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From 1,4-Disaccharide to 1,3-Glycosyl-Carbasugar: Synthesis of a Bespoke Inhibitor of Family GH99 Endo- α -Mannosidase

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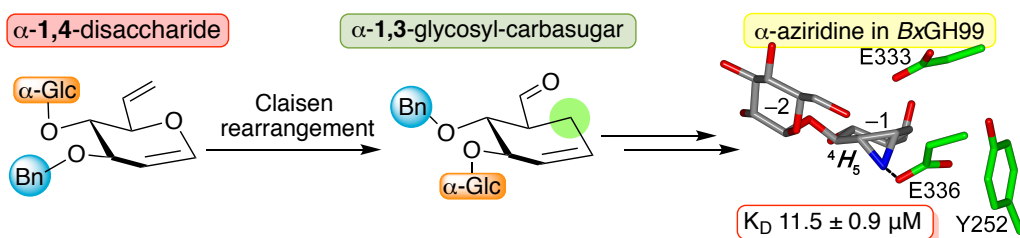
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Supporting Information Placeholder



ABSTRACT: Understanding of enzyme reaction mechanism can lead to the design of enzyme inhibitors. A Claisen rearrangement was used to allow the conversion of an α -1,4 disaccharide into an α -1,3-linked glycosylcarbasugar to target the endo- α -mannosidase from glycosidase family GH99 which, unusually, is believed to act through a 1,2 anhydrosugar “epoxide” intermediate. Using NMR and X-ray crystallography, it is shown that glucosylcarbasugar- α -aziridines can act as reasonably potent endo- α -mannosidase inhibitors, likely by virtue of their shape-mimicry and the interactions of the aziridine nitrogen with the conserved catalytic acid/base of the enzyme active-site.

Understanding enzyme reaction mechanisms is essential for the rational design of inhibitors. Glycoside hydrolases (GH), also called glycosidases, are a vast group of enzymes that are classified, based upon their amino-acid sequence similarity, into over 150 distinct families in the Carbohydrate Active Enzymes database (CAZy).^{1,2} The vast majority of these enzymes can be divided into two classes according to their reaction mechanism and the relative configuration of substrate and product: those that act with retention or inversion of anomeric configuration. The retaining glycosidases utilize a double nucleophilic displacement which results in net retention of configuration at the anomeric center. The retaining reaction classically involves nucleophilic amino-acid side-chain and proceeds via a covalent glycosyl-enzyme intermediate. Some enzymes also use neighboring group participation reactions (notably those active on beta linked GlcNAc).³ Recently, a new neigh-

bouring-group retaining mechanism involving an intermediate epoxide and hence no covalent bond to the enzyme was proposed for endo- α -mannosidases and endomannanases (α -Glc-1,3- α -Man-OR and α -Man-1,3- α -Man-OR substrates) of the GH99 family.^{4,5} In this putative mechanism, nucleophilic attack by the manno-O2 atom, allows catalysis via a 1,2 anhydro sugar “epoxide” intermediate. (Scheme 1). In terms of shape and charge changes, a positive charge is very likely to be developed and the conformation of the pyranose changes from 4C_1 of the substrate to the ${}^4E/{}^4H_5$ of the epoxide. Based on these considerations, inhibitors that have been designed for this important class of enzymes⁶ are α -1,3-disaccharide analogues and these fall into two categories: those which mimic the charge and those which mimic its shape. In the first class, we find the usual deoxymannojirimycin (1), isofagomine (2) or neuromycin derivatives.^{4,7} A clear influence of the amine

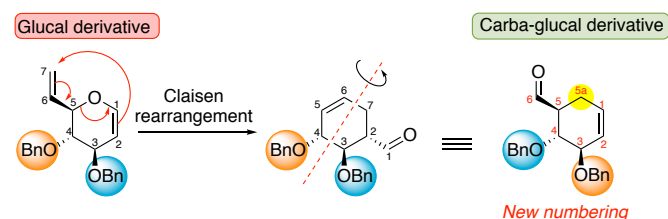
position has been demonstrated. It has also been shown that the glucal derivative **3** is an inhibitor of the enzyme due to its shape, which corresponds to a 4H_5 conformation.⁷ Here, we decided to combine both properties in a new inhibitor and create an amine with a likely ${}^4E/{}^4H_5$ conformation. To achieve both the half-chair conformation and introduce an amino group, we decided to synthesize an aziridine on a carbasugar for obvious stability reason. Furthermore, according to the mechanism, the catalytic acid E336 of *Bacteroides xylanisolvens* GH99 (*BxGH99*) should be in its acidic form and therefore prone to release a proton to the aziridine, hence we decided to investigate the synthesis and inhibitory ability of glucosyl-carbasugar- α -aziridine **4**. (Scheme 1)

There are many ways to synthesize carbasugars,⁸ including those containing a disaccharide motif.⁹ The most direct way would be to start directly from the corresponding disaccharide,¹⁰ but the necessary 1,3-disaccharides (like Glc- α -1,3-Glc) are rare and expensive. The synthesis of the two monosaccharides and their subsequent assembly seemed tedious. We therefore decided to use a trick based on the use of Claisen rearrangement to make carbasugars nicely developed by Nagarajan on monosaccharides.¹¹ In this reaction, a glucal functionalized with an alkene in position 6 affords

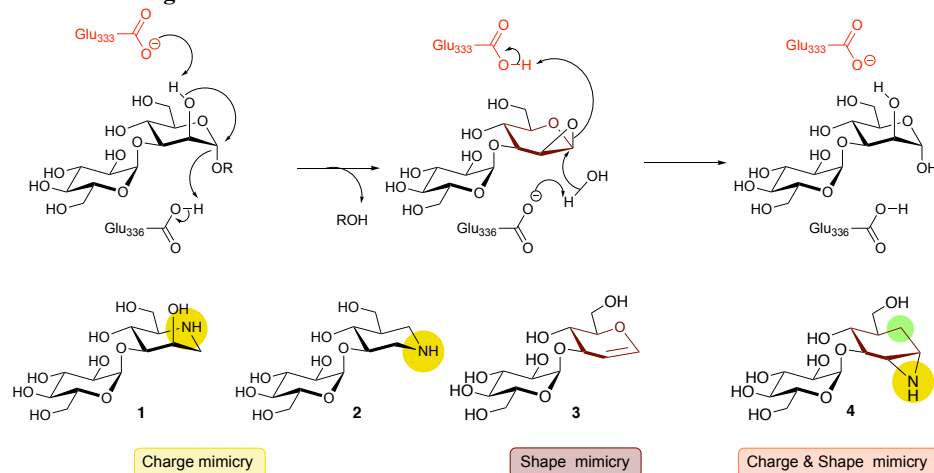
a carba-glucal with an aldehyde in position 6. This is due to the symmetry of the molecule, and in the operation, the protecting group in position 4 becomes the one in position 3 in the new numbering of the carbasugar. (Scheme 2)

This approach was used to synthesize a 1,3-pseudo-disaccharide from a 1,4-disaccharide. The synthesis would start from maltose, that would be converted into the doubly unsaturated maltal **5**, which in turn could undergo Claisen rearrangement to give keto-alkene **6**. This operation is key to the total synthesis as we not only obtain the desired 1,3-pseudo-disaccharide but also the alkene which will be converted into the desired epoxide **7** and aziridine **4**. (Scheme 3)

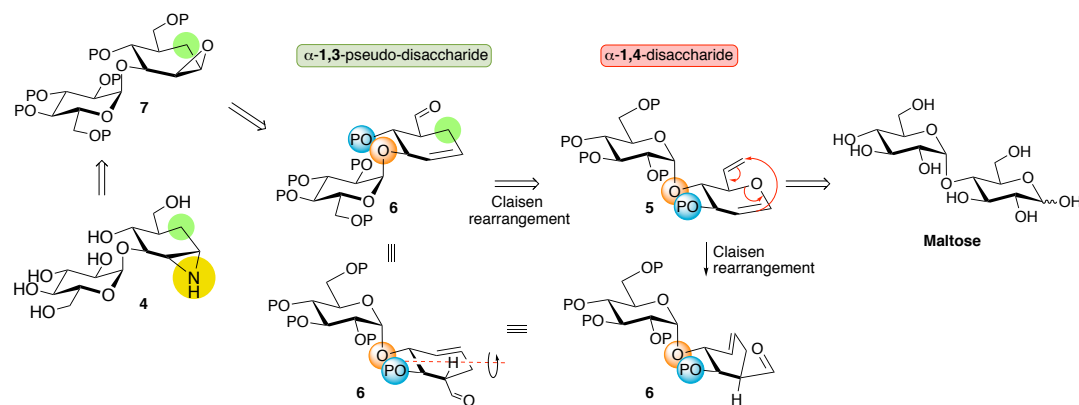
Scheme 2. Claisen rearrangement leading to interchange of position 3 and 4 from the glucal to the carba-glucal.



Scheme 1. Postulated enzymatic hydrolytic mechanism for *Bacteroides xylanisolvens* GH99 (*BxGH99*), in which catalysis occurs through a 1,2 anhydrosugar intermediate (top). Known, and designed (this work) substrate analogues (below) Residue numbering as in *BxGH99*.



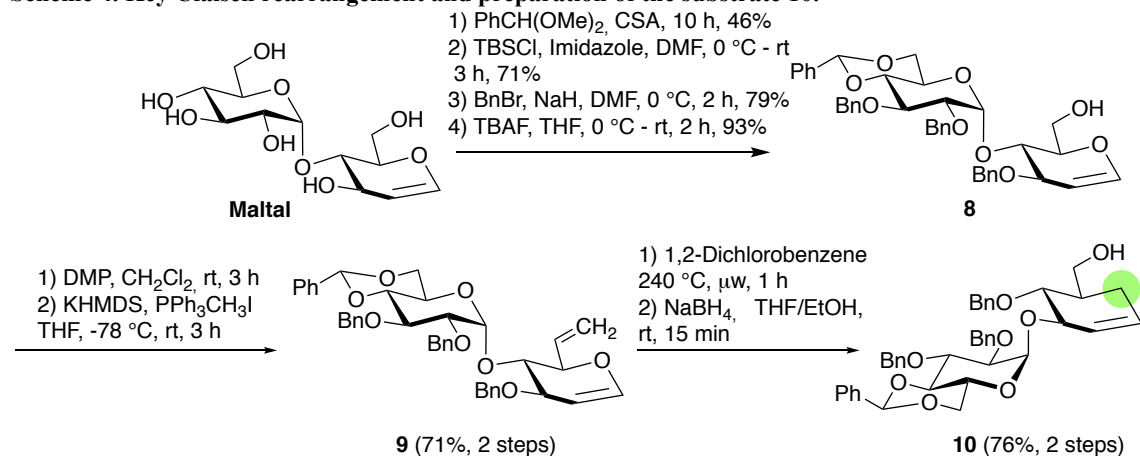
Scheme 3. Retrosynthetic analysis for the synthesis of aziridine **4.**



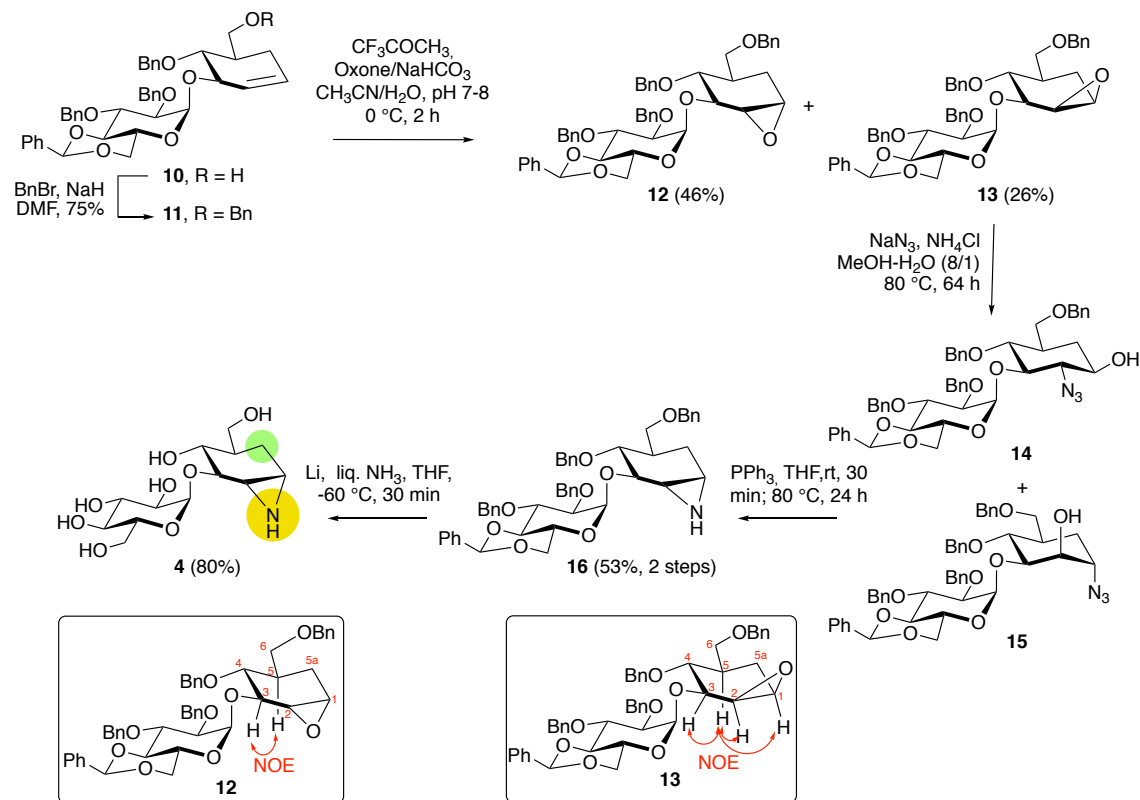
Maltol was prepared according to Haworth,¹² with a slight modification in the deprotection step (see SI). Next, the differentiation of the two primary hydroxyl groups was required. For this, we decided to protect the glucoside with a benzylidene group. This reaction had to be performed with great care (Scheme 4) by controlling the amount of camphor sulfonic acid used (0.1 equiv). Then, the other

primary alcohol could be selectively silylated, subsequent benzylation and de-silylation afforded the alcohol **8**. Oxidation using Dess-Martin periodinane (DMP) followed by olefination produced dialkene **9**. Claisen rearrangement on **9** was performed under micro-wave irradiation at 240 °C and was immediately followed by the aldehyde reduction to afford alcohol **10** in 76% yield. (Scheme 4)

Scheme 4. Key Claisen rearrangement and preparation of the substrate 10.



Scheme 5. Synthesis of aziridine carbasugars 4.



With the alkene **10** in hand, we investigated its aziridination. We opted for a three-step route: epoxidation, azide-

ring opening, and Staudinger azide-reduction-aziridination;¹³ this strategy also operates with overall inversion of configuration of the three-membered ring. We first benzylation the remaining hydroxyl in **10** to give

11, which was epoxidized using methyl(trifluoromethyl)dioxirane generated *in situ*¹⁴ to give the α -epoxide **12** in 46% yield and the β -epoxide **13** in 26% yield, which were separated by silica gel flash chromatography. The stereochemistry of these epoxides was determined thanks to a NOESY experiment. α -Epoxide **12** displayed cross correlations between H-5 and H-3, while in β -epoxide **13** H-5 showed cross-correlations with H-3 but also with H-1 and H-2 (Scheme 5 and SI). This reaction therefore produced the undesired α -epoxide as the major product, but all our attempts to improve the situation with various epoxidation reactions gave lower yields. For instance, mCPBA which was previously often used to synthesize epoxides from cyclohexenes polyols¹⁵ did not improve the selectivity nor the yield. Further treatment of the isolated epoxide **13** with sodium azide furnished a mixture of 1,2-*trans* azido alcohols **14/15**, which under the action of PPh₃ afforded α -aziridine **16** in 52% yield after these two steps. Final deprotection, using Birch conditions,¹⁶ afforded the unprotected aziridinated pseudo-sugar **4** (Scheme 5).

The properties of aziridine **4** were investigated with GH99 family members *Bt* (*Bacteroides thetaiotaomicron*) and *Bx*GH99. A crystal structure of the *Bx*GH99 enzyme in complex with the aziridine **4** at a resolution of 1.27 Å (PDB accession: **6HMG**) was obtained by soaking of

*Bx*GH99 crystals with an aqueous solution of the compound. The nitrogen atom of the aziridine makes a 2.6 Å hydrogen bond with one of the side chain oxygen atoms of *Bx*GH99 residue E336, which acts as an acid/base in the proposed GH99 mechanism (Scheme 1). In a ternary complex with the aziridine and α -1,2-mannobiose (obtained at a resolution of 1.03 Å, PDB **6HMH**), this distance changes to 2.7 Å. The interaction is shown in Figure 1A. In both cases, the pseudopyranose ring is in a ⁴H₅ conformation, as predicted for the aziridine and likely resembles the intermediate/transition-state of the reaction. Indeed, it binds similarly to ManGlucal and GlcChex¹⁷ (Figure 1B), which was also observed in a ⁴H₅ conformation in complex with *Bx*GH99 (PDB **5M5D** and **5MC8**).⁷

Using NMR we have determined the dissociation constant (K_D) of the aziridine with ¹⁵N-labelled *Bt*GH99 to be $11.5 \pm 0.9 \mu\text{M}$ (Figure 1C). The compound, therefore, binds to *Bt*GH99 with an affinity 2-fold higher than GlcDMJ (24 μM) and slightly higher than ManGlucal (15 μM). However, it is not a potent inhibitor in comparison with GlcIFG, ManIFG or ManNOE, all of which bind to *Bt*GH99 with a K_D in the nanomolar range.⁷

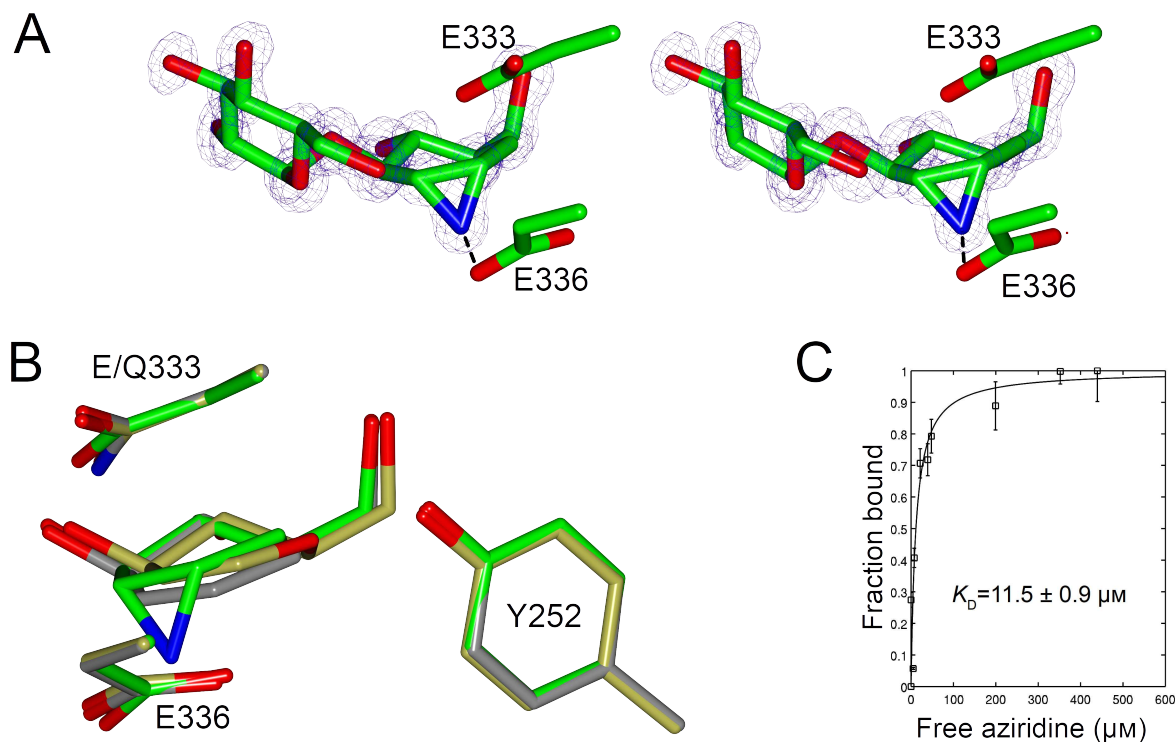


Figure 1. (A) Divergent stereo (“wall-eyed”) view of **4** in complex with *Bx*GH99. Catalytic residues surrounding the –1 subsite and the interaction between the aziridine N and E336 O_ε are shown. The $2mF_o - DF_c$ synthesis contoured at $1 e/\text{Å}^3$. (B) Comparison of the ligands observed in the ⁴H₅ conformation in complex with *Bx*GH99.⁷ Green: the aziridine **4**, gold: ManGlucal (from PDB **5M5D**), grey: GlcChex (from PDB **5MEL**). The sugar moiety in the –2 subsite is not shown. (C) The result of NMR titration of *Bt*GH99 with **4**. The protein concentration in the solution was 58 μM .

In conclusion, we have devised a strategy to synthesize 1,3-glycosyl-carbasugars designed to interact with the family GH99 endo- α -mannosidases. The obtained inhibitor has improved activity in comparison with the shape mimics, but does not reach the affinity of isofagomine or noeuromycin analogues. This probably indicates that the aziridine is not charged owing to its lower pKa. In future work, we will be able to use this synthetic strategy to synthesize other analogues to study the mechanism of this family of glycosidases which is not yet firmly established.

ASSOCIATED CONTENT

Supporting Information

Experimental details, spectra and X-ray crystallography. The Supporting Information is available free of charge on the ACS Publications website.

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Author Contributions

All authors have given approval to the final version of the manuscript. / ‡These authors contributed equally.

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