Expanding the Thioglycoligase Strategy to the Synthesis of \( \alpha \)-Linked Thioglycosides Allows Structural Investigation of the Parent Enzyme/Substrate Complex

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The \( \alpha \)-retaining glycosidases of family 31 include pharmaceutically important human enzymes, such as the human lysosomal \( \alpha \)-glucosidase (GAA), endoplasmic reticulum glucosidase II, and sucrase isomaltase (SUIS). The first structural analysis of a family 31 enzyme was that on the \( \alpha \)-xylosidase (EC 3.2.1.) from \( E\). coli (YicI), thereby providing a useful structural framework for the understanding of the enzymes from higher order organisms. However, delineation of the substrate specificities and mechanism requires structural information on the enzyme—substrate complex. Nonhydrolyzable substrate analogues, such as thioglycosides, are useful probes for such studies due to their resistance to chemical or enzymatic hydrolysis and their similar conformation to that of their natural O-linked counterparts. Thioglycosidases are recently developed mutants of retaining glycosidases in which the acid/base carboxylic acid residue has been replaced by an inert amino acid. These mutants catalyze the formation of S-glycosidically linked oligosaccharides by coupling of acceptors bearing a suitably leaving group, such as dinitrophenol or fluoride (Figure 1). However, no \( \alpha \)-thioglycosidases have yet been reported, raising the question of whether such mutants will function as described. This report describes the generation of the first \( \alpha \)-thioglycosidases, the use of one such mutant to form an inhibitor of the wild-type enzyme, and elucidation of the three-dimensional structure of a YicI/xthioglycosidase complex.

Two family 31 glycosidases, an \( \alpha \)-xylosidase from \( E\). coli (YicI) and an \( \alpha \)-glucosidase from \( S\). solfataricus (MalA), were selected for investigation as potential \( \alpha \)-thioglycosidases. Previous kinetic studies have revealed a preference for YicI for the \( \alpha \)-(1,6)-xylosidic linkage over the \( \alpha \)-(1,4)-glucosidic linkage. By contrast, \( \alpha \)-glucosidases, such as MalA, prefer \( \alpha \)-(1,4)-glucosides.

The acid/base catalysts of YicI (Asp482) and MalA (Asp416) were mutated to alanine using the mega primer method, and the variants were purified by Ni+NTA affinity chromatography (see Supporting Information). The thioglycoligase reactions catalyzed by YicId482A and MalA D416A were first investigated using pNP 4-deoxy-4-thio-\( \beta \)-d-glucopyranoside as acceptor and an equimolar amount of the glycosyl donor, \( \alpha \)-d-xylopyranosyl fluoride for YicI D482A, and \( \alpha \)-d-glucopyranosyl fluoride for MalA D416A. YicI D482A catalyzed the thioglycoligase reaction more efficiently than did MalA D416A, the thio-linked disaccharides, 1 and 2, being formed in 79 and 48% yields, respectively. Thioglycoligase reactions using pNP 4-deoxy-4-thio-\( \beta \)-d-xylopyranoside as acceptor were less efficient than those with the gluco-version, MalA D416A not accepting 4-thioglycoside at all, while YicI D482A yielded 3 in 41% yield. Analysis of the isolated products after acetylation by \( ^{1} H \) and \( ^{13} C \) NMR and mass spectrometry confirms these structures.

Given the preference of YicI for \( \alpha \)-(1,6)-linkages, our interest moved to the synthesis of \( \alpha \)-1,6-thioglycosidic linkages using these thioglycoligases. Using pNP 6-deoxy-6-thio-\( \beta \)-d-glucopyranoside as acceptor, YicI D482A generated a single product 4 in 86% yield. MalA D416A, consistent with the specificity of its parent enzyme, gave no product.

Next, the thiosaccharides, 1, 3, and 4, were tested as potential inhibitors for YicI. Compound 4, which differs from isoprimeverose (xylopyranosyl-\( \alpha \)-(1,6)-glucopyranose), the natural substrate of YicI, only in the nature of the intersugar linkage where a sulfur atom replaces an oxygen atom, acted as a potent competitive inhibitor with a \( K_{i} \) value of 2 \( \mu M \). \( \alpha \)-(1,4)-Linked thiosugars, 1 and 3, also inhibited YicI well with \( K_{i} \) values of 27 and 14 \( \mu M \), respectively.

The three-dimensional structure of YicI in complex with 4 was determined by X-ray crystallography at 1.95 \( \AA \) resolution. Six active sites in the hexameric quaternary structure of YicI are fully occupied by 4 (see Supporting Information). A seventh molecule of 4 is found between the N-terminal end of the \( \alpha \beta \) barrel and the C-terminal domain of monomer F. This may be an artifact of crystal packing as it is only observed for the one monomer, but it is intriguing to speculate that carbohydrate-binding motifs may exist outside of the active site. The electron density is clear for the ligands, with xylolide moieties in the \(-1\) subsites being in the 4 \( C_{1} \) chair conformation, consistent with the conformational itinerary (\( C_{1} \) \( \rightarrow \) \( H_{1} \) \( \rightarrow \) \( S_{2} \)) previously proposed. The Michaelis complex structure reveals the +1 subsite and suggests the role of the N-terminus of YicI (N-domain) in the formation of the aglycon sites (Figure 2A). The +1 subsite of each monomer in the hexameric unit consists of four residues from its own \( \alpha \beta \) barrel (Trp380, Phe417, Arg466, and Phe515) and three residues from the N-domains of three different monomers: Asp815 from its own monomer (black in Figure 2A), Asp49 from an adjacent monomer (orange), and Trp8 from a second distinct monomer (pink). Importantly, the hexameric structure has also been demonstrated in solution. Although Asp815,
Trp380, and Arg466 are conserved in most family 31 glycosidases. Trp8 and Asp49 are conserved only in α-xylosidases that are highly homologous to YicI. Consequently, the N-domain-mediated hexamer structure of YicI constructs an efficient 2C). However, both GAA and SUIS are expected to have space around O-6- of the fructose (Figure 2B). In addition, Asp511 of YicI interacts with Arg538, which corresponds to Arg672 of GAA, another severe mutation (-1 subsite as the basis for 1 subsite in the Michaelis complex of its parent enzyme has been determined, as has been proposed for Arg227 and His327 of GAA that cause Pompe’s disease, G643R (Ser509 in 1 subsite would clash with Trp380 (Figure 1B). Therefore, Arg466 appears to stabilize the partial negative charge that accumulates on O-2 of the sugar at the transition state, much as has been proposed for Arg227 and His327. In conclusion, the generation of α-thioglycosides has been successfully achieved, for the first time, using the thioglycoligase strategy. Using the product of one of these thioglycoligases as a nonhydrolyzable substrate analogue, the three-dimensional structure of the Michaelis complex of its parent enzyme has been determined, providing valuable structural insights into the specificity and mechanism of family 31 α-glycosidases and the molecular basis of an associated genetic disease. This general strategy could be applied to other glycosidases.

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Supporting Information Available: Experimental procedures, NMR data, crystallization, data collection, additional structure figures, and complete refs 7 and 8. This material is available free of charge via the Internet at http://pubs.acs.org.

References


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Figure 2. (A) Interactions of YicI with compound 4. Green dotted lines and numbers represent H-bonds and their lengths. Black dotted lines and numbers represent distances between 4 and residues within 4Å.

(A) Interactions of YicI with compound 4. Green dotted lines and numbers represent H-bonds and their lengths. Black dotted lines and numbers represent distances between 4 and residues within 4Å. (B) Superposition of the Michaelis complex (2F2H) and the 5-fluoroxyllosyl (eq5FX) enzyme intermediate (1XSK). Site for Pompe’s disease.8 Therefore, the mutations mentioned above which corresponds to Arg672 of GAA, another severe mutation 4 complex have been deposited in the Protein Data Bank with accession code 2F2H.

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