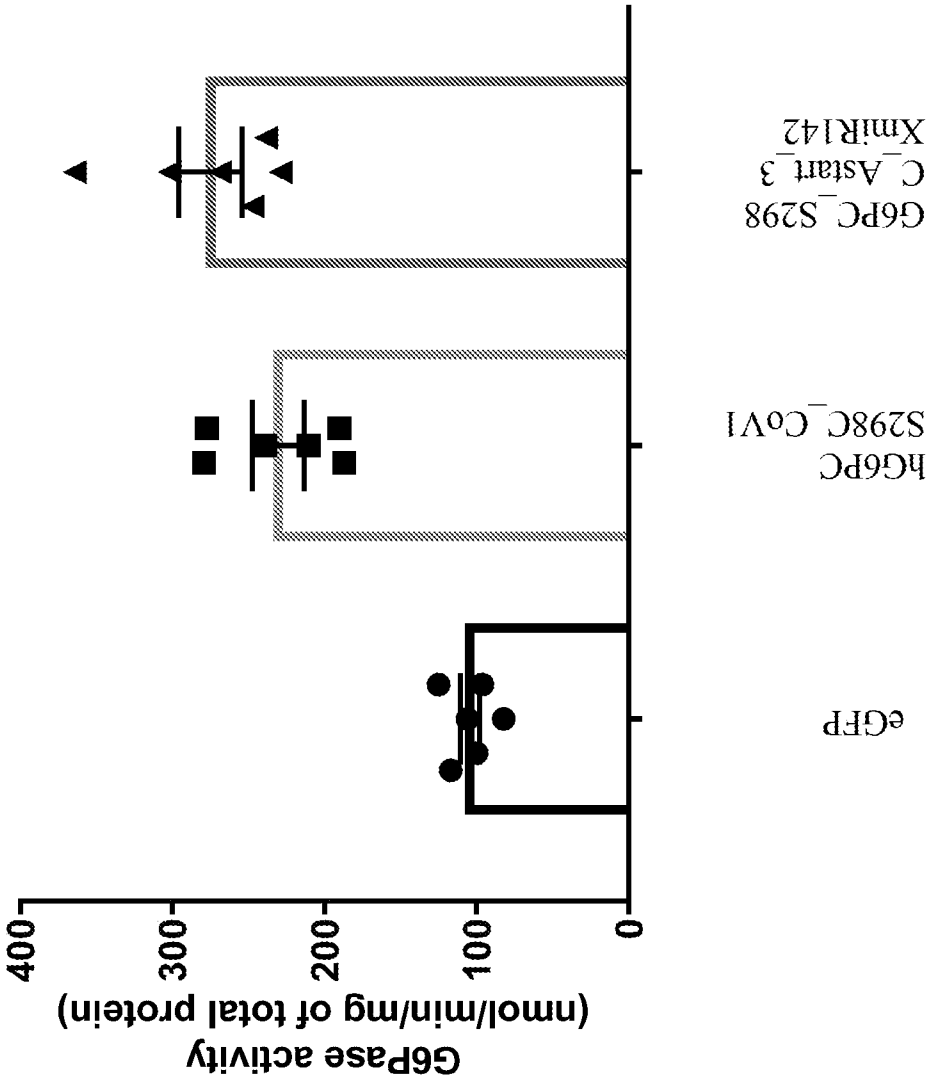

G6Pase activity
(nmol/min/mg of total protein)
0
100
200
300
400
eGFP
hG6PC
S298C_CoV1
G6PC_S298
C_Astart_3
XmiR142

FIGURE 16

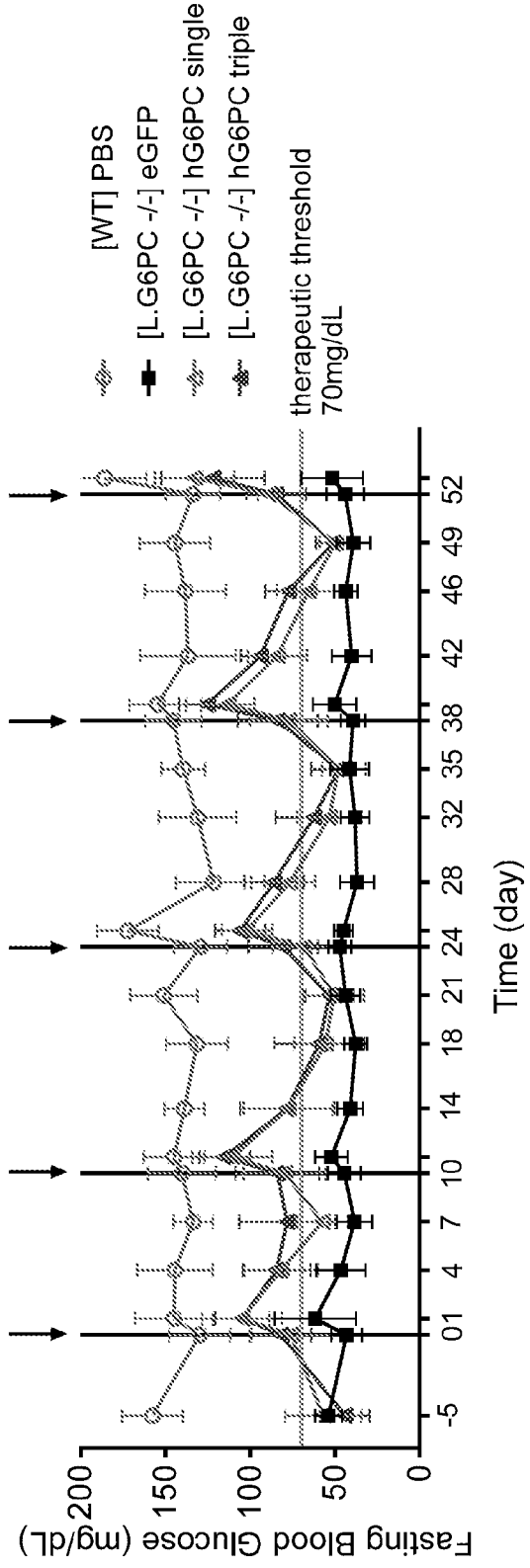

[WT] PBS
[L.G6PC -/-] eGFP
[L.G6PC -/-] hG6PC single
[L.G6PC -/-] hG6PC triple
therapeutic threshold
70mg/dL
1st Dose
2nd Dose
3rd Dose
4th Dose
5th Dose
Fasting Blood Glucose (mg/dL)
Time (day)
0
50
100
150
200
-5
01
4
7
10
14
18
21
24
28
32
35
38
42
46
49
52

FIGURE 17

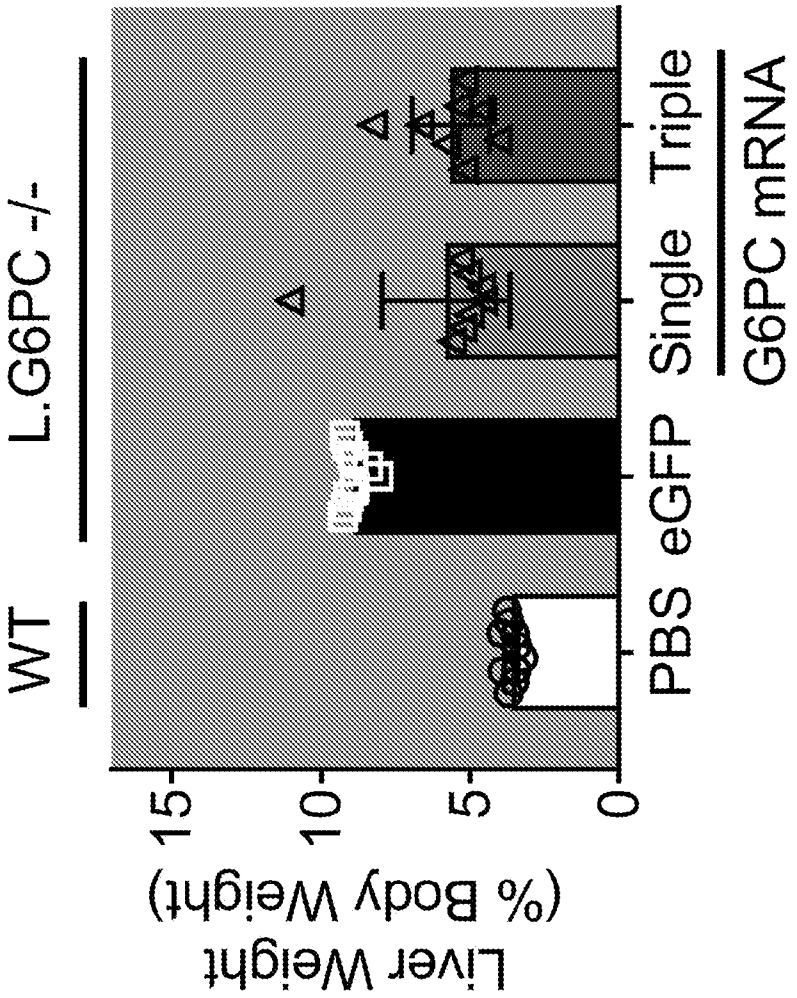

WT
L.G6PC -/-
Liver Weight
(% Body Weight)
15
10
5
0
PBS
eGFP
Single
Triple
G6PC mRNA

FIGURE 18

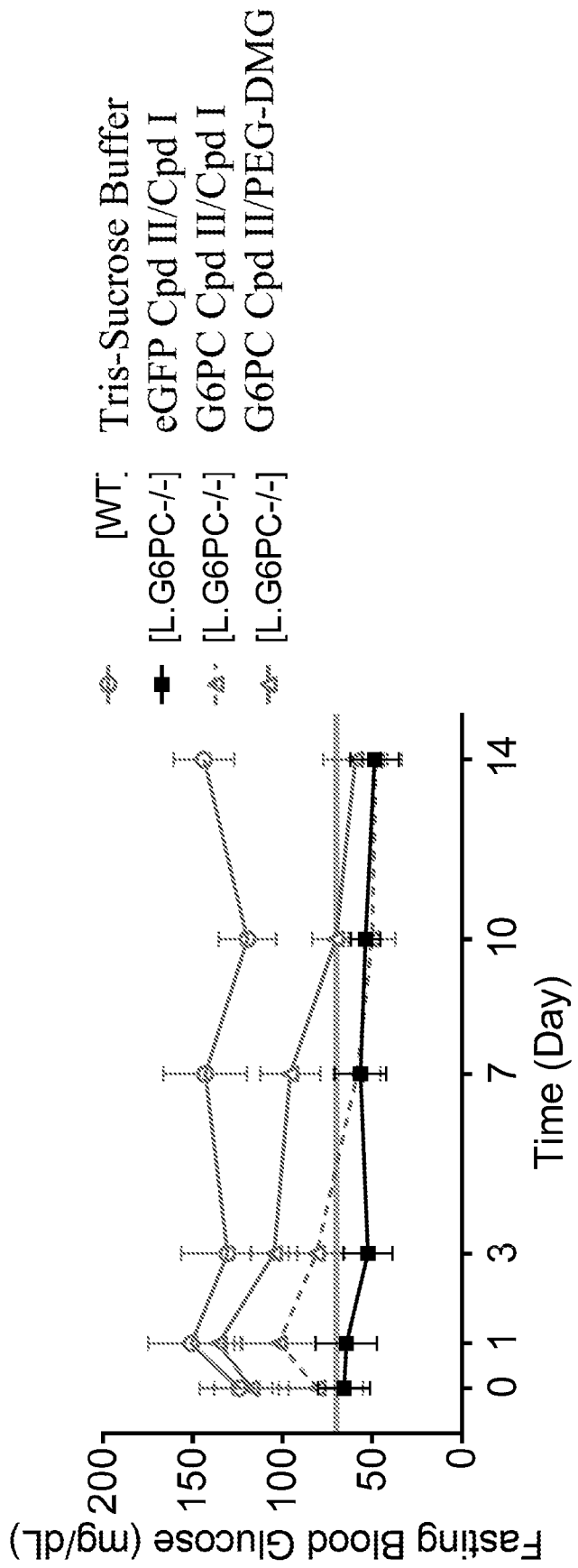

[WT Tris-Sucrose Buffer
[L.G6PC-/-] eGFP Cpd II/Cpd I
[L.G6PC-/-] G6PC Cpd II/Cpd I
[L.G6PC-/-] G6PC Cpd II/PEG-DMG
Fasting Blood Glucose (mg/dL)
200
150
100
50
0
0 1 3 7 10 14
Time (Day)

FIGURE 19A

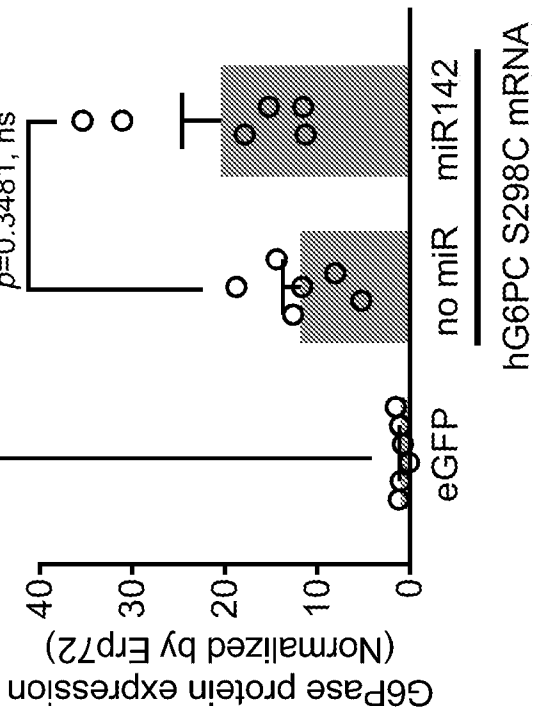


FIGURE 19B

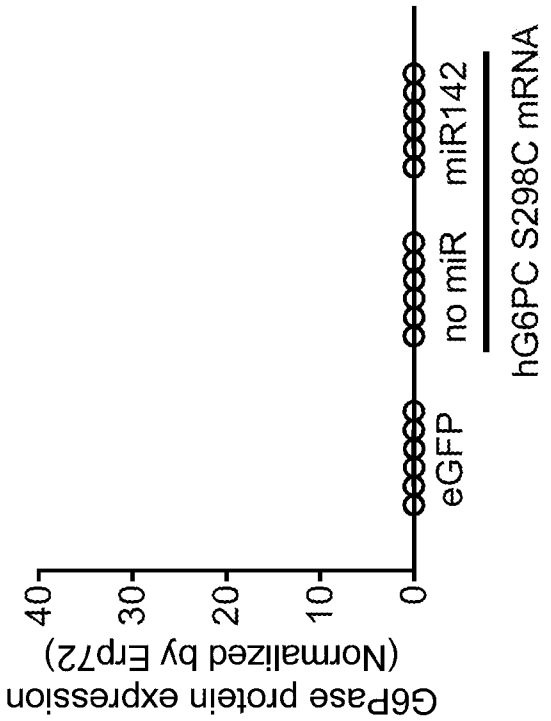


FIGURE 20A

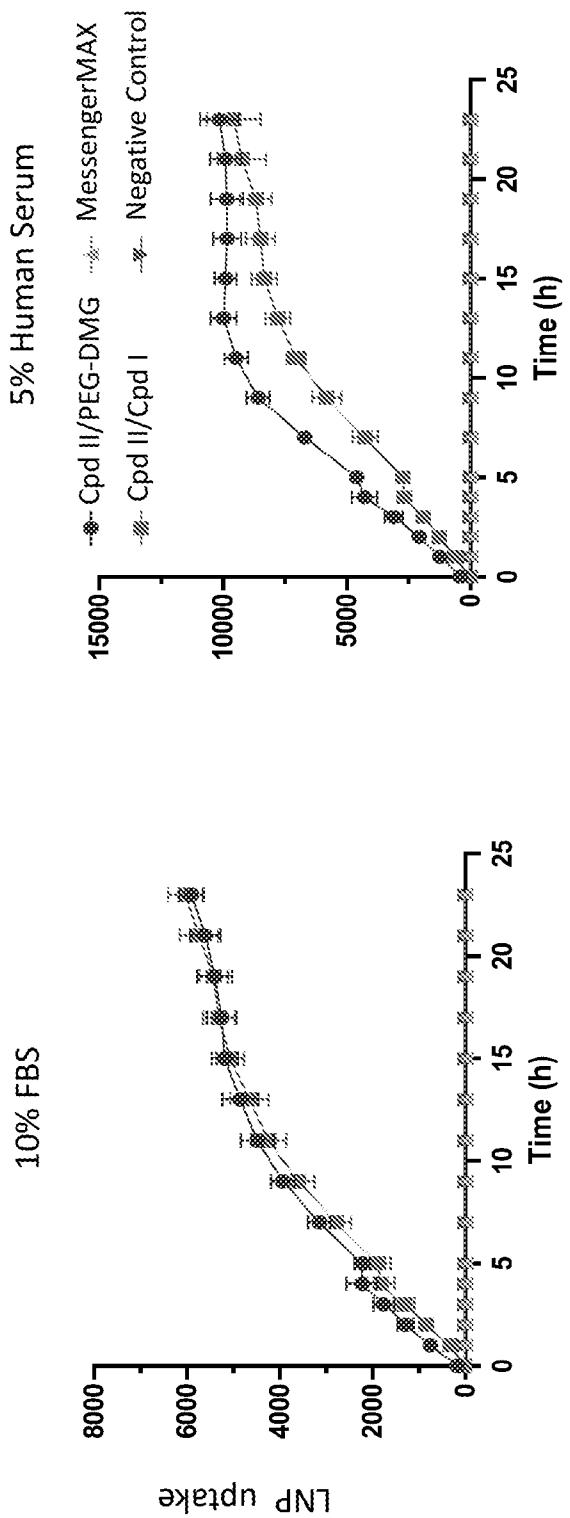

5% Human Serum
Cpd II/PEG-DMG
MessengerMAX
Cpd II/Cpd I
Negative Control
15000
10000
5000
0
0
5
10
15
20
25
Time (h)
10% FBS
LNP uptake
8000
6000
4000
2000
0
0
5
10
15
20
25
Time (h)

FIGURE 20B

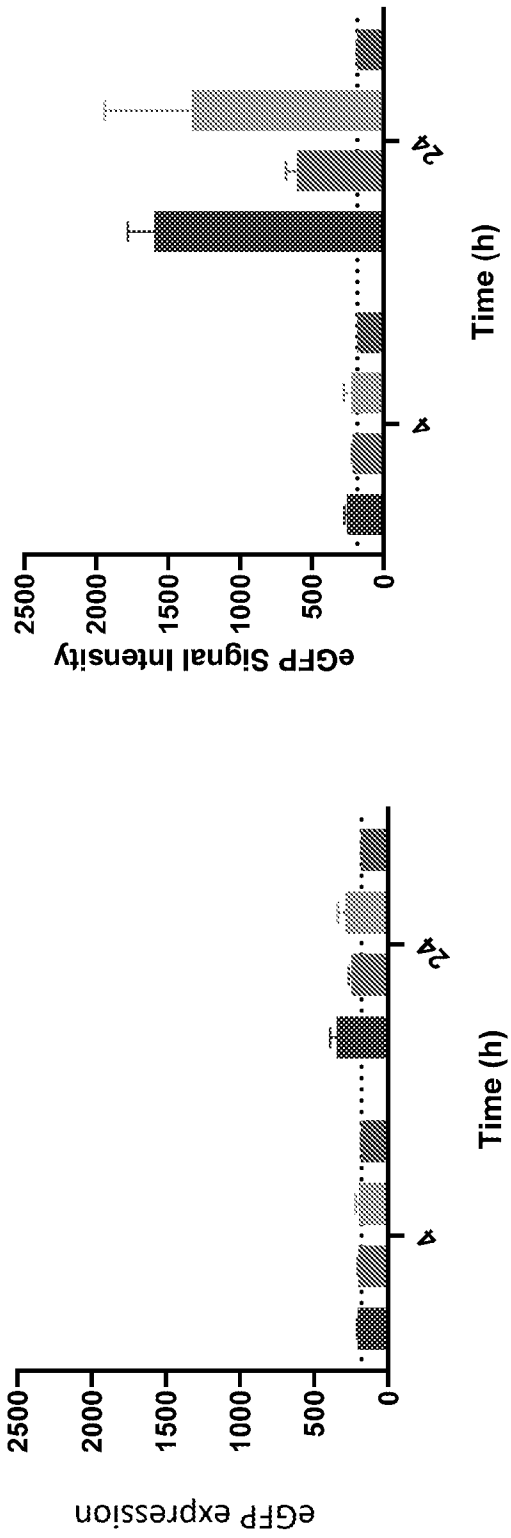

10% FBS
eGFP expression
2500
2000
1500
1000
500
0
4
24
Time (h)
5% Human Serum
eGFP Signal Intensity
2500
2000
1500
1000
500
0
4
24
Time (h)

FIGURE 20C

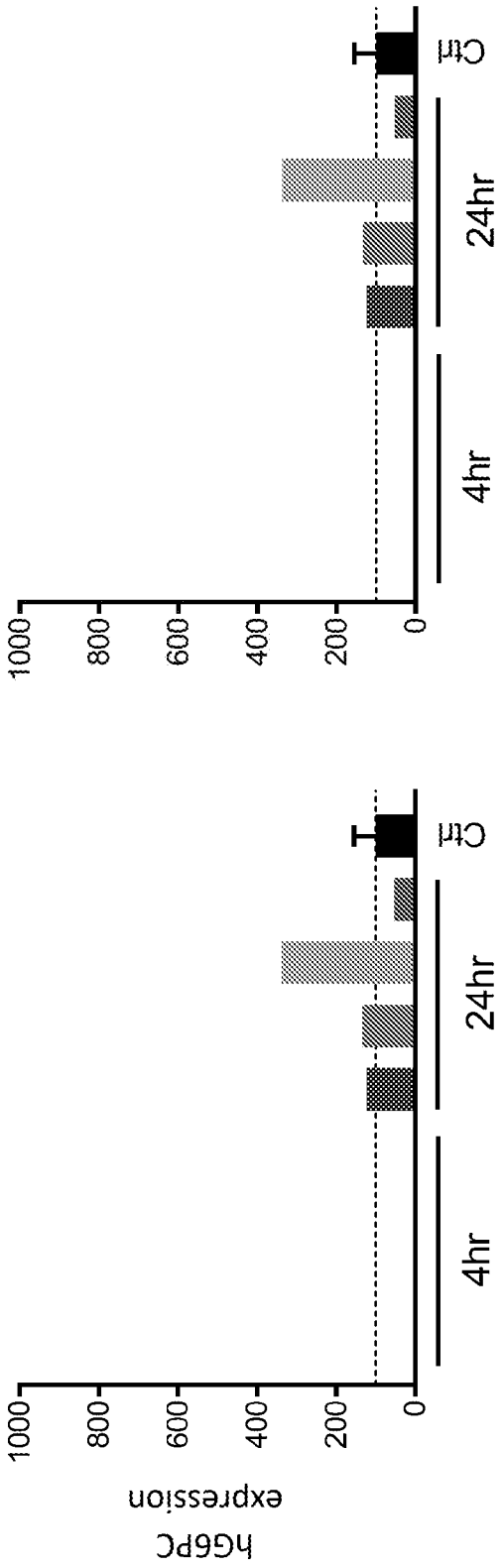

hG6PC expression
1000
800
600
400
200
0
10% FBS
4hr
24hr
Ctl
5% Human Serum
4hr
24hr
Ctl

FIGURE 20D

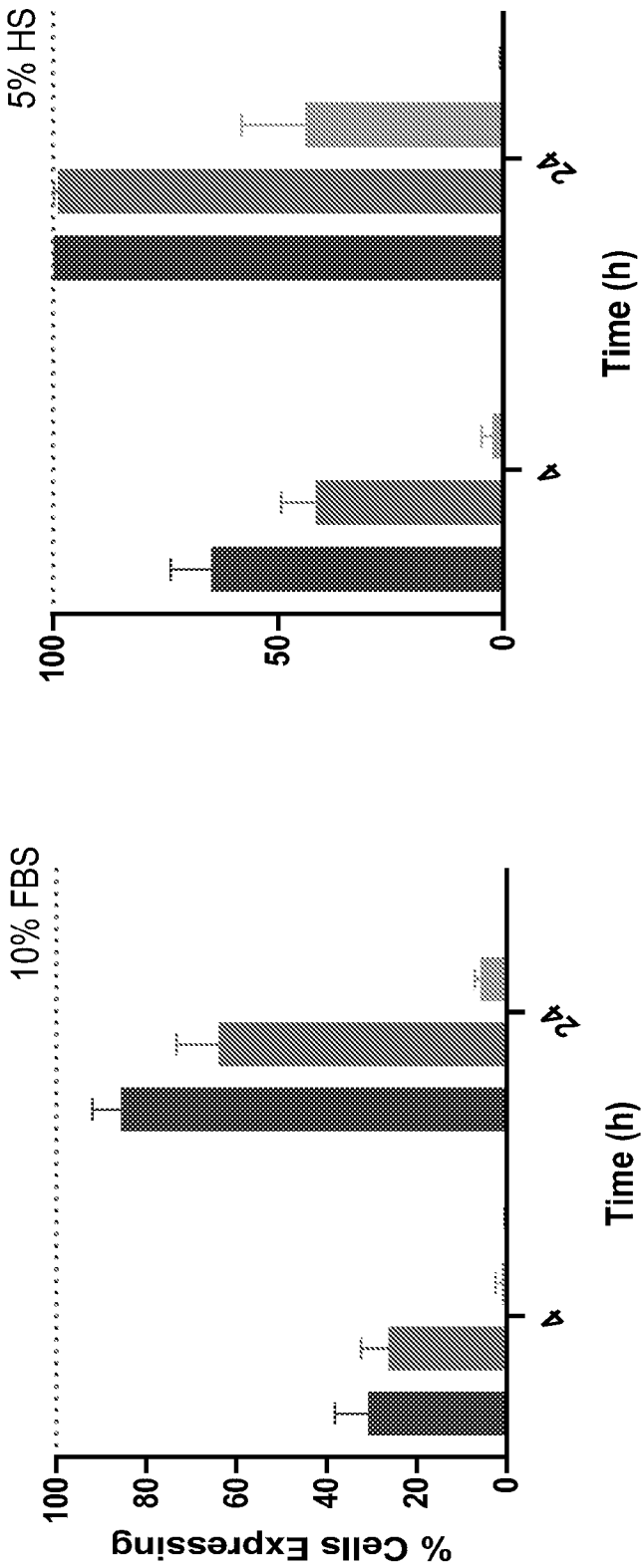


FIGURE 21A

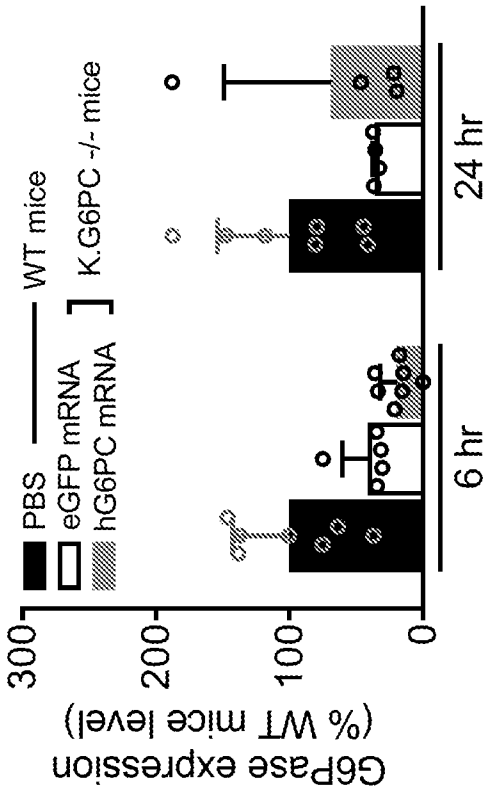


FIGURE 21B

