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Synthesis of $C$-glycosyl phosphonate derivatives of 4-amino-4-deoxy-$\alpha$-L-arabinose

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Abstract

Incorporation of basic substituents into the structurally conserved domains of cell-wall lipopolysaccharides has been identified as a major mechanism contributing to antimicrobial resistance of Gram-negative pathogenic bacteria. Inhibition of the corresponding enzymatic steps, specifically the transfer of 4-amino-4-deoxy-L-arabinose would thus restore the activity of cationic antimicrobial peptides and several antimicrobial drugs. $C$-glycosidically linked phospholipid derivatives of 4-amino-4-deoxy-L-arabinose have been prepared as hydrolytically stable and chain-shortened analogues of the native undecaprenyldonor. The $C$-phosphonate unit was installed via a Wittig-type reaction of benzyl-protected 1,5-arabinonic acid lactone with the lithium salt of dimethyl methylphosphonate followed by an elimination step of the resulting hemiketal leading to the corresponding exo- and endo-glycal
derivatives. The ensuing selective mono-demethylation and hydrogenolysis of the benzyl groups and reduction of the 4-azido group gave the α-L-anomeric arabino- and ribo-configured methyl phosphonate esters. In addition, the monomethyl phosphonate glycal intermediates were converted into n-octyl derivatives followed by subsequent selective removal of the methyl phosphonate ester group and hydrogenation to give the octyl-phosphono derivatives. These intermediates thus will be of value for future conversion into transition state analogues as well as for introduction of various lipid extensions at the anomeric phosphonate moiety.

**Keywords**

Glycosyl phosphonate; lipid A; lipopolysaccharide; glycosyl transferase; antibiotic resistance

**Introduction**

Glycosyltransferases are important enzymes to accomplish the transfer of activated sugar phosphates onto their respective acceptor molecules [1]. In most cases, nucleotide diphosphate sugars serve as the reactive species, but lipid-linked diphosphate derivatives are equally important, e.g. when connected to dolichol in mammalian systems or to undecaprenol in prokaryotic donor substrates for bacterial glycosyltransferases [2]. 4-Amino-4-deoxy-L-arabinose (Ara4N) is an important microbial carbohydrate in bacterial lipopolysaccharides (LPS) and has been implicated in resistance mechanisms of pathogenic Gram-negative bacteria against antibiotics such as polymyxin B and colistin, respectively [3]. The main effect of Ara4N incorporation into the lipid A part - and less frequently into the inner core region of LPS [4] – is thought to originate from blocking the electrostatic interaction of
cationic antimicrobial peptides with the negatively charged phosphate and carboxylate groups in LPS domains. Suitable inhibitors intercepting the attachment of Ara4N units to the lipid A and inner core region might restore sensitivity towards the cationic antimicrobial drugs as a novel approach to combat the looming antibiotic crisis [5]. 4-Amino-4-deoxy-L-arabinose units are activated as the phosphodiester linked undecaprenyl derivative [6] that is then transferred by the action of several Ara4N-transferases (ArnT) [7]. Synthesis of potential inhibitors of the biosynthesis of Ara4N and the glycosyl transfer has not been fully explored yet. Previously, Kline et al. reported on the synthesis of acetylated 4-azido-arabinose phosphate and UDP-derivatives, respectively. In addition, a 4-aminophosphoamidate UDP-derivative was also obtained [8]. Whereas these compounds were inactive towards enzyme upstream of the biosynthetic pathway to undecaprenyl Ara4N, the peracetylated 4-azido-derivative showed a modest reduction of Ara4N incorporation into the lipid A part of a *Salmonella typhimurium* strain [8].

We have recently set out to study the substrate specificity of ArnT enzymes in more detail using phosphodiester linked derivatives in both anomeric configurations and containing short lipid appendages in order to define the minimum structural requirements for Ara4N glycosyltransferase substrates [9]. In parallel, we have started to develop C-phosphonate analogues as these have frequently been exploited as potential inhibitors for glycosyl transferases, since the carbon-phosphorus bond is not hydrolyzed in the active site of glycosyl transferases [10-13]. Herein we report on the synthesis of α-anomeric C-arabinosyl methylphosphonate ester derivatives as model compounds to allow for future incorporation of different lipid chains and options towards glycal analogues as potential transition state analogues to inhibit 4-amino-4-deoxy-L-arabinose transfer to bacterial LPS.
Results and Discussion

The previously synthesized [14] methyl 4-azido-4-deoxy-\(\beta\)-L-arabinopyranoside 1, available in multigram amounts, was benzylated in 79% yield using benzyl bromide and sodium hydride in DMF, followed by hydrolysis of the anomic aglycon of 2 with 5 M HCl and SrCl\(_2\) in acetic acid which afforded the hemiacetal 3 as a 2:3 \(\alpha/\beta\) anomic mixture in 60% yield (Scheme 1). Conversion of benzylated reducing sugars such as glucose [15], \(N\)-acetyl-\(D\)-glucosamine [16] and galactose [17] into the corresponding C-glycosyl phosphonates using a Wittig reaction with methylenetriphenylphosphorane furnished the respective glycoenitols which were then subjected to mercuriocyclization followed by iodination and phosphonate introduction by an Arbusov reaction. Alternative approaches were elaborated from olefinic C-glycosides which were converted into the corresponding C-linked hydroxymethyl derivatives and processed to give the glycosyl methylphosphonic acid derivatives [18,19]. As the Ara4N transferase reacts with the equatorial phosphodiester lipid, we opted to use lactone 4 as suitable precursor for C-glycosyl phosphonates based on literature precedents [10, 20, 21]. Thus, modified Swern oxidation of lactol 3 with acetic anhydride in DMSO afforded a near quantitative yield of lactone 4 which, however, was unstable upon chromatographic purification and was used as a crude material after lyophilization to remove solvent. Coupling of 4 with the lithium salt of dimethyl methylphosphonate in THF afforded the \(\beta\)-anomeric ketol phosphate derivative 5 in 57% yield. The axial orientation of the anomic hydroxy group was proven by NOesy correlations of the H-2 proton to both geminal protons of the CH\(_2\)P unit. A strong NOe was also observed for the upfield shifted doublet of doublets at 1.71 ppm of the methylene group, which had an additional NOe correlation to the broad signal of the anomic OH group at 5.79 ppm.
Next, the ensuing elimination step was carried out to explore the access to transition state analogues [22], potentially mimicking the sp$^2$ character of the oxocarbenium intermediate in the enzymatic transfer reaction. In addition, exo-glycals are versatile precursors for introduction of fluoromethyl phosphonates which are the better bioisosters of phosphates [23,24]. Exo-glycal 8 was obtained in 74% yield when using methyl oxalyl chloride [25], which proved to be superior to the use of trifluoroacetic anhydride, which gave 8 in 57% yield [26]. The Z-configuration of the hexenitol unit was derived from the $^3J_{C,P}$ coupling constant (14.1 Hz) in good agreement with reported data [22]. Next, selective de-O-methylation of the phosphonate diester was elaborated. Using sodium iodide in acetone afforded the mono de-O-methylated derivative 9 in very good yield [27, 28]. At this stage, the Z-configuration of the enol ether was unambiguously assigned on the basis of NOesy experiments. Specifically, NOe-correlations were seen from the olefinic proton to the methylene protons of a benzyl group. The location of the 2-O-benzyl group was assigned on the basis of an HMBC-correlation to H-2 of the pyranose ring. Additional interactions were also found from the olefinic proton to H-2 and the O-methyl group, respectively. The preferred formation of Z-configured Wittig products fully agrees with similar results in the literature for 2-O-benzyl protected gluco- and galacto-derivatives [20, 26, 29, 30].

Full deprotection including the conversion of the 4-azido-group into the amino function was accomplished upon hydrogenation of 9 on palladium hydroxide in a 1:1 mixture of methanol - acetic acid to deliver the target derivative 11 in 15% yield after final HILIC purification. The equatorial arrangement of the C-glycosyl linkage was supported from the large value of the coupling constant $J_{1,2}$ (9.5 Hz) indicating a 1,2-trans orientation of the respective protons.
In addition to exo-cyclic glycals mimicking putative planar transition states of substrates involved in enzymatic reactions such as glycosyl transfer, mutase and epimerization, endo-glycals are also of interest [21, 22, 31-34].

Scheme 1: Phosphonate formation and deprotection, reagents and conditions: a) BnBr, NaH, DMF, 79%; b), SrCl₂, 5 M HCl, AcOH, 60%; c) DMSO, Ac₂O, then d) 5, n-BuLi, THF, 57%; e) 7, pyridine, DCM, 74%; f), NaI, acetone, 98% for 9, 82% for 13, 75% for 15; g) 1 M NaOH, MeOH, 80%; h) H₂/Pd(OH)₂-C, MeOH, AcOH, 50 bar, flow conditions, HILIC, 15% for 11, 38% for 16, 41% for 17; i) 1-bromo-octane, Cs₂CO₃, DMF, 78-86% for 12 and 14, for product ratios see Text.

Under basic conditions, such as by treatment of 9 with 1 M NaOH in MeOH, the exo-double bond of compound 9 could be shifted to produce the corresponding endo-glycal 10 in 80% yield. The m/z value of the high-resolution mass spectrum indicated that 10 was an isomer of 9. In the 600 MHz ¹H NMR spectrum of 10, the olefinic
proton was absent, whereas the $^{13}$C NMR spectrum showed downfield shifts for the anomeric carbon (146.48 ppm) and the adjacent ring carbon (133.95 ppm). Evidence by an HMBC experiment additionally indicated a correlation from benzylic protons to the latter carbon as well as signals of two upfield shifted deoxy-protons at 2.72 and 2.55 ppm, respectively, to the anomeric carbon. In addition, for compound 10 lacking the conjugation to the phosphorus atom, a significant downfield shift of the $^{31}$P NMR signal was observed (17.97 ppm in 10, versus 10.95 ppm in compound 9).

The monomethyl phosphonate ester 9 was then subjected to alkylation reactions in order to allow for a selective introduction of longer-chain alkyl ester groups as better mimics of the native prenyl-activated donor substrate. Since the reaction of 9 with 1-bromo-octane as a model system also involved basic conditions, mixtures of endo- and exo-glycal derivatives 12 and 14 were obtained. Highest yields were obtained for reactions performed in DMF, whereas DMSO and MeCN as solvents were less effective. Reaction times of 2-5 h at 80 °C were sufficient, the ratio of isomers obtained, however, critically depended on the ensuing work-up procedure. Addition of MeOH prior to chromatography in combination with dry-loading of the column mainly produced the endo-glycal 14 as the major product (55%) as well as the exo-isomer (31%). In contrast, direct separation of a concentrated reaction mixture by HPLC afforded 12 as the major product (61%) with isolation of 14 as the minor isomer in 17% yield. The HPLC-based purification step even allowed for further separation of the diastereomers on phosphorus of both compounds.

Selective methyl ester cleavage of 12 and 14 - again using NaI in a minimum volume of acetone – was straightforward and furnished the octyl phosphonate ester derivatives 13 and 15 in good yields. Cleavage of the benzyl protecting groups with concomitant reduction of the azido group afforded the $\alpha$-anomeric octyl 4-amino-4-deoxy-L-arabinopyranosyl-methylphosphonate 16 in 38% yield. Deprotection of the
endo-glycal ester 15 was also investigated. An intermediate enol resulting from hydrogenolytic cleavage of the benzyl ether was envisaged to be prone to facile formation of tautomers which would then lead to different reduction products. Notably, however, the reduction proved to be highly selective, and hence might involve a fast hydrogen addition from the bottom face of the pyranose ring. After full hydrogenation and azide to amine conversion, compound 17 was isolated as the 4-amino-4-deoxy-L-ribopyranosyl derivative. The configuration was determined from the NMR characteristics. H-2 at 4.02 ppm appeared as broadened doublet with small homonuclear coupling constants as would be expected for a manno-type spin system. In addition NOesy correlations were observed from H-3 to H-5ax and the anomeric proton which is consistent with the equatorial position of the C-phosphonate entity.

Conclusion

In conclusion, Wittig-type reactions of the suitably protected arabinonic lactone allow for a straightforward implementation of a phosphonate dimethyl ester, that can be readily converted into exo- and endo-glycal ester derivatives. The selective cleavage of one of the methyl groups opens various options for introduction of a separate alkyl chain, since the remaining methyl group is amenable for a selective ester cleavage at a later stage. The glycal ester derivatives themselves offer additional options towards the preparation of transition state analogues but can also be fully deprotected to provide hydrolytically stable substrate derivatives in the correct anomeric configuration.
Supporting Information

Supporting Information File 1: Experimental section, spectral data and copies of $^1$H, $^{13}$C and $^{31}$P NMR spectra of compounds 4, 6, 8-17.

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