Structural Analysis of β-Fructofuranosidase from *Xanthophyllomyces dendrorhous* Reveals Unique Features and the Crucial Role of N-Glycosylation in Oligomerization and Activity*

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*Xanthophyllomyces dendrorhous* β-fructofuranosidase (XdINV) is a highly glycosylated dimeric enzyme that hydrolyzes sucrose and releases fructose from various fructooligosaccharides (FOS) and fructans. It also catalyzes the synthesis of FOS, prebiotics that stimulate the growth of beneficial bacteria in human gut. In contrast to most fructosylating enzymes, XdINV produces neo-FOS, which makes it an interesting biotechnology target. We present here its three-dimensional structure, which shows the expected bimodular arrangement and also a long extension of its C-terminus that together with an N-linked glycan mediate the formation of an unusual dimer. The two active sites of the dimer are connected by a long crevice, which might indicate its potential ability to accommodate branched fructans. This arrangement could be representative of a group of GH32 yeast enzymes having the traits observed in XdINV. The inactive D80A mutant was used to obtain complexes with relevant substrates and products, with its crystals structures showing at least four binding subsites at each active site. Moreover, two different positions are observed from subsite +2 depending on the substrate, and thus, a flexible loop (Glu-334–His-343) is essential in binding sucrose and β(2→1)-linked oligosaccharides. Conversely, β(2→6) and neo-type substrates are accommodated mainly by stacking to Trp-105, explaining the production of neokestose and the efficient fructosylating activity of XdINV on α-glycosides. The role of relevant residues has been investigated by mutagenesis and kinetics measurements, and a model for the transfructosylating reaction has been proposed. The plasticity of its active site makes XdINV a valuable and flexible biocatalyst to produce novel bioconjugates.

Invertases, or β-fructofuranosidases (EC 3.2.1.26), are biotechnologically important enzymes secreted by many fungi showing wide applications in the food and pharmaceutical industries. β-Fructofuranosidase (XdINV) from the basidiomycota yeast *Xanthophyllomyces dendrorhous* (also *Phaffia rhodozyma*) is a highly glycosylated protein with a content of 60–70% N-linked carbohydrates and a homodimeric active form of 320–380 kDa (1, 2). As other β-fructofuranosidases, XdINV catalyzes the release of β-fructose from the nonreducing termini of various β-d-fructofuranoside substrates such as sucrose, 1-kestose, or nystose (1), and it may also catalyze the synthesis of short-chain fructooligosaccharides (FOS), in which one or two fructosyl moieties are linked to the sucrose skeleton. Most fructosylating enzymes, such as levansucrases (3), inulinsucrases (4), and the majority of fructofuranosidases (5), are able to form either β(2→1) (1F-FOS) or β(2→6) (6F-FOS) linkages between fructose units. However, the distinctive property of XdINV is its ability to transfer fructosyl moieties to the glucose unit of sucrose, thus forming neokestose and neonytose as the main transglycosylation products, both neo-FOS with a levan-type structure (6G-FOS series). FOS are considered as functional food, acting as prebiotics that selectively stimulate growth/activity of lactobacilli and bifidobacteria in the digestive tract and prevent growth of pathogenic microorganisms. They exert a beneficial effect on human health, contributing to the prevention of cardiovascular diseases, colon cancer, or osteoporosis (6). Neo-FOS possess enhanced properties and chemical stability compared with 1F-FOS series (e.g. 1-kestose or nystose) typically used as prebiotics (7–9). In addition, XdINV is also able to fructosylate a variety of carbohydrates containing glucose, in particular disaccharides (maltose, 1,3-bis[tris(hydroxymethyl)methylamino]propane; FOS, fructooligosaccharides; GlCNac, N-acetylgalactosamine; AaeL, *A. awamori* exo-ulinulase; AjFT, *A. japonicus* fructosyltransferase; GH, glycosyl hydrolase; ScINV, *S. cerevisiae* invertase; SoFase, *S. occidentalis* β-fructofuranosidase; Endo H, endoglycosidase H.

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§3 The abbreviations used are: XdINV, *X. dendrorhous* invertase; Bistris propane, 1,3-bis[tris(hydroxymethyl)methylamino]propane; FOS, fructooligosaccharides; GlCNac, N-acetylgalactosamine; AaeL, *A. awamori* exo-ulinulase; AjFT, *A. japonicus* fructosyltransferase; GH, glycosyl hydrolase; ScINV, *S. cerevisiae* invertase; SoFase, *S. occidentalis* β-fructofuranosidase; Endo H, endoglycosidase H.
Structure and Function of Dimeric XdINV

isomaltulose, isomalto, and trehalose) and higher oligosaccharides (maltotriose, raffinose, and maltotetraose) but not monosaccharides (glucose, fructose, and galactose) (10). Thus, novel heterofructooligosaccharides with potential applications in the functional food and pharmaceutical industries can be obtained with this enzyme, which makes it an interesting biotechnological target. Further structural analysis of XdINV is necessary to fully understand its particular specificity and to improve its biosynthetic potential.

On the basis of its amino acid sequences, XdINV was classified into the glycoside hydrolase (GH) family GH32 (1), a family including other invertases, β-fructofuranosidases, inulinases, and fructosyltransferases (11), which along with proteins in family GH68 constitute clan GH-J. The enzymes within this clan share a five-blade β-propeller N-terminal domain, in which β-sheets are arranged around a central pocket that accommodates the active site (catalytic domain). Three key acidic residues located in the active site and surrounded by conserved sequences in the GH32 family, NDPNG (D acts as a nucleophile), RDP (D acts as a stabilizer of the transient state), and EC (E acts as an acid base catalyst), are implicated in substrate binding and hydrolysis (12). These enzymes can have hydrolytic or synthetic activity, depending on whether water or an oligosaccharide (the acceptor substrate) accepts the fructose released by fructan hydrolysis (the donor substrate). A distinctive trait of GH32 enzymes is the presence of an additional β-sandwich structure fused at the C terminus of the catalytic domain. In the last decade, GH32 enzymes from bacteria (13, 14), fungi (15–17), and plants (18–20) have been the subject of many structural studies that showed that all are monomeric enzymes. Surprisingly, studies on Schwanniomyces occidentalis fructofuranosidase (SoFfase) (21) and Saccharomyces cerevisiae invertase (ScINV) (22) showed that these enzymes from yeast are unique in that they form dimers mediated by their β-sandwich domain, which in the case of ScINV associate into higher oligomers. The structural studies have delineated the main structural determinants of activity, the loops and turns connecting the different elements of the β-propeller domain, in the surroundings of the active site, defining the specificity unique to each enzyme (21, 23). However, and remarkably, oligomerization of the yeast enzymes regulates their functionality. Thus, the analysis of the structure of a complex of SoFfase with the polymeric inulin showed the direct involvement of the β-sandwich domain of an adjacent subunit in substrate binding (24).

In this study, we report the three-dimensional structure of the yeast XdINV that discloses novel features and an unusual dimer. Complexes with relevant substrates and products have been obtained, which allowed mapping the catalytic pocket. The role of identified leading residues has been investigated by mutagenesis. Our results give the experimental evidence of the direct involvement of glycosylation in dimerization and activity of a GH32 enzyme and suggest the role of a flexible loop surrounding the catalytic pocket in discriminating substrate specificity.

Experimental Procedures

Organisms, Media, Plasmids, and Mutagenesis—X. dendrorhous ATCC MYA-131 was grown in MMM medium as referred previously (1). Pichia pastoris GS115 (his4-, Invitrogen) was used as expression host for the different enzyme variants. It was grown in YEPl medium (yeast extract 1%, peptone 2%, and glucose 2%) at 30 °C. MD media (YNB 1.34%, biotin 4 × 10^{-5} %, and glucose 2%) was used to select transformants. Expression and induction of proteins in P. pastoris were analyzed using BMG and BMM media (both are the same as MD but in potassium phosphate, pH 6.0, and glycerol 1% or methanol 0.5% as carbon sources, respectively). Yeast growth was followed spectrophotometrically at a 600-nm wavelength. Escherichia coli DH5α was used for DNA manipulation and amplification using the standard techniques.

The β-fructofuranosidase Xd-INV gene from X. dendrorhous (GenBank™ accession number FJ539193.2) comprises an open reading frame of 1995 bp corresponding to a 665-amino acid protein. The QDNS-pIB4 construction contained the 1902-bp fragment of Xd-INV responsible for the synthesis of the last 634 amino acids of this protein fused to the 267-bp fragment of the S. cerevisiae MFα secretion signal sequence (25). This construct was used as a template to obtain all the mutants generated in this work. Site-directed mutagenesis was carried out using specific primers, including substitutions responsible for mutations N58S, D80A, N107S, E303A, E334S/E334N/E334V/E334Q, N342S, H343A/H343T, N471S, and Y659STOP, and following the method described previously (21). DNA sequencing was used to verify that only the desired mutations were present in all the obtained constructs.

Protein Expression and Purification—X. dendrorhous β-fructofuranosidase was purified using tangential concentration and DEAE-Sephalchromatography, as described previously (26). Active fractions were concentrated using Microcon YM-10 (Amicon) filters and stored at −70 °C. β-Fructofuranosidase variants from X. dendrorhous (wild type and mutants) expressed in P. pastoris were purified as described elsewhere (25). ProtoBlue Safe Colloidal Coomassie-stained (National Scientific) denaturing gel electrophoresis (SDS-PAGE; 8% polyacrylamide) of the samples confirmed the purity of the protein fractions. Protein concentration was determined using the Bio-Rad microprotein determination assay and bovine serum albumin as standard.

After purification of XdINV, protein deglycosylation was performed using endoglycosidase H (Endo H; New England Biolabs) as described previously (26). Deglycosylated protein was subjected to a 1-min ultrafiltration using an Amicon Ultra-4 50K device (Millipore) to remove the Endo H present in the sample and was subsequently concentrated to about 8 mg ml^{-1} using a 10-kDa cutoff membrane.

β-Fructofuranosidase Activity and Kinetics Assays—β-Fructofuranosidase hydrolytic activity was determined by the dinitrosalicylic acid method adapted to a 96-well microplate scale as described previously (25). Standard enzymatic assays were performed using 2% sucrose in 100 mM sodium phosphate buffer, pH 5.5, at 60 °C. Inulin (Orafti GR, BENE0 Ibérica, S.L. Barcelona, Spain) and fructans from Agave tequilana (kindly supplied by Dr. Georgina Sandoval, CIATEJ, Guadalajara, Mexico) at 1% were also used in this assay. One unit of activity was defined as that catalyzing the formation of 1 μmol of reducing sugar/min. Hydrolase activity analysis at 55-80 °C was carried...
out under the aforementioned conditions. The thermostability refers to the temperature required for 50% enzyme inactivation (enzyme half-life) after heating at 0.5 units of the pure enzyme at different temperatures (60–85 °C) during 10-120 min and was determined by removing samples at regular intervals and estimating the residual β-fructofuranosidase activity.

For kinetic analysis, the initial velocity was measured in triplicate with 45–30 μg ml⁻¹ of enzyme and using 0–80 mM sucrose, 0–60 mM raffinose, or 0–60 mM nystose. The reaction time was 20 min. The plotting and analysis of the curves were carried out using the kinetic module of SigmaPlot software (version 12). Kinetic parameters were calculated by fitting the initial rate values to the Michaelis-Menten equation.

The activity of several XdINV mutants was analyzed by zymogram analysis using non-denaturing gels (6% polyacrylamide) stained with 2,3,5-triphenyltetrazolium-chloride, as described previously (1). Bovine serum albumin was used as reference and negative control.

**Crystallization and Data Collection—** Crystals of XdINV were grown as described before (26). For data collection, native crystals were transferred to cryoprotectant solutions consisting of mother liquor plus 10% (v/v) glycerol before being cooled in liquid nitrogen. Diffraction data were collected using synchrotron radiation at the European Synchrotron Radiation Facility (ESRF, Grenoble) on ID23-1 beamline. Diffraction images were processed with iMOSFLM (27) and merged using the CCP4 package (28). Crystals of XdINV were indexed in the P2_1_2_1 space group with two molecules in the asymmetric unit and 70% solvent content within the unit cell (Table 1).

The inactive D80A mutant was deglycosylated as reported for native enzyme and tested for crystallization. The best diamond-shaped crystals were grown by mixing equal amounts of 8 mg ml⁻¹ protein (in 20 mM Tris-HCl, pH 7) and a solution consisting of 1.3 M sodium citrate and 0.1 M Hepes or 0.1 M BisTris propane, pH 7.5. D80A complexes with 1-kestose or nystose (both from TCI-Europe), raffinose (Sigma), neokestose or neoerlose, both purified from reactions including XdINV as described in references (1, 10), were obtained by removing samples at regular intervals and estimating the residual β-fructofuranosidase activity.

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**Structure Solution and Refinement—** The structure of XdINV was solved by molecular replacement using the MOLREP program (30). The best model was the inverase from Aspergillus japonicus (PDB code 3LF7), having 34% identity (48% similarity), from which a template was prepared using the program Chainsaw (31) and a protein sequence alignment of XdINV onto the template. A solution containing two molecules in the asymmetric unit was found using reflections up to 2.5 Å resolution and a Patterson radius of 40 Å, which after rigid body fitting led to an R-factor of 49%. Crystallographic refinement was performed using the program REFMAC (32) within the CCP4 suite with flat bulk-solvent correction, maximum likelihood target features, and local non-crystallographic symmetry. Free R-factor was calculated using a subset of 5% randomly selected structure-factor amplitudes that were excluded from automated refinement. Extensive model building using the program COOT (33) was combined with several rounds of refinement leading to a model showing a continuous density for the whole polypeptide chain. At the later stages, carbohydrate and water molecules were included in the model, combined with more rounds of restrained refinement that led to a final R-factor of 16.9 (R_free = 18.8) for all data set up to 2.14 Å resolution. Refinement parameters are reported in Table 1.

The structure of D80A mutant was solved by molecular replacement using Molrep and the coordinates of native XdINV as the search model. A first cycle of refinement was performed using REFMAC, with model building with COOT. The structures of the D80A complexes were solved by difference Fourier synthesis using these refined coordinates. The ligands were manually built into the electron density maps and were refined similarly, to reach the R-factors listed in Table 1. For neokestose and neoerlose, not present in the Protein Data Bank, a model was built by the on-line biomolecule building program GLYCAM (34). Then PRODRG (35) was used to automatically generate molecular topologies suitable for REFMAC refinement.

**Results**

**XdINV Folding Displays Novel Structural Features—** We have purified and crystallized the deglycosylated form of the wild-type β-fructofuranosidase XdINV, as reported previously (26). Treatment of this highly glycosylated protein (monomer of 160–200 kDa) with Endo H led to a molecular mass of 66 kDa and reduced drastically the heterogeneity of the sample, allowing us to obtain high quality crystals. We present here its three-dimensional structure solved by molecular replacement at 2.14 Å resolution. Experimental and structural determination details are given under “Experimental Procedures” and in Table 1. It should be noted that the deglycosylated enzyme retains activity, and heterogeneity was reduced without affecting protein quaternary structure, as shown below. The crystals belong...
### Table 1
Crystallographic statistics (Values in parentheses are for the high resolution shell)

| Crystal data | XdINV | XdINV-D80A / sucrose | XdINV-D80A / 1-kestose | XdINV-D80A / nystose | XdINV-D80A / neokestose | XdINV-D80A / neocerlose | XdINV-D80A / raffinose | XdINV-D80A / Fru/Hepes | XdINV-D80A / Fru/BTpro |
|--------------|-------|----------------------|------------------------|----------------------|------------------------|------------------------|------------------------|------------------------|------------------------|
| Space group  | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ | $P_{2_1} \times 2$ |
| Unit cell parameters |
| a (Å) | 74.62 | 74.73 | 74.53 | 74.55 | 74.75 | 74.57 | 74.62 | 74.57 | 74.43 |
| b (Å) | 205.18 | 205.22 | 205.019 | 205.41 | 205.27 | 205.07 | 204.77 | 205.07 | 204.74 |
| c (Å) | 145.77 | 146.07 | 145.99 | 146.36 | 145.90 | 146.59 | 146.50 | 146.26 | 145.56 |
| Data collection |
| Beamline | ID23-1 (ESRF) | XALOC (ALBA) | P13 (DESY) | XALOC (ALBA) | XALOC (ALBA) | P13 (DESY) | XALOC (ALBA) | P13 (DESY) | ID23-1 (ESRF) |
| Temperature (K) | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 | 100 |
| Wavelength (Å) | 1.0395 | 1.10704 | 0.9786 | 0.97949 | 1.10704 | 0.9786 | 0.9786 | 0.96863 |
| Resolution (Å) | 6.1971 | 2.14 | 119.00-2.01 | 118.92-1.78 | 119.20-1.78 | 118.92-2.05 | 119.25-1.88 | 119.15-1.82 | 119.07-1.91 | 118.63-1.84 |
| (2.26-1.89) | (2.01-1.78) | (1.81-1.78) | (2.05-1.88) | (1.91-1.82) | (1.85-1.82) |
| Multiplicity | 7.1 (6.8) | 6.8 (6.8) | 6.7 (6.8) | 6.7 (6.7) | 6.7 (6.8) | 6.7 (6.9) | 6.8 (6.8) | 6.8 (7.0) | 4.5 (4.6) |
| Completeness (%) | 99.9 (99.9) | 99.0 (98.1) | 99.9 (99.8) | 100.0 (100.0) | 99.9 (99.8) | 99.3 (98.6) | 99.6 (99.5) | 100.0 (100.0) | 99.9 (99.9) |
| Mean $I/\sigma (I)$ | 12.8 (4.3) | 9.6 (3.4) | 19.5 (3.3) | 16.2 (3.6) | 10.1 (3.4) | 7.4 (3.0) | 9.6 (2.6) | 10.1 (2.1) | 7.4 (1.9) |
| $R_{merge}$ (%) | 10.8 (38.9) | 11.8 (55.6) | 6.2 (53.4) | 6.9 (55.2) | 11.5 (59.8) | 15.8 (59.1) | 6.8 (58.8) | 7.2 (48.0) | 6.4 (34.3) |
| $R_{work}^{*}$ (%) | 3.3 (16.8) | 4.9 (22.4) | 2.6 (22.1) | 2.9 (23.0) | 4.8 (24.8) | 6.6 (24.2) | 2.8 (24.0) | 3.0 (19.4) | 3.2 (17.4) |
| Molecules per ASU | 2 | 2 | 2 | 2 | 2 | 2 | 2 | 2 |
| Refinement |
| $R_{work} / R_{free}^{**}$ (%) | 16.9/18.8 | 16.6/18.5 | 17.0/18.4 | 16.7/17.8 | 16.2/18.3 | 16.8/18.3 | 16.9/18.5 | 17.0/18.5 | 16.3/17.9 |
| N° of atoms/average B (Å$^2$) |
| Protein | 9616 / 18.6 | 9610 / 29.82 | 9610 / 24.75 | 9610 / 24.08 | 9610 / 30.86 | 9610 / 25.78 | 9610 / 24.07 | 9610 / 25.55 | 9610 / 23.34 |
| Carbohydrate | 820 / 38.53 | 913 / 53.94 | 889 / 45.75 | 911 / 45.90 | 842 / 54.79 | 889 / 47.61 | 878 / 44.13 | 875 / 48.03 | 897 / 46.64 |
| Water molecules | 1217 / 27.4 | 1514 / 41.42 | 1642 / 36.71 | 1445 / 36.50 | 1316 / 41.50 | 1623 / 39.18 | 1597 / 35.50 | 1491 / 35.68 | 1496 / 34.56 |
| All atoms | 11065 / 20.93 | 12037 / 33.11 | 12141 / 27.90 | 11966 / 27.24 | 11768 / 33.765 | 12122 / 29.18 | 12085 / 27.04 | 11976 / 28.45 | 12003 / 26.48 |
| Ramachandran plot (%) |
| Favored | 95.43 | 96.00 | 96.00 | 96.00 | 96.00 | 96.00 | 96.00 | 96.00 | 96.00 |
| Outliers | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| RMS deviations |
| Bonds (Å) | 0.009 | 0.008 | 0.007 | 0.007 | 0.008 | 0.007 | 0.007 | 0.007 | 0.007 |
| Angles (°) | 1.372 | 1.506 | 1.402 | 1.371 | 1.494 | 1.405 | 1.402 | 1.450 | 1.413 |
| PDB accession codes | 5ANN | 5FIX | 5FKB | 5FMD | 5FK7 | 5FK8 | 5FKC | 5FMB | 5FMC |
to the space group P2₁2₁2, with two molecules per asymmetric unit that are related by a non-crystallographic 2-fold symmetry axis. No significant structural differences were observed between the two chains (Ser-38–Tyr-665) that present a root mean square deviation of 0.325 Å on 627 matched Ca atoms.

In common with the GH32 enzymes, XdINV is bimodular (Fig. 1) and folds into a catalytic β-propeller domain and a β-sandwich domain. Two well-ordered glycan chains are observed at positions Asn-58 and Asn-107, whereas Asn-471 is the only position conserved within GH32 from fungi. Each molecular surface of XdINV with the same color code, and the N-glycan chains shown as sticks. C. XdINV is a highly glycosylated protein containing 18 potential glycosylation sites; apart from the two long N-glycans, GlcNAc units have been found in the crystal N-attached to other Asn residues, their location being indicated by small yellow spheres.

FIGURE 1. Structure of XdINV. A, ribbon diagram of XdINV monomer showing the bimodular molecular architecture, colored in rainbow sequence-code, which folds into a catalytic β-propeller domain and a β-sandwich domain. Two well-ordered glycan chains are observed at positions Asn-58 and Asn-107, whereas Asn-471 is the only position conserved within GH32 from fungi. B, molecular surface of XdINV with the same color code, and the N-glycan chains shown as sticks. C, XdINV is a highly glycosylated protein containing 18 potential glycosylation sites; apart from the two long N-glycans, GlcNAc units have been found in the crystal N-attached to other Asn residues, their location being indicated by small yellow spheres.
N471S substitutions on the thermostability of XdINV was evaluated. Accordingly, these protein variants were purified, and their sensitivities to different temperatures were studied as described under “Experimental Procedures.” The N471S variant behaved similarly to the wild-type protein, displaying maximum activity levels (75–100%) in the range of 60–75 °C, whereas the N58S variant activity decreased to about 40 and 20%, at 70 and 75 °C, respectively (Fig. 4C).

FIGURE 2. Structural alignment of GH32 from fungi. The catalytic domain of XdINV is superimposed to the A. japonicus fructosyltransferase, AjFT (PDB code 3LEM), A. awamori exo-inulinase, AaEI (PDB code 1Y9G), S. cerevisiae invertase, ScINV (PDB code 4EQV), and S. occidentalis/H9252-fructofuranosidase, SoFfase (PDB code 3U75), as given by the DALI server (44). The red squares indicate amino acid similarity as calculated by ESPript (45). A–D indicate the β-strands, from the axis toward the outside of the propeller, within each blade. Loops L1–L4, connecting the different blades of the propeller, and turns T1–T4, linking strands B–C within each blade, are underlined with an orange bar. XdINV glycosylation sites are highlighted in yellow. Catalytic residues are marked with a black asterisk. Residues of XdINV involved in polar (angle) and hydrophobic (asterisk) interactions within the dimer interface are indicated in green. Residues of XdINV mutated in this work are marked with a cyan asterisk.
results indicate that glycosylation at Asn-58 contributes to stabilize the structure of XdINV, whereas the role of that attached to Asn-471 is unclear.

XdINV Presents Unique Traits within Known GH32 Enzymes from Fungi—The three-dimensional structure of five other GH32 members from fungi has been reported as follows: the Aspergillus awamorii exo-inulinase (AaEI (15)), the A. japonicus fructosyltransferase (AjFT (16)), the Aspergillus ficuum endo-inulinase (17), the S. occidentalis β-fructofuranosidase (SoFase (21)), and the S. cerevisiae invertase (ScINV (22)). XdINV shares the highest identity (37%) with AjFT, whereas 20–25% identity is found with respect to the other enzymes, a value equivalent to that observed when XdINV is compared with the GH32 enzymes from bacteria and plants. Structural alignment of the catalytic domain of XdINV onto the fungal exo-enzymes (all but A. ficuum endo-inulinase) is shown in Fig. 2. As observed, both XdINV and AjFT present long loops linking the different β-sheets, and remarkably, AjFT presents also the inserted region at the N terminus described in XdINV but not the C-terminal extension. In contrast, Asn-471, at the segment linking the β-structure to its catalytic domain, is the sole conserved Asn residue in all the sequences, although Asn-52 and Asn-58 are present in AjFT and Asn-107 is only conserved in AaEI. Consequently, the glycosylation pattern of XdINV and the concurrence of Asn-107 and the C-terminal extension, both

| Molecule A          | Molecule B          | Distance Å |
|---------------------|---------------------|------------|
| Molecule A          | Molecule B          | Distance Å |
| Intermolecular      |                     |            |
| Tyr-200 (OH)        | Thr-655 (OG1)       | 2.66       |
| Ala-202 (N)         | Trp-651 (O)         | 3.04       |
| Trp-651 (O)         | Ala-202 (N)         | 2.92       |
| Trp-134 (NE1)       | MAN111 (O3)         | 2.77       |
| Gln-137 (OE1)       | NAG107 (O3)         | 2.60       |
| Gln-137 (NE2)       | NAG107 (O4)         | 3.00       |
| Gln-137 (NE2)       | NAG108 (O7)         | 3.01       |
| Asp-155 (OD2)       | MAN115 (O4)         | 2.70       |
| Lys-197 (N)         | MAN111 (O6)         | 3.05       |
| Intramolecular      |                     |            |
| Glu-128 (OE1)       | NAG108 (N2)         | 2.85       |
| Glu-128 (OE1)       | NAG107 (O6)         | 2.70       |
| Lys-131 (NZ)        | NAG107 (O5)         | 2.74       |
| Lys-131 (NZ)        | NAG107 (O6)         | 2.96       |
| Lys-405 (N)         | NAG059 (O6)         | 2.90       |
| NAG108 (O3)         | BMA109 (O6)         | 2.98       |
| NAG58 (O6)          | NAG59 (O6)          | 3.45       |
| NAG59 (O3)          | BMA60 (O6)          | 3.30       |
shown to be key determinants in oligomerization, are unique to this protein. Moreover, as can be observed in Fig. 2, loops L1, T1, and L2 are highly variable in sequence, and the residues involved in polar links or interatomic interactions at the dimer interface are not conserved.

It is worth noting that from all GH32 enzymes, only the reported structures from yeasts, SoFfase and ScINV, revealed the existence of dimeric forms. However, the dimerization pattern of those enzymes was different from that observed in XdINV (Fig. 5), with blades 4 and 5 involved in the interface of this protein. Moreover, as can be observed in Fig. 2, loops L1, T1, and L2 are highly variable in sequence, and the residues involved in polar links or interatomic interactions at the dimer interface are not conserved.

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**FIGURE 4. Role of glycosylation in XdINV.** A, PAGE analysis of the XdINV variants. Culture filtrates (200 ml) of the P. pastoris transformants expressing the wild-type (wt) or the indicated XdINV mutants were concentrated through 50000 M cutoff (PVDF membrane) and applied to DEAE-Sephacel chromatography. Active fractions eluting at 0.1 M NaCl were concentrated to 0.2–0.5 mg/ml. The enzyme activities (100 μl; about 2–5 μg of proteins) were revealed in situ (left) and subsequently stained using colloidal Coomassie blue (right). The mutant N107S was processed in the same way, but 20 μl of the final concentrate was analyzed. B, referred proteins were analyzed by SDS-PAGE. Numbers at the right in A and B indicate the positions of the bovine serum albumin (BSA) molecular masses and the weight markers used as control (in kDa), respectively. C, temperature dependence profiles. Activity of the wild-type (filled triangles), N58S (filled circles), and N471S (open circles) protein variants were evaluated at the indicated temperatures. Each value represents the average for four independent measurements. Standard errors are indicated.

**FIGURE 5. XdINV versus SoFfase dimers.** A, molecular surface of XdINV dimer showing the two catalytic domains in light green, and the corresponding β-sandwich domains are colored in sand. Each active site cavity is filled with a fructosyl-nystose molecule (in spheres) as found in the complexes obtained in this work and described below. A modelled β(2–6)-linked fructose chain (in sticks) has been manually docked into the long crevice that encompasses the two active sites and beyond. Inset, a detail of this crevice as viewed from the position indicated by the arrow, i.e., along the modeled levan-type fructose chain; a nystose molecule (spheres) is visible at one of the active sites. B, molecular surface of the SoFfase dimer showing the catalytic domains in cyan and their β-sandwich domains in yellow. Each active site is located at one face of the dimer, and only a fructosyl-nystose molecule found in the crystal (24) is visible as spheres.

**TABLE 3**

| XdINV variant | Sucrose | Raffinose | Nystose | Sucrose | Raffinose | Nystose | Sucrose | Raffinose | Nystose | Sucrose | Raffinose | Nystose |
|---------------|---------|-----------|---------|---------|-----------|---------|---------|-----------|---------|---------|-----------|---------|
| WT            | 1.9 ± 0.3 | 7.7 ± 1.6 | 21.7 ± 3.6 | 911 ± 47 | 291 ± 51 | 148 ± 25 | 479 ± 68 | 56 ± 10 | 72 ± 12 |
| N58S          | 6.1 ± 1.7 |          |         | 711 ± 200 |          |         | 117 ± 33 |          |         |         |
| E343Q         | 3.6 ± 0.6 | 15.5 ± 3.4 | 1.9 ± 0.4 | 91 ± 15 | 7 ± 1 | 46 ± 10 | 25 ± 4 | 0.4 ± 0.1 | 23 ± 5 |
| Q341N         | 2.2 ± 0.1 | 8.4 ± 2.0 | 30.6 ± 6.9 | 377 ± 76 | 171 ± 40 | 71 ± 16 | 212 ± 43 | 20 ± 5 | 2 ± 1 |
| N342S         | 7.3 ± 1.3 | 2.9 ± 0.7 | 12.7 ± 2.1 | 1070 ± 189 | 165 ± 41 | 65 ± 11 | 147 ± 26 | 58 ± 14 | 5 ± 1 |
| H343A         | 8.3 ± 1.5 | 7.5 ± 1.5 | ND      | 105 ± 19 | 32 ± 6 | ND      | 13 ± 2 | 4 ± 1 | ND      |
| H343T         | 12.6 ± 3.3 | 12.9 ± 3.5 | ND      | 855 ± 223 | 101 ± 27 | ND      | 68 ± 18 | 8 ± 2 | ND      |
| N471S         | 3.4 ± 0.9 |          |         | 1151 ± 307 |          |         | 339 ± 91 |          |         |         |
the SoFfase and ScINV dimers through loops L₃, L₄, and the segment linking CS-D5 (Fig. 2). Nevertheless, the β-sandwich domain is involved in the assembly of the two types of dimers and also in shaping the active site cavity, although through an evident different pattern. Thus, this feature seems to be a common trend of the GH32 enzymes from yeast, and remarkably, it is the only case in which a role has been assigned to the β-sandwich domain within the GH32 family enzymes. A notable feature of XdINV is that, unlike SoFfase, the particular architecture of its dimer houses the active sites at the same face and separated 30 Å away. A close inspection of the molecular surface reveals a long crevice encompassing the two active sites and beyond. Therefore, it is appealing to suggest that this disposition might be conferring the XdINV dimer the additional ability to accommodate a putative polymeric substrate, as illustrated in Fig. 5. In this way, the XdINV cleft could be able to allocate, for instance, a linear β(2–6)-linked fructose chain with β(2–1) branches, characteristic of mixed fructans or graminans (38), its β(2–1) branches being hydrolyzed at each catalytic pocket. In agreement with this, XdINV showed a small but significant hydrolase activity on fructan polymers, showing a moderate activity on inulin (72 units/mg versus 960 units/mg on sucrose) as is observed in other GH32 enzymes. Furthermore, XdINV is also able to hydrolyze agave fructans (15 units/mg), which consist of a complex mixture of fructose-based polymers containing principally β(2–1) linkages but also β(2–6) (39). Although this is not evidence that branched substrates can bind to both sites, the channel-like topology of the dimer surface is, indeed, a remarkable trait of XdINV.

**Structure of the XdINV Complexes Discloses the Binding Subsites**—Inactivated XdINV D80A mutant was crystallized and used for soaking experiments with different donor substrates, including sucrose, 1-kestose, neokestose, and nystose; from these, 1-kestose and neokestose are also transfructosylation products. As has been reported recently (10), XdINV is able to fructosylate glucose-containing oligosaccharides as maltose, which produces neoerlose. To depict this activity, this trisaccharide has been also included in the soaking experiments. Furthermore, in an attempt to capture a putative intermediate complex, mutant crystals were pre-incubated with fructose before being transferred to solutions containing tested acceptors as maltotriose, isomaltulose, isomaltose, and raffinose. However, only raffinose was found at the active site, but it was occupying the catalytic pocket in a donor substrate mode. Interestingly, fructose and a molecule of Hapes or Bistris propane, both from the buffer, were found at the active site in some of the crystals mimicking the location of an acceptor for the fructose in a putative intermediate complex, as will be commented on below.

The catalytic site is buried in a deep pocket 20 Å deep and about 10 × 10 Å width. All the obtained complexes show clear electron density at the active site, which allowed unambiguous ligand modeling that situated the terminal fructose at the bottom and in contact with the catalytic residues (Fig. 6). This fructose at subsite −1 keeps the same atomic interactions and features previously described for other complexes from GH32 enzymes (16, 19, 24). Remarkably, a net of well ordered water molecules is found in all complexes, contributing to substrate recognition. The only significant differences found in the protein upon substrate binding are located at loop T₁ᵥ (residues Glu-334–His-343), which is rather disordered in the free enzyme and is stabilized in the complexes making crucial interactions, as explained below.

The glucose bound at subsite +1 in the complex with sucrose, neokestose, and raffinose is tightly fixed by direct hydrogen bonds to the acid/base catalytic Glu-303 (through its O2), and to Glu-334 (O2 and O3) and His-343 (O2), both from loop T₁ᵥ. The glucose is further stabilized by interaction of its O4 to Leu-170 and Ala-172 from loop T₁ᵥ through several water molecules. In the case of 1-kestose and nystose, having fructose at this subsite, and neoerlose, with a β(2–6) linked glucose, a water molecule is occupying the position of the glucose O2 observed in the sucrose complex, therefore mediating the same interaction pattern to Glu-334 and His-343.

Subsite +2 is made by stacking interaction to Trp-105 from loop Tₚ, which allocates the corresponding sugar unit in a variety of modes revealing a noticeable plasticity. Apart from this hydrophobic interaction, there are many hydrogen bonds to different residues, always through numerous water molecules and, also, many intramolecular hydrogen bonds fixing each substrate in a precise conformation. First of all, the C-terminal extension is making a polar interaction through Leu-661 main chain, conserved in all the complexes. But additional interactions depend on the type of substrate and, thus, neokestose, raffinose, and neoerlose are accommodated between loop T₁ᵥ and the C-terminal segment by multiple polar interactions to Leu-170, Ala-172, Gly-660, and Leu-661. By contrast, 1-kestose and nystose are oriented toward loop T₁ᵥ and interacting with Gln-341. Finally, subsite +3, occupied only in the nystose complex, is mainly shaped by loop T₁ᵥ and shows direct polar interactions of the terminal glucose with Gln-335, Asn-342, and His-343.

In summary, subsite +1 is common to all complexes, being shaped by T₁ᵥ through Leu-170 and Ala-172 main chain and by T₁ᵥ through Glu-334 and His-343. Subsite +2 is created by stacking to Trp-105 and polar interaction to the main chain of the C-terminal region (Gly-660 and Leu-661), but two different sites are observed depending on the substrate, and thus, inulin-type ligands are accommodated by loop T₁ᵥ through Gln-341, whereas levan or neo-type ligands are allocated by loop T₁ᵥ (Leu-170 and Ala-172) (Fig. 7). An additional subsite +3 for inulin-type substrate is formed essentially by loop T₁ᵥ (Gln-335, Gln-341, Asn-342, and His-343). In summary, the structure of the XdINV complexes presented here suggests a role of loop T₁ᵥ in regulating specificity of this protein, especially through residues Glu-334, Gln-341, and His-343.

**Mutational Analysis Unveils Key Residues Modulating Activity**—The role of the leading residues identified in the complex analysis was investigated by kinetic analysis of mutants that is given in Table 3. First, the expected lack of activity observed in mutants D80A and E303A is consistent with its catalytic role as nucleophile and electron/proton donors, respectively. But, remarkably, the presence of glutamate at position 334 seems to be also required for sucrose hydrolysis, as its replacement by serine, valine, or glutamine is deleterious for activity and leads to fully inactivated enzymes, revealing the
The major role of Glu-334 in binding sucrose. Only the mild E334Q replacement gives detectable activity on sucrose, and interestingly, this mutant keeps a significant activity on nystose. As described above, Glu-334 is making a direct hydrogen bond to the sucrose O2 and O3 atoms, from the glucose located at site +1, and therefore, some effect upon its replacement was expected in the hydrolysis of this substrate. The presence of an acidic residue at this position seems necessary for sucrose binding and hydrolysis in many clan GH-J enzymes, and as such, the equivalent D239A mutant of *Arabidopsis thaliana* invertase exhibited a 6-fold increase in $K_m$ against sucrose (40). Moreover, although this Asp/Ala replacement compromised its ability to degrade sucrose, the enzyme retained fructan hexohydrolase activity (41). Thus, from sequence analysis of plant GH32 enzymes, it was proposed that members having an Asp-239 homolog have sucrose as the preferential donor, whereas enzymes lacking this residue use fructans as their donor substrate (41). In agreement with this role in regulating specificity, kinetic analysis of SoFfase mutants at Gln-228 position, structurally related to the *A. thaliana* Asp-239, revealed that Gln-228 is essentially responsible for conferring affinity for acceptor sucrose in the transfructosylation reaction (24). However, in that case, even the SoFfase Q228V and Q228T mutants retained moderate invertase activity. Consequently, the XdINV Glu-334 plays a more essential role in sucrose binding than that reported for the yeast SoFfase, resembling the behavior described for plant GH32 enzymes.

Mutants H343A/H343T also showed drastically reduced activity on the assayed substrates. As observed in the complexes, His-343 establishes a direct hydrogen bond to the glucose O2 from sucrose, and in agreement, there is a concomitant decreased binding affinity as revealed by its $K_m$ value. This position is equivalent to Lys-242 from the *A. thaliana* invertase that, together with Asp-239, is proposed to be a hot spot mediating sucrose binding. Furthermore, this position is structurally equivalent to the SoFfase Asn-254, which was reported to be involved in altering the hydrolase/transferase ratio but mostly in modulating the transfructosylating product specificity (24). According to this observation, preliminary analysis shows that the H343T mutant shows an altered profile of FOS production.
Fructans, the fructose-rich polymers biosynthetically derived from sucrose, are important non-structural storage compounds in many bacteria, fungi, and numerous plant species. They are associated with essential biological processes as sucrose metabolism and vegetative development. Also, fructans are biotechnologically relevant compounds due to its prebiotic effect. Consequently, the enzymes involved in fructans and sucrose processing are subject to extensive research with the aim to investigate new potential acceptors yielding novel bioconjugates of biomedical interest.

**Discussion**

Fructans, the fructose-rich polymers biosynthetically derived from sucrose, are important non-structural storage compounds in many bacteria, fungi, and numerous plant species. They are associated with essential biological processes as sucrose metabolism and vegetative development. Also, fructans are biotechnologically relevant compounds due to its prebiotic effect. Consequently, the enzymes involved in fructans and sucrose processing are subject to extensive research with the aim to investigate new potential acceptors yielding novel bioconjugates of biomedical interest.
Structure and Function of Dimeric XdINV
aim to uncover their biological role and also to improve their enzymatic efficiency.

In this study, we described the structure of $\beta$-fructofuranosidase from \(X.\ dendrorhous\) that exhibits novel features regulating activity. The crystal structure shows a bimodular arrangement of its polypeptide chain similar to that described previously for other GH32 enzymes. However, a long 30-residue extension at its C terminus, together with a glycan chain attached to a non-conserved Asn, mediates the formation of an unusual dimer. This dimer shows a peculiar topology that locates both active sites in the same face of the protein and apparently connected by a long crevice. A blast search shows about 60 entries showing 30–60% similarity with XdINV sequence. These sequences include hydrolases or transferases mostly from \(B.\) globiformis but also from \(A.\) nidulans (data not shown). Alignment of the corresponding sequences reveals that they all display both determinants that are shown to be key determinants for the oligomerization state in XdINV, i.e., the C-terminal extension and the N-glycosylation sequon at a position equivalent to Asn-107, and consequently, the dimer described here would be a model for this group of GH32 enzymes.

The unusual topology of the XdINV dimer suggests the possibility that it might be able to accommodate a putative polymeric substrate as it could be a mixed fructan with branched $\beta$(2–1)/$\beta$(2–6) linkages (42, 43) that, indeed, is degraded by XdINV. Fructans present in plants are a mixture of heterogeneous oligomers and polymers with a composition depending on the environment and the development state of the plant (38). Key enzymes for plant fructan metabolism include a complex set of biosynthetic and hydrolytic enzymes, including different types of fructan exohydrolases and fructosyltransferases. Consequently, fungi isolated from plant environments may have evolved to interact or degrade these molecules, by expressing their own enzymes. Nevertheless, the possibility of the yeast producing these compounds cannot be discarded, as the biological function of microbial fructans has also been related to the creation of a physical barrier, by enhancing resistance to environmental stresses and assisting in nutrient assimilation (Ref. 43 and references therein). In fact, fructans have proved to act as virulence factors in colonization and to play a critical factor in the pathogenesis of disease-related microorganisms. Many studies have shown that bacteria that synthesize fructans interact with eukaryotic hosts and can be found in symbiotic association with plants. Although the reported studies have been performed in bacteria, it is conceivable that fungi develop a similar mechanism to adapt to the environment. In this sense, it should be noted that the yeast \(X.\ dendrorhous\) has been isolated from soil samples collected at different cold ecosystems. The exact physiological role of XdINV and the target substrate at its natural habitat are difficult to envisage, and therefore, it is hard to derive precise structure-function relationships, but clearly, the peculiar topology of its molecular surface is a remarkable and distinctive trait within GH32 enzymes.

The crystallographic analysis of the XdINV complexes described here gives a detailed picture of the main features of its active site. Previous studies showed that most GH32 enzymes present a shallow pocket with residues shaping a conserved fructose-binding pocket, at subsite $-1$, and several hydrophobic and polar interactions defining subsite $+1$. In contrast, the active site is loosely bound at subsite $+2$ or dangling into the solvent. In agreement with this observation, short substrates such as 1-ketose or raffinose are accommodated in essentially the same conformation in all the formerly described complexes (13, 16, 18, 19). In contrast, XdINV presents a deep cavity showing multiple binding sites, a feature that has been previously described for two other fungal enzymes, the AjFF and the SoFfase. The reported complexes of these enzymes with long substrates reveal that the active sites diverge from subsite $+2$, and thus, nystose bound at the AjFF catalytic pocket locates its terminal glucose at a position that is occupied by Trp-105 in XdINV (see Fig. 7A) (16). A very short T$_\text{1}$ loop and the absence of the C-terminal segment make a very different active site pocket in AjFF, which is remarkable taking into account that this enzyme is phylogenetically close to XdINV. Likewise, Fru-nystose bound at the SoFfase accommodates in a precise conformation that mimics the polymeric substrate inulin (24). As commented previously, both yeast enzymes are distinctive within GH32 in being dimers, and both active sites must be considered in terms of their oligomeric state. In the case of SoFfase, the topology of the active site seemed designed to optimally fit the conformation of the polymer inulin, which is consistent with its high inulinase activity. Thus, the topology of sites $+2$ and further from the cleaved linkage must reflect the profile of its natural substrate and consequently the physiological role of each enzyme in its environment. Remarkably, XdINV presents both inulin- and levan-type binding sites and an open pocket filled with many water molecules that confer plasticity to the enzyme. Therefore, this plasticity might be consistent with a more complex substrate as are the mixed fructans commented on previously, in contrast to the rigid active site of SoFfase adapted to accommodate the more regular inulin.

Carbohydrate-processing enzymes have developed intricate molecular mechanisms to cope with their highly heterogeneous substrates. Understanding these mechanisms will be of general interest to uncover essential biological processes at the molecular level. In the case of XdINV, an insertion at the C-terminal polypeptide and glycosylation have been shown to be essential in oligomerization and thus in shaping a peculiar active site. In turn, the plasticity of its active site makes the enzyme a valuable and flexible biocatalyst to produce novel compounds with bio-

**FIGURE 8. Proposed mechanism of XdINV transfructosylating activity.** A, view of the XdINV-neoerlose complex, showing the ligand in cyan and the main interactions of its Glu-\(\alpha\)1–4Glu moiety located at subsites $+1$ and $+2$. The position of this portion represents a potential binding mode of acceptor maltose for the formation of a $\beta$(2–6) linkage of glucose to fructose. The position of a putative sucrose acceptor molecule (white) has been inferred by superposition of its glucose to the glucose found at subsite $+1$ of neoerlose and adjusting manually the fructose unit by small torsion of its glycosidic bond to best packing to Trp-105 side chain. B, scheme of the proposed mechanism. The donor-substrate sucrose is hydrolyzed by nucleophilic attack forming the fructosyl-XdINV intermediate; a subsequent sucrose molecule (the acceptor substrate) can enter the active site pocket with binding mode 1 generating 1-ketose or, alternatively, with binding mode 2 and generating neokestose.
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technological relevance. The potential of XdINV to produce novel bioconjugates is underway.

Author Contributions—J. S. A. and M. F. L. conceived and coordinated the study. D. L. performed the production of the native protein. M. G. P., Z. M., and M. F. L. conducted molecular biology assays and kinetics measurements and analyzed the results. M. R. E., B. G., and J. S. A. designed and performed the crystallographic work and interpreted the results. J. S. A. wrote the paper, and all authors read and commented on the manuscript.

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