

Structural Enzymology of *Cellvibrio japonicus* Agd31B Protein Reveals α -Transglucosylase Activity in Glycoside Hydrolase Family 31^{*[5]}

Received for publication, September 13, 2012, and in revised form, November 5, 2012. Published, JBC Papers in Press, November 6, 2012, DOI 10.1074/jbc.M112.416511

Johan Larsbrink^{†1}, Atsushi Izumi^{§1}, Glyn R. Hemsworth[§], Gideon J. Davies^{§2}, and Harry Brumer^{†1,3}

From the [†]Division of Glycoscience, School of Biotechnology, Royal Institute of Technology, AlbaNova University Centre, 106 91 Stockholm, Sweden, [§]York Structural Biology Laboratory, Department of Chemistry, The University of York, York YO10 5DD, United Kingdom, and [¶]Michael Smith Laboratories and Department of Chemistry, University of British Columbia, 2185 East Mall, Vancouver, British Columbia V6T 1Z4, Canada

Background: Transglucosylases are important enzymes in bacterial glycogen metabolism.

Results: The tertiary structure and function of a novel α -transglucosylase have been defined.

Conclusion: In addition to previously known activities, glycoside hydrolase family 31 (GH31) contains a group of enzymes with 1,4- α -glucan 4- α -glucosyltransferase activity.

Significance: This gives new insight into bacterial glycogen utilization and will inform future bioinformatics analyses of (meta)genomes.

The metabolism of the storage polysaccharides glycogen and starch is of vital importance to organisms from all domains of life. In bacteria, utilization of these α -glucans requires the concerted action of a variety of enzymes, including glycoside hydrolases, glycoside phosphorylases, and transglucosylases. In particular, transglucosylases from glycoside hydrolase family 13 (GH13) and GH77 play well established roles in α -glucan side chain (de)branching, regulation of oligo- and polysaccharide chain length, and formation of cyclic dextrans. Here, we present the biochemical and tertiary structural characterization of a new type of bacterial 1,4- α -glucan 4- α -glucosyltransferase from GH31. Distinct from 1,4- α -glucan 6- α -glucosyltransferases (EC 2.4.1.24) and 4- α -glucanotransferases (EC 2.4.1.25), this enzyme strictly transferred one glucosyl residue from $\alpha(1\rightarrow4)$ -glucans in disproportionation reactions. Substrate hydrolysis was undetectable for a series of malto-oligosaccharides except maltose for which transglucosylation nonetheless dominated across a range of substrate concentrations. Crystallographic analysis of the enzyme in free, acarbose-complexed, and trapped 5-fluoro- β -glucosyl-enzyme intermediate forms revealed extended substrate interactions across one negative and up to three positive subsites, thus providing structural rationalization for the unique, single monosaccharide transferase activity of the enzyme.

Glycogen is a highly branched, mixed linkage $\alpha(1\rightarrow4)/\alpha(1\rightarrow6)$ -glucan polymer that serves as a readily accessible, osmotically neutral, cellular energy reserve in all domains of life (1–3). Glycogen is structurally related to amylopectin, which together with the linear polysaccharide amylose ($\alpha(1\rightarrow4)$ -glucan) comprises the plant storage reserve starch (4). Prokaryotic and eukaryotic glycogen biosynthesis and degradation are a complex, highly conserved, and tightly controlled process involving a myriad of enzymes and regulatory factors (2, 3). In bacteria such as *Escherichia coli*, glycogen is synthesized from ADP-glucose by the combined action of glycogen synthase, which builds linear $\alpha(1\rightarrow4)$ -glucan chains, and glycogen branching enzyme, which catalyzes chain rearrangement via $\alpha(1\rightarrow4)$ -to- $\alpha(1\rightarrow6)$ transglucosylation, thereby yielding poly-disperse molecules with molar masses of up to 10^7 – 10^8 Da (2, 3). In turn, catabolism under carbon-limited conditions occurs via the sequential action of glycogen phosphorylase and debranching enzyme to yield glucose 1-phosphate.

In bacteria, glycogen metabolism is closely linked to the metabolism of storage maltodextrins that involves the buildup and rearrangement of linear $\alpha(1\rightarrow4)$ -gluco-oligosaccharides by transglucosylation (5). This process has been well described in *E. coli* in which the amyломaltase MalQ, a member of glycoside hydrolase family 77 (GH77⁴; Ref. 6), catalyzes the transfer of a 4- α -glucanosyl fragment from the non-reducing end of malto-oligosaccharide donor substrates and possibly the disaccharide maltose to malto-oligosaccharide acceptors (4- α -glucanotransferase activity; EC 2.4.1.25) (5, 7, 8). The bacterial amyломaltases are structurally and functionally related to the plant disproportionating enzymes (“D-enzymes”) of GH77, which transfer maltosyl and longer 4- α -glucanosyl units from maltotriose and higher congeners (9). Likewise, certain thermophilic

* Work in Stockholm was supported by the Mizutani Foundation for Glycoscience and The Swedish Research Council Formas (via the European Union WoodWisdom-Net ERA-NET project “FiberSurf: New biotechnological tools for wood fiber surface modification and analysis” and via “CarboMat—the KTH Advanced Carbohydrate Materials Centre”), and work in York was supported by Biotechnology and Biological Sciences Research Council Grant BB/I014802/1.

[5] This article contains supplemental Figs. S1–S5.

The atomic coordinates and structure factors (codes 4b9y, 4b9z, and 4ba0) have been deposited in the Protein Data Bank (<http://www.pdb.org/>).

¹ Both authors contributed equally to this work.

² To whom correspondence may be addressed. Tel.: 44-1904-328260; Fax: 44-904-328266; E-mail: gideon.davies@york.ac.uk.

³ To whom correspondence may be addressed. Tel.: 604-827-3738; Fax: 604-822-2114; E-mail: brumer@msl.ubc.ca.

⁴ The abbreviations used are: GH, glycoside hydrolase family; HPAEC-PAD, high performance anion exchange chromatography with pulsed amperometric detection; *Cj*, *C. japonicus*; Bistris propane, 1,3-bis[tris(hydroxymethyl)methylamino]propane; pNP, *p*-nitrophenyl; *Ro*, *R. obcum*.

bacterial 4- α -glucanotransferases of GH13 catalyze the disproportionation of maltotriose (10, 11), maltotetraose (12, 13), and longer 4- α -glucan chains.

Indeed, transglycosylation reactions leading to the rearrangement of α -glucans are widespread among bacteria. Glycogen branching and debranching aside, diverse enzymes with 4- α -glucanotransferase activity ($\alpha(1\rightarrow4)$ -glucan: $\alpha(1\rightarrow4)$ -glucan transferase activity) can produce a range of linear and cyclic maltodextrin products via freely reversible disproportionation and cyclization reactions, respectively (9, 14). The production of six-, seven-, and eight-membered $\alpha(1\rightarrow4)$ -linked cyclodextrins by the cyclodextrin glucanotransferases (EC 2.4.1.19) of GH13 represents an especially important process in industrial starch valorization (15). Analogous $\alpha(1\rightarrow6)$ -linked cyclisomalto-oligosaccharides are the main products of some GH66 enzymes (16–18). Likewise, certain bacterial members of GH77 catalyze the production of large cyclic α -glucans (degrees of polymerization ≥ 22) through intramolecular transglycosylation (9). The GH31 enzymes CtsY and CtsZ from *Arthrobacter* and *Sporosarcina* species generate commercially interesting cycloalternan tetrasaccharides (cyclo[$\rightarrow 6$]- α -D-Glcp-(1 $\rightarrow 3$)- α -D-Glcp-(1 $\rightarrow 6$)- α -D-Glcp-(1 $\rightarrow 3$)- α -D-Glcp-(1 \rightarrow)] from $\alpha(1\rightarrow4)$ -glucans via a remarkable three-step, coupled reaction involving $\alpha(1\rightarrow4)$ -to- $\alpha(1\rightarrow6)$ transglycosylation, intermolecular isomaltosyl transfer, and cyclization (19, 20). In addition to these predominant transglycosylases, a number of retaining α -glucoside hydrolases with degrees of transglycosylation ability have also been identified (Refs. 21–23 and references therein).

The Gram-negative soil saprophyte *Cellvibrio japonicus* is best known for its ability to efficiently utilize a plethora of plant cell wall polysaccharides as energy sources (24). Additionally, the genome sequence of this organism has revealed a large number of predicted α -glucan-active enzymes. In total, the *C. japonicus* genome encodes 22 enzymes from GH13, GH15, GH31, GH57, and GH77 (25) that may be predicted to act on starch and/or glycogen. However, none of these have been biochemically or structurally characterized (6, 25).

GH31 in particular is one of the major α -glucosidase-containing glycoside hydrolase families. This family is functionally diverse; it also contains α -xylosidases and α -glucan lyases in addition to the aforementioned CtsY and CtsZ α -transglycosylases. A phylogenetic analysis has recently been presented that partially delineates these activities in clades, although sequence-based functional prediction is not absolute (26). The generally *exo*-acting GH31 enzymes, which are members of Clan GH-D together with GH27 and GH36, have been suggested to share a common ancestor with members of clan GH-H, which comprises generally *endo*-acting α -glucan-active enzymes of GH13, GH70, and GH77 (27).

Building upon our interest in the postgenomic characterization of GH31 enzymes from *C. japonicus* (26), we present here a detailed structural enzymology study of *CjAgd31B*, whose coding sequence resides within a gene cluster encoding predicted α -glucan-active enzymes and sugar transporters. Biochemical analysis revealed that *CjAgd31B* is a predominant transglucosylase with strict $\alpha(1\rightarrow4)$ linkage specificity, which represents a previously undiscovered activity in GH31. Crystal-

lography of the enzyme in free, acarbose-complexed, and trapped 5-fluoro- β -D-glucopyranosyl-enzyme intermediate forms has highlighted the structural basis for the strict transfer of a single glucosyl residue and preference for maltotriose and longer substrates. Taken together, the data suggest a biological role for *CjAgd31B* in glycogen or maltodextrin metabolism that may be complementary to that predicted for the GH77 homologue *CjMal77Q*.

EXPERIMENTAL PROCEDURES

Curve fitting and processing of kinetics data were performed using Origin 8 software (OriginLab). *p*-Nitrophenyl (*p*NP) α -glycosides, sucrose, D-maltose, and starch from corn were purchased from Sigma. Malto-oligosaccharides (maltotriose to maltohexaose), isomaltose, melibiose, and acarbose were purchased from Carbosynth. α -Glucosyl fluoride and 5-fluoro- α -D-glucopyranosyl fluoride were kind gifts from Professor Stephen Withers (Department of Chemistry, University of British Columbia, Canada). Ultrapure water was used in all experiments and refers to water purified on a Milli-Q system (Millipore) with a resistivity (ρ) > 18.2 megaohms \cdot cm.

Cloning of *CjAgd31B*

The open reading frame encoding *CjAgd31B* (GenBank accession number ACE84782.1) was amplified by PCR from genomic DNA of *C. japonicus* Ueda107 using Phusion polymerase (Finnzymes) and the following primers (Thermo Fischer Scientific): 5'-CACCATGAATCCGGTCAAACG-3' and 5'-ATGCAACCTGAGGTTAAGCGCTTC-3' with the forward primer incorporating the CACC overhang (underlined) needed for TOPO cloning and excluding the predicted signal peptide (cleavage site between amino acid residues 24 and 25). The PCR product was cloned into the pENTR/SD/D-TOPO entry vector (Invitrogen) and recombined into the pET-DEST42 destination vector (Invitrogen) as described previously (26).

Gene Expression and Protein Purification

Plasmids harboring the *CjAgd31B* gene were transformed into *E. coli* BL21(DE3) by electroporation, the gene was expressed and the resulting protein was purified by immobilized metal affinity chromatography following an established protocol (26). Analysis by SDS-PAGE showed the protein to be electrophoretically pure. LC electrospray ionization MS was used for protein molar mass determination as described previously (28). For crystallization studies, the protein was further purified by size exclusion chromatography and ion exchange chromatography. The eluted protein solution was concentrated to 5 ml by a Vivaspin 20 concentrator (Sartorius Stedim Biotech) and loaded onto a HiLoad 16/60 Superdex 200 prep grade column (GE Healthcare) equilibrated with 20 mM Tris (pH 8.0), 300 mM sodium chloride. The eluted protein solution was dialyzed into 20 mM Tris (pH 8.0) at 4 °C for 16 h. The dialyzed protein solution was loaded onto a Resource Q column (GE Healthcare) equilibrated with 20 mM Tris (pH 8.0) and eluted with a linear gradient of 20 mM Tris (pH 8.0), 400 mM sodium chloride. Two major peaks of *CjAgd31B* were obtained, and the peak eluted in lower salt concentration was collected and used for protein crystallization. Protein concentrations were deter-

A Bacterial α -Transglucosylase from GH31

mined from A_{280} values of suitably diluted samples using an extinction coefficient of $139,245 \text{ M}^{-1}\text{cm}^{-1}$ as calculated by the ProtParam tool on the ExPASy server (29).

Thin Layer Chromatography (TLC)

TLC was performed using normal phase silica on aluminum plates eluted with acetonitrile-water (2:1). Analytes were visualized by immersion in 8% H_2SO_4 in ethanol followed by charring.

High Performance Anion Exchange Chromatography with Pulsed Amperometric Detection (HPAEC-PAD)

Oligo- and monosaccharides were analyzed on a Dionex ICS-3000 HPLC system operated by Chromelion software version 6.80 (Dionex) using a Dionex CarboPac PA200 column. Solvent A was water, solvent B was 1 M sodium hydroxide, and solvent C was 1 M sodium acetate.

Gradient A—Conditions used were 0–5 min, 10% B, 2% C; 5–12 min, 10% B and a linear gradient from 2 to 30% C; 12–12.1 min, 50% B, 50% C; 12.1–13 min, an exponential gradient of B and C back to the initial conditions; 13–17 min, initial conditions.

Gradient B—Conditions used were 0–4 min, 10% B, 5% C; 4–8 min, 10% B and a linear gradient from 5 to 25% C; 8–8.1 min, 50% B, 50% C; 8.1–9 min, an exponential gradient of B and C back to the initial conditions; 9–13 min, initial conditions.

Gradient C—Conditions used were 0 to 4 min, 10% B, 6% C; 4 to 17 min, 10% B and a linear gradient from 6–25% C; 17 to 17.1 min, 50% B, 50% C; 17.1 to 18 min, an exponential gradient of B and C back to initial conditions; 18 to 22 min, initial conditions.

Gradient D—Conditions used were 0–4 min, 10% B, 6% C; 4–10 min, 10% B and a linear gradient from 5 to 25% C; 10–10.1 min, 50% B, 50% C; 10.1–11 min, an exponential gradient of B and C back to the initial conditions; 11–15 min, initial conditions.

Gradient E—Conditions used were 0–4 min, 10% B, 6% C; 4–15 min, 10% B and a linear gradient from 5 to 25% C; 15–15.1 min, 50% B, 50% C; 15.1–16 min, an exponential gradient of B and C back to the initial conditions; 16–20 min, initial conditions.

pH-Rate Profile

Measurements of the pH dependence of *CjAgd31B* catalysis were carried out using maltose as substrate in an HPAEC-PAD-based transglycosylation assay described below. The buffers used (50 mM) were sodium citrate (pH 3–6.5), sodium phosphate (pH 6.5–8), glycylglycine (pH 8–9), and glycine (pH 9–10) (supplemental Fig. S1).

Enzyme Assays

Activity on *p*NP-glycosides was analyzed by a stopped assay as described previously (26) using an enzyme concentration of $6.5 \mu\text{M}$. The transglycosylation activity of *CjAgd31B* on various oligosaccharides was performed in $100\text{-}\mu\text{l}$ reactions at 25°C in 50 mM citrate buffer (pH 6). For initial rate saturation kinetics experiments, *CjAgd31B* was added to a final concentration of $1.6 \mu\text{M}$ for maltose and 270 pM for maltotriose, maltotetraose, and maltopentaose. Reactions typically proceeded for 10 min and were stopped by addition of $4 \mu\text{l}$ of 5 M sodium hydroxide.

HPAEC-PAD was used for product analysis using Gradients A, B, D, and E for reactions on maltose, maltotriose, maltotetraose, and maltopentaose, respectively. Commercial malto-oligosaccharides (maltose to maltohexaose) were used as standards.

To test different acceptors, starch was used as a glucosyl donor. Starch from corn was dissolved to 1% (w/v) in water followed by dialysis in deionized water using a 5-kDa-cut-off membrane to remove monosaccharides and small oligosaccharides. $50\text{-}\mu\text{l}$ reactions containing 0.4% starch as donor, 1 mM acceptor (glucose or isomaltose), and $2 \mu\text{M}$ enzyme were incubated at 25°C for 10 min and terminated by addition of $2 \mu\text{l}$ of 5 M sodium hydroxide. Products were analyzed by HPAEC-PAD using Gradient C.

IC_{50} Measurements

The inhibition of *CjAgd31B* by acarbose ($0\text{--}1050 \mu\text{M}$) was determined by using maltotriose ($100 \mu\text{M}$) as a substrate in reactions as described above. Product formation was analyzed by HPAEC-PAD (Gradient B).

Crystallization and Data Collection

CjAgd31B was stored in 5 mM Bistris propane (pH 8.5) and concentrated to 7 mg/ml by using a Vivaspin 20 concentrator. In initial crystal screens using Crystal Screen HT, Index HT, SaltRx HT (Hampton Research), and modified Newcastle Screen prepared at the York Structural Biology Laboratory, small single crystals were obtained in several conditions. Well diffracting crystals were obtained after 3–4 days in 1.8 M ammonium sulfate, 0.1 M HEPES (pH 7.0), 2% PEG 400 at 20°C by the sitting drop vapor diffusion method. The structure was solved using experimental phasing with an iodine derivative. This was prepared by placing $\sim 1 \mu\text{l}$ of a 0.25 g/ml potassium iodide solution into a $2\text{-}\mu\text{l}$ crystallization droplet to allow slow diffusion of iodine into the crystal. Crystallization droplets with iodine solution were left at 20°C for 16 h prior to freezing and data collection. For the complex structures, crystals were soaked in 1.8 M ammonium sulfate, 0.1 M HEPES (pH 7.0), 2% PEG 400 with either 5 mM 5-fluoro- α -D-glucopyranosyl fluoride or 5 mM acarbose for 1 h at 20°C .

All crystals were cryoprotected by 2.0 M lithium sulfate, 0.1 M HEPES (7.0), 2% PEG 400. The x-ray data for the free enzyme and iodine-soaked crystals were collected at 100 K on an ADSC Q315 charge-coupled device detector on beamline I02 of the Diamond Light Source. X-ray data for the complex with acarbose and the 5F β Glc-enzyme were collected at 100 K on an ADSC Q315 charge-coupled device detector at BL-ID14-4 and BL-ID29 at the European Synchrotron Radiation Facility, respectively. The details of the data collections are listed in Table 1. All data were processed using iMOSFLM (30) and programs from the CCP4 suite (31) unless otherwise stated. The statistics of the data processing and structure refinement are listed in Table 1.

Experimental phasing was performed by single wavelength anomalous diffraction methods at a wavelength of 1.8 Å. Heavy atom substructure solution and initial phasing was performed on the 2.9-Å resolution iodine-derivatized crystal data with autoSHARP (32) followed by phase extension with the 1.9-Å free enzyme data set using DM (33). The 1.9-Å data were used

TABLE 1

X-ray data collection and refinement statistics

OXL, oxalate; EDO, 1,2-ethanediol; PEG, 2-(2-hydroxyethoxy)ethanol; PGE, 2-[2-(2-hydroxyethoxy)ethoxy]ethanol; PG4, 2-[2-(2-hydroxyethoxy)ethoxy]ethoxy]ethanol; r.m.s.d., root mean square; 5F β Glc, 5-fluoro- β -glucosyl; ESRF, European Synchrotron Radiation Facility; SO₄, sulfate.

	Name (Protein Data Bank code)		
	Free enzyme (4b9y)	5F β Glc (4ba0)	Acarbose (4b9z)
Data collection			
Beamline	Diamond I02	ESRF ID29	ESRF14-4
Space group	P622	P622	P622
Cell dimensions			
<i>a</i> , <i>b</i> , <i>c</i> (Å)	197.3, 197.3, 103.0	197.2, 197.2, 103.1	196.9, 196.9, 102.8
α , β , γ (°)	90, 90, 120	90, 90, 120	90, 90, 120
Resolution (Å) (outer shell)	49.85–1.90 (2.00–1.90)	44.48–1.85 (1.95–1.85)	51.38–2.0 (2.11–2.00)
<i>R</i> _{merge}	0.10 (0.52)	0.10 (0.52)	0.11 (0.46)
<i>I</i> / σ <i>I</i>	21.5 (6.0)	19.8 (5.9)	22.2 (7.4)
Completeness (%)	100.0 (100.0)	100.0 (100.0)	100.0 (100.0)
Redundancy	22.0	20.8	21.5
Refinement			
Resolution (Å)	49.3–1.90	44.5–1.85	49.7–2.00
No. reflections	92,667	100,216	79,078
<i>R</i> _{work} / <i>R</i> _{free}	0.16/0.19	0.17/0.19	0.16/0.19
No. atoms			
Protein	6,273	6,270	6,251
Ligand	93	80	142
Water	647	557	444
B-factors (Å ²)			
Protein (TLS refinement)	27.9	31.2	29.3
Water	33.8	36.8	32.4
Ligands	SO ₄ , 80.8; PG4, 44.8; OXL, 42.7; EDO, 48.2	5F β Glc, 21.2; SO ₄ , 71.3; PGE, 47.7; Arg, 48.2; PEG, 34.4; EDO, 53.8	Acarbose, 40.2; SO ₄ , 55.3; OXL, 55.1; PEG, 37.5; EDO, 53.1
r.m.s.d.			
Bond lengths (Å)	0.007	0.006	0.007
Bond angles (°)	1.1	1.1	1.1

TABLE 2

Iodine single wavelength anomalous diffraction (SAD) phasing statistics

FOM, figure of merit.

Data collection	
Beamline	Diamond I02
Wavelength (Å)	1.80
Space group	P622
Cell dimensions	
<i>a</i> , <i>b</i> , <i>c</i> (Å)	197.7, 197.7, 102.6
α , β , γ (°)	90, 90, 120
Resolution (Å) (outer shell)	71.17–2.9 (3.06–2.90)
Total no. of measured reflections	1,127,119
No. of unique reflections	26,692
<i>R</i> _{merge}	0.16 (0.74)
<i>I</i> / σ <i>I</i>	24.9 (6.7)
Anomalous completeness (%)	100.0 (100.0)
Anomalous redundancy	22.6 (22.5)
FOM SAD/density modification	0.17/0.84

as a starting point for automatic model building using ARP/wARP (34) (Table 2). Structure refinement, including TLS refinement of molecular motions, was performed using PHENIX (35) interspersed with manual rebuilding using COOT (36). Complex structures were solved by molecular replacement using MOLREP (37) with the free enzyme structure as a search model and refined as above.

RESULTS

Bioinformatics Analyses—*CjAgd31B* is found among putative receptors, transporters, and α -glucan-active enzymes in the genome of *C. japonicus* (supplemental Fig. S2). Notably, these genomic neighbors include a predicted α -amylase, cyclomaltodextrin glucanotransferase, 6-phospho- β -glucosidase, and a glucokinase. *CjAgd31B* has a predicted secretion signal peptide and is thus likely to be localized in the periplasm or to be secreted extracellularly. In addition to *CjAgd31B*, the genome of *C. japonicus* encodes two other GH31 members, the putative

α -glucosidase *CjAgd31A* and the biochemically and structurally characterized α -xylosidase *CjXyl31A* (26). Although all are members of GH31, *CjAgd31B* has a low sequence similarity to both *CjAgd31A* and *CjXyl31A* with amino acid identities of 28 and 27% and similarities of 43 and 45%, respectively. From our recent phylogenetic analysis (26), the biochemically characterized member of GH31 most similar to *CjAgd31B* is YihQ from *E. coli* (sequence identity, 28%; similarity, 44%). *E. coli* YihQ has been annotated as an α -glucosidase based on a weak ability to hydrolyze the artificial substrate α -glucosyl fluoride, although the enzyme was impotent toward a range of other α -glucosides (38).

Gene Expression—A gene construct encoding *CjAgd31B* with a C-terminal hexahistidine tag and lacking the predicted native signal peptide was expressed in *E. coli* BL21(DE3) cells. The protein product was purified by immobilized metal affinity chromatography for kinetics analyses and additionally by size exclusion chromatography and ion exchange for crystallization studies; purity in both cases was confirmed by SDS-PAGE (data not shown). The molar mass of *CjAgd31B*, corresponding to the C-terminal His-tagged enzyme starting at Asn-25 (the natural site of signal peptide cleavage), was verified by LC electrospray ionization MS (expected, 94,478.8 Da; observed, 94,478.1 Da; supplemental Fig. S3). The overall yield was typically around 100 mg/liter of culture broth.

Transglycosylation Activity on Malto-oligosaccharides—Based on membership in GH31, the substrate specificity of *CjAgd31B* was initially tested using *p*NP- α -glucoside and *p*NP- α -xyloside; the enzyme showed no apparent liberation of the aglycone from either of these substrates after extended incubation (1 mM substrate and 6.5 μ M enzyme and up to 4-h incubation). The enzyme also displayed no detectable activity on sucrose, meli-

A Bacterial α -Transglucosylase from GH31

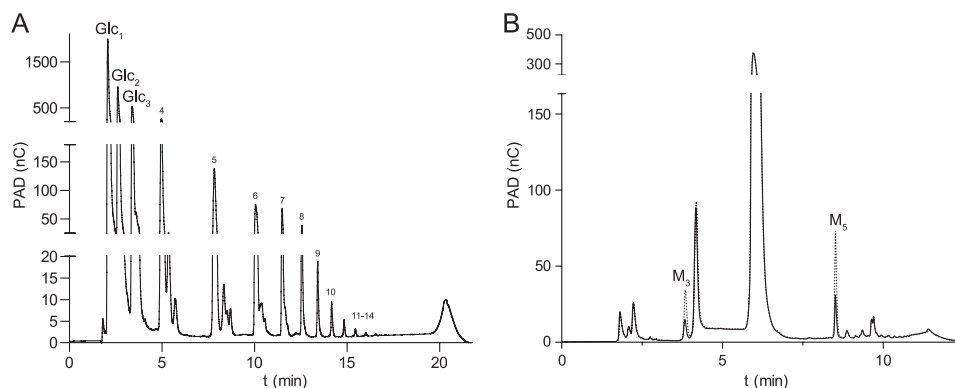


FIGURE 1. HPAEC-PAD chromatograms of reaction products of malto-oligosaccharides, catalyzed by *CjAgd31B*. *A*, incubation with maltose (10 mM; 6.5 mM enzyme; 30 min). Shoulder peaks are assumed to be transglycosylation products arising from contaminants in the commercial maltose preparation. *B*, incubation with maltotetraose (0.5 mM; 10-min reaction). *Solid line*, commercial maltotetraose sample prior to addition of enzyme; *dotted line*, enzyme reaction indicating production of maltotriose (M_3) and maltopentaose (M_5) by disproportionation. Commercial malto-oligosaccharides were used as standards to determine retention times. nC, nanocoulomb.

biose, isomaltose, or α -glucosyl fluoride (a substrate for *E. coli* YihQ (38)). On maltose (Glc α (1 \rightarrow 4)Glc), however, formation of both glucose and longer oligosaccharides could be observed using TLC (data not shown). The transglycosylation potential of *CjAgd31B* was confirmed by HPAEC-PAD following incubation of 6.5 μ M enzyme with 10 mM maltose for 30 min, which led to a buildup of malto-oligosaccharides. Products with a degree of polymerization of up to 14 glucose residues could be detected (Fig. 1A).

To further analyze the catalytic properties of *CjAgd31B*, HPAEC-PAD was used to measure product formation from reactions on malto-oligosaccharides. Under conditions of low substrate conversion (<10% of substrate consumed), the enzyme was shown to transfer a single glucose moiety from a donor to an acceptor molecule. Incubation of the enzyme with linear malto-oligosaccharides (maltotriose to maltopentaose) exclusively yielded Glc $_{n-1}$ and Glc $_{n+1}$ products via transglycosylation (Fig. 1B). On maltotriose, maltotetraose, and maltopentaose, the production of glucose, which would indicate competing substrate hydrolysis, was not observed under initial rate conditions. Thus, apparent Michaelis-Menten kinetics parameters for these substrates could be directly determined from plots of $v_0/[E]_t$ versus [Glc $_n$] (Fig. 2A) where the rate of transglycosylation product formation is given by Equation 1 (in this case, the rate of Glc $_{n-1}$ formation can also be used). The best substrate for the enzyme was maltotriose with a $(k_{cat}/K_m)_{app}$ value of 196 s $^{-1}$ ·mM $^{-1}$, whereas maltotetraose and maltopentaose displayed comparable $(k_{cat}/K_m)_{app}$ values of 72 and 58 s $^{-1}$ ·mM $^{-1}$, respectively (Table 3).

$$v_{\text{transglycosylation}} = \frac{d[\text{Glc}_{n+1}]}{dt} \quad (\text{Eq. 1})$$

Hydrolysis could, however, be detected when maltose was used as a substrate. Here, the production of Glc was measurably higher than the 1:1 stoichiometric ratio of Glc to Glc $_3$ expected for disproportionation (Glc $_2 \rightarrow$ Glc $_3$ + Glc). In this case, the velocity of the transglycosylation reaction was directly measured according to Equation 1 by quantifying the maltotriose produced. Determination of the hydrolytic rate required subtraction of the amount of glucose co-produced by disproportionation

from that arising from hydrolysis, including accounting for the stoichiometry of glucose release by hydrolysis (2 mol of Glc/mol of maltose) (Equation 2). From the specific transglycosylation activity on 0.5 mM maltose, the pH profile was found to be broad with an optimum of pH 6.5 (supplemental Fig. S1).

$$v_{\text{hydrolysis}} = \frac{d\left(\frac{[\text{Glc}] - [\text{maltotriose}]}{2}\right)}{dt} \quad (\text{Eq. 2})$$

Fig. 2B shows plots of $v_0/[E]_t$ versus [Glc $_2$] for both the hydrolytic and transglycosylation reactions, which clearly indicate the predominance of transglycosylation over a wide range of substrate concentrations. The $(k_{cat}/K_m)_{app}$ value determined for the transglycosylation reaction was 39 s $^{-1}$ ·mM $^{-1}$ (Table 3). Although the $(k_{cat})_{app}$ value for maltose transglycosylation is actually higher than that of maltotriose, the lower $(k_{cat}/K_m)_{app}$ value on the disaccharide is due to a high $(K_m)_{app}$ value. This high value may reflect the observation of up to four enzyme subsites (−1 to +3; nomenclature according to Ref. 39) in the crystal structure (see below) of which only two would be occupied by maltose acting as a donor substrate.

Across the range of maltose concentrations examined (50 μ M to 50 mM), the rate of hydrolysis was consistently low (Fig. 2B). At 50 μ M maltose, the rate of hydrolysis was equal to the rate of transglycosylation ($v_0/[E]_t = 2$ s $^{-1}$) after which the hydrolytic rate increased to a maximum of 10 s $^{-1}$ at 2 mM substrate. Nonetheless, the hydrolytic rate was only 13% of the transglycosylation rate at this maximum. The rate of hydrolysis steadily decreased to ~ 8 s $^{-1}$ at 50 mM maltose as the transglycosylation rate continued to increase, resulting in a relative hydrolysis rate of <2% at this concentration.

Starch and Isomaltose as an Alternate Donor/Acceptor Pair—Starch was also tested as a donor substrate with high degree of polymerization with glucose and isomaltose as alternate acceptor substrates. Despite dialysis to reduce the content of short malto-oligosaccharides, the commercial starch sample contained a minor amount of α (1 \rightarrow 4)-glucans with a degree of polymerization of 2 and higher. When *CjAgd31B* was incubated with starch alone, the amounts of shorter malto-oligosac-

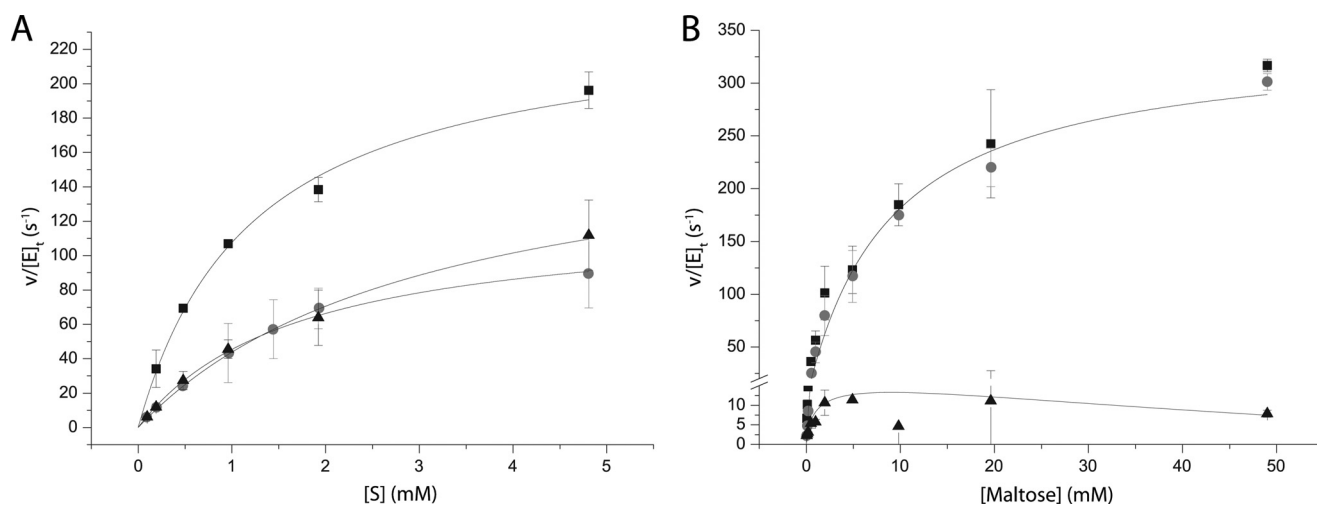


FIGURE 2. A, initial rate kinetics curves fitted to the Michaelis-Menten equation for maltotriose (black squares), maltotetraose (gray circles), and maltopentaose (black triangles). B, initial rate kinetics curves fitted to the Michaelis-Menten equation for maltose. Black squares represent formed glucose, gray circles represent maltotriose, and black triangles represent the rate of hydrolysis. Error bars represent two standard deviations from the mean value.

TABLE 3
Transglycosylating activity of CjAgd31B on various substrates

$\Delta\Delta G^\ddagger$ was calculated using the following formula: $\Delta\Delta G^\ddagger = RT \ln(k_{\text{cat}}/K_m[\text{maltotriose}]/k_{\text{cat}}/K_m[\text{maltooligo}])$. No activity was detected on pNP- α -Glc, pNP- α -Xyl, α -glucosyl fluoride, isomaltose, sucrose, or melibiose.

Substrate	$(k_{\text{cat}})_{\text{app}}$ s^{-1}	$(K_m)_{\text{app}}$ mM	$(k_{\text{cat}}/K_m)_{\text{app}}$ $s^{-1} \cdot \text{mM}^{-1}$	$\Delta\Delta G^\ddagger$ kJ mol^{-1}
Maltose	341 ± 12	8.8 ± 0.87	38.8	4.17
Maltotriose	239 ± 8.3	1.2 ± 0.13	195.5	
Maltotetraose	123 ± 9.0	1.7 ± 0.29	72.3	2.57
Maltopentaose	181 ± 17	3.1 ± 0.55	57.8	3.14

charides (maltose to maltotetraose) still present in the substrate mixture decreased significantly, whereas longer malto-oligosaccharides (maltopentaose to maltododecaose) increased in keeping with previous observations on purified malto-oligosaccharides during kinetics analyses (Fig. 3A). In contrast, when the reaction was supplemented with 1 mM glucose, a large increase of malto-oligosaccharides (maltose to maltododecaose) could be observed (Fig. 3B). Here, glucose (which as a monosaccharide cannot act as a glycosyl donor) acted as viable acceptor substrate in the breakdown of the glycosyl-enzyme intermediate formed via initial attack of starch chain ends. This reaction is the reverse of glycosyl-enzyme formation when maltose acts as a donor substrate, consistent with the principle of microscopic reversibility. The large increase in medium length malto-oligosaccharides compared with the glucose-free reaction both demonstrates how the non-reducing ends present in starch are utilized as glucose donors and suggests that glucose moieties are mainly “shuffled” between the non-reducing ends of the starch molecules when no additional acceptor is included in the reactions.

The disaccharide isomaltose (Glc α (1 \rightarrow 6)-Glc) on its own was not a substrate for CjAgd31B, indicating that the enzyme was unable to utilize α (1 \rightarrow 6)-linked glucosides as donor substrates. Glycogen and starch both contain α (1 \rightarrow 6)-linked branch points, which prompted us to test whether such branches might be extended by CjAgd31B using isomaltose as a model. Indeed, the enzyme transferred glucosyl moieties from starch to isomaltose (1 mM) as indicated by the appearance of an alternate series of peaks on the chromatogram, each slightly

preceding the corresponding all- α -linked congener, which suggests the formation of Glc $_n$ - α (1 \rightarrow 4)-Glc- α (1 \rightarrow 6)-Glc saccharides (Fig. 3B). As such, these data indicated that the +2 subsite of the active site is not strictly specific for α (1 \rightarrow 4)-linked sugars but will accommodate α (1 \rightarrow 6)-linked isomers.

Inhibition by Acarbose—The inhibitory effect of the pseudotetrasaccharide acarbose, a common α -glucanase and α -glucosidase inhibitor, was assayed using maltotriose as the substrate at a fixed concentration of 100 μM . The IC_{50} value was determined to be $75.1 \pm 3.4 \mu\text{M}$ by plotting the relative activity versus the concentration of acarbose and fitting Equation 3 by non-linear regression (Fig. 4). With reactions performed at a substrate concentration of 100 μM , which is much lower than the apparent K_m value (1.2 mM), the IC_{50} value is approximately equal to the K_i value. The K_i may be more accurately calculated using Equation 4 (40), which yielded 81.4 μM .

$$k_{\text{obs}} = \frac{k_{\text{max}}}{1 + \frac{[I]}{\text{IC}_{50}}} \quad (\text{Eq. 3})$$

$$K_i = \frac{\text{IC}_{50}}{1 + \frac{(S)}{K_m}} \quad (\text{Eq. 4})$$

The calculated K_i value of CjAgd31B is similar to K_i values for acarbose with other GH31 enzymes such as the human maltase-glucoamylase N-terminal subunit (62 μM ; Ref. 41) and sucrose N-terminal subunit (14 μM ; Ref. 42), whereas it is sig-

A Bacterial α -Transglucosylase from GH31

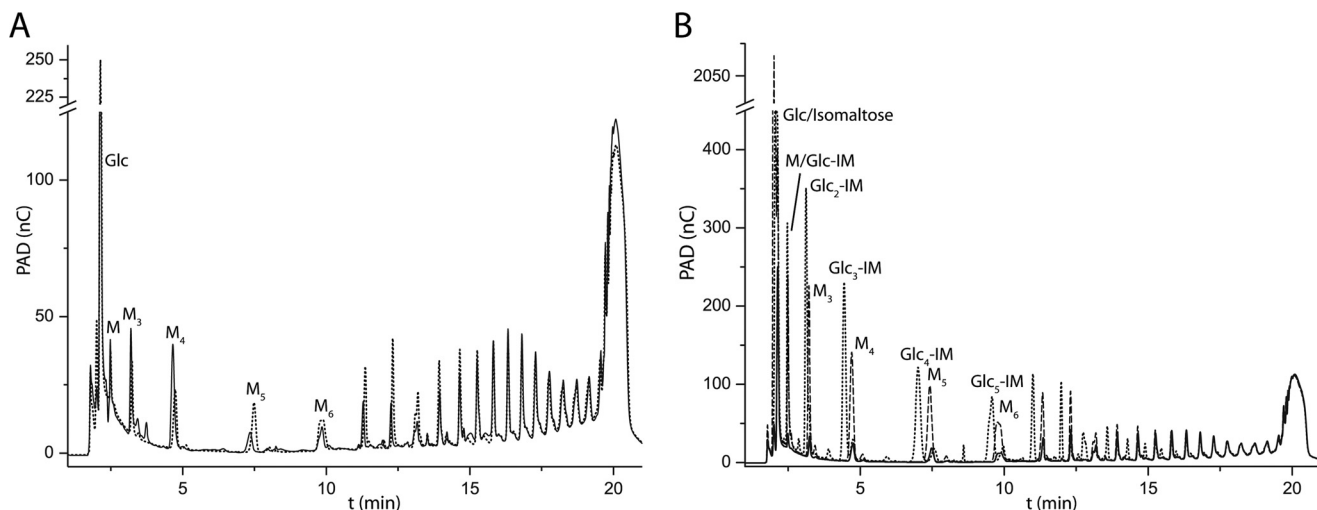


FIGURE 3. HPAEC-PAD chromatograms of donor/acceptor specificity experiments using starch, isomaltose (IM), and glucose. *A*, starch blank (0.4%, w/v; solid line) and following incubation with CjAgd31B (dotted line) showing a reduction of short malto-oligosaccharides (M_1 to M_5) and an increased amount of medium length malto-oligosaccharides (M_7 to M_{12}). *B*, starch (0.4%) following incubation with CjAgd31B with no additional acceptor substrate added (solid line), with added isomaltose (1 mM; dashed line), and with added glucose (1 mM; dotted line). With both alternate acceptor substrates, there is a clear increase in the amount of short to medium length oligosaccharides formed compared with starch alone. nC, nanocoulomb.

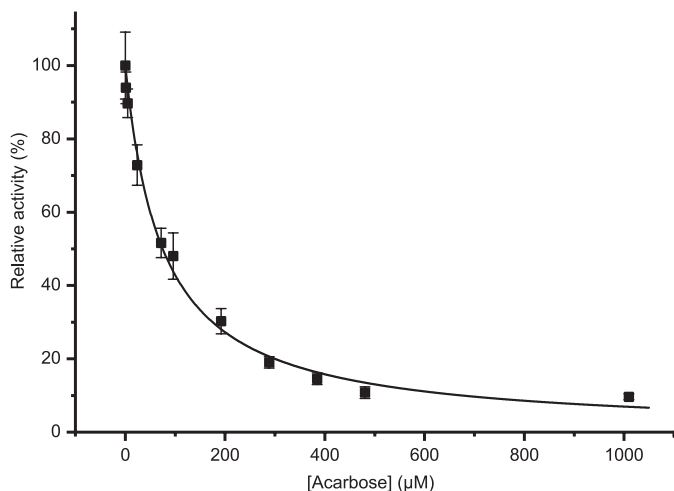


FIGURE 4. Inhibition of CjAgd31B by acarbose. Maltotriose was used as substrate (100 μ M; 10-min reaction), and the inhibitory effect of acarbose on transglucosylation (formation of maltotetraose) was calculated by non-linear regression, fitting Equation 3 to the data. nC, nanocoulomb. Error bars represent two standard deviations from the mean value.

nificantly higher than the K_i values for the human maltase-glucoamylase C-terminal subunit (1.72 μ M; Ref. 43), the *Gracilariopsis lemaneiformis* α -glucan lyase (0.02 μ M; Ref. 44), and the *Thermoplasma acidophilum* AglA α -glucosidase (2.99 μ M; Ref. 23).

Tertiary Structures of CjAgd31B—Three tertiary structures of CjAgd31B were obtained at 1.9-, 2.0-, and 1.85- \AA resolution, respectively: the free enzyme, a non-covalent complex with the inhibitor acarbose, and a trapped 5-fluoro- β -D-glucopyranosyl-enzyme intermediate. All crystals contained one molecule in the asymmetric unit. The CjAgd31B structures consisted of a typical GH31 fold comprising four domains with two insertions (Fig. 5A): the N-terminal domain (N-terminal; residues 35–240), the catalytic (β/α)₈ domain (residues 241–586) with insertion domain 1 (Insert 1; residues 345–384) and insertion domain 2 (Insert 2; residues 415–435), the C-terminal proximal

domain (C-proximal; residues 587–667), and the C-terminal distal domain (C-distal; residues 668–817). In all structures, the electron density map of the 10 N-terminal residues from 25 to 34 and C-terminal residues from 818 to the end (859), including the V5 epitope and His tag provided by the expression vector, were disordered. The free and acarbose structures had a disordered region from 137 to 140, and the 5-fluoro- β -glucosyl-enzyme structure had a disordered region from 139 to 140. The visible secondary structures of the four domains, including the insertion domains, in CjAgd31B were well conserved with the human sucrase-isomaltase (Protein Data Bank code 3lpp) of GH31 with a root mean square deviation of 1.9–2.4 \AA .

The free enzyme structure reveals a water-lined pocket where the conserved catalytic aspartic acid residues (Asp-412 and Asp-480) are located (45, 46). The pocket was 318 \AA^3 as calculated by the Pocket-Finder server (47) with a depth of \sim 11 \AA . The active site pocket was composed of residues Tyr-179, Phe-271, Asp-299, Leu-300, Ile-307, Met-311, Ile-341, Glu-343, Tyr-376, Phe-377, Trp-410, Asp-412, Leu-413, Glu-417, Arg-463, Trp-477, Asp-480, Asp-509, Phe-513, Arg-538, His-540, and Gln-542 (supplemental Fig. S4). To define enzyme-substrate interactions, two complexes, with the inhibitory tetrasaccharide acarbose (see above) and the covalent 5-fluoro- β -D-glucopyranosyl-enzyme intermediate, were obtained.

Structure of the Acarbose Complex—The CjAgd31B complex with acarbose revealed clear, unambiguous density for the inhibitor in the -1 to $+3$ subsites (Fig. 5B; subsite nomenclature according to Ref. 39). In the -1 subsite, the enzyme-derived nucleophile, Asp-412, is indeed poised for nucleophilic attack, lying 3.2 \AA “above” the pseudo-anomeric carbon of acarbose and with a nucleophile-C1-NH angle of 164.1°. The catalytic acid, Asp-480, lies 2.5 \AA from the “interglycosidic” nitrogen of acarbose as expected. The hydrophobic residues Leu-300, Ile-341, Trp-410, Trp-477, Phe-271, and Phe-513 all lie within a 4- \AA distance from the -1 subsite pseudosugar of acarbose (supplemental Fig. S4).

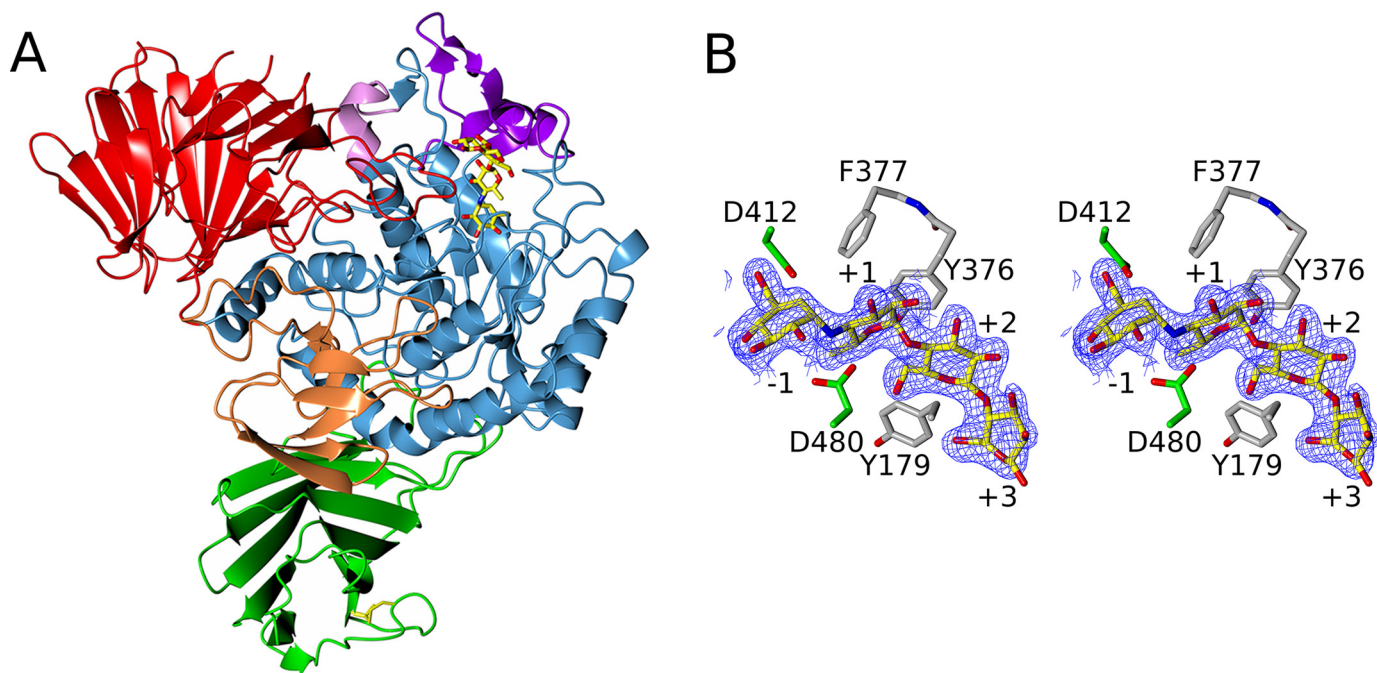


FIGURE 5. **The three-dimensional structure and ligand binding site of CjAgd31B.** *A*, three-dimensional structure of CjAgd31B as a protein schematic. The N-terminal β -sandwich domain is colored red; the central catalytic (β/α)₈ domain is shown in blue with Inserts 1 and 2 colored purple and pink, respectively; the C-terminal proximal β -sandwich is orange; and the C-terminal distal β -sandwich domain is green. *B*, observed electron density for the acarbose complex of CjAgd31B (map is at 1 σ in divergent stereo). The nucleophile Asp-412, acid-base Asp-480, and hydrophobic “clamp” residues (discussed in text) are shown. This figure was drawn with CCP4MG (66).

The +1 subsite contains the 6-deoxyglucosyl moiety of acarbose. Hydrogen bonds are made to Arg-463, Glu-417, and a water molecule, and enzyme-substrate distances suggest van der Waals contacts to Phe-377. In most other solved structures of GH31, the hydrogen bonds provided here by Glu-417 (Insert 2) are made instead by an aspartate residue from a loop in the N-terminal domain (42, 48–50); the other exception to this is found in CjXyl31A in which a PA14 domain insert in the N-terminal domain extends the active site (26, 42, 48–50).

The substrate-interacting residues in the –1 and +1 subsites of the CjAgd31B structure are essentially homologous to those in the human maltase-glucoamylase and the *Ro*- α G1 α -glucosidase from *Ruminococcus obeum* with the exception of Phe-271, Leu-300, Phe-377, and Leu-413, which instead are Trp, Ile, Trp, and Met, respectively, in the maltase-glucoamylase and *Ro*- α G1 structures (supplemental Fig. S5 and Refs. 48 and 49). These side chains all make van der Waals contacts to acarbose as well as the 5-fluoro- β -glucosyl residue (see below). It is possible that Phe-271 of CjAgd31B, corresponding to Trp-169 and Tyr-299 in *Ro*- α G1 and maltase-glucoamylase, respectively, contributes to substrate specificity as wild type *Ro*- α G1 prefers isomaltose to maltose as a substrate, whereas the W169Y mutant inverts this preference (49).

Notable features of the +2 and +3 subsites are the tyrosine clamp of Tyr-179 (part of a long loop extending from the N-terminal domain) and Tyr-376 of Insert 1, which together form van der Waals contacts to the internal glucose moiety (third ring) of acarbose (cf. Fig. 5B and supplemental Fig. S4). A similar hydrophobic clamp has not been found in other GH31 structures apart from the PA14-mediated protein-sugar interaction in CjXyl31A (26).

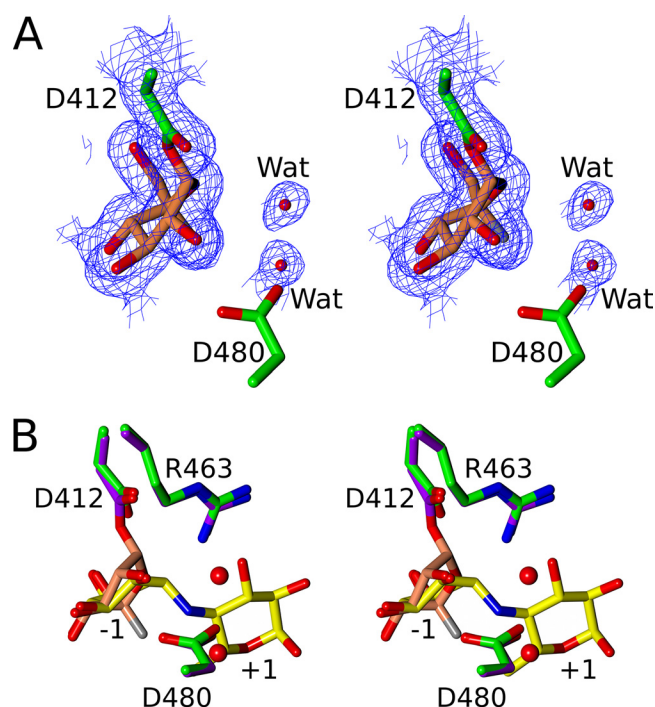


FIGURE 6. **The covalent intermediate of CjAgd31B.** *A*, observed electron density for the trapped covalent 5-fluoro- β -glucosyl-enzyme intermediate of CjAgd31B (map is contoured at 1 σ in divergent stereo). The nucleophile Asp-412, acid-base Asp-480, and a pair of solvent water (Wat) molecules are shown. *B*, overlay of the –1 and +1 subsites of the covalent intermediate complex (pink; waters are shown as red spheres) with the acarbose (yellow) complex (Fig. 5B). Of particular note is that neither (nearby) water of the intermediate complex is in an appropriate position for nucleophilic attack of the intermediate by hydrolysis, implying that hydrogen bonding is optimized to prevent hydrolysis and facilitate transglycosylation. This figure was drawn with CCP4MG (66).

A Bacterial α -Transglucosylase from GH31

Structure of the Trapped Covalent 5-Fluoro- β -glucosyl-enzyme Intermediate—To assess factors that may lead to the strict transglycosylation activity of *CjAgd31B*, a near mimic of the covalent glycosyl-enzyme intermediate was accessed using a classic “Withers” reagent, 5-fluoro- α -D-glucopyranosyl fluoride. The electron density map clearly reveals the trapped 5-fluoro- β -D-glucopyranosyl-enzyme observed in 1S_3 skew boat conformation (conformational aspects of catalysis are reviewed in Ref. 51) formed via covalent linkage to the O δ 2 atom of Asp-412 with 1.34-Å distance (Fig. 6A). The majority of interactions of this sugar are the same as previously observed for the -1 subsite sugar of the acarbose pseudotetrasaccharide (supplemental Fig. S4), but in addition, two water molecules (Fig. 6A) bind to O δ 2 atom of Asp-480, the acid-base residue. These water molecules form a hydrogen bond network with Asp-480, Arg-463, Glu-417, and a third water molecule binding to Gln-542. Although transglycosylation is always kinetically favored over hydrolysis (52), what is unusual about *CjAgd31B* and indeed other transglucosylases is how they overcome the thermodynamically favored hydrolysis reaction in 55 M water as discussed below.

DISCUSSION

Through a combination of enzymological and structural analysis, we have revealed that *CjAgd31B* from the soil saprophyte *C. japonicus* possesses the ability to exclusively transfer single glucosyl units from $\alpha(1\rightarrow4)$ -glucans to the non-reducing terminal 4-OH of glucose and $\alpha(1\rightarrow4)$ - and $\alpha(1\rightarrow6)$ -linked glucosyl residues; weak hydrolysis activity is only observed on the disaccharide maltose. As outlined in the Introduction and discussed below, this type of transglucosylase has not previously been described in GH31 nor any other CAZyme family to our knowledge.

GH31 enzymes utilize a double displacement mechanism involving a covalent glycosyl-enzyme intermediate, which, as was the case here for *CjAgd31B*, can be trapped and directly observed using kinetic probes derived from fluorosugars (46). In the natural reactions catalyzed by GH31 members, the glycosyl-enzyme is most commonly decomposed by water, yielding substrate hydrolysis. However, this intermediate can also be intercepted by saccharide acceptor substrates to generate transglycosylation products with varying efficiencies in a substrate- and enzyme-dependent manner (53, 54). Indeed, several members of GH31 have been shown to possess transglycosylation ability, although yields are typically low due to dominating hydrolytic reactions (21–23, 55, 56). In this context, the strict transglycosylating activity of *CjAgd31B* on malto-oligosaccharide substrates with degree of polymerization ≥ 3 is particularly noteworthy.

CjAgd31B is also distinct from homologues of bacterial *ctsY* and *ctsZ* gene products, which are the only other predominant transglucosylases to have been identified in GH31 thus far. Working in concert, *CtsY* and *CtsZ* generate cycloalternan tetrasaccharides from $\alpha(1\rightarrow4)$ -glucans via a three-step reaction (19, 20). In the first step, *CtsZ* acts as an $\alpha(1\rightarrow4)$ -to- $\alpha(1\rightarrow6)$ transglucosylase to generate isomaltosyl moieties at the end of $\alpha(1\rightarrow4)$ -glucan chains. Our HPAEC-PAD data indicate that in contrast *CjAgd31B* effects $\alpha(1\rightarrow4)$ -to- $\alpha(1\rightarrow4)$ transglycosyla-

tion and moreover cannot address $(1\rightarrow6)$ linkages; isomaltose is not a donor substrate. In the second and third steps of cycloalternan tetrasaccharide synthesis, *CtsY* catalyzes an intermolecular isomaltosyl transfer to yield a α -D-Glcp-(1 \rightarrow 6)- α -D-Glcp-(1 \rightarrow 3)- α -D-Glcp-(1 \rightarrow 6)- α -D-Glcp-(1 \rightarrow 4)- α -glucan structure followed by intramolecular cyclization to yield cyclo[\rightarrow 6]- α -D-Glcp-(1 \rightarrow 3)- α -D-Glcp-(1 \rightarrow 6)- α -D-Glcp-(1 \rightarrow 3)- α -D-Glcp-(1 \rightarrow)]. These two reactions are clearly distinct from that catalyzed by *CjAgd31B*. Unfortunately, the three-dimensional structures of both *CtsZ* and *CtsY* are currently unknown, which precludes comparison with *CjAgd31B* to understand the structural basis for these divergent transglycosylation activities in GH31.

The reactions catalyzed by *CjAgd31B* bear some similarity to, but are again distinct from, those catalyzed by 4- α -glucanotransferases (EC 4.2.1.25) of GH13 and GH77. GH13 encompasses a huge diversity of α -glucan-hydrolyzing and -transglycosylating enzymes (57) of which the *Thermotoga* spp. 4- α -glucanotransferases are perhaps the most relevant to the present study (12, 13, 58). *Thermotoga maritima* and *Thermotoga neapolitana* 4- α -glucanotransferases catalyze disproportionation reactions of malto-oligosaccharides, utilizing maltotetraose as the smallest donor substrate (12, 13). The structure of the *T. maritima* enzyme has revealed that the active site is an open cleft comprising at least 5 subsites (-2 to +3), which provides clear rationalization for the ability of the enzyme to randomly transfer longer α -glucan chains. In contrast, the *T. maritima* maltosyltransferase, also of GH13, strictly transfers maltosyl (Glc₂) units from maltotriose and longer malto-oligosaccharides to the 4-position of α -glucan acceptor substrates (58) due to the presence of a unique protein motif that blocks the active site cleft (59).

GH77 enzymes are structurally related to GH13 enzymes in clan GH-H and thus generally possess open cleft-shaped active sites (60–62) that confer specificity for longer glucan donor substrates (9). For example, the eukaryotic *Solanum tuberosum* (potato) starch disproportionating (“D”) enzyme has such an extended active site (Protein Data Bank code 1x1n), transfers long α -glucan chains, and does not use maltose as a donor substrate (14, 63, 64). The disaccharide is also not a substrate for the *Thermus aquaticus* amyloamylase of GH77 that is distinguished by its propensity to form large cyclic α -glucans from long $\alpha(1\rightarrow4)$ -glucan donors (14, 60). In the context of bacterial malto-oligosaccharide metabolism, the *E. coli* GH77 amyloamylase MalQ appears to favor the transfer of longer α -glucan chains, although there appears to be some debate whether this enzyme can utilize maltose as a donor, thereby transferring a single glucosyl residue to longer congeners.

CjAgd31B thus occupies a unique catalytic place as a 4- α -glucosyltransferase among the broader spectrum of 4- α -glucanotransferases. As a member of GH31, *CjAgd31B* belongs to clan GH-D, which is also composed of GH27 and GH36. Clan GH-D and GH-H members are built on a common triose isomerase (β/α)₈ barrel scaffold and may share a distant evolutionary relationship (27). However, in contrast to the clefted clan GH-H members, clan GH-D members are typified by shallow, pocket-shaped active sites comprising only one negative subsite accommodating a monosaccharide residue, the glyco-

sidic bond of which undergoes catalysis. *CjAgd31B* likewise presents an active site pocket as revealed by acarbose and 5-fluoro-glycosyl-enzyme complex structures, allowing speculation regarding its strict glucosyl transfer capacity.

The inherent challenge for transglycosylases is overcoming the thermodynamic preference for water as a nucleophile *versus* saccharide acceptor substrates. It has long been established by Withers *et al.* (52) through analysis of the reactivation of trapped intermediates that transglycosylation is kinetically favored over hydrolysis. Crystallographic analysis of the trapped covalent glycosyl-enzyme intermediate of *CjAgd31B* suggests that the hydrogen-bonding scheme does not place a water molecule with appropriate geometry or interaction with the catalytic acid-base residue Asp-480 to facilitate hydrolysis of the intermediate. Instead, two water molecules (one of them disordered) appear to lie on either side of the position expected of a catalytically competent nucleophile. However, the interglycosidic nitrogen of acarbose does interact with the acid-base (and with its O3 and C5 groups, binding in positions corresponding to the observed waters of the trapped intermediate).

This solvent hydrogen-bonding arrangement suggests that the *CjAgd31B* active site has evolved to avoid deprotonation and activation of water, whereas optimization of hydrogen bonds to O6 and O3 of acceptor glucosides in the +1 subsite allows a favorable placement of the O4 atom for deprotonation and concerted electrophilic migration of C1 of the β -glucosyl-enzyme. In this process, Glu-417 and Arg-463 play particularly important roles in binding the O3 of the +1 sugar and legislating against a water molecule positioned to enable hydrolysis. A caveat is that the trapped intermediate observed is that of a 5-fluoroglycoside, so it is possible that the observed solvent network is perturbed by the unnatural 5-fluoro substituent. However, the solvent network of the free enzyme structure is similar to that of the trapped intermediate complex especially in context of the O6- and O3-mimicking water molecules, where there is no suitably poised "nucleophilic" water molecule bound to Asp-480.

It is currently unclear what role(s) *CjAgd31B* might play in the biology of *C. japonicus*. Interestingly, the gene encoding *CjAgd31B* is located among a cluster of genes predicted to encode α -glucan-active enzymes (α -amylase, cyclomaltodextrin glucanotransferase, 6-phospho- β -glucosidase, and glucokinase) and transporter proteins (TonB-dependent receptors and ATP-binding cassette transporters; supplemental Fig. S2). Whether these genes are co-regulated or comprise an operon is currently not known. This genomic association together with the observation that *CjAgd31B* is encoded with a native secretion signal peptide hints toward a role in glycogen or starch metabolism in the periplasm. Indeed, *CjAgd31B* could possibly have a function similar to the GH77 amylomaltase MalQ from *E. coli*, which creates longer α -glucan chains from shorter malto-oligosaccharides as substrates for maltodextrin phosphorylases; these phosphorylases require maltopentaose as a minimal substrate to generate glucose 1-phosphate for further metabolism (5, 65, 66). *C. japonicus*, however, does possess a predicted GH77 homologue (*CjMal77Q*, CJA_1882; Ref. 25), which is located elsewhere in the genome and in proximity to other predicted glycogen/starch-active enzymes. This would

suggest that *CjAgd31B* and *CjMal77Q* most likely have independent or perhaps complementary functions.

Another possible clue to the physiological function of *CjAgd31B* can be gleaned from analysis of potential GH31 orthologs. The biochemically characterized GH31 member closest to *CjAgd31B* is the α -glucosidase YihQ from *E. coli* K12 MG1655 (38). Biological data on *E. coli* YihQ are currently lacking; however, a reverse genetics analysis of a YihQ orthologue in *Salmonella enterica* serovar Enteritidis indicates that $\Delta yihQ$ mutants are deficient in capsular polysaccharide formation (67). Notably, the LPS of this organism consists of a repeating core glycan comprising tyvelose, L-rhamnose, galactose, and mannose that is appended with extended $\alpha(1\rightarrow4)$ -glucan chains. It is therefore tempting to speculate that YihQ and by extension *CjAgd31B* may act as a transglucosylase to extend or restructure these chains. In this context, it is interesting to note that *E. coli* YihQ has previously been designated as an α -glucosidase based on a weak activity on α -glucosyl fluoride but no other α -glucosides (38). A reassessment of YihQ activity both *in vitro* and *in vivo* in light of the transglycosylation capacity of GH31 enzymes demonstrated in the present study may well be warranted. In conclusion, the detailed enzyme structure-function analysis of Agd31B from the model soil bacterium *C. japonicus* presented here that has defined a previously unknown α -transglucosylase activity in GH31 will inform future functional genomics studies in bacteria and other microorganisms.

Acknowledgments—We are grateful to Prof. Harry J. Gilbert (Newcastle University) for insightful discussions on *C. japonicus* and early access to genomic sequence data. We thank Prof. Stephen G. Withers (University of British Columbia) for gifts of α -glucosyl fluoride and 5-fluoro- α -D-glucopyranosyl fluoride and Dr. Gustav Sundqvist (KTH Glycoscience) for assistance with protein mass spectrometry. The staffs of the Diamond Light Source and European Synchrotron Radiation Facility are thanked for provision of data collection facilities.

REFERENCES

1. Elbein, A. D. (2009) in *Microbial Glycobiology: Structures, Relevance and Applications* (Moran, A., Holst, O., Brennan, P. J., and von Itzstein, M., eds) pp. 185–201, Academic Press, London
2. Wilson, W. A., Roach, P. J., Montero, M., Baroja-Fernández, E., Muñoz, F. J., Eydallin, G., Viale, A. M., and Pozueta-Romero, J. (2010) Regulation of glycogen metabolism in yeast and bacteria. *FEMS Microbiol. Rev.* **34**, 952–985
3. Roach, P. J., Depaoli-Roach, A. A., Hurley, T. D., and Tagliabracci, V. S. (2012) Glycogen and its metabolism: some new developments and old themes. *Biochem. J.* **441**, 763–787
4. Ball, S., Colleoni, C., Cenci, U., Raj, J. N., and Tirtiaux, C. (2011) The evolution of glycogen and starch metabolism in eukaryotes gives molecular clues to understand the establishment of plastid endosymbiosis. *J. Exp. Bot.* **62**, 1775–1801
5. Boos, W., and Shuman, H. (1998) Maltose/maltodextrin system of *Escherichia coli*: transport, metabolism, and regulation. *Microbiol. Mol. Biol. Rev.* **62**, 204–229
6. 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 glycogenomics. *Nucleic Acids Res.* **37**, D233–D238
7. Palmer, T. N., Ryman, B. E., and Whelan, W. J. (1976) Action pattern of

A Bacterial α -Transglucosylase from GH31

- amylomaltase from *Escherichia coli*. *Eur. J. Biochem.* **69**, 105–115
- Dippel, R., and Boos, W. (2005) The maltodextrin system of *Escherichia coli*: metabolism and transport. *J. Bacteriol.* **187**, 8322–8331
 - Takaha, T., and Smith, S. M. (1999) The functions of 4- α -glucanotransferases and their use for the production of cyclic glucans. *Biotechnol. Genet. Eng. Rev.* **16**, 257–280
 - Roujeinikova, A., Raasch, C., Burke, J., Baker, P. J., Liebl, W., and Rice, D. W. (2001) The crystal structure of *Thermotoga maritima* maltosyltransferase and its implications for the molecular basis of the novel transfer specificity. *J. Mol. Biol.* **312**, 119–131
 - Roujeinikova, A., Raasch, C., Sedelnikova, S., Liebl, W., and Rice, D. W. (2002) Crystal structure of *Thermotoga maritima* 4- α -glucanotransferase and its acarbose complex: implications for substrate specificity and catalysis. *J. Mol. Biol.* **321**, 149–162
 - Berezina, O. V., Zverlov, V. V., Lunina, N. A., Chekanovskaya, L. A., Dubinina, E. N., Liebl, W., and Velikodvorskaya, G. A. (1999) Gene and properties of thermostable 4- α -glucanotransferase of *Thermotoga neapolitana*. *Mol. Biol.* **33**, 801–806
 - Liebl, W., Feil, R., Gabelsberger, J., Kellermann, J., and Schleifer, K. H. (1992) Purification and characterization of a novel thermostable 4- α -glucanotransferase of *Thermotoga maritima* cloned in *Escherichia coli*. *Eur. J. Biochem.* **207**, 81–88
 - Terada, Y., Fujii, K., Takaha, T., and Okada, S. (1999) Thermus aquaticus ATCC 33923 amylomaltase gene cloning and expression and enzyme characterization: production of cycloamylose. *Appl. Environ. Microbiol.* **65**, 910–915
 - Leemhuis, H., Kelly, R. M., and Dijkhuizen, L. (2010) Engineering of cyclodextrin glucanotransferases and the impact for biotechnological applications. *Appl. Microbiol. Biotechnol.* **85**, 823–835
 - Kim, Y. M., Kiso, Y., Muraki, T., Kang, M. S., Nakai, H., Saburi, W., Lang, W., Kang, H. K., Okuyama, M., Mori, H., Suzuki, R., Funane, K., Suzuki, N., Momma, M., Fujimoto, Z., Oguma, T., Kobayashi, M., Kim, D., and Kimura, A. (2012) Novel dextranase catalyzing cycloisomaltooligosaccharide formation and identification of catalytic amino acids and their functions using chemical rescue approach. *J. Biol. Chem.* **287**, 19927–19935
 - Kim, Y. M., Yamamoto, E., Kang, M. S., Nakai, H., Saburi, W., Okuyama, M., Mori, H., Funane, K., Momma, M., Fujimoto, Z., Kobayashi, M., Kim, D., and Kimura, A. (2012) *Bacteroides thetaiotaomicron* VPI-5482 glycoside hydrolase family 66 homolog catalyzes dextranolytic and cyclization reactions. *FEBS J.* **279**, 3185–3191
 - Suzuki, R., Terasawa, K., Kimura, K., Fujimoto, Z., Momma, M., Kobayashi, M., Kimura, A., and Funane, K. (2012) Biochemical characterization of a novel cycloisomaltooligosaccharide glucanotransferase from *Paenibacillus* sp. 598K. *Biochim. Biophys. Acta* **1824**, 919–924
 - Mukai, K., Maruta, K., Satouchi, K., Kubota, M., Fukuda, S., Kurimoto, M., and Tsujisaka, Y. (2004) Cyclic tetrasaccharide-synthesizing enzymes from *Arthrobacter globiformis* A19. *Biosci. Biotechnol. Biochem.* **68**, 2529–2540
 - Kim, Y. K., Kitaoka, M., Hayashi, K., Kim, C. H., and Côté, G. L. (2003) A synergistic reaction mechanism of a cycloalternan-forming enzyme and a D-glucosyltransferase for the production of cycloalternan in *Bacillus* sp. NRRL B-21195. *Carbohydr. Res.* **338**, 2213–2220
 - Wang, Y. H., Jiang, Y., Duan, Z. Y., Shao, W. L., and Li, H. Z. (2009) Expression and characterization of an α -glucosidase from *Thermoanaerobacter ethanolicus* JW200 with potential for industrial application. *Biologia* **64**, 1053–1057
 - Zhou, C., Xue, Y., Zhang, Y., Zeng, Y., and Ma, Y. (2009) Recombinant expression and characterization of *Thermoanaerobacter tengcongensis* thermostable α -glucosidase with regioselectivity for high-yield isomaltooligosaccharides synthesis. *J. Microbiol. Biotechnol.* **19**, 1547–1556
 - Seo, S. H., Choi, K. H., Hwang, S., Kim, J., Park, C. S., Rho, J. R., and Cha, J. (2011) Characterization of the catalytic and kinetics properties of a thermostable *Thermoplasma acidophilum* α -glucosidase and its transglucosylation reaction with arbutin. *J. Mol. Catal. B Enzym.* **72**, 305–312
 - Hazlewood, G. P., and Gilbert, H. J. (1998) Structure and function analysis of *Pseudomonas* plant cell wall hydrolases. *Prog. Nucleic Acid Res. Mol. Biol.* **61**, 211–241
 - DeBoy, R. T., Mongodin, E. F., Fouts, D. E., Tailford, L. E., Khouri, H., Emerson, J. B., Mohamoud, Y., Watkins, K., Henrissat, B., Gilbert, H. J., and Nelson, K. E. (2008) Insights into plant cell wall degradation from the genome sequence of the soil bacterium *Cellvibrio japonicus*. *J. Bacteriol.* **190**, 5455–5463
 - Larsbrink, J., Izumi, A., Ibatullin, F. M., Nakhai, A., Gilbert, H. J., Davies, G. J., and Brumer, H. (2011) Structural and enzymatic characterization of a glycoside hydrolase family 31 α -xylosidase from *Cellvibrio japonicus* involved in xyloglucan saccharification. *Biochem. J.* **436**, 567–580
 - Janecek, S., Svensson, B., and MacGregor, E. A. (2007) A remote but significant sequence homology between glycoside hydrolase clan GH-H and family GH31. *FEBS Lett.* **581**, 1261–1268
 - Sundqvist, G., Stenvall, M., Berglund, H., Ottosson, J., and Brumer, H. (2007) A general, robust method for the quality control of intact proteins using LC-ESI-MS. *J. Chromatogr. B Analyt. Technol. Biomed. Life Sci.* **852**, 188–194
 - Gasteiger, E., Hoogland, C., Gattiker, A., Duvaud, S., Wilkins, M. R., Appel, R. D., and Bairoch, A. (2005) in *The Proteomics Protocols Handbook* (Walker, J. M., ed.) pp. 571–607, Humana Press, Totowa, NJ
 - Leslie, A. G. (1999) Integration of macromolecular diffraction data. *Acta Crystallogr. D Biol. Crystallogr.* **55**, 1696–1702
 - Winn, M. D., Ballard, C. C., Cowtan, K. D., Dodson, E. J., Emsley, P., Evans, P. R., Keegan, R. M., Krissinel, E. B., Leslie, A. G., McCoy, A., McNicholas, S. J., Murshudov, G. N., Pannu, N. S., Potterton, E. A., Powell, H. R., Read, R. J., Vagin, A., and Wilson, K. S. (2011) Overview of the CCP4 suite and current developments. *Acta Crystallogr. D Biol. Crystallogr.* **67**, 235–242
 - Vonrhein, C., Blanc, E., Roversi, P., and Bricogne, G. (2007) Automated structure solution with autoSHARP. *Methods Mol. Biol.* **364**, 215–230
 - Cowtan, K. D. (1994) 'dm': an automated procedure for phase improvement by density modification. *Joint CCP4 ESF-EACBM Newslett. Protein Crystallogr.* **31**, 34–38
 - Perrakis, A., Morris, R., and Lamzin, V. S. (1999) Automated protein model building combined with iterative structure refinement. *Nat. Struct. Biol.* **6**, 458–463
 - Adams, P. D., Afonine, P. V., Bunkóczi, 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., Richardson, J. S., Terwilliger, T. C., and Zwart, P. H. (2010) PHENIX: a comprehensive Python-based system for macromolecular structure solution. *Acta Crystallogr. D Biol. Crystallogr.* **66**, 213–221
 - Emsley, P., and Cowtan, K. (2004) Coot: model-building tools for molecular graphics. *Acta Crystallogr. D Biol. Crystallogr.* **60**, 2126–2132
 - Vagin, A., and Teplyakov, A. (1997) MOLREP: an automated program for molecular replacement. *J. Appl. Crystallogr.* **30**, 1022–1025
 - Okuyama, M., Mori, H., Chiba, S., and Kimura, A. (2004) Overexpression and characterization of two unknown proteins, YicI and YihQ, originated from *Escherichia coli*. *Protein Expr. Purif.* **37**, 170–179
 - Davies, G. J., Wilson, K. S., and Henrissat, B. (1997) Nomenclature for sugar-binding subsites in glycosyl hydrolases. *Biochem. J.* **321**, 557–559
 - Burlingham, B. T., and Widlanski, T. S. (2003) An intuitive look at the relationship of K_i and IC_{50} : a more general use for the Dixon plot. *J. Chem. Educ.* **80**, 214–218
 - Rossi, E. J., Sim, L., Kuntz, D. A., Hahn, D., Johnston, B. D., Ghavami, A., Szczepina, M. G., Kumar, N. S., Sterchi, E. E., Nichols, B. L., Pinto, B. M., and Rose, D. R. (2006) Inhibition of recombinant human maltase glucoamylase by salacinol and derivatives. *FEBS J.* **273**, 2673–2683
 - Sim, L., Willemsma, C., Mohan, S., Naim, H. Y., Pinto, B. M., and Rose, D. R. (2010) Structural basis for substrate selectivity in human maltase-glucoamylase and sucrase-isomaltase N-terminal domains. *J. Biol. Chem.* **285**, 17763–17770
 - Ren, L., Cao, X., Geng, P., Bai, F., and Bai, G. (2011) Study of the inhibition of two human maltase-glucoamylases catalytic domains by different α -glucosidase inhibitors. *Carbohydr. Res.* **346**, 2688–2692
 - Lee, S. S., Yu, S., and Withers, S. G. (2003) Detailed dissection of a new mechanism for glycoside cleavage: α -1,4-glucan lyase. *Biochemistry* **42**, 13081–13090
 - Lee, S. S., He, S., and Withers, S. G. (2001) Identification of the catalytic nucleophile of the family 31 α -glucosidase from *Aspergillus niger* via trapping of a 5-fluoroglycosyl-enzyme intermediate. *Biochem. J.* **359**, 381–386

46. Lovering, A. L., Lee, S. S., Kim, Y. W., Withers, S. G., and Strynadka, N. C. (2005) Mechanistic and structural analysis of a family 31 α -glucosidase and its glycosyl-enzyme intermediate. *J. Biol. Chem.* **280**, 2105–2115
47. Hendlich, M., Rippmann, F., and Barnickel, G. (1997) LIGSITE: automatic and efficient detection of potential small molecule-binding sites in proteins. *J. Mol. Graph. Model.* **15**, 359–363, 389
48. Sim, L., Quezada-Calvillo, R., Sterchi, E. E., Nichols, B. L., and Rose, D. R. (2008) Human intestinal maltase-glucoamylase: crystal structure of the N-terminal catalytic subunit and basis of inhibition and substrate specificity. *J. Mol. Biol.* **375**, 782–792
49. Tan, K., Tesar, C., Wilton, R., Keigher, L., Babnigg, G., and Joachimiak, A. (2010) Novel α -glucosidase from human gut microbiome: substrate specificities and their switch. *FASEB J.* **24**, 3939–3949
50. Ernst, H. A., Lo Leggio, L., Willemoës, M., Leonard, G., Blum, P., and Larsen, S. (2006) Structure of the *Sulfolobus solfataricus* α -glucosidase: implications for domain conservation and substrate recognition in GH31. *J. Mol. Biol.* **358**, 1106–1124
51. Davies, G. J., Planas, A., and Rovira, C. (2012) Conformational analyses of the reaction coordinate of glycosidases. *Acc. Chem. Res.* **45**, 308–316
52. Withers, S. G., Warren, R. A. J., Street, I. P., Rupitz, K., Kempton, J. B., and Aebersold, R. (1990) Unequivocal demonstration of the involvement of a glutamate residue as a nucleophile in the mechanism of a retaining glycosidase. *J. Am. Chem. Soc.* **112**, 5887–5889
53. Sinnott, M. L. (1990) Catalytic mechanisms of enzymatic glycosyl transfer. *Chem. Rev.* **90**, 1171–1202
54. Stick, R., and Williams, S. (2009) *Carbohydrates: the Essential Molecules of Life*, pp. 262–263 and 273–276, Elsevier, Oxford
55. Kato, N., Suyama, S., Shirokane, M., Kato, M., Kobayashi, T., and Tsukagoshi, N. (2002) Novel α -glucosidase from *Aspergillus nidulans* with strong transglycosylation activity. *Appl. Environ. Microbiol.* **68**, 1250–1256
56. Yamamoto, T., Unno, T., Watanabe, Y., Yamamoto, M., Okuyama, M., Mori, H., Chiba, S., and Kimura, A. (2004) Purification and characterization of *Acremonium implicatum* α -glucosidase having regioselectivity for α -1,3-glucosidic linkage. *Biochim. Biophys. Acta* **1700**, 189–198
57. Stam, M. R., Danchin, E. G., Rancurel, C., Coutinho, P. M., and Henrissat, B. (2006) Dividing the large glycoside hydrolase family 13 into subfamilies: towards improved functional annotations of α -amylase-related proteins. *Protein. Eng. Des. Sel.* **19**, 555–562
58. Humphry, D. R., Black, G. W., and Cummings, S. P. (2003) Reclassification of '*Pseudomonas fluorescens* subsp. *cellulosa*' NCIMB 10462 (Ueda *et al.* (1952) as *Cellvibrio japonicus* sp. nov., and revival of *Cellvibrio vulgaris* sp. nov., nom. rev., and *Cellvibrio fulvus* sp. nov., nom. rev. *Int. J. Syst. Evol. Microbiol.* **53**, 393–400
59. Meissner, H., and Liebl, W. (1998) *Thermotoga maritima* maltosyltransferase, a novel type of maltodextrin glycosyltransferase acting on starch and malto-oligosaccharides. *Eur. J. Biochem.* **258**, 1050–1058
60. Przyas, I., Tomoo, K., Terada, Y., Takaha, T., Fujii, K., Saenger, W., and Sträter, N. (2000) Crystal structure of amylomaltase from *Thermus aquaticus*, a glycosyltransferase catalysing the production of large cyclic glucans. *J. Mol. Biol.* **296**, 873–886
61. Jung, J. H., Jung, T. Y., Seo, D. H., Yoon, S. M., Choi, H. C., Park, B. C., Park, C. S., and Woo, E. J. (2011) Structural and functional analysis of substrate recognition by the 250s loop in amylomaltase from *Thermus brockianus*. *Proteins* **79**, 633–644
62. Barends, T. R., Bultema, J. B., Kaper, T., van der Maarel, M. J., Dijkhuizen, L., and Dijkstra, B. W. (2007) Three-way stabilization of the covalent intermediate in amylomaltase, an α -amylase-like transglycosylase. *J. Biol. Chem.* **282**, 17242–17249
63. Jones, G., and Whelan, W. J. (1969) Action pattern of D-enzyme, a trans-maltodextrinylase from potato. *Carbohydr. Res.* **9**, 483
64. Takaha, T., Yanase, M., Okada, S., and Smith, S. M. (1993) Disproportionating enzyme (4- α -glucanotransferase; EC 2.4.1.25) of potato. Purification, molecular-cloning, and potential role in starch metabolism. *J. Biol. Chem.* **268**, 1391–1396
65. Schwartz, M., and Hofnung, M. (1967) Maltodextrin phosphorylase of *Escherichia coli*. *Eur. J. Biochem.* **2**, 132–145
66. Becker, S., Palm, D., and Schinzel, R. (1994) Dissecting differential binding in the forward and reverse reaction of *Escherichia coli* maltodextrin phosphorylase using 2-deoxyglucosyl substrates. *J. Biol. Chem.* **269**, 2485–2490
67. Gibson, D. L., White, A. P., Snyder, S. D., Martin, S., Heiss, C., Azadi, P., Surette, M., and Kay, W. W. (2006) *Salmonella* produces an O-antigen capsule regulated by AgfD and important for environmental persistence. *J. Bacteriol.* **188**, 7722–7730

SUPPLEMENTAL INFORMATION

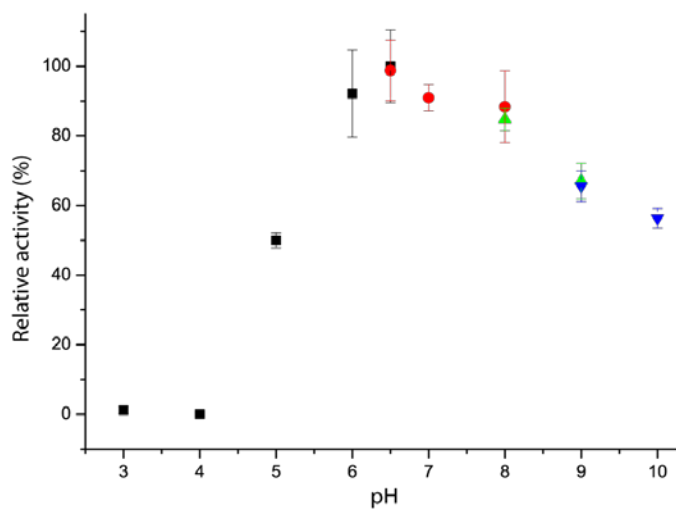


Figure S1: pH profile of *CjAgd31B* from the reaction on maltose (0.5 mM, 10 min). The rate was calculated for the transglycosylation reaction (formation of maltotriose). The following buffers were used: citrate (black squares), sodium phosphate (red circles), glycylglycine (green triangles) and glycine (blue triangles).

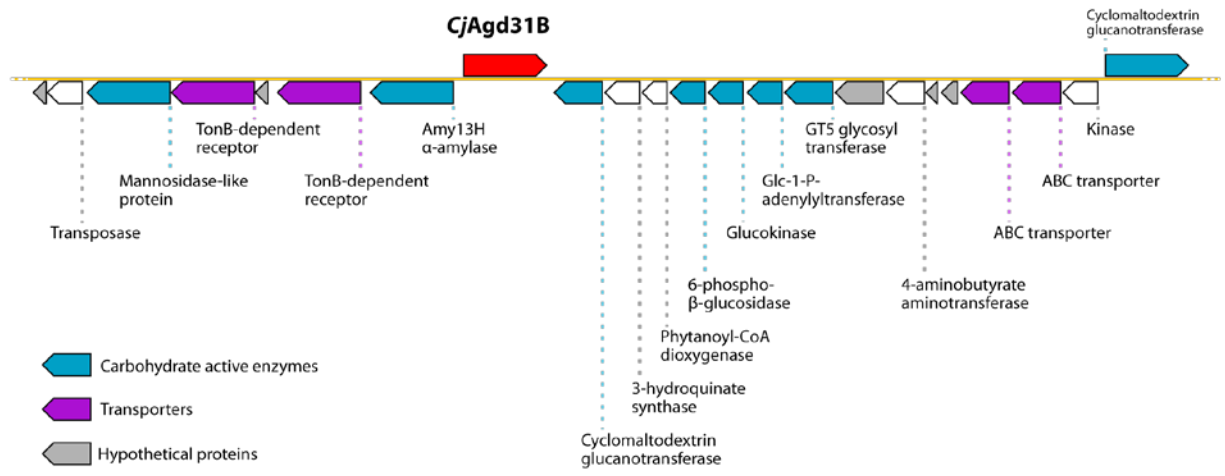


Figure S2: The genomic map surrounding *CjAgd31B* (red). Carbohydrate-active enzymes are colored blue, transporters purple, other activities white, and hypothetical proteins grey.

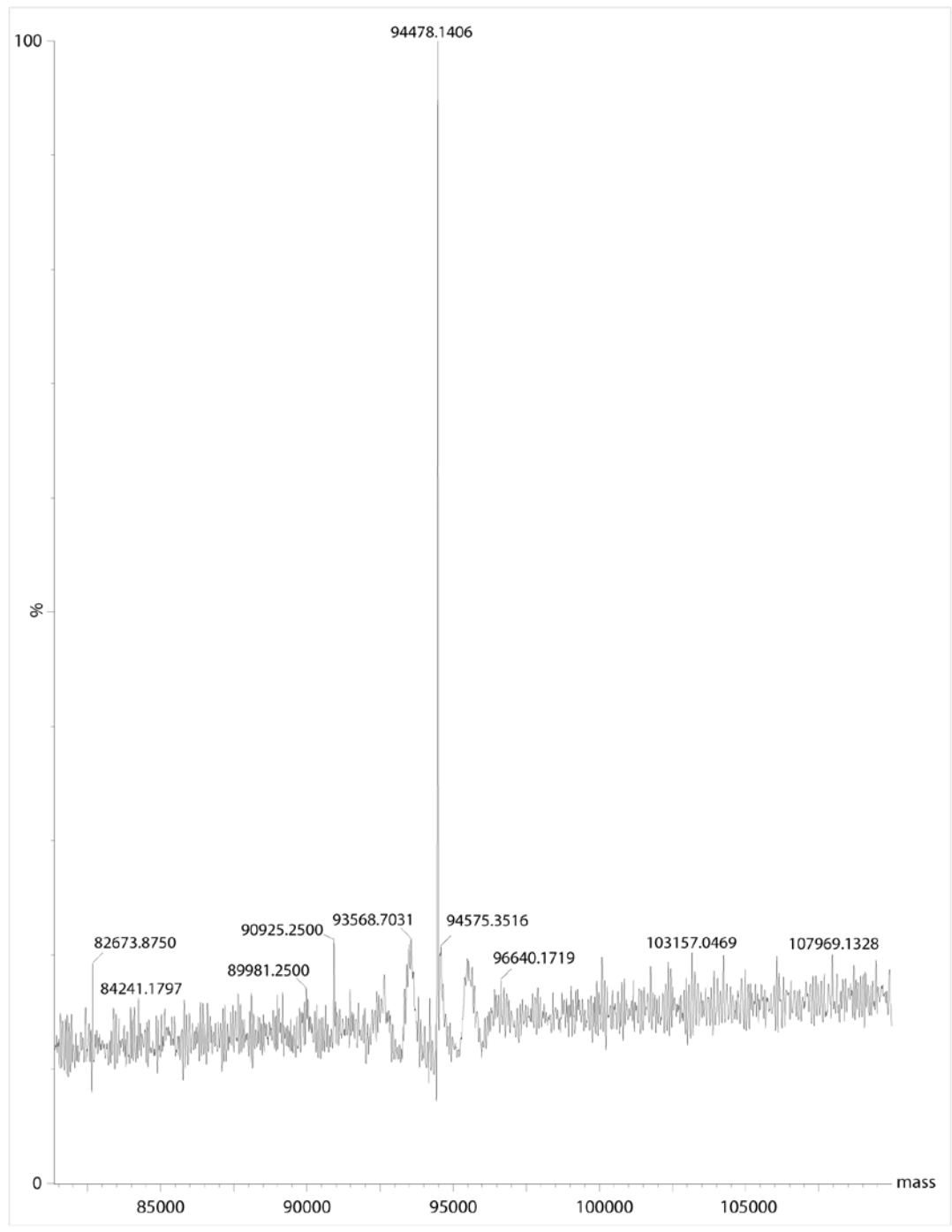


Figure S3: LC-ESI-MS of *CjAgd31B*.

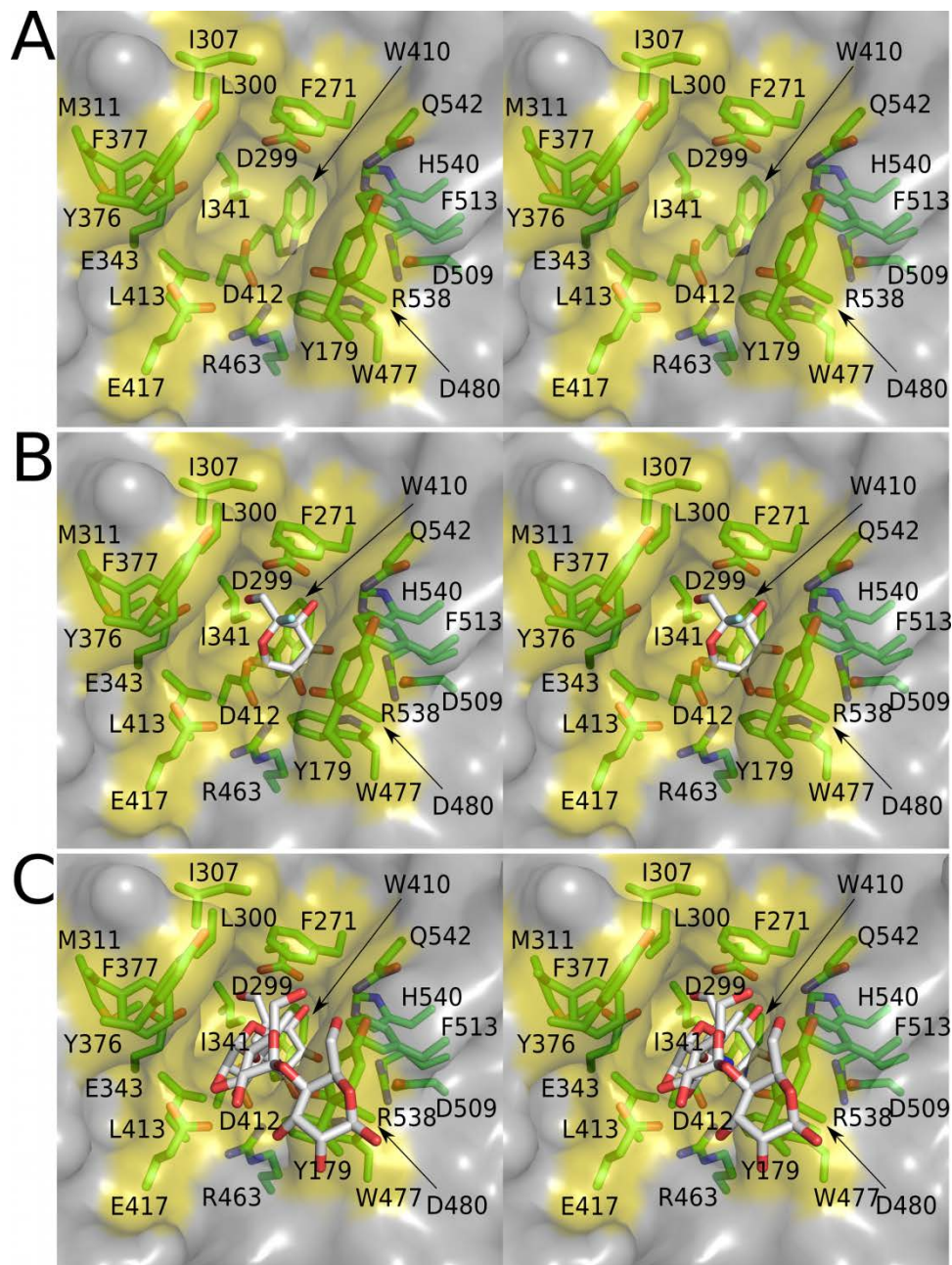


Figure S4: The active site pocket of *CjAgd31B* in (A) the native structure, (B) the 5FGlc Complex and (C) the acarbose complex. The surface of the pocket is coloured yellow with the residues forming the site shown as sticks with green carbons and ligands coloured with white carbon atoms.

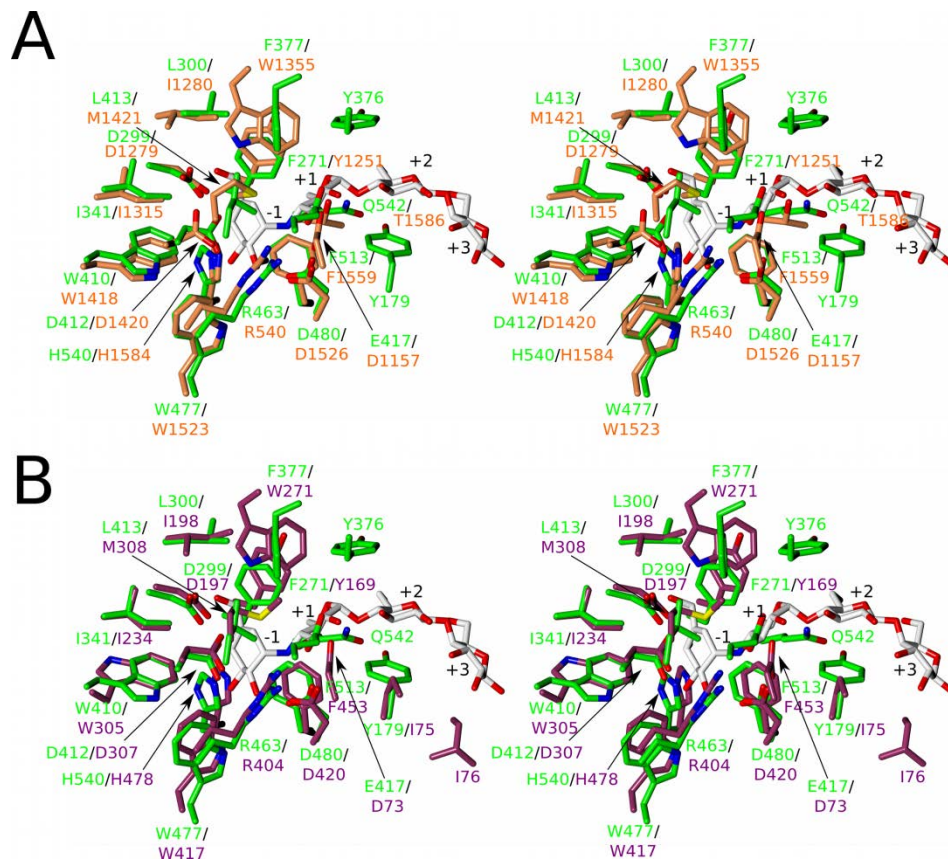


Figure S5: Divergent stereo figures for the superposition of the active site residues in *CjAgd31B* (green carbon atoms) with those in (A) Human maltase-glucoamylase (orange carbon atoms) and (B) *Ro-αG1 α-glucosidase* (purple carbon atoms). Only the acarbose (white carbon atoms) from the *CjAgd31B* structure is shown for clarity.

Structural Enzymology of *Cellvibrio japonicus* Agd31B Protein Reveals α -Transglucosylase Activity in Glycoside Hydrolase Family 31
Johan Larsbrink, Atsushi Izumi, Glyn R. Hemsworth, Gideon J. Davies and Harry Brumer

J. Biol. Chem. 2012, 287:43288-43299.

doi: 10.1074/jbc.M112.416511 originally published online November 6, 2012

Access the most updated version of this article at doi: [10.1074/jbc.M112.416511](https://doi.org/10.1074/jbc.M112.416511)

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

Supplemental material:

<http://www.jbc.org/content/suppl/2012/11/06/M112.416511.DC1.html>

This article cites 64 references, 20 of which can be accessed free at <http://www.jbc.org/content/287/52/43288.full.html#ref-list-1>