Nanovesicles displaying functional linear and branched oligomannose self-assembled from sequence-defined Janus glycodendrimers

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Cell surfaces are often decorated with glycoconjugates that contain linear and more complex symmetrically and asymmetrically branched carbohydrates essential for cellular recognition and communication processes. Mannose is one of the fundamental building blocks of glycans in many biological membranes. Moreover, oligomannoses are commonly found on the surface of pathogens such as bacteria and viruses as both glycolipids and glycoproteins. However, their mechanism of action is not well understood, even though this is of great potential interest for translational medicine. Sequence-defined amphiphilic Janus glycodendrimers containing simple mono- and disaccharides that mimic glycolipids are known to self-assemble into glycodendrimerosomes, which in turn resemble the surface of a cell by encoding carbohydrate activity via supramolecular multivalency. The synthetic challenge of preparing Janus glycodendrimers containing more complex linear and branched glycans has so far prevented access to more realistic cell mimics. However, the present work reports the use of an isothiocyanate-amine “click”-like reaction between isothiocyanate-containing sequence-defined amphiphilic Janus dendrimers and either linear or branched oligosaccharides containing up to six monosaccharide units attached to a hydrophobic amino-pentyl linker, a construct not expected to assemble into glycodendrimerosomes. Unexpectedly, these oligoMan-containing dendrimers, which have their hydrophobic linker connected via a thiourea group to the amphiphilic part of Janus glycodendrimers, self-organize into nanoscale glycodendrimerosomes. Specifically, the mannose-binding lectins that best agglutinate glycodendrimerosomes are those displaying hexamannose. Lamellar “raft-like” nanomorphologies on the surface of glycodendrimerosomes, self-organized from these sequence-defined glycans, endow these membrane mimics with high biological activity.

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Significance

Synthetic macromolecules that mimic glycolipids, named Janus glycodendrimers (JGDs), have been shown to self-assemble into nanovesicles displaying glycans on their outer surface, similar to the glycolalyx coating of eukaryotic cells, bacteria, and viruses. Specifically, both linear and branched oligosaccharides synthesized by automated glycan assembly, with hydrophobic linkers, have been used to create JGDs via an isothiocyanate–amine coupling reaction. Surprisingly, in spite of the surface of the hydrophobic linker, these JGDs self-organize into nanovesicles exhibiting lamellar surface morphologies, which mimic the recognition structures of cell-surface glycans and viral glycoproteins. Therefore, they are likely to be useful in helping elucidate mechanisms of significance for translational medicine such as the camouflage functionality employed by viruses to evade recognition.
the surface of GDSs were shown to bind more effectively at low sugar concentrations (18).

To date, most JGDs (14–20) have been synthesized via copper-catalyzed azide–alkyne cycloadditions (CuAAC) usually known as the conventional click reaction (21, 22) involving simple mono- or disaccharides bearing azido or alkyne moieties (14–20). A detailed investigation of the design principles of JGDs demonstrated through seven libraries containing 51 JGDs that hydrophilic linkers containing three to four oligo(oxyethylene)s must be connected directly to the sugar while the triazole group formed by click should be attached directly to the hydrophilic part of the dendron (14). These principles were used to design and predict the primary structure of self-assembling JGDs (15–20). Longer oligosaccharides have not been conjugated to JGDs due to the difficulties associated with their synthesis and further conjugation by conventional CuAAC click reactions.

Automated glycan assembly (AGA) (23–25) has expedited access to well-defined complex glycans up to 100-mer (24) that can serve as molecular tools for glycoscience (26). Synthetic oligosaccharides obtained by AGA are stereochemically pure and are equipped with an N-pentyl amino hydrophobic linker that facilitates the conjugation to surfaces or biomolecules (23–25). This hydrophobic N-pentyl amino linker attached to oligosaccharides prepared by AGA in milligram quantities does not fulfill the most fundamental structural requirement for the construction of JGDs by the classic click chemistry already elaborated (14). Fortunately, control experiments have demonstrated that a thiourea group incorporated between the hydrophilic part of the dendron (14). These principles go(oxyethylene)s must be connected directly to the sugar while the triazole group formed by click should be attached directly to the hydrophilic part of the dendron (14). These principles were used to design and predict the primary structure of self-assembling JGDs (15–20). Longer oligosaccharides have not been conjugated to JGDs due to the difficulties associated with their synthesis and further conjugation by conventional CuAAC click reactions.

Results and Discussion
A representative example for the synthesis of a disaccharide Lac containing a triethylene glycol linker with an azido group is presented in Fig. 1A (19). Glycosylation of 2-(2-[2-chloroethoxy]ethoxy)ethanol (1) catalyzed by the Lawesson reagent (27) started when lactose octaacetate (1) was functionalized with benzyl (5-hydroxypentyl)carbamate (5–21). The acetyl groups were removed via Zemplén deacetylation with sodium methoxide in methanol to give compound 5. Hydrogenolysis in the presence of Pd/C provided Lac bearing a triethylene glycol chain with an amino group (6). The synthesis of Lac functionalized with the hydrophobic amino-pentyl linker (10) (27) started when lactose octaacetate (1) was first reacted with 1,2-diazobicyclo[5.4.0]undec-7-ene (DBU) to give 7. Intermediate 7 was functionalized with benzyl (5-hydroxypentyl)carbamate (8) to give 9 (Fig. 1B). After Zemplén deacetylation and hydrogenolysis of the carboxylbenzyl (Cbz) group of 9, Lac containing the amino-pentyl linker (10) was obtained.

AGA of hexaMan (Fig. 1C) is representative for the synthesis of all oligoMan (23–25) (SI Appendix, Figs. S1–S32). AGA was performed with thiomannoside (11) on a polymeric resin functionalized with a photo-cleavable linker (12). Six cycles of glycosylation and deprotection were repeated to obtain the desired hexa-saccharide. Upon completion of the assembly, the compound was cleaved form the solid support and subjected to Zemplén methanolyis and hydrogenolysis to give compound hexaMan (13).

Fig. 1. Synthetic route for preparation of sugars containing triethylene glycol azido and amino linkers (A). Synthetic route for sugar containing the hydrophobic N-pentyl amino linkers (B). Automated glycan assembly of oligoMan containing N-pentyl amino linkers (C). Reagents and conditions: (i) BF₃·Et₂O, CH₂Cl₂, 0 °C to 23 °C, 15 h, 80%; (ii) Na₂CO₃, DMF, 80 °C, 15 h, 74%; (iii) MeONa, methanol, pH = 10, 23 °C, 7 h, 95%; (iv) H₂, Pd/C, methanol, 23 °C, 15 h, 95%; (v) H₂NNH₂·HOAc, DMF, then CCl₄/N₂, DBU, CH₂Cl₂, 79%; (vi) TMSOTf, CH₂Cl₂, 39%; (vii) MeONa, methanol, then H₂, Pd/C, 82%.
The AGA of oligoMan is usually performed on a 0.0125-mmol scale and therefore requires a highly efficient methodology to conjugate this small quantity of oligosaccharides to the JGDs. In order to selectively conjugate the primary amino group to the JGDs in the presence of multiple hydroxyl groups and to potentially diminish the hydrophobic character of oligoMan linker, the incorporation of an alkyl ITC on the hydrophilic part of JD was designed. ITC reacts with amines fast and irreversibly, while the reaction between ITC and alcohols and water, which can be used also as a solvent, is negligible and reversible (28), being below the level that can be detected even by analytical methods. Due to this selectivity, ITC functionalized compounds are widely used for orthogonal bioconjugation (28, 29) and Sharpless and coworkers (21) considered this reaction as part of the large protocol of click methods. In addition, it will be discussed later, the Lac-containing hydrophilic and hydrophobic linkers were used to produce the model compounds 5, 6, and 10. They all formed JGDs that self-assemble in GDSs with similar stability in buffer and efficiency to agglutination. Therefore, the ITC-amine “click”-like reaction was preferred since it did not require any change to the current AGA technology.

The synthesis of ITC functionalized JD is presented in Fig. 2. The 2-(2-[2-chloroethoxy]ethoxy)ethanol (2) was reacted with sodium azide to obtain 2-[2-(2-azidoethoxy)ethoxy] ethanol (14), followed by chlorination with thionyl chloride to generate 1-azido-2-(2-[2-chloroethoxy]ethoxy)ethane (15) (30). Alkylation of methyl 4-(benzyloxy)-3,5-dihydroxybenzoate (16), followed by deprotection and alkylation, provided the azido-functionalized hydrophilic minidendron (21). The hydrophilic minidendron (22)
was connected to the acetonide-protected Tris(hydroxymethyl)-
aminomethane (Tris) core (23) with 2-chloro-4,6-dimethoxy-
1,3,5-triazine (CDMT) and N-methylmorpholine (NMM), followed
by conjugation with the azido functionalized hydrophilic mini-
dendron (21) via N,N'-dicyclohexylcarbodiimide (DCC) and 4-
(dimethylamino)pyridinium 4-toluenesulphonate (DPTS) (31).

**Library of oligosaccharides prepared by organic synthesis
and by automated glycan assembly (AGA)**

**Fig. 4.** Oligosaccharides with pentyl-amino linker used for the preparation of JGDs via ITC-amine “click”-like reaction. Man and Lac were prepared by solution phase synthesis. Man, Gal, Glc, GlcNAc, and oligoMan were prepared by AGA.

**Fig. 3.** Sequence-defined JGDs obtained by conventional click reaction via CuAAC (20) (A) and by ITC-amine “click”-like reaction (B), reported here.
Upon deprotection of the acetonide and conjugation to the hydrophobic minidendron (27) with DCC and DPTS, the azido containing amphiphilic JD [JD(5/1)$_{N3}$, 28] was obtained. The nomenclature of JD(5/1)$_{N3}$ was adapted from that of sequence-defined JGDs elaborated by our laboratory (18). For example, 5/1 corresponds to five methyl triethylene glycol chains and one azido (N$_3$) functionalized triethylene glycol chain on the hydrophilic dendrons of sequence-defined JD. The azido group of JD(5/1)$_{N3}$ was reduced to amine by hydrogenation to obtain JD(5/1)$_{ITC}$ (29). Finally, an efficient method using carbon disulfide (32), triethyl amine, and tosyl chloride was employed to transform the amino group to ITC under mild conditions and produce the ITC functionalized sequence-defined JD JD(5/1)$_{ITC}$ (30). Tosyl chloride was used to mediate decomposition of the corresponding dithiocarbamate salt which was generated in situ by treatment of the amine JD(5/1)$_{ITC}$ with carbon disulfide and triethylamine (32). The formation of the azido and ITC-containing JGs was confirmed by NMR and matrix-assisted laser desorption ionization time-of-flight (MALDI-TOF) mass analysis (SI Appendix, Fig. S33).

Disaccharides 6 and 10 were selected as models to test the ITC-amine “click” conjugation with ITC functionalized JD.
**JD(5/1_{ITC})** (30), in the presence of triethyl amine and DMF. The conventional click reaction via CuAAC (18) was previously utilized to synthesize sequence-defined JGDs (Fig. 3A) (14–20). Copper sulfate was reduced in situ with sodium ascorbate to catalyze the azide–alkyne cycloaddition and form a 1,2,3-triazole ring that connected the JD with the sugar. Similarly, the ITC–amine “click”-like reaction occurred between the ITC group of the JD(5/1_{ITC}) (30) and the amine of the Lac analogs 6, containing a triethylene glycol linker (3EO), and 10, containing the hydrophobic N-pentyl linker (CS) (Fig. 3B) where thiourea connected the hydrophilic JD and the hydrophobic part of the sugar. Two JGDs containing sequence-defined Lac 3EO hydrophilic chain JD(5/1_{Lac}^{3EO}) and sequence-defined Lac 3EO hydrophilic chain JD(5/1_{Lac}^{3EO}) were successfully synthesized in 57% and 46% isolated yields from Lac analogs 6 or 10, respectively. This reaction is competitive with the conventional click chemistry (compare isolated yields in Fig. 3A and B) except that the as obtained from AGA hydrophobic N-pentyl linked amine is used directly in the ITC-amine “click”-like reaction and therefore it does not require any additional synthetic steps.

The ITC–amine “click”-like methodology was employed to build a library of JGDs containing oligosaccharides (Fig. 4). Amino-pentyl functionalized Man and Lac were prepared by conventional solution phase synthesis (Fig. 1B). Man, Gal, Glc, N-acetylglucosamine (GlcNAc), and oligoMan, including (α1→3)-type diMan, triMan, pentaMan, hexaMan; (α1→2)-type diMan and triMan; and (α1→6) (α1→6)-branched-triMan were prepared by AGA (SI Appendix, Figs. S1–S32). The sugars were mixed with JDG(5/1_{ITC}) in DMF in the presence of triethyl amine. DMF was selected as the solvent due to good solubility for both ITC-containing JD JDG(5/1_{ITC}) and amino-containing sugars. The target JGDs were purified on silica gel chromatography with CH$_2$Cl$_2$/methanol (for JGDs containing mono- to trisaccharides) or CH$_2$Cl$_2$/methanol/water (for JGDs containing more than trisaccharides) as eluents to give the desired products in 39 to 61% isolated yields without any optimization (Fig. 4).

This reaction can be efficiently performed even with only few milligrams of sugar as obtained by AGA and confirmed by MALDI-TOF analysis (SI Appendix, Figs. S34 and S35). Due to the solubility of the less-polar JD part, the amphiphilic JGDs can be dissolved in less-polar NMR solvents such as CDCl$_3$, where the JGDs did not self-assemble. The chemical structures of sequence-defined amphiphilic JGDs containing monosaccharide (Man, Gal, Glc, and GlcNAc), disaccharide as Lac, and linear or branched oligoMan are summarized and compared in Fig. 5.

GDSs were assembled by injecting tetrahydrofuran solutions of JGDs in water or buffer, as reported previously (14–20). Cryogenic transmission electron microscopy analysis (SI Appendix, Fig. S36), of these GDSs self-organized from JGDs containing Lac and oligoMan via ITC-amine “click” reaction demonstrated the formation of unilamellar vesicles with a diameter of about 100 nm. The self-assembled GDSs are similar to those previously reported GDSs when they were prepared via the CuAAC click method (14–20). This is a remarkable result considering that JDSs containing hydrophobic linkers attached to the sugar do not form GDSs (14). This demonstrates the extraordinary contribution of the thiourea interconnecting group between the hydrophobic and hydrophilic fragments of JDSs to their self-assembly. Therefore, these experiments demonstrated the efficiency of the ITC–amine “click”-like reaction for the synthesis of amphiphilic JGDs containing complex oligosaccharides.

In order to investigate the activity of sugars on the surface of JGDs, we self-assembled GDSs from JDG(5/1_{Gal}^{Cs}), JDG(5/1_{Lac}^{Cs}), and JDG(5/1_{Man}^{3EO}) in 1x phosphate-buffered saline (Fig. 6). The agglutination assays were performed with these GDSs with two adhesion/growth-regulatory human galectins: galectin-1 (Gal-1) (18) and galectin-8S (Gal-8S) (17, 18). Human galectins bind to the disaccharide Lac much stronger than monosaccharide Gal. Gal-containing GDSs did not show increase of turbidity, most probably due to the weak interaction between the two components. In contrast, the two GDSs with Lac showed similar agglutination behavior with Gal-1 and Gal-8S. These data indicated the similar activity of the sugars on the surface of GDSs regardless of the presence of the hydrophilic triethylene glycol or hydrophobic...
N-pentyl linker connected to the hydrophilic fragment via thiourea (see ovoidal dashed structures in Figs. 3A and B and 6). Thus, complex carbohydrates containing the hydrophobic pentyl amine linker attached to hydrophobic fragments via thiourea can be used to design self-assembling JGDs (14). This demonstrates that the thiourea group incorporated during the ITC–amine “click”-like reaction enhances the hydrophilic character of the N-pentyl linker and facilitates the self-assembly process of the JGDs.

Fig. 7. Agglutination assays of GDSs from amphiphilic JGDs containing oligoMan with ConA. (A–C) Agglutination assays of GDSs from JGDs containing Man (A), diMan (B), and triMan (C). The GDSs were prepared by self-assembly of the JGDs (100%) or coassembled with JD-3 (20 and 50 mol % of JGDs). (D and E) Agglutination assays of GDSs from JGDs containing (α1–2) oligoMan (D), (α1–6) oligoMan (E), and comparison of three JGDs containing isomeric linear or branched trimannose (F). The GDSs were prepared by coassembled with JD-3 (20 mol % of JGDs).

Fig. 8. Surface topography of GDS self-organized from Lac-containing sequence-defined JGDs including JGD(5/1Lac)(A–C) with a hydrophobic C5 linker and JGD(5/1Lac)(D–F) with a hydrophilic 3EO linker deposited on mica. AFM height images (A and D), phase images (B and E) with inserted FFT, and the corresponding height profiles (C and F) of GDSs on mica measured at ambient humidity (40%). Arrows in A and D indicate the directions for analysis of the height profiles in C and F.
GDSs self-organized from JGDs containing oligoMan were assembled by injection into 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (Hepes) buffer containing CaCl$_2$ and MnCl$_2$. Agglutination experiments of the GDSs from Man (Fig. 7A), (α1–6) diMan (Fig. 7B), and (α1–6) triMan (Fig. 7C) with concanavalin A (ConA) (14, 16) were performed. Tetrameric leguminous lectin ConA binds to Man- and oligosaccharides containing terminal Man units, in the presence of Ca$^{2+}$ and Mn$^{2+}$ (7, 9, 10, 12). These GDSs aggregated with ConA, indicating the existence of (α1–6)-type oligoMan on the surface of GDSs. However, the binding curves (blue curves, Fig. 7A–C) did not correlate the different numbers of the (α1–6) oligoMan units. Therefore, JGDs were coassembled in the presence of a nonsugar JD (JD-3, Fig. 5, top left row). The ratio of JDGs was diluted from 100 to 50% (red curves, Fig. 7A–C) and to 20% (green curves, Fig. 7A–C). For Man (Fig. 7A), the green curve (20%) is much lower than the blue curve (100%). In contrast, for triMan (Fig. 7C), the green curve (20%) maintained at a relative high level.

The coassembled GDSs with 20% JDGs showed an increased trend with an increasing number of (α1–6) oligoMan units (green curves in Fig. 7A–C, and in Fig. 7E). Thus, the coassembled GDSs containing 20% of JDGs can be employed to investigate the relative activity of JDGs in the biological membrane mimic with the function of oligoMan units. Based on these results, we prepared the coassembled GDSs including 20% JDGs and 80% of JD-3 (see structure of JD-3 in Fig. 5, top left row) for the entire oligoMan-containing JDGs library (Fig. 7F–D). For (α1–2) oligoMan, the slopes of binding curves increased with each lengthening of the sugar from Man to (α1–2) diMan and then to (α1–2) triMan (Fig. 7D). Fig. 7E demonstrated the hexaMan containing JGD is the most active among the (α1–6) oligoMan. Three triMan-containing JDGs with linear (α1–2) triMan, linear (α1–6) triMan, or (α1–2) (α1–6) branch-triMan, showed the highest slopes as well as plateau (Fig. 7F). It should be noted that all oligoMan-presenting GDSs demonstrated higher activity toward ConA whether linear or branched than the monosaccharide Man-presenting GDSs. Therefore, the synthesis of JDGs from unprotected sugars performed under the ITC-amine “click”–like reaction conditions provides an efficient methodology to combine AGA-generated small quantities of complex-architecture oligosaccharides with GDSs to yield synthetic cell-like objects decorated with sugar moieties.

Analysis using a combination of atomic force microscopy (AFM) and fast Fourier transform (FFT) with a diffraction-like approach to JGD and GDSs. Oligosaccharides prepared by AGA (23–25), including a linear oligoMan with six mannose units and a branched trimannose, were conjugated to JDs via ITC-amine “click”–like reaction balances the hydrophobicity of the linker. This is a surprising result considering that the hydrophobic linkers attached to sugars do not form JDGs that self-assemble into GDSs (14). Therefore, additional experiments are needed to generalize this concept, which may endow a new architectural approach to JGD and GDSs. Oligosaccharides prepared by AGA (23–25), including a linear oligoMan with six mannose units and a branched trimannose, were conjugated to JDs via ITC-amine “click”–like without any additional synthetic modification of the carbohydrate. The resulting JDGs self-assembled into unilamellar GDSs. OligoMan-containing GDSs bind the mannose-binding lectin ConA, with the hexaMan binding the most tightly. Some GDSs adopt raft-like nanomorphologies on their cell membrane mimic surface to support the enhanced activity previously observed for other sequence-defined GDSs (17, 18, 19) and hence shed light on the relationship between glycan structure and activity. The combination of AGA and the self-organization of sequence-defined JDGs has created models to start illuminating why biological cell membranes carry long and branched glycans, which are bound more tightly by sugar-binding proteins. These results provide a platform to develop cell-membrane mimics and the tools required to conjugate amino containing biomacromolecules to JDs, including not only more complex carbohydrates synthesized by AGA but also peptides and oligonucleotides.

The combined synthetic ITC-amine conjugation (28, 29) and AGA (23–25) with supramolecular multivalency (35, 36) methodology reported here is likely to help elucidate the function of glycan camouflage employing by viruses (40, 41).

**Methods**

**Glycan Synthesis.** AGA was performed on a home-built synthesizer developed at the Max Planck Institute of Colloids and Interfaces, Golm, Germany (25). All details concerning automation steps, building blocks, AGA modules, and post-synthesis manipulations can be found in SI Appendix.

**Data Availability.** All data associated with this work are available either in the main text or in SI Appendix.

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