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Reagent controlled stereoselective synthesis of teichoic acid α -(1,2)-glucans†

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The stereoselective construction of 1,2-cis-glycosidic linkages is key in the assembly of biologically relevant glycans, but remains a synthetic challenge. Reagent-controlled glycosylation methodologies, in which external nucleophiles are employed to modulate the reactivity of the glycosylation system, have become powerful means for the construction of 1,2-cis-glycosidic linkages. Here we establish that nucleophilic additives can support the construction of α -1,2-glucans, and apply our findings in the construction of a D-alanine kojibiose functionalized glycerol phosphate teichoic acid fragment. This latter molecule can be found in the cell wall of the opportunistic Gram-positive bacterium, *Enterococcus faecalis* and represents a structural element that can possibly be used in the development of therapeutic vaccines and diagnostic tools.

Introduction

α-Glucans, polysaccharides composed of glucose monosaccharides connected through α-glycosidic linkages, are widely found in nature. The most abundant α -glucans are the α -1,4glucans, that occur as starch and glycogen, and can be found as extracellular polysaccharides surrounding bacteria, such as Mycobacterium tuberculosis.² α-1,3-Glucans are prominent components of fungal cell walls and play a role in the interaction between the pathogenic fungus Aspergillus fumigatus and the host immune system.³ α-1,2-Glucans, also referred to as kojioligosaccharides, have been found in koji extract, sake, beer and honey and they have also been encountered in bacterial cell walls.4 Kojibiose, kojitriose and kojitetraose have been found as part of the lipoteichoic acid (LTA) of Enterococcus faecalis, which is a commensal Gram-positive bacterium.⁵ Although this bacterium is generally of low virulence, it is the causative agent of life-threatening infections in immunocomKojibiose is a prominent appendage of *E. faecalis* LTA, and it can be decorated with p-alanyl esters at both C6-hydroxyl groups of the disaccharide. P-Alanylation plays a critical role in the virulence of the bacteria and it renders the bacteria less susceptible to (cationic) antimicrobial peptides. *E. faecalis* LTA fragments, containing a single α-glucosyl substituent have previously been explored as components in synthetic vaccine conjugates, and antibodies raised by these conjugates have been shown to be opsonophagocytic and capable of inducing killing of the bacteria in both active and passive immunization strategies. The microheterogeneity of naturally occurring LTA makes it challenging if not impossible- to establish clear structure-activity relationships for these molecules, and therefore organic synthesis is the method of choice to generate well-defined single LTA melecules (2.10 Only a few shorpical synthesis).

promised patients and one of the most prominent hospital

bacteria.⁶ LTA is an anionic polymer featuring a glycolipid tail,

through which it is anchored in the cell wall of the bacteria. The polyanionic part is composed of glycerol phosphate

repeating units, which can be functionalised with glycans and

p-alanine esters at C2 of the glycerol moieties (Fig. 1).7

The microheterogeneity of naturally occurring LTA makes it challenging if not impossible- to establish clear structure–activity relationships for these molecules, and therefore organic synthesis is the method of choice to generate well-defined single LTA molecules. Only a few chemical syntheses of α -(1,2)-glucans have been reported. Takeo and Suzuki reported a procedure to assemble kojitriose, kojitetraose and kojipentaose based on Koenings-Knorr-type glycosylations, using anomeric chlorides under the activation of silver per-

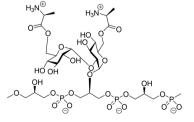


Fig. 1 The structure of $\it E. faecalis \, p$ -alanine kojibiose functionalized LTA.

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chlorate. 11 Hogendorf et al. relied on Crich's α-glucosylation protocol, which involves the use of benzylidene glucose donors for the synthesis of a kojibiose glycerol building block.9 We have previously shown that additive-controlled glycosylations¹² of per-benzylated glucosyl imidate donors can be used effectively for the introduction of α -(1,4)- and α -(1,3)-glucosyl linkages, which we used to assemble a set of M. tuberculosis α -(1,4)-glucans and A. fumigatus α -(1,3)-glucans.¹³ In this strategy we exploited a suite of orthogonal benzyl ethers to discriminate between different hydroxyl functionalities that had to be protected permanently (using a benzyl (Bn) group), and hydroxyl groups that were used for chain elongation (protected with 2-methylnaphthyl (Nap) ethers) and hydroxyl groups from which branching occurred (using para-methoxybenzyl (PMB) ethers). The use of solely benzylic ethers to protect the hydroxyls ensured that the nature of the protecting groups does not induce variation in the reactivity of the building blocks and that we could rely on reactivity tuning by the used external nucleophile to control the stereoselective construction of the α-glucan linkages. This strategy could provide an effective entry to construct α -(1,2)-linkages and generate (substituted) kojioligosaccharides such as encountered in E. faecalis LTA. We here describe the use of additive-controlled glycosylation methodology for the construction of α -(1,2)glucans and we show the application of the methodology in the preparation of a p-alanine-kojibiose functionalized LTA fragment.

Results and discussion

We first explored the assembly of spacer-functionalized kojioligosaccharides. To achieve this aim, donor 6 (Scheme 1) was designed and assembled. The building block bears three permanent Bn-ethers to mask the C3, C4 and C6 hydroxyls and a Nap-ether as a temporary protecting group at the C2-OH. The synthesis of this building block used benzylidene protected glucose building block 1 as starting compound. Selective protection of the C3-OH as the benzyl ether was achieved through the formation of an intermediate tin ketal which was treated with benzyl bromide and cesium fluoride to give compound 2. Protection of the remaining C2-OH with a Nap ether then provided compound 3. Selective opening of the benzylidene and protection of the liberated primary alcohol provided the fully benzylated thiodonor 5. The anomeric thioacetal in this build-

Ph
$$\stackrel{\circ}{\text{NapO}}$$
 SPh $\stackrel{\circ}{\text{R}^2\text{O}}$ SPh $\stackrel{\circ}{\text{BnO}}$ SPh $\stackrel{\circ$

Scheme 1 Synthesis of donor 6. (a) (i) Bu_2SnO , toluene, 120 °C; (ii) CsF, BnBr, 76%. (b) NapBr, NaH, DMF, 91%. (c) BF_3 : Et_2O , TES, DCM. (d) BnBr, NaH, DMF, 88%. (e) (i) NIS, acetone/ H_2O (10:1, v:v); (ii) 2,2,2-trifluoro-N-phenylacetimidoyl chloride, Cs_2CO_3 , acetone, 81% (over two steps).

ing block was hydrolyzed using *N*-iodosuccinimide (NIS) in acetone/water to liberate the anomeric hydroxyl group, which could then be turned into the required *N*-phenyltrifluoroacetimidate **6**.

With donor 6 in hand, the assembly of kojioligosaccharides was explored as depicted in Scheme 2. Firstly, donor 6 was glycosylated with azidopropanol, a primary and relatively reactive alcohol, using the TMSI-Ph₃P=O activation couple. TMSI transforms the anomeric imidate into the α-glucosyl iodide, which under the agency of Ph₃P=O is isomerized into its less stable but more reactive β-counterpart. These conditions formed azidopropyl glucoside 7 in good yield and stereoselectivity $(\alpha:\beta = 10:1)$, in line with results previously described. 13a,14 Deprotection of the C2-O-Nap ether under oxidative conditions (dichlorodicyanobenzoquinone (DDQ) in DCM/H₂O, 10:1 v/v) provided alcohol 8 which was purified by silica gel column chromatography to deliver the pure α-anomer. Next the secondary and less reactive alcohol acceptor 8 was glycosylated with donor 6 under the agency of TfOH and DMF. This latter additive generates a mixture α/β-glucosyl imidinium ions, which are in rapid equilibrium. Substitution of the more reactive β-ion then leads to the stereoselective formation of α-glucosyl linkages. This additive has found wide application in the construction of various 1,2-cis-linked oligosaccharides. 15 The condensation of 8 and 6 provided disaccharide 9 in 95% yield and a 15:1 α:β-ratio. Deprotection of the C2-O-Nap ether by DDQ provided disaccharide acceptor 10 and the third glycosylation, again performed under DMF mediated glycosylation conditions, then provided trisaccharide 11. The stereoselectivity of this glycosylation diminished slightly and 11 was formed as a 10:1 α: β-mixture. The erosion of stereoselectivity developed further upon elongation of the trisaccharide, as the DMF-mediated glycosylation of acceptor 12, obtained after DDQ treatment of 11, with per-benzyl donor

Scheme 2 The assembly of kojioligosaccharides. (a) TMSI, Ph₃P=O, DCM, 36 h, 75%, α : β = 10 : 1. (b) DDQ, DCM : H₂O = 10 : 1 (v : v), 8: 75%; 10: 49%; 12: 47%. (c) DMF, TfOH, DCM, 9: 95%, α : β = 15 : 1; 11: 67%, α : β = 10 : 1; 14: 67%, α : β = 2 : 1.

13 delivered tetrasaccharide 14 as a 2:1 $\alpha:\beta$ -mixture. Possibly the overall structure of the twisted α -(1,2)-oligosaccharide is at the root of the erosion of stereoselectivity, with increasing steric demands leading to lower yields and poorer selectivity.

Although a limitation of the DMF-mediated glycosylation protocol come forward in the above described kojitetraose assembly, the synthesis also showed that the additive controlled glycosylation approach can provide a rapid entry into the assembly of smaller kojibiose derivatives. Therefore, the assembly of an E. faecalis D-Ala-kojibiose LTA fragment was next explored. The synthesis of D-Ala-kojibiose LTA requires the availability of compound 19 featuring the kojibiose, decorated with two p-alanine esters at both primary alcohols, attached to the glycerol through an α -bond (Scheme 3). Compound 19 can be assembled from four building blocks: two benzyl type protected glucosyl donors, 15 and 16, glycerol 17 protected with a benzoyl ester (Bz) and allyl ether and N-benzyloxycarbonyl protected p-alanine 18 (Scheme 3). The glucosyl donors are protected with the triad of Bn, PMB and Nap-ethers to discriminate the alcohol groups that required permanent protection or needed to be removed to introduce the next glucose residue or introduce the D-alanine esters.

The synthesis of the used building blocks is depicted in Scheme 4. Firstly, donor 15 was synthesized from compound 1 by benzylation of the C2 and C3-hydroxyls, regioselective opening of the benzylidene acetal and protection of the liberated primary alcohol with a Nap-ether, to give thio donor 22. Hydrolysis of the thioacetal and introduction of the 2,2,2-trifluoro-*N*-phenylacetimidate formed donor 15. Donor 16 was obtained from compound 2 following a similar sequence of reactions: protection of the C2-OH with a PMB-ether, regioselective opening of the benzylidene acetal and installation of the C6-O-Nap ether provided thioacetal 25, which was transformed into the corresponding imidate donor 16.

With all the required building blocks available, we explored the assembly of a kojibiose LTA fragment as depicted in

Scheme 3 Building blocks applied in the here-presented studies in the synthesis of D-alanine kojibiose functionalized LTA-fragments of *Enterococcus faecalis*.

Scheme 4 Synthesis of donor **15** and **16**. (a) BnBr, NaH, DMF, **20**: 96%. (b) BH₃.THF, Cu(OTf)₂, **21**: 80%; **24**: 76%. (c) NapBr, NaH, DMF, **22**: 91%; **25**: 93%. (d) 1) NIS, acetone: $H_2O = 10:1 \ (v:v)$; (2) 2,2,2-trifluoro-*N*-phenylacetimidoyl chloride, Cs_2CO_3 , acetone, yield **15**: 86%; **16**: 84%. (e) PMBCl, NaH, DMF, 94%.

Scheme 5. First, glycerol building block 19 was assembled. To this end glycerol 17 was glycosylated with donor 16, of which the primary alcohol groups were protected by a benzoate ester and an allyl ether, using the TMSI-Ph3P=O activation conditions to give the compound 26 in good yield and excellent selectivity. Next, the C2-O-PMB was selectively removed using a catalytic amount of HCl in a mixture of hexafluoro-iso-propanol (HFIP) and DCM to give C2-OH acceptor 27 in 95% yield.16 All other benzyl ethers remained unaffected under these conditions. Acceptor 27 was then glycosylated with donor 15 using the DMF-mediated glycosylation conditions to form the disaccharide 28 with good selectivity and excellent yield. To set the stage for the introduction of the p-alanyl esters, first the glycerol benzoate was replaced by a TBDPS ether, after which the two C-6-O-Nap ethers were removed to give diol 31. The introduction of the two D-alanine esters was accomplished using PyBOP as a condensing agent to form 6,6-di-alanyl-α-kojibiose 32 in 90% yield. To transform this building block into a synthon suitable for the assembly of TAfragments, the allyl ether was replaced by a 4,4'-dimethoxytrityl (DMTr) group. Therefore, the allyl group was isomerized to the corresponding enol ether using an iridium catalyst, which was subsequently hydrolyzed using I2 in combination with sodium bicarbonate. The liberated primary alcohol was treated with DMTrCl in DCM to give compound 34, which was treated with HF-pyridine to give the 6,6-di-alanyl- α -kojibiose **19** in 86% yield. For the construction of the kojibiobiosyl glycerol trimer 40, we chose to use benzyl phosphoramidites, as opposed to the more commonly employed cyanoethyl reagents.10 The latter group however requires basic conditions for its removal, that may jeopardize the labile p-alanine esters in the target compound. Thus, compound 19 and phosporamidite 35 were coupled under the agency of dicyanoimidazole (DCI), to provide the intermediate phosphite triester, which was immediately oxidized to the phosphate using (1S)-(+)-(10-camphorsulfonyl)-oxaziridine (CSO), and the resulting dimer 36 was treated with a catalytic amount of p-toluenesulfonic acid (TsOH) in DCM/MeOH to provide compound 37 in 93% over three steps. A similar coupling of phophoramidite 38 and alcohol 37 then provided

Scheme 5 Synthesis of a di-alanyl-α-kojibiose containing LTA fragment. (a) TMSI, Ph₃P=O, DCM, 36 h, 76%, α : β > 20:1. (b) TES, HCl/HFIP, DCM: HFIP = 1:1 (v:v), 93%. (c) DMF, TfOH, 24 h, 95%, α : β > 10:1. (d) CH₃ONa, DCM: CH₃OH = 1:1 (v:v), 92%. (e) TBDPSCI, imidazole, DMF, 99%. (f) DDQ, DCM: H₂O = 10:1 (v:v), 77%. (g) PyBOP, NMI, DCM, 90%. (h) (1) Ir(COD)(Ph₂MeP)₂PF₆: (2) I₂, NaHCO₃, H₂, THF, 85%. (i) DMTrCI, DIPEA, DCM, 95%. (j) HF-pyridine, THF/pyridine, 86%. (k) 1) DCI, CH₃CN; (2) CSO. (l) TsOH, DCM/CH₃OH = 1:1 (v:v), 93%. (m) DCI, CH₃CN; (2) CSO, 88%. (n) H₂, Pd/C, dioxane/H₂O = 2:1 (v:v), 3 day, 95%.

trimer 39 in 88% yield. Removal of all benzyl ethers, benzyl carbamates and benzyl phosphotriesters in 39 in a single hydrogenolysis event delivered 40.

Conclusion

In conclusion, we have here explored additive controlled glycosylations for the construction of α -1,2-glucosyl linkages. After the successful construction of α -1,3-glucosyl and α -1,4-glucosyl linkages, described previously, we have shown that conditions, involving the activation of per-benzylated glucosyl imidate donors with stoichiometric TfOH in the presence of an excess DMF, can be profitably used for the construction of α -1,2-glucosyl linkages. The assembly of a linear kojitetraose however has brought to light that the stereoselectivity of the glycosylation reactions decrease with growing chain length. A kojibiose was assembled with excellent stereoselectivity ($\alpha:\beta = 15:1$) and elongation of this disaccharide proceeded with good selectivity (α : β = 10:1), but extension of the trisaccharide with a fourth residue led to an anomeric mixture ($\alpha:\beta=2:1$). Using the methodology and building on a triad of benzyl-type protecting groups, a p-alanine-kojibiose functionalized LTA from the E. faecalis cell wall was synthesized. Through the use of mild global deprotection conditions the labile p-alanine esters were maintained in the final product. The generation of this structure will enable binding studies with antibodies, which may pave the way to the generation of a synthetic vaccine, based on this structure, directed at this important nosocomial pathogen.

Experimental

General experimental procedures

All reagents were of commercial grade and used as received. All moisture sensitive reactions were performed under an argon atmosphere. DCM used in the glycosylation reactions was dried with flamed 4 Å molecular sieves before being used. Reactions were monitored by TLC analysis with detection by UV (254 nm) and where applicable by spraying with 20% sulfuric acid in EtOH or with a solution of (NH₄)₆Mo₇O₂₄·4H₂O (25 g L^{-1}) and $(NH_4)_4Ce(SO_4)_4 \cdot 2H_2O$ (10 g L^{-1}) in 10% sulfuric acid (aq.) followed by charring at ~150 °C. Column chromatography was carried out using silica gel (0.040-0.063 mm). Sizeexclusion chromatography was carried out using Sephadex LH-20.¹H and ¹³C spectra were recorded on a Bruker AV 400 and Bruker AV 500 in CDCl₃ or D₂O. Chemical shifts (δ) are given in ppm relative to tetramethylsilane as internal standard (1H NMR in CDCl₃) or the residual signal of the deuterated solvent. Coupling constants (1) are given in Hz. All ¹³C spectra are proton decoupled. NMR peak assignments were made using COSY and HSQC experiments, where applicable Clean TOCSY, HMBC and GATED experiments were used to further elucidate the structure. The anomeric product ratios were analysed through integration of proton NMR signals.

Experimental procedures and characterization data of products

For the synthesis procedure and data of known compounds 13, 13 17, 18 22, 19 23 20 see references. We used "a", "b", "c" and "d" to specify the H-1 and C-13 NMR signals of sugar rings

from sp.

St.

mac (1 cu - 7) th sh

from the "reducing" to the "non-reducing" end and "o" to specify the H-1 and C-13 NMR signals of the spacer.

Standard procedures

Communication

Procedure A for the glycosylation of secondary alcohols. A mixture of donor (1.0 eq.), acceptor (0.7 eq.) (donors and acceptors co-evaporated with toluene three times), DMF (16 eq.) in dry DCM were stirred over fresh flame-dried molecular sieves 3A under nitrogen. The solution was cooled to -78 °C, after which TfOH (1.0 eq.) was added. After 30 min, the reaction was stirred at 0 or -10 °C until TLC-analysis showed complete conversion of the acceptor. The reaction was quenched with Et₃N, filtered and concentrated *in vacuo*. The products were purified by size exclusion and silica gel column chromatography.

Procedure B for the glycosylation of primary alcohols. A mixture of donor (1.0 eq.), acceptor (0.7 eq.) (donor and acceptor co-evaporated with toluene three times), $Ph_3P = O$ (6 eq.) in dry DCM were stirred over fresh flame-dried molecular sieves 3A under nitrogen. Then TMSI (1.0 eq.) was added slowly in the mixture. The reaction was stirred at room temperature until TLC-analysis indicated the reaction to be complete. The solution was diluted and the reaction quenched with saturated $Na_2S_2O_3$. The organic phase was washed with water and brine, dried with anhydrous $MgSO_4$, filtered and concentrated *in vacuo*. The products were purified by size exclusion and silica gel column chromatography.

Procedure C for deprotection of the PMB protecting group. The starting material (1 eq.) was dissolved in DCM: HFIP (1:1, 0.1 M). TES (1.0 eq.) and 0.1 M HCl/HFIP (0.1 eq.) were added to the mixture. The reaction stirred until TLC-analysis indicated full consumption of the starting material (15 min-2 h). Then the mixture was diluted with DCM and the reaction quenched with saturated NaHCO₃. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated *in vacuo*. The product was purified by silica gel column chromatography.

Procedure D for deprotection of the Nap protecting group. The starting material (1 eq.) was dissolved in DCM: H_2O (10:1, 0.1 M). DDQ (1.1 eq.) was added to the mixture. The reaction stirred until TLC-analysis indicated full consumption of the starting material (± 2 h). Then the mixture was diluted with DCM and the reaction quenched with saturated $Na_2S_2O_3$. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated *in vacuo*. The product was purified by silica gel column chromatography.

Phenyl 4,6-O-benzelidine-2-O-(naphthalen-2-ylmethyl)-3-O-benzyl-1-thio-β-p-glucopyranoside 3. Compound 1 (7.7 g, 21.4 mmol) and Bu_2SnO (5.9 g, 23.7 mmol) was dissolved in dry toluene. The mixture was refluxed at 120 °C for 3 h. Then CsF (3.6 g, 23.7 mmol) and BnBr (2.8 mL, 23.7 mmol) were added in the mixture. The mixture was refluxed at 120 °C until the reaction was completed by TLC monitoring. Then the mixture was dilutted with EtOAc and washed with 1 mol L^{-1} KF and brine. The organic phase was dried with anhydrous

MgSO₄, filtered and concentrated in vacuo, and the product purified by column chromatography. Compound 2 was obtained with 76% yield. Then compound 2 (7.3 g, 16.2 mmol) was dissolved in DMF at 0 °C. Then NaH (1.95 g, 48.6 mmol) and NapBr (5 g, 20.6 mmol) were added the mixture under N2. The reaction was stirring at 0 °C untill the reaction was completed by TLC monitoring. Then the mixture was dilutted with EtOAc and washed with H2O and brine. The organic phase was dried with anhydrous MgSO4, filtered and concentrated in vacuo, and the product purified by column chromatography. Compound 3 was obtained with 91% yield. ¹H-NMR (CDCl₃, 400 MHz) δ 7.80–7.77 (m, 4 H, aromatic H), 7.55–7.22 (m, 18 H, aromatic H), 5.54 (s, 1 H), 5.02-4.94 (m, 3 H, 3 CHH), 4.79-4.76 (m, 2 H, H-1, 1 CHH), 4.35 (dd, J_1 = 4.8 Hz, J_2 = 10.4 Hz, 1 H, H-6_a), 3.86-3.66 (m, 3 H, H-6, H-4, H-3), 3.56 (t, $J = 8.4 \text{ Hz}, 1 \text{ H}, \text{H-2}, 3.47-3.41 (m, 1 \text{ H}, \text{H-5}).^{13}\text{C-APT (CDCl}_3,$ 100 MHz) δ 138.34, 137.29, 135.56, 133.29, 133.18, 133.08 (aromatic C), 132.31, 129.05, 128.99, 128.41, 128.28, 128.12, 128.99, 128.87, 127.78, 127.73, 126.90, 126.26, 126.07, 126.03, 125.94 (aromatic CH), 101.08 (PhCH), 88.27 (C-1), 82.99 (C-3), 81.45 (C-4), 80.48 (C-2), 75.92, 75.28 (CH₂), 70.22 (C-5), 68.67 (C-6).

Phenyl 3,4,6-tri-O-benzyl-2-O-(naphthalen-2-ylmethyl)-1-thioβ-D-glucopyranoside 5. Compound 3 (8.7 g, 14.7 mmol) was dissolved in dry DCM under N2 at 0 °C. Then TES (23 mL, 144 mmol) and BF₃·Et₂O (3.7 mL, 29 mmol) were added in the solution. The reaction was stirring at t = 0 °C untill the reaction was completed by TLC monitoring. Then the mixture was dilutted with EtOAc and washed with H2O and brine. The organic phase was dried with anhydrous MgSO₄, filtered and concentrated in vacuo. Crude compound 4 was dissolved in DMF at 0 °C. Then NaH (1.8 g, 45 mmol) and BnBr (2.6 mL, 21.9 mmol) were added the mixture under N2. The reaction was stirring at 0 °C untill the reaction was completed by TLC monitoring. Then the mixture was dilutted with EtOAc and washed with H₂O and brine. The organic phase was dried with anhydrous MgSO4, filtered and concentrated in vacuo, and the product purified by column chromatography. Compound 5 was obtained with 88% yield. 1 H-NMR (CDCl₃, 400 MHz) δ 7.81-7.75 (m, 4 H, aromatic H), 7.62-7.18 (m, 23 H, aromatic H), 5.04 (d, J = 10.4 Hz, 1 H, 1 CHH), 4.93-4.82 (m, 4 H, 4 CHH), 4.71 (d, J = 9.6 Hz, 1 H, H-1), 4.62-4.52 (m, 3 H, 3 CHH), 3.81-3.65 (m, 4 H, H-6, H-4, H-3), 3.60-3.50 (m, 2 H, H-2, H-5). 13 C-APT (CDCl₃, 100 MHz) δ 138.49, 138.37, 138.11, 135.62, 133.94, 133.37, 133.16 (aromatic C), 132.01, 129.02, 128.55, 128.45, 128.25, 128.07, 128.04, 127.92, 127.85, 127.79, 127.76, 127.67, 127.54, 127.00, 126.34, 126.12, 126.00 (aromatic CH), 87.57 (C-1), 86.86 (C-3), 80.92 (C-2), 79.18 (C-4), 77.89 (C-5), 75.93, 75.59, 75.16, 73.50 (CH₂), 69.08 (C-6).

3,4,6-Tri-O-benzyl-2-O-(naphthalen-2-ylmethyl)- α/β -D-glucopyranosyl N-phenyltrifluoroacetimidate 6. Compound 5 (11.5 g, 16.8 mmol) was dissolved in acetone (200 mL). N-Iodosuccinimide (NIS, 7.5 g, 33.6 mmol) was added in one portion and the reaction was stirred at room temperature for 2 hours. The solution was diluted with DCM and the reaction was quenched with saturated aqueous Na₂S₂O₃. Then the

organic layer was washed with water and brine. The organic layer was dried with anhydrous MgSO₄, filtered and concentrated in vacuo, and the product purified by column chromatography. The lactol (8.7 g, 90% yield) was obtained as a white solid. Next, the lactol (8.7 g, 15.1 mmol) was dissolved in acetone (150 mL). Cs₂CO₃ (7.4 g, 22.6 mmol) and 2,2,2-trifluoro-N-phenylacetimidoyl chloride (3.5 mL, 25.1 mmol) were added to the solution respectively. The reaction was stirred overnight, then quenched with Et₃N, filtered and concentrated in vacuo. The product was purified by column chromatography (pentane: EA = 40:1-20:1). Compound 6 (9.6 g, 92% yield, $\alpha:\beta=1:1$) was obtained as yellow syrup. ¹H-NMR (CDCl₃, 500 MHz, 60 °C) δ 7.79–7.74 (m, aromatic H), 7.49–7.01 (m, aromatic H), 6.83-6.73 (m, aromatic H), 6.40 (bs, 1 H, H-1 α), 5.58 (bs, 1 H, H-1β), 4.96-4.53 (m), 4.02-3.95 (m), 3.79-3.59 (m), 3.40 (bs, 1 H). 13 C-APT (CDCl₃, 125 MHz, 60 °C) δ 159.77, 159.73, 143.96, 143.73, 143.42, 138.99, 138.77, 138.38, 138.31, 135.81, 135.70, 133.57, 133.32, 130.33, 130.26 (aromatic C), 129.86, 129.43, 128.80, 128.46, 128.43, 128.33, 128.28, 128.06, 127.98, 127.94, 127.91, 127.84, 127.78, 127.69, 127.65, 127.60, 126.78, 126.72, 126.22, 126.20, 126.01, 125.97, 124.41, 124.28, 119.69, 119.58, 114.20, 114.17 (aromatic CH), 117.62 (q, CF₃), 97.71, 94.00, 84.78, 81.75, 80.86, 79.41, 77.61, 77.36, 76.06, 75.31, 75.63, 75.55, 75.27, 75.02, 74.72, 73.82, 73.73, 73.48, 73.24, 68.73, 68.66, 55.83, 55.36. HR-MS: calculated for $C_{46}H_{42}F_3NO_6 [M - [O(C=NPh)CF_3] + OH + NH_4]^{+}$: 608.30066, found: 608.29853.

Synthesis of 7. The reaction was carried out according to the standard procedure B, using 6 (2000 mg, 2.6 mmol, 0.1 M in DCM), 3-aminopropanol (370 μ L, 3.9 mmol), Ph₃P=O (4.4 g, 15.8 mmol) and TMSI (371 µL, 2.6 mmol). The product was purified by silica gel column chromatography (pentane: EA = 10:1). Compound 7 (1300 mg, 75% yield, $\alpha:\beta = 10:1$) was obtained as a colourless syrup. IR (neat, cm $^{-1}$) ν 697, 735, 818, 1027, 1155, 1358, 1454, 2095, 2866, 2922. ¹H-NMR (CDCl₃, 400 MHz) δ 7.83–7.74 (m, 4 H, aromatic H), 7.49–7.12 (m, 18 H, aromatic H), 5.02 (d, J = 11.2 Hz, 1 H, 1 CHH), 4.92 (d, J =12.4 Hz, 1 H, 1 CHH), 4.86-4.77 (m, 3 H, 3 CHH), 4.74 (d, J = 3.6 Hz, 1 H, H-1), 4.59 (d, J = 11.6 Hz, 1 H, 1 CHH), 4.48-4.41 (m, 2 H, 2 CHH), 3.99 (t, J = 9.2 Hz, 1 H, H-2), 3.75-3.59 (m, 6)H, H-6, H-5, H-4, H-2, H- 1_a°), 3.48–3.35 (m, 3 H, H- 3° , H- 1_b°), 1.97–1.81 (m, 2 H, H-2°). ¹³C-APT (CDCl₃, 100 MHz) δ 138.93, 138.25, 138.14, 127.98, 135.72, 133.29, 133.17 (aromatic C), 128.52, 128.49, 128.46, 128.40, 128.06, 128.00, 127.84, 127.79, 127.70, 127.03, 126.29, 126.13, 126.06 (aromatic CH), 97.31 (C-1), 82.10 (C-3), 80.10 (C-2), 77.75 (C-4), 75.81, 75.24, 73.57, 73.53 (CH₂), 70.43 (C-5), 68.50 (C-6), 64.83 (C-1°), 48.41 (C-3°), 28.96 (C-2°). HR-MS: calculated for $C_{41}H_{43}N_3O_6 [M + NH_4]^+$: 691.34901, found: 691.34807.

Synthesis of 8. The reaction was carried out according to the general procedure D, using 7 (1.3 g, 1.93 mmol, 0.1 M in DCM: H₂O) and DDQ (482 mg, 2.1 mmol). The product was purified by silica gel column chromatography. Compound 8 (770 mg, 75% yield) was obtained as a colourless syrup. $[\alpha]_{D}^{20}$ +90.5 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 697, 735, 1027, 1066, 1038, 1359, 1454, 1497, 2095, 2870, 2923. ¹H-NMR (CDCl₃,

400 MHz) δ 7.39–7.13 (m, 15 H, aromatic H), 4.94–4.80 (m, 4 H, H-1, 3 CHH), 4.63 (d, J = 12.4 Hz, 1 H, 1 CHH), 4.52-4.48 (m, 2 H, 2 CHH), 3.85-3.61 (m, 7 H, H-6, H-5, H-4, H-3, H-2, $H-3_{a}^{\circ}$), 3.56–3.51 (m, 1 H, $H-3_{b}^{\circ}$), 3.43–3.32 (m, 2 H, $H-1^{\circ}$), 2.18 (d, J = 7.6 Hz, 1 H, OH), 1.94–1.82 (m, 2 H, H-2°). ¹³C-APT (CDCl₃, 100 MHz) δ 138.69, 138.11, 137.93 (aromatic C), 128.50, 128.47, 128.02, 127.97, 127.96, 127.84, 127.80, 127.77 (aromatic CH), 98.66 (C-1), 83.31 (C-3), 77.46 (C-4), 75.43, 75.11, 73.59 (CH₂), 72.94 (C-2), 70.82 (C-5), 68.47 (C-6), 65.15 (C-1°), 48.53 (C-3°), 28.86 (C-2°). HR-MS: calculated for $C_{30}H_{35}N_3O_6 [M + NH_4]^+$: 551.28641, found: 551.28604.

Synthesis of 9. The reaction was carried out according to the standard procedure B, using 6 (845 mg, 1.11 mmol), 8 (455 mg, 0.85 mmol, 0.05 M in DCM), DMF (1.4 mL, 17.7 mmol) and TfOH (98 µL, 1.11 mmol). The reaction was stirred at -78-0 °C until TLC-analysis showed complete conversion of the acceptor. The reaction was quenched with Et₃N, filtered and concentrated in vacuo. The product was purified by size exclusion chromatography (DCM: MeOH = 1:1). Compound 9 (1160 mg, 95% yield, $\alpha:\beta=15:1$) was obtained as a colourless syrup. $\left[\alpha\right]_{D}^{20}$ +75.4 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 697, 736, 1029, 1069, 1359, 1454, 1497, 2098, 2866, 2922. ¹H-NMR (CDCl₃, 400 MHz) δ 7.83–7.71 (m, 4 H, aromatic H), 7.49-7.45 (m, 3 H, aromatic H), 7.34-7.21 (m, 23 H, aromatic H), 7.16-7.03 (m, 7 H, aromatic H), 5.07 (d, J = 3.2 Hz, 1 H, H-1b), 5.05 (d, J = 3.2 Hz, 1 H, H-1a), 5.01-4.79 (m, 8 H, 8 CHH), 4.63-4.39 (m, 5 H, 5 CHH), 4.31 (d, J = 12.4 Hz, 1 H, 1 CHH), 4.11 (t, J = 91.6 Hz, 1 H, H-3b), 4.06–3.99 (m, 2 H, H-3a, H-5a), 3.83 (dd, J_1 = 3.2 Hz, J_2 = 9.6 Hz, 1 H, H-2a), 3.78-3.61 (m, 7 H, H-6, H-5b, H-4b, H-4a, H-2b, H-3_a°), 3.55–3.39 (m, 3 H, H-6, H- 3_b°), 3.29–3.19 (m, 2 H, H- 1°), 1.84–1.70 (m, 2 H, H-2°). ¹³C-APT (CDCl₃, 100 MHz) δ 138.81, 138.72, 138.32, 138.25, 138.00, 137.98, 135.69, 133.29, 133.11 (aromatic C), 128.49, 128.43, 128.42, 128.39, 128.29, 128.09, 128.03, 128.01, 127.97, 127.90, 127.81, 127.80, 127.71, 127.66, 127.62, 127.52, 126.87, 126.33, 126.12, 125.89 (aromatic CH), 96.06 (C-1a), 94.72 (C-1b), 82.17 (C-3b), 80.79 (C-3a), 79.14 (C-2b), 78.07 (C-4), 77.68 (C-4), 76.18, 75.71 (CH₂), 75.45 (C-2a), 75.20, 74.95, 73.60, 73.45, 72.99 (CH₂), 70.61 (C-5b), 70.44 (C-5a), 68.52 (C-6), 68.04 (C-6), 65.15 (C-1°), 48.36 (C-3°), 28.02 (C-2°). HR-MS: calculated for $C_{68}H_{71}N_3O_6 [M + NH_4]^{\dagger}$: 1123.54269, found: 1123.54104.

Synthesis of 10. The reaction was carried out according to the general procedure D, using 9 (970 mg, 0.88 mmol, 0.1 M in DCM: H₂O) and DDQ (217 mg, 0.96 mmol). The product was purified by silica gel column chromatography. Compound 10 (416 mg, 49% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +90.8 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 697, 735, 1027, 1050, 1132, 1361, 1454, 2098, 2866, 2929. H-NMR (CDCl₃, 400 MHz) δ 7.38-7.06 (m, 30 H, aromatic H), 5.02-4.97 (m, 3 H, H-1a, H-1b, 1 CHH), 4.87-4.75 (m, 5 H, 5 CHH), 4.63-4.44 (m, 5 H, 5 CHH), 4.32 (d, J = 12.0 Hz, 1 H, 1 CHH), 3.94-3.65 (m, 11 H), 3.53-3.33 (m, 5 H, H-6), 1.91-1.84 (m, 2 H, H-2°). ¹³C-APT $(CDCl_3, 100 \text{ MHz}) \delta 138.69, 138.61, 137.97, 137.86, 737.83$ (aromatic C), 128.41, 128.39, 128.34, 128.30, 128.28, 128.16, 128.08, 127.97, 127.95, 127.87, 127.82, 127.78, 127.72, 127.65,

127.60, 127.55, 127.41 (aromatic *CH*), 96.21 (C-1b), 85.58 (C-1a), 83.28 (C-3b), 80.50 (C-3a), 77.79 (C-4), 76.94 (C-4), 76.00 (*CH*₂), 75.37 (C-2a), 75.12, 74.75, 73.48, 73.29 (*CH*₂), 72.79 (C-2b), 70.62 (C-5), 70.58 (C-5), 68.28 (C-6), 67.86 (C-6), 64.83 (C-1°), 48.33 (C-3°), 28.62 (C-2°). HR-MS: calculated for $C_{68}H_{71}N_3O_{11} [M + NH_4]^{\dagger}$: 983.48003, found: 983.47815.

Synthesis of 11. The reaction was carried out according to the standard procedure B, using 6 (547 mg, 0.72 mmol), 10 (347 mg, 0.36 mmol, 0.05 M in DCM), DMF (900 μL, 11.4 mmol) and TfOH (64 µL, 0.72 mmol). The reaction was stirred at -78-0 °C until TLC-analysis showed complete conversion of the acceptor. The reaction was quenched with Et₃N, filtered and concentrated in vacuo. The product was purified by size exclusion chromatography (DCM: MeOH = 1:1). Compound 11 (372 mg, 67% yield, $\alpha:\beta=10:1$) was obtained as a colourless syrup. $\left[\alpha\right]_{D}^{20}$ +96.5 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 697, 735, 1029, 1069, 1359, 1454, 1497, 2098, 2845, 2921. ¹H-NMR (CDCl₃, 400 MHz) δ 7.82 (s, 1 H, aromatic H), 7.71-7.62 (m, 3 H, aromatic H), 7.52-7.50 (m, 1 H, aromatic H), 7.41-7.37 (m, 1H, aromatic H), 7.32-6.96 (m, 46 H, aromatic H), 5.49 (d, J = 3.6 Hz, 1 H, H-1c), 5.41 (d, J = 3.6 Hz, 1 H, H-1b), 5.23 (d, J = 2.0 Hz, 1 H, H-1a), 5.16 (d, J = 12.0 Hz, 1 H, 1 CHH), 4.99-4.34 (m, 18 H, 18 CHH), 44.18-3.52 (m, 21 H), 3.31–3.22 (m, 2 H, H-3°), 1.93–1.82 (m, 2 H, H-2°). ¹³C-APT (CDCl₃, 100 MHz) δ 138.78, 138.76, 138.74, 138.45, 138.26, 138.16, 138.09, 137.98, 137.90, 136.09, 133.35, 132.94 (aromatic C), 128.72, 128.45, 128.44, 128.40, 128.38, 128.29, 128.26, 128.13, 128.06, 128.04, 127.93, 127.79, 127.72, 127.65, 127.60, 127.56, 127.51, 127.46, 126.11, 125.84, 125.77, 125.55 (aromatic CH), 95.84 (C-1c), 92.77 (C-1b), 92.09 (C-1a), 81.96 (C-3c), 81.45 (C-3a), 80.99 (C-3b), 79.97 (C-2c), 77.37 (2 C-4), 77.05 (C-4), 76.08, 75.99, 75.65, 74.94, 74.84 (CH₂), 74.71 (C-2), 74.60 (C-2), 73.58, 73.50, 71.60 (CH₂), 70.94 (C-5), 70.80 (C-5), 70.61 (C-5), 68.61 (C-6), 68.34 (C-6), 68.07 (C-6), 65.16 (C-1°), 48.31 (C-3°), 29.20 (C-2°). HR-MS: calculated for C₉₅H₉₉N₃O₁₆ $[M + NH_4]^+$: 1555.73636, found: 1555.73599.

Synthesis of 12. The reaction was carried out according to the general procedure D, using 11 (700 mg, 0.45 mmol, 0.1 M in DCM: H₂O) and DDQ (140 mg, 0.6 mmol). The product was purified by silica gel column chromatography. Compound 12 (300 mg, 47% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +122.1 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 697, 735, 1027, 1050, 1132, 1361, 1454, 2098, 2866, 2929. H-NMR (CDCl₃, 400 MHz) δ 7.37-6.98 (m, 45 H, aromatic H), 5.18 (d, J = 3.2 Hz, 1 H, H-1b), 5.06 (d, J = 3.2 Hz, 1 H, H-1a), 5.00–4.90 (m, 3 H, H-1a, 2 CHH), 4.84-4.30 (m, 16 H, 16 CHH), 4.01-3.34 (m, 22 H), 2.41 (s, 1 H, OH), 2.06-1.85 (m, 2 H, H-2°). C-APT (CDCl₃, 100 MHz) δ 138.89, 138.85, 138.56, 138.24, 138.22, 138.02, 137.96, 137.95, 137.89 (aromatic C), 128.56, 128.53, 128.51, 128.50, 128.42, 128.38, 128.36, 128.28, 128.08, 128.07, 127.86, 127.82, 127.79, 127.74, 127.67, 127.63, 127.60, 127.50 (aromatic CH), 96.03 (C-1c), 95.04 (C-1a), 91.66 (C-1b), 83.42 (C-3c), 80.90 (C-3a), 80.62 (C-3b), 77.89 (C-4), 77.73 (C-4), 76.84 (C-4), 76.30, 76.25, 75.20, 75.15, 74.94, 74.92 (CH₂), 74.67 (C-2b), 74.00 (C-2a), 73.65, 73.59, 73.45 (CH₂), 72.67 (C-2c), 70.69 (C-5), 70.64 (C-5), 70.57 (C-5), 68.38 (C-6), 68.00 (C-6), 67.97

(C-6), 64.50 (C-1°), 48.51 (C-3°), 28.77 (C-2°). HR-MS: calculated for $C_{84}H_{91}N_3O_{16}$ [M + NH₄]⁺: 1415.67376, found: 1415.67378.

Synthesis of 14. The reaction was carried out according to the standard procedure B, using 13 (190 mg, 0.27 mmol), 12 (70 mg, 0.05 mmol, 0.05 M in DCM), DMF (78 μ L, 0.99 mmol) and TfOH (23 μ L, 0.27 mmol). The reaction was stirred at -78-0 °C until TLC-analysis showed complete conversion of the acceptor. The reaction was quenched with Et₃N, filtered and concentrated *in vacuo*. The product was purified by size exclusion chromatography (DCM:MeOH = 1:1). Compound 14 (64 mg, 67% yield, $\alpha:\beta=2:1$) was obtained as a colourless syrup. IR (neat, cm $^{-1}$) ν 695, 734, 1027, 1069, 1209, 1359, 1454, 1497, 1729, 2096, 2859, 2923. HR-MS: calculated for $C_{118}H_{125}N_3O_{21}$ [M + NH₄] † : 1937.91438, found: 1937.91440.

Synthesis of 15. Compound 22 (1.0 g, 1.46 mmol) was dissolved in acetone: H₂O (10:1, 15 mL). N-Iodosuccinimide (NIS) (660 mg, 2.93 mmol) was added in one portion and the reaction was stirred at room temperature for 2 hours. The solution was diluted with DCM and the reaction was quenched with saturated aqueous Na2S2O3. Then the organic layer was washed with water and brine. The organic layer was dried with anhydrous MgSO₄, filtered and concentrated in vacuo, and the product purified by column chromatography. The lactol was obtained as a white solid. Next, the lactol was dissolved in acetone (15 mL). Cs₂CO₃ (713 mg, 2.19 mmol) and 2,2,2-trifluoro-N-phenylacetimidoyl chloride (355 μL, 2.19 mmol) were added to the solution respectively. The reaction was stirred overnight, then quenched with Et3N, filtered and concentrated in vacuo. The product was purified by column chromatography. Compound 15 (959 mg, 86% yield over two steps, $\alpha:\beta=1:1$) was obtained as yellow syrup. IR (neat, cm⁻¹) ν 695, 734, 1027, 1073, 1153, 1208, 1314, 1454, 1597, 1716, 2869, 2915. ¹H-NMR (CDCl₃, 400 MHz) δ 7.82–6.50 (m, aromatic H), 6.53 (bs, 1 H, H-1), 5.67 (bs, 1 H, H-1), 5.00-4.61 (m), 4.52-4.46 (m), 4.06–3.97 (m), 3.81–3.66 (m). 13 C-APT (CDCl₃, 125 MHz) δ 143.78, 143.55, 138.67, 138.64, 137.92, 137.83, 135.45, 135.28, 133.35, 133.19, 133.16 (aromatic C), 129.51, 128.82, 128.64, 128.61, 128.57, 128.55, 128.49, 128.43, 128.36, 128.33, 128.13, 128.05, 127.99, 127.92, 127.88, 127.84, 127.81, 127.78, 127.00, 126.88, 126.49, 126.30, 126.26, 126.10, 126.08, 126.05, 124.39, 120.57, 119.44 (aromatic CH), 84.64, 81.62, 81.04, 79.37, 77.27, 76.94, 75.91, 75.82, 75.74, 75.44, 75.31, 75.17, 73.76, 73.65, 73.44, 73.19, 68.12.

Synthesis of 16. Compound **23** (2.57 g, 4.5 mmol) was dissolved in dry DCM under N₂ at 0 °C. Then TES (7.2 mL, 45 mmol) and BF₃·Et₂O (1.2 mL, 9 mmol) were added in the solution. The reaction was stirring at t 0 °C untill the reaction was completed by TLC monitoring. Then the mixture was dilutted with EtOAc and washed with H₂O and brine. The organic phase was dried with anhydrous MgSO₄, filtered and concentrated *in vacuo*. The product purified by column chromatography. Compound **24** was dissolved in DMF at 0 °C. Then NaH (540 g, 13.05 mmol) and NapBr (1.4 g, 6.7 mmol) were added the mixture under N₂. The reaction was stirring at 0 °C untill the reaction was completed by TLC monitoring. Then the mixture was dilutted with EtOAc and washed with H₂O and

brine. The organic phase was dried with anhydrous MgSO₄, filtered and concentrated in vacuo, and the product purified by column chromatography. Compound 25 was obtained with 83% yield over two steps. Compound 25 (4.8 g, 6.73 mmol) dissolved in acetone: H₂O (10:1,77 N-Iodosuccinimide (NIS) (3.0 g, 13.3 mmol) was added in one portion and the reaction was stirred at room temperature for 2 hours. The solution was diluted with DCM and the reaction was quenched with saturated aqueous Na₂S₂O₃. Then the organic layer was washed with water and brine. The organic layer was dried with anhydrous MgSO4, filtered and concentrated in vacuo, and the product purified by column chromatography. The lactol was obtained as a white solid. Next, the lactol was dissolved in acetone (70 mL). Cs₂CO₃ (6.3 g, 19.5 mmol) and 2,2,2-trifluoro-N-phenylacetimidoyl chloride (3.2 mL, 19.5 mmol) were added to the solution respectively. The reaction was stirred overnight, then quenched with Et₃N, filtered and concentrated in vacuo. The product was purified by column chromatography. Compound 16 (4.48 g, 84% yield over two steps, $\alpha:\beta=1:1$) was obtained as yellow syrup. ¹H-NMR (CDCl₃, 500 MHz, 60 °C) δ 7.79–7.74 (m, aromatic *H*), 7.49–7.01 (m, aromatic H), 6.83–6.73 (m, aromatic H), 6.40 (bs, 1 H, H-1 α), 5.58 (bs, 1 H, H-1 β), 4.96–4.53 (m), 4.02–3.95 (m), 3.79–3.59 (m), 3.40 (bs, 1 H). ¹³C-APT (CDCl₃, 125 MHz, 60 °C) δ 159.77, 159.73, 143.96, 143.73, 143.42, 138.99, 138.77, 138.38, 138.31, 135.81, 135.70, 133.57, 133.32, 130.33, 130.26 (aromatic C), 129.86, 129.43, 128.80, 128.46, 128.43, 128.33, 128.28, 128.06, 127.98, 127.94, 127.91, 127.84, 127.78, 127.69, 127.65, 127.60, 126.78, 126.72, 126.22, 126.20, 126.01, 125.97, 124.41, 124.28, 119.69, 119.58, 114.20, 114.17 (aromatic CH), 117.62 (q, CF₃), 97.71, 94.00, 84.78, 81.75, 80.86, 79.41, 77.61, 77.36, 76.06, 75.31, 75.63, 75.55, 75.27, 75.02, 74.72, 73.82, 73.73, 73.48, 73.24, 68.73, 68.66, 55.83, 55.36. HR-MS: calculated for $C_{47}H_{44}F_3NO_7$ [M - [O(C=NPh)CF₃] + OH + Na]⁺: 643.26662, found: 643.26659.

(*S*)-3-(Allyloxy)-2-hydroxypropyl benzoate 17. 1 H-NMR (CDCl₃, 400 MHz) δ 8.07–8.04 (m, 2 H, aromatic *H*), 7.58–7.53 (m, 1 H, aromatic *H*), 7.45–7.41 (m, 2 H, aromatic *H*), 5.95–5.85 (m, 1 H, H-5), 5.31–5.17 (m, 2 H, H-6), 4.45–4.36 (m, 2 H, H-1), 4.19–4.14 (m, 1 H, H-2), 4.05–4.03 (m, 2 H, H-4), 3.63–3.53 (m, 2 H, H-3), 3.01 (bs, 1 H, O*H*). 13 C-APT (CDCl₃, 100 MHz) δ 166.69 (*C*=O), 134.30, 133.18 (aromatic *CH*), 129.87 (aromatic *C*), 129.80 (aromatic *CH*), 128.43 (C-5), 117.54 (C-6), 772.43, 71.02, 68.93 (C-2), 66.07. HR-MS: calculated for $C_{13}H_{16}O_4$ [M + Na]⁺: 259.0941, found: 259.0948.

Synthesis of 26. The reaction was carried out according to the standard procedure B, using **15** (1.5 g, 1.9 mmol), **17** (410 mg, 1.74 mmol, 0.1 M in DCM), Ph₃P=O (3.16 g, 11.4 mmol) and TMSI (270 μL, 1.9 mmol). The product was purified by silica gel column chromatography. Compound **26** (1100 mg, 76% yield, $\alpha: \beta > 20: 1$) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +38.9 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 713, 820, 1037, 1072, 1095, 1249, 1272, 1452, 1513, 1720, 2865, 2912. ¹H-NMR (CDCl₃, 400 MHz) δ 7.99–7.97 (m, 2 H, aromatic H), 7.76–7.63 (m, 4 H, aromatic H), 7.46–7.08 (m, 16 H, aromatic H), 6.92 (bd, 2 H, aromatic H), 6.85–6.82 (m, 2 H,

matic H), 5.93-5.84 (m, 1 H, H-5°), 5.31-5.25 (m, 1 H, H-6_a°), 5.19-5.16 (m, 2 H, H-1a, H-6_b°), 4.99 (d, J = 10.8 Hz, 1 H, CHH), 4.81-4.77 (m, 2 H, 2 CHH), 4.70-4.63 (m, 3 H, 3 CHH), 4.57 (dd, $J_1 = 3.6$ Hz, $J_2 = 11.6$ Hz, 1 H, H-1_a°), 4.45-4.35 (m, 3 H, 2 CHH, H-1_b°), 4.28-4.24 (m, 1 H, H-2°), 4.04-3.99 (m, 4 H, H-3a, H-5a, H-4°), 3.74 (s, 3 H, OCH₃), 3.73-3.58 (m, 4 H, H-2a, H-4a, H-3°), 3.50 (dd, 1 H, J_1 = 2.8 Hz, J_2 = 11.8 Hz, H-6a_a), 3.37 (dd, 1 H, J_1 = 2.0 Hz, J_2 = 10.4 Hz, H-6a_b). ¹³C-APT (CDCl₃, 100 MHz) δ 166.19 (C=O), 159.30, 138.91, 138.22, 135.16 (aromatic C), 134.38 (C-5°), 133.11, 132.96 (aromatic C), 132.88 (aromatic CH), 130.27, 129.84 (aromatic C), 129.62, 129.60, 128.31, 128.28, 128.11, 127.85, 127.82, 127.67, 127.48, 127.44, 126.80, 126.03, 125.97, 125.83 (aromatic CH), 117.19 (C-6°), 113.78 (aromatic CH), 96.13 (C-1a), 81.83 (C-5a), 79.26 (C-2a), 77.47 (C-4a), 75.59, 74.83 (CH₂), 73.59 (C-2°), 73.52, 72.43 (CH_2) , 72.35 (C-4°), 70.23 (C-3a), 69.59 (C-3°), 67.97 (C-6a), 65.01 (C-1°), 55.16 (OCH₃). HR-MS: calculated for $C_{52}H_{54}O_{10}$ $[M + Na]^+$: 861.3609, found: 861.3647.

Synthesis of 27. The reaction was carried out according to the general procedure C, using 26 (1000 mg, 1.19 mmol, 0.1 M in DCM: HFIP), triethylsilane (190 µL, 1.19 mmol) and 0.1 M HCl/HFIP (1.2 ml, 0.12 mmol). The product was purified by silica gel column chromatography (PE: EA = 6:1). Compound 27 (800 mg, 93% yield) was obtained as a colourless syrup. $[\alpha]_{D}^{20}$ +73.0 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 710, 713, 738, 1027, 1070, 1095, 1272, 1452, 1721, 2866, 2915. ¹H-NMR (CDCl₃, 400 MHz) δ 8.02–8.00 (m, 2 H, aromatic H), 7.78–7.66 (m, 4 H, aromatic H), 7.49-7.09 (m, 14 H, aromatic H), 6.96-6.94 (m, 4 H, aromatic H), 5.93-5.84 (m, 1 H, H-5°), 5.32-5.26 (m, 1 H, $H-6^{\circ}_{a}$, 5.23-5.19 (m, 1 H, $H-6^{\circ}_{b}$), 5.12 (d, J = 3.2 Hz, 1 H, H-1a), 4.99 (d, J = 11.2 Hz, 1 H, 1 CHH), 4.81 (d, J = 11.2 Hz, 1 H, 1 CHH), 4.78 (d, J = 11.2 Hz, 1 H, 1 CHH), 4.69 (d, J = 12.4Hz, 1 H, 1 CHH), 4.49-4.38 (m, 4 H, 3 CHH, H-1°),4.24-4.18 (m, 1 H, H-2°), 4.07-3.98 (m, 2 H, H-4°), 3.97-3.93 (m, 1 H, H-3a), 3.80-3.73 (m, 2 H, H-2a, H-5a), 3.68-3.57 (m, 4 H, H-3°, H-4a, H-6a_a), 3.42 (dd, 1 H, $J_1 = 2.0$ Hz, $J_2 = 10.4$ Hz, H-6a_b), 2.77 (d, 1 H, J = 8.0 Hz, OH). ¹³C-APT (CDCl₃, 100 MHz) δ 166.29 (C = O), 138.93, 138.25, 135.30 (aromatic C), 134.03 (C-5°), 133.24 (aromatic C), 133.18 (aromatic CH), 133.07, 129.83 (aromatic C), 129.75, 128.47, 128.41, 128.26, 127.99, 127.94, 127.80, 127.77, 127.61, 127.59, 126.86, 126.16, 126.05, 125.95 (aromatic CH), 117.92 (C-6°), 99.40 (C-1a), 83.53 (C-5a), 77.02 (C-4a), 76.14 (C-2°), 75.33, 74.95, 73.65 (CH_2), 73.44 (C-2a), 72.46 (C-4°), 71.08 (C-3a), 69.23 (C-3°), 68.11 (C-6a), 64.80 (C-1°). HR-MS: calculated for $C_{44}H_{46}O_9$ [M + Na]⁺: 741.3034, found: 741.3062.

Synthesis of 28. The reaction was carried out according to the standard procedure B, using **15** (1210 mg, 1.59 mmol), **27** (760 mg, 1.05 mmol, 0.1 M in DCM), DMF (1.3 mL, 16.8 mmol) and TfOH (140 μ L, 1.59 mmol). The reaction was stirred at -78-0 °C until TLC-analysis showed complete conversion of the acceptor. The reaction was quenched with Et₃N, filtered and concentrated *in vacuo*. The product was purified by size exclusion chromatography (DCM: MeOH = 1:1). Compound **28** (2.06 g, 95% yield, $\alpha: \beta > 10:1$) was obtained as a colourless syrup. Then the compound was purified by silica

gel column chromatography to get pure α product. $[\alpha]_D^{20}$ +75.1 $(c = 1, CHCl_3)$. IR (neat, cm⁻¹) ν 698, 713, 820, 1046, 1070, 1095, 1272, 1359, 1454, 1721, 2863, 2919. H-NMR (CDCl₃, 400 MHz) δ 8.00–7.81 (m, 2 H, aromatic H), 7.78–7.63 (m, 8 H, aromatic H), 7.44-6.91 (m, 34 H, aromatic H), 5.81-5.76 (m, 1 H, H-5°), 5.46 (d, I = 3.6 Hz, 1 H, H-1a), 5.29 (d, I = 3.6 Hz, 1 H, H-1b), 5.22-5.10 (m, 2 H, H-6°), 4.98-4.67 (m, 10 H, 10 CHH), 4.56-4.33 (m, 6 H, 4 CHH, H-1°), 4.29-4.23 (m, 1 H, H-2°), 4.14-4.04 (m, 4 H, H-5a, H-5b, H-3a, H-3b), 3.91-3.88 (m, 3 H, H-3a, H-4°), 3.80-3.43 (m, 9 H, H-3°, H-2b, H-4a, H-4b, H-6a, H-6b). ¹³C-APT (CDCl₃, 100 MHz) δ 166.23 (C=O), 138.69, 138.46, 138.34, 138.31, 138.27, 135.33, 135.22 (aromatic C), 134.27 (C-5°), 133.18, 133.15, 133.01, 132.99 (aromatic C), 132.91 (aromatic CH), 129.92 (aromatic C), 129.62, 128.41, 128.38, 128.35, 128.31, 128.19, 128.17, 128.15, 128.09, 127.92, 127.87, 127.74, 127.69, 127.63, 127.54, 127.51, 127.46, 127.43, 127.34, 126.88, 126.80, 126.09, 126.06, 125.98, 125.85, 125.84 (aromatic CH), 117.41 (C-6°), 99.70 (C-1b), 94.53 (C-1a), 82.10 (C-5b), 80.73 (C-5a), 79.35 (C-2b), 77.80 (C-4), 77.56 (C-4), 76.03, 75.54 (CH₂), 75.39 (C-2a), 74.85, 74.82, 73.54, 73.53 (CH₂), 73.14 (C-2°), 72.36, 72.22 (CH₂), 70.46 (C-3a and 3b), 69.23 (C-3°), 68.98 (C-6a and 6b), 64.96 (C-1°). HR-MS: calculated for $C_{82}H_{82}O_{14} [M + Na]^+$: 1313.5597, found: 1313.5624.

Synthesis of 29. Compound 28 (1100 mg, 0.85 mmol) was dissolved in DCM: CH₃OH (1:1/v:v, 8.5 mL) stirring at room temperature. Then 5 drops of solution of CH₃ONa in CH₃OH (5.4 M) was added in the mixture. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial (3 h). Then the mixture was diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO4, filtered and concentrated in vacuo. The product was purified by silica gel column chromatography. Compound 29 (930 mg, 92% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +69.4 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 737, 818, 1069, 1209, 1359, 1454, 2865, 2921. H-NMR (CDCl₃, 400 MHz) δ 7.83–7.69 (m, 8 H, aromatic H), 7.49–7.09 (m, 25 H, aromatic H), 7.03-6.92 (m, 6 H, aromatic H), 5.86-5.76 (m, 1 H, H-5°), 5.31 (d, J = 3.6 Hz, 1 H, H-1b), 5.22–5.10 (m, 3 H, H-6°, H-1a), 4.94-4.35 (m, 14 H, 14 CHH), 4.09-3.98 (m, 4 H, H-5a, H-5b, H-3a, H-3b), 3.89–3.45 (m, 14 H). ¹³C-APT (CDCl₃, 100 MHz) δ 138.73, 138.52, 138.31, 138.21, 137.98135.41, 135.20 (aromatic C), 134.56 (C-5°), 133.33, 133.29, 133.18, 133.12 (aromatic C), 128.60, 128.45, 128.42, 128.31, 128.22, 128.07, 128.04, 128.01, 127.95, 127.84, 127.81, 127.78, 127.74, 127.67, 127.64, 127.49, 126.98, 126.26, 126.19, 126.06, 126.04, 125.97 (aromatic CH), 117.25 (C-6°), 95.62 (C-1b), 95.08 (C-1a), 82.22 (C-5a), 80.73 (C-5b), 79.24 (C-2b), 78.89 (C-2b), 78.28 (C-4), 77.71 (C-4), 76.15 (CH₂), 75.73 (C-2°), 75.69, 75.16, 74.99, 73.70, 73.61, 72.86 (CH₂), 72.34 (C-4°), 70.86 (C-3b), 70.59 (C-3a), 69.67 (C-3°), 68.64, 68.08 (C-6a and 6b), 63.61 (C-1°). HR-MS: calculated for $C_{75}H_{78}O_{13}$ [M + Na]⁺: 1209.5335, found: 1209.5356.

Synthesis of 30. Compound 29 (350 mg, 0.3 mmol) was dissolved in DMF (6 mL) stirring at room temperature. Then TBDPSCl (153 μ L, 0.6 mmol) and imidazole (120 mg, 1.8 mmol) were added in the mixture. The reaction was stirred

at rt until TLC-analysis showed complete conversion of the starting martial. Then the mixture was diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated in vacuo. The product was purified by silica gel column chromatography. Compound 29 (423 mg, 99% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +62.1 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 738, 818, 1047, 1070, 1361, 1454, 2858, 2928. ¹H-NMR (CDCl₃, 400 MHz) δ 7.80-7.62 (m, 12 H, aromatic H), 7.48-7.06 (m, 31 H, aromatic H), 6.97-6.94 (m, 6 H, aromatic H), 5.87-5.77 (m, 1 H, H-5°), 5.48 (d, J = 3.2 Hz, 1 H, H-1a), 5.34 (d, J = 3.2 Hz, 1 H, H-1b), 5.22-5.10 (m, 2 H, H-6°), 4.95-4.35 (m, 14 H, 14 CHH), 4.11-4.00 (m, 4 H, H-5a, H-5b, H-3a, H-2°), 3.89-3.84 (m, 4 H, H-3b, H-2a, H-4°), 3.78-3.40 (m, 11 H), 1.01 (s, 9 H, 3 CH₃). 13 C-APT (CDCl₃, 100 MHz) δ 138.86, 138.70, 138.58, 138.54, 138.44 (aromatic C), 135.68 (aromatic CH), 135.52 (aromatic C), 134.70 (C-5°), 133.44, 133.43, 133.32, 133.13 (aromatic C), 129.77, 129.76, 128.68, 128.43, 128.40, 128.38, 128.31, 128.23, 128.19, 128.13, 128.05, 128.01, 127.98, 127.86, 127.84, 127.73, 127.60, 127.58, 127.45, 127.41, 126.98, 126.89, 126.23, 126.21, 126.15, 125.98, 125.93 (aromatic CH), 117.33 (C-6°), 94.22 (C-1a), 93.60 (C-1b), 82.16 (C-5b), 81.01 (C-5a), 79.44 (C-2b), 77.76 (C-4), 77.57 (C-4), 76.29, 75.67 (CH₂), 75.12 (C-2°), 74.94, 74.85 (CH₂), 74.79 (C-2a), 73.70, 72.28, 71.91 (CH_2) , 70.43 (C-3b), 70.34 (C-3a), 69.29, 68.22, 63.77 (CH_2), 27.00 (3 CH₃), 19.30. HR-MS: calculated for C₉₁H₉₆O₁₃Si $[M + Na]^+$: 1447.6512, found: 1447.6539.

Synthesis of 31. The reaction was carried out according to the general procedure D, using 30 (460 mg, 0.32 mmol, 0.1 M in DCM: H₂O) and DDQ (170 mg, 0.75 mmol). The product was purified by silica gel column chromatography. Compound 31 (280 mg, 77% yield, pentane: EA = 4:1, R_f = 0.33) was obtained as a colourless syrup. $\left[\alpha\right]_{D}^{20}$ +72.3 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 737, 1027, 1072, 1209, 1361, 1454, 2858, 2929. 1 H-NMR (CDCl₃, 400 MHz) δ 7.66–7.63 (m, 4 H, aromatic H), 7.41-7.07 (m, 31 H, aromatic H), 5.88-5.78 (m, 1 H, H-5°), 5.44 (d, J = 3.2 Hz, 1 H, H-1a), 5.34 (d, J = 3.6 Hz, 1 H, H-1b), 5.23-5.13 (m, 2 H, H-6°), 4.99-4.58 (m, 10 H, 10 CHH), 4.11-4.00 (m, 3 H, H-5a, H-5b, H-2°), 3.92-3.88 (m, 3 H, H-3b, H-4°), 3.76-3.51 (m, 14 H), 1.76 (s, 1 H, OH), 1.64 (s, 1 H, OH), 1.04 (s, 9 H, 3 C H_3). ¹³C-APT (CDCl₃, 100 MHz) δ 138.69, 138.58, 138.44, 138.38, 138.35 (aromatic C), 135.62, 135.61 (aromatic CH), 134.52 (C-5°), 133.27, 133.24 (aromatic C), 129.81, 128.46, 128.44, 128.41, 128.38, 128.36, 128.05, 127.87, 127.84, 127.77, 127.70, 127.67, 127.62, 127.58 (aromatic CH), 117.28 (C-6°), 94.19 (C-1a), 93.40 (C-1b), 81.80 (C-5b), 80.69 (C-5a), 79.65 (C-2b), 77.31 (C-4), 77.25 (C-4), 76.09, 75.57 (CH₂), 75.04 (C-2°), 74.93 (CH₂), 74.90 (C-2a), 72.24, 71.93 (CH₂), 71.25 (C-3b), 70.94 (C-3a), 69.84, 63.67, 61.68, 61.50 (CH₂), 26.93 (3 CH_3), 19.22. HR-MS: calculated for $C_{69}H_{80}O_{13}Si$ [M + Na]⁺: 1167.5260, found: 1167.5280.

Synthesis of 32. Compound 31 (650 mg, 0.57 mmol) and 18 (633 mg, 2.83 mmol) were dissolved in DCM (6 mL) stirring at room temperature. Then TPyBOP (1.48 g, 2.85 mmol) and NMI (454 μ L, 5.7 mmol) were added in the mixture. The reaction was stirred at rt until TLC-analysis showed complete conver-

sion of the starting martial. Then the mixture was diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO4, filtered and concentrated in vacuo. The product was purified by silica gel column chromatography. Compound 29 (800 mg, 90% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +66.3 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 738, 1029, 1070, 1171, 1208, 1251, 1454, 1498, 1725, 2858, 2931. 1 H-NMR (CDCl₃, 400 MHz) δ 7.67–7.63 (m, 4 H, aromatic H), 7.41-7.06 (m, 41 H, aromatic H), 5.85-5.75 (m, 1 H, H-5°), 5.42-5.34 (m, 3 H, H-1a, H-1b, H-6°), 5.21-4.73 (m, 13 H), 4.59-4.50 (m, 3 H), 4.35-3.45 (m, 22 H), 1.36 (d, J = 6.8 Hz, 1 H, CH_3), 1.25 (d, J = 7.2 Hz, 1 H, CH_3), 1.04 (s, 9 H, 3 CH_3). ¹³C-APT (CDCl₃, 100 MHz) δ 172.78, 172.69, 155.62 (4 C=O), 138.54, 138.32, 138.26, 138.19, 136.33 (aromatic C), 135.62, 135.60 (aromatic CH), 134.46 (C-5°), 133.20 (aromatic C), 129.86, 128.57, 128.46, 128.40, 128.35, 128.19, 128.10, 128.03, 127.88, 127.76, 127.70, 127.66, 127.60 (aromatic CH), 117.39 (C-6°), 96.64 (C-1a), 93.00 (C-1b), 81.86 (C-5b), 80.79 (C-5a), 79.60, 77.36, 76.12, 75.63, 74.96, 74.68, 72.22, 71.84, 69.30, 69.12, 68.71, 66.95, 63.73, 63.66, 63.50, 49.69 (2 CHNH), 29.34 (CH₃), 26.95 (3 CH₃), 19.23, 18.73 (CH₃). HR-MS: calculated for $C_{91}H_{102}N_2O_{19}Si [M + Na]^+$: 1577.67383, found: 1577.67285.

Synthesis of 33. A solution of 32 (600 mg, 0.38 mmol) and freshly activated molecular sieves 3A in freshly distilled THF (4 ml) was stirred under argon for 30 min. After the addition of Ir(COD)(Ph₂MeP)₂PF₆ (33 mg, 10 mol%) the solution turned red and the mixture was purged with H2 until the solution turned colourless again (5-15 seconds). After stirring under argon for 4 h, the solution was diluted with THF and satd aq NaHCO₃. After the addition of I₂ (73 mg, 0.57 mmol), the mixture was allowed to stir overnight at room temperature. The mixture was diluted with EtOAc and washed with satd aq NaS2O3 and brine, respectively. The organic layer was dried over MgSO4 and concentrated in vacuo. Column chromatography afforded 33 (734 mg, 85%) as colourless syrup. $[\alpha]_D^{20}$ +55.9 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 697, 737, 1027, 1069, 1168, 1208, 1258, 1454, 1498, 1720, 2858, 2929. ¹H-NMR (CDCl₃, 400 MHz) δ 7.66–7.64 (m, 4 H, aromatic H), 7.43–7.07 (m, 41 H, aromatic H), 5.33 (d, J = 7.2 Hz, 1 H, NH), 5.26 (d, J =7.6 Hz, 1 H, NH), 5.11-4.73 (m, 14 H, 12 CHH, H-1a, H-1b), 4.52 (bt, 2 H, 2 CHH), 4.32-4.25 (m, 2 H, 2 CHNH), 4.17-3.99 (m, 6 H, H-5a, H-5b, H-3, 3 CHH), 3.91-3.74 (m, 5 H, H-3, H-2°, 3 CHH), 3.67-3.61 (m, 3 H, H-2a, 2 CHH), 3.50-3.42 (m, 3 H, H-2b, H-4a, H-4b), 1.35 (d, J = 6.8 Hz, 1 H, CH_3), 1.28 (d, $J = 7.2 \text{ Hz}, 1 \text{ H}, \text{ C}H_3$), 1.06 (s, 9 H, 3 C H_3). ¹³C-APT (CDCl₃, 100 MHz) δ 172.82, 172.65, 155.65, 155.57 (4 C=O), 138.38, 138.15, 137.92, 137.83, 137.55, 136.29 (aromatic C), 135.65, 135.62 (aromatic CH), 133.17, 133.09 (aromatic C), 128.71, 128.58, 128.52, 128.48, 128.43, 128.39, 128.22, 128.20, 128.18, 128.16, 127.98, 127.95, 127.93, 127.90, 127.77, 127.73 (aromatic CH), 96.63 (C-1a), 95.90 (C-1b), 82.27 (C-5b), 80.14 (C-5a), 79.57 (C-2°), 77.97 (C-2b), 77.67 (C-4), 77.59 (C-4), 76.41 (C-2a), 75.95, 75.61, 75.08, 75.03, 73.81 (CH₂), 69.21 (C-3b), 69.06 (C-3a), 69.97, 63.84, 63.09, 63.03, 62.12 (CH₂), 49.68 (CHNH), 49.60 (CHNH), 26.94 (3 CH₃), 19.26, 18.70 (CH₃),

18.57 (CH₃). HR-MS: calculated for $C_{88}H_{98}N_2O_{19}Si [M + Na]^+$: 1537.64253, found: 1537.64262.

Synthesis of 34. Compound 32 (610 mg, 0.4 mmol) and DMTrCl (273 mg, 0.8 mmol) were dissolved in DCM (4 mL) stirring at room temperature. Then DIPEA (283 µL, 1.6 mmol) was added in the mixture. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial. Then the mixture was diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated in vacuo. The product was purified by silica gel column chromatography. Compound 34 (691 mg, 95% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +118.8 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 738, 827, 1029, 1070, 1175, 1209, 1249, 1454, 1508, 1607, 1724, 2931. ¹H-NMR (d-acetone, 400 MHz) δ 7.74–7.68 (m, 4 H, aromatic H), 7.55-7.12 (m, 50 H, aromatic H), 6.89 (bd, 4 H, aromatic H), 6.82 (d, J = 7.6 Hz, 1 H, NH), 6.79 (d, J = 8.0 Hz, 1 H, NH), 5.58 (d, J = 2.8 Hz, 1 H, H-1a), 5.25 (d, J = 2.8 Hz, 1 H, H-1b),5.15-5.05 (m, 5 H, 5 CHH), 4.97-4.71 (m, 6 H, 2 CHH), 4.56-4.11 (m, 13 H), 3.91 (dd, $J_1 = 2.8$ Hz, $J_2 = 9.6$ Hz, 1 H, H-2a), 3.82-3.66 (m, 10 H), 3.59-3.52 (m, 2 H, H-2b, H-6a), 3.34-3.31 (m, 1 H, H-6_b), 1.49-1.45 (m, 6 H, 2 CH₃), 1.04 (s, 9 H, 3 C H_3). ¹³C-APT (d-acetone, 100 MHz) δ 173.52, 173.41 (2 C=O), 159.55 (aromatic C), 156.90, 156.87 (2 C=O), 146.16, 139.99, 139.74, 139.68, 138.48, 139.32, 138.03, 137.98, 136.85, 136.69 (aromatic C), 136.40, 136.38 (aromatic CH), 134.06, 134.03 (aromatic C), 131.04, 131.00, 130.67, 130.65, 129.10, 129.07, 129.01, 128.95, 128.83, 128.67, 128.55, 128.40, 128.33, 128.26, 128.24, 128.17, 127.58, 114.08, 114.06 (aromatic CH), 94.97 (C-1a), 93.94 (C-1b), 87.37 (quaternary C), 82.55 (C-5a), 81.79 (C-5b), 80.87 (C-2b), 78.59 (C-4), 78.26 (C-4), 76.90 (C-2°), 76.58 (CH₂), 76.03 (C-2a), 75.86, 75.64, 75.53, 72.41 (CH₂), 70.29 (C-3b), 70.11 (C-3a), 66.93, 66.90, 64.59, 64.54, 63.86 (CH₂), 55.53 (2 OCH₃), 50.72 (2 CHNH), 27.39 (3 CH₃), 19.73 (quaternary C), 17.97 (CH₃), 17.93 (CH₃). HR-MS: calculated for $C_{109}H_{116}N_2O_{21}Si [M + Na]^+: 1839.77321, found: 1839.77281.$

Synthesis of 19. Compound 33 (730 mg, 0.4 mmol) was dissolved in THF/pyridine (1:1/v:v, 6 mL) stirring at room temperature. Then HF-pyridine (0.4 mL) was added in the mixture. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial (10 h). Then the mixture was diluted with EtOAc. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated in vacuo. The product was purified by silica gel column chromatography. Compound 19 (548 mg, 86% yield) was obtained as a colourless syrup. $[\alpha]_D^{20}$ +64.0 (c = 1, CHCl₃). IR (neat, cm⁻¹) ν 698, 751, 830, 1029, 1070, 1176, 1213, 1251, 1302, 1454, 1508, 1608, 1724, 2928. H-NMR (d-acetone, 400 MHz) δ 7.49–7.10 (m, 44 H, aromatic H), 6.87-6.73 (m, 5 H, 4 aromatic H, 1 NH), 6.74 (d, J = 7.6 Hz, 1 H, NH), 5.52 (d, J = 2.8 Hz, 1 H, H-1a), 5.26 (d, J = 2.8 Hz, 1 H, H-1b), 5.10-5.02 (m, 5 H, 5 CHH), 4.93-4.80 (m, 4 H, 2 CHH), 4.70-4.62 (m, 3 H), 4.56-4.48 (m, 2 H), 4.42-4.21 (m, 7 H), 4.17-4.01 (m, 4 H), 3.88-3.86 (m, 2 H), 3.75-3.47 (m, 12 H), 3.19 (dd, J_1 = 5.6 Hz, J_2 = 10.0 Hz, 1 H), 1.47-1.38 (m, 6 H, 2 CH₃). ¹³C-APT (d-acetone, 100 MHz) δ 173.50, 173.43 (2 C=O),

159.60, 159.58 (aromatic C), 156.99, 156.89 (2 C=O), 146.16, 140.03, 139.77, 139.58, 139.50, 138.07, 138.04, 136.93, 136.71 (aromatic C), 131.07, 130.99, 129.25, 129.14, 129.13, 129.09, 129.07, 129.05, 129.04, 128.99, 128.83, 128.76, 128.67, 128.64, 128.43, 128.34, 128.25, 128.19, 128.62, 114.06, 114.04 (aromatic CH), 95.52 (C-1a), 93.96 (C-1b), 87.35 (quaternary C), 82.63, 81.72, 80.85, 78.83, 78.59, 78.29, 76.48, 76.06, 75.86, 75.63, 72.42, 70.34, 70.20, 66.92, 65.28, 64.32, 63.92, 63.43, 55.54 (2 OCH₃), 50.78 (CHNH), 50.74 (CHNH), 17.98 (2 CH₃). HR-MS: calculated for C₉₃H₉₈N₂O₂₁ [M + NH₄][†]: 1596.70003, found: 1596.69828.

Synthesis of 36. Compound 19 (40 mg, 0.025 mmol) and DCI (1.5 eq., 0.25 M in CH₃CN) was dissolved in dry CH₃CN (1 mL) stirring at room temperature. Then compound 35 (1.1 eq.) was added in the solution. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial (2–3 h). Then CSO (0.5 M in CH₃CN) was added in the mixture. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial (10–30 min). Then the mixture was diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated *in vacuo*. Crude compound 36 (50 mg) was obtained as a colourless syrup.

Synthesis of 37. Crude compound 36 (50 mg, 0.025 mmol) was dissolved in CH₃OH/DCM = 1:1 (v:v, 1 mL) stirring at room temperature. Then TsOH (0.1 eq.) was added in the solution. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial (2-3 h). Then the reaction was quenched with Et₃N, diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO4, filtered and concentrated in vacuo. The product was purified by size exclusion chromatography (DCM: MeOH = 1:1). Compound 37 (40 mg, 93%) was obtained as a colourless syrup. ¹H-NMR (d-acetone, 400 MHz) δ 7.47–7.05 (m, 50 H, aromatic H), 6.95 (d, J = 7.6 Hz, 1 H, 1 NH), 6.74 (d, J = 7.6 Hz, 1 H, NH), 5.42–5.35 (m, 2 H, 2 H-1), 5.12-3.58 (m, 44 H), 2.89 (s, 1 H), 1.42-1.37 (m, 6 H, 2 CH₃). 13 C-APT (d-acetone, 100 MHz) δ 173.52, 173.33, 156.93, 156.82 (4 C=O), 138.91, 139.64, 139.49, 139.39, 139.33, 138.03 (aromatic C), 129.36, 129.17, 129.09, 129.04, 129.02, 128.87, 128.85, 128.77, 128.71, 128.68, 128.60, 128.53, 128.52, 128.46, 128.35, 128.24, 128.22, 128.17, 128.14 (aromatic CH), 96.03 (C-1), 85.08 (C-1), 82.74, 81.47, 79.96, 78.89, 78.53, 77.80, 77.71, 77.67, 77.60, 76.43, 76.09, 75.93, 75.56, 73.72, 73.07, 72.50, 70.22, 70.15, 69.87, 69.84, 67.89, 67.73, 66.81, 64.14, 63.64, 61.32, 61.25, 63.43, 55.54, 50.76, 50.65 (CHNH), 50.74 (CHNH), 17.85, 17.79 (2 CH₃). 31 P (d-acetone, 400 MHz) δ 0.77,

Synthesis of 39. Compound 37 (40 mg, 0.023 mmol) and DCI (1.5 eq., 0.25 M in CH₃CN) was dissolved in dry CH₃CN (1 mL) stirring at room temperature. Then compound 38 (1.1 eq.) was added in the solution. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial (2–3 h). Then CSO (0.5 M in CH₃CN) was added in the mixture. The reaction was stirred at rt until TLC-analysis showed complete conversion of the starting martial

(10-30 min). Then the mixture was diluted with DCM. The organic phase was washed with water and brine, dried with anhydrous MgSO₄, filtered and concentrated in vacuo. The product was purified by size exclusion chromatography (DCM: MeOH = 1:1). Compound 39 (44 mg, 88%) was obtained as a colourless syrup. ¹H-NMR (d-acetone, 400 MHz) δ 7.45–7.06 (m, 65 H, aromatic H), 6.94–6.92 (m, 1 H, 1 NH), 6.73 (d, J = 7.6 Hz, 1 H, NH), 5.38-5.34 (m, 2 H, 2 H-1), 5.12-3.54 (m, 57 H), 1.42-1.37 (m, 6 H, 2 CH₃). ¹³C-APT (d-acetone, 100 MHz) δ 173.39, 173.22, 156.79, 156.68 (4 C=O), 139.87, 139.54, 139.51, 139.49, 139.33, 139.23, 137.88, 137.19, 137.14, 137.07, 137.02 (aromatic C), 129.26, 129.23, 129.11, 129.06, 129.05, 129.04, 129.00, 129.96, 129.93, 128.90, 128.87, 128.85, 128.83, 128.72, 128.68, 128.62, 128.55, 128.53, 128.45, 128.38, 128.35, 128.26, 128.20, 128.11, 128.07, 127.99, 127.97 (aromatic CH),96.37, 96.23 (C-1), 94.95, 94.80 (C-1), 82.60, 81.25, 80.24, 78.55, 78.24, 77.53, 77.50, 77.47, 77.45, 77.38, 76.27, 76.24, 76.07, 75.95, 75.79, 75.49, 75.43, 73.57, 72.97, 72.93, 72.87, 72.37, 72.35, 72.33, 72.31, 70.38, 70.07, 69.83, 69.79, 69.76, 69.73, 69.71, 69.67, 67.85, 67.80, 67.76, 67.69, 67.65, 66.68, 65.92, 65.87, 63.91, 63.55, 50.61 (CHNH), 50.51 (CHNH), 17.71, 17.64 (2 CH₃). 31 P (d-acetone, 400 MHz) δ 0.53, 0.57, 0.62, 0.65, 0.68, 0.78, 0.82. HR-MS: calculated for $C_{120}H_{130}N_2O_{29}P_2[M + Na]^+$: 2147.81267, found: 2147.81610.

Synthesis of 40. Crude compound 39 (11 mg, 0.005 mmol) was dissolved in dioxane/H₂O = 2:1 (v:v, 1 mL) stirring at room temperature. Then Pd/C was added in the solution. The reaction was stirred at rt for 3 days under a H₂ atmosphere, filtered and concentrated in vacuo. Compound 40 (~5 mg, 93%) was obtained as a white solid. 1 H-NMR (d-acetone, 400 MHz) δ 5.42 (d, 1 H, H-1), 5.11 (d, 1 H, H-1), 4.63-4.57 (m, 2 H), 4.42-4.38 (m, 2 H), 4.21-4.15 (m, 5 H), 3.98-3.56 (m, 20 H), 3.50-3.38 (m, 2 H), 1.87 (s, 1 H), 1.57-1.53 (m, 6 H, 2 CH₃). ¹³C-APT (d-acetone, 100 MHz) δ 170.46, 170.38 (2 C=O), 96.00, 94.42, 75.04, 74.70, 72.37, 72.17, 71.71, 71.54, 71.04, 70.85, 70.72, 70.68, 70.64, 70.61, 69.82, 69.43, 69.20, 66.42, 66.37, 66.32, 65.22, 64.90, 64.50, 62.47, 62.04, 62.01, 60.38, 50.45, 48.72, 16.06, 15.20. ³¹P (d-acetone, 400 MHz) δ 1.22, 1.13. HR-MS: calculated for $C_{27}H_{52}N_2O_{25}P_2$ [M + H]⁺: 867.24071, found: 867.24106.

Conflicts of interest

There are no conflicts to declare.

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