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Stereoselective Synthesis of 2-Deoxy-β-Glycosides Using Anomeric O-Alkylation/Arylation

William J. Morris and Matthew D. Shair*

Department of Chemistry and Chemical Biology, Harvard University
12 Oxford Street, Cambridge, Massachusetts 02138
shair@chemistry.harvard.edu

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ABSTRACT

Anomeric O-alkylation/arylation is applied to the synthesis of 2-deoxy-β-glycosides. Treatment of lactols with NaH in dioxane followed by the addition of electrophiles leads to the formation of 2-deoxy-β-glycosides in high yield and high selectivity. The high β-selectivity observed here demonstrates a powerful stereoelectronic effect for the stereoselective formation of acetics under kinetic control.

2-Deoxy-β-glycosides are present in biologically active natural products such as the lomaiviticins, olivomycin A, OSW-1, and durhamycin. The stereoselective preparation of 2-deoxy-β-glycosides is difficult because substituents at C2 often serve as directing groups during the glycosylation event. The synthesis challenge posed by 2-deoxy-β-glycosides coupled with their presence in nature has inspired a variety of approaches aimed at accessing these important glycosides. The most common methods involve the use of a heterocycle or dinitrogen at C2 of the glycosyl donor followed by its reductive removal after glycosylation. Other methods include the use of α-glycosyl phosphites, displacement of α-glycosyl halides, palladium catalyzed glycosylation reactions, utilization of alkoxy-substituted anomeric radicals, and the use of glycosyl imidates as glycosyl donors under oxidative conditions.


We became interested in the synthesis of 2-deoxy-β-glycosides since they are present in the lomaiviticins, molecules we are targeting for synthesis (Figure 1). In particular, our recent synthesis of the central ring system of the lomaiviticins involves incorporation of the C4 and C4′ carbinols by S₈ displacement of allylic sulfones with methoxide anions. This raised the possibility that the 2-deoxy-β-glycosides at C4 and C4′ might be incorporated via an anomic O-alkylation using glycosyl-1-aloxides.

Anomic O-alkylation with glycosyl-1-alkoxides generally affords high levels of β-glycosides. An explanation for this selectivity involves rapid equilibrium between axial and equatorial alkoxides with the enhanced nucleophilicity of the equatorial alkoxide leading to selective generation of β-glycosides (Scheme 1). It has been proposed that the enhanced nucleophilicity of the equatorial alkoxide (II), compared to the axial alkoxide (I), is due to increased electron-electron repulsion resulting from the alkoxide of II being gauche to both electron lone pairs of the ring oxygen compared to a single gauche interaction in I. This phenomenon has been referred to as the kinetic anomic effect, or the β-effect, and may be similar to the α-effect, which is observed in molecules where the nucleophilic atom is directly attached to another heteroatom. To date, most examples of anomic O-alkylation have been performed with substituents in the C2 position. This has made it difficult to assess the contribution of the β-effect versus steric (or electronic) influences of C2 substituents in the selective formation of β-glycosides by anomic O-alkylation. In this paper we report the first examples of anomic O-alkylation/arylation in the absence of a C2 substituent, stereoselectively forming 2-deoxy-β-glycosides.

**Scheme 1. Kinetic Anomeric Effect**

Before beginning our investigation into anomic O-alkylation/arylation, a synthesis of a protected form of the N,N-dimethylpyrolosamine sugar of lomaivitin B was accomplished (Scheme 2). Beginning from the known glycal 1, triflation was followed by displacement with NaN₃. The azido-glycal 2 was hydrated in a 2-step procedure involving addition of AcOH to catalyzed by triphenylphosphine hydrogen bromide and cleavage of the resulting anomic acetate with Me₃SiH. Lactol 3 was isolated as a 4:1 (α:β) mixture of anomers.

Our next goal was to establish general reaction conditions to access 2-deoxy-β-glycosides directly from an anomic mixture of lactols (Table 1). During the course of our optimization studies we found that treatment of lactol 3 with KHMDS in THF at –78 °C followed by the addition of allyl bromide led to the exclusive formation of the α-anomer (Entry 1). Changing the base to NaH and raising the reaction temperature to 0 °C led to a 1:1 mixture of α and β anomers. The addition of LiBr, an additive known to enhance β-selectivity in anomic O-alkylation provided only minor improvements in selectivity (Entry 3). Gratifyingly, we found that changing the solvent from

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**Scheme 2. Synthesis of protected N,N-dimethylpyrolosamine**

**Figure 1. Lomaivitin B**

![Figure 1. Lomaivitin B](image-url)
DMF to dioxane and increasing the reaction temperature to 23 °C led to a dramatic increase in selectivity. These conditions afforded the 2-deoxy-β-glycoside exclusively in 91% yield.

With optimal reaction conditions in hand, the scope of this reaction was evaluated (Table 2). We considered nucleophilic aromatic substitution as a valuable application of this methodology due to its potential use in the synthesis of the aureolic acid family of natural products. Towards that end, when lactol 3 was treated with NaH in dioxane followed by 1-bromo-2,4-dinitrobenzene, the 2-deoxy-β-glycoside product 8 was isolated as an 18:1 mixture of anomers favoring the β-anomer. Primary triflates such as 9αβ proved to be suitable electrophiles for anomic O-alkylation, as disaccharide 10 was isolated in 90% yield exclusively as the β-diastereomer. Following the successful anomic O-alkylation of a C6 triflate, we determined if secondary triflates were competent electrophiles for this reaction. Despite previous reports by Schmidt of anomic O-alkylations on similar systems,17 triflates 11 and 12 failed to undergo displacement (Entries 4 & 5). In each case, addition of the alkoxide took place at sulfur resulting in detriflation of 11 and 12.

Table 1. Optimization of Reaction Conditions

<table>
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<tr>
<th>Entry</th>
<th>conditions</th>
<th>entries</th>
<th>α:β[α]</th>
<th>yield (%)</th>
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<tbody>
<tr>
<td>1</td>
<td>KHMD, THF, −78 °C, allyl bromide</td>
<td>95:5</td>
<td>91% (20:1)</td>
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<tr>
<td>2</td>
<td>NaH, DMF, allyl bromide, 0 °C</td>
<td>1:1</td>
<td>87% (18:1)</td>
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</tr>
<tr>
<td>3</td>
<td>NaH, dioxane, allyl bromide, 0 °C</td>
<td>1:1.5</td>
<td>87% (18:1)</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>NaH, dioxane, allyl bromide, 23 °C</td>
<td>5:95</td>
<td>87% (18:1)</td>
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* Reactions performed at 0.1 M. † Isolated yield after chromatographic purification. ‡ As determined by 1H NMR (600 MHz)

Table 2. Electrophile Scope

<table>
<thead>
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<th>electrophile</th>
<th>2-deoxy-β-glycoside</th>
<th>yield (%)</th>
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<tr>
<td>1</td>
<td>[6]</td>
<td>91% (20:1)</td>
</tr>
<tr>
<td>2</td>
<td>[7]</td>
<td>87% (18:1)</td>
</tr>
<tr>
<td>3</td>
<td>[8]</td>
<td>90% (20:1)</td>
</tr>
<tr>
<td>4</td>
<td>[9]</td>
<td>n/a</td>
</tr>
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Table 3. Lactol and Electrophile Scope

18 (18) This methodology has been applied in an iterative manner to access polysaccharides. See Supporting Information.
We next devised a competition experiment between lactol 3 and trans-4-tert-butylcyclohexanol\(^{20}\) to determine whether the increased reactivity of equatorial alkoxides derived from lactols (II, Scheme 1) can be attributed to steric effects or the aforementioned \(\beta\)-effect. An equimolar mixture of lactol 3 and trans-4-tert-butylcyclohexanol (26) was dissolved in dioxane and treated with 2 equivalents of NaH (Scheme 3). After 10 minutes, 1 equivalent of allyl bromide was added to the reaction mixture. After workup, \(^1\)H NMR of the crude reaction mixture showed only 2 products: 2-deoxy-\(\beta\)-glycoside 5 and unreacted trans-4-tert-butylcyclohexanol 26. The mixture was purified by flash column chromatography yielding 81% of 5 and quantitative recovery of trans-4-tert-butylcyclohexanol.

This experiment is the best support to date that the high \(\beta\)-selectivity in anomic O-alkylations/arylations is due to the increased nucleophilicity of the \(\beta\)-configured anomic alkoxides.

In conclusion, we report that anomic O-alkylation/arylation can be used to form 2-deoxy-\(\beta\)-glycosides with high stereoselectivity. The high \(\beta\)-selectivity afforded with 2-deoxy-1-lactols supports the theory that the \(\beta\)-effect, rather than the substituent at C2, is the stereocontrol element in anomic O-alkylations/arylations. This conclusion has been obscured in previous reports concerning anomic O-alkylations/arylations since they all involved substrates with C2 substituents which may have influenced the stereoselectivity. The competition experiment reported here demonstrates that the nucleophilicity of \(\beta\)-configured anomic alkoxides is enhanced over similar cyclohexyl alkoxides, presumably due to the proximity of the alkoxide and the lone pair electrons of the ring oxygen. These results suggest that the \(\beta\)-effect is a powerful stereoelectronic effect that may be useful in designing other stereoselective reactions.

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**Supporting Information Available:** Experimental procedures and spectral data for all new compounds. This material is available free of charge via the Internet at http://pubs.acs.org.

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**Scheme 3. Competition Experiment**

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