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(54) **OLIGOSACCHARIDES COMPRISING AN AMINOOXY GROUP AND CONJUGATES THEREOF**

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(71) Applicant: **Genzyme Corporation**, Cambridge, MA (US)

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(72) Inventors: **Yunxiang ZHU**, Bridgewater, NJ (US); **Seng H. CHENG**, Natick, MA (US); **Canwen JIANG**, Southborough, MA (US); **Luis Z. AVILA**, Bridgewater, NJ (US)

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(57) **ABSTRACT**

Related U.S. Application Data

(63) Continuation of application No. 14/272,960, filed on May 8, 2014, now Pat. No. 9,469,850, which is a continuation of application No. 12/523,631, filed on Aug. 20, 2009, now Pat. No. 8,759,501, filed as application No. PCT/US2008/051429 on Jan. 18, 2008.

The invention provides methods for the synthesis of oligosaccharides comprising an aminooxy group. The invention further provides oligosaccharides comprising an aminooxy group, methods for coupling oligosaccharides comprising an aminooxy group to glycoproteins, and oligosaccharide-protein conjugates. Also provided are methods of treating a lysosomal storage disorder in a mammal by administration of an oligosaccharide-protein conjugate.

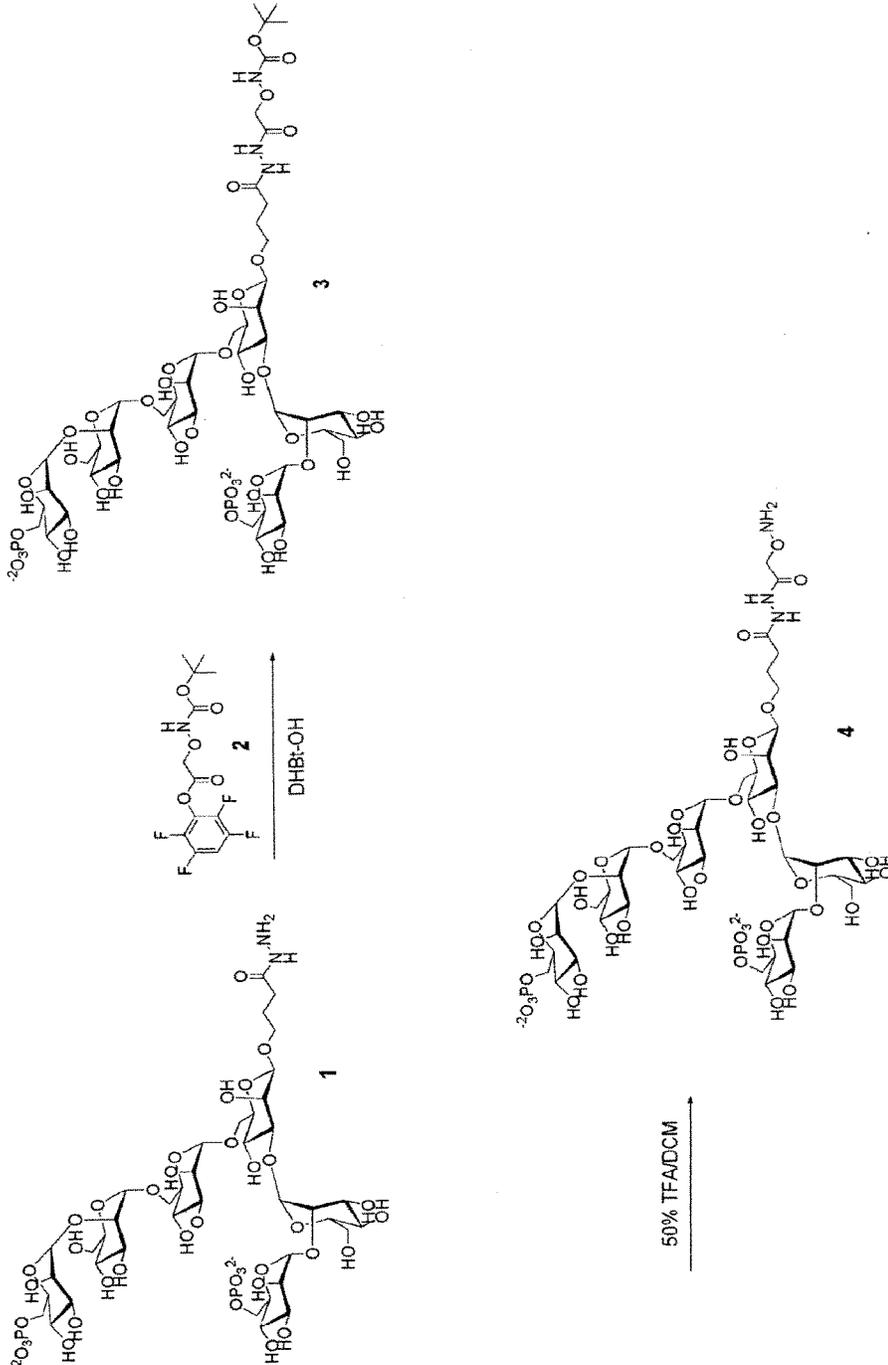


FIG. 1

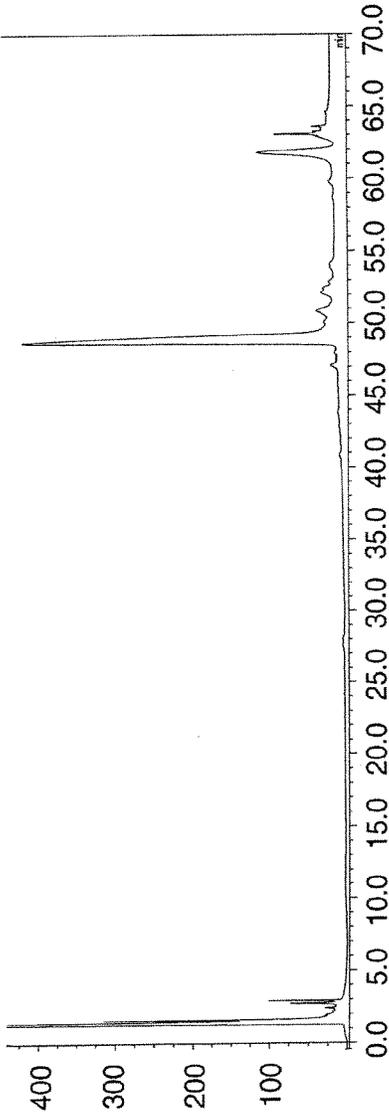


FIG. 2A

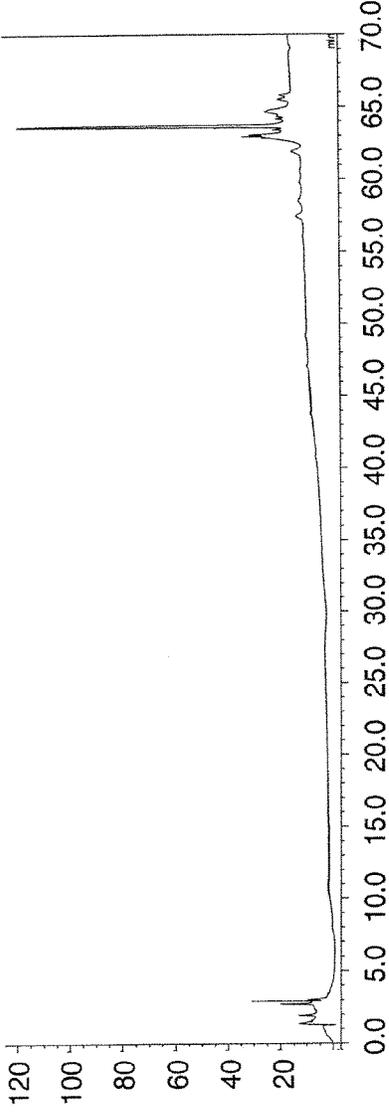


FIG. 2B

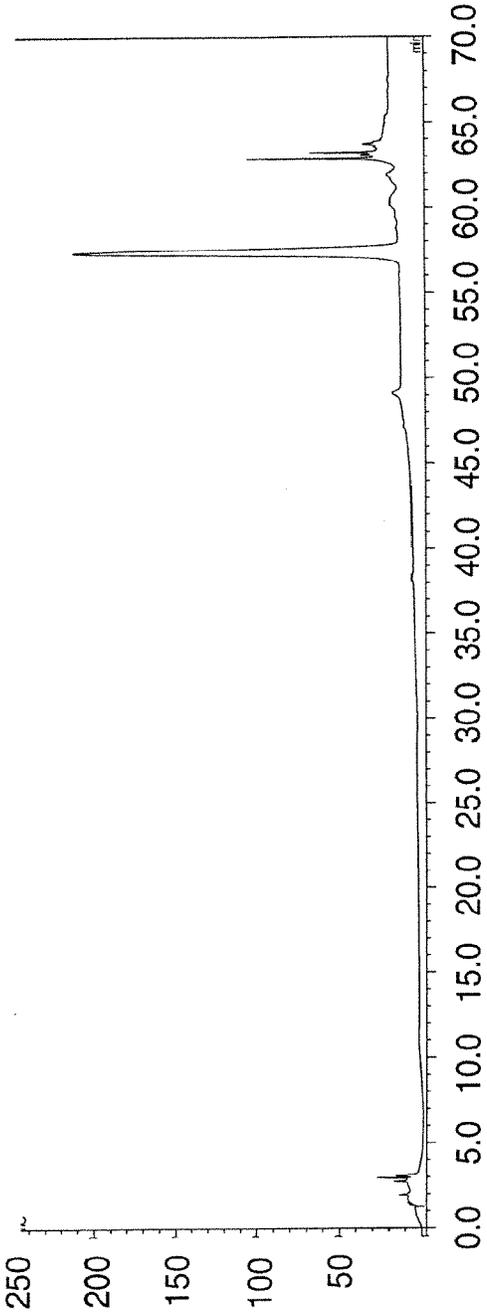


FIG. 2C

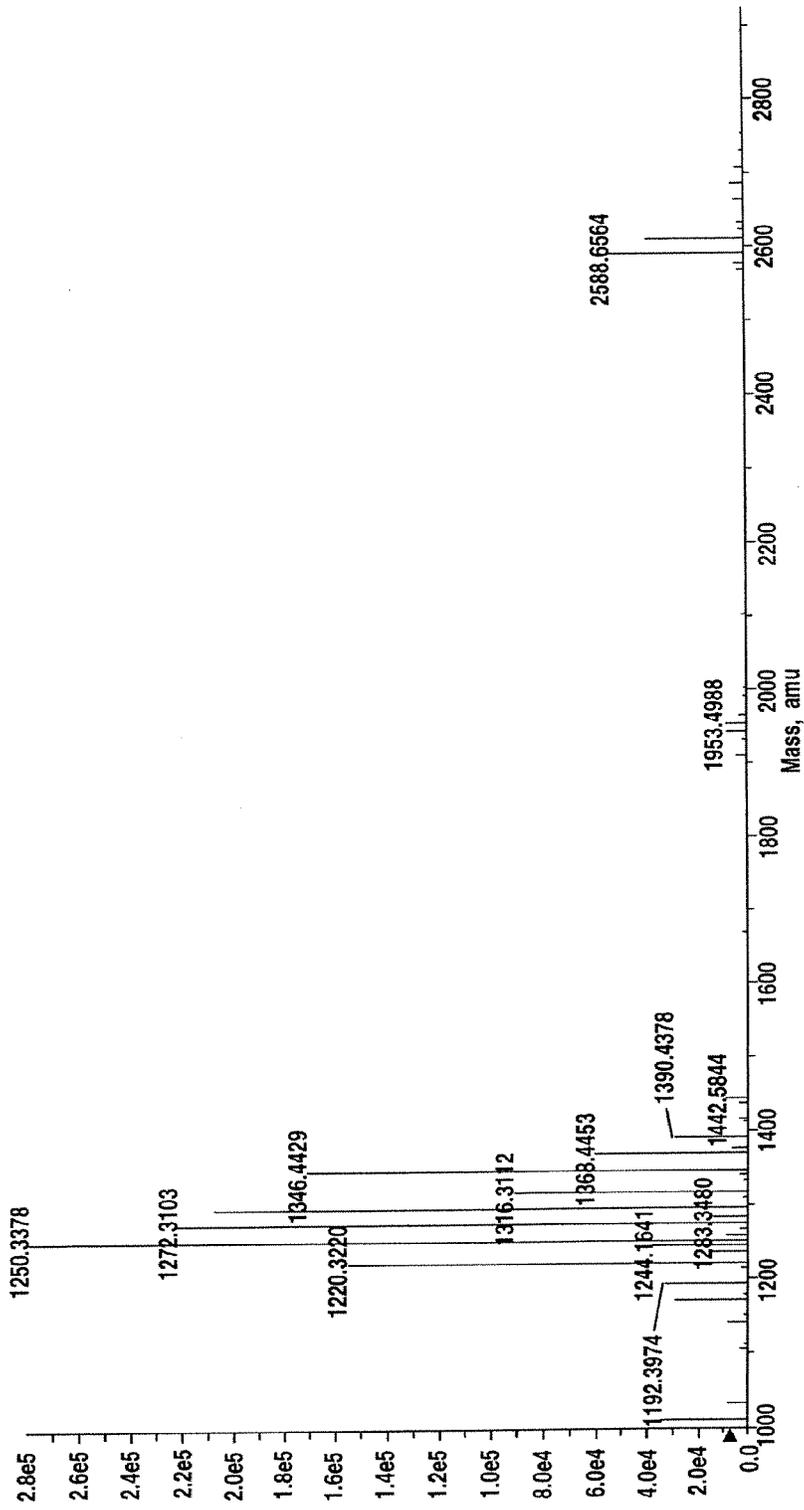


FIG. 3A

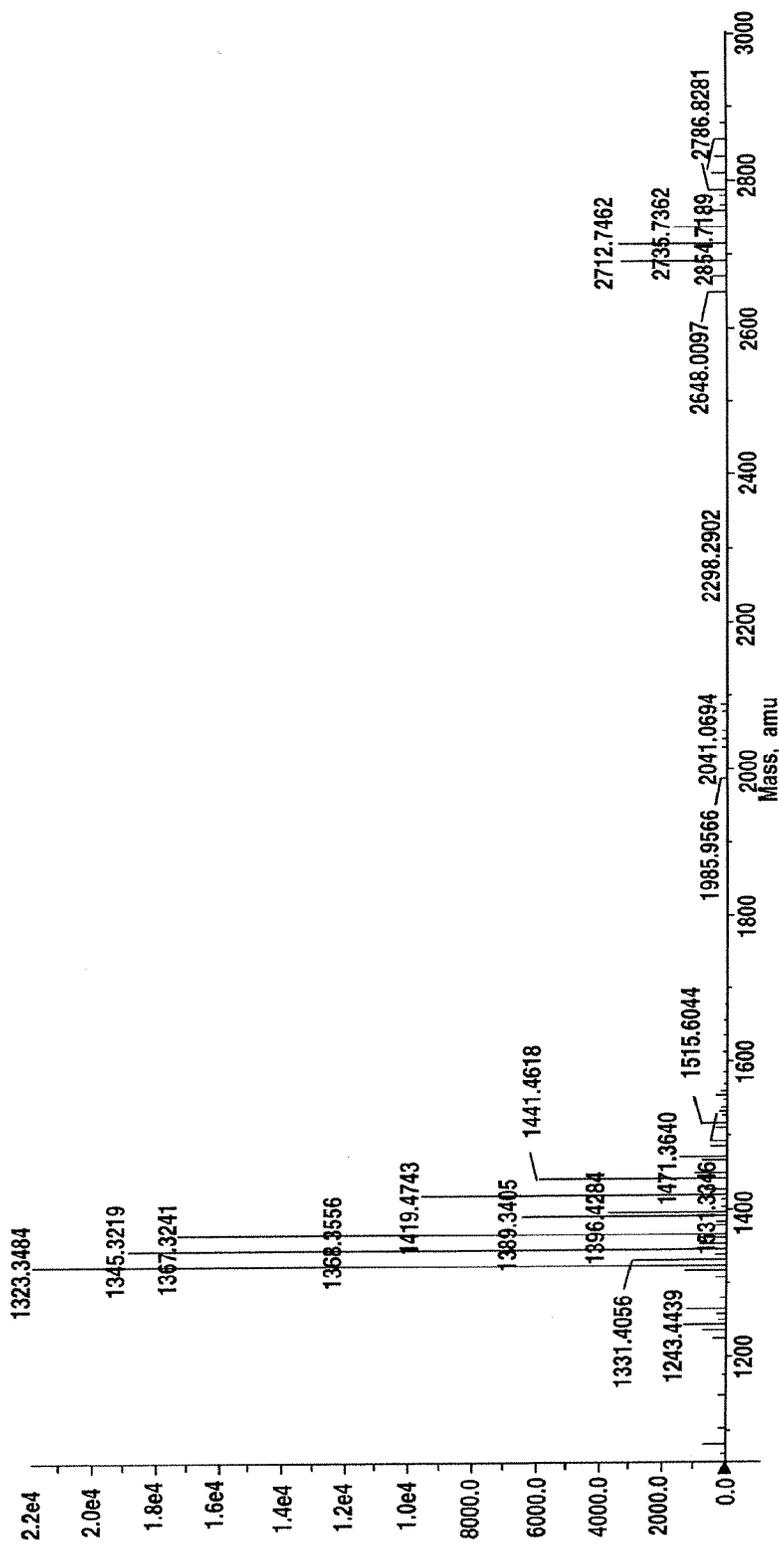


FIG. 3B



FIG. 4

**OLIGOSACCHARIDES COMPRISING AN
AMINOXY GROUP AND CONJUGATES
THEREOF**

[0001] This application claims the benefit of U.S. Provisional Application No. 60/885,471, filed Jan. 18, 2007, the disclosure of which is incorporated herein by reference.

[0002] The invention relates generally to methods for the synthesis of oligosaccharides comprising an aminoxy group from oligosaccharides comprising a reactive group. In another embodiment, the invention further relates to oligosaccharides comprising an aminoxy group. The invention also relates to methods of conjugating oligosaccharides comprising an aminoxy group to proteins, including glycoproteins (such as, e.g., lysosomal enzymes), and to compositions of oligosaccharide-protein conjugates, including oligosaccharide-glycoprotein conjugates. Another embodiment of the invention relates to methods of treating lysosomal storage disorders using such oligosaccharide-lysosomal enzyme conjugates.

[0003] Lysosomal storage disorders (LSDs) are a class of rare metabolic disorders comprising over forty genetic diseases involving a deficiency in the activity of lysosomal hydrolases. A hallmark feature of LSDs is the abnormal accumulation of lysosomal metabolites, which leads to the formation of large numbers of distended lysosomes.

[0004] LSDs can be treated by administration of the active version of the enzyme deficient in the patient, a process termed enzyme replacement therapy (ERT). The administered replacement enzyme bearing a terminal mannose-6-phosphate (M6P) is taken up by target cells through cell-surface-associated cation-independent M6P receptor (CI-MPR)-mediated endocytosis, and directed to the lysosome.

[0005] In general, poorly phosphorylated replacement enzymes are not internalized by the M6P receptor on cell surfaces, and therefore cannot be directed to the lysosome where they function. Consequently, a low degree of mannose phosphorylation can have a significant and deleterious effect on the therapeutic efficacy of a replacement enzyme.

[0006] Methods thus have been developed for increasing the M6P content of replacement enzymes. U.S. Pat. No. 7,001,994, for example, describes a method for coupling oligosaccharides comprising M6P with glycoproteins. The oligosaccharides of the glycoproteins are first oxidized with periodate or galactose oxidase to result in the formation of carbonyl groups, which are then chemically conjugated with an oligosaccharide functionalized at the reducing end with a carbonyl-reactive group (such as, e.g., a hydrazine, hydrazide, aminoxy, thiosemicarbazide, semicarbazide, or amine group) to yield an oligosaccharide-glycoprotein conjugate.

[0007] A conjugate of the lysosomal enzyme acid α -glucosidase (GAA) with a bis-M6P oligosaccharide was prepared by the above-described method, and found to be more effective in reducing skeletal and cardiac muscle glycogen than recombinant human GAA in a murine model of Pompe disease, an autosomal recessive muscular disease resulting from a metabolic deficiency of GAA, and characterized by the accumulation of lysosomal glycogen.

[0008] Aminoxy groups are particularly useful carbonyl-reactive groups for the conjugation reactions described above, as the resulting conjugates comprise a relatively stable oxime linkage. Therefore, there is a need for methods for the preparation of aminoxy functionalized oligosaccharides.

[0009] The present invention provides methods of preparing oligosaccharides comprising an aminoxy group. These methods are generally applicable to a broad range of protected and unprotected oligosaccharides, such as, e.g., branched and unbranched, and phosphorylated and unphosphorylated, oligosaccharides. In certain embodiments, the oligosaccharide may be a disaccharide, trisaccharide, tetrasaccharide, pentasaccharide, hexasaccharide, heptasaccharide, or greater. The oligosaccharide may, in certain embodiments, comprise at least one M6P residue. In some embodiments, the oligosaccharide may comprise at least 1, 2, 3, 4, 5, 6, or 7 terminal M6P residues.

[0010] The invention provides a method of preparing an oligosaccharide comprising an aminoxy group from an oligosaccharide comprising a reactive group. The method comprises:

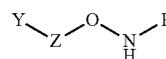
[0011] (a) providing an oligosaccharide comprising a first reactive group;

[0012] (b) providing an aminoxy compound comprising an aminoxy group and a second reactive group; and

[0013] (c) reacting the first reactive group of the oligosaccharide with the second reactive group of the aminoxy compound, thereby preparing the oligosaccharide comprising an aminoxy group.

[0014] The first and second reactive groups may be chosen from, e.g., hydrazine, hydrazide, thiosemicarbazide, semicarbazide, amine, carboxyl, activated ester, acyl halide, acyl azide, alkyl halide, anhydride, isothiocyanate, isocyanate, and sulfonyl halide groups.

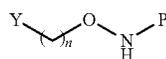
[0015] In some embodiments, the aminoxy compound is chosen from compounds of Formula II:



Formula II

wherein Y is the second reactive group, Z is chosen from alkyl, alkenyl, alkynyl, aryl, heteroaryl, and heterocyclyl, and P is chosen from amino protecting groups (such as, e.g., carbamate protecting groups). For example, in some embodiments, Y may be a carboxyl, activated ester, acyl halide (such as, e.g., an acyl fluoride or acyl chloride), acyl azide, alkyl halide, anhydride, isothiocyanate, isocyanate, or sulfonyl halide (such as, e.g., a sulfonyl chloride or sulfonyl bromide). In other embodiments, Y may be, e.g., a hydrazine, hydrazide, thiosemicarbazide, semicarbazide, or amine group.

[0016] In certain embodiments, the aminoxy compound of Formula II is chosen from compounds of Formula III:

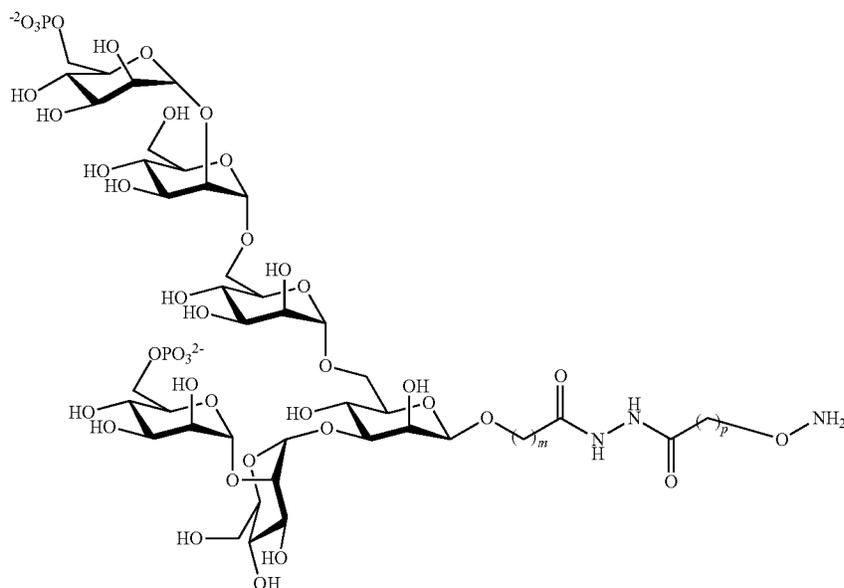


Formula III

wherein Y is the second reactive group, n is chosen from integers ranging from 1 to 10, and P is chosen from amino protecting groups.

[0017] In certain embodiments, the aminoxy compound comprises an amino protecting group, and the method further comprises a step (d), deprotecting the oligosaccharide comprising an aminoxy group.

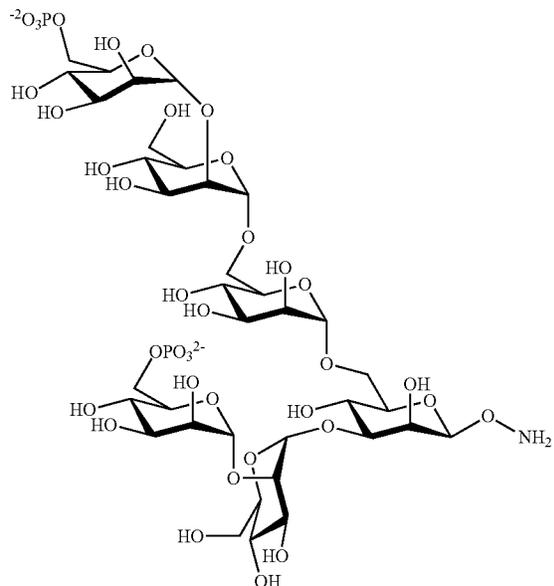
[0018] The invention further provides an oligosaccharide comprising (1) an aminoxy group and (2) mannose-6-phosphate. In some embodiments, that oligosaccharide is prepared by the methods described above. For example, in some embodiments, the invention provides an oligosaccharide comprising an aminoxy group of Formula IV:



Formula IV

wherein m and p are independently chosen from integers ranging from 1 to 10.

[0019] In another embodiment, the invention provides an oligosaccharide of Formula V:



Formula V

[0020] In another embodiment, the invention provides methods of coupling an oligosaccharide to a protein. In one embodiment, the method comprises:

[0021] (a) providing an oligosaccharide comprising an aminoxy group;

[0022] (b) providing a protein having at least one carbonyl group; and

[0023] (c) reacting the aminoxy group of the oligosaccharide with the at least one carbonyl group of the protein,

thereby coupling the oligosaccharide to the protein.

[0024] In other embodiments, the invention further provides an oligosaccharide-protein conjugate comprising (1) a protein, (2) an oligosaccharide, and (3) an oxime group connecting the protein and the oligosaccharide. For example, in some embodiments, the invention provides an oligosaccharide-protein conjugate prepared by the methods disclosed above. In certain embodiments, the oligosaccharide-protein conjugate is an oligosaccharide-glycoprotein conjugate. In certain embodiments, the oligosaccharide-glycoprotein conjugate is the conjugate of an oligosaccharide comprising at least one M6P and of a lysosomal enzyme such as, e.g., a lysosomal hydrolase. In some embodiments, the invention provides pharmaceutical compositions comprising an oligosaccharide-protein conjugate of the invention.

[0025] Another embodiment of the invention provides methods of treating a lysosomal storage disorder such as, e.g., those disclosed in Table 1. In some embodiments, the methods comprise administering to a mammal an oligosaccharide-glycoprotein conjugate of the invention, wherein the oligosaccharide comprises at least one M6P and the glycoprotein is a lysosomal hydrolase. This disclosure further provides the use of a conjugate of the invention for treating a lysosomal storage disorder in a subject in need thereof, and in the manufacture of a medicament for treating a lysosomal storage disorder.

BRIEF DESCRIPTION OF THE FIGURES

[0026] FIG. 1 is a reaction scheme depicting an illustrative embodiment of the methods of the invention. Oligosaccharide 1, having a first reactive group (a hydrazide group), is reacted with aminoxy compound 2 in presence of the catalyst 3-hydroxy-1,2,3-benzotriazin-4(3H)-one (DHBt-OH), to yield oligosaccharide 3. The tert-butyloxycarbonyl (t-Boc) amino protecting group of oligosaccharide 3 is then removed with 50% trifluoroacetic acid/dichloromethane (TFA/DCM) to yield oligosaccharide 4.

[0027] FIG. 2 depicts a series of gel chromatographs of intermediates in the synthetic scheme described in FIG. 1. FIG. 2A is a Dionex analytical chromatograph of starting oligosaccharide 1. FIG. 2B is a Dionex analytical chromatograph of oligosaccharide 3. FIG. 2C is a Dionex analytical chromatograph of oligosaccharide 4.

[0028] FIG. 3A is a mass spectrum of oligosaccharide 1 (calculated molecular weight=1250; calculated molecular weight of sodium salt=1338). FIG. 3B is a mass spectrum of oligosaccharide 4 (calculated molecular weight=1323; calculated molecular weight of sodium salt=1411).

[0029] FIG. 4 is a reaction scheme depicting an illustrative embodiment of the methods of the invention. Oligosaccharide 5 having a first reactive group (a carboxyl group) is reacted with aminoxy compound 6 in presence of the coupling agent 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC) and the catalyst N-hydroxysuccinimide (NHS), to yield aminoxy-containing oligosaccharide 3. The Boc amino protecting group of oligosaccharide 3 is then removed with 50% TFA/DCM to yield oligosaccharide 4.

I. PREPARATION OF AN OLIGOSACCHARIDE COMPRISING AN AMINOXY GROUP

A. Oligosaccharide Comprising a Reactive Group

[0030] The methods of the invention are applicable to a broad range of oligosaccharides comprising a reactive group. As used herein, an oligosaccharide refers to a disaccharide, trisaccharide, tetrasaccharide, pentasaccharide, hexasaccharide, heptasaccharide, or larger oligosaccharide (such as, e.g., an oligosaccharide comprising 2-50, 2-10, 8-25, or 8-50 saccharide units). Accordingly, in various embodiments, an oligosaccharide may be, e.g., a disaccharide, trisaccharide, tetrasaccharide, a pentasaccharide, a hexasaccharide, a heptasaccharide, or a larger oligosaccharide. An oligosaccharide may be mono-, bi-, tri-, tetra-, or pentamery in structure. An oligosaccharide may comprise 0, 1, 2, 3, 4, or more branch points.

[0031] The reactive group on the oligosaccharide, also referred to as a first reactive group, may be, in some embodiments, e.g., a hydrazine group, hydrazide group, semicarbazide group, thiosemicarbazide, or amine group. In some embodiments, the first reactive group may be, e.g., a carboxyl, ester (such as, e.g., an activated ester), acyl halide (such as, e.g., acyl fluoride or acyl chloride), acyl azide, alkyl halide, anhydride, isothiocyanate, isocyanate, or sulfonyl halide (such as, e.g., sulfonyl chloride or sulfonyl bromide) group.

[0032] The first reactive group may be connected to the reducing end of the oligosaccharide or may be located anywhere in the oligosaccharide. The first reactive group may, in certain embodiments, be connected through one or more linkers to the oligosaccharide. A linker, as used herein,

may be chosen from, e.g., a combination of optionally substituted alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, acyloxy, alkoxy, aryloxy, and heterocyclyloxy groups. A linker may be interrupted or terminated by one or more heteroatoms such as, e.g., nitrogen, sulfur, and oxygen. For example, a linker, in some embodiments, may comprise one or more ether, ester, or amide group.

[0033] Any chemical group of the linker (such as, e.g., alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, acyloxy, alkoxy, aryloxy, and heterocyclyloxy) may be substituted or unsubstituted, unless otherwise stated. Substituents may be chosen from, e.g., acyl, acylamino, acyloxy, alkenyl, alkoxy, alkyl, alkynyl, amido, amino, aryl, aryloxy, azido, carbamoyl, carboalkoxy, carboxy, cyano, cycloalkyl, formyl, guanidino, halo, heteroaryl, heterocyclyl, hydroxy, iminoamino, nitro, oxo, phosphonamino, sulfinyl, sulfonamino, sulfonate, sulfonyl, thio, thioacylamino, thioureido, and ureido. The substituents may themselves be substituted or unsubstituted, and may be interrupted or terminated by one or more heteroatoms such as, e.g., nitrogen, sulfur, and oxygen.

[0034] In certain embodiments, an oligosaccharide may comprise at least one protecting group. The term "protecting group" refers to any substituent that may be used to prevent a functional group (such as, e.g., an amine group, a carboxyl group, a hydroxyl group, a hydrazine group, a hydrazide group, a semicarbazide group, or a thiosemicarbazide group) on a molecule from undergoing a chemical reaction while chemical change occurs elsewhere in the molecule. A protecting group can be removed under the appropriate chemical conditions. Numerous protecting groups are known to those skilled in the art, and examples of protecting groups, methods for their addition, and methods for their removal can be found in, e.g., Greene et al., *Protective Groups in Organic Synthesis*, 3rd ed., John Wiley and Sons: New York, 1999 and Kocienski, *Protecting Groups*, 3rd ed., Georg Thieme Verlag: Stuttgart, Germany, 2005, the disclosures of which are herein incorporated by reference. In certain embodiments, the oligosaccharide may comprise at least one protecting group chosen from hydroxyl protecting groups, carboxyl protecting groups, and amino protecting groups. In other embodiments, an oligosaccharide may be "unprotected," and may not comprise any protecting groups.

[0035] An oligosaccharide may be isolated from a natural source or may be prepared by chemical or enzymatic synthesis. An oligosaccharide isolated from a natural source may be homogeneous or may be a heterogeneous mixture of related oligosaccharides. In some embodiments, an oligosaccharide may be prepared by chemical or enzymatic modification of an oligosaccharide isolated from a natural source ("semi-synthesis"). In some embodiments, the oligosaccharide may be a synthetic oligosaccharide having the chemical structure of a naturally occurring oligosaccharide.

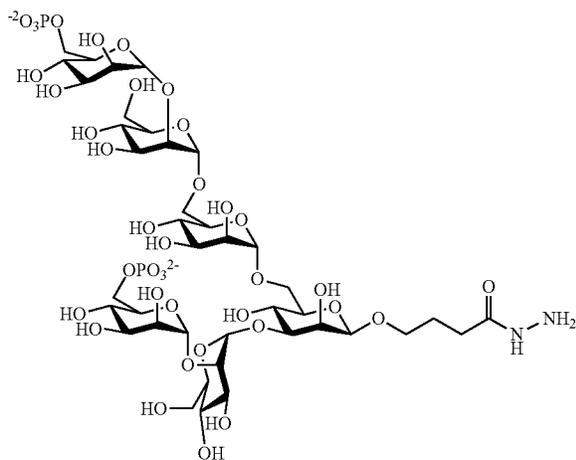
[0036] In some embodiments, an oligosaccharide may comprise a monosaccharide that is recognized by a particular receptor. The monosaccharide recognized by a particular receptor may be chosen from, e.g., galactose, GalNAc, mannose, M6P, glucose, GlcNAc, sialic acid, or sulfated sialic acid residue. An oligosaccharide may, in certain embodiments, comprise at least one M6P residue, such as, e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 M6P residues.

[0037] The monosaccharide recognized by a particular receptor may be, in some embodiments, a penultimate monosaccharide or a terminal monosaccharide. In some

embodiments, the monosaccharide recognized by a particular receptor may be a terminal galactose, mannose, M6P, glucose, GlcNAc, or sialic acid residue. An oligosaccharide may, in some embodiments, contain at least 1, 2, 3, 4, 5, 6, 7 terminal M6P residues.

[0038] In certain embodiments, the oligosaccharide comprising a reactive group may be an M6P-containing hexasaccharide of Formula Ia:

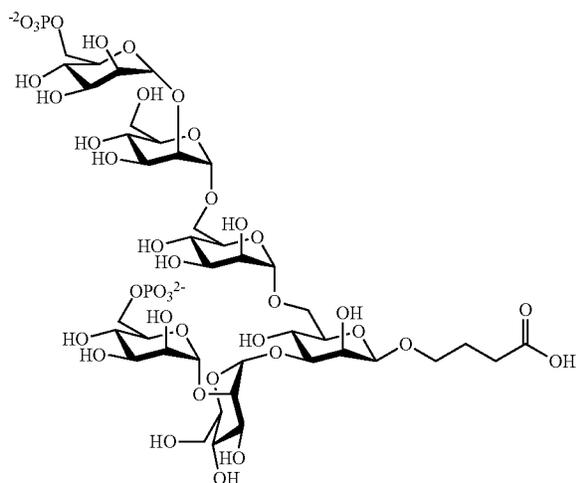
Formula Ia



[0039] The oligosaccharide of Formula Ia can be described as butyrylhydrazone-4-yl 6-O-phosphono- α -D-mannopyranosyl-(1 \rightarrow 2)- α -D-mannopyranosyl-(1 \rightarrow 6)- α -D-mannopyranosyl-(1 \rightarrow 2)- α -D-mannopyranosyl-(1 \rightarrow 3)- β -D-mannopyranoside.

[0040] In certain embodiments, the oligosaccharide comprising a reactive group may be an M6P-containing hexasaccharide of Formula Ib:

Formula Ib

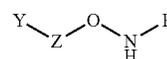


B. Aminoxy Compound

[0041] As used herein, an aminoxy compound may be any compound comprising an aminoxy group and a second reactive group, wherein the second reactive group may react with a first reactive group on an oligosaccharide to form a covalent bond. For example, in some embodiments, the second reactive group may be a carboxyl, ester (such as, e.g., an activated ester), acyl halide (such as, e.g., an acyl fluoride or acyl chloride), acyl azide, anhydride, isothiocyanate, isocyanate, or sulfonyl halide (such as, e.g., a sulfonyl chloride or sulfonyl bromide) group. In other embodiments, the second reactive group may be, e.g., a hydrazine group, hydrazide group, semicarbazide group, thiosemicarbazide, or amine group.

[0042] In certain embodiments, the nitrogen of the aminoxy group of the aminoxy compound is protected with an amino protecting group. Numerous amino protecting groups are known to those skilled in the art, and examples of amino protecting groups, methods for their addition, and methods for their removal can be found in pp. 494-653 of Greene et al., *Protective Groups in Organic Synthesis*, 3rd ed., John Wiley and Sons: New York, 1999; Chapter 8 of Kocienski, *Protecting Groups*, 3rd ed., Georg Thieme Verlag: Stuttgart, Germany, 2005; Bodanszky, *Principles of Peptide Synthesis*, Springer Verlag: New York, 1993; Lloyd-Williams et al., *Chemical Approaches to the Synthesis of Peptides and Proteins*, CRC Press: Boca Raton, Fla., 1997; and Stewart et al., *Solid Phase Peptide Synthesis*, 2nd ed., Pierce Chemical Co.: Rockford, Ill., 1984, the inventions of which are incorporated herein by reference.

[0043] In some embodiments, the aminoxy compound is chosen from compounds of Formula II:

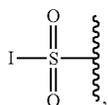
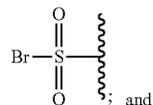
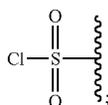
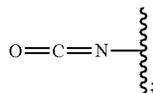
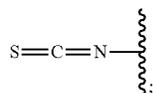
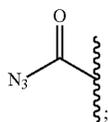
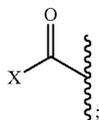
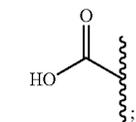


Formula II

wherein Y is the second reactive group, Z is chosen from alkyl, alkenyl, alkynyl, heteroaryl, aryl, and heterocyclyl, and P is chosen from amino protecting groups.

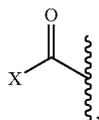
[0044] As used herein, any chemical group on the aminoxy compound (such as, e.g., alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, acyloxy, alkoxy, aryloxy, and heterocyclyloxy) may be substituted or unsubstituted, and may be interrupted by one or more chemical groups, unless otherwise stated. Substituents and interrupting chemical groups may be chosen from, e.g., acyl, acylamino, acyloxy, alkenyl, alkoxy, alkyl, alkynyl, amido, amino, aryl, aryloxy, azido, carbamoyl, carboalkoxy, carboxy, cyano, cycloalkyl, formyl, guanidino, halo, heteroaryl, heterocyclyl, hydroxy, iminoamino, nitro, oxo, phosphonamino, sulfinyl, sulfonamino, sulfonate, sulfonyl, thio, thioacylamino, thioureido, and ureido. The substituents may themselves be substituted or unsubstituted, and may be interrupted or terminated by one or more heteroatoms such as, e.g., nitrogen, sulfur, and oxygen.

[0045] In certain embodiments, Y may be chosen from, for example:

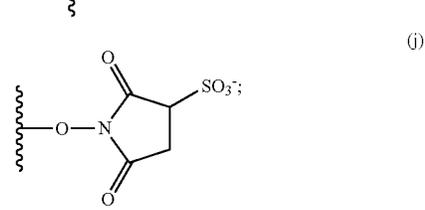
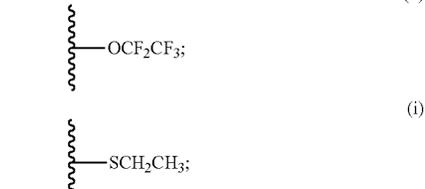
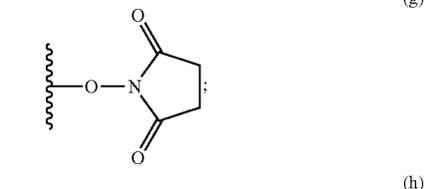
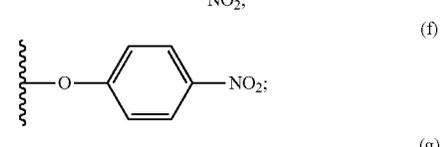
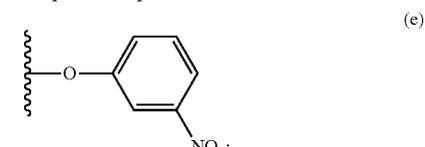
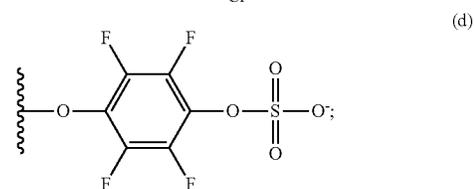
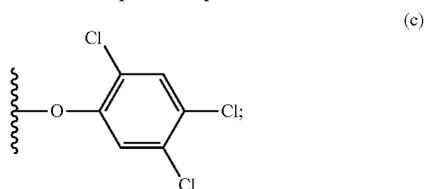
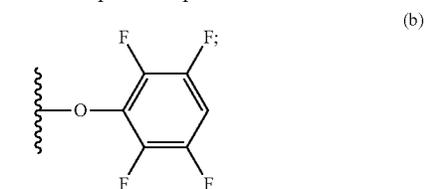
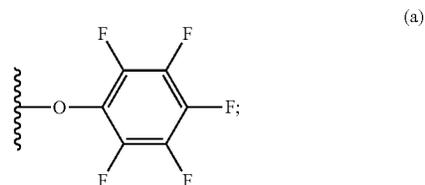


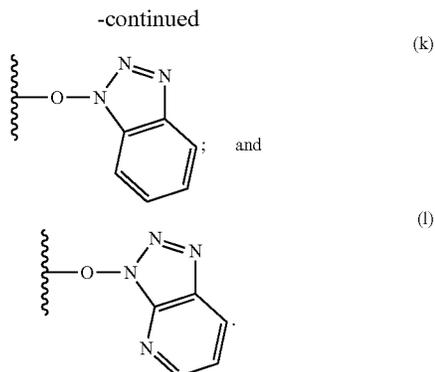
[0046] wherein X is chosen from halogens, azide, acyloxy, alkoxy, aryloxy, heteroaryloxy, and heterocycloxy.

[0047] In certain embodiments, the aminoxy compound is an activated ester. As used herein, an activated ester is an ester that reacts to form an amide bond under mild conditions. In general, an activated ester is an ester of a relatively acidic alcohol. In certain embodiments, the aminoxy compound of Formula II is an activated ester of formula



and X is chosen from alkoxy, aryloxy, heteroaryloxy, and heterocycloxy. For example, X may be chosen from:





[0048] In other embodiments, Y is chosen from, e.g., hydrazide, hydrazine, thiosemicarbazide, semicarbazide, and amine groups.

[0049] In some embodiments, Z may comprise, for example, a carbonyl, ether, ester, or amide group. In some embodiments, Z may be, for example, alkyl interrupted by one or more heteroatoms, such as an oligoethyleneglycol. For example, Z may be monoethyleneglycol, diethyleneglycol, triethyleneglycol, tetraethyleneglycol, or larger oligoethyleneglycol.

[0050] In some embodiments, Z may be, for example, alkyl substituted with oxo and interrupted by one or more heteroatoms, such as an oligopeptide. For example, the oligopeptide may comprise one, two, three, four, five, six, or more component amino acids. The amino acids may be, for example, α -amino acids, β -amino acids, γ -amino acids, δ -amino acids, and co-amino acids. An amino acid may have R or S chirality at any chiral atom. An amino acid may be chosen from, e.g., alanine, β -alanine, α -amino adipic acid, 2-aminobutanoic acid, 4-aminobutanoic acid, 1-aminocyclopentanecarboxylic acid, 6-aminohexanoic acid, 2-aminoheptanedioic acid, 7-aminoheptanoic acid, 2-aminoisobutyric acid, aminomethylpyrrole carboxylic acid, 8-amino-3,6-dioxaoctanoic acid, aminopiperidinecarboxylic acid, 3-amino-propionic acid, aminoserine, aminotetrahydropyran-4-carboxylic acid, arginine, asparagine, aspartic acid, azetidine carboxylic acid, benzothiazolylalanine, butylglycine, carnitine, 4-chlorophenylalanine, citrulline, cyclohexylalanine, cyclohexylstatine, cysteine, 2,4-diaminobutanoic acid, 2,3-diaminopropionic acid, dihydroxyphenylalanine, dimethylthiazolidine carboxylic acid, glutamic acid, glutamine, glycine, histidine, homoserine, hydroxyproline, isoleucine, isonipecotic acid, leucine, lysine, methanoproline, methionine, norleucine, norvaline, ornithine, p-aminobenzoic acid, penicillamine, phenylalanine, phenylglycine, piperidinyllalanine, piperidinyglycine, proline, pyrrolidinyllalanine, sarcosine, selenocysteine, serine, statine, tetrahydropyrynglycine, thienylalanine, threonine, tryptophan, tyrosine, valine, allo-isoleucine, allo-threonine, 2,6-diamino-4-hexanoic acid, 2,6-diaminopimelic acid, 2,3-diaminopropionic acid, dicarboxidine, homoarginine, homocitrulline, homocysteine, homocystine, homophenylalanine, homoproline, and 4-hydrazinobenzoic acid.

[0051] P may be chosen from amino protecting groups known to those of skill in the art. In some embodiments, P may be a carbamate protecting group, such as, e.g., a (9-fluorenylmethyl)carbamate (Fmoc), (tert-butoxy)carbamate (t-Boc), (trichloroethyl)carbamate (Troc), or allyl-carbamate (Alloc) protecting group. In other embodiments, P may be a non-carbamate protecting group, such as, e.g., an

amide protecting group such as a phthalimide or a trifluoroacetamide protecting group.

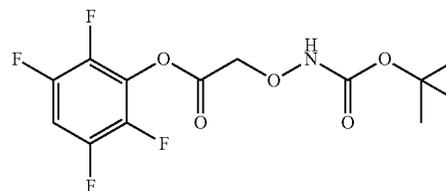
[0052] In some embodiments, the aminoxy compound of Formula II is chosen from compounds of Formula III:



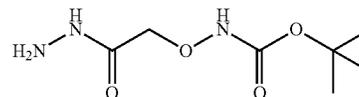
wherein Y and P are as disclosed above, and n is chosen from integers ranging from 1 to 10.

[0053] In certain embodiments, n may be chosen from integers from the following ranges: 1-4, 2-6, 2-8, 3-6, and 4-10. In illustrative embodiments, n is 1.

[0054] In one illustrative embodiment, the aminoxy compound is t-Boc-aminoxy acetic acid tetrafluorophenyl ester, the structure of which is depicted below.



[0055] In another illustrative embodiment, the aminoxy compound has the structure depicted below.



C. Methods of Preparing an Oligosaccharide Comprising an Aminoxy Group

[0056] In another embodiment, the invention provides a method of preparing an oligosaccharide comprising an aminoxy group from an oligosaccharide comprising a reactive group. The method comprises:

[0057] (a) providing an oligosaccharide comprising a first reactive group;

[0058] (b) providing an aminoxy compound comprising a second reactive group; and

[0059] (c) reacting the first reactive group of the oligosaccharide with the second reactive group of the aminoxy compound,

thereby preparing the oligosaccharide comprising an aminoxy group.

[0060] The oligosaccharide comprising a first reactive compound may be, e.g., any oligosaccharide comprising a reactive group as described supra. In illustrative embodiments, the oligosaccharide comprising a first reactive group is an oligosaccharide of Formula Ia or an oligosaccharide of Formula Ib. The aminoxy compound comprising a second reactive group may be any aminoxy compound comprising a reactive group, as described supra.

[0061] The terms “first reactive group” and “second reactive group,” as used herein, do not denote any particular experimental sequence. I.e., step (c), reacting the first reactive group of the oligosaccharide with the second reactive group of the aminoxy compound, may be accomplished by any order of addition of the reactants. For example, the oligosaccharide comprising a first reactive group may be added to the aminoxy compound comprising the second reactive group, or vice versa. In another example, both the

oligosaccharide and the aminoxy compound may be added simultaneously to a reaction vessel.

[0062] Step (c) may occur under any suitable conditions (e.g., solvent and temperature) known to those of ordinary skill in the art. In certain embodiments, one or more additional reagents, such as, e.g., coupling reagents and catalysts, may be present during step (c). A coupling reagent, as used herein, is a reagent that may be used to form a covalent bond between the first reactive group and the second reactive group.

[0063] In some embodiments, such as, e.g., when the first or second reactive group is a carboxyl group, the reaction conditions may comprise a coupling reagent. Coupling reagents may be chosen from, e.g., phosphonium coupling reagents such as, e.g., BOP (benzotriazole-1-yl-oxy-tris(dimethylamino)-phosphonium hexafluorophosphate), PyBOP® (benzotriazole-1-yl-oxy-tris-pyrrolidino-phosphonium hexafluorophosphate), and PyBroP® (bromo-tris-pyrrolidino-phosphonium hexafluorophosphate), and from aminium (uronium) coupling reagents such as, e.g., HBTU (2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate), HATU (2-(7-Aza-1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate), TBTU (2-(1H-Benzotriazole-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate). Coupling reagents may also be chosen from, e.g., carbodiimide coupling reagents such as, e.g., DIC (1,3-diisopropylcarbodiimide), CDI (1,1' carbonyl diimidazole), and EDC (1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide). For example, in some illustrative embodiments, the coupling reagent is EDC. In certain embodiments, the reaction conditions comprise both a coupling reagent and a catalyst.

[0064] The reaction conditions may, in certain embodiments, comprise a catalyst. The catalyst may be chosen from any suitable catalyst known to those of skill in the art, such as, e.g., DHBt-OH (3-hydroxy-1,2,3-benzotriazin-4(3H)-

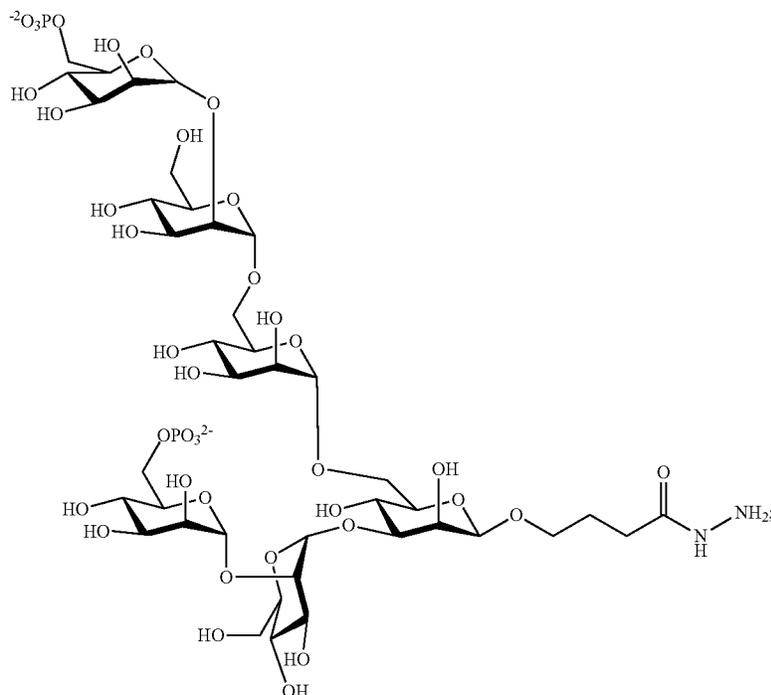
one), HOBt (N-hydroxybenzotriazole), DMAP (4-dimethylaminopyridine), NHS (N-hydroxysuccinimide), N-hydroxy-sulfosuccinimide, HONB (N-hydroxy-5-norbornene-endo-2,3-dicarboximide), or a tetrabutylammonium salt such as, e.g., TBAI (tetrabutylammonium iodide). In some illustrative embodiments, the reaction conditions comprise the catalyst DHBt-OH or the catalyst NHS.

[0065] In some embodiments, step (c), reacting the first reactive group of the oligosaccharide with the second reactive group of the aminoxy compound results in the formation of an amide bond. Conditions suitable for the formation of an amide bond are well known to those of ordinary skill in the art, and are described in, e.g., Chan et al., eds., *Fmoc Solid Phase Peptide Synthesis: A Practical Approach*, Oxford University Press: New York, 2000; Bodanszky, *Principles of Peptide Synthesis*, Springer Verlag: New York, 1993; Lloyd-Williams et al., *Chemical Approaches to the Synthesis of Peptides and Proteins*, CRC Press: Boca Raton, Fla., 1997; and the Novabiochem® (San Diego, Calif.) Catalog.

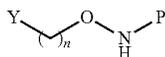
[0066] In certain embodiments, the aminoxy compound comprises an amino protecting group, and the method comprises a further step (d), deprotecting the oligosaccharide comprising an aminoxy group to remove the amino protecting group. Deprotection may occur under any suitable conditions known to those of skill in the art, such as, e.g., those taught in pp. 494-653 of Greene et al., *Protective Groups in Organic Synthesis*, 3rd ed., John Wiley and Sons: New York, 1999 and Kocienski, *Protecting Groups*, 3rd ed., Georg Thieme Verlag: Stuttgart, Germany, 2005, the inventions of which are incorporated herein by reference.

[0067] An illustrative embodiment of the method of the invention provides a method of preparing an M6P-containing oligosaccharide comprising an aminoxy group. The method comprises:

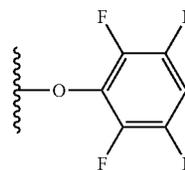
[0068] (a) providing an oligosaccharide comprising a first reactive group, wherein the oligosaccharide is



[0069] (b) providing an aminoxy compound comprising a second reactive group, wherein the aminoxy compound is chosen from compounds of Formula III:



Formula III

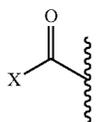


[0070] wherein n is chosen from integers ranging from 1 to 10, P is chosen from amino protecting groups, and Y is a second reactive group; and

[0071] (c) reacting the first reactive group of the oligosaccharide with the second reactive group of the aminoxy compound,

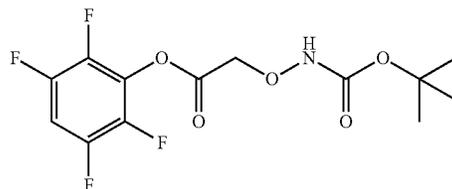
thereby preparing the oligosaccharide comprising an aminoxy group.

[0072] In certain embodiments, Y in Formula III is



where X is chosen from hydroxy, aryloxy, heteroaryloxy, and heterocyclyloxy. For example, in certain illustrative embodiments, X is

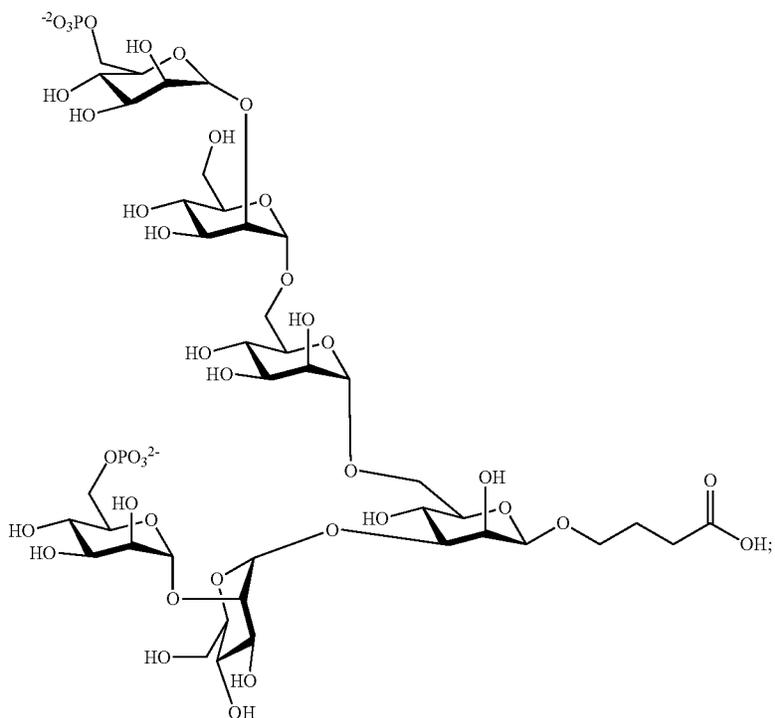
[0073] In illustrative embodiments, the aminoxy compound is



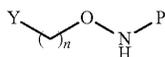
In certain embodiments, the first reactive group of the oligosaccharide may be reacted with the second reactive group of the aminoxy compound in the presence of a coupling agent, such as, e.g., EDC, and/or a catalyst, such as, e.g., DHBt-OH.

[0074] Another illustrative embodiment of the method of the invention comprises:

[0075] (a) providing an oligosaccharide comprising a first reactive group, wherein the oligosaccharide is



[0076] (b) providing an aminoxy compound comprising a second reactive group, wherein the aminoxy compound is chosen from compounds of Formula III:



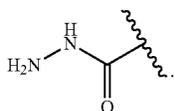
Formula III

[0077] wherein n is chosen from integers ranging from 1 to 10, P is chosen from amino protecting groups, and Y is a second reactive group; and

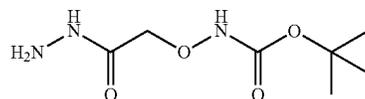
[0078] (c) reacting the first reactive group of the oligosaccharide with the second reactive group of the aminoxy compound,

thereby preparing the oligosaccharide comprising an aminoxy group.

[0079] In certain embodiments, Y in Formula III is a hydrazine, hydrazide, aminoxy, thiosemicarbazide, semicarbazide, or amine group. In certain embodiments, Y in Formula III is



In illustrative embodiments, the aminoxy compound is



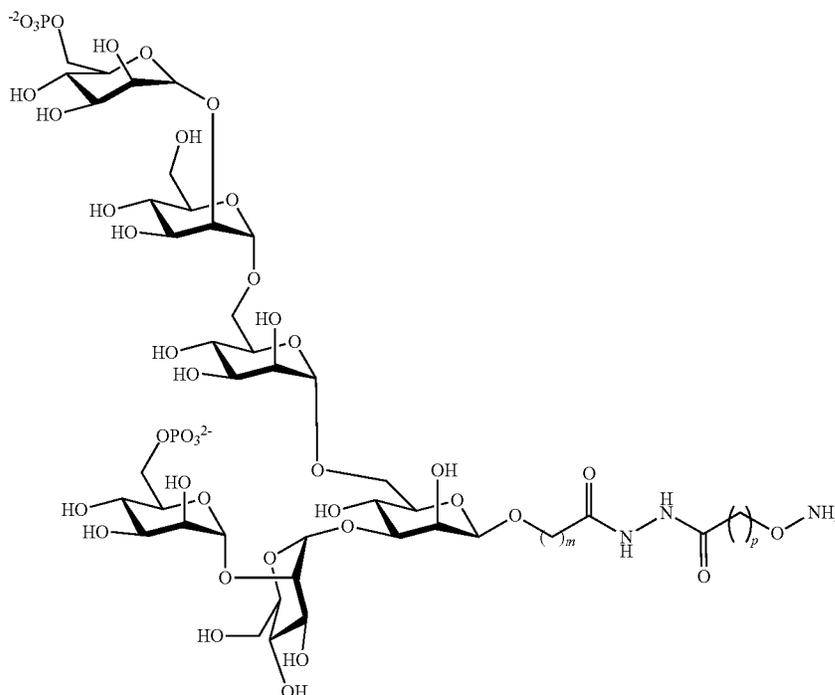
In certain embodiments, the first reactive group of the oligosaccharide may be reacted with the second reactive group of the aminoxy compound in the presence of a coupling agent, such as, e.g., EDC, and/or a catalyst, such as, e.g., NHS.

II. OLIGOSACCHARIDES COMPRISING AN AMINOXY GROUP

[0080] The present invention also provides oligosaccharides comprising an aminoxy group. In some embodiments, the invention provides oligosaccharides comprising an aminoxy group prepared by the methods disclosed above. The oligosaccharide comprising an aminoxy group may comprise, for example, at least 2, 3, 4, 5, 6, or more monosaccharides, including, e.g., at least one galactose, GalNAc, mannose, M6P, glucose, GlcNAc, sialic acid, or sulfated sialic acid residue. Such an oligosaccharide may be mono-, bi-, tri-, tetra-, or penta-antennary in structure, and may contain 0, 1, 2, 3, 4, or more branch points.

[0081] In some embodiments, the present invention provides an oligosaccharide comprising (1) an aminoxy group and (2) mannose-6-phosphate. The oligosaccharide comprising an aminoxy group may, in some embodiments, comprise, e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 M6P residues. In some embodiments, the oligosaccharide comprising an aminoxy group may comprise at least 1, 2, 3, 4, or more terminal or penultimate M6P residues.

[0082] The oligosaccharides comprising an aminoxy group are, in certain embodiments, chosen from oligosaccharides of Formula IV:

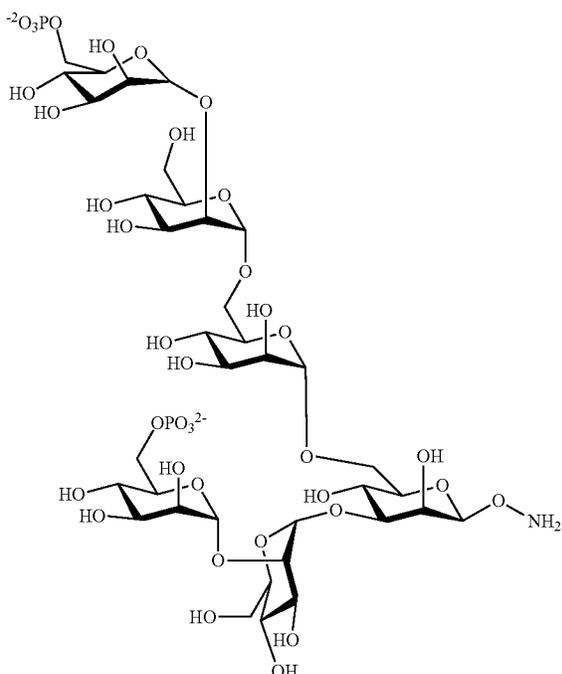


Formula IV

wherein m and p are independently chosen from integers ranging from 1 to 10. For example, in certain embodiments, m and p may be independently chosen from integers selected from the following ranges: 1-4, 2-6, 2-8, 3-6, and 4-10. In illustrative embodiments, m is 3 and p is 1.

[0083] In other embodiments, the aminoxy group is directly linked to the reducing end of the oligosaccharide. For example, in some embodiments, the oligosaccharide comprising an aminoxy group may be an oligosaccharide of Formula V:

Formula V



III. CONJUGATION OF AN OLIGOSACCHARIDE COMPRISING AN AMINOXY GROUP WITH A PROTEIN

A. Oligosaccharide

[0084] The oligosaccharide to be conjugated with a protein may be chosen from any oligosaccharide comprising a reactive group, as discussed supra, and from any oligosaccharide comprising an aminoxy group, as discussed supra. For example, in some embodiments, the oligosaccharide to be conjugated may be an oligosaccharide of Formula Ia, Formula Ib, Formula IV or Formula V.

B. Protein

[0085] The conjugation methods described herein are broadly applicable to any pure protein, partially purified protein, or fragment thereof, having at least one carbonyl group (where a carbonyl group is a ketone or an aldehyde), including isolated proteins and recombinantly or synthetically produced proteins. The terms “pure,” “purified,” and “isolated” refer to a molecule that is substantially free of its natural environment. For instance, a pure protein is substantially free of cellular material and/or other proteins from the cell or tissue source from which it is derived. The term refers

to preparations that are, for example, at least 70% to 80%, 80% to 90%, 90 to 95%; or at least 95%, 96%, 97%, 98%, 99%, or 100% (w/w) pure.

[0086] In other embodiments, the protein may be an enzyme that has optimal activity, as measured by an activity assay, at a pH ranging from 1-7, such as, e.g., 1-3, 2-5, 3-6, 4-5, 5-6, or 4-6. For example, the enzyme may have a pH optimum at a pH ranging from 4-6.

[0087] In some embodiments, the protein may be an enzyme that has an isoelectric point (pI), ranging from 1 to 8, such as, e.g., from 1-3, 2-5, 3-8, 4-5, 5-6, 4-6, 5-8, 6-8, or 7-8. The pI of a protein may be measured using, e.g., isoelectric focusing gel electrophoresis.

[0088] In some embodiments, the protein containing a carbonyl group is obtained by the use of an expression system having an expanded genetic code, as described in, e.g., Wang et al., *Proc. Natl. Acad. Sci. USA* 100:56-61 (2003). In such a case, the carbonyl group may be located on amino acid side chain, as translated.

[0089] In certain embodiments, the protein having at least one carbonyl group is a protein having at least one oligosaccharide (i.e., a glycoprotein). For example, a glycoprotein having at least one carbonyl group may be obtained by oxidation of that glycoprotein by any means known to those of skill in the art. In some embodiments, e.g., a glycoprotein having at least one carbonyl group may be obtained by oxidation of that glycoprotein with periodate (e.g., sodium periodate) or with galactose oxidase. In such a case, the carbonyl group may be located at a protein glycosylation site.

[0090] In certain embodiments, the protein having at least one carbonyl group is a glycoprotein, such as a therapeutic glycoprotein. A therapeutic glycoprotein may be targeted to the lysosome by conjugation with an oligosaccharide comprising mannose-6-phosphate. For example, the glycoprotein may be a lysosomal enzyme, including an ERT enzyme. The enzyme may be a lysosomal hydrolase, including those listed in Table 1. In certain embodiments, the lysosomal hydrolase is chosen from, e.g., α -glucosidase, α -galactosidase A, and acid sphingomyelinase. In certain embodiments, the lysosomal hydrolase is GAA.

TABLE 1

Examples of LSDs and Corresponding Lysosomal Hydrolases	
Lysosomal Storage Disorder	Defective Enzyme
Fabry	α -Galactosidase A
Farber	Acid ceramidase
Fucosidosis	Acid α -L-fucosidase
Gaucher types 1, 2, and 3	Acid β -glucosidase
G_{M1} gangliosidosis	Acid β -galactosidase
Hunter (Mucopolysaccharidosis (MPS) II)	Iduronate-2-sulfatase
Hurler-Scheie, Hurler, Scheie (MPS I)	α -L-Iduronidase
Krabbe	Galactocerebrosidase
α -Mannosidosis	Acid α -mannosidase
β -Mannosidosis	Acid β -mannosidase
Maroteaux-Lamy (MPS VI)	Arylsulfatase B
Metachromatic leukodystrophy	Arylsulfatase A
Morquio A (MPS IV)	N-Acetylgalactosamine-6-sulfate sulfatase
Morquio B (MPS IV)	Acid β -galactosidase
Niemann-Pick A and B	Acid sphingomyelinase (ASM)
Pompe	Acid α -glucosidase (α -glucosidase; GAA)
Sandhoff	β -Hexosaminidase B
Sanfilippo A (MPS III)	Heparan N-sulfatase
Sanfilippo B (MPS III)	α -N-Acetylglucosaminidase

TABLE 1-continued

Examples of LSDs and Corresponding Lysosomal Hydrolases	
Lysosomal Storage Disorder	Defective Enzyme
Sanfilippo C (MPS III)	Acetyl-CoA: α -glucosaminide N-acetyltransferase
Sanfilippo D (MPS III)	N-Acetylglucosamine-6-sulfate sulfatase
Schindler-Kanzaki	α -N-acetylgalactosaminidase
Sialidosis	Sialidase
Sly (MPS VII)	β -Glucuronidase
Tay-Sachs	β -Hexosaminidase A

[0091] In certain embodiments, the glycoprotein may be a glycoprotein having at least 1, 2, 3, 4, 5, or more N-linked or O-linked glycosylated amino acid residues. In other embodiments, the protein may have 1, 2, 3, 4, 5 or more consensus sites for N-linked or O-linked glycosylation, at least one of which is glycosylated.

[0092] In certain embodiments, the protein may be a ligand for a receptor. For example, in some embodiments the protein may be a glycoprotein that binds to a receptor that recognizes a sugar such as, e.g., mannose or mannose-6-phosphate. In some embodiments, the glycoprotein may bind to, e.g., the asialoglycoprotein receptor, the cation-dependent mannose-6-phosphate receptor, the insulin-like growth factor II/cation-independent mannose-6-phosphate receptor, or the macrophage mannose receptor.

[0093] In certain embodiments, the protein is a glycoprotein that, when conjugated to an oligosaccharide comprising mannose-6-phosphate, is internalized more efficiently by a target cell (e.g., via CI-MPR-mediated endocytosis) than is the corresponding unconjugated glycoprotein. For example, the conjugated glycoprotein may be internalized more efficiently than the unconjugated glycoprotein by, e.g., at least 10%, 15%, 20%, 25%, 30%, 35%, 40%, 50%, 60%, 70%, 80%, or 90% (w/w) in a given time period. In other embodiments, at least 2, 3, 4, 5, 6, 7, 8, 9, or 10 fold (w/w) as much of the conjugated glycoprotein may be internalized, relative to the unconjugated glycoprotein, in a given time period. The referenced time period may be, for example, 10, 30, 45 minutes or 1, 2, 3, 5, 6, 12, 24, 48, or 72 hours, or more.

C. Methods of Coupling an Oligosaccharide to a Protein

[0094] The invention provides methods of coupling an oligosaccharide to a protein, such as, e.g., a glycoprotein. In one embodiment, the method comprises:

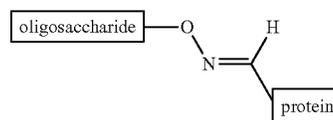
- [0095]** (a) providing an oligosaccharide comprising an aminoxy group;
- [0096]** (b) providing a protein having at least one carbonyl group; and
- [0097]** (c) reacting the aminoxy group of the oligosaccharide with the at least one carbonyl group of the protein,

thereby coupling the oligosaccharide to the protein.

[0098] In certain embodiments, the methods further comprise adding a reducing agent to the coupled lysosomal enzyme. The reducing agent may be any reducing agent known to those of skill in the art, such as, e.g., sodium cyanoborohydride or sodium triacetoxyborohydride (STAB).

IV. OLIGOSACCHARIDE-PROTEIN CONJUGATES

[0099] The invention further provides an oligosaccharide-protein conjugate, comprising (1) a protein, (2) an oligosaccharide, and (3) an oxime group connecting the protein and the oligosaccharide. In some embodiments, the invention provides an oligosaccharide-protein conjugate prepared by the methods disclosed above. The oligosaccharide and protein components of the conjugate may be, for example, any oligosaccharide and protein described herein, wherein a conjugate thereof comprises an oxime group, as depicted below. (The oxime group depicted below is formally derived by reaction of an aminoxy group and an aldehyde group; oxime groups formally derived by reaction of an aminoxy group and a ketone group are also encompassed by this invention.)



[0100] In certain embodiments, the oligosaccharide-protein conjugate is an oligosaccharide-glycoprotein conjugate. In certain embodiments, the oligosaccharide-protein conjugate is the conjugate of an oligosaccharide comprising at least one M6P and of a lysosomal hydrolase.

V. PHARMACEUTICAL COMPOSITIONS

[0101] This disclosure provides the use of a conjugate of the invention in the manufacture of a medicament for treating a lysosomal storage disorder. It also provides pharmaceutical compositions comprising an oligosaccharide-protein conjugate of the invention. In some embodiments, the pharmaceutical compositions of the invention comprise a conjugate of an oligosaccharide comprising at least one M6P and a lysosomal enzyme.

[0102] Pharmaceutical compositions of the invention may comprise one or more suitable pharmaceutical excipients. Standard pharmaceutical formulation techniques and excipients are well known to persons skilled in the art (see, e.g., 2005 *Physicians' Desk Reference*[®], Thomson Healthcare: Montvale, N.J., 2004; Remington: The Science and Practice of Pharmacy, 20th ed., Gennado et al., Eds. Lippincott Williams & Wilkins: Philadelphia, Pa., 2000). The compositions may or may not contain preservatives. In some embodiments, pharmaceutical compositions comprising α -galactosidase A conjugates may comprise one or more excipients such as, e.g., mannitol, sodium phosphate monobasic monohydrate, and/or sodium phosphate dibasic heptahydrate. In some embodiments, pharmaceutical compositions comprising conjugates of α -glucosidase may comprise one or more excipients such as, e.g., mannitol, polysorbate 80, sodium phosphate dibasic heptahydrate, and sodium phosphate monobasic monohydrate.

[0103] The pharmaceutical composition may comprise any of the conjugates described herein either as the sole active compound or in combination with another compound, composition, or biological material. For example, the pharmaceutical composition may also comprise one or more small molecules useful for the treatment of a LSD and/or a side effect associated with the LSD. In some embodiments, the composition may comprise miglustat and/or one or more compounds described in, e.g., U.S. Patent Application Publication Nos. 2003/0050299, 2003/0153768; 2005/0222244; 2005/0267094.

[0104] The formulation of pharmaceutical compositions may vary depending on the intended route of administrations

and other parameters (see, e.g., Rowe et al. *Handbook of Pharmaceutical Excipients*, 4th ed., APhA Publications, 2003.) In some embodiments, the composition may be a sterile, non-pyrogenic, white to off-white lyophilized cake or powder to be administered by intravenous injection upon reconstitution with Sterile Water for Injection, USP.

[0105] Administration of a pharmaceutical composition of the invention is not limited to any particular delivery system and may include, without limitation, parenteral (including subcutaneous, intravenous, intracranial, intramedullary, intraarticular, intramuscular, intrathecal, or intraperitoneal injection), transdermal, or oral (for example, in capsules, suspensions, or tablets). Administration to an individual may occur in a single dose or in repeat administrations, and in any of a variety of physiologically acceptable salt forms, and/or with an acceptable pharmaceutical carrier and/or additive as part of a pharmaceutical composition.

[0106] The conjugates described herein are administered in therapeutically effective amounts. Generally, a therapeutically effective amount may vary with the subject's age, condition, and sex, as well as the severity of the medical condition in the subject. The dosage may be determined by a physician and adjusted, as necessary, to suit observed effects of the treatment. Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in vitro (i.e., cell cultures) or in vivo (i.e., experimental animal models), e.g., for determining the LD₅₀ (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index (or therapeutic ratio), and can be expressed as the ratio LD₅₀/ED₅₀. Conjugates that exhibit therapeutic indices of at least 1, 1.5, 2, 3, 4, 5, 6, 7, 8, 9, 10, and 20 are described herein. Conjugates that exhibit a large therapeutic index are preferred.

[0107] The data obtained from in vitro assays and animal studies, for example, can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED₅₀ with low, little, or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. For any conjugate used in the present invention, the therapeutically effective dose can be estimated initially from in vitro assays. A dose may be formulated in animal models to achieve a circulating plasma concentration range that includes the IC₅₀ (i.e., the concentration of the test conjugate which achieves a half-maximal inhibition of symptoms) as determined in in vitro experiments. Levels in plasma may be measured, for example, by high performance liquid chromatography or by an appropriate enzymatic activity assay. The effects of any particular dosage can be monitored by a suitable bioassay of endpoints.

[0108] Unless otherwise indicated, conjugates of the invention may be administered at a dose of approximately from 1 µg/kg to 500 mg/kg, depending on the severity of the symptoms and the progression of the disease. For example, proteinaceous compounds may be administered by slow intravenous infusion in an outpatient setting every, e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more days, or by, e.g., weekly, biweekly, monthly, or bimonthly administration. The appropriate therapeutically effective dose of a compound is selected by a treating clinician and would range approximately from 1 µg/kg to 500 mg/kg, from 1 µg/kg to 10 mg/kg, from 1 µg/kg to 1 mg/kg, from 10 µg/kg to 1 mg/kg, from 10 µg/kg to 100 µg/kg, from 100 µg to 1 mg/kg, and from 500 µg/kg to 5 mg/kg.

[0109] For example, conjugates of α-galactosidase A may be administered by intravenous infusion at a dose of, e.g., 1.0 mg/kg body weight every two weeks at an infusion rate

of, e.g., less than or equal to 10, 13, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, or 33 mg/hour). In another example, conjugates of α-glucosidase may be administered intravenous injection at a dose of, e.g., 20 mg/kg or 40 mg/kg every two weeks, over approximately, e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 hours. In some embodiments, the rate of administration of α-glucosidase may be started at, e.g., 1 mg/kg/hr and then increased by, e.g., 2 mg/kg/hr every 30 minutes, after establishing patient tolerance to the infusion rate, until a maximum of, e.g., 7 mg/kg/hr. Additionally, examples of specific dosages may be found in the *Physicians' Desk Reference*®.

VI. METHODS OF TREATING LYSOSOMAL STORAGE DISORDERS

[0110] The invention provides methods of treating lysosomal storage disorders, such as, e.g., those disclosed in Table 1. In some embodiments, the invention provides the use of a conjugate described herein for treating a lysosomal storage disorder in a subject in need thereof. The invention further provides methods of targeting proteins to the lysosome by conjugation with oligosaccharides comprising mannose-6-phosphate.

[0111] In one embodiment, the method comprises administering to a mammal having a lysosomal storage disorder an oligosaccharide-glycoprotein conjugate of the invention in a therapeutically effective amount. The oligosaccharide-glycoprotein conjugate may be a conjugate of a lysosomal enzyme, such as a lysosomal enzyme listed in Table 1, with an oligosaccharide comprising mannose-6-phosphate. In one embodiment, the method comprises administering to a subject in need thereof a pharmaceutical composition comprising at least one of the conjugates described herein.

[0112] In certain embodiments, conjugates of the invention may be administered with one or more other therapies. The one or more other therapies may be administered concurrently with (including concurrent administration as a combined formulation), before, or after the administration of the conjugates of the invention.

[0113] In some embodiments, a patient may be treated (before, after, or during treatment with a conjugate of the invention) with an antipyretic, antihistamine, and/or immunosuppressant. In some embodiments, a patient may be treated with an antipyretic, antihistamine, and/or immunosuppressant prior to treatment with an oligosaccharide-glycoprotein conjugate of the invention in order to decrease or prevent infusion associated reactions. For example, patients may be pretreated with one or more of acetaminophen, azathioprine, cyclophosphamide, cyclosporin A, methotrexate, mycophenolate mofetil, oral steroids, or rapamycin.

[0114] In some embodiments, patients may be treated with one or more of acetaminophen, azathioprine, cyclophosphamide, cyclosporin A, methotrexate, mycophenolate mofetil, oral steroids, or rapamycin at or about, e.g., t=0 (the time of administration of the conjugate of the invention) and/or t=12, 24, 36, 48, 60, 72, 96, 120, and 144 hours for, e.g., the first 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, or more incidences of treatment with a conjugate of the invention. For example, in some embodiments a patient with Fabry disease or Pompe disease may be treated with methotrexate (e.g., with 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9, 1, 2, 3, 4, 5, 6, 8, 10, 12, 15, 25, 30, 35, 40, 50, 60, 70, 80 mg/kg methotrexate, or more) at or about, e.g., t=0, 24, and 48 hours for, e.g., the first 1, 2, 3, 4, 5, 6, 7, 8 weeks of treatment with a conjugate of the invention. In some embodiments, immune tolerance toward conjugates of the invention may be induced in a patient with a lysosomal storage disorder such as, e.g., mucopolysaccharidosis I, by treatment with cyclosporin A and azathioprine.

For example, the patient may be treated with cyclosporine A and azathioprine as described in Kakkis et al., *Proc. Natl. Acad. Sci. U.S.A.* 101:829-834 (2004).

[0115] In some embodiments, a patient may be treated (before, after, or during treatment with a conjugate of the invention) with e.g., small molecule therapy and/or gene therapy, including small molecule therapy and gene therapy directed toward treatment of a lysosomal storage disorder. Small molecule therapy may comprise administration of one or more compounds described in, e.g., U.S. Patent Application Nos. 2003/0050299, 2003/0153768; 2005/0222244; and 2005/0267094. Gene therapy may be performed as described in, e.g., U.S. Pat. Nos. 5,952,516; 6,066,626; 6,071,890; and 6,287,857 and U.S. Patent Application Publication No. 2003/0087868.

[0116] The terms "treatment," "therapeutic method," and their cognates refer to both therapeutic treatment and prophylactic/preventative measures. Thus, those in need of treatment may include individuals already having a particular lysosomal storage disease as well as those at risk for the disease (i.e., those who are likely to ultimately acquire the disorder or certain symptoms of the disorder).

[0117] A therapeutic method results in the prevention or amelioration of symptoms or an otherwise desired biological outcome, and may be evaluated by improved clinical signs or delayed onset of disease, increased activity of the metabolically defective enzyme, and/or decreased levels of the accumulated substrate of the metabolically defective enzyme.

[0118] The conjugates of the present invention are useful to treat various lysosomal storage disorders in humans or animals. For example, administration of the conjugates can be used to increase the deficient enzymatic activity in a patient, for example, by at least 10%. The increased enzymatic activity may be determined by, e.g., a reduction in clinical symptoms or by an appropriate clinical or biological assay.

[0119] GAA conjugates may be administered for the treatment of Pompe disease (also known as acid α -glucosidase deficiency, acid maltase deficiency, glycogen storage disease type II, glycogenosis II, and lysosomal α -glucosidase deficiency). Increased GAA activity may be determined by biochemical (see, e.g., Zhu et al., *J. Biol. Chem.* 279:50336-50341 (2004)) or histological observation of reduced lysosomal glycogen accumulation in, e.g., cardiac myocytes, skeletal myocytes, or skin fibroblasts. GAA activity may also be assayed in, e.g., a muscle biopsy sample, in cultured skin fibroblasts, in lymphocytes, and in dried blood spots. Dried blood spot assays are described in e.g., Umphathisviam et al., *Clin. Chem.* 47:1378-1383 (2001) and Li et al., *Clin. Chem.* 50:1785-1796 (2004). Treatment of Pompe disease may also be assessed by, e.g., serum levels of creatinine kinase, gains in motor function (e.g., as assessed by the Alberta Infant Motor Scale), changes in left ventricular mass index as measured by echocardiogram, and cardiac electrical activity, as measured by electrocardiogram. Administration of GAA conjugates may result in a reduction in one or more symptoms of Pompe disease such as cardiomegaly, cardiomyopathy, daytime somnolence, exertional dyspnea, failure to thrive, feeding difficulties, "floppiness," gait abnormalities, headaches, hypotonia, organomegaly (e.g., enlargement of heart, tongue, liver), lordosis, loss of balance, lower back pain, morning headaches, muscle weakness, respiratory insufficiency, scapular winging, scoliosis, reduced deep tendon reflexes, sleep apnea, susceptibility to respiratory infections, and vomiting.

[0120] In another aspect, conjugates of α -galactosidase A with oligosaccharides comprising M6P are administered for the treatment of Fabry disease. Fabry disease, or Anderson-Fabry disease, is a rare, X-linked, lysosomal storage disorder

marked by a deficiency of α -galactosidase A, and results in accumulation of globotriaosylceramide (GL3) and other neutral glycosphingolipids in the lysosomes of visceral tissues and endothelial, perithelial, and muscle cells. Accumulation of the neutral glycosphingolipids in the vasculature results in narrowing and dilatation of the blood vessels, and ultimately to ischemia and infarction.

[0121] Administration of α -galactosidase A conjugates may result in a reduction in one or more clinical symptoms of Fabry disease including, e.g., acroparesthesia, angina, angiokeratoma, arrhythmia, ataxia of gait, burning and/or tingling pain in the hands and feet, cataracts, cold intolerance, conduction abnormalities, corneal whorling, coronary artery disease, dementia, depression, diarrhea, dilated cardiac chambers, dizziness, cardiomegaly, cardiomyopathy, diplopia, dysarthria, fatigue, fever with elevated erythrocyte sedimentation rate, hearing problems, heart disease, heart valve problems, heat intolerance, hemiataxia, hemiparesis, hypohidrosis, impaired sweating, infarction, ischemia, joint pain, kidney disease, left ventricular hypertrophy, lenticular abnormalities, lenticular opacity, lipiduria, muscle weakness, myocardial infarction, nausea, nystagmus, pain (e.g., intense pain radiating throughout the body), polydipsia, proteinuria, post-prandial pain, renal failure, retinal abnormalities, ringing in ears, stomach pain, ST-T wave changes, stroke, uremia, valvular disease, vertigo, vomiting, and weakness. Administration of α -galactosidase A conjugates may result in increased α -galactosidase A activity in, e.g., plasma, tears, leukocytes, biopsied tissues, or cultured skin fibroblasts. Administration of α -galactosidase A conjugates may also result in a histologic finding of a reduction (e.g., of at least 10%) or lack of increase of birefringent lipid globules. It may also result in a decrease in lipid globules in urinary sediment, improved renal function as measured by serum creatinine levels or creatinine clearance, and reduced proteinuria. Administration of α -galactosidase A conjugates may also result in a reduction in GL3 inclusions in the capillary endothelium of the kidney, heart, and skin. Additional assays for measuring efficacy of treatment for Fabry disease can be found in, e.g., MacDermott et al., *J. Med. Genet.* 38:750-760 (2001).

[0122] In yet another aspect, conjugates of acid sphingomyelinase are administered for treatment of Niemann-Pick disease, or acid sphingomyelinase deficiency. Administration of acid sphingomyelinase conjugates may result in a reduction in one or more clinical symptoms of Niemann-Pick disease including, e.g., abnormal cholesterol levels, abnormal lipid levels, ataxia, blood abnormalities, cherry red spots in the eye, frequent lung infections, growth retardation, hepatosplenomegaly, low numbers of platelets, lymphadenopathy, peripheral neuropathy, problems with lung function, shortness of breath, skin pigmentation changes, or xanthomas. In some embodiments, conjugates may be administered intracranially.

[0123] An alternative embodiment relates to treatment of mucopolysaccharidosis I (including, e.g., Hurler and Hurler-Scheie forms of MPS I) with conjugates comprising α -L-iduronidase. Administration of α -L-iduronidase conjugates may result in a reduction in one or more clinical symptoms of MPS I including, e.g., aortic regurgitation, aortic stenosis, carpal tunnel syndrome, chronic rhinitis, conductive hearing loss, constipation, corneal clouding, developmental delay, diarrhea, distended abdomen, dorsolumbar kyphosis, gibbus deformity of the back, hepatosplenomegaly, hydrocephalus, inguinal hernia, kyphosis, mental retardation, mitral regurgitation, mitral stenosis, night-blindness, open-angle glaucoma, poor hand function, progressive arthropathy, recurrent respiratory infections, respiratory insufficiency, retinal degeneration, scoliosis, sensorineural hearing loss, severe

back pain, rhinorrhea, sleep apnea, spinal cord compression, thenar atrophy, umbilical hernia, and upper airway complications.

[0124] The foregoing and the following description are exemplary and explanatory only and are not restrictive of the invention, as claimed.

EXAMPLES

[0125] Examples 1-4 below describe the synthetic route depicted in FIG. 1. Compounds 1, 2, 3, and 4, as used below, have the chemical structures depicted in FIG. 1.

Example 1

Synthesis of Oligosaccharide 3

[0126] 100 mg of oligosaccharide 1 (MW=1250; bisM6P-hydrazide, supplied by Biomira Inc., Edmonton, Canada) was dissolved in 15 ml of DMSO/H₂O (50:50 in volume), yielding a 5.3 μmol/ml solution. 100 mg of t-Boc-aminoxy acetic acid tetrafluorophenyl ester 2 (Invitrogen Corp.; Carlsbad, Calif.; catalog # B3030) was dissolved in 7.5 ml of DMSO. 15 ml of the oligosaccharide solution was then mixed with 7.5 ml of the solution of 2 in a glass bottle, such that the molar ratio of compound 2:compound 1 in the resulting solution was 4:1. 744 μl of DHBt-OH (from a 32.06 mg/ml stock in DMSO) was added to the reaction mixture in a glass bottle, such that the final ratio of compound 2:DHBt-OH is 1:0.5. The mixture was gently shaken at room temperature (25° C.) at 100 RPM overnight for about 18 hours.

[0127] The following morning, 10 μl of the reaction mixture was removed for Dionex analysis to confirm completion of the reaction. The results, depicted in FIG. 2, indicated 100% conversion from 1 to 3.

Example 2

Purification of Oligosaccharide 3

[0128] Method A.

[0129] The reaction mixture was diluted with an equal volume of H₂O and dialyzed in dialysis tubing with molecular weight cutoff of 1000 Dalton (SpectraPor Inc.) twice against 4 L of H₂O at 4° C. for at least 3 hours each. The samples were then lyophilized.

[0130] Method B.

[0131] A Sephadex G-10 gel permeation chromatography column with a bed volume of 225 ml was packed and equilibrated with deionized water. The reaction mixture was loaded onto the column, drained by gravity, and then eluted with deionized water at a flow rate of 75 ml per hour. 4.5 ml fractions were collected with a fraction collector. Fractions 10-23, which contained oligosaccharide 3, were collected, combined and lyophilized. The other small molecules, including t-Boc-AOAA, DHBt-OH, and DMSO, eluted in the later fractions, and were discarded.

Example 3

Deprotection of Oligosaccharide 3

[0132] The t-Boc group of the lyophilized sample was deprotected in 5 ml of 50% trifluoroacetic acid (TFA) in dichloromethane (DCM) in a glass bottle for 30 min with gentle shaking at 100 RPM. The TFA/DCM was then removed by a stream of N₂ in a chemical hood.

Example 4

Purification of Oligosaccharide 4

[0133] Method A.

[0134] After removing the TFA/DCM, the residue was dissolved in 10 ml of 0.5 M sodium acetate buffer, pH 5, and transferred to dialysis tubing with a molecular weight cutoff of 1000 Dalton. The bottle was washed with 4 ml of the same buffer, which was then transferred to the dialysis tubing. The sample was dialyzed twice against 3 L of 25 mM sodium acetate buffer, pH 7, for at least 3 hours, and then transferred to 4 L ice-cold H₂O for overnight dialysis. The sample was recovered from the dialysis tubing and lyophilized.

[0135] Method B.

[0136] After removing the TFA/DCM, the residue was dissolved in 5 ml of 0.5 M sodium acetate buffer, pH 7.5, and loaded onto a Sephadex G-10 gel permeation chromatography column as in Example 2, Method B. The reaction mixture was loaded onto the column, drained by gravity, and then eluted with deionized water at a flow rate of 75 ml per hour. 4.5 ml fractions were collected with a fraction collector. Fractions 10-23, which contained purified oligosaccharide 4, were collected and lyophilized. A higher yield of oligosaccharide 4 was obtained upon purification by Method B than by Method A.

[0137] The final product obtained either from method B was analyzed by Dionex chromatography (FIG. 2C), and the identity of the product was confirmed by mass spectrometry (FIG. 3B). Some impurities were present in the spectra of FIG. 2C and FIG. 3B.

Example 5

Coupling of Oligosaccharide 4 to GAA

[0138] Oxidation of GAA.

[0139] Lyophilized recombinant human GAA (rhGAA) was reconstituted in H₂O and dialyzed against 4 L of 100 mM acetate buffer (pH 5.6) 4 times to completely remove mannitol. After dialysis, the rhGAA was oxidized with 7.5 mM sodium periodate from 100 mM stock in 100 mM acetate buffer. After 30 minutes at 4° C. on ice, glycerol was added, and the sample was mixed on ice for 10 minutes to decompose excess sodium periodate. The oxidized material was then dialyzed against aqueous buffer (e.g., 100 mM sodium acetate) overnight.

[0140] Coupling.

[0141] A solution of oligosaccharide 4 in aqueous buffer (e.g., 100 mM sodium acetate, pH 5.6) was mixed with oxidized GAA and incubated at 37° C. for 4 hours to yield oligosaccharide-GAA conjugate 5. The reaction mixture was then diafiltered against 25 mM sodium phosphate buffer, pH 6.25, to remove unconjugated bisM6P glycan, and then adjusted with GAA formulation buffer (25 mM sodium phosphate buffer, pH 6.25, 2% mannitol, 0.005% Tween-80).

Example 6

Characterization of the GAA Conjugate

[0142] Detection of M6P.

[0143] The extent of oligosaccharide conjugation was measured by assaying conjugate 5 for binding to a M6P receptor column to which glycoproteins lacking M6P do not bind. Five micrograms of conjugate 5 were loaded onto a pre-equilibrated CI-MPR-Sepharose column (the column was prepared by coupling CI-MPR isolated from fetal

bovine serum to Affigel-10), which was then washed with CI-MPR binding buffer for 11×2 mL fractions and eluted with CI-MPR binding buffer containing 5 mM M6P for 7×2 mL fractions. A total of 18 fractions were collected and assayed for enzymatic activity.

[0144] Monosaccharide Analysis.

[0145] Conjugate 5 is treated with 4N trifluoroacetic acid to hydrolyze the oligosaccharides, followed by high pH anion exchange chromatography with pulsed amperometric detection (PAD) on a BioLC liquid chromatography system (Dionex). The monosaccharide content is extrapolated from a monosaccharide standard curve using premixed monosaccharide standards (Dionex).

[0146] Specific Activity.

[0147] GAA activity is measured using a fluorometric assay in black 96-well microplates using 4-methylumbelliferyl- α -D-glucoside as a substrate. Dilutions of conjugate 5 are added in triplicate to a microtiter plate. 4-methylumbelliferyl- α -D-glucoside is added to each sample. The 96-well plate is incubated in a 37° C. incubator for 30 minutes. The release of product is detected fluorometrically, and compared to standard curves generated by measuring the fluorescence of a known quantity of a standard. The reaction is quenched by the addition of 125 μ L of 1.0 M glycine-carbonate buffer, pH 10.5 to all wells. The specific activity is defined as nmol product released/hr/mg.

[0148] Internalization by L6 Myoblasts.

[0149] Cells (ATCC CRL-1458) were seeded into 6-well plates at 5.0×10^5 cells/well in growth media (DMEM+10% FBS) and grown to confluency. Cells were incubated with 0-100 nM GAA (conjugate 5 or unconjugated rhGAA) for 16 hours in DMEM+1% heat-inactivated-FBS+10 mM HEPES pH 6.7. After uptake, cells were washed with 3×PBS containing 5 mM M6P and lysed with 0.25% Triton X-100 for 1 hour on ice. Lysates were centrifuged at 18000 g for 5 minutes and tested for specific activity. See, e.g., Zhu et al., *J. Biol. Chem.* 279:50336-50341 (2004); Zhu et al., *Biochem. J.* 389:619-628 (2005).

[0150] All references cited herein are incorporated herein by reference in their entirety. To the extent publications and patents or patent applications incorporated by reference contradict the invention contained in the specification, the specification is intended to supercede and/or take precedence over any such contradictory material.

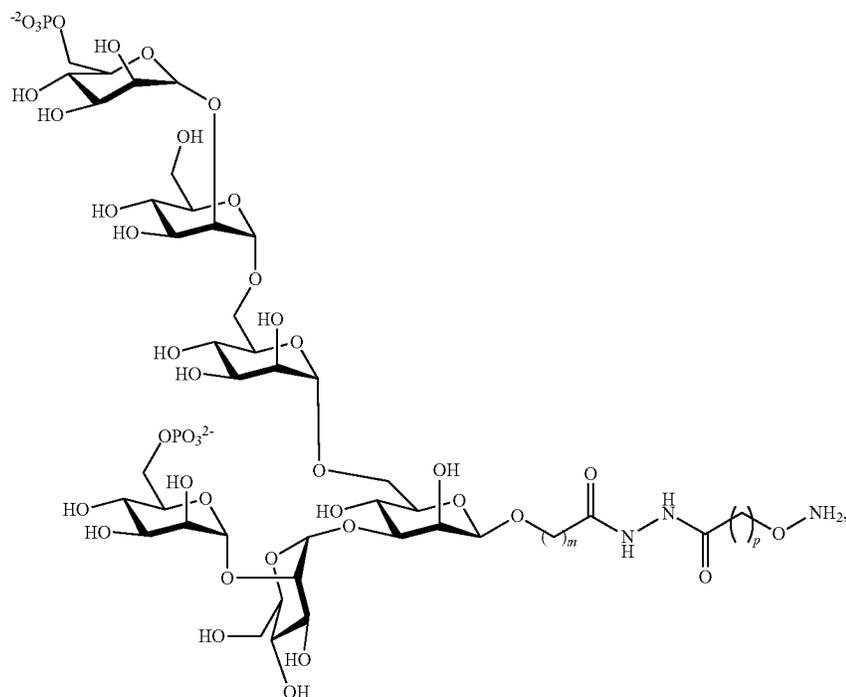
[0151] All numbers expressing quantities of ingredients, reaction conditions, and so forth used in the specification and claims are to be understood as being modified in all instances by the term "about." Accordingly, unless indicated to the contrary, the numerical parameters set forth in the specification and attached claims are approximations that may vary depending upon the desired properties sought to be obtained by the present invention. At the very least, and not as an attempt to limit the application of the doctrine of equivalents to the scope of the claims, each numerical parameter should be construed in light of the number of significant digits and ordinary rounding approaches.

[0152] Many modifications and variations of this invention can be made without departing from its spirit and scope, as will be apparent to those skilled in the art. The specific embodiments described herein are offered by way of example only and are not meant to be limiting in any way. It is intended that the specification and examples be considered as exemplary only, with a true scope and spirit of the invention being indicated by the following claims.

1-55. (canceled)

56. A method of coupling an oligosaccharide to a protein, comprising:

- (a) providing an oligosaccharide comprising an aminoxy group;
- (b) providing a protein having at least one carbonyl group; and
- (c) reacting the aminoxy group of the oligosaccharide with the at least one carbonyl group of the protein, thereby coupling the oligosaccharide to the protein, wherein the oligosaccharide comprising an aminoxy group is



wherein m is an integer from 1 to 10, and p is an integer from 1 to 10.

57. The method of claim **56**, wherein the protein is a glycoprotein and the at least one carbonyl group is obtained by oxidation of the glycoprotein with periodate.

58. The method of claim **56**, wherein the protein is a lysosomal enzyme.

59. The method of claim **58**, wherein the lysosomal enzyme is acid α -glucosidase, α -galactosidase A, acid sphingomyelinase, or α -L-iduronidase.

60. The method of claim **58**, wherein the lysosomal enzyme is acid α -glucosidase.

61. The method of claim **56**, wherein m is 3.

62. The method of claim **56**, wherein p is 1.

63. The method of claim **56**, wherein m is 3 and p is 1.

64. The method of claim **63**, wherein the protein is acid α -glucosidase.

65. An oligosaccharide-protein conjugate produced by the method of claim **56**.

66. A pharmaceutical composition comprising the oligosaccharide-protein conjugate of claim **65** and an excipient.

* * * * *