Structural insights into the carbohydrate binding ability of an alpha-(1 -> 2) branching sucrase from glycoside hydrolase family 70
Yoann Brison, Yannick Malbert, Georges Czaplicki, Lionel Mourey, Magali Remaud Simeon, Samuel Tranier

To cite this version:
Yoann Brison, Yannick Malbert, Georges Czaplicki, Lionel Mourey, Magali Remaud Simeon, et al.. Structural insights into the carbohydrate binding ability of an alpha-(1 -> 2) branching sucrase from glycoside hydrolase family 70. Journal of Biological Chemistry, American Society for Biochemistry and Molecular Biology, 2016, 291 (14), pp.7527-7540. 10.1074/jbc.M115.688796. hal-01886416

HAL Id: hal-01886416
https://hal.archives-ouvertes.fr/hal-01886416
Submitted on 20 Jun 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
Structural Insights into the Carbohydrate Binding Ability of an $\alpha$-(1→2) Branching Sucrase from Glycoside Hydrolase Family 70*

Received for publication, September 9, 2015, and in revised form, February 7, 2016. Published, JBC Papers in Press, February 10, 2016, DOI 10.1074/jbc.M115.688796

Yoann Brison§†, Yannick Malbert§‡‡, Georges Czaplicki§‡‡, Lionel Mourey§‡‡, Magali Remaud-Simeon§¶‡‡, and Samuel Tranier§¶‡‡‡

From the§Université de Toulouse, INSA, UPS, INP, LISBP, 135 avenue de Rangueil, F-31077 Toulouse, the§INRA, UMR792 Ingénierie des Systèmes Biologiques et des Procédés, F-31400 Toulouse, the†CNRS, UMR5504, F-31400 Toulouse, the‡Institut de Pharmacologie et de Biologie Structurale (IPBS), Centre National de la Recherche Scientifique (CNRS), 205 route de Narbonne, BP 64182, F-31077, Toulouse, and the¶Université de Toulouse, Université Paul Sabatier, IPBS, F-31077 Toulouse, France

The $\alpha$-(1→2) branching sucrase $\Delta$N$_{123}$-GBD-CD2 is a trans-glucosylase belonging to glycoside hydrolase family 70 (GH70) that catalyzes the transfer of D-glucosyl units from sucrose to dextrins or gluco-oligosaccharides via the formation of $\alpha$-(1→2) glucosidic linkages. The first structures of $\Delta$N$_{123}$-GBD-CD2 in complex with D-glucose, isomalto-, or isomaltotriosyl residues were solved. The glucose complex revealed three glucose-binding sites in the catalytic gorge and six additional binding sites at the surface of domains B, IV, and V. One aromatic residue is systematically identified at the bottom of these pockets in stacking interaction with one glucosyl moiety. The carbohydrate is also maintained by a network of hydrogen bonds and van der Waals interactions. The sequence of these binding pockets is conserved and repeatedly present in domain V of several GH70 glucansucrases known to bind $\alpha$-glucans. These findings provide the first structural evidence of the molecular interaction occurring between isomalto-oligosaccharides and domain V of the GH70 enzymes.

The glucosyltransferases of glycoside hydrolase family 70 (GH70),§ namely the glucansucrases, the 4,6-$\alpha$-glucanotransferases, and the branching sucrases, are $\alpha$-retaining transglycosylases produced by various lactic acid bacteria from Leuconostoc, Streptococcus, Weissella, and Lactobacillus genera (1–3). Glucansucrases from Leuconostoc spp. have been mainly studied for their industrial applications in the production of dextrins or prebiotic gluco-oligosaccharides (4–7), whereas investigations on streptococcal glucansucrases were mostly motivated by their involvement in the accumulation of streptococci on tooth enamel and subsequent human dental caries formation (8). Glucansucrases catalyze the transfer of $\alpha$-D-glucopyranosyl moieties from sucrose to acceptor molecules through the formation of a $\beta$-D-glucosyl-enzyme intermediate. They synthesize $\alpha$-glucans, which vary in terms of size, osidic linkages as well as degree and arrangement of branching. Devoid of polymerase activity, the $\alpha$-(1→2) branching sucrases are specialized in dextran branching through the formation of $\alpha$-(1→2) glucosidic linkages from sucrose to dextran acceptor (Fig. 1) (9, 10). In the absence of acceptor, these enzymes essentially catalyze sucrose hydrolysis.

Glucansucrases are also known to bind dextrins via a carbohydrate-binding domain remote from the active site (2, 11). This domain is referred to as the glucan-binding domain (GBD) and is not classified in the CAZy database. Carbohydrate-protein interactions often play critical roles in the catalytic activities of carbohydrate active enzymes involved in polysaccharide synthesis, degradation, or decoration. Carbohydrate-binding sites may be intrinsic components of catalytic sites, but may also exert essential functions when located in non-catalytic modules (12). Evidence of the presence of non-catalytic domains showing $\alpha$-glucan binding ability in GH70 enzymes were first provided by analyzing dextran interactions with non-catalytic polypeptides isolated from trypsin digestion of various streptococcal glucansucrases (13). Based on sequence and biochemical analyses, the streptococcal glucan binding regions were shown to consist of a series of repeats that were classified into four different groups (A, B, C, and D) (14–19). These repeats were proposed to have evolved from a common motif, the so called YG repeat, characterized by conserved aromatic amino acids and glycine ((N/X)(D/E)(X)(G/Y)ψβφxxxxψψx(G/N/L)(X)(Ψ/Ψ)(Ψ/Ψ), where $\psi$ represents a non-conserved residue, $\Psi$ a hydrophobic residue, and $\phi$ an aromatic residue) for a full description, see Ref. 20.

They were also identified at the N-terminal or C-terminal domains of glucansucrases from Leuconostoc spp. and shown to share homology with the cell wall binding repeats of choline-
Carbohydrate Binding Ability of a GH70 Branching Sucrase

binding proteins or toxins (Pfam CW_binding_1 (PF01473)) (21–23). In particular, the five repeats found at the C-terminal extremity of the GBD of DsrP glucansucrase are capable of binding the cell surface of Leuconostoc cells (24). Mutagenesis of conserved aromatic or basic residues of the repeats found in Streptococcus downei GtfI was performed and led to a reduction of the glucan binding capability (25). The elimination of some repeats by truncation was also shown to influence transglycosylation, modify glucan binding properties, and affect the polymer size (26–29). In particular, the GBD was suggested to provide a glucan anchoring platform, which may affect the single-chain elongation in a semi-processive manner (29). A few studies focused on isolated GBDs disconnected from their catalytic domain, and further demonstrated that truncated forms can autonomously fold and bind α-glucans with very low $K_d$ values estimated in the nanomolar range (30–34).

The crystal structures of several GH70 enzymes truncated at their N-terminal ends shed light on the GBD-fold, although no structures with entire GBDs have been solved yet (9, 35–37). These enzymes follow a U-path course and are organized in five domains, A, B, C, IV, and V. Domains A, B, and C are related to their counterparts found in GH13 α-amylases. In the GH70 family, domains A, B, IV, and V are formed by non-contiguous sequences distributed at the N terminus or C terminus. Only domain C at the basis of the U-fold is built up by a contiguous amino acid network constituting the glucan binding pockets of the catalytic domain. Further, the GBD of DsrP glucansucrase are capable of binding proteins or toxins (Pfam CW_binding_1 (PF01473)) (21–23). In particular, the five repeats found at the C-terminal extremity of the GBD of DsrP glucansucrase are capable of binding the cell surface of Leuconostoc cells (24). Mutagenesis of conserved aromatic or basic residues of the repeats found in Streptococcus downei GtfI was performed and led to a reduction of the glucan binding capability (25). The elimination of some repeats by truncation was also shown to influence transglycosylation, modify glucan binding properties, and affect the polymer size (26–29). In particular, the GBD was suggested to provide a glucan anchoring platform, which may affect the single-chain elongation in a semi-processive manner (29). A few studies focused on isolated GBDs disconnected from their catalytic domain, and further demonstrated that truncated forms can autonomously fold and bind α-glucans with very low $K_d$ values estimated in the nanomolar range (30–34).

The crystal structures of several GH70 enzymes truncated at their N-terminal ends shed light on the GBD-fold, although no structures with entire GBDs have been solved yet (9, 35–37). These enzymes follow a U-path course and are organized in five domains, A, B, C, IV, and V. Domains A, B, and C are related to their counterparts found in GH13 α-amylases. In the GH70 family, domains A, B, IV, and V are formed by non-contiguous sequences distributed at the N terminus or C terminus. Only domain C at the basis of the U-fold is built up by a contiguous amino acid network constituting the glucan binding pockets of the catalytic domain. Further, the GBD of DsrP glucansucrase are capable of binding proteins or toxins (Pfam CW_binding_1 (PF01473)) (21–23). In particular, the five repeats found at the C-terminal extremity of the GBD of DsrP glucansucrase are capable of binding the cell surface of Leuconostoc cells (24). Mutagenesis of conserved aromatic or basic residues of the repeats found in Streptococcus downei GtfI was performed and led to a reduction of the glucan binding capability (25). The elimination of some repeats by truncation was also shown to influence transglycosylation, modify glucan binding properties, and affect the polymer size (26–29). In particular, the GBD was suggested to provide a glucan anchoring platform, which may affect the single-chain elongation in a semi-processive manner (29). A few studies focused on isolated GBDs disconnected from their catalytic domain, and further demonstrated that truncated forms can autonomously fold and bind α-glucans with very low $K_d$ values estimated in the nanomolar range (30–34).

The crystal structures of several GH70 enzymes truncated at their N-terminal ends shed light on the GBD-fold, although no structures with entire GBDs have been solved yet (9, 35–37). These enzymes follow a U-path course and are organized in five domains, A, B, C, IV, and V. Domains A, B, and C are related to their counterparts found in GH13 α-amylases. In the GH70 family, domains A, B, IV, and V are formed by non-contiguous sequences distributed at the N terminus or C terminus. Only domain C at the basis of the U-fold is built up by a contiguous amino acid network constituting the glucan binding pockets of the catalytic domain. Further, the GBD of DsrP glucansucrase are capable of binding proteins or toxins (Pfam CW_binding_1 (PF01473)) (21–23). In particular, the five repeats found at the C-terminal extremity of the GBD of DsrP glucansucrase are capable of binding the cell surface of Leuconostoc cells (24). Mutagenesis of conserved aromatic or basic residues of the repeats found in Streptococcus downei GtfI was performed and led to a reduction of the glucan binding capability (25). The elimination of some repeats by truncation was also shown to influence transglycosylation, modify glucan binding properties, and affect the polymer size (26–29). In particular, the GBD was suggested to provide a glucan anchoring platform, which may affect the single-chain elongation in a semi-processive manner (29). A few studies focused on isolated GBDs disconnected from their catalytic domain, and further demonstrated that truncated forms can autonomously fold and bind α-glucans with very low $K_d$ values estimated in the nanomolar range (30–34).
Carbohydrate Binding Ability of a GH70 Branching Sucrase

| Soaking | Observed ligand | Data collection |
|---------|-----------------|----------------|
| Complex A | d-Glucose | ID29 |
| Complex B | Isomaltotriose residue | 1D23-1 |
| Complex C | Gluco-oligosaccharides | ID23-1 |

Diffraction data

| Complex A | Complex B | Complex C |
|-----------|-----------|-----------|
| Beamline | ID23-1 | ID23-1 | ID23-1 |
| Wavelength (Å) | 0.9763 | 0.8726 | 0.9763 |
| Unit cell parameters (Å) | a = 67.77 b = 98.98 c = 181.91 | a = 68.29 b = 100.10 c = 184.43 | a = 68.31 b = 100.14 c = 186.24 |
| Space group | P2₁2₁2₁ | P2₁2₁2₁ | P2₁2₁2₁ |
| Molecules per AU* | 1 | 1 | 1 |
| Resolution (Å) | 47.8–2.3 (2.36–2.30) | 44.0–2.2 (2.30–2.18) | 46.920 (9.657) |
| No. of unique reflections | 54,661 (3,770) | 100.0 (100.0) | 109,405 (15,715) |
| Completeness (%) | 99.5 (94.2) | 5.4 (5.3) | 99.7 (99.1) |
| Multiplicity | 6.5 (4.6) | 5.4 (5.3) | 3.5 (3.2) |
| Rmerge b | 0.108 (0.735)* | 0.111 (0.439) | 0.090 (0.497) |
| Mean | 13.7 (2.1)* | 11.0 (3.6)* | 8.7 (2.0)* |
| Overall B factor–Wilson plot (Å²) | 34.2 | 23.8 | 24.7 |

| Refinement | |
| No. of reflections | 51,888 (2,772)* | 63,475 (3,388)* | 103,857 (5,458)* |
| Rwork (%) | 16.9 | 16.8 | 17.0 |
| Rsym (%) | 22.6 | 22.0 | 21.0 |
| Mean B values (Å²) | 40.0 | 28.0 | 30.0 |
| No of non-H atoms | 8,113 | 8,196 | 8,189 |
| Protein | 8,113 | 8,196 | 8,189 |
| Ligands | 127 | 42 | 64 |
| Water molecules | 395 | 715 | 922 |
| R.m.s. deviation from ideal | 0.013 | 0.015 | 0.019 |
| Bond lengths (Å) | 1.531 | 1.591 | 1.840 |
| Bond angles (°) | 95.0 | 96.6 | 96.9 |
| Ramachandran plot (%) | 99.8 | 99.9 | 100.0 |
| Favored | 0.2 | 0.1 | 0.0 |

**Note:** AU stands for asymmetric unit; values in parentheses are for the highest resolution shell.

*As defined in SCALA (43).

**Table 1** Data collection and refinement statistics for structures of ΔN₁₂₃-GBD-CD2 in complex with different sugars

purified as previously described were dissolved at 40 mM (39). Soaking of crystals with isomaltotriose Sigma-Aldrich Chemie S.a.r.l., Saint Quentin-Fallavier, France) was performed at 50 mM ligand. The concentration of PEG 3,350 was increased to 28.5% (w/v) for cryoprotection. D-Glucose (Sigma) was dis-

plex B, the groove search from subsite −1 was done using CAVER 3.01 as a PyMOL plugin (48).

**Molecular Dynamics**—The starting point for the molecular dynamics (MD) simulation was the crystal structure of complex A, composed of the protein and 10 D-glucose molecules. The calculations were performed with the Amber 14 software (49–51) using the FF03 force field for the protein, the GAFF force field for the ligands, and the TIP3P water model for the solvent (52). The complex was placed at the center of a periodic box. To hydrate the system, the volume occupied by the complex was extended by 10 Å on each side, and 40,251 water molecules were added to the box. To neutralize the charge of the system, 37 Na⁺ ions were added by random replacement of the equivalent number of solvent molecules. The final system had the dimensions of 101 × 141 × 113 Å and contained 136,985 atoms. The equilibration of the entire system was achieved in several steps. Initially, the energy of the system was minimized by 1,000 cycles of the steepest descent algorithm, with the solute held fixed, by constraining its Cartesian coordinates using a harmonic potential with the force constant k equal to 100 kcal/mol/Å². In the second step, the energy was minimized by 500 cycles of steepest descent and 1,500 cycles of the conjugate gradient algorithm, with weakly restrained solute (k = 10 kcal/mol/Å²). Next, a short 20-ps run of NVT-MD (constant volume and temperature) was performed on weakly restrained solute with temperature varying linearly from 0 to 300 K. The temperature control was achieved using the Langevin dynamics with the collision frequency parameter γ set to 1.0 ps⁻¹. The inte-

**Data Collection, Phasing, and Refinement**—Diffraction data were collected at the ESRF (Grenoble, France). Intensities were indexed and integrated using MOSFLM or XDS and scaled using SCALA or XSCALE (40–42). Crystals belong to space group P2₁2₁2₁ with one molecule per asymmetric unit, which corresponds to a solvent content of 52.5% and Matthews coefficient of 2.59 Å³ Da⁻¹. Structures were solved using molecular replacement with Phaser (43) and the atomic coordinates of the apo-form of ΔN₁₂₃-GBD-CD2 (PDB entry 3TTQ (9)) as a search model. Models were manually constructed in σA weighted 2Fo – Fc electron density maps using Coot and refined using Refmac5 (44, 45). Carbohydrate fitting in electron density maps was also guided by simulated annealing (Fc – Fc) omit maps were calculated with PHENIX (46). Data collection and refinement statistics are shown in Table 1.

**Structural Analysis**—Electrostatic potential of the ΔN₁₂₃-GBD-CD2 surface, without any ligand, was calculated via the PDB2PQR server using the program APBS, specifying the pH at 5.75, which is the optimum pH for activity (47). From complex APRIL 1, 2016 • VOLUME 291 • NUMBER 14 • JOURNAL OF BIOLOGICAL CHEMISTRY 7529
Carbohydrate Binding Ability of a GH70 Branching Sucrase

Determination of Binding Constants—Migration distances of BSA (dBSA) or \( \Delta N_{123} \)-GBD-CD2 (dENZ) were measured relative to the top and bottom of gels using the software associated with the GelDoc™ EZ System (Bio-Rad). Relative mobilities \( R \) were calculated as the migration distance of dextran divided by the migration distance of BSA \( R = d_{\text{ENZ}}/d_{\text{BSA}} \). In the absence of dextran, the relative mobility is defined as \( R_{\text{MAX}} \). This relative mobility is normalized to 1.0 \( (R'_{\text{MAX}}) \). The normalized relative mobilities are \( R' = R/R_{\text{MAX}} \). In the presence of a large excess of dextran, the relative normalized mobility is defined as \( R'_{\text{MIN}} \). Assuming that the total concentration of dextran in gels greatly exceeds the total polypeptide concentration, the affinity constant \( (K_a) \) of \( \Delta N_{123} \)-GBD-CD2 for dextran was calculated by plotting the normalized relative mobility, expressed as \( 1/(R'_{\text{MAX}} - R') \) of \( \Delta N_{123} \)-GBD-CD2 versus the molar polysaccharide concentration expressed as \( 1/[\text{Dex}] \). As described by Abbot and Boraston (65), the relationship between normalized relative mobilities and polysaccharide concentrations is,

\[
\frac{1}{R_{\text{MAX}} - R} = \frac{1}{(R_{\text{MAX}} - R_{\text{MIN}})K_a} \times \frac{1}{[\text{Dex}]} + \frac{1}{R_{\text{MAX}} - R_{\text{MIN}}}
\]

(Eq. 1)

where \( R'_{\text{MIN}} \), \( R'_{\text{MAX}} \), and \( K_a \) are constants.

Accession Numbers—The atomic coordinates were deposited in the Protein Data Bank with accession codes 4TVD, 4TTU and 4TVC for structures in complex with D-glucose, isomaltotriose, or isomaltotriosyl residues, respectively.

Results

Evidence of the Dextran Binding Ability of \( \Delta N_{123} \)-GBD-CD2—The \( \alpha -(1 \rightarrow 2) \) branching sucrase \( \Delta N_{123} \)-GBD-CD2 is a truncated enzyme that was designed from DSR-E, a GH70 glucansucrase with two catalytic domains (CD1 and CD2) separated by a putative GBD rich in repeats proposed to be involved in the glucan binding ability of several enzymes from the GH70 family (27, 57). The \( \Delta N_{123} \)-GBD-CD2 enzyme catalyzes transglucosylation reaction from sucrose to dextran or isomaltosyl-maltose, and a mixture of isomalto-oligosaccharides.

Affinity Gel Electrophoresis—For affinity gel electrophoresis, \( \Delta N_{123} \)-GBD-CD2 fused with a His\textsubscript{6} tag at the N terminus and a Strep-\textsubscript{tag} at the C terminus was expressed in a soluble form and purified as previously described. The dissociation constant of the purified \( \alpha -(1 \rightarrow 2) \) branching sucrase \( \Delta N_{123} \)-GBD-CD2 with dextrans was determined by affinity gel electrophoresis in 6.5% polyacrylamide native gels containing increasing amounts of dextran. Two types of dextran were tested, a high molar mass dextran with an estimated molar mass of 22.5 MDa and a medium molar mass dextran with an estimated molar mass of 68.4 kDa. Their concentrations ranged from 0 to 0.9% (w/v). Bovine serum albumin (BSA, Sigma) was used as a standard. Electrophoresis migration was performed for 2 h at 20 °C, pH 8.3, in a mini PROTEAN system (Bio-Rad) at constant intensity (10 mA per gel). The gels were stained with Coomassie Blue.

Identification of Glucan Binding Pockets by Multiple Sequence Alignments—Sequence alignment of the three repeats forming the sugar-binding pockets, V-J, V-K, and V-L in domain V of \( \Delta N_{123} \)-GBD-CD2, was first performed using Clustal Omega. This alignment was inspected and corrected using the structural superimposition of each of the three binding pockets. Then, the primary structure of the full-length GBD of DSR-E was analyzed using the online repeat detection tool TRUST, and this allowed detecting 12 sequence repeats in the studied system, nine simulations of this type have been performed, for the total time length of 450 ns. The resulting MD trajectories were analyzed with the AmberTools 14 programs as well as with in-house software. The last 20 ns of each equilibrated trajectory were used to calculate inter-molecular interaction energy for all protein-ligand pairs with the MM-GBSA approach (55).

Migration step used in this run was 1 fs. Throughout the calculations, a cut-off of 12 Å was used for electrostatic interactions, the particle mesh Ewald method was used to treat long-range electrostatic interactions and the SHAKE algorithm was used to constrain hydrogen bond lengths. The MD simulation continued for 100 ps with constant pressure of 1 bar (NPT-MD) at 300 K with no restraints, with the integration step of 2 fs. Finally, a 50-ns run with constant pressure of 1 bar was launched, with atomic coordinates saved every 10 ps. The Langevin dynamics was used to control the temperature, with \( \gamma = 1.0 \) ps\(^{-1} \), whereas the pressure was controlled by the Berendsen barostat with the pressure relaxation time \( \tau_p = 2 \) ps. Calculations were performed using a GeForce GTX TITAN Black GPU card, which worked at the speed of about 8.22 ns/day. To obtain a detailed view of the studied system, nine simulations of this type have been performed, for the total time length of 450 ns. The resulting MD trajectories were analyzed with the AmberTools 14 programs as well as with in-house software. The last 20 ns of each equilibrated trajectory were used to calculate inter-molecular interaction energy for all protein-ligand pairs with the MM-GBSA approach (55).

For affinity gel electrophoresis, (from T-coffee suite), providing structure-based multiple sequence alignment. In our case, the structural information of the binding pockets V-J, V-K, and V-L was used as templates. From this alignment, we deduced the LOGO sequence of one repeat using WebLogo 3.

Affinity Gel Electrophoresis—For affinity gel electrophoresis, \( \Delta N_{123} \)-GBD-CD2 fused with a His\textsubscript{6} tag at the N terminus and a Strep-\textsubscript{tag} at the C terminus was expressed in a soluble form and purified as previously described. The dissociation constant of the purified \( \alpha -(1 \rightarrow 2) \) branching sucrase \( \Delta N_{123} \)-GBD-CD2 with dextrans was determined by affinity gel electrophoresis in 6.5% polyacrylamide native gels containing increasing amounts of dextran. Two types of dextran were tested, a high molar mass dextran with an estimated molar mass of 22.5 MDa and a medium molar mass dextran with an estimated molar mass of 68.4 kDa. Their concentrations ranged from 0 to 0.9% (w/v). Bovine serum albumin (BSA, Sigma) was used as a standard. Electrophoresis migration was performed for 2 h at 20 °C, pH 8.3, in a mini PROTEAN system (Bio-Rad) at constant intensity (10 mA per gel). The gels were stained with Coomassie Blue.
Carbohydrate Binding Ability of a GH70 Branching Sucrase

Among the nine glucose binding sites, site A-1 corresponds to subsite \(-1\) of GH70 transglucosylases, which was shown to accommodate the glucopyranosyl ring of sucrose (66). The \(\beta\)-D-glucopyranose in A-1 adopts a \(4\C_1\) conformation, similar to that observed for the \(\beta\)-glucopyranosyl unit of sucrose described for the GTF180-\(\Delta N\)-sucrase complex (35). The oxygen atom O1 of glucose is hydrogen bound to the side chain atoms of the catalytic residues Glu-2248 and Asp-2210 at a distance of 2.35 and 2.67 \(\AA\), respectively. The C2 atom also forms a hydrogen bond to the side chain of Ser-2266. These values of 2.9 kcal/mol). MM-GBSA calculations were performed to estimate the interaction energy between \(\Delta N\)-GBD-CD2 and \(\delta\)-glucose molecules. Interaction energy values for glucose molecules at subsites A-1 and A2 were comparable (about \(-20\) kcal/mol). Altogether, these observations reveal that the binding sites A-1 and A2 of the catalytic gorge have strong affinities for glucose molecules. The A3 site was found to bind the ligand for a limited time (about 90 ns), after which the ligand left for the solvent.

The \(\delta\)-glucose molecule at site A4 is bound in a pocket at the rear of the \((\beta/\alpha)_{8}\) barrel of domain A (Fig. 3A) and establishes hydrogen bonds with Asn-2241, Asn-2588, Asn-2683, and Gln-2685. The oxygen atoms of the hydroxyl groups O6 and O1 are oriented toward the enzyme. Molecular dynamics simulations suggest that the molecule is loosely bound to the protein. In domain IV, two glucose molecules that are located 4.2 Å from each other occupy the glucose-binding sites IV1 and IV2.

\(\delta\)-Glucose at site IV1 is stacked over Trp-2014 and is stabilized by a hydrogen bond network involving side chains of Asp-2022, His-2023, and Gln-2102. Finally, additional glucose molecule is located at the interface of two symmetry-related protein molecules. This binding site will not be described further due to its biological inconsistency.

Among the nine glucose binding sites, site A-1 corresponds to subsite \(-1\) of GH70 transglucosylases, which was shown to accommodate the glucopyranosyl ring of sucrose (66). The \(\beta\)-D-glucopyranose in A-1 adopts a \(4\C_1\) conformation, similar to that observed for the \(\beta\)-glucopyranosyl unit of sucrose described for the GTF180-\(\Delta N\)-sucrase complex (35). The oxygen atom O1 of glucose is hydrogen bound to the side chain atoms of the catalytic residues Glu-2248 and Asp-2210 at a distance of 2.35 and 2.67 \(\AA\), respectively. The C2 atom also forms a hydrogen bond to the side chain of Ser-2266. These values of 2.9 kcal/mol). MM-GBSA calculations were performed to estimate the interaction energy between \(\Delta N\)-GBD-CD2 and \(\delta\)-glucose molecules. Interaction energy values for glucose molecules at subsites A-1 and A2 were comparable (about \(-20\) kcal/mol). Altogether, these observations reveal that the binding sites A-1 and A2 of the catalytic gorge have strong affinities for glucose molecules. The A3 site was found to bind the ligand for a limited time (about 90 ns), after which the ligand left for the solvent.

The \(\delta\)-glucose molecule at site A4 is bound in a pocket at the rear of the \((\beta/\alpha)_{8}\) barrel of domain A (Fig. 3A) and establishes hydrogen bonds with Asn-2241, Asn-2588, Asn-2683, and Gln-2685. The oxygen atoms of the hydroxyl groups O6 and O1 are oriented toward the enzyme. Molecular dynamics simulations suggest that the molecule is loosely bound to the protein. In domain IV, two glucose molecules that are located 4.2 Å from each other occupy the glucose-binding sites IV1 and IV2.

\(\delta\)-Glucose at site IV1 is stacked over Trp-2014 and is stabilized by a hydrogen bond network involving side chains of Asp-2022, His-2023, and Gln-2102. Finally, additional glucose molecule is located at the interface of two symmetry-related protein molecules. This binding site will not be described further due to its biological inconsistency.

Among the nine glucose binding sites, site A-1 corresponds to subsite \(-1\) of GH70 transglucosylases, which was shown to accommodate the glucopyranosyl ring of sucrose (66). The \(\beta\)-D-glucopyranose in A-1 adopts a \(4\C_1\) conformation, similar to that observed for the \(\beta\)-glucopyranosyl unit of sucrose described for the GTF180-\(\Delta N\)-sucrase complex (35). The oxygen atom O1 of glucose is hydrogen bound to the side chain atoms of the catalytic residues Glu-2248 and Asp-2210 at a distance of 2.35 and 2.67 \(\AA\), respectively. The C2 atom also forms a hydrogen bond to the side chain of Ser-2266. These values of 2.9 kcal/mol). MM-GBSA calculations were performed to estimate the interaction energy between \(\Delta N\)-GBD-CD2 and \(\delta\)-glucose molecules. Interaction energy values for glucose molecules at subsites A-1 and A2 were comparable (about \(-20\) kcal/mol). Altogether, these observations reveal that the binding sites A-1 and A2 of the catalytic gorge have strong affinities for glucose molecules. The A3 site was found to bind the ligand for a limited time (about 90 ns), after which the ligand left for the solvent.

The \(\delta\)-glucose molecule at site A4 is bound in a pocket at the rear of the \((\beta/\alpha)_{8}\) barrel of domain A (Fig. 3A) and establishes hydrogen bonds with Asn-2241, Asn-2588, Asn-2683, and Gln-2685. The oxygen atoms of the hydroxyl groups O6 and O1 are oriented toward the enzyme. Molecular dynamics simulations suggest that the molecule is loosely bound to the protein. In domain IV, two glucose molecules that are located 4.2 Å from each other occupy the glucose-binding sites IV1 and IV2. \(\delta\)-Glucose at site IV1 is stacked over Trp-2014 and is stabilized by a hydrogen bond network involving side chains of Asp-2022, His-2023, and Gln-2102. Finally, additional glucose molecule is located at the interface of two symmetry-related protein molecules. This binding site will not be described further due to its biological inconsistency.

Among the nine glucose binding sites, site A-1 corresponds to subsite \(-1\) of GH70 transglucosylases, which was shown to accommodate the glucopyranosyl ring of sucrose (66). The \(\beta\)-D-glucopyranose in A-1 adopts a \(4\C_1\) conformation, similar to that observed for the \(\beta\)-glucopyranosyl unit of sucrose described for the GTF180-\(\Delta N\)-sucrase complex (35). The oxygen atom O1 of glucose is hydrogen bound to the side chain atoms of the catalytic residues Glu-2248 and Asp-2210 at a distance of 2.35 and 2.67 \(\AA\), respectively. The C2 atom also forms a hydrogen bond to the side chain of Ser-2266. These values of 2.9 kcal/mol). MM-GBSA calculations were performed to estimate the interaction energy between \(\Delta N\)-GBD-CD2 and \(\delta\)-glucose molecules. Interaction energy values for glucose molecules at subsites A-1 and A2 were comparable (about \(-20\) kcal/mol). Altogether, these observations reveal that the binding sites A-1 and A2 of the catalytic gorge have strong affinities for glucose molecules. The A3 site was found to bind the ligand for a limited time (about 90 ns), after which the ligand left for the solvent.

The \(\delta\)-glucose molecule at site A4 is bound in a pocket at the rear of the \((\beta/\alpha)_{8}\) barrel of domain A (Fig. 3A) and establishes hydrogen bonds with Asn-2241, Asn-2588, Asn-2683, and Gln-2685. The oxygen atoms of the hydroxyl groups O6 and O1 are oriented toward the enzyme. Molecular dynamics simulations suggest that the molecule is loosely bound to the protein. In domain IV, two glucose molecules that are located 4.2 Å from each other occupy the glucose-binding sites IV1 and IV2. \(\delta\)-Glucose at site IV1 is stacked over Trp-2014 and is stabilized by a hydrogen bond network involving side chains of Asp-2022, His-2023, and Gln-2102. Finally, additional glucose molecule is located at the interface of two symmetry-related protein molecules. This binding site will not be described further due to its biological inconsistency.
by the constant distance of their center of mass to the neighboring protein residues and by the formation of only 1 intra-molecular contact. Their interaction energies are similar to those found for the A-1 and A2 sites (about $-20$ kcal/mol), probably because of abundance of OH groups capable of forming electrostatic interactions with neighboring residues, nearly independently of the orientation of the ligand within the binding site, revealing for the first time a possible carbohydrate binding function of GH70 glucansucrase domain IV. The glucose bound in site IV-V is labile in half of the simulated trajectories. In the site B-IV-V, D-glucose is orientated with O6 and O1 exposed to the solvent and was shown to be labile by molecular dynamics.

Finally, one glucose molecule was bound to the surface of domain V. This is the first carbohydrate molecule ever identified in domain V of a GH70 enzyme. The corresponding binding site expanding from Ala-1906 to Asp-1981 was named the V-L pocket (Fig. 3A). The p-glucose molecule in the 4C1 conformation is stacked over Tyr-1914, with its oxygen atoms O2, O3, and O4 engaged in hydrogen bonds with the side chains of Gln-1942, Gln-1951, Lys-1953 and with the main chain carbonyl group of Lys-1969 (not shown). Glucose binding in the V-L site is labile because the r.m.s. deviation value was about 46.6 ± 6.6 Å over the last 10 ns of molecular dynamics simulation. A non-interpretable electron density was also observed in domain V over residue Tyr-1834, suggesting the presence of another binding site, named V-K, expanding from Met-1828 to Val-1905.

Structures of $\Delta N_{125}$-GBD-CD2 in Complex with Isomaltosyl or Isomaltotriosyl Residues—$\Delta N_{125}$-GBD-CD2 was soaked with pure isomaltotriose or with a mixture of gluco-oligosaccharides (mainly composed of isomaltosyl-maltose and isomaltotriosyl-maltose). The crystal structures of the corresponding complexes B and C were solved at 2.2 and 1.85 Å, respectively. These structures did not reveal any interpretable electron density in the catalytic gorge or in the previously described glucose-binding sites (A-1, A2, A3, A4, IV1, IV2, B-IV-V, or IV-V). In contrast, electron density corresponding to glucose, isomaltosyl, or isomaltotriosyl groups were visible in pockets V-K and V-L of domain V. Both binding pockets are made up of three consecutive super-secondary elements, either $\beta$-hairpins or three-stranded $\beta$-sheets. In site V-K, a clear electron density was observed, which corresponds to an isomaltosyl group or an isomaltotriosyl group in complexes B and C, respectively (Fig. 6). The electron density maps indicate that the oligosaccharides may possess an additional glucosyl residue oriented toward the solvent. In the V-L binding site, one glucosyl
residue was visible in the electron density maps of both complexes (Fig. 7).

A Rearrangement of the Catalytic Gorge in Complex B—Compared with the previously published apo structure, a rearrangement of loop 7 (from His-2319 to Ser-2369) is observed in complex B (Fig. 8). This loop connects strand 7 and helix 7 of the (β/α)8 barrel. The mean B-factor of the corresponding Cα atoms is 39.7 Å², which is almost twice as much as the value observed for Cα atoms of domains A, B, and C (21.5 Å²). A geometric exploration of the grooves expanding from subsite −1 of the catalytic gorge was performed with CAVER, and four different grooves were found. Grooves A2 and A3 include glucose binding sites A2 and A3, respectively (Fig. 3B). The groove named “groove loop 7” is delineated by the cleft occurring upon

FIGURE 4. Glucose-binding sites found over the ∆N123-GBD-CD2 enzyme. The simulated annealing omit (Fo-Fc) electron density maps (in green) around the nine glucose molecules bound to ∆N123-GBD-CD2 were contoured at 2σ. ∆N123-GBD-CD2 residues in the neighborhood of each glucose molecule are shown as lines and colored by atom types. Stereo view of subsites A-1, A2, A3, A4, B-IV-V, IV1, IV2, IV-V, and VL are shown on panels A–I, respectively.

FIGURE 5. Glucose-binding sites A2 (left panel) and A3 (right panel). The simulated annealing omit (Fo-Fc) electron density maps around glucose were contoured at 3σ or 2.3σ for A2 and A3, respectively. Interacting water molecules are depicted as red spheres. Residues of domains A or B are represented as blue or green sticks, respectively. For residues that are involved in sugar binding through their backbone atoms, side chains are omitted. In the left panel, residues forming the catalytic triad (Asp-2210, Glu-2248, and Asp-2322) are shown as turquoise sticks.
loop 7 rearrangement. Finally, groove 4 expands over loop 7. As shown in Fig. 3C, the movement of loop 7 reveals a positively charged patch in the vicinity of subsite −1 in groove 4, which is absent in the apo structure.

The V-K Binding Pocket—In complex C, seven residues of the V-K pocket are involved in isomaltotriosyl binding (Fig. 6). The α-D-glucopyranosyl unit 2 (Glc-2) is stacked in the C1 conformation over Tyr-1834. The α-D-glucosyl units 1 and 3 (Glc-1 and Glc-3) are also probably stabilized in the C1 conformation, although the electron density map is less well defined. The Glc-1 and Glc-2 moieties are essentially stabilized by hydrogen bonding with side chain atoms; the O2 atom of Glc-2 interacts with Gln-1879 and Lys-1881 and the O3 atom with Gln-1870 and Lys-1898 (via the oxygen atom of its main chain carbonyl group). The O5 atom of Glc-1 could make a long-range hydrogen bond with the side chain of Lys-1881, O2 with the oxygen atom of Gly-1897 carbonyl function, and O4 with Lys-1898 and a water molecule. The side chain of Lys-1881 also interacts with the oxygen atom of the osidic bond between Glc-1 and Glc-2. The Glc-3 is weakly stabilized by hydrogen bonding through two water molecules and through van der Waals interactions with Tyr-1834 and Trp-1849.

In complex B, one isomaltosyl residue is found in the V-K pocket. The backbone atoms of the V-K pocket of complexes B and C superimpose well. The glucopyranosyl ring at the non-reducing end is stacked in the C1 conformation over the side chain of Tyr-1834, similar to Glc-2 in complex C (Fig. 6). The ring is also maintained by hydrogen bonds between O3 and the main chain carbonyl group of Lys-1898 and the side chain of Gln-1870. Of note, in complexes B and C, the side chains of these two amino acids adopt different conformations. Finally, water-mediated hydrogen bonds with residues Asn-1900 and Thr-1868 complete the network of interactions. The second glucosyl unit (equivalent to Glc-1 in complex C) is weakly stabilized by a hydrogen bond between O5 and the side chain of Lys-1881.

The V-L Binding Pocket—Electron density corresponding to one D-glucopyranosyl unit in C1 conformation was found in the binding pocket V-L of complexes B and C (Fig. 7). The interaction network is similar to that described for the complex with D-glucose (Fig. 3A). Interestingly, residues Tyr-1914, Gln-1942, Gln-1951, Lys-1953, and Lys-1969 of the V-K pocket are conserved in the V-L pocket. The glucopyranosyl residue is bound so that the O1 and O6 atoms are pointing toward the
In this pocket, there is no tryptophan equivalent to Trp-1849 of V-K pocket that could stabilize additional glucosyl residues.

Discussion

In recent years, major advances have been made in the comprehension of structure-function relationships of GH70 sucrose-active enzymes thanks to the three-dimensional structure resolution of GTF180-ΔN GTFα-ΔN, GTF-SI, and ΔN123-GBD-CD2 glucansucrases in the apo form (9, 35–37). These structures revealed the unique U-fold type of these enzymes organized in five structural domains. In addition, several structures in complex with sucrose donor, maltose acceptor, or acarbose (a known inhibitor of various glycoside-hydrolases) were solved that allowed further description of the -1, +1, +2, and +2′ subsites of these enzymes (35, 36). However, structural data are still missing to draw up the full scenario of polymerization at the molecular level and identify the structural determinants controlling substrate and product specificity. In particular, three-dimensional structures in complex with either isomaltotriose ligands or dextrans, the natural acceptors of these enzymes, were heavily lacking, thus preventing molecular descriptions of isomaltotriose or dextran-binding sites. In particular, no complexes were previously disclosed in which carbohydrates were present in domain V, although this domain was biochemically proven to act as a glucan-binding domain in several GH70 enzymes (25, 31, 32) and was suggested to participate in the processivity of polymerization (26, 28, 29).

Regarding the α-(1→2) branching sucrase (ΔN123-GBD-CD2) enzymes, structural data are even scarcer as only two structures in the apo form have been solved thus far (9).

In this context, solving the structure of the α-(1→2) branching sucrase ΔN123-GBD-CD2 in complex with a glucoligosaccharide acceptor was quite challenging, and we have obtained the first three-dimensional structures of this enzyme in complex with D-glucose, isomaltosyl, or isomaltotriosyl residues. The complex with D-glucose clearly showed that one glu-
Carbohydrate Binding Ability of a GH70 Branching Sucrase
cose molecule is accommodated in −1 subsite in a conformation similar to that described for the GTF180-ΔN-sucrose complex, corroborating the α-retaining mechanism proposed for GBD-CD2. In addition, the glucose molecules occupying sites A2 and A3, in proximity of the catalytic center, both expose their O1 and O6 atoms to the solvent in a proper orientation to be engaged in α-(1→6) osidic linkages. Such positioning suggests that A2 and A3 sites may provide binding patches delineating a gorge capable of accommodating longer α-(1→6) linked gluco-oligosaccharides or dextrans. The question remains of whether binding sites A2 and A3 define acceptor subsites connected to −1 and +1 subsites of the catalytic center. Additionally, it is unknown if the binding sites could be connected to the other glucose-binding sites found in domains B, IV, and V. Only complexes with longer oligosaccharides could help answer this question (66). Interestingly, residues Glu-2265, Ser-2266, and Glu-2270 of binding site A2 are also conserved in the recently discovered α-(1→2) branching sucrase BRS-A, suggesting that they could be determinants of α-(1→2) branching specificity (10).

Structural analysis of complexes B and C also provide the first structural evidence of interactions between domain V of ΔN123-GBD-CD2 and glucose, as well as isomaltosyl and isomaltotriosyl groups. Surprisingly, comparison of complex B with the apo form revealed that loop 7 of the (β/α)8 barrel moves and opens a cleft toward subsite −1. Thus, it may be tempting to propose that such a rearrangement could assist oligodextran accommodation (or release) in (or from) the observed grooves and could play a role in the ping-pong bi-bi mechanism as well as in the previously described stochastic branching of short dextran chains (39, 67). In domain V of ΔN123-GBD-CD2, the amino acid residues of binding pockets V-K and V-L that interact with carbohydrates are well conserved, with the exception of Trp-1849 and Thr-1868, which are replaced by a threonine and alanine in the pocket V-L, respectively. These differences may be sufficient to modulate affinity and only permit glucose binding in pocket V-L. Furthermore, by inspecting the structures of GTFA-ΔN and GTF180-ΔN, we identified a pocket similar to the V-K and V-L binding pockets of ΔN123-GBD-CD2 in each (Fig. 9). These four pockets contain from 76 to 81 residues and are also constituted by a sequence of three super-secondary motifs that can be either β-hairpins or three-stranded antiparallel β-sheets (Figs. 9 and 10). Most of the conserved aromatic residues are mainly oriented toward the hydrophobic core of the pocket and belong to β-strands. The four pockets are exposed to the solvent and delineated by acidic or basic residues (Asp, Glu, Lys, and Arg), by aromatic residues (Trp or Tyr), and by polar residues (Thr, Ser, Gln, and Asn). They all contain one aromatic residue (Phe or Tyr) at the bottom, suggesting that they could also promote dextran binding. However, structural data obtained by Vujčić-Žagar et al. from GTF180-ΔN crystals soaked with D-glucose, isomaltose, panose, or nigerotriose did not reveal any sugar binding in domain V (35). It remains to be understood what subtle differences in the experimental conditions or in the putative binding pocket of GTF180-ΔN render this pocket non-functional. However, the complete truncation of domain V in GTF180-ΔN strongly reduces its ability to synthetize α-glucans from sucrose in favor of oligosaccharide synthesis (69). This suggests that domain V of GTF180-ΔN may have a positive impact on α-glucan synthesis, possibly by promoting the binding to α-glucan chains.

In addition, structural alignments with other domains V of several GH70 glucansucrases, which were experimentally proved to bind α-glucans, were performed with EXPRESSO to check the possible conservation of the residues delineating the binding pockets. The alignment reveals that several sequences putatively corresponding to sugar binding pockets are present. These putative binding pockets include four YG repeats, as defined by Giffard and Jacques (20), two complete repeats flanked by two partial ones (Fig. 10). The residues of the V-K pockets interacting with carbohydrates are conserved, i.e. the aromatic residues acting as a stacking platform (logo number 19 -Fig. 10), Trp-34, Gln-71, and Lys-73 (respectively, equivalent to Tyr-1834, Trp-1849, Gln-1879, and Lys-1881 in the V-K pocket). Then the alignment was extended to the entire GBD of DSR-E, from which ΔN123-GBD-CD2 is derived. As seen in Fig. 11, 10 sequences of putative binding pockets (named V-A to V-I), containing 46 to 84 residues each, aligned with the sequences of V-K and V-L pockets. As for the domains that were shown to bind dextrans, these sequences contain the residues involved in sugar binding in V-L and V-K pockets (i.e. aromatic residue 11, Trp-26, Gln-62, and Lys-64, LOGO sequence numbering, Fig. 11), indicating that these pockets are
likely to be functional. In 2004, Shah et al. (25) carried out a site-directed mutagenesis study focusing on the conserved residues of GBD1A, a truncated domain V disconnected from the enzymatic core. A total of nine amino acids belonging to binding pocket repeats 1, 2, 3, and 4 were targeted, including Phe-1122, Trp-1163, Trp-1227, Trp-1292, Tyr-1294, Tyr-1314, Phe-1315, Gln-1321, and Lys-1323 (Fig. 11). Eight mutants lost their ability to bind biotinylated dextrans, whereas eight others retained this ability, showing that some positions are tolerant to mutations and others are more critical for this function (25).

The sugar-binding sites identified in domain V are likely involved in the α-glucan binding ability of \( \Delta N_{123} \)-GBD-CD2, even if we cannot exclude that dextran binding also occurs in other domains. We determined \( K_d \) values for dextrans in the micromolar range. In comparison, the full-length DSR-S, a dextranase produced by a constitutive mutant of \( L. mesenteroides \) B-512F, has a higher affinity for dextran than the truncated domains. Sequences are GBD1A_GTFI_rp# (repeats 1 to 4 of the GBD1A of GTFI from \( S. downei \)), GBD-4RS_GTFI_rp# (repeats 1 to 4 of the GBD-4RS of GTFI from \( S. sobrinus \)), GBD5_DSR-S_rp# (repeats 1 to 3 of the GBD5 of DSR-S from \( L. mesenteroides \) B-512F), GBD_GTF180_rp1 (repeat 1 of the GBD of GTF180-JN from \( Lactobacillus reuteri \)), GBD_GTFA_rp1 (repeat 1 of the GBD of GTFIA-JN from \( L. reuteri \)), and GBD_DSR-E_rp# (repeats J to L of the GBD of \( \Delta N_{123} \)-GBD-CD2 from \( L. citreum \) NRRL-1299). Residues highlighted in red are well conserved, whereas those trending toward blue are not. Residues framed in gray delineate the sugar binding pockets and are pointed out in the LOGO sequence. Residues framed in dashed line on the LOGO sequence correspond to YG repeats. In the repeats of GBD1A, residues circled in white, were mutated by Shah et al. (25) (see “Discussion”). Secondary structure elements, as determined by DSSP from PDB entries 3TTQ, 3KLK, and 4AMC, are displayed below the LOGO sequence.
teroides NRRL B-512F, was shown to bind 170-kDa dextran with a $K_d$ value of 18 nm using a quartz crystal microbalance (68). The whole domain V of DSR-S, devoid of the catalytic domain, showed a $K_d$ value for biotinylated dextran of 70 nm (32). Although the methods and the dextran used for determining the $K_d$ value vary among these studies, our results suggest that the truncated branching sucrase $\Delta N_{23}$-GBD-CD2 has a weaker affinity for dextran than the full-length domain V of DSR-S dextransucrase, which seems to contain a higher number of putative binding pockets. Using isothermal titration calorimetry, Komatsu et al. (31) studied the binding of 25-kDa dextran to a set of independent truncated domain V derived from GTFI, including the truncated form GBD-4RS and the full-length GBD (Fig. 11). The exothermic binding of dextran to the full-length domain V gave a $K_d$ value of 20 nm at 25 °C. The binding stoichiometry was as low as 0.12, which indicates that several molecules bind to stretches of dextran chains. Moreover, $-\Delta H^0$ and $-\Delta G^0$ were found to be proportional to the number of residues of the truncated domain V, which substantiated the fact that the binding sites are independent and similar. The affinity of this GBD with dextran was proposed to be the sum of weak interactions with different glucose-binding sites, which is consistent with our structural data.

Our findings reveal for the first time that binding of isomaltooligosaccharides to domain V occurs via sugar binding pockets comprising between 60 and 81 residues. In the V-K and V-L pockets, disclosed herein, eight residues are interacting with the sugar rings. Only four of them, one aromatic residue acting as a stacking platform, one tryptophan, one glutamine, and one lysine, are conserved in the putative binding pockets of other domain V shown to bind $\alpha$-glucans. It remains to be understood what specific features in these binding pockets confer, or do not confer, a strong affinity for $\alpha$-glucans. The x-ray structures of other domain V, in complex with longer isomaltooligosaccharides and containing variable numbers of sugar pockets, could help to elucidate the possible synergy between the binding pockets and its influence on glucan binding strength. In addition, glucan binding in domain V probably contribute to increasing the local concentration of carbohydrate chains in the vicinity of the active site and may play a key role in presenting these chains to the active site for their branching or elongation. This assumption is corroborated by the fact that DSR-S and ASR glucansucrases partially truncated of their domain V synthesized glucans of lower molar mass than those synthesized by the entire enzymes (29). Efforts should be pursued to solve the structure of glucansucrases or branching sucrases with short dextran chains bound in the active site as well as in domains B, IV, and V and combine structural analyses to mutagenesis and detailed biochemical characterization to get further insight into these mechanisms.

Acknowledgments—We greatly acknowledge the staff from the European Synchrotron Radiation Facility (ESRF, Grenoble, France) and the Integrated Screening Platform of Toulouse (PICT, Toulouse, France) for providing access to macromolecular crystallography equipment.

References

1. Cantarel, B. L., Coutinho, P. M., Rancurel, C., Bernard, T., Lombard, V., and Henrissat, B. (2009) The Carbohydrate-Active EnZymes database (CAZy): an expert resource for Glycomics. Nucleic Acids Res. 37, D233–238
2. Leemhuis, H., Pijning, T., Dobruchowska, J. M., van Leeuwen, S. S., Kralj, S., Dijkstra, B. W., and Dijkhuizen, L. (2013) Glucansucrases: three-dimensional structures, reactions, mechanism, $\alpha$-glucan analysis and their implications in biotechnology and food applications. J. Biotechnol. 163, 260–272
3. Leemhuis, H., Dijkman, W. P., Dobruchowska, J. M., Pijning, T., Grijspeer, P., Kralj, S., Kamerling, J. P., and Dijkhuizen, L. (2013) 4,6-\alpha-Glucanosyltransferase activity occurs more widespread in Lactobacillus strains and constitutes a separate GH70 subfamily. Appl. Microbiol. Biotechnol. 97, 181–193
4. Côté, G. L., and Leathers, T. D. (2005) A method for surveying and classifying Leucosostoc spp. glucansucrases according to strain-dependent acceptor product patterns. J. Ind. Microbiol. Biotechnol. 32, 53–60
5. Naessens, M., Cerdobbel, A., Soetart, W., and Vandamme, E. J. (2005) Leucosostoc dextransucrase and dextran: production, properties and applications. J. Chem. Technol. Biotechnol. 80, 845–860
6. Falconer, D. J., Muckerjee, R., and Robyt, J. F. (2011) Biosynthesis of dextrins with different molecular weights by selecting the concentration of Leucosostoc mesenteroides B-512FMC dextranosucrase, the sucrose concentration, and the temperature. Carbohydr. Res. 346, 280–284
7. Vettori, M. H. P. B., Blanco, K. C., Cortez, M., De Lima, C. J. B., and Contiero, J. (2012) Dextran: effect of process parameters on production, purification and molecular weight and recent applications. Diálogo Ciên. 2012, 171–186
8. Bowen, W. H., and Koo, H. (2011) Biology of Streptococcus mutans-derived glucosyltransferases: role in extracellular matrix formation of cariogenic biofilms. Caries Res. 45, 69–86
9. Brison, Y., Pijning, T., Malbert, Y., Fabre, É., Mourey, L., Morel, S., Potocki-Véronèse, G., Monsan, P., Tranier, S., Remaun-Siméon, M., and Dijkstra, B. W. (2012) Functional and structural characterization of $\alpha$-(1→2) branching sucrose derived from DSR-E glucansucrase. J. Biol. Chem. 287, 7915–7924
10. Passerini, D., Vuillemin, M., Ufarté, L., Morel, S., Loux, V., Fontagné-Faucher, C., Monsan, P., Remaun-Siméon, M., and Moulin, C. (2015) Inventory of the GH70 enzymes encoded by Leucosostoc citreum NRRL B-1299: identification of three novel $\alpha$-transglycosylases. FEBS J. 282, 2115–2130
11. Lombard, V., Golacancla Ramulu, H., Drula, E., Coutinho, P. M., and Henrissat, B. (2014) The carbohydrate-active enzymes database (CAZy) in 2013. Nucleic Acids Res. 42, D490–D495
12. Boraston, A. B., Bolam, D. N., Gilbert, H. J., and Davies, G. J. (2004) Carbohydrate-binding modules: fine-tuning polysaccharide recognition. Biochem. J. 382, 769–781
13. Mooser, G., and Wong, C. (1998) Isolation of a glucan-binding domain of glucosyltransferase (1,6-$\alpha$-glucan synthase) from Streptococcus sobrinus. Infect. Immun. 66, 880–884
14. Ferratti, J. J., Gilpin, M. L., and Russell, R. R. (1987) Nucleotide sequence of a glucosyltransferase gene from Streptococcus sobrinus MFe28. J. Bacteriol. 169, 4271–4278
15. Banas, J. A., Russell, R. R., and Ferratti, J. J. (1990) Sequence analysis of the gene for the glucan-binding protein of Streptococcus mutans Ingritt. Infect. Immun. 58, 667–673
16. Gilmore, K. S., Russell, R. R., and Ferratti, J. J. (1990) Analysis of the Streptococcus downei gfs gene, which specifies a glucosyltransferase that syn-
Carbohydrate Binding Ability of a GH70 Branching Sucrase

Ikuo, I., Ito, S., Shimamura, T., Weyand, S., Kawarasaki, Y., Misaka, T., Abe, K., Kobayashi, T., Cameron, A. D., and Iwata, S. (2011) Crystal structure of glucan binding from the dental caries pathogen Streptococcus mutans. J. Biol. Chem. 408, 177–186

Pijning, T., Vujicic-Zagar, A., Kralj, S., Dijkstra, B. W. (2012) Structure of the α-1,6-α-1,4-specific glucanase GTFA from Lactobacillus reuteri 121. Acta Crystallogr. Sect. F Struct. Biol. Cryrm. 68, 1448–1454

Pijning, T., Vujicic-Zagar, A., Kralj, S., Dijkstra, B. W., and Bovent, T. (2014) Flexibility of truncated and full-length glucanase GTF180 enzymes from Lactobacillus reuteri 180. FEBS J. 281, 2159–2171

Stron, Y., Laguerre, S., Fereol, M., Morel, S., Monties, N., Potok-Mirenow, G., Monsan, P., and Remaud-Simeon, M. (2013) Branching pattern of gluco-oligosaccharides and 1.5-kDa dextran grafted by the α-1,2 branching succrase GBD-CD2. Carbohydr. Polym. 94, 567–576

Kabsch, W. (2010) XDS. Acta Crystallogr. D Biol. Crystallogr. 66, 125–132

Bartey, T. G., Kontogiani, L., Johnson, O., Powell, H. R., and Leslie, A. G. (2011) iMOSFLM: a new graphical interface for diffraction-image processing with MOSFLM. Acta Crystallogr. D Biol. Crystallogr. 67, 271–281

Evans, P. (2006) Scaling and assessment of data quality. Acta Crystallogr. D Biol. Crystallogr. 62, 72–82

McCoy, A. J., Grosse-Kunstleve, R. W., Adams, P. D., Winn, M. D., Storoni, L. C., and Read, R. J. (2007) Phaser crystallographic software. J. Appl. Crystallogr. 40, 658–674

Emsley, P., Lohkamp, B., Scott, W. G., and Cowtan, K. (2010) Features and development of Coot. Acta Crystallogr. D. 66, 486–501

Muschudov, G. N., Skubak, P., Lebedev, A. A., Pannu, N. S., Steiner, R. A., Nicholls, R. A., Winn, M. D., Long, F., and Vagin, A. A. (2011) REFMAC5 for the refinement of macromolecular crystal structures. Acta Crystallogr. D Biol. Crystallogr. 67, 355–367

Adams, P. D., Afonine, P. V., Bunkoczi, G., Chen, V. B., Davis, I. W., Echols, N., Headd, J. J., Hung, L.-W., Kapral, G. J., Grosse-Kunstleve, R. W., McCoy, A. J., Moriarty, N. W., Oeffner, R., Read, R. J., Richardson, D. C., et al. (2010) PHENIX: a comprehensive Python-based system for macromolecular structure solution. Acta Crystallogr. D Biol. Crystallogr. 66, 213–221

Dolinsky, T. J., Nielsen, E. J., McCammon, J. A., and Baker, N. A. (2004) PDBePQR: an automated pipeline for the setup of Poisson–Boltzmann electrostatic calculations. Nucleic Acids Res. 32, W665–W667

Kozlikova, B., Sebestova, E., Sustr, V., Brezovicky, I., Strnad, O., Daniel, L., Bednar, D., Pavelka, A., Manak, M., Bezdekova, M., Benes, P., Kotry, M., Gora, A., Damborsky, J., and Sochor, J. (2013) CAVER Analyst 1.0: a graphic tool for interactive visualization and analysis of tunnels and channels in protein structures. Bioinformatics 29, 2684–2685

Case, D. A., Cheatham, T. E., 3rd, Darden, T., and Brooks, C. L. (2004) Development and testing of a general amber forcefield. J. Chem. Theory Comput. 25, 1157–1174

Salomon-Ferrer, R., Götz, A. W., Poole, D., LeGrand, S., and Walker, R. C. (2013) Routine microsecond molecular dynamics simulations with AMBER on GPUs: 2. explicit solvent particle mesh Ewald. J. Chem. Theory Comput. 9, 3878–3888

Salomon-Ferrer, R., Case, D. A., and Walker, R. C. (2013) An overview of the Amber biomolecular simulation package. WIREs Comput. Mol. Sci. 3, 198–210

Wang, J., Wolf, R. M., Caldwell, J. W., Kollman, P. A., and Case, D. A. (2004) Development and testing of a general amber force field. J. Comput. Chem. 25, 1157–1174

Darden, T., York, D., and Pedersen, L. (1993) Particle mesh Ewald: an N-log(N) method for Ewald sums in large systems. J. Chem. Phys. 98, 10089–10092

Ryckaert, J.-P., Cicotti, G., and Berendsen, H. J. (1977) Numerical integration of the cartesian equations of motion of a system with constraints: molecular dynamics of n-alkanes. J. Comput. Phys. 23, 327–341

Jayaram, B., Sprous, D., and Beveridge, D. L. (1998) Solvation free energy of biomacromolecules: parameters for a modified generalized born model consistent with the AMBER force field. J. Phys. Chem. B 102, 9571–9576

Sievers, F., Wilm, A., Dineen, D., Gibson, T. J., Karplus, K., Li, W., Lopez, R., McWilliam, H., Remmert, M., Soding, I., Thompson, J. D., and Higgins,
D. G. (2011) Fast, scalable generation of high-quality protein multiple sequence alignments using Clustal Omega. Mol. Syst. Biol. 7, 537
57. Bozonnet, S., Dols-Laffargue, M., Fabre, E., Pizzut, S., Remaud-Simeon, M., Monsan, P., and Willemot, R.-M. (2002) Molecular characterization of DSR-E, an α-1,2 linkage-synthesizing dextranucrase with two catalytic domains. J. Bacteriol. 184, 5753–5761
58. Szklarczyk, R., and Heringa, J. (2004) Tracking repeats using significance and transitivity. Bioinformatics 20, i311–317
59. Armougom, F., Moretti, S., Poirot, O., Audic, S., Dumas, P., Schaeli, B., Keduas, V., and Notredame, C. (2006) Expresso: automatic incorporation of structural information in multiple sequence alignments using 3D-Coffee. Nucleic Acids Res. 34, W604–W608
60. Di Tommaso, P., Moretti, S., Xenarios, I., Orobitg, M., Montanyola, A., Chang, J.-M., Taly, J.-F., and Notredame, C. (2011) T-Coffee: a web server for the multiple sequence alignment of protein and RNA sequences using structural information and homology extension. Nucleic Acids Res. 39, W13–W17
61. Crooks, G. E., Hon, G., Chandonia, J.-M., and Brenner, S. E. (2004) WebLogo: a sequence logo generator. Genome Res. 14, 1188–1190
62. Vuillemin, M., Malbert, Y., Laguerre, S., Remaud-Siméon, M., and Moulis, C. (2014) Optimizing the production of an α-(1→2) branching sucrase in Escherichia coli using statistical design. Appl. Microbiol. Biotechnol. 98, 5173–5184
63. Takeo, K. (1995) Advances in affinity electrophoresis. J. Chromatogr. A 698, 89–105
64. Tomme, P., Boraston, A., Kormos, J. M., Warren, R. A., and Kilburn, D. G. (2000) Affinity electrophoresis for the identification and characterization of soluble sugar binding by carbohydrate-binding modules. Enzyme Microb. Technol. 27, 453–458
65. Abbott, D. W., and Boraston, A. B. (2012) Quantitative approaches to the analysis of carbohydrate-binding module function. Methods Enzymol. 510, 211–231
66. Davies, G. J., Wilson, K. S., and Henrissat, B. (1997) Nomenclature for sugar-binding subsites in glycosyl hydrolases. Biochem. J. 321, 557–559
67. Brison, Y., Fabre, E., Moulis, C., Portais, J.-C., Monsan, P., and Remaud-Siméon, M. (2010) Synthesis of dextrans with controlled amounts of α-1,2 linkages using the transglucosidase GBD–CD2. Appl. Microbiol. Biotechnol. 86, 545–554
68. Nihira, T., Mori, T., Asakura, M., and Okahata, Y. (2011) Kinetic studies of dextranucrase enzyme reactions on a substrate- or enzyme-immobilized 27 MHz quartz crystal microbalance. Langmuir 27, 2107–2111
69. Meng, X., Dobruchowska, J. M., Pijning, T., Gerwig, G. J., Kamerling, J. P., and Dijkhuizen, L. (2015) Truncation of domain V of the multidomain glucansucrase GTF180 of Lactobacillus reuteri 180 heavily impairs its polysaccharide-synthesizing ability. Appl. Microbiol. Biotechnol. 99, 5885–5894
Structural Insights into the Carbohydrate Binding Ability of an α-(1→2) Branching Sucrase from Glycoside Hydrolase Family 70
Yoann Brison, Yannick Malbert, Georges Czaplicki, Lionel Mourey, Magali Remaud-Simeon and Samuel Tranier

J. Biol. Chem. 2016, 291:7527-7540.
doi: 10.1074/jbc.M115.688796 originally published online February 10, 2016

Access the most updated version of this article at doi: 10.1074/jbc.M115.688796

Alerts:
• When this article is cited
• When a correction for this article is posted

Click here to choose from all of JBC's e-mail alerts

This article cites 69 references, 15 of which can be accessed free at
http://www.jbc.org/content/291/14/7527.full.html#ref-list-1