Understanding How Noncatalytic Carbohydrate Binding Modules Can Display Specificity for Xyloglucan*

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Ana S. Luís, Immacolata Venditto, Max J. Temple, Artur Rogowski, Arnaud Baslé, Jie Xue, J. Paul Knox, José A. M. Prates, Luís M. A. Ferreira, Carlos M. G. A. Fontes, Shabir Najmudin, and Harry J. Gilbert

From the 1CISFA, Faculdade de Medicina Veterinária, Universidade Técnica de Lisboa, Avenida da Universidade Técnica, 1300-477 Lisboa, Portugal, the 2Institute for Cell and Molecular Biosciences, The Medical School, Newcastle University, Newcastle-upon-Tyne NE 4HH, United Kingdom, and the 3Centre for Plant Sciences, Faculty of Biological Sciences, University of Leeds, Leeds LS2 9JT, United Kingdom

Background: Carbohydrate binding modules (CBMs) contribute to the enzymatic degradation of complex polysaccharide structures.

Results: New CBMs display specificity for decorated glucans through an extensive hydrophobic platform that interacts with both backbone and side chain structures.

Conclusion: CBMs that bind to complex β-glucans exploit different components of these ligands as specificity determinants.

Significance: CBMs can utilize the side chains of decorated glucans as specificity determinants.

Plant biomass is central to the carbon cycle and to environmentally sustainable industries exemplified by the biofuel sector. Plant cell wall degrading enzymes generally contain non-catalytic carbohydrate binding modules (CBMs) that fulfil a targeting function, which enhances catalysis. CBMs that bind β-glucan chains often display broad specificity recognizing β1,4-glucans (cellulose), β1,3-β1,4-mixed linked glucans and xyloglucan, a β1,4-glucan decorated with α1,6-xylose residues, by targeting structures common to the three polysaccharides. Thus, CBMs that recognize xyloglucan target the β1,4-glucan backbone and only accommodate the xylose decorations. Here we show that two closely related CBMs, CBM65A and CBM65B, derived from EcCel5A, a Eubacterium cellulosolvens endoglucanase, bind to a range of β-glucans but, uniquely, display significant preference for xyloglucan. The structures of the two CBMs reveal a β-sandwich fold. The ligand binding site comprises the β-sheet that forms the concave surface of the proteins. Binding to the backbone chains of β-glucans is mediated primarily by five aromatic residues that also make hydrophobic interactions with the xylose side chains of xyloglucan, conferring the distinctive specificity of the CBMs for the decorated polysaccharide. Significantly, and in contrast to other CBMs that recognize β-glucans, CBM65A utilizes different polar residues to bind cellulose and mixed linked glucans. Thus, Gln106 is central to cellulose recognition, but is not required for binding to mixed linked glucans. This report reveals the mechanism by which β-glucan-specific CBMs can distinguish between linear and mixed linked glucans, and show how these CBMs can exploit an extensive hydrophobic platform to target the side chains of decorated β-glucans.

The plant cell wall represents a major nutrient for numerous microbial ecosystems, exemplified by bacterial and fungal communities established in the rumen and large bowel of mammals, where they play an important role in animal nutrition and human health, respectively (1, 2). It is also evident that these composite structures are of increasing industrial significance, particularly in the environmentally relevant bioenergy and bioprocessing sectors (3, 4). The complex physical and chemical structure of the plant cell wall restricts its access to degradative enzymes. Microorganisms that utilize plant biomass as a significant nutrient express extensive repertoires of degradative enzymes, primarily, glycoside hydrolases but also lyases and esterases, which attack the structural polysaccharides of the plant cell wall (5).

A common feature of plant cell wall degrading enzymes is their complex modular architecture in which the catalytic module is appended to one or more noncatalytic carbohydrate binding modules (CBMs) (6) (see Ref. 6 for review), which are grouped into sequence-based families on the CAZy database (7). The general function of CBMs is to direct the cognate catalytic modules to their target substrate within the plant cell wall, thereby increasing the efficiency of catalysis (8–10). The majority of CBMs display a β-sandwich fold with the ligand binding site located in either the concave surface presented by one of the β-sheets, a topography that facilitates the targeting of the inter-

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This article contains Supplemental Table S1 and Fig. S1.

The atomic coordinates and structure factors (codes 4ACK, 2YPJ, and 4AFM) have been deposited in the Protein Data Bank (http://wwpdb.org/).

1 These authors contributed equally to this work.

2 To whom correspondence may be addressed. Tel.: 351-213652876; Fax: 351-213652889; E-mail: cafontes@fmv.utl.pt.

3 To whom correspondence may be addressed. Tel.: 441912228800; Fax: 441912227424; E-mail: h.j.gilbert@ncl.ac.uk.

4 The abbreviations used are: CBM, carbohydrate binding module; ITC, isothermal titration calorimetry; PDB, Protein Data Bank.
nal regions of glycan chains (11–13), or in the loops that connect the two sheets (14, 15). This latter binding site can either target the end (15) or, less frequently, the internal regions of glycan chains (14).

The majority of CBMs that target the plant cell wall bind to crystalline cellulose, single chains of β-glucans and xylan (see Ref. 6 for review). Binding to crystalline cellulose by Type A CBMs is mediated by a planar hydrophobic surface, which makes apolar contacts with exposed cellulose chains (16). Ligand recognition in CBMs that bind to the internal regions of single polysaccharide chains can be highly specific, exemplified by the CBM6 from the clostridial xylanase CjXyn10B, which exclusively targets xylan (14), whereas examples of promiscuous specificity include CmCBM6 and the Cellvibrio endoglucanase CmLic5A that binds to both β1,4-glucans and mixed linked β1,3-β1,4-glucans (17), and the CBM62 from the Cellvibrio xylanase CjXyn11A that recognizes β-glucans, xylans, and even β-galactans (15). In both examples plasticity in ligand recognition is achieved through binding to a conserved element of the target glycan, demonstrated by the primary binding site of CmCBM6, which is specific for cellobiose (Glc-β1,4-Glc), a structure found in both cellulose and the mixed linked glucan (17). It is unclear, however, whether a cohort of CBMs exist that recognize diverse glycans by binding to distinct structures in the different target ligands. In addition, all CBMs that recognize β-glucans, also bind to xyloligase, a β1,4-glucan that is decorated with α1,6-linked xylose residues. It would appear that these proteins accommodate, but do not target the xylose side chains. Indeed, to date no CBMs have been described that display a preference for xyloligase over other β-glucans (12).

A recent report has identified two modules in a Eubacterium cellulosolvens endoglucanase (EcCel5A), designated hereafter as CBM65A and CBM65B, that bind to both disordered cellulose and mixed linked glucans (18). In this study we have exploited the two CBM65s as a model system to understand the mechanistic basis for the diverse specificities displayed by some CBMs. We show that the CBM65s, uniquely, display a significant preference for xyloligase. The structure of CBM65B in complex with a xyloligase-derived oligosaccharide, in combination with mutagenesis studies on CBM65A, revealed the mechanism by which these proteins display a preference for xyloligase. The ligand binding cleft contains an unusually large number of aromatic residues that are optimized to not only make apolar contacts with the glucan backbone, but also make hydrophobic interactions with the xylose side chains. In addition to the dominant apolar contacts, CBM65A contains two polar residues that play an important role in binding undecorated β-glucans. Gln106 confers specificity for β1,4-glucan (cellobiose), whereas Gln110 interacts with both cellulose and mixed linked β1,3-β1,4-glucans.

**MATERIALS AND METHODS**

**Protein Production and Purification**—DNA encoding the CBM65A (residues 37–170 of EcCel5A) and CBM65B (residues 581–713 of EcCel5A) were synthesized (NZYTech Ltd., Portugal) with codon usage optimized for expression in Escherichia coli. The synthesized genes contained engineered NheI and Xhol recognition sequences at the 5′ and 3′ ends, respectively, which were used for subsequent subcloning into the E. coli expression vector pET28a (Novagen), generating pCBMAL1 and pCBMAL2, which encode CBM65A and CBM65B, respectively. Both CBMs contain an N-terminal His8 tag. E. coli Tuner DE3 cells harboring pCBMAL1 and pCBMAL2 were cultured in Luria-Bertani broth containing kanamycin (50 μg/ml) at 37 °C to mid-exponential phase (A560nm = 0.6) and recombinant protein expression was induced by the addition of 0.2 mM isopropyl β-D-1-thiogalactopyranoside and incubation for a further 16 h at 19 °C. The His8-tagged recombinant CBMs, and their respective mutants (see below), were purified from cell-free extracts by immobilized metal-ion affinity chromatography as described previously (19).

For crystallization, CBM65A was further purified by size exclusion chromatography. Following immobilized metal-ion affinity chromatography, fractions containing the purified proteins were buffer exchanged, using PD-10 Sephadex G-25M gel-filtration columns (GE Healthcare), into 50 mM HEPES-Na buffer, pH 7.5, containing 200 mM NaCl and 5 mM CaCl2, and was then subjected to gel filtration using a HiLoad 16/60 Superdex 75 column (GE Healthcare) at a flow rate of 1 ml/min. Purified CBM65A was concentrated using an Amicon 10-kDa molecular mass centrifugal concentrator and washed three times with 1 mM CaCl2. Preparation of E. coli to generate selonomethionine CBM65A was performed as described in Ref. 20 and the protein was purified using the same procedures as employed for the native CBM. Purified CBM65A was concentrated using an Amicon 10-kDa molecular mass centrifugal concentrator and washed three times with 5 mM DTT. SDS-PAGE showed that all the recombinant proteins were more than 95% pure after Coomassie Blue staining.

**Site-directed Mutagenesis**—Site-directed mutagenesis was carried out employing a PCR-based NZYMutagenesis site-directed mutagenesis kit (NZYTech Ltd.) according to the manufacturer’s instructions, using pCBMAL1 as the template. The sequence of the primers used to generate these mutants is displayed in **supplemental Table S1**. The mutated DNA sequences were sequenced to ensure that only the appropriate mutations had been incorporated into the amplified DNA.

**Source of Sugars Used**—All soluble polysaccharides and celloligosaccharides were purchased from Megazyme International (Bray, County Wicklow, Ireland), except apple and citrus pectin, konjak galactomannan, and hydroxyethylcellulose, which were obtained from Sigma, and pustullan, which was obtained from Calbiochem. Catalog numbers of polysaccharides where more than one version exists are: wheat arabinoylan, P-WAXYM; rye arabinoylan, P-RAXY; galactomannan, Carob (P-GALML); galactomannan, Guar (P-GGMMV).

**Affinity Gel Electrophoresis**—Affinity gel electrophoresis was used to screen CBM65A and CBM65B for binding to soluble polysaccharides. The method used was essentially that described by Ref. 17, using the polysaccharide ligands at a concentration of 0.3% (w/v), unless stated otherwise. Electrophoresis was carried out for 4 h at room temperature in native 10% (w/v) polyacrylamide gels. The nonbinding negative control protein was BSA.

**Isothermal Titration Calorimetry (ITC)**—The thermodynamic parameters of the binding of the CBM65s to soluble
polysaccharides and cellooligosaccharides were determined by ITC using a VP-ITC calorimeter (MicroCal, Northampton, MA), as described by Ref. 17. Briefly, titrations were performed at 25 °C by injecting 10-μl aliquots of 5–20 mM oligosaccharide or 10 mg/ml of polysaccharide, in 50 mM Na-HEPES buffer, pH 7.5, into the cell containing 100 μM CBM diazoyed into the Na-HEPES buffer, and the release of heat was recorded. The stochiometry of binding (n), the association constant Kₐ, and the binding enthalpy ΔH were evaluated by using MicroCal Origin 7.0 software. The standard Gibbs energy change ΔG⁰ and the standard entropy change ΔS⁰ were calculated from ΔG⁰ = −RT lnKₐ and ΔG° = ΔH° − TΔS°, where R is the gas constant and T the absolute temperature. The polysaccharide at 10 mg/ml was converted into a molarity that gave a stochiometry of 1 to determine the molar concentration of CBM65 binding sites on the polymer.

**RESULTS**

**Quantitative Evaluation of the Binding of CBM65A to Its Ligands**—The endoglucanase from *E. cellulosolvens*, EcCel5A, consists of two GH5 modules and two CBMs designated hereafter as CBM65A and CBM65B, which are located at the N terminus and between the two catalytic modules, respectively (18) (Fig. 1). The two CBMs display 73% sequence identity. Previous qualitative studies showed that both CBM65A and CBM65B bound to acid swollen cellulose, lichenan (β1,3-β1,4 mixed linked glucan), but did not bind to laminarin (β1,3-glucan), Avicel or α-glucans (18). Here we have explored further the specificities of the two protein modules. Recombinant forms of CBM65A and CBM65B, comprising residues 37–170 and 581–713, respectively, of full-length *Ec* Cel5A were purified to electrophoretic homogeneity by immobilized metal-ion affinity chromatography. Initially affinity gel electrophoresis was used to screen potential polysaccharide ligands of the two proteins. The data, presented in Table 2, with example gels displayed in Fig. 2A, show that both protein modules, in addition to binding β1,3-β1,4 mixed linked glucans, also bound to highly decorated β1,4-glucans such as xyloglucan and hydroxyethylcellulose, displayed weak affinity for gluco- mannan, but did not exhibit significant binding to other β1,4-glycans such as xylans, galactomannans, or galactans; no binding to pectin backbone structures or α-glycans were observed. The specificity of the two CBMs appeared to be identical. These data indicate that the CBM65s target β-glucans containing β1,4-linkages.

To provide a quantitative assessment of glucan recognition, the thermodynamic parameters of ligand binding were determined by ITC. Example titrations are shown in Fig. 3, and the full data set is displayed in Tables 3 and 4. The CBM65s displayed the highest affinity for xyloglucan, with a Kₐ of ~10⁵ M⁻¹, whereas binding to barley β-glucan, a β1,3-β1,4 mixed linked glucan, and hydroxyethylcellulose was ~10-fold weaker. With respect to oligosaccharides, the CBM65A displayed highest affinity for XXXG (X comprises glucose decorated at O6 with xylose and G corresponds to undecorated glucose), the repeating unit of xyloglucan, with a Kₐ of 5.6 × 10⁹ M⁻¹, and bound with a similar affinity to cellohexaose (Kₐ 3.6 × 10⁹ M⁻¹). Although binding to cellopentaose had an estimated Kₐ value of 1.2 × 10³ M⁻¹, no quantifiable binding to celloctetraose or smaller celluloooligosaccharides were observed. Similar to the binding of CBMs to soluble ligands (11, 33, 34) (see Ref. 6 for review), the interaction of the CBM65s with their target polysaccharides and oligosaccharides was driven by enthalpic changes, whereas the decrease in entropy had a negative impact on affinity. The stoichiometry of binding, assuming a single

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**Xyloglucan Recognition by Noncatalytic Proteins**
XYLOGLUCAN RECOGNITION BY NONCATALYTIC PROTEINS

TABLE 1

| Dataset         | CBM65A       | CBM65B       | CBM65B-XXXG |
|-----------------|--------------|--------------|-------------|
| Source          | Soleil–Proxima 1 | Diamond–I04  |             |
| Detector        | Quantum 315 CCD | Quantum 315 CCD |             |
| Wavelength (Å)  | 0.9795       | 0.9795       | 0.9795      |
| Space group     | P6,22        | P6,3         | P4,2,2      |

Unit cell parameters

- a = b (Å) 48.74 83.75 57.92
- c (Å) 193.70 36.75 116.74
- α, β, γ (°) 90.90, 120 90.90, 90 90.90, 90
- Resolution limits (Å) 38.74–1.25 (1.25–1.25) 38.75–1.42 (1.42–1.42) 58.37–2.35 (2.35–2.35)
- No. of unique observations 35,557 27,755 8,856
- Multiplicity 21.6 (6.2) 5.3 (5.4) 6.2 (4.7)
- Completeness (%) 91.2 (62.4) 99.8 (99.9) 100.0 (100.0)
- Rmerge (%) 15.80 16.98 22.44
- Rwork (%) 17.30 19.73 28.03
- No. non-H atoms 1097 1060 955
- No. protein atoms 1097 1060 955
- No. water molecules 130 166 1
- No. other 34 6 NA
- No. ligand atoms NA NA 72
- Root mean square deviations from ideal values (Å) Bond length 0.031 0.025 0.012 Angle distance 2.873 2.495 1.56
- Average B factor (Å²) Protein 11.5 21.6 50.4 Water 28.5 36.9 39.5 Other 31.4 62.3 NA
- Ramachandran plot, residues in allowed and most favoured regions (%) 100 99.2 100
- PDB accession code 4afm 4ba6 2ypj

*Values in parentheses are for the high resolution shell.

The binding site for each CBM protomer, indicated that, at saturation, each protein molecule occupied ~11 sugar residues arrayed in tandem in the backbone of the various polysaccharides. These data indicate that the two CBM65s bind to the internal regions of β-glucans. The specificity of the CBMs for, predominantly, β1,4- and β1,3-β,1,4 mixed linked glucans is entirely consistent with the activity of the parent enzyme, EcCel5A, which displays much higher activity against lichenan and carboxymethylcellulose than oat spellet xylan (18).

Structure of CBM65A—The crystal structure of the apo form of CBM65A was solved previously using the selenomethionine-SAD method (28), whereas the structure of apo-CBM65B was determined by molecular replacement to 1.42-Å resolution, using CBM65A as the search model. The two CBMs adopt a β-sandwich fold in which the β-sheets, comprising the convex (β-sheet 1) and concave surface (β-sheet 2) of the protein, contain five and four anti-parallel β-strands, respectively. The order of the β-strands in β-sheet 1 and β-sheet 2 are β1, β9, β3, β7, β6, and β2, β8, β4, β5, respectively. The β-strands are connected primarily by loops, although there is a small helix extending from Lys125 to Tyr132 in CBM65A and Lys668 to Tyr675 in CBM65B (Fig. 4A). The structures of CBM65B and CBM65A are very similar with an root mean square deviation over the 127 α-carbons of only 0.1 Å. In the majority of CBMs that adopt a β-sandwich fold the structure of these proteins is stabilized by a calcium bound to the loops connecting β3 and β4 (35), however, this conserved metal ion site is absent in both CBM65s.

The Ligand Binding Site in CBM65—The ligand binding sites in CBMs that display a β-sandwich fold are, typically, located in the concave surface presented by one of the β-sheets, or at the end of the elliptical protein, within the loops connecting the two β-sheets (see Ref. 6 for review). Inspection of the concave surface of CBM65A and CBM65B reveals a cleft-like structure 20–25 Å long, rich in tryptophan residues. Substituting these aromatic residues with alanine caused a substantial reduction in ligand binding (described in detail below) (Table 3 and Fig. 2B), confirming that the concave surface presented by β-sheet 2 comprises the β-glucan binding site in CBM65A and, by inference, CBM65B.

To explore the mechanism of ligand recognition both CBM65A and CBM65B were co-crystallized with a variety of oligosaccharides. Clear electron density corresponding to XXXG was evident when CBM65B was crystallized in the presence of the xyloglucan-derived oligosaccharide. Despite extensive screening, no crystals of either CBM bound to celulooligosaccharides, or CBM65A in complex with XXXG,

![Schematic of EcCel5A](image)
were obtained. The structure of the CBM65B-XXXG complex, at a resolution of 2.35 Å, shows that the backbone of the ligand, comprising /H9251,1,4-cellotetraose, makes extensive hydrophobic contacts with the four tryptophans that line the cleft: Glc-1 (reducing end unsubstituted glucose) makes parallel apolar contacts with Trp646, Glc-2 and Glc-3 make extensive hydrophobic interactions with Trp651, whereas Trp602 interacts with Glc-4 through parallel hydrophobic contacts. Perpendicular apolar contacts between Trp607 and Glc-3 completes the interactions between the aromatic residues and the glucan tetrasaccharide backbone. Hydrophobic interactions between the tryptophan residues assist in fixing the orientation of the aromatic residues that bind to the glucan ligand. The topology of the tryptophans imposes a twisted conformation on the cellotetraose between Glc-2 and Glc-4, whereas Glc-2 and Glc-1 are orientated 180° with respect to each other. The only hydrogen bond between the tetrasaccharide backbone of XXXG and CBM65B is between O2 and O3 of Glc-4 with O/H9280 and N/H9280 of Gln653 (Fig. 4B).

FIGURE 2. Examples of affinity gel electrophoresis of CBM65A and CBM65B against soluble polysaccharides. Panel A, the two CBM65 proteins were electrophoresed on nondenaturing polyacrylamide gels containing no ligand (control) or 0.3 mg/ml of the target polysaccharide (HEC, hydroxyethylcellulose). BSA was used as a nonpolysaccharide binding control. Panel B, wild type (WT) and mutants of CBM65A were electrophoresed in the presence or absence of the stated polysaccharides.

TABLE 2

Affinity gel electrophoresis of CBM65A and CBM65B

| Polysaccharide (0.3%) | CBM65A  | CBM65B |
|-----------------------|---------|---------|
| Celluloses            |         |         |
| HEC                   | + + +   | + + +   |
| Lichenan              | + + +   | + + +   |
| Curdlan               | -       | -       |
| CMC                   | + +     | + +     |
| Xylans                |         |         |
| Arabinobioxylan (rye) | -       | -       |
| 4-O-Methyl-1-3-Glucurono-1,6-xylan | - | - |
| Xylan (birchwood)     | -       | -       |
| Arabinobioxylan (wheat medium viscosity) | + | + |
| Arabinobioxylan (wheat; insoluble) | - | - |
| Other hemicelluloses  |         |         |
| β-Glucan (barley)     | + + +   | + + +   |
| Xyloglucan (Tamarind) | + + +   | + + +   |
| Mannan (ivy nut)      | -       | -       |
| Galactomannan (locust bean) | + | + |
| Galactomannan (guar gum) | - | - |
| Galactomannan (carob) | -       | -       |
| Arabinogalactan (larchwood) | - | - |
| Galactan (lupin)      | +       | +       |
| Arabinan (sugar beet) | -       | -       |
| Konjac glucosaminan   | + + +   | + + +   |
| Pectins               |         |         |
| Rhamnogalacturonan I (soybean) | - | - |
| Rhamnogalacturonan I (potato) | - | - |
| Pectic galactan (lupin) | +       | -       |
| Pectic galactan (potato) | -       | +       |
| Polygalacturonic Acid (citrus) | - | - |
| Pectin (apple)        | -       | -       |
| Pectin (citrus)       | -       | -       |
| Other polysaccharides |         |         |
| Pustulan              | -       | -       |
| Pullulan              | -       | -       |

Symbols represent: +++, tight binding, + +, significant binding, +, marginal binding, -, no binding.

*HEC, hydroxyethylcellulose.
is likely to be very similar in the two proteins, although CBM65A may make an additional polar contact with the glucan backbone (see below).

Site-directed Mutagenesis of CBM65A—Site-directed mutagenesis was used to further investigate ligand recognition by CBM65A. The capacity of the mutants to bind to ligands was assessed by ITC (cellohexaose and polysaccharides) and affinity gel electrophoresis (polysaccharides). Examples of the affinity gels are shown in Fig. 2B, with the full dataset reported in Tables 3 and 4. Alanine substitution of Trp55, Trp108, Trp99, or Trp60 in CBM65A, which are structurally equivalent to Trp602, Trp607, Trp646, and Trp651 in CBM65B, abrogated cellohexaose and \( \beta \)-glucan recognition, confirming the importance of these aromatic residues in binding the \( \beta \)-linked glucan backbone in both CBM65A and CBM65B. With respect to polar contacts, mutating Gln110 in CBM65A (Q110A mutant), equivalent to Gln653 in CBM65B, significantly reduced, but did not abrogate, binding to both cellohexaose and \( \beta \)-glucan. It would appear, therefore, that Gln110 and Gln653 contribute to ligand recognition.

Both CBM65A and CBM65B bind to cellohexaose considerably more tightly than cellulotetraose (see above), and thus it is possible that the CBM65s make more interactions with the hexasaccharide than the tetrasaccharides. Inspection of the binding cleft downstream of the four tryptophan residues failed to identify obvious ligand binding residues, although it is possible that a tyrosine, at the entrance to the binding cleft (Tyr70 in CBM65A and Tyr617 in CBM65B), and a glutamine (Gln67 in CBM65A and Gln659 in CBM65B) are potential candidates. However, as the Q67A and Y70A mutants of CBM65A displayed similar affinities to the wild type protein (Table 3), it is unlikely that Tyr70/Tyr617 or Gln67/Gln659 contribute to cellulose recognition.

Sequence alignment of CBM65A and CBM65B revealed 73% sequence identity and, as described above, XXXG recognition in CBM65B is conserved in CBM65A (Fig. 4C). A potentially biologically significant difference between the proteins is the loop connecting \( \beta \)4 and \( \beta \)5, which is longer in CBM65A (Trp99 to Gln106) than in CBM65B. Inspection of an overlay of the two proteins indicates that Oe1 and Ne2 of Gln106 will make hydrogen bonds with O2 and O3 of Glc-2 in the cellotetraose backbone, whereas the equivalent residue in CBM65B, Asp649, will be too distant from the ligand to make a polar contact. To test
### TABLE 3
Affinity and thermodynamic parameters of the binding of CBM65A and its variants to polysaccharide and oligosaccharide ligands

The molar concentration of a 1% solution of polysaccharide was iteratively adjusted to give a stoichiometry \(~1\). In general each protein covered \(~11\) sugar residues at saturation.

| CBM65A | Ligand          | $K_a$ ($M^{-1}$) | $\Delta G$ (kcal mole$^{-1}$) | $\Delta H$ (kcal mole$^{-1}$) | $T\Delta S$ (kcal mole$^{-1}$) | $n$   |
|--------|-----------------|-----------------|------------------------------|-----------------|-----------------|-------|
| Wild type | β-Glucan        | 1.5 (±0.2) x 10$^4$ | -5.7                          | -11.8 ± 0.1      | -6.1             | 1.01 ± 0.0 |
| Wild type | Xyloglucan      | 1.7 (±0.1) x 10$^3$ | -7.0                          | -18.0 ± 0.3      | -11.0            | 1.03 ± 0.0 |
| Wild type | HEC$^a$         | 1.2 (±0.02) x 10$^4$ | -5.5                          | -11.1 ± 0.2      | -5.6             | 1.02 ± 0.0 |
| Wild type | XXXG            | 5.62 (±0.1) x 10$^4$ | -5.1                          | -9.6 ± 0.4       | -4.5             | 1.03 ± 0.0 |
| Wild type | Cellohexaose    | 2.1 (±0.3) x 10$^3$ | -4.3                          | -11.3 ± 5.5      | -6.9             | 1.0 ± 0.1 |
| T58A   | β-Glucan        | 8.3 (±1.0) x 10$^3$ | -5.3                          | -12.9 ± 0.2      | -7.6             | 1.0 ± 0.1 |
| Q106A  | β-Glucan        | 1.4 (±0.02) x 10$^4$ | -5.6                          | -10.0 ± 0.1      | -4.4             | 1.0 ± 0.0 |
| Q110A  | β-Glucan        | 3.9 (±0.4) x 10$^3$ | -4.9                          | -10.0 ± 0.2      | -5.1             | 1.0 ± 0.1 |
| K146A  | β-Glucan        | 5.3 (±0.2) x 10$^4$ | -5.1                          | -9.9 ± 0.5       | -4.8             | 1.0 ± 0.0 |
| Y70A   | β-Glucan        | 8.3 (±0.1) x 10$^3$ | -5.3                          | -11.6 ± 0.2      | -6.3             | 1.0 ± 0.0 |
| Y114A  | β-Glucan        | 1.1 (±0.2) x 10$^4$ | -5.5                          | -12.8 ± 0.3      | -7.3             | 1.0 ± 0.0 |
| Y142A  | β-Glucan        | 8.9 (±0.04) x 10$^3$ | -5.4                          | -13.0 ± 0.6      | -7.6             | 1.0 ± 0.0 |
| W55A   | β-Glucan        | No binding       |                               |                  |                  |       |
| W60A   | β-Glucan        | No binding       |                               |                  |                  |       |
| W99A   | β-Glucan        | No binding       |                               |                  |                  |       |
| W108A  | β-Glucan        | No binding       |                               |                  |                  |       |
| Q106A  | Xyloglucan      | 3.5 (±0.5) x 10$^5$ | -7.5                          | -26.7 ± 0.7      | -19.2            | 1.0 ± 0.0 |
| Q110A  | Xyloglucan      | 1.3 (±0.2) x 10$^3$ | -6.9                          | -16.6 ± 0.3      | -9.7             | 1.0 ± 0.0 |
| W55A   | Xyloglucan      | 9.6 (±0.3) x 10$^4$ | -6.8                          | -22.7 ± 0.2      | -15.9            | 1.0 ± 0.0 |
| W60A   | Xyloglucan      | 3.1 (±0.06) x 10$^3$ | -4.7                          | -22.8 ± 0.3      | -18.1            | 1.0 ± 0.0 |
| W99A   | Xyloglucan      | 5.0 (±0.9) x 10$^3$ | -5.0                          | -22.7 ± 0.3      | -17.7            | 1.0 ± 0.0 |
| W108   | Xyloglucan      | No binding       |                               |                  |                  |       |
| Q110A  | Cellohexaose    | -6.9 (±0.6) x 10$^2$ | Binding too weak to quantify thermodynamics |
| K146A  | Cellohexaose    | -9.0(±1.2) x 10$^2$ | Binding too weak to quantify  |
| T58A   | Cellohexaose    | 2.5 (±0.4) x 10$^3$ | -4.6                          | -11.4 ± 5.5      | -6.8             | 1.01 ± 0.4 |
| Y70A   | Cellohexaose    | 2.9 (±0.3) x 10$^3$ | -4.7                          | -11.8 ± 3.8      | -7.1             | 1.0 ± 0.2 |
| Y142A  | Cellohexaose    | 1.6 (±0.01) x 10$^3$ | -4.3                          | -14.7 ± 3.0      | -10.4            | 1.01 ± 0.1 |
| W55A   | Cellohexaose    | No binding       |                               |                  |                  |       |
| W60A   | Cellohexaose    | No binding       |                               |                  |                  |       |
| W99A   | Cellohexaose    | No binding       |                               |                  |                  |       |
| W108A  | Cellohexaose    | No binding       |                               |                  |                  |       |

$^a$ HEC: hydroxyethylcellulose.

### TABLE 4
Affinity and thermodynamic parameters of the binding of CBM65B and its variant D69A to polysaccharide and oligosaccharide ligands

The molar concentration of a 1% solution of polysaccharide was iteratively adjusted to give a stoichiometry \(~1\). In general each protein covered \(~11\) sugar residues at saturation.

| CBM65B | Ligand          | $K_a$ ($M^{-1}$) | $\Delta G$ (kcal mole$^{-1}$) | $\Delta H$ (kcal mole$^{-1}$) | $T\Delta S$ (kcal mole$^{-1}$) | $n$   |
|--------|-----------------|-----------------|------------------------------|-----------------|-----------------|-------|
| Wild type | β-Glucan        | 8.2 (±0.2) x 10$^3$ | -5.3                          | -12.1 ± 0.1      | -6.8             | 1.0 ± 0.0 |
| Wild type | Xyloglucan      | 3.3 (±0.1) x 10$^3$ | -7.4                          | -15.1 ± 0.1      | -7.7             | 1.0 ± 0.0 |
| Wild type | Cellohexaose    | 2.3 (±0.2) x 10$^3$ | -4.6                          | -12.8 ± 1.0      | -8.2             | 1 ± 0.3  |
| Wild type | XXXG$^a$        | 1.7 (±0.01) x 10$^3$ | -4.4                          | -10.2 ± 0.0      | -5.8             | 1.0 ± 0.0 |
| Wild type | HEC$^{bc}$      | 1.42 (±0.4) x 10$^4$ | -5.6                          | -7.0 ± 0.1       | -1.4             | 1.0 ± 0.0 |
| D649A  | Cellohexaose    | 1.5 (±0.05) x 10$^3$ | -4.3                          | -8.1 ± 0.2       | -3.8             | 1.0 ± 0.2 |

$^a$ Heptasaccharide derived from xyloglucan in which X is Glc decorated with Xyl, and G is undecorated Glc.

$^{bc}$ HEC, hydroxyethylcellulose.
this hypothesis the specificity of the Q106A mutant of CBM65A and the D649A variant of CBM65B were analyzed. The affinities of the two variants for xyloglucan and barley \(1,4\)-glucan were similar to the corresponding wild type proteins. Although the D649A mutation did not affect the capacity of CBM65B to bind to cellohexaose, the Q106A variant of CBM65A did not display any detectable affinity for cellohexaose, indicating that Glu106 plays a critical role in the recognition of the hexasaccharide, and likely cellulose.

To provide further support for the view that the Q106A mutation has a significant impact on binding to cellulose but not to mixed linked \(1,4\)-1,3-glucans, the capacity of the mutant to bind to transverse sections of \textit{M. x giganteus} stem was explored. The data (Fig. 5A) showed that the wild type protein bound specifically to phloem cell walls, before or after treatment with a lichenase, which specifically degrades mixed linked \(1,4\)-1,3-glucans; the enzyme does not attack \(1,4\)-glucans (36). These data are consistent with the view that CBM65A is able to bind to both cellulose and \(1,4\)-1,3-glucans, polysaccharides, which are particularly abundant in the phloem cell walls of \textit{Miscanthus} stems. Although the Q106A mutant bound to phloem cell walls in untreated \textit{Miscanthus} stems, section treatment with the lichenase abrogated the binding of the CBM65A variant to these cell walls. These data are consistent with the ITC results in showing that the Q106A mutation influences binding to \(1,4\)-glucans, but has no effect on binding to mixed linked \(1,4\)-1,3-glucans. A comparison of the capacity of the wild type and mutant protein to bind to tobacco stem cell walls, which contain no mixed linked \(1,4\)-1,3-glucans (Fig. 5B), and in which xyloglucan had been exposed by a prior treatment with pectate lyase, indicated that the wild type protein bound more strongly than Q106A and showed some differential labeling in relationship to cell walls at intercellular spaces (around which xyloglucans are known to be differentially regulated (23)). Section treatment with a xyloglucan-specific endo-\(1,4\)-\(\beta\)-glucanase resulted in loss of Q106A binding to cell walls, whereas binding of the wild type protein bound to xyloglucan-depleted cell walls was retained.
Structural Similarity of CBM65 to Other Proteins—Interrogation of the UNIPROT database revealed only two proteins, an endoglucanase from *Cellulosilyticum ruminicola* and one from *Clostridium lentocellum*, which displayed limited sequence similarity with the two CBM65s. The putative *C. ruminicola* endoglucanase contains two tandem repeated sequences and *C. lentocellum* a single sequence that displayed ~30% sequence identity with CBM65A, corresponding to a Z-score of ~0.085. Of potential significance is the observation that three of the four CBMs bind specifically to cell walls of the phloem (Panel A) and anatomy of a vascular bundle. In untreated sections both CBMs bind to adhered cell walls and particularly to cell wall regions lining intercellular spaces (*) as indicated by arrows; x = xylem. After lichenase pre-treatment of the section, before immunofluorescence analysis, wild type CBM65A (WT) still binds to the phloem cell walls but Q106A does not. All fluorescence micrographs have equivalent exposure times. Panel B, transverse sections of tobacco stem showing cell walls in the region of the pith parenchyma after pre-treatment withpectate lyase to remove pectic homogalacturonan. WT and Q106A displayed differential binding to parenchyma cell walls with WT binding strongly to cell walls and particularly to cell wall regions lining intercellular spaces (*) as indicated by arrows; exposure time 25 ms. Q106A bound less strongly to cell walls (exposure time 200 ms) and displayed some preferential binding to adhered cell wall regions at the corners of intercellular spaces; xyloglucan is known to be preferentially located in these regions (23). After a section pre-treatment with a xyloglucan-specific xyloglucanase, WT bound evenly to all cell walls with no differential binding in relationship to intercellular spaces, whereas Q106A did not bind (exposure time for both + xyloglucanase micrographs, 600 ms). Scale bars = 100 μm.

**DISCUSSION**

This report describes the structure of CBM65A and CBM65B, the founding members of a new CBM family that targets β-glucans. Similar to other CBMs that target β-glucans, the CBM65s display no significant binding to xylan. Such specificity can be achieved by the targeting of O6 groups (productive binding in the case of cellulose binders and through steric clashes in xylan-specific CBMs) that distinguishes gluco- from xyla-configured ligands (11, 14). In the CBM65B-XXXG complex C6 of all the backbone glucose moieties make extensive hydrophobic interactions with the surface tryptophans, and thus are likely to make a significant contribution to overall affinity. Furthermore, the CBM65s are not optimized to bind xylan, which adopts a 3-fold screw axis conformation (37). Xylan-specific CBMs often contain a pair of tryptophans, oriented at 120° with respect to each other, which bind to xylene residues n and n + 2 in the polysaccharide (38). Disruption of the orientation of these aromatic residues can convert a xylan binding CBM into a cellulose-specific protein (38). In the CBM65s the ligand binding tryptophans are not optimized to bind to a polysaccharide that has a regular 3-fold helical structure. Indeed Glc-1 and Glc-2 are oriented at 180° to each other, and the hydrogen bond between O3 and O6, which would be absent in a xylan chain, plays a critical role in stabilizing the conformation adopted by these two sugars.

CBM65A and CBM65B display higher affinity for oligosaccharides (e.g. cellohexaose), and particularly polysaccharides (xyloglucan and barley β-glucan), than cellotetraose, which fully occupies the core component of the substrate binding cleft. Although it is formally possible that the two additional residues in cellohexaose provide additional contacts with the protein, mutagenesis of residues in the vicinity of the two glucose moieties in CBM65A did not influence affinity. In cello-oligosaccharides with a degree of polymerization > 4 the Glc that interacts with Trp in CBM65A is internal and thus the pyranose ring is fixed. In cellotetraose the equivalent Glc is at the reducing end of the tetrasaccharides, and hence adopts multiple conformations through mutarotation. Thus, the reduction in entropy upon binding the tetrasaccharide may explain the weak affinity. It is evident, however, that polysaccharides, in which the backbone is either mixed linked or a β1,4-linked polymer binds more tightly than cellohexaose, indicating that fixing the conformation of the terminal sugar is not the sole...
reason for the tighter binding of ligands with a degree of polymerization >4. There are examples of ligands that extend outside the CBM binding region, which bind more tightly to the protein than smaller glycans that, nevertheless, fully occupy the sugar binding sites (11, 39, 40). It has been suggested that the longer ligands adopt a more fixed conformation, through extensive intra-chain hydrogen bonds, which is optimized to recognize the target CBM (39). An alternative possibility is that the CBMs physically associate, resulting in increased affinity for multivalent ligands through avidity effects (15, 40, 41). However, size exclusion chromatography indicated that CBM65A is a monomer (data not shown), although ligand-induced oligomerization is possible; a phenomenon, which would not be observed by studying the molecular mass of the apoprotein (42).

The observation that the Q106A mutation destroys binding to cellohexaose and cellulose, but not xyloglucan or β1,3-β1,4 mixed linked glucans, is intriguing. These data suggest that CBM65A may display flexibility in ligand recognition, with its binding site capable of recognizing both linear β1,4-glucans and β1,3-β1,4 mixed linked glucans, and that Glc106 only contributes to cellulose recognition. It is also interesting that CBM65B, despite lacking a functionally equivalent residue to Glc106 displays affinity for cellohexaose, although the mechanism by which the protein module retains this specificity is unclear.

This report provides insights into how a CBM can specifically recognize xyloglucan in preference to other β-glucans. Previously, Najmudin et al. (12) showed that β-glucan binding CBMs can accommodate, but do not display a preference for, xyloglucan. The primary mechanism by which CBM65B binds to the xylose side chains is through apolar interactions with the surface aromatic amino acids. In particular, Tyr685 makes extensive hydrophobic contacts with Xyl-2, although the sugar also makes apolar contacts with Trp607, Trp646, and Trp651. The CBM65s are similar to many CBMs (11, 38, 43) (see Ref. 6 for review), where binding to glycans is dominated by hydrophobic interactions with aromatic residues. In typical β-glucan binding CBMs there are three aromatic residues that stack against the sugar rings, or against both faces of the same pyranose. In the CBM65s, however, the binding cleft contains five aromatic residues. Only Glc-4 aligns perfectly with a tryptophan to maximize planar hydrophobic contacts with a tryptophan (Trp602). The side chains of the other aromatic residues, except Trp602, make no apolar contact with the xylose side chains, and hence its contribution to xyloglucan recognition is considerably less than the other aromatic residues in the ligand binding cleft. Thus, xyloglucan recognition is dominated by aromatic residues that recognize both the glucan backbone and the xylose side chains.

To summarize, this report describes the biochemical properties of two CBMs that are the founding members of CBM65. The protein modules bind to mixed linked β1,4-β1,3-linear and decorated β1,4-glucans, but displays a preference for the decorated β-glucan, xyloglucan. Specificity for decorated glucans is achieved through an extensive hydrophobic platform that contacts both the glucan backbone and the xylose side chains. Significantly, one of few hydrogen bonds between CBM65A and its ligands confers specificity for cellulose. Thus, this article shows that in CBM65, specificity for diverse β1,4-glucans is not achieved through the targeting of conserved features of these glycans, whereas the work also reveals how the orientation of hydrophobic residues can be optimized to recognize backbone and side chain sugars, providing a model for the recognition of decorated polysaccharides.

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