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Synthesis of orthogonally protected and functionalized bacillosamines†

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2,4-Diamino-2,4,6-trideoxyglucose (bacilosamine) is a monosaccharide found in many pathogenic bacteria, variation in the functionalities appended to the amino groups occurs depending on the species the sugar is derived from. We here report the first synthesis of bacilosamine synthons that allow for the incorporation of two different functionalities at the C-2-N-acetyl and C-4-amines. We have developed chemistry to assemble a set of conjugation ready *Neisseria meningitidis* C-2-N-acetyl bacilosamine saccharides, carrying either an acetyl or (R)- or (S)-glyceroyl at the C-4 amine. The glyceroyl bacilosamines have been further extended at the C-3-OH with an α -D-galactopyranose to provide structures that occur as post-translational modifications of *N. meningitidis* PilE proteins, which make up the bacterial pili.

Introduction

Bacilosamine (**1**, Bac, Fig. 1) is a characteristic bacterial monosaccharide that is present in various bacterial capsular polysaccharides and glycoproteins.¹ It is a 2,4-diamino-2,4,6-trideoxyhexose (DATDH), that is most commonly functionalized with *N*-acetyl groups at C-2 and C-4 (2,4-di-NAcBac, **2**). This residue is found at the reducing end of oligosaccharide chains of both *O*- and *N*-linked glycoproteins of *Campylobacter jejuni*² and *Neisseria gonorrhoeae*.³ Recently, a differently modified bacilosamine has been discovered in *Neisseria meningitidis*,⁴ where it is decorated, besides a single acetamide, with an *N*-glyceroyl group, thought to reside at the C4-amine (**3**, Fig. 1).⁵ This Bac-residue represents one of various post-translational modifications (PTMs) that can occur on *N. meningitidis* PilE proteins, the constituent protein monomers of the type IV pili. It can be further decorated with

different mono- or disaccharides (see Fig. 1). The role of these PTMs is currently not known but it is clear that the type IV pili play an important role in the pathogenicity of the bacterium. Also, the exact structure of the Bac-residue present on the *N. meningitidis* pili remains elusive as the stereochemistry of the glyceroyl group has not been established.⁵

To aid in the structure elucidation of the functionalized Bac residues in the *N. meningitidis* pili, its detection and the role of these differentially decorated bacilosamines in virulence, we reasoned that the availability of synthetic Bac-residues would provide powerful tools. We therefore set out to develop an effective route of synthesis for an orthogonally protected bacilosamine synthon, of which all functional groups can be site selectively addressed. We here report the first synthesis of such an orthogonally protected bacilosamine building block and we show the synthesis of spacer-equipped bacilosamines bearing different acyl functionalities (**4b**–**4c**, Fig. 1) and the extension of these towards a relevant conjugation-ready disaccharide (**5**, Fig. 1).

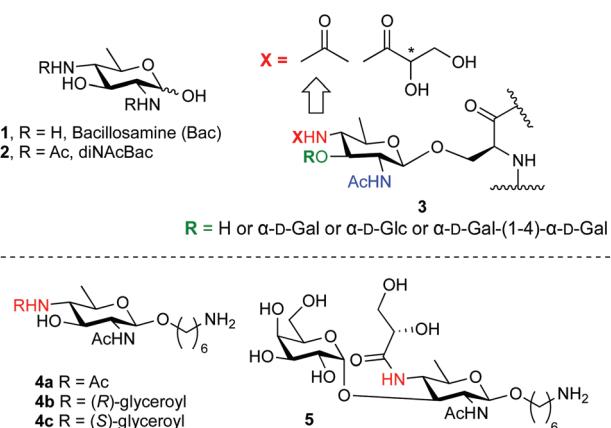


Fig. 1 Bacilosamine (**1**), di-*N*-acetylbacilosamine (**2**), bacilosamine PTMs found in *N. meningitidis* pili (**3**) and the synthetic targets of the current study (**4** and **5**).

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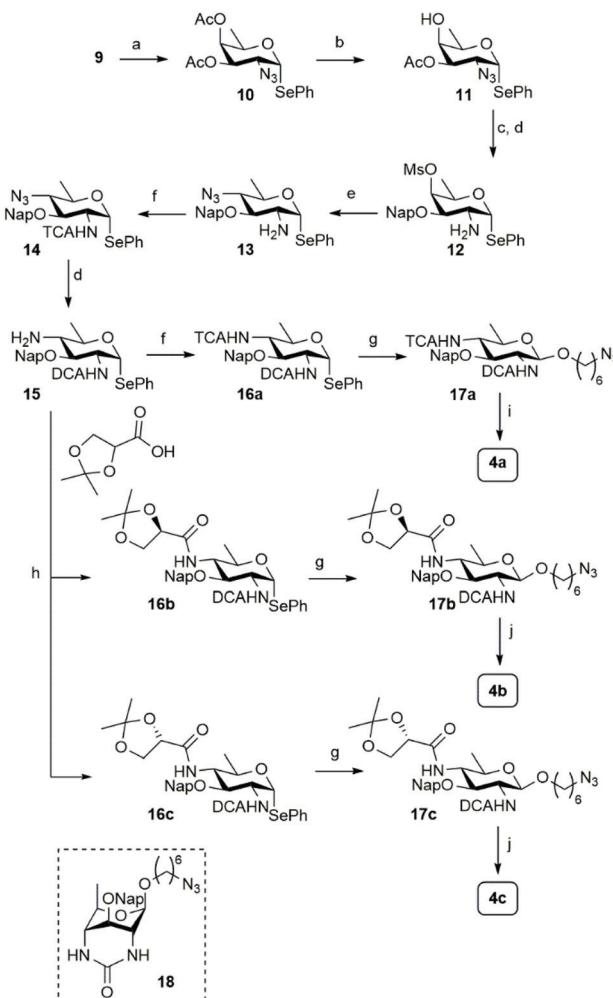
† Electronic supplementary information (ESI) available: Synthetic procedures and experimental data. See DOI: 10.1039/d0ob00256a



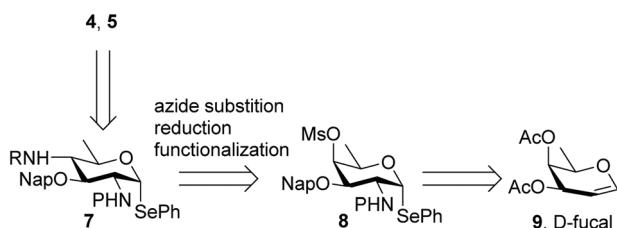
Results and discussion

Different syntheses of bacillosamine⁶ and bacilosamine containing oligosaccharides⁷ and glycopeptides⁸ have been reported, but none of these allow for the discrimination of the C-2 and C-4 amino groups and therefore the development of a new synthetic route was required. We thus set out to target key synthon 7, bearing two orthogonally protected amino groups at C-2 and C-4, as well as a 2-naphthylmethyl ether that masks the C-3-OH and an anomeric selenophenyl group for glycosylation purposes (see Scheme 1). Retrosynthetically, this building block can be derived from a suitably protected 6-deoxy galactosamine derivative 8, by a substitution reaction of the C-4-O-mesylate. Selenophenyl galactosamine 8 can be readily accessed from D-fucal through an azidoselenation event⁹ and subsequent transformation of the azide into the protected C-2-aminosugar.

Accordingly, 3,4-di-O-acetyl-D-fucal 9¹⁰ was subjected to azidoselenation conditions to provide azidofucose 10¹¹ (see Scheme 2). Saponification of the two acetates in 10 was then followed by regioselective installation of the C-3-naphthylmethyl ether and subsequent mesylation of the C-4-OH. Because we aimed to use azide as nucleophile for the introduction of the C-4-amine functionality, we had to transform the C-2-azide in 11 to the corresponding amine and invert the C-4-O-mesylate in the resulting fucosamine, without masking the C-2-amine. Thus, fucosazide 11 was exposed to 1,3-propanedithiol to deliver fucosamine 12, which was treated with NaN₃ and TBABr in DMF at elevated temperature to provide the desired bacilosamine 13, bearing a fully orthogonal functional group pattern, in 81%. Next, the C-2-amine was masked with a trichloroacetyl (TCA) group to provide fully protected bacilosamine 14. Reduction of the azide with 1,3-propanedithiol then set the stage for the differentiation of the C-4-amino group into either an acetamide (towards 4a) or the two glyceroyl substituted bacilosamines 4b and 4c. During the reduction of the azide in 14, the TCA moiety was partially reduced as well to the corresponding C-2-dichloroacetamide (DCA), yielding compound 15. As the DCA group can be



Scheme 2 Synthesis of differentially functionalized bacilosamines. Reagents and conditions: (a) (Ph₃)₂, PhI(OAc)₂, TMS-N₃, DCM, -20 °C to rt, 93%. (b) (i) NaOMe, MeOH. (ii) Bu₂SnO, toluene, 110 °C. (iii) TBABr, NapBr, 82%. (c) MsCl, pyr. 0 °C to rt, 93%. (d) 1,3-Propanedithiol, TEA, DMF, 92% for 12, 82% for 15. (e) NaN₃, TBABr, DMF, 80 °C, 81%. (f) TCA-Cl, DCM : pyr., 0 °C, 89% for 14, 88% for 16a. (g) 6-Azidohexanol, NIS, TMSOTf, DCM, 0 °C, 70% for 17a, 74% for 17b, 74% for 17c. (h) 2,2-Dimethyl-1,3-dioxolane-4-carboxylic acid (R for 16b, S for 16c), HCTU, DIPEA, DMF, 0 °C to rt, 83% for 16b, 80% for 16c. (i) Pd(OH)₂/C 20 wt%, H₂, H₂O/THF/tBuOH, 63%. (j) (i) H₂O/THF/AcOH, 80 °C. (ii) Pd(OH)₂/C 20 wt%, H₂, H₂O/THF/tBuOH, 79% for 4b, 81% for 4c.



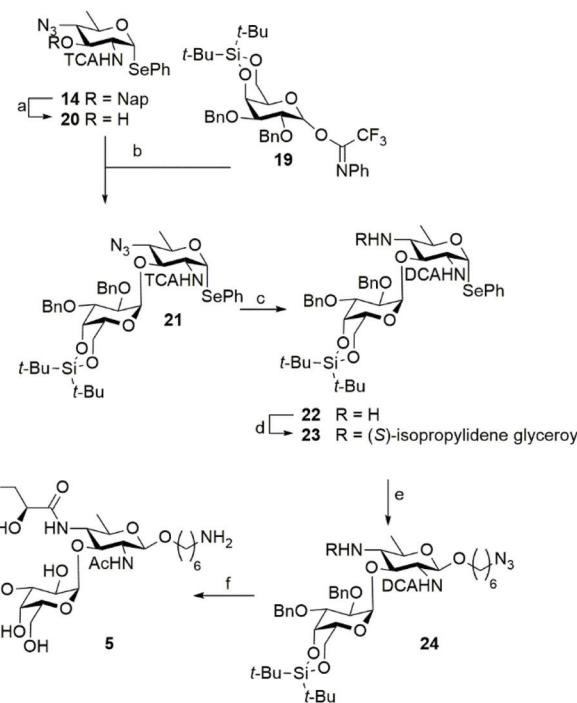
Scheme 1 Retrosynthetic analysis for the assembly of target compounds 4 and 5.

removed under similar conditions as the TCA, the synthetic strategy did not need any adjustment and we continued with 4-N-acylation. Trichloroacetylation provided bacilosamine synthon 16a while reaction of the C-4-amine with either *R*- or *S*-isopropylidene-glyceric acid under the agency of HCTU, provided bacilosamines 16b and 16c. Next, the three building blocks were equipped with an azidohexanol spacer, in glycosylation reactions promoted by *N*-iodosuccinimide (NIS) and trimethylsilyl trifluoromethanesulfonate (TMSOTf) to provide 17a, 17b and 17c. The products were obtained as single diastereoisomers and no oxazoline side products were detected, indicating that the dichloroacetyl groups provided effective



anchimeric assistance.¹² With the fully protected spacer-equipped bacillosamines in hand, deprotection of the intermediates was undertaken. We first tried to unmask the TCA and DCA groups in **17a** using Cs_2CO_3 in DMF,¹³ but this did not lead to the desired C-2,4-diamine. Instead bicyclic urea **18** was formed.¹⁴ We therefore switched to a hydrogenation reaction, aiming to unmask the spacer azide, C-2-N-DCA, C-3-O-Nap and C-4-N-TCA groups in one step. After the first hydrogenation event using Pearlman's catalyst, LC-MS analysis of the crude reaction mixture revealed the presence of three chlorides and therefore the mixture was resubjected to the hydrogenation conditions until all chlorides had been removed. This delivered spacer-equipped di-N-acetyl bacillosamine **4a** in 63% yield. After the acidic hydrolysis of the glyceroyl isopropylidene ketal in **17b** and **17c**, deprotection under the same conditions delivered the C-2-N-acetyl-C-4-N-glyceroyl bacilosamines **4b** and **4c** in 79% and 81% yield, respectively.

With a successful route of synthesis towards differentially functionalized bacilosamines established, we set out to probe the generation of a *N. meningitidis* disaccharides, featuring an additional α -D-galactopyranose attached to the C-3-OH of the bacilosamine. To install the required 1,2-cis-galactosyl linkage we used a 4,6-di-*tert*-butyl silylidene protected galactose donor **19** as 4,6-silylidene functionalized galactopyranosides are amongst the most reliable donors for the installation of *cis*-galactosyl linkages.¹⁵ Because the presence of amides in glycosyl acceptors can lead to problematic glycosylation reactions¹⁶ we decided to attach the α -galactoside prior to the introduction of the *N*-glyceroyl. Thus, we prepared C-4-azide acceptor **20** by DDQ-mediated unmasking of the C-3-hydroxyl function in **14**. In a chemoselective glycosylation reaction, imidate donor **19**¹⁷ and bacilosamine selenoglycoside acceptor **20** were then reacted to provide, in a completely stereoselective manner, the desired disaccharide **21** in 91% yield. Before installation of the azidohexanol spacer, the C-4-N-glyceroyl groups had to be incorporated and we therefore reduced the C-4-azide in **21**, which again led to the conversion of the C-2-N-TCA amide into the corresponding DCA-group. Amine **22** was obtained in 48% yield and condensed with (S)-isopropylidene-glyceric acid using HCTU as the condensation agent to furnish the glyceroyl bacilosamine disaccharide **23** in 30%. It was observed that both the reduction and the amide bond forming reactions proceeded significantly slower in the disaccharide than the corresponding reactions on the monosaccharides described above. This may be attributed to increased steric hindrance of the appended galactopyranose with the neighboring C4-nitrogen. Next the azidohexanol spacer was introduced using NIS/TMSOTf to activate the seleno disaccharide to afford the fully protected disaccharide **24** in 57% yield. At this stage, all protecting groups in disaccharide **24** were removed by the following procedure. First, the silylidene group was removed by treatment of HF-Pyr in THF yielding the diol in 64% yield. Next, the isopropylidene group in the glyceroyl moiety was hydrolysed with $\text{AcOH}/\text{H}_2\text{O}/\text{THF}$. Finally, the hydrogenation procedure, as described above, was undertaken to remove the benzyl ethers and transform the dichloroacetamide into the



Scheme 3 Synthesis of galactosylated C4-N-glyceroyl bacilosamines. Reagents and conditions: (a) DDQ, DCM/H₂O, 80%. (b) Compound **20**, TMSOTf, DCM, 0 °C, 91%. (c) 1,3-Propanedithiol, TEA, DMF, 48%. (d) (S)-2,2-Dimethyl-1,3-dioxolane-4-carboxylic acid for, HCTU, DIPEA, DMF, 0 °C to rt, 30%. (e) 6-Azidohexanol, NIS, TMSOTf, DCM, 0 °C, 57%. (f) i) HF-pyr, THF, 64% (ii) H₂O/THF/AcOH, 80 °C. (iii) Pd(OH)₂/C 20 wt%, H₂, H₂O/THF/tBuOH, 41%.

corresponding acetamide and the spacer azide into a primary amine. The hydrogenation reaction was monitored by LC-MS analysis, and repeated when required. In this manner disaccharide **5** was obtained in 41% yield (Scheme 3).

Conclusions

In conclusion, we have here disclosed the first synthesis towards an orthogonally protected bacilosamine synthon, that allows for the differentiation of the two amino functionalities, and its use in the generation of a *N. meningitidis* PilE disaccharide bearing a C-3- α -D-galactose appendage and a C-4-N-glyceroyl. The chemistry developed here can be used to generate differentially functionalised bacilosamines and pave the way to the assembly of larger bacilosamine containing oligosaccharides as well as glycopeptides. The set of conjugation ready bacilosamines can be used to generate antibodies directed at these *N. meningitidis* PilE post-translational modifications for both therapeutic and diagnostic purposes.

Conflicts of interest

There are no conflicts to declare.



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