NMR and Fluorescence Spectroscopies Reveal the Preorganized Binding Site in Family 14 Carbohydrate-Binding Module from Human Chitotriosidase

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ABSTRACT: Carbohydrate-binding modules (CBM) play important roles in targeting and increasing the concentration of carbohydrate active enzymes on their substrates. Using NMR to get the solution structure of CBM14, we can gain insight into secondary structure elements and intramolecular interactions with our assigned nuclear overhauser effect peaks. This reveals that two conserved aromatic residues (Phe437 and Phe456) make up the hydrophobic core of the CBM. These residues are also responsible for connecting the two β-sheets together, by being part of β2 and β4, respectively, and together with disulfide bridges, they create CBM14’s characteristic “heven-like” fold. Most CBMs rely on aromatic residues for substrate binding; however, CBM14 contains just a single tryptophan (Trp465) that together with Asn466 enables substrate binding. Interestingly, an alanine mutation of a single residue (Leu454) located behind Trp465 renders the CBM incapable of binding. Fluorescence spectroscopy performed on this mutant reveals a significant blue shift, as well as a minor blue shift for its neighbor Val455. The reduction in steric hindrance causes the tryptophan to be buried into the hydrophobic core of the structure and therefore suggests a preorganized binding site for this CBM. Our results show that both Trp465 and Asn466 are affected when CBM14 interacts with both (GlcNAc)3 and β-chitin, that the binding interactions are weak, and that CBM14 displays a slightly higher affinity toward β-chitin.

INTRODUCTION

Protein–carbohydrate interactions are involved in numerous biological processes, such as cell–cell recognition, fertilization, embryogenesis, and tumor metastasis among others.1 Proteins involved in such interactions often have noncatalytic modules called carbohydrate-binding modules (CBMs). CBMs are subdivided into families according to their amino acid sequence similarity. They are currently classified into seven “fold families”, which are further divided into three types. Type A binds to crystalline surfaces, B to glycan chains, and C to linear and water-insoluble polymer that is an abundant component in the cell walls of fungi and the exoskeleton of crustaceans (i.e., crab, shrimp, and insects).4 Chitin is not produced by mammals, making the polymer a likely candidate to trigger an innate immune response.3 Lectins and other defense proteins are known to play an important role in initiating this immune response. Carbohydrate active enzymes contain many members, among them are chitinases that catalyze the hydrolysis of chitin. Chitinases are synthesized by both animals and plants that are not able to produce chitin, which underlines their importance in their immune defense and thus their ability to survive.6

In humans, two active chitinases have been reported: chitotriosidase-1 (CHIT1) and acidic mammalian chitinase. In addition, a chitobiase is synthesized that catalyzes the hydrolysis of a soluble GlcNAc dimer.7,8 CHIT1 is synthesized as a 50 kDa soluble protein able to hydrolyze chitin.9 This human chitinase is classified as belonging to family 18 of glycosyl hydrolases (GH18) in the Carbohydrate-Active EnZymes (CAZy) database (http://www.cazy.org).10 It consists of two domains: a catalytic domain that adopts the conserved (α/β)8 TIM barrel form typical for GH18 and a noncatalytic domain in the N-terminal belonging to a CBM in family 14 (CBM14), which many of its members are known to
bind chitin according to the CAZy database. These two domains are separated by a proline-rich and extended linker (PT-linker) that comprises 31 residues. The flexible nature of this linker could suggest that CBM14 works as a guiding system for CHIT1.11 CBM family 14 has a lectin-like property of binding small sugars (mono-, di-, and trisaccharides) and is characterized by its highly conserved “hevein-like” fold.3 The structure has three disulfide bonds that are important for its structural integrity as found by site-directed mutagenesis of either of the cysteins involved. It has also been shown that removal or mutagenesis of its only tryptophan has a detrimental effect on the CBMs ability to bind chitin.12,13 Moreover, CHIT1 with its CBM degrades chitin faster and much more efficiently than its isoform without the CBM.14

According to CAZy, there are currently seven structures of CBM14s. Only two of the structures have been solved using NMR and correspond to tachycitin (PDB ID: 1DQC)15 and dust mite allergen Blo t 12 (PDB ID: 2MFK). The remaining structures have been solved using X-ray crystallography, among which are CHIT1 (PDB ID: 5HBF) that includes the CBM14 elucidated by NMR here, Der p 23 (PDB ID: 4ZCE), a dust mite allergen from Dermatophagoides pteronyssinus,16 and Pf Avr4 (PDB ID: 4Z4A), an effector protein from the tomato pathogen Pseudocercospora fuligena.17 A structural homolog of PfAvr4 and CfAvr4, an effector protein from the tomato pathogen Cladosporium fulvum, has recently18 been co-crystalized with a chitin hexa oligosaccharide (GlcNAc)₆ (PDB ID: 6BN0). This is the first co-crystal structure of a CBM14 with an oligosaccharide. CfAvr4 has an extended binding surface that can accommodate longer polysaccharide chains and the interaction with the substrate is through aromatic residues and hydrogen bonds, which is more reminiscent of a type B CBM than the type C common for CBM14s.3 This protein contains an extended loop that hydrogen-bonds with the oligosaccharide, as well as linking an additional N-terminal α-helix to the first β-sheet. It also contains two residues (Gln69 and Lys84) that through mutation studies are shown to have a great effect on the binding. These aspects are not conserved in other CBM14s and can therefore explain the observed difference in binding mode.

We have used NMR spectroscopy to determine the solution structure of CBM14 from CHIT1. As a result, we have developed further insight into the structural elements important in keeping CBM14s binding affinity for chitin. This CBM shows a binding affinity toward both (GlcNAc)₃ and β-chitin, and through our studies, we have found that the same residues (Trp465 and Asn466) are involved in binding both substrates. However, a single mutation of Leu454 to alanine renders the CBM unable to bind. Leu454 does not show a significant change in chemical shift perturbation when CBM14 (wild type, wt) interacts with either substrates. Interestingly, with fluorescence spectroscopy, we observe that Leu454Ala displays a significant blue shift, meaning that the position of Leu454 in the protein structure is significant for the correct orientation of Trp465 and therefore the protein’s ability to bind substrates.

## RESULTS

**Solution Structure of CBM14.** The solution structure of CBM14 (PDB ID: 6SO0) was determined by NMR spectroscopy. Assignment of the backbone and side-chain chemical shifts for CBM14 are essentially complete (N, N, C, H, and C > 92%; H and C side chains > 76%) (Table S2). The mature protein contains the following amino acids for purification purposes: three extra amino acids (Gly-Ser-Pro) at the N-terminus and five extra amino acids (Gly-Leu-Val-Pro-Arg) at the C-terminus. Gly-Ser-Pro-Arg in the N-terminus has not been assigned. Due to their location in a flexible-loop region, Cys460 and Cys462 are not observed by line
broadening. Chemical shift data have been deposited at the Biological Magnetic Resonance Data Bank (BMRB) under the accession number 27277. The restraints and structural statistics for the 20 best conformers of CBM14 are given in Table S1.

The nuclear overhauser effect spectroscopy (NOESY) experiments show that the CBM14 fold comprises two β-sheets, three antiparallel β-strands in the N-terminal region (β1, β2, β3; residues 426–430, 436–442, and 444–449), and two antiparallel β-strands in the C-terminal region (β4, β5; residues 455–456 and 463–465). This fold pattern matches the previously described distorted β-sandwich fold in the crystal structure of the same protein,11 and is known as the “hevein-fold”. However, the two structures differ in the amount of amino acids involved in each β-strand (Table S2).

By aligning the NMR structure with the crystal structure (Figure 1), we obtain a root-mean-square deviation (RMSD) of 1.59 Å over the secondary structure (3.78 Å over 48 CA residues). The highest deviations are observed in the loop regions of the CBM.

Connecting β2 and β3 is a type I turn (residues 442–444). The protein contains six cysteins that form three disulfide bridges (Cys420-Cys440, Cys450-463, and Cys460-462) that have a stabilizing effect on the β-sandwich, also described by Fadel et al.11 Through the NOESY experiments, we observed NOEs between two aromatic residues Phe437 and Phe456 involved in β2 and β4, respectively. These residues create a hydrophobic core holding the two β-sheets together. This core is probably essential for the β-sandwich fold given the conservation of aromatic residues in this position in other CBM14s.

Interaction with (GlcNAc)₃. The binding properties of CBM14 and (GlcNAc)₃ were investigated using isothermal titration calorimetry (ITC). The shape of the ITC binding curve is described by the so-called Wiseman c value,19 which can be expressed as follows

$$c = nK[M]$$

where n is the stoichiometry of the reaction, Kₐ is the equilibrium binding association constant, and [M]ₐ is the protein concentration. It is well established that c values within the range of 10 < c < 1000 are a prerequisite for meaningful calculations of Kₐ.19 A Kₐ of 9.9 mM (Kₐ = 100 M⁻¹) has previously been observed by Crasson et al.13 With a protein concentration of 45 μM, typical for an ITC experiment, it is suggested that the c value would be close to 0.05. It has been shown that binding thermodynamics can be obtained even if c is in the range of 0.01 < c < 10 if a sufficient portion of the binding isotherm is used for analysis.20 This is achieved by ensuring a high molar ratio of ligand versus protein at the end of the titration, accurate knowledge of the concentrations of both ligand and receptor, an adequate level of signal-to-noise ratio in the data, and known stoichiometry. Using this approach, the fit of theoretical data to the experimental data (Figure S2) for four independent measurements yielded a Kₐ of 3.1 ± 0.2 mM and a ΔHₐ of −1.6 ± 0.2 kcal/mol. Moreover, a ΔGₐ of −3.3 ± 0.1 kcal/mol and a ΔSₐ of 6 ± 1 cal/kmol (−TΔSₐ = −1.7 ± 0.2 kcal/mol) can be calculated from eq 2. The obtained Kₐ is close to that previously observed.

$$ΔGₐ = ΔHₐ - TΔSₐ = -RT \ln Kₐ = RT \ln K_d$$

Interaction Studies by NMR. To investigate whether binding of a different substrate, β-chitin, behaved in the same way as (GlcNAc)₃, we performed ¹⁵N-heteronuclear single-
quantum correlation (HSQC) experiments on CBM14 alone and CBM14 together with \( \beta \)-chitin. This allowed us to investigate the chemical shift perturbation to probe the residues involved by comparing the \( ^{15}\text{N}-\text{HSQC} \) spectra of the protein before and after addition of substrate (Figure S3).

The \( ^{15}\text{N}-\text{HSQC} \) spectrum shows the cross-peaks of the amide nitrogens and amide hydrogens and is often referred to as the protein’s “fingerprint” spectrum. Upon substrate binding, the environment around the nuclei that interact changes and, as a result, the corresponding cross-peak in \( ^{15}\text{N}-\text{HSQC} \) will change. The residues with the highest changes in chemical shift perturbation often belong to the binding site. The biggest differences observed in chemical shifts belong to the side chain of Trp465 (H\( \varepsilon_1/N\varepsilon_1 \)) and Asn466 (H\( \delta_1/N\delta_1 \) and H\( \delta_2/N\delta_2 \)), corresponding to the interaction surface of the protein. Both have previously been described as most perturbed upon titration with (GlcNAc)\(_3\). An overlay of \( ^{15}\text{N}-\text{HSQC} \) for the side chain of Trp465 and Asn466 for CBM14 interacting with (GlcNAc)\(_3\) and \( \beta \)-chitin is given by Figure 2 (the diagram comparing all residues is given in Figure S3). The chemical shift perturbation for CBM14 interacting with 6 mg \( \beta \)-chitin is not included as the signals in \( ^{15}\text{N}-\text{HSQC} \) were broadened beyond detection. As the chemical shift perturbation is more pronounced for \( \beta \)-chitin, it could indicate a stronger binding affinity compared to the soluble (GlcNAc)\(_3\).

Another residue that shows a chemical shift perturbation is Thr464. Mutational studies have been done on this threonine and showed medium impact on binding activity for Thr464, which could indicate that this residue could be one of the key polar residues that are directly involved in binding. Furthermore, Cys450 and Cys463 also show a minor change in chemical shift. These cysteins create a disulfide bridge linking the two \( \beta \)-sheets together.

To allow us to obtain detailed information about the bound sugar, we performed T\(_{1\text{rho}}\) NMR spectroscopy on the CBM14–(GlcNAc)\(_3\) system using different protein:ligand concentrations. The resulting one-dimensional (1D) \( ^{1}\text{H} \) NMR spectra are given in Figure S4. Compared to the reference spectrum, the T\(_{1\text{rho}}\) experiment will display a reduction in signal intensity if the protein and substrate are bound because the protein’s relaxation will be observed on the substrate. We observe indications of substrate binding by getting a reduction in signal intensity for the methyl groups, H1, H2–H6 for (GlcNAc)\(_3\). It is evident that CBM14 interacts with (GlcNAc)\(_3\) and the nonreducing end of the trimer seems to be preferred (Figure S4). By knowing the binding site (Trp465 and Asn466) in CBM14, we were able to use a docking algorithm (HADDOCK, high-ambiguity driven docking). This showed that the lowest-energy cluster of CBM14 and (GlcNAc)\(_3\) coincides with the affected chemical shifts in the T\(_{1\text{rho}}\) experiments (Figure S4). We also tried to probe the interaction using a water LOGSY\(_{21,22}\) experiment, but due to the H\( \beta \) of the reducing end of (GlcNAc)\(_3\) being too close to the frequency for the saturation of the water signal, the spectrum was inconclusive. To support the indicated interactions in the T\(_{1\text{rho}}\) experiment, we also did a saturation transfer difference (STD)\(_{23,24}\) experiments varying the protein:ligand concentrations, temperature, saturation times, and on-resonances. However, no detailed information could be extracted from these spectra.

**Leu454Ala Mutant.** Looking from a structural point of view, this leucine is positioned close to the binding site (Trp465 and Asn466) by \( \beta \)-sheet 2 (\( \beta \)4 and \( \beta \)5). According to a multiple sequence alignment done using Clustal Omega (1.2.4) given by Figure 3, we observe that this particular leucine is conserved in tachycitin, PfAvr4 and CfAvr4. These

![Figure 3](https://www.rcsb.org/)
proteins also have an aromatic residue (tryptophan or tyrosine) in the same position as CBM14’s Trp465.

To obtain insight into the differences between CBM14 wild type and CBM14 Leu454Ala mutant, we produced a 15N-labeled mutant and recorded a 15N-HSQC spectrum. The differences in chemical shift compared to the wild type are shown in Figure 4. With a threshold of $\Delta \delta 0.2$ ppm, five residues showed a significant change in chemical shift. Divided into two groups: $\Delta \delta 0.2$–0.4 ppm (Thr452, side chains of Trp465 and Asn466 and $\Delta \delta 0.4$–0.8 ppm (green): Val455 and Thr464.

Fluorescence Spectroscopy. Taking advantage of the fact that CBM14 only has one tryptophan (Trp465), we monitored the structural impact of alanine substitutions on key binding residues using fluorescence spectroscopy. The maximum emission wavelength ($\lambda_{\text{max}}$) provides a measure of the protein’s water exposure. The resulting $\lambda_{\text{max}}$ for native CBM14 wild type (wt), denatured (6 M guanidinium chloride), and its mutants are given in Figure 5.

The $\lambda_{\text{max}}$ of the native CBM14 is 353.21 nm, which is explained by the high exposure of Trp465 to the aqueous solvent. Most of the mutants have a similar $\lambda_{\text{max}}$ compared to the native form of CBM14, indicating that an alanine substitution does not modify significantly the direct environment of the Trp465.

A significant blue shift is observed for CBM14-Leu454Ala and to a lesser extent for CBM14-Val455Ala. Both residues are located behind Trp465. A slight blue shift is also observed for CBM14-Pro451A, which can be explained by its position behind the Leu454 and/or its implication in the loop formation (Pro451, Thr452, Gly453, and Leu454). A slight
red shift is observed for CBM14-Thr464Ala and CBM14-Asn466Ala. This suggests that Trp465 is more exposed to the aqueous solvent in the mutant compared to the wild type due to the smaller side chain of alanine. The highest $\lambda_{\text{max}}$ (354.01 nm) is obtained for the partially denatured sample of CBM14 in 6 M guanidinium chloride.

### DISCUSSION

Here, we have focused on the solution structure of CBM14 solved by NMR and how this protein interacts with (GlcNAc)$_3$ and $\beta$-chitin. This has led to information about structural elements of this CBM that are vital for its binding affinity. Of particular interest is Leu454, which is important for orienting the main binding residue Trp465 correctly. Both residues, as well as the two aromatic residues creating the hydrophobic core are conserved in other CBM14s, indicating that these residues create a preorganized binding site for the substrate.

Use of NOEY experiments enables observation of a cross-peak between two hydrogen atoms if they are separated by a distance up to 5 Å. This gives rise to a direct determination of the secondary structure of the protein and therefore differences in the residues involved in the $\beta$-strands for the X-ray and NMR structures. Together with the three disulphide bridges, the $\beta$-sheets maintain the functional conformation of the CBM. Furthermore, the NOEY experiments show NOEs between Phe437 and Phe456. Phe437 is conserved in Der p 23 and tachycitin, but is replaced by a tyrosine in CfAvr4, PfAvr4, and Bt t 12, whereas Phe456 is conserved in Bt t 12, replaced by tyrosine in tachycitin and tryptophan in the three other proteins. These hydrophobic residues that are part of $\beta$2 and $\beta$4 are commonly found at these positions (except for Bt t 12 where they are part of $\beta$4 and $\beta$6) and presumably help linking the $\beta$-sheets together. This suggests that the two aromatic residues are creating a hydrophobic core that helps stabilize the structure. Altogether, these aspects give rise to the hevein-like fold characteristic for CBM14.

Another trait in CBM14, as with most other CBMs, is that it relies on aromatic residues (often tryptophans) for substrate binding.

This CBM14 contains a single tryptophan (Trp465) located on the protein surface in the C-terminus and makes up the binding site for chitin together with Asn466. This tryptophan plays a second role for this CBM by being part of the secondary structure of the protein and therefore participating in hydrogen bonding with the polar groups of the crystalline substrate. By comparing the $\lambda_{\text{max}}$ of its ability to bind $\beta$-chitin, however, the ability to bind crystalline substrate is a feature that is characteristic of type A CBMs that exhibit a platform-like binding site able to accommodate a crystalline substrate. Interestingly, a comparison of the substrates (GlcNAc)$_3$ and $\beta$-chitin shows that the same residues in CBM14 are affected upon substrate binding.

The importance of the polar residues below Trp465, which participate in hydrogen bonding with the polar groups of the polysaccharide, for substrate interaction is supported by the observed change in chemical shift for Thr464. We also observe a minor change in chemical shift for Cys450 and Cys463 (disulphide bridge). Even though these amino acids are not located directly below the binding site, the disulphide bridge they create is pivotal for keeping the structural integrity of the protein intact for binding.

Studies of CBM14’s ability to bind chito-oligosaccharides have shown that CBM14 have the best affinity for (GlcNAc)$_3$ ($K_d = 9.9 \pm 0.8 \text{ mM}$). Our investigation using ITC demonstrated a dissociation constant in the same range with a measured $K_d$ of 3.1 ± 0.2 mM. These $K_d$’s indicate a weak binding affinity to (GlcNAc)$_3$. By comparing the $^{13}$N-HSQC of the side chains of Trp465 and Asn466 of CBM14 when it interacts with (GlcNAc)$_3$ or $\beta$-chitin (Figure 2), we observe an increase in chemical shift perturbation for the latter. This indicates that CBM14 has a stronger binding affinity toward crystalline substrate compared to the previously observed soluble oligosaccharides.

The relatively weak (GlcNAc)$_3$ binding to CBM14 enabled investigation using $T_{\text{1rho}}$ experiments. In Figure S4, we observe a decrease in signal intensity for the methyl groups, H1, and H2–H6 for (GlcNAc)$_3$ in the $T_{\text{1rho}}$ experiment, all indicating binding. Ideally, the $T_{\text{1rho}}$ spectra could indicate a difference in the trimer’s orientation upon binding with CBM14 by giving differences in intensity for the sugar signals. From the experiments shown in Figure S4, we see that the nonreducing end seems to be preferred. This was further tested by using the restraints from the chemical shift perturbation, where Trp465 and Asn466 compose the binding site of CBM14 and model of the trimer as input in a HADDOCK algorithm. The best model generated supports the observations from the $T_{\text{1rho}}$ experiments and suggests a CH–π interaction between the second sugar ring and the side chain of Trp465 (Figure S5).

Even though weak interactions are favored in saturation transfer-based experiments like STD-NMR, detailed information about the (GlcNAc)$_3$ interaction with CBM14 could not be acquired due to spillover from the saturation of the methyl...
region of the protein to the methyl groups of \((\text{GlcNAc})_3\). We have also tried to quantify the binding kinetics \((\text{on/off rate})\) using stopped-flow spectrophotometry, but the reaction was too fast for obtaining kinetic details.

It has previously been suggested\(^{11}\) that this CBM14 functions as a guiding system for its catalytic domain. Having a CBM help concentrate enzymes onto their substrate is a trait that can be found in other enzymes—CBM systems.\(^{27-30}\) This can explain the weak binding affinity showed by this CBM as well as its fast on/off rate. A relatively weak binding and an innate fast on/off rate would be beneficial for CBM action. CBMs cannot disrupt the crystalline surface and by this alter the binding interactions resulting in a decreased binding affinity as observed for, e.g., GHs. Here, productive binding yielding hydrolysis reduces the number of binding interactions and greatly weakens the affinity.\(^{31-33}\)

**MATERIALS AND METHODS**

**Protein Production and Purification.**\(^{13}\)C, \(^{15}\)N, and \(^{15}\)N CBM14 samples were expressed in *Escherichia coli* BL21 (DE3) cells. Precultures were grown in Lysogeny broth (LB) medium \((10 \text{ g/L tryptone}, 5 \text{ g/L yeast extract}, \text{and } 5 \text{ g/L NaCl})\) supplemented with 50 μL of kanamycin \((50 \text{ mg/mL})\) in a shaking incubator at 30 °C, 225 rpm overnight. Four flasks with 500 mL of LB media and 500 μL of kanamycin \((50 \text{ mg/mL})\) in each were inoculated with 1% \((v/v)\) of the overnight culture and grown in a shaking incubator at 30 °C, 225 rpm to \(O_D_{200} \approx 0.9\) before cooled on ice for 10 min. The cultures were centrifuged \((\text{Sorvall})\) at 4 °C, 6,150g, 10 min, and the cells were resuspended in 450 mL of MQ media \((6 \text{ g/L Na}_2\text{HPO}_4, 3 \text{ g/L KH}_2\text{PO}_4, 0.5 \text{ g/L NaCl})\) in a shaking incubator at 30 °C, 225 rpm for 2 h before adding cOmplete ULTRA protease inhibitor \((\text{Roche}).\)

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An AKTA FPLC instrument equipped with a size-exclusion chromatography (SEC) column \((\text{HiLoad Superdex } 75 \text{ pg})\) was equilibrated with SEC-buffer at pH 7.5 \((20 \text{ mM TRIS}, 50 \text{ mM NaCl})\) for 20 h. The flow rate was 1 mL/min. Fractions containing CBM14 were concentrated and buffer-exchanged into the NMR buffer \((50 \text{ mM sodium phosphate, pH } 5.5 \text{ or pH } 7.0)\) for interaction studies. Samples for NMR were made with CBM14 in NMR buffer with \(D_2O\) added to a final ratio of \(90\% \text{H}_2\text{O}/10\% \text{D}_2\text{O}\) by centrifugation using Vivaspin 6 protein spin concentrators \((\text{MWCO } 3 \text{ kDa, Sartorius})\) at 10 °C, 7,000g.

**Generation of Mutants of CBM14.** Alanine mutants of CBM14 were generated as previously described by Crasson and co-workers.\(^{35}\) The \(^{15}\)N CBM14 Leu454Ala mutant was produced and purified as described above.

**NMR Spectroscopy.** All CBM14 NMR samples were prepared in 50 mM sodium phosphate buffer and 10% \(D_2O\), pH 5.5 or pH 7.0 \((\text{interaction studies}).\)

The spectra were recorded at 25 °C on a Bruker Ascend 800 MHz spectrometer Avance III HD (Bruker Biospin) equipped with a 5 mm Z-gradient CP-TCI \((H/C/N)\) cryoprobe at the NV-NMR-Centre/Norwegian NMR Platform at NTNU Norwegian University of Science and Technology, Trondheim, Norway. \(^1H\) shifts were referenced internally to the water signal, while \(^13\)C and \(^15\)N chemical shifts were referenced indirectly to HDO based on the absolute frequency ratios.\(^{34}\) Backbone and side-chain assignments of CBM14 were achieved using \(^15\)N-HSQC, \(^13\)C-HSQC, HNCA, HN(CA)CO, H(C)CACB, H(C)NO, H(N)CO, H(CA)CO, HNCACB, and \(^15\)N-edited NOESY-HSQC, and \(^13\)C-edited aliphatic NOE-HSQC. The NMR data were recorded and processed with TopSpin version 3.5, and the data were analyzed with CARA version 1.5.\(^{35}\) Backbone torsion angles were predicted using TALOS-N \((https://spinnidd.nihs.nih.gov/bax/software/TALOS-N/)\)\(^{36}\) and chemical shifts of N, H\(^\text{N}\), C\(^\alpha\), C\(^\beta\), H\(^\text{C}^\beta\), and C\(^\gamma\).

**Structure Elucidation.** NOE distance restraints were obtained using \(^{1}\text{H}\)-edited NOE-HSQC and \(^{13}\)C-edited aliphatic NOE-HSQC. Backbone torsion angles predicted by TALOS-N\(^{36}\) were used as constraints for structure calculation, as were the three disulfide bridges Cys420-Cys440, Cys450-Cys463, and Cys460-Cys462. This information enabled structure calculation using CYANA 3.98\(^{37,38}\) by generating 100 structures that were optimized using 10,000 steps of simulated annealing. Using YASARA\(^{39}\) with a YASARA force field,\(^{40}\) energy minimization was done on the 20 conformers with lowest CYANA target function values \((\text{Table S1}).\)

**Substrate Binding.** The interaction between CBM14 and β-chitin \((\text{Mahtani Chitosan, Veraval, Gujarat, India, milled and}}

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sieved to a particle size of ca. 0.5 mm) was investigated using NMR spectroscopy. A $^{15}$N-HSQC spectrum of 90 μM CBM14 in 50 mM sodium phosphate buffer and 10% D$_2$O, pH 7.0, was recorded at 25 °C as reference. New $^{15}$N-HSQC spectra were recorded after addition of β-chitin (1, 2, and 6 mg). The change in chemical shift $\Delta \delta$ (in ppm) was calculated using the following formula: $\Delta \delta = (\Delta \delta_{\text{obs}})^2 + (\Delta \delta_{\text{ref}})^2)^{1/2}$, where $\Delta \delta_{\text{obs}}$ and $\Delta \delta_{\text{ref}}$ are the change in chemical shift for the amide proton and amide nitrogen (ppm), respectively. The interaction between CBM14 and (GlcNAc)$_3$ was done as described by Crasson and co-workers.13

In addition, interaction studies were carried out with the mutant, 90 μM $^{15}$N CBM14 Leu454Ala with two substrates: (GlcNAc)$_3$ (Seikagaku Corporation, Tokyo, Japan) (12 mM) and β-chitin (6 mg).

The interaction between CBM14 and (GlcNAc)$_3$ was studied by saturation transfer difference (STD)-NMR using the following pulse programs: water LOGSY,21,22 STD,23,24 and $T_{1\text{rho}}$ (a spinlock element to measure the $T_{1\text{rho}}$ was incorporated in a normal proton experiment with excitation sculpting for water suppression). The water LOGSY was performed at 25 °C in a 3 mm NMR tube containing 26 μM CBM14 and 400 μM (GlcNAc)$_3$ (protein:ligand ratio, 1:15) in 50 mM sodium phosphate buffer and 10% D$_2$O, pH 7.0. For the STD and $T_{1\text{rho}}$ experiments, four different protein:ligand concentrations (1:100: 3 μM CBM14 and 300 μM (GlcNAc)$_3$; 1:50: 5 μM CBM14 and 260 μM (GlcNAc)$_3$; 3:50: 15 μM CBM14 and 260 μM (GlcNAc)$_3$; and 1:15: 26 μM CBM14 and 400 μM (GlcNAc)$_3$) were tested at 25 °C. The STD experiments were also done at 15 °C and tested at different saturation times (0.40, 0.50, 0.75, 0.80, 1.20, 1.60, 2.00, 2.50, 3.00, 4.00, 5.00 s). The on-resonance frequency was set in the aliphatic region (0.28, 0.37, and 0.82 ppm), and the off-resonance frequency at −30 ppm.

Docking. High-ambiguity driven biomolecular docking (HADDOCK) was used to get further insight into the interaction between CBM14 and (GlcNAc)$_3$. The docking was carried out using the Easy interface of HADDOCK 2.2 (https://haddock.science.uu.nl/services/HADDOCK2.2/haddockserver-easy.html).42 The PDB file containing the NMR structure of CBM14 and a model of (GlcNAc)$_3$ was uploaded after being generated by GLYCAM Carbohydrate builder (glycam.org). Active residues were Trp465 and Asn466, and passive residues were determined automatically by HADDOCK as those within a radius of 6.5 Å.

Isothermal Titration Calorimetry. Isothermal titration calorimetry (ITC) was performed using the VP-ITC system from Microcal, Inc. (Northampton, MA).17 Buffered solutions of 20 mM TRIS and 50 mM NaCl, pH 7.5, were thoroughly degassed prior to experiments to avoid air bubbles in the calorimeter. An enzyme concentration of 45 μM was used in the reaction cell with a volume of 1.42 mL and 12 mM (GlcNAc)$_3$ was used in the ITC syringe. Aliquots of 8 μL were injected into the reaction cell at 200 s intervals. The stirring speed was set to be 260 rpm. The titrations were performed at $t = 15$ °C and were completed after 40 injections. ITC data were collected automatically using the Microcal Origin version 7.0 software associated with the VP-ITC system. Prior to further analysis, all data were corrected for heat of dilution by subtracting the heat produced by titrating 12 mM (GlcNAc)$_3$ into plain buffer. The fitting of data took place by utilizing a nonlinear least-squares algorithm using a single-site binding model employed by the Origin software, yielding the equilibrium binding association constant ($K_a$) and the enthalpy change ($\Delta H_m$). The stoichiometry (n) was set to be 1 based on the knowledge from the NMR experiments. Errors in $K_a$ and $\Delta H_m$ were obtained as standard deviations from four individual experiments. $K_a$, $\Delta G_m$, $\Delta S_m$, and $-\Delta T\Delta S_m$ were calculated from eq 2, and errors in these parameters were obtained from propagation of error.

Fluorescence Spectroscopy. The partially denatured CBM14 was prepared by incubating the protein in 6 M guanidinium chloride during 24 h at 4 °C. Proteins were used at 16 μM in 8 mM phosphate (Na$_2$HPO$_4$/NaH$_2$PO$_4$) and 16 mM NaCl buffer, pH 7.5. A Varian Cary Eclipse fluorimeter was used with the software Scan (version 1.1) to analyze the intrinsic fluorescence of all of the samples. The device was set up as follows: Excitation wavelength: 295 nm, measured wavelengths: 300–400 nm, excitation fast filter: 2.5 nm, emission fast filter: 5.0 nm, temperature: 25 °C, with 20 scans for each sample.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsomega.9b03043.

**1H,$^{15}$N-HSQC spectrum of $^{13}$C,$^{15}$N-labeled CBM14 (0.1 mM) (Figure S1); restraints and structural statistics for the 20 best conformers of CBM14 (PDB ID: 6SO0) (Table S1); residues involved in β-strands from the crystal (PDB ID: 5HBF) and NMR structure of CBM14 (Table S2); binding isotherm with theoretical fits obtained for the titration of (GlcNAc)$_3$ against CBM14 (Figure S2); chemical shift perturbation for CBM14 interacting with either β-chitin or (GlcNAc)$_3$ (Figure S3); $T_{1\text{rho}}$ 1D $^1$H NMR spectra of CBM14 and (GlcNAc)$_3$ (Figure S4); and proposed model of the interaction between CBM14 and (GlcNAc)$_3$ using HADDOCK (Figure S5) (PDF)

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Notes
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