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A Ferrier glycosylation/*cis*-dihydroxylation strategy to synthesize *Leishmania* spp. lipophosphoglycanassociated βGal(1,4)Man disaccharide†

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The Gal $\beta(1 \rightarrow 4)$ Man disaccharide, found in the cell surface lipophosphoglycan (LPG) of *Leishmania* species, has been synthesized by a Ferrier glycosylation/*cis*-dihydroxylation strategy. This stereoselective method proved efficient for synthesizing the target saccharide in good yield. In addition, we prepared two clickable *O*-glycoside and phospho-glycoside versions of Gal $\beta(1 \rightarrow 4)$ Man to enable conjugation to protein carriers for further immunological and antibody-binding studies.

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Introduction

Leishmaniasis is a poverty-associated parasitic disease that imposes a heavy burden on the public health systems of least developed countries, particularly those of South Asia, East Africa, and Latin America. The protozoan Leishmania spp. (Trypanosomatidae), the etiological agent of leishmaniasis, is a vector-borne obligate intracellular parasite transmitted by hematophagous sandflies. More than 20 human-infective Leishmania species have been described to date, together with 70 different types of sandflies (Phlebotomus and Lutzomyia) identified as vectors.^{1,2} Depending on the infecting Leishmania species, individuals can present one of the three disease phenotypes: self-resolvent cutaneous leishmaniasis (CL), disfiguring mucocutaneous leishmaniasis (MCL), or potentially fatal visceral leishmaniasis (VL).² Although leishmaniasis is an underreported infection, current estimates for CL range from 700 000 to 1.2 million worldwide cases each year, and yearly VL cases are about 100 000.3 Available anti-leishmaniasis treatments include pentavalent antimonials, liposomal amphotericin B, miltefosine, and paromomycin.4 These drugs are effective in treating leishmaniasis; however, side effects, poor patient compliance, and parasite resistance motivate the search for new and improved drugs. Moreover, the lack of an effective anti-leishmaniasis vaccine to prevent and treat the infection is a critical need.

A dense glycocalyx covers the *Leishmania* spp. cell membrane playing a pivotal role in the parasite's survival and infectivity. The chemical structure of this sugar coat varies according to the

parasite species, its life cycle stage, and the host type.5 Fig. 1A shows the structure of Leishmania's lipophosphoglycan (LPG), the predominant membrane-glycan in promastigotes, the human-infective form.6 Its structure comprises a phosphatidylinositol anchor, a conserved oligosaccharide core, a linear phosphoglycan (PG), and a neutral mannose-rich oligosaccharide cap. The chemical structure of LPG varies among Leishmania species. The differences are typically found in the structures of the capping oligosaccharides and the type and sequence of the branching sugars connected to the conserved linear PG, a phospho-polysaccharide comprising the [-6)- β -D- $Gal(1 \rightarrow 4)\alpha$ -D-Man- $(1 \rightarrow PO_3O]$ repeating unit. For example, in Leishmania donovani, the etiological agent of VL in the Old World, promastigotes PG is unsubstituted and remains linear.7 In contrast, in L. infantum (L. chagasi), the causative agent of New World's VL, PG carries $Glc\beta(1 \rightarrow 3)$ substitutions.⁸

Given its role as an immune determinant, LPG is an attractive target for developing glycoconjugate vaccine candidates against leishmaniasis.⁹ The groups of Nikolaev, Vishwakarma, and Seeberger have accomplished the synthesis of different LPG fragments and proved the suitability of protein-conjugated LPG subunits as immunogens.¹⁰ However, despite the promising preliminary immunization results,¹¹ an effective antileishmaniasis vaccine remains elusive.

Because of our interest in anti-parasitic immunotherapies¹² and the development of virus-like particles (VLP) glycoconjugates to boost anti-carbohydrate immune responses,¹³ we have centered our attention on LPG to explore VLP-conjugated anti-leishmaniasis vaccine candidates. Here, we report an expeditious synthetic route to prepare Gal $\beta(1 \rightarrow 4)$ Man, a disaccharide exclusively found in the linear phosphoglycan and the capping oligosaccharides of *Leishmania's* LPG. Two clickable versions of Gal $\beta(1 \rightarrow 4)$ Man (Fig. 1B), namely 3-azidopropyl 4-*O*- β -D-galactopyranosyl- α -D-mannopyranoside (1) and 3-azidopropyl 4-*O*- β -D-galactopyranosyl- α -D-mannopyranosyl- α

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Fig. 1 (A) Structure of *Leishmania donovani* lipophosphoglycan; (B) structures of the clickable $Gal\beta(1 \rightarrow 4)Man O$ - and phospho-glycosides 1 and 2 synthesized in this work.

phosphate (2), were prepared to enable conjugation to the protein carrier and subsequent performance of immunological studies.

Results and discussion

Synthesis of Gal $\beta(1 \rightarrow 4)$ Man disaccharide

Scheme 1 shows the synthetic strategy to prepare $Gal\beta(1 \rightarrow 4)$ Man. We envisioned the combination of the Ferrier glycosylation¹⁴ and osmium-catalyzed *cis*-dihydroxylation,¹⁵ both highyielding and stereoselective reactions, as a convenient sequence to synthesize the target disaccharide from the inexpensive and readily available starting material *D*-lactose *per*acetate. First. we synthesized the Ferrier glycosyl donor hexa-*O*acetyl-*D*-lactal 3 in 42% yield by adjusting the procedure of Zhao *et al.*¹⁶ which involves forming the lactosyl bromide from lactose *per*-acetate followed by reduction with Zn dust. The subsequent Ferrier glycosylation^{14,17} of benzyl alcohol with lactal 3, catalyzed by BF₃·Et₂O, afforded a mixture of 2,3-unsaturated benzyl glycosides 4α and 4β ($\alpha/\beta = 6:1$) in 92% yield. The osmiumcatalyzed *cis*-dihydroxylation of the anomeric mixture gave diols 5 (major product) and 6 in excellent yield, which were readily separated by normal-phase flash chromatography. The treatment of diols 5 and 6 with Ac₂O in pyridine yielded the respective *per*-acetates 7 and 8. Given the convenient peak separation in their ¹H NMR spectra, we performed the stereochemical configuration analysis of the oxidation adducts on their respective *per*-acetylated derivatives.

In the ¹H NMR spectrum of acetate 7, derived from the major *cis*-dihydroxylation product 5, proton H3 (δ 5.32 ppm) appears as a doublet of doublets with coupling constants ${}^{3}J_{\text{H3-H4}}$ and ${}^{3}J_{\text{H3-H2}}$ of 8.8 and 3.6 Hz respectively.

The large value for the ${}^{3}J_{\rm H3-H4}$ unambiguously identifies a *trans*-diaxial correlation between H3 and H4. The anomeric proton H1 (δ 4.74 ppm) appears as a doublet with a small ${}^{3}J_{\rm H1-H2}$ (1.8 Hz). These attributes are consistent with an α -mannose configuration.¹⁸ In acetate **8**, derived from the minor *cis*-dihydroxylation product **6**, H3 (δ 5.76 ppm) appears as a triplet with



Scheme 1 Synthesis of Gal $\beta(1 \rightarrow 4)\alpha$ Man disaccharide *via* Ferrier glycosylation/*cis*-dihydroxylation sequence: (a) synthesis of D-lactal: (i) HBr (34% in AcOH), CH₂Cl₂, 0 °C to room temperature. (ii) Zn, NaH₂PO₄ (sat.), acetone, (42% two steps); (b) BnOH, BF₃·Et₂O, dioxane, room temperature. (66%, $\alpha/\beta = 6 : 1$); (c) OsO₄, NMO, H₂O, acetone, room temperature (92%); (d) Ac₂O, pyridine, CH₂Cl₂, room temperature (88% for **7**, 81% for **8**).

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 ${}^{3}J_{\rm H3-H4}$ and ${}^{3}J_{\rm H3-H2}$ values of ~2.5 Hz, indicating a *cis* correlation with H4. The proton H1 (δ 4.83 ppm) showed as a doublet with a large ${}^{3}J_{\rm H1-H2}$ value (8.3 Hz), confirming a β -allose configuration for the minor oxidation product.

The results show that the osmium-catalyzed *cis*-dihydroxylation of disaccharide derivatives 2,3-dideoxy-4-*O*- β -D-galactopyranosyl-D-*erythro*-hex-2-enopyranosides exhibit the same stereospecificity reported for the oxidation of the monosaccharide 2,3-dideoxy-D-*erythro*-hex-2-enopyranoside congeners: oxidation of 2,3-unsaturated α -glycosides leads to the formation of α -mannosides whereas the oxidation of the respective β -anomers yields β -allosides.¹⁹

Since the formation of the β -glycoside in the Ferrier glycosylation limits the yield of the target Gal $\beta(1 \rightarrow 4)$ Man disaccharide 5, we attempted to improve the α -stereoselectivity of this reaction by testing different solvents (MeCN, dioxane, THF, and Et₂O), common Lewis acid catalysts (FeCl₃,²⁰ TMSOTf,²¹ and I₂ (ref. 22)), and Lewis acids reported to afford high α -glycoside yields in Ferrier glycosylations (Y(OTf)₃,²³ Gd(OTf)₃ (ref. 24)). However, in our hands, these attempts did not significantly improve the α -selectivity of this step using D-lactal 3 as a glycosyl donor.

Synthesis of azide-functionalized Gal $\beta(1 \rightarrow 4)\alpha$ Man *O*-glycoside 1. To facilitate immobilization onto surfaces and conjugation with protein carriers for further immunological studies, we installed a clickable²⁵ azide-functionalized aglycone on Gal $\beta(1 \rightarrow 4)\alpha$ Man (Scheme 2). The Pd-catalyzed hydrogenolysis of benzyl glycoside 7, afforded the respective anomeric alcohol 9 in quantitative yield. The subsequent treatment of this intermediate with Cl₃CCN and DBU in MeCN generated the respective imidate 10 which was coupled to 3-azidopropanol by TMSOTf-catalyzed imidate activation in Et₂O at 0 °C. The azide-functionalized glycoside 11 was obtained in 70% yield as a single anomer. The removal of the acetyl protective groups *via* Zemplén transesterification²⁶ yielded the target azidefunctionalized Gal $\beta(1 \rightarrow 4)$ Man α -glycoside 1.

Synthesis of azide-functionalized $Gal\beta(1 \rightarrow 4)\alpha Man$ phospho-glycoside 2. The H-phosphonate 12 was prepared by treating the anomeric alcohol 9 with a freshly prepared solution of the phosphorylating reagent triimidazolylphosphine (PIm₃),²⁷ made by reacting PCl₃ with imidazole in MeCN (Scheme 3). The addition of alcohol 9 to a PIm₃ solution forms the respective phosphorodiamidite intermediate which upon



Scheme 3 Synthesis of azide-functionalized Gal $\beta(1 \rightarrow 4)\alpha$ Man phospho-glycoside 2. (a) PCl₃, imidazole, Et₃N, MeCN (42%); (b) (i) 3-azidopropanol, PivCl, THF; (ii) I₂, H₂O/pyridine (5 : 95) (30% two steps); (c) NaOMe, MeOH, Amberlite IR 120(+) (67%).

hydrolysis with triethylammonium bicarbonate buffer (TEAB, pH 7.2) affords the triethylammonium Gal $\beta(1 \rightarrow 4)\alpha$ Man Hphosphonate **12** in 42% yield. The successful Hphosphonylation was confirmed by the ³¹P NMR spectrum of **12**, which showed a characteristic H-phosphonate signal at δ_P -0.10 ppm (dd, ${}^{1}J_{P-H} = 638$ Hz, ${}^{1}J_{P-H1} = 9.8$ Hz). After confirming H-phosphonate identity, we coupled **12** to 3-azidopropanol in THF using pivaloyl chloride as a coupling reagent. The H-phosphonate diester intermediate, that is formed in this reaction, was oxidized *in situ* with I₂ and H₂O/pyridine to afford the respective phosphate diester **13** in 30% yield. The ³¹P NMR spectrum of **13** showed phosphorus at $\delta_P - 3.09$ (d, ${}^{3}J_{P-H1} = 6.9$ Hz) confirming the H-phosphonate oxidation to phosphate. Removal of the acetyl protecting groups afforded the target azide-functionalized Gal $\beta(1 \rightarrow 4)\alpha$ Man phospho-glycoside **2**.

Conclusions

We have synthesized the $\beta Gal(1 \rightarrow 4) \alpha Man$ disaccharide, the phospho-polysaccharide repeating unit found in the cell surface



Scheme 2 Synthesis of clickable Gal $\beta(1 \rightarrow 4)$ Man O-glycoside. (a) H₂, Pd/C, EtOH (96%); (b) Cl₃CCN, DBU, CH₂Cl₂ (68%); (c) 3-azidopropanol, TMSOTf, CH₂Cl₂ (70%); (d) NaOMe, MeOH, Amberlite IR120 (H⁺) (97%).

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lipophosphoglycan of all Leishmania species. The Ferrier glycosylation/*cis*-dihydroxylation sequence proved efficient in affording the target β Gal($1 \rightarrow 4$)Man core in a stereoselective fashion from readily available lactose *per*-acetate. Two clickable versions of β Gal($1 \rightarrow 4$)Man, azide-functionalized *O*-glycoside and phospho-glycoside, were also prepared to enable coupling to protein carriers and immobilization onto surfaces for further immunological and binding studies.

Experimental

General information

D-Lactose and all reagents were purchased to the highest purity available and used without previous purification. Yields refer to isolated yields after chromatographic purification and anomeric ratios were determined from the ¹H NMR spectra of reaction crudes. Reactions were monitored by thin-layer chromatography (TLC) using SiliCycle SiliaPlate glass-backed TLC 250 mm with F-254 UV indicator. TLC visualization was performed under short wave UV light (254 nm) and charring with 5% H₂SO₄ in CH₃OH. Normal phase flash chromatography was performed using SiliCycle SiliaFlash P60 silica gel (40-63 µm) and mixtures of EtOAc-hexanes and CH2Cl2-MeOH (ACS-grade solvents) as eluents. 1D and 2D NMR spectra were recorded on a Bruker Avance 400 spectrometer at 298 K. ¹H NMR chemical shifts are reported in parts per million vs. tetramethylsilane (TMS) internal standard ($\delta_{\rm H}$ 0.00 ppm). ¹³C{¹H} NMR chemical shifts are reported in parts per million vs. the residual peak of the solvent.28 LCMS data were collected in a Waters e2695 HPLC system coupled to a Waters Acquity QDa single quadrupole mass detector. HRMS data for new compounds were collected in a Waters high-resolution Xevo G2-XS Q-ToF mass spectrometer coupled with an Acquity UPLC H-Class in the positive electrospray ionization mode.

3,6-Di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-1,5-anhydro-2-deoxy-D-arabino-hex-1-enitol

(3).

Following the procedure described by Zhao et al.16 In a roundbottomed flask, a solution of HBr (147 ml, 0.62 moles, 34% in HOAc) in dry CH₂Cl₂ (200 mL) was cooled to 0 °C with an ice bath. Then, a solution of lactose per-acetate (141.2 g, 0.21 moles) in dry CH₂Cl₂ (100 mL) was added dropwise through addition funnel. After completing the addition, the system was allowed to warm to room temperature and stirring was kept until TLC showed total consumption of the sugar (\sim 3 hours). The mixture was then diluted with water, transferred to a separation funnel, and extracted with CH₂Cl₂. The organic layer was then successively washed with H₂O (3 times), and NaHCO₃ (sat. 2 times). The organic phase was collected, dried over Na₂SO₄, filtered, and the volatiles removed in rotary evaporator. The resulting material (hepta-O-acetyl lactose bromide) was then re-dissolved in acetone (420 mL), and a saturated aqueous solution of NaH₂PO₄ (830 mL) was added. To the resulting biphasic system, Zn (170 g, 2.6 moles) was added portion-wise, and the suspension was maintained at room temperature under strong stirring for 12 hours. After that period, TLC showed total consumption of the starting bromide and the predominance of a slightly less polar spot. The solids were then removed by filtration through

a Celite pad, and the organic filtrate was concentrated in rotary evaporator. The residue was re-dissolved in EtOAc, washed with water (2 times), dried over Na₂SO₄, filtered, and the volatiles removed in rotary evaporator. Flash chromatography purification of the residue afforded hexa-O-acetyl D-lactal 2 (49.4 g, 88.2 mmol, 42%) as a colourless syrup. The identity of peracetylated lactal 3 was confirmed by comparing its ¹H and ¹³C 1 H NMR spectra with reported data.²⁹ $R_{\rm f} = 0.56$ (EtOAc/*n*-Hex 3:2), ¹H NMR (500 MHz, CDCl₃): 6.35 (dd, J = 1.0 and 6.1 Hz, H-1), 5.34 (t, J = 4.3 Hz, H-3), 5.30 (dd, J = 1.0 and 3.3 Hz, H-4'), 5.13 (dd, *J* = 7.9 and 10.5 Hz, H-2′), 4.94 (dd, *J* = 3.4 and 10.5 Hz, H-3′), 4.77 (dd, *J* = 3.4 and 6.1 Hz, H-2), 4.59 (d, *J* = 8.0 Hz, H-1′), 4.37 (dd, J = 2.6 and 11.7 Hz, H-6), 4.13 (dd, J = 6.2 and 11.7 Hz, H-6), 4.08 (m, H-6', H-5), 4.01 (dd, *J* = 7.3 and 11.2 Hz, H-6'), 3.93 (dd, J = 5.4 and 7.4 Hz, H-4), 3.84 (ddd, J = 1.0, 6.0 and 7.0 Hz,H-5'), 2.09 (s, 3H), 2.05 (s, 3H), 2.02 (s, 3H), 1.99 (s, 3H), 1.98 (s, 3H), 1.91 (s, 3H); APT ¹³C{¹H} NMR (125 MHz, CDCl₃): 170.4, 170.4, 170.2, 170.1, 169.9, 169.3, 145.4 (C-1), 101.0 (C-1'), 98.9 (C-2), 74.6 (C-4), 74.1 (C-5), 70.8 (C-3'), 70.7 (C-5'), 68.8 (C-3), 68.8 (C-4), 66.6 (C-4'), 61.8 (C-6), 60.9 (C-6'), 21.1, 20.8, 20.6, 20.6, 20.6, 20.5.

Benzyl 2,3-dideoxy-4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-6-O-acetyl- α -D-erythro-hex-2-enopyranoside (4 α and 4 β). p-Lactal (2.0 g, 3.5 mmol) and benzylic alcohol (563 µL, 1.5 equiv.) were dissolved in dry CH_2Cl_2 (10.0 mL mmol⁻¹) and cooled to 0 °C under nitrogen atmosphere. Then boron trifluoride (230 µL, 0.5 equiv.) was added dropwise, and the reaction was allowed to warm to room temperature. Reaction was stirred until TLC analysis showed total consumption of starting lactal (2 hours). Then the reaction is cooled down to 0 °C and quenched with Et₃N (500 µL). Crude was diluted with CH₂Cl₂, transferred to a separatory funnel, and washed with water. Aqueous layer was re-extracted with CH₂Cl₂, and organic fractions were gathered, dried over Na2SO4, filtered, and concentrated in rotary evaporator. Silica gel flash chromatography (EtOAc/n-Hex) afforded 1.4 g (2.3 mmol, 66%) of a mixture of 4α and 4β . Physical data for major product $4a R_f = 0.5$ (EtOAc/n-Hex, 1 : 3), ¹H NMR (500 MHz, CDCl₃): 7.35–7.26 (m, 5H), 6.11 (d, J = 10.2 Hz, H-3), 5.78 (ddd, J = 2.2, 2.2 and 10.2 Hz, H-2), 5.39 (d, J = 2.7 Hz, H-4'), 5.21 (dd, J = 8.0 and 10.4 Hz, H-2'), 5.08 (d, J = 2.2 Hz, H-1), 5.00 (dd, J = 3.4 and 10.4 Hz, H-3'), 4.79 (d, J = 11.8 Hz, OC \underline{H}_2 Ph), 4.57 (d, J = 8.9 Hz, H-1'), 4.58 (d, J = 11.8 Hz, OC H_2 Ph), 4.20 (m, H-4, H-6', H-6 and H-6), 4.11 (m, H-6' and H-5), 3.92 (dd, *J* = 6.1 and 6.1 Hz, H-5'), 2.16 (s, 3H), 2.16 (s, 3H), 2.08 (s, 3H), 2.04 (s, 3H), 1.98 (s, 3H); APT ¹³C{¹H} NMR (125 MHz, CDCl₃): 170.7, 170.4, 170.3, 170.1, 169.4, 137.7, 131.6 (C-3), 128.4 (2C), 127.9 (2C), 127.8, 127.1 (C-2), 102.2 (d, C-1'), 93.6 (d, C-1), 73.4, 70.9 (C-3'), 70.8 (C-5'), 70.1 (O<u>C</u>H₂Ph), 68.8 (C-2'), 67.6 (C-5), 66.9 (C-4'), 63.0 (C-6), 61.3 (C-6'), 20.9, 20.7, 20.7, 20.7, 20.6

Benzyl 4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-6-Oacetyl-α-D-mannopyranoside (5) and Benzyl 4-O-(2,3,4,6-tetra-Oacetyl-β-D-galactopyranosyl)-6-O-acetyl-β-D-allopyranoside (6). The mixture of Ferrier adducts 4α and 4β (6.38 g, 10.5 mmol, α/ $\beta = 6:1$) and *N*-methylmorpholine *N*-oxide (1.7 g, 12.5 mmol), were dissolved in acetone (40 mL) at room temperature. Then, OsO₄ (1.3 ml, 4% sol. in water, 0.20 mmol) was added and the

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system was maintained for 12 hours at room temperature under vigorous stirring. After that period, TLC showed total consumption of the starting olefin and the predominance of two more polar spots. Then, the volatiles were removed in rotary evaporator and the resulting syrup was purified by normal phase flash chromatography using an isocratic EtOAc/n-Hex (7:3) elution system. Chromatographic separation afforded the diol 5 (3.6 g, 5.6 mmol, 53%) as a white foam, the diol 6 (758.3 mg, 1.18 mmol, 11%) as a white foam, and a mixed fraction of 5 and 6 (1.83 g, 5/6 = 12 : 1). The combined isolated yield for this reaction was 92%. Physical data for major derivative 5 (Gal β (1 \rightarrow 4)Man): $R_{\rm f}$ = 0.57 (EtOAc), ¹H NMR (500 MHz, $CDCl_3$: 7.30–7.22 (m, 5H), 5.33 (dd, J = 0.7 and 3.4 Hz, H-4'), 5.17 (dd, J = 8.0 and 10.5 Hz, H-2'), 4.93 (dd, J = 3.4 and 10.5, H-3'), 4.88 (d, J = 1.0 Hz, H-1), 4.64 (d, J = 11.8 Hz, 1H), 4.45 (d, J = 8.0 Hz, H-1'), 4.44 (d, J = 11.8 Hz, 1H), 4.22 (dd, J = 2.0 and 11.8 Hz, H-6'), 4.11 (dd, J = 4.8 and 11.6 Hz, H-6), 4.06 (dd, J = 7.6 and 11.3 Hz, H-6), 4.01 (dd, J = 5.5 and 11.7 Hz, H-6'), 4.03 (m, H-2 and H-5'), 3.94 (dd, J = 1.0 and 8.7 Hz, H-3), 3.96 (m, 2H, J)H-2 and H-5'), 3.88 (dd, J = 3.5 and 8.6 Hz, H-3), 3.82 (ddd, J =1.6, 5.4, and 9.9 Hz, H-5), 3.67 (dd, *J* = 8.9 and 9.6 Hz, H-4), 2.10 (s, 3H), 2.06 (s, 3H), 2.03 (s, 3H), 2.02 (s, 3H), 1.91 (s, 3H); APT ¹³C{¹H} NMR (125 MHz, CDCl₃): 170.6, 170.2, 169.9, 169.8, 169.4, 136.6, 128.4, 127.9 (x 2Cs), 127.8 (x 2Cs), 101.7 (C-1'), 97.9 (C-1), 80.4, 71.3, 70.7, 70.1, 69.4, 69.2, 68.5, 67.7, 66.7, 62.8, 61.6, 20.8, 20.5, 20.4, 20.4, 20.3; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for C₂₉H₃₈NaO₁₆ 665.2058; found 665.2137; physical data for minor derivative 6 (Gal $\beta(1 \rightarrow 4)$ All): $R_f = 0.48$ (EtOAc/n-Hex, 4:1), ¹H NMR (400 MHz, CDCl₃): 7.35 (m, 5H), 5.39 (d, J =2.8 Hz, H-4'), 5.24 (dd, J = 8.2 and 10.3 Hz, H-2'), 5.01 (dd, J = 3.3 and 10.5 Hz, H-3'), 4.91 (d, J = 11.8 Hz, 1H, PhCH₂O), 4.73 (d, J = 7.8 Hz, H-1), 4.64 (d, J = 11.8 Hz, 1H, PhCH₂O), 4.57 (d, J =8.0 Hz, H-1'), 4.36 (d, J = 2.1 Hz, H-3), 4.32 (dd, J = 1.6 and 12.2 Hz, H-6), 4.21 (dd, J = 7.8 and 11.2 Hz, H-6'), 4.06 (m, 2H, H-6', H-6), 3.96 (m, 2H, H-5, H-5'), 3.64 (dd, J = 2.3 and 9.5 Hz, H-4), 3.51 (ddd, J = 3.0, 7.6, and 10.0 Hz, H-2), 2.78 (s, 1H, OH), 2.59 (d, J = 7.6 Hz, 1H, OH) 2.17 (s, 3H), 2.11 (s, 3H), 2.08 (s, 3H), 2.07 (s, 3H), 1.99 (s, 3H); APT ${}^{13}C{}^{1}H$ NMR (100 MHz, CDCl₃): 170.7, 170.5, 170.1, 170.0, 169.5, 137.1, 128.5, 128.1, 127.9, 101.8 (C-1'), 99.5 (C-1), 78.1 (C-4), 71.4 (C-5'), 71.2 (O<u>C</u>H₂Ph), 70.6 (C-2), 70.5 (C-3'), 69.6 (C-5), 69.5 (C-3), 68.4 (C-2'), 66.9 (C-4'), 63.2 (C-6), 61.4 (C-6'), 20.9, 20.6, 20.6, 20.5, 20.5; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for C₂₉H₃₈NaO₁₆ 665.2058; found 665.2141.

General procedure for acetylation

Diol, Ac_2O (6.0 equiv.), and pyridine (3.0 equiv.) were dissolved in dry CH_2Cl_2 (5.0 mL mmol⁻¹) at room temperature and stirred until TLC showed total consumption of the starting diol (~15 min). Then, the mixture was cooled to 0 °C with an ice bath and the excess of Ac_2O was quenched by addition of EtOH (30 equiv.). The volatiles were then removed in rotary evaporator and the residue was re-dissolved in EtOAc, transferred to a separation funnel, and washed with H₂O followed by 1 M HCl, and NaHCO₃. The organic layer was collected, dried over Na₂SO₄, filtered, and concentrated in rotary evaporator. The crude acetate was purified by normal phase flash chromatography using EtOAc/n-Hex (1 : 1 to 7 : 3) gradient as eluent system.

Benzyl 2,3-di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-6-O-acetyl-α-D-mannopyranoside (7). Following the general procedure for acetylation, diol 5 (2.11 g, 3.3 mmol), Ac2O (1.9 mL, 19.8 mmol), and pyridine (797 µL, 9.9 mmol), afforded the respective octaacetate 7 (2.13 g, 2.9 mmol, 88%) as a white solid. $R_f = 0.44$ (EtOAc/*n*-Hex, 3 : 2), ¹H NMR (400 MHz, $CDCl_3$: 7.31–7.22 (m, 5H), 5.32 (dd, J = 3.6 and 8.8 Hz, H-3), 5.27 (dd, J = 1.0 and 3.4 Hz, H-4'), 5.20 (dd, J = 1.9 and 3.6 Hz, H-2), 5.07 (dd, J = 7.9 and 10.5 Hz, H-2'), 4.89 (dd, J = 3.4 and 10.4 Hz, H-3'), 4.74 (d, J = 1.8 Hz, H-1), 4.62 (d, J = 11.9 Hz, 1H, OC H_2 Ph), 4.48 (d, J = 7.9 Hz, H-1'), 4.46 (d, J = 11.9 Hz, 1H, OCH₂Ph), 4.29 (dd, J = 1.7 and 11.9 Hz, H-6a), 4.10 (m, 2H, H-6b and H-6a'), 3.96 (dd, J = 7.7 and 11.9 Hz, H-6b'), 3.86 (m, 2H, H-5 and H-4), 3.80 (dd, J = 1.0 and 7.7 Hz, H-5'), 2.08 (s, 3H), 2.07 (s, 3H), 2.04 (s, 3H), 1.99 (s, 3H), 1.98 (s, 3H), 1.97 (s, 3H), 1.90 (s, 3H); ¹³C{¹H} NMR (100 MHz, CDCl₃): 170.5, 170.4, 170.2, 170.1, 169.8, 169.3, 169.2, 136.2, 128.6 (×2C), 128.2 (×2C), 128.2, 101.1 (C-1), 96.4 (C-1), 74.0, 71.0, 70.4, 69.7, 69.6, 69.6, 69.2, 69.1, 66.6, 62.6, 60.8, 20.9, 20.8, 20.8, 20.6, 20.6, 20.6, 20.5; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for $C_{33}H_{42}NaO_{18}$ 749.2269; found 749.2399.

Benzyl 2,3-di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-6-O-acetyl-β-D-allopyranoside (8). Following the general procedure for acetylation, diol 6 (124.2 mg, 0.19 mmol), Ac₂O (107 µL, 1.14 mmol), and pyridine (45 µL, 0.57 mmol) afforded the respective octaacetate 8 (109.0 mg, 0.15 mmol, 81%) as a white solid. $R_{\rm f} = 0.45$ (EtOAc/n-Hex, 3 : 2), ¹H NMR $(400 \text{ MHz}, \text{CDCl}_3)$: 7.33 (m, 5H), 5.76 (t, J = 2.5 Hz, H-3), 5.33 (d, J = 2.6 Hz, H-4'), 5.15 (dd, J = 7.9 and 10.3 Hz, H-2'), 4.98 (dd, J = 3.2 and 10.4 Hz, H-3'), 4.89 (d, J = 12.1 Hz, 1H, OC \underline{H}_2 Ph), 4.83 (d, J = 8.3 Hz, H-1), 4.77 (dd, J = 2.5 and 8.2 Hz, H-2), 4.63 (d, J =12.1 Hz, 1H, OCH_2Ph), 4.56 (d, J = 7.8 Hz, H-1'), 4.38 (d, J =10.1 Hz, H-6), 4.03 (m, 4H, H-5, H-6', H-6Gal, H-6'Gal), 3.91 (d, J = 6.4 and 6.4 Hz, H-5'), 3.80 (d, *J* = 2.5 and 9.3 Hz, H-4), 2.12 (s, 3H), 2.11 (s, 3H), 2.08 (s, 3H), 2.07 (s, 3H), 2.06 (s, 3H), 1.98 (s, 3H), 1.97 (s, 3H); ¹³C{¹H} NMR (100 MHz, CDCl₃): 170.6, 170.6, 170.1, 169.9, 169.6, 169.6, 169.5, 137.1, 128.3 (×2C), 127.8, 127.6 (×2C), 100.7 (C-1'), 97.4 (C-1), 74.9 (C-4), 70.9 (O<u>C</u>H₂Ph), 70.8 (C-5'), 70.7 (C-3'), 70.6 (C-5), 70.1 (C-2), 68.9 (C-3), 68.7 (C-2'), 66.8 (C-4'), 62.9 (C-6), 61.3 (C-6'), 20.8, 20.8, 20.6, 20.6, 20.6, 20.5, 20.5; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for $C_{33}H_{42}NaO_{18}$ 749.2269; found 749.2330.

2,3-Di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl β -D-galactopyranosyl)-6-O-acetyl- α -D-mannopyranose (9). In a round-bottomed flask, benzyl glycoside 7 (1.92 g, 2.6 mmol) was dissolved in EtOH (300 mL) and Pd/C (260 mg) was added under inert atmosphere (N₂). The system was closed with a septum and the N₂ replaced by a reducing H₂ atmosphere *via* four vacuum/H₂ cycles. The reaction was vigorously stirred at room temperature until TLC showed complete consumption of the starting benzyl glycoside (~3 hours). The catalyst was removed by passing the suspension through a Celite® pad and the volatiles were removed by rotary evaporation. The resulting crude product was used in the next step without further purification. This procedure afforded the alcohol 9 (1.59 g, 2.5 mmol, 96%, α : β = 8 : 1) as a colorless syrup. Physical data for major anomer (α): $R_f = 0.62$ (EtOAc/n-Hex, 19 : 1), ¹H NMR (400 MHz, CDCl₃): 5.34 (dd, J = 3.5 and 9.5 Hz, H-3), 5.28 (d, J = 2.5 Hz, H-4'), 5.15 (dd, J = 1.5 and 3.3 Hz, H-2), 5.08 (d, J = 1.5 Hz, H-1), 5.05 (dd, J = 7.8 and 10.4 Hz, H-2'), 4.90 (dd, J = 3.3 and 10.5 Hz, H-3'), 4.49 (d, J = 7.9 Hz, H-1'), 4.38 (dd, J = 3.5 and 13.2 Hz, H-6_{Man}), 4.10 (m, 3H, H-5, H-6_{Gal}, and H-6'_{Man}), 3.96 (dd, J = 7.6 and 11.0 Hz, H-6'_{Gal}), 3.84 (m, 2H, H5' and H4), 2.09 (s, 3H), 2.07 (s, 3H), 2.06 (s, 3H), 1.99 (s, 3H), 1.99 (s, 3H), 1.97 (s, 3H), 1.90 (s, 3H); ¹³C{¹H} NMR (100 MHz, CDCl₃): 170.8, 170.5, 170.3, 170.2, 170.1, 169.6, 169.3, 100.9 (C-1'), 91.8 (C-1), 74.2 (C-4), 70.9 (C-3'), 70.4 (C-5'), 70.3 (C-2), 69.2 (C-2'), 69.1 (C-3), 68.9 (C-5), 66.7 (C-4'), 62.6 (C-6), 60.9 (C-6'), 20.9, 20.8, 20.6, 20.6, 20.6, 20.5; HRMS (ESI/Q-TOF) *m*/*z*: $[M + Na]^+$ calcd for C₂₆H₃₆NaO₁₈ 659.1799; found 659.1939.

4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-1-(2,2,2-trichloroethanimidate)-a-p-mannopyranose 2,3,6-triacetate (10). To a solution of alcohol 9 (535.2 mg, 0.84 mmol), and Cl_3CCN (400 mL, 5.04 mmol) in dry CH₂Cl₂ (4.0 mL), 1,8-diazabicyclo [5.4.0]undec-7-ene (DBU, 37 mL, 0.25 mmol) was added and the system was stirred at room temperature for 18 hours. After that period, TLC showed total consumption of the starting material. The volatiles were then removed in rotary evaporator and the residue was directly purified by normal phase flash chromatography. The imidate 10 (446.1 mg, 0.57 mmol, 68%) was obtained as a yellowish foam. $R_{\rm f} = 0.47$ (EtOAc/*n*-Hex/Et₃N, 60: 40: 1), ¹H NMR (400 MHz, CDCl₃): 8.68 (s, 1H), 6.15 (d, J =2.0 Hz, H-1), 5.37 (m, 2H, H-2, H-3), 5.29 (dd, J = 0.8 and 3.3 Hz, H-4'), 5.09 (dd, J = 8.0 and 10.5 Hz, H-2'), 4.91 (dd, J = 3.4 and 10.5 Hz, H-3'), 4.52 (d, J = 7.8 Hz, H-1'), 4.37 (dd, J = 1.9 and 11.9 Hz, H-6), 4.12 (dd, J = 6.1 and 11.1 Hz, H-6'), 4.10 (dd, J = 5.2 and 11.1 Hz, H-6), 4.05 (m, H-5), 3.97 (dd, J = 7.7 and 11.2 Hz, H-6'), 3.93 (dd, J = 9.6 and 9.6 Hz, H-4), 3.83 (dd, J = 7.4and 7.4 Hz, H-5'), 2.11 (s, 3H), 2.09 (s, 3H), 2.04 (s, 3H), 1.99 (s, 3H), 1.98 (s, 3H), 1.98 (s, 3H), 1.91 (s, 3H). APT ${}^{13}C{}^{1}H$ NMR (100 MHz, CDCl₃): 170.4, 170.4, 170.2, 170.1, 169.5, 169.3, 169.2, 160.0 (<u>C</u>=NH), 101.4 (C-1'), 94.5 (C-1), 90.5 (<u>C</u>Cl₃), 74.2 (C-4), 71.5 (C-5), 71.0 (C-3'), 70.5 (C-5'), 69.2 (C-3), 69.2 (C-2'), 67.9 (C-2), 66.5 (C-4'), 62.2 (C-6), 60.8 (C-6'), 20.8, 20.8, 20.8, 20.7, 20.7, 20.6, 20.6; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for C₂₈H₃₆Cl₃NNaO₁₈ 802.0896; found 802.0851.

3-Azidopropyl 2,3-di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl-β-Dgalactopyranosyl)-6-O-acetyl-a-D-mannopyranoside (11). Imidate 10 (367.0 mg, 0.47 mmol) and 3-azidopropanol (142.5 mg, 1.41 mmol) were placed in a round-bottomed flask. Prior to glycosyl donor activation, the system was dried via azeotropic removal of H_2O traces with toluene (3 cycles of ~2.0 mL) by rotary evaporation. The reactants were then dissolved in dry CH₂Cl₂ (3.0 mL) and cooled to 0 °C with an ice bath. Imidate was activated by addition of BF₃·Et₂O (30 mL, 0.24 mmol) and the reaction was allowed to warm at room temperature under stirring. After 3 hours, TLC showed a significant amount of remaining imidate and more BF₃Et₂O (50 mL, 0.4 mmol) was added. After 1 hour, TLC showed reaction completion and the excess of Lewis acid was quenched with Et₃N (1.0 mL). Then, the volatiles were removed by rotary evaporation and the residue directly purified by normal phase flash chromatography using

EtOAc/n-Hex (3 : 7 to 3 : 2) gradient as eluent system. Glycoside 11 (240.0 mg, 0.33 mmol, 70%) was obtained as a colorless oil. $R_{\rm f} = 0.43$ (EtOAc/*n*-Hex, 7 : 3), ¹H NMR (500 MHz, CDCl₃): 5.28 (dd, *J* = 1.0 and 3.4 Hz, H-4′), 5.25 (dd, *J* = 3.6 and 9.1 Hz, H-3), 5.15 (dd, *J* = 1.9 and 3.6 Hz, H-2), 5.07 (dd, *J* = 7.9 and 10.5 Hz, H-2'), 4.90 (dd, J = 3.4 and 10.4 Hz, H-3'), 4.68 (d, J = 1.9 Hz, H-1), 4.48 (dd, J = 8.0 Hz, H-1′), 4.35 (dd, J = 1.2 and 11.9 Hz, H- 6_{Man}), 4.12 (dd, J = 6.0 and 11.1 Hz, H- $6'_{\text{Man}}$), 4.11 (dd, J = 1.3and 11.6 Hz, H-6_{Gal}), 3.96 (dd, J = 7.8 and 11.1 Hz, H-6'_{Gal}), 3.82 (m, 3H, H-4, H-5, and H-5'), 3.71 (m, 1H), 3.43 (m, 1H), 3.35 (t, J = 6.5 Hz, 2H), 2.09 (s, 3H), 2.07 (s, 3H), 2.06 (s, 3H), 2.00 (s, 3H), 1.99 (s, 3H), 1.97 (s, 3H), 1.91 (s, 3H), 1.79 (m, 2H); APT ¹³C{¹H} NMR (125 MHz, CDCl₃): 170.5, 170.4, 170.2, 170.1, 169.8, 169.3, 169.2, 101.2 (C-1'), 97.4 (C-1), 74.4, 70.9 (C-3'), 70.4, 69.6 (C-2), 69.4 (C-3), 69.2 (C-2'), 69.1, 66.6 (C-4'), 64.8 (OCH₂CH₂CH₂N₃), 62.5 (C-6), 60.8 (C-6'), 48.1 (OCH₂CH₂CH₂N₃), 28.6 (OCH₂CH₂-CH₂N₃), 20.9, 20.8, 20.8, 20.7, 20.6, 20.6, 20.5; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for $C_{29}H_{41}N_3NaO_{18}$ 742.2283; found 742.2336.

3-Azidopropyl 4-O-β-p-galactopyranosyl-α-p-mannopyranoside (1). In a round-bottomed flask provided with a magnetic bar, per-acetylated disaccharide 11 (221.1 mg, 0.31 mmol) and NaOMe (\sim 10 mg) were dissolved in CH₃OH (3.0 mL) at room temperature. The solution was stirred until TLC showed total consumption of the starting disaccharide (~12 h). The reaction was then diluted with CH₃OH (3.0 mL) and neutralized with Amberlite IR-120 (H form). The mixture was filtered, and the volatiles removed by rotary evaporation. This procedure afforded the target azide-derived $Gal\beta(1 \rightarrow 4)\alpha$ Man disaccharide 1 as a white foam (127.6 mg, 0.30 mmol, 97%). $R_{\rm f} = 0.50$ (MeOH/ CH₂Cl₂, 1 : 4), ¹H NMR (400 MHz, CD₃OD): 4.78 (s, H-1), 4.36 (d, J = 7.4 Hz, H-1'), 3.90 (m, 4H), 3.81 (m, 4H), 3.73 (dd, J = 4.3 and 11.5 Hz, H-6), 3.63 (m, 2H), 3.53 (m, 3H), 3.41 (m, 2H), 1.88 (m, 2H); ¹³C{¹H} NMR (100 MHz, CD₃OD): 103.8 (C-1'), 100.0 (C-1), 76.9, 75.8, 73.4, 71.7, 71.2, 70.1, 69.8, 69.0, 64.1, 61.2, 60.7, 48.1, 28.5; HRMS (ESI/Q-TOF) m/z: $[M + Na]^+$ calcd for $C_{15}H_{27}N_3NaO_{11}$ 448.1543; found 448.1654.

2,3-Di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-6-O-acetyl-a-D-mannopyranosyl H-phosphonate (12). In a round-bottomed flask provided with a magnetic bar, imidazole (2.6 g, 38.2 mmol) and Et₃N (5.4 mL, 38.7 mmol) were suspended in MeCN (30.0 mL) and cooled to 0 °C. PCl₃ (1.1 mL, 12.6 mmol) was then added, and the resulting suspension was vigorously stirred at 0 °C for 20 minutes. Then, a solution of alcohol 9 (1.0 g, 1.6 mmol) in MeCN (5.0 mL) was added dropwise over 30 minutes under vigorous stirring. The reaction was allowed to warm to room temperature. After 4 h, TLC showed complete consumption of the starting disaccharide, and the system was quenched by addition of 1 M TEAB buffer (27 mL, pH 7.2) and left stirring for 20 minutes. The volatiles were then removed by rotary evaporation and the residue was re-dissolved in CHCl₃, transferred to a separation funnel, and washed with 1 M TEAB buffer. The organic layer was collected, dried over Na₂SO₄, filtered, concentrated in rotary evaporator, and the resulting crude was purified by normal phase flash chromatography using a MeOH/CH₂Cl₂ gradient (1: 24 to 1: 9) containing 1% Et₃N as mobile phase. Purification afforded H-

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phosphonate 12 (409.1 mg, 0.50 mmol, 42%) as a white solid. R_f = 0.50 (MeOH/CH₂Cl₂, 1 : 4), ¹H NMR (400 MHz, CDCl₃): 6.92 $(d, J = 638 \text{ Hz}, P-\underline{H}), 5.47 (dd, {}^{3}J_{H1-H2} = 2.3 \text{ and } {}^{3}J_{H-P} = 10.5 \text{ Hz},$ H-1), 5.32 (dd, *J* = 3.4 and 9.5 Hz, H-3), 5.29 (d, *J* = 3.0 Hz, H-4'), 5.21 (m, H-2), 5.07 (dd, I = 7.4 and 10.0 Hz, H-2'), 4.89 (dd, I =3.5 and 10.0 Hz, H-3'), 4.47 (d, J = 10.5 Hz, H-1'), 4.35 (dd, J = 3.2 and 12.0 Hz, H-6_{Man}), 4.12 (m, 3H, H-5, H-6_{Gal}, and H-6'_{Man}), 3.96 (dd, J = 7.7 and 11.0 Hz, H-6[']_{Gal}), 3.84 (t, J = 9.3 Hz, H-5[']), 3.82 (dd, J = 7.2 and 7.2 Hz, H-4), 3.02 (q, J = 7.3 Hz, 6H, N(CH₂CH₃)₃), 2.09 (s, 3H), 2.06 (s, 3H), 2.05 (s, 3H), 1.99 (s, 3H), 1.98 (s, 3H), 1.94 (s, 3H), 1.91 (s, 3H), 1.30 (t, J = 7.3 Hz, 9H, $N(CH_2CH_3)_3$; ¹³C{¹H} NMR (100 MHz, CDCl₃): 170.6, 170.4, 170.2, 170.1, 169.8, 169.6, 169.1, 101.1 (C-1'), 92.6 (d, ${}^{2}J_{C-P} =$ 3.7 Hz, C-1), 74.0 (C-5'), 70.9 (C-3'), 70.4 (C-4), 70.1 (d, ${}^{3}J_{C-P} =$ 7.2 Hz, C-2), 69.9 (C-5), 69.4 (C-3), 69.1 (C-2'), 66.7 (C-4'), 62.4 (C-6), 60.8 (C-6'), 45.7 (N(CH2CH3)3), 20.8, 20.8, 20.8, 20.6, 20.6, 20.6, 20.5, 8.6 (N(CH₂<u>C</u>H₃)₃); ³¹P NMR (162 MHz, CDCl₃): -0.10 $(dd, {}^{1}J_{P-H} = 638 \text{ Hz}, {}^{3}J_{P-H1} = 9.8 \text{ Hz}); \text{HRMS} (\text{ESI/Q-TOF}) m/z: [M]$ - H]⁻ calcd for C₂₆H₃₆O₂₀P 699.1543; found 699.1545.

Triethylammonium 3-azidopropyl 2,3-di-O-acetyl-4-O-(2,3,4,6-tetra-O-acetyl-β-D-galactopyranosyl)-6-O-acetyl-α-D-mannopyranosyl phosphate (13). In a round-bottomed flask provided with a magnetic bar, H-phosphonate 12 (100.1 mg, 0.12 mmol) and 3-azidopropanol (14 µL, 0.15 mmol) were dissolved in dry THF (2.8 mL) and cooled to 0 °C with an ice bath. To the resulting solution, PivCl (37 µL, 0.3 mmol) was added and the system was stirred allowing it to warm to room temperature. After 3 hours, TLC showed complete consumption of the starting H-phosphonate and a solution of I₂ (121.8 mg, 0.48 mmol) in pyridine/H₂O (19:1, 1.0 mL) was added. TLC showed the oxidation of H-phosphonate to phosphate completes in less than 30 minutes. The reaction was then quenched by addition of 10% $Na_2S_2O_3$ (6.0 mL), transferred to a separation funnel, and extracted with CHCl₃. The organic laver was then successively washed with brine, and 1 M TEAB buffer. The collected organic fraction was then dried over Na₂SO₄, filtered, and concentrated in rotary evaporator. The resulting crude was purified by normal phase flash chromatography using a MeOH/CH₂Cl₂ gradient (1:99 to 1:9) containing 1% of Et₃N. This procedure afforded the phosphodiester 13 (30.2 mg, 0.03 mmol, 30%) as a white solid. $R_{\rm f} = 0.5$ (MeOH/ CH₂Cl₂, 15:85), ¹H NMR (400 MHz, CDCl₃): 11.9 (br s, 1H, $HN^{+}(CH_{2}CH_{3})_{3})$, 5.41 (dd, ${}^{3}J_{H1-H2} = <1.0$ Hz and ${}^{3}J_{H1-P} = 7.9$ Hz, H-1), 5.31 (dd, J = 3.5 and 9.4 Hz, H-3), 5.28 (d, J = 3.3 Hz, H-4'), 5.24 (s, H-2), 5.07 (dd, J = 8.2 and 10.4 Hz, H-2'), 4.89 (dd, J = 3.2 and 10.4 Hz, H-3'), 4.47 (d, J = 8.0 Hz, H-1'), 4.36 (d, J = 10.9 Hz, H-6_{Man}), 4.12 (m, 3H, H-5, H-6'_{Man}, H-6_{Gal}), 3.95 (dd, J = 7.9 and 11.0 Hz, H-6'_{Gal}), 3.92 (m, 2H, OCH₂CH₂CH₂N₃), 3.87 (t, J =9.7 Hz, H-4), 3.81 (dd, J = 6.8 and 8.0 Hz, H-5'), 3.37 (t, J =6.9 Hz, OCH₂CH₂CH₂CH₂N₃), 3.04 (m, 6H, HN⁺(CH₂CH₃)₃), 2.09 (s, 3H), 2.06 (s, 3H), 2.06 (s, 3H), 1.99 (s, 3H), 1.98 (s, 3H), 1.94 (s, 3H), 1.91 (s, 3H), 1.83 (m, 2H, OCH₂CH₂CH₂N₃), 1.33 (t, J =7.3 Hz, 9H, $HN^{+}(CH_2CH_3)_3$). APT ${}^{13}C{}^{1}H{}$ (100 MHz, $CDCl_3$): 170.5, 170.4, 170.2, 170.1, 169.8, 169.6, 169.1, 101.1 (C-1'), 93.4 $(d, {}^{2}J_{C1-P} = 6.0 \text{ Hz}, \text{C-1}), 73.8 (\text{C-5}'), 70.9 (\text{C-3}'), 70.4 (\text{C-4}), 69.8 (d, C-4))$ ${}^{2}J_{C2-P} = 6.8$ Hz, C-2), 69.8 (C-5), 69.5 (C-3), 69.1 (C-2), 66.6 (C-4'), 62.7 (d, ${}^{2}J_{C1-P} = 5.3$ Hz, OCH₂CH₂CH₂N₃), 62.5 (C-6), 60.8 (C-6'),

48.2 (OCH₂CH₂CH₂N₃), 45.8 (HN+(CH₂CH₃)₃), 30.1 (d, ${}^{2}J_{C-P} =$ 7.4 Hz, OCH₂CH₂CH₂N₃), 20.9, 20.9, 20.8, 20.6, 20.6, 20.5, 8.6 (HN⁺(CH₂CH₃)₃); ³¹P NMR (162 MHz, CDCl₃): -3.09 (d, ${}^{3}J_{P-}$ H₁ = 6.9 Hz); HRMS (ESI/Q-TOF) *m/z*: [M - H]⁻ calcd for C₂₉H₄₁N₃O₂₁P 798.1976, found 798.1972.

3-Azidopropyl 4-O-β-D-galactopyranosyl-α-D-mannopyranosyl phosphate (2). In a round-bottomed flask provided with a magnetic bar, phosphodiester 13 (83.0 mg, 0.09 mmol) and NaOMe (14.6 mg, 0.27 mmol) were suspended in CH₃OH (5.0 mL) and stirred at room temperature for 48 hours. Then, the reaction was diluted with CH₃OH (5.0 mL) and NH₄Cl was added dropwise until reaching pH \sim 7. The volatiles were then removed in rotary evaporator and the crude was subjected to purification by flash reverse phase chromatography using C-18 as stationary phase and MeCN/H₂O gradient (1:99 to 1:1) as eluent system. This procedure afforded the deprotected azidoderived Gal $\beta(1 \rightarrow 4)\alpha$ Man phospho-glycoside 2 (33.7 mg, 0.06 mmol, 67%) as a white solid. ¹H NMR (400 MHz, CD_3OD): 5.34 (dd, ${}^{3}J_{H1-H2} = 1.5$ Hz, ${}^{3}J_{H1-P} = 7.9$ Hz, H-1), 4.25 (d, J =7.5 Hz, H-1'), 3.85 (m, 2H), 3.83 (m, 2H), 3.79 (m, 5H), 3.70 (m, 3H), 3.61 (dd, J = 4.1 and 11.5 Hz, H-6), 3.50 (dd, J = 4.1 and 7.3 Hz, 1H), 3.45 (dd, *J* = 7.6 and 9.7 Hz, 1H), 3.40 (d, *J* = 2.6 Hz, H-4'), 3.36 (m, 4H), 1.79 (m, 2H), 1.26 (t, J = 7.2 Hz, 3H); APT ¹³C ${^{1}H}$ (100 MHz, CD₃OD): 103.9 (C-1'), 96.1 (d, ${^{2}J_{C1-P}} = 5.5$ Hz, C-1), 76.6, 75.7, 73.3, 72.6, 71.1, 70.5 (d, $J_{C-P} = 8.6$ Hz), 69.1, 69.0, 62.3 (d, $J_{C-P} = 5.8$ Hz, OCH₂CH₂CH₂N₃), 61.3, 60.4, 52.2, 29.7 (d, $J_{C-P} = 7.6$ Hz, OCH₂CH₂CH₂N₃); ³¹P NMR (162 MHz, CD₃OD): -1.82 (q, J = 7.1 Hz); HRMS (ESI/Q-TOF) m/z: $[M - H]^-$ calcd for C15H27N3O14P 504.1236; found 504.1239.

Conflicts of interest

The authors declare no competing financial interests or personal relationships that could have influenced the performance of this work.

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