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Crystal Structure of the GalNAc/Gal-Specific Agglutinin from the Phytopathogenic Ascomycete *Sclerotinia sclerotiorum* Reveals Novel Adaptation of a β -Trefoil Domain

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A lectin from the phytopathogenic ascomycete *Sclerotinia sclerotiorum* that shares only weak sequence similarity with characterized fungal lectins has recently been identified. *S. sclerotiorum* agglutinin (SSA) is a homodimeric protein consisting of two identical subunits of ~17 kDa and displays specificity primarily towards Gal/GalNAc. Glycan array screening indicates that SSA readily interacts with Gal/GalNAc-bearing glycan chains. The crystal structures of SSA in the ligand-free form and in complex with the Gal- β 1,3-GalNAc (T-antigen) disaccharide have been determined at 1.6 and 1.97 Å resolution, respectively. SSA adopts a β -trefoil domain as previously identified for other carbohydrate-binding proteins of the ricin B-like lectin superfamily and accommodates terminal non-reducing galactosyl and *N*-acetylgalactosaminyl glycans. Unlike other structurally related lectins, SSA contains a single carbohydrate-binding site at site α . SSA reveals a novel dimeric assembly markedly dissimilar to those described earlier for ricin-type lectins. The present structure exemplifies the adaptability of the β -trefoil domain in the evolution of fungal lectins.

Keywords: fungal lectin; crystallography; galactose; β -trefoil domain; glycan array

Edited by I. Wilson

Introduction

Numerous mushrooms and other fungi contain carbohydrate-binding proteins commonly known as lectins.^{1–3} Though not all previously identified lectins have been purified and characterized, the available biochemical, sequence and structural data provide ample evidence that fungi, and more specifically the largest and diverse phylum *Ascomycota*, express a heterogeneous mixture of carbohydrate-binding proteins.

Fungal lectins can be assigned as storage proteins^{4,5} and are involved in morphogenesis and development of the fungi.^{6,7} Contrary to established roles of bacterial lectins in host–parasite interactions, the functional roles of lectins expressed by phytopathogenic fungi are poorly understood. Although many believe that fungal lectins mediate host–parasite interactions⁸ similar to bacterial adhesins, they have been implicated in the process of specific recognition in mycoparasitism and mediate interactions between parasitic and pathogenic fungi and their hosts.^{5,9} However, none of these assigned roles are clearly established.

To further corroborate the diversity of fungal lectins, a cytoplasmic lectin has recently been characterized from the phytopathogenic ascomycete *Sclerotinia sclerotiorum*, a fungal pathogen that has the broadest host range of all known fungi.¹⁰

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Abbreviations used: SSA, *Sclerotinia sclerotiorum* agglutinin; EW29, earthworm 29-kDa lectin; MOA, *Marasmius oreades* agglutinin.

The so-called *S. sclerotiorum* agglutinin (SSA) can be considered the prototype of a small lectin family, which hitherto has been documented exclusively in the *Sclerotiniaceae* family. Although orthologs of SSA have been isolated from sclerotes of several *Sclerotinia* species,¹¹ complete sequences are available only for SSA and the orthologs from *Botryotinia fuckeliana*, *Pyrenophora tritici-repentis* and *Cochliobolus heterostrophus*. Sequence comparison reveals that the SSA orthologs represent a novel family of fungal lectins that shares limited sequence similarity with other fungal lectins. Preliminary biochemical and structural characterization has revealed that SSA readily interacts with glycolipid glycans harboring terminal non-reducing Gal or GalNAc residues and galactosylated N-glycans with highest affinity for α 1–3 branched mono- and multi-antennary chains.¹² Since SSA appears to exhibit sugar-binding properties distinct from other previously characterized fungal lectins, it might identify a new lectin subfamily, more widespread within the phylum *Ascomycota*.

A molecular modeling study using threading algorithms has shown that SSA is remotely related to the β -trefoil domain of the non-toxin hemagglutinin HA33/A from *Clostridium botulinum*¹³ that is classified in the ricin B-like lectin superfamily. The ricin B chain, which belongs to the ricin-like or R-type lectin family, is characterized by the presence of a triple (QXW)₃ motif¹⁴ and is widespread in bacteria, animals and plants, but has not been studied extensively in fungi. Only a few ricin B-type lectins have been isolated from fungi, the best known examples being the multi-modular lectins from the mushrooms *Marasmius oreades*¹⁵ and *Polyporus squamosus*,¹⁶ and the pore-forming lectin from the mushroom *Laetiporus sulphureus*.¹⁷ However, we may anticipate that the ricin B domain is also fairly widespread in the newly available fungal genomes.

In a further step towards understanding the carbohydrate specificity and the oligomeric assembly of this novel fungal lectin, we report the 1.60 and 1.97 Å resolution crystal structure of SSA in the absence or presence of bound Gal- β 1,3-GalNAc (T-antigen). The monomer structure of SSA closely resembles that of the ricin B lectin domain, a trefoil-based fold observed in many lectins and carbohydrate-binding domains, but the shape and hydrophobic character of the unique carbohydrate-binding site are markedly modified compared to that of other members of the β -trefoil fold family, consistent with the carbohydrate-binding specificity of the lectin. SSA shares highest structural similarity with the β -trefoil domain of the multi-modular Gal α 1,3Gal-specific lectin from the mushroom *M. oreades*,¹⁸ the hemagglutinin HA1 and HA-17 subcomponents from the *C. botulinum* type C progenitor toxin,¹⁹ and a serotype D toxin complex from *C. botulinum*.²⁰ SSA exhibits a novel dimeric assembly that is rarely observed within members of the ricin B-type lectin family and may participate to multivalent lectin-carbohydrate cross-linking interactions. Altogether, these results suggest that SSA identifies a new lectin subfamily with specific

sequence and carbohydrate-binding properties and might give indications towards the possible involvement of this subfamily of fungal lectins in the regulation of morphogenesis or pathogenesis.

Results and Discussion

Quality and overall view of the structure

The structures of SSA in the apo form and in complex with the T-antigen disaccharide were solved from crystals grown in two distinct space groups. They show well-defined electron densities for most of the protein regions and bound sugar and exhibit excellent stereochemistry (Materials and Methods; Table 1).

The SSA monomer with overall dimensions of 30 Å × 30 Å × 25 Å belongs to the β -trefoil fold family and adopts a typical three-lobed organization that consists of 3 four-stranded β -sheets (β 1– β 4, β 5– β 8, and β 9– β 12), referred to as subdomains α , β , and γ and displaying characteristic pseudo-3-fold symmetry²² (Fig. 1). Structure superposition of the three subdomains reveals that the overall structure of these domains is quite similar to each other with rmsd between subdomains in the 1.55 to 1.66 Å range for ~40 C α atoms, with subdomain β being the smallest. In contrast to most of the extracellular R-type lectins that have been extensively studied, SSA lacks the two disulfide bridges that stabilize the β -trefoil fold, a feature that is also observed in most cytoplasmic ricin B lectin domains. Moreover, the key/characteristic signature QXW motif reminiscent of the R-type lectins is present only in the third subdomain of SSA.¹⁴

A DALI search for close structural homologs of SSA revealed top-ranked hits (Z-score values >20) for the N-terminal β -trefoil domain of the *M. oreades* agglutinin (MOA),¹⁸ the C-terminal β -trefoil domain of the HA1 subcomponent of botulinum type C toxin,¹⁹ and the 17-kDa HA-17 subcomponent of a serotype D toxin complex from *C. botulinum*.²⁰ Surprisingly, SSA appears to possess higher structural homology with multi-modular proteins harboring a β -trefoil domain rather than with individual/isolated R-type lectins. The rmsd's between the SSA structure and those of the three top-ranked homologs above are 1.24, 0.94 and 1.21 Å for 125, 124 and 118 C α atoms, respectively. The major structural differences between SSA and the three lectin homologs are confined to the shape and dimensions of loop regions connecting β -strands and forging the architecture of the carbohydrate-binding sites.

Glycan array screening

Analysis of the carbohydrate-binding specificity of SSA by glycan array screening revealed strong interaction of the lectin with GalNAc-containing oligosaccharides (Table 2), consistent with previously

Table 1. Data collection and refinement

	apo	Gal β 1,3GalNAc
<i>Data collection</i>		
Beamline	ESRF ID23-1	ESRF ID23-2
Resolution (\AA) ^a	55–1.60 (1.69–1.60)	60–1.97 (2.08–1.97)
Space group	$P6_1$	$P6_122$
Cell dimensions: a, b, c (\AA)	127.32, 127.32, 86.65	51.46, 51.46, 295.99
Number of observations	460,339	97,652
Number of unique reflections	100,976	17,040
R_{sym} (%) ^b	0.072 (0.450)	0.140 (0.499)
$\langle I \rangle / \sigma(I)$	16.8 (2.6)	9.0 (3.3)
Redundancy	4.6 (4.0)	5.7 (5.7)
Completeness (%)	96.2 (80.2)	97.88 (98.9)
B -factor from Wilson plot (\AA^2)	16.7	15.7
<i>Refinement</i>		
Resolution (\AA)	55–1.60 (1.64–1.60)	49–1.97 (2.02–1.97)
R_{cryst} ^c	14.00 (21.2)	16.30 (21.40)
R_{free} (%) ^d	16.20 (21.0)	19.51 (24.80)
Number of reflections used in refinement	96,905 (5289)	15,811 (1127)
Number of reflections for R_{free}	4061 (237)	1228 (98)
Number of water molecules	710	169
<i>rmsd</i> ^e		
Bond length (\AA)	0.014	0.014
Bond angles ($^\circ$)	1.51	1.45
Chiral volume (\AA^3)	0.099	0.092
Mean B -factors (\AA^2)		
Main/side chain	19.8/22.9	27.0/29.4
Bound disaccharide	—	27.9
Solvent/other ligands	29.3/30.8	30.1/43.7
<i>rmsd</i> 's on B -factors (\AA^2)		
Main chain	1.11	0.81
Side chain	1.63	1.64
<i>Ramachandran plot statistics</i> (%) ^f		
Residues in favored regions	98.5	98.1
Residues in allowed regions	100	99.4
Outliers	0	0

^a Values in parentheses are for the highest-resolution shell.

^b $R_{\text{sym}} = \sum_{hkl} (\sum_i |I_{hkl} - \langle I_{hkl} \rangle|) / \sum_{hkl} \langle I_{hkl} \rangle$.

^c $R_{\text{cryst}} = \sum_{hkl} ||F_o| - |F_c|| / \sum_{hkl} |F_o|$.

^d R_{free} is calculated for randomly selected reflections excluded from refinement.

^e rmsd from ideal geometry.

^f Ramachandran plot statistics have been calculated with the MolProbity server.²¹

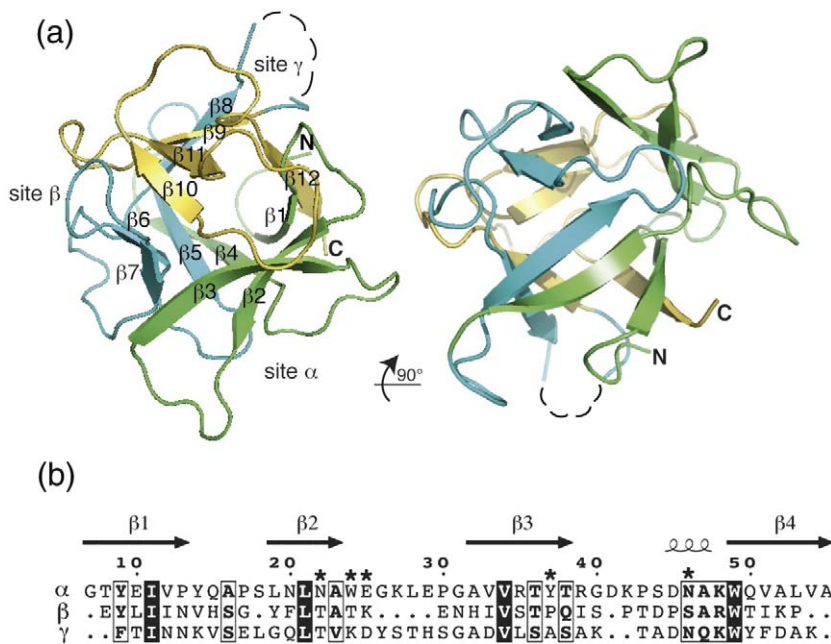


Fig. 1. Overall view of the structure of SSA. (a) Overall structure of the SSA monomer showing the three subdomains, denoted α , β , and γ , viewed in two orientations rotated by 90° and colored green, cyan and orange, respectively. β -Strands of a subunit are labeled $\beta 1$ to $\beta 12$. Broken lines indicate the disordered $\beta 8$ – $\beta 9$ loop region in the apo SSA structure. (b) Structure-based sequence alignment of the three individual subdomains of SSA; residues involved in carbohydrate binding at site α are shown by \star .

Table 2. Glycan array binding of 0.25 µg/ml SSA: the 15 glycan structures are ranked from highest RFU to lowest

Glycan number	Glycan	Mean RFU ^a	STDEV ^b	SEM ^c	%CV ^d
9	GalNAc α -Sp8	23,843	2458	1229	10
84	GalNAc α 1-3Gal β -Sp8	23,589	221	111	1
90	GalNAc β 1-4GlcNAc β -Sp0	21,092	348	174	2
85	GalNAc α 1-4(Fuca1-2)Gal β 1-4GlcNAc β -Sp8	21,025	1327	664	6
329	GalNAc β 1-3Gal α 1-4Gal β 1-4GlcNAc β 1-3Gal β 1-4Glc β -Sp0	19,851	810	405	4
19	GalNAc β -Sp8	19,428	1347	673	7
88	GalNAc β 1-3Gal α 1-4Gal β 1-4GlcNAc β -Sp0	16,809	1706	853	10
298	GalNAc α -Sp15	16,684	634	317	4
91	GalNAc β 1-4GlcNAc β -Sp8	14,934	582	291	4
300	GalNAc β 1-3Gal β -Sp8	14,658	2082	1041	14
346	Gal β 1-4GlcNAc β 1-2Man α 1-3(Man α 1-6)Man β 1-4GlcNAc β 1-4GlcNAc β -Sp12	11,530	555	278	5
4	Gal β 1-3GlcNAc β 1-2Man α 1-3(Gal β 1-3GlcNAc β 1-2Man α 1-6)Man β 1-4GlcNAc β 1-4GlcNAc β -Sp19	7656	571	286	7
83	GalNAc α 1-3GalNAc β -Sp8	7575	54	27	1
55	Fuca1-2Gal β 1-3GalNAc β 1-3Gal α 1-4Gal β 1-4Glc β -Sp9	7122	1485	742	21
59	Fuca1-2Gal β 1-3GalNAc β 1-4(Neu5Ac α 2-3)Gal β 1-4Glc β -Sp9	7074	913	456	13

^a Relative fluorescence unit.^b Standard deviation.^c Standard error measurement.^d Coefficient of variation.

reported binding data using various pNP-oligosaccharides as potential ligands.¹² Although there is a clear preference for terminal non-reducing Gal/GalNAc residues, the specificity of SSA is rather broad and glycans with internal Gal/GalNAc residues are also recognized. SSA also reacts with branched oligosaccharides, but addition of extra sugar moieties can lead to low-affinity glycans. For instance, glycans with α 1,2-fucosyl residues linked to the Gal moiety of the GalNAc α 1,4Gal core show a significant response, while those with fucose linked to the terminal Gal, as found in the core H (type 3) blood group antigen, show only a moderate response. Such an interaction with fucosylated glycans was revealed in a previous binding analysis using version 2 of the printed glycan array.²³ While the profile of binding specificity is similar for the three (0.25, 0.5 and 1.0 µg/ml) lectin concentrations tested, complex N-glycans were recognized only at the two higher concentrations (Supplementary Fig. S1).

The carbohydrate-binding site

The structure of SSA in complex with the Gal β 1,3GalNAc disaccharide reveals a well-ordered carbohydrate at only one (site α) out of the three possible sites (Fig. 2) and provides detailed information about lectin-carbohydrate interactions (Table 3). The non-reducing Gal moiety is tightly bound at the primary binding site within the α site, consistent with the requirement of unsubstituted non-reducing terminal Gal/GalNAc residues for binding activity.¹² The aromatic ring of Tyr37 establishes a stacking interaction against the B face of Gal, while the perpendicularly oriented Trp24 indole ring makes additional Van der Waals contacts with the C6-O6 atom pair. The axial O4 hydroxyl group is tightly anchored to the nitrogen backbone atom of Glu25 and the oxygen atom of the Asn22 side chain. The neighboring O3 hydroxyl is hydrogen

bound to the side chains of Asn22 and Asn46 and to Ser44 via a water molecule. The O2 hydroxyl is bound to Asn46 via a water-mediated interaction. The GalNAc moiety is exposed to the solvent and interacts only weakly with residues in the carbohydrate-binding site (Fig. 2), consistent with the diversity of sugar linkages and moieties allowed beyond the requirement of terminal Gal or GalNAc residues (Table 2). Indeed, only the O4 hydroxyl of GalNAc acts as hydrogen-bond donor to the carbonyl oxygen of Glu25, while the O6 hydroxyl weakly interacts with the Glu25 carboxyl group. The overall conformation of the bound Gal β 1,3GalNAc ($\Phi/\Psi=44^\circ/-10^\circ$) falls within the minimum energy area calculated for the disaccharide.²⁶ Structural comparison of the apo and sugar-bound form reveals no major conformational changes upon binding of the disaccharide. In the apo form, a glycerol molecule is observed in site α and partly mimics a bound Gal moiety (Fig. 2).

Examination of the amino-acid sequence of SSA reveals that the key aromatic side chains (Tyr37 and Trp24) for sugar binding in site α are absent in sites β and γ and may explain the lack of sugar-binding activity in these two sites (Fig. 1). The carbohydrate-binding site of SSA is shallow and solvent-exposed and may permit accommodation of both the α and β conformation of the 1,3- or 1,4-glycosidic linkage. The carbohydrate-binding site of SSA is filled with 14 well-ordered water molecules that are mostly clustered in the vicinity of the O2 and O3 positions of Gal, suggesting that SSA could accommodate the larger acetamido group of GalNAc, other sugar substituents, or bisecting/multi-antennary glycan chains. Indeed, models of the α -GalNAc complex generated using Autodock Vina²⁷ showed that the docked orientation of α -GalNAc is nearly identical with that of Gal in the T-antigen complex observed in the X-ray crystal structure. The N-acetyl group lies in the vicinity of the Glu25-Gly26 backbone

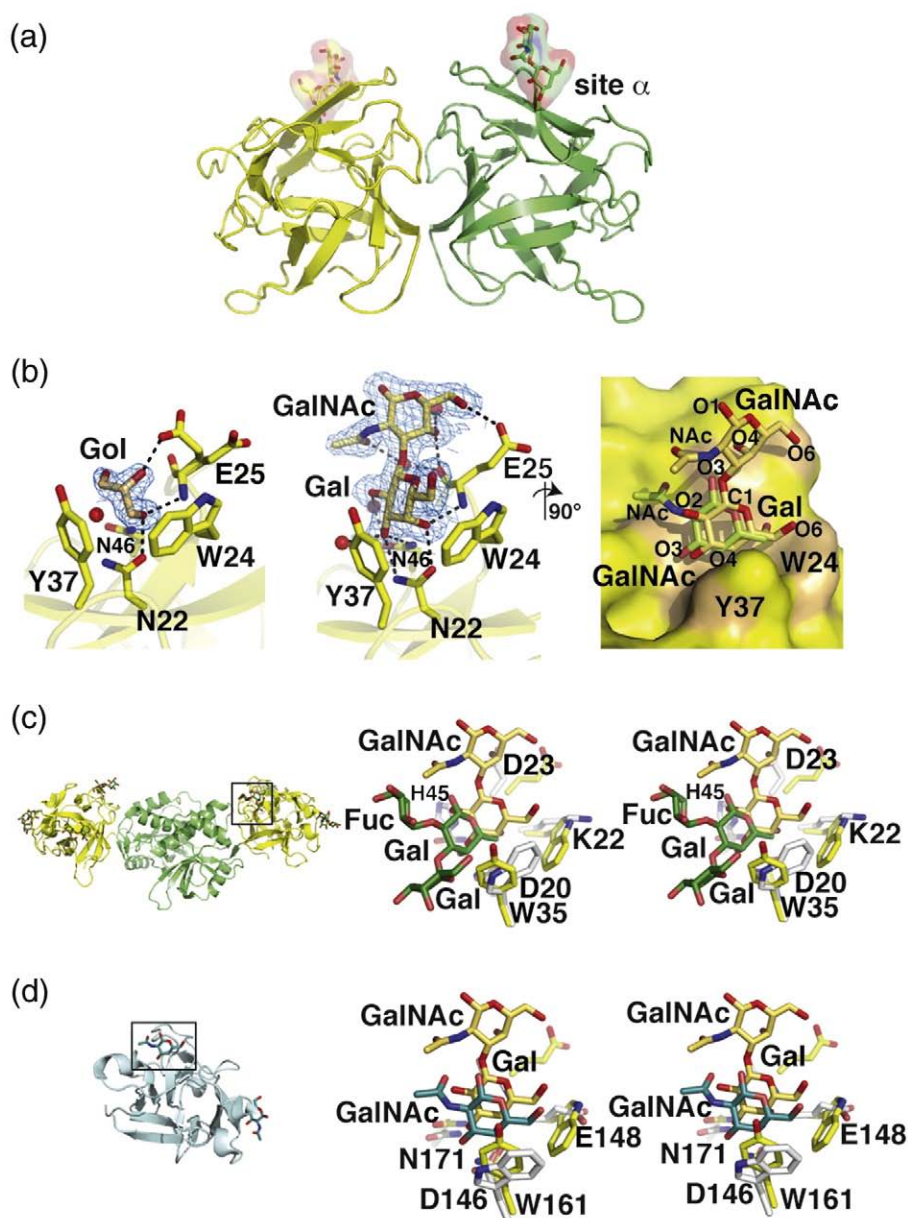


Fig. 2. Carbohydrate-binding site at site α . (a) Overall view of the dimer with bound Gal β 1,3GalNAc in each subunit. (b) Close-up views of the carbohydrate-binding site in (left) the apo form with bound glycerol (orange) and (middle) the complex with bound Gal β 1,3GalNAc (orange), viewed in a similar orientation. The 1.95 Å resolution $2mF_o - DF_c$ electron density map (cyan) is contoured at 1σ . Molecular surface (right) showing the shallow binding site. The position of the docked α -GalNAc (green) is displayed and overlaid with bound Gal β 1,3GalNAc. Side chains that directly interact with the disaccharide are labeled. (c) Overall view (left) of the MOA dimer with bound blood group B Gal α 1,3(Fuc α 1,2)Gal trisaccharide (green)²⁴ (accession code: 3EF2). The N- and C-terminal domains from each subunit are colored yellow and green, respectively. Close-up stereo view (right) of the overlay of the carbohydrate-binding sites in SSA and MOA. (d) Overall view (left) of the EW29 lectin with bound GalNAc (blue)²⁵ (accession code: 2ZQO). Close-up stereo view (right) of the overlay of the carbohydrate-binding sites, oriented as in (c), in SSA and EW29. Side chains that directly interact with the bound carbohydrate in EW29 and MOA are labeled.

atoms and the side chains of Ser44 and Asn46 (Fig. 2), indicating that terminal *N*-acetylgalactosaminyl moieties could also be easily accommodated into the carbohydrate-binding site of SSA, consistent with the glycan array data (Table 2).

Despite low sequence similarity, the carbohydrate-binding site of SSA reveals structural similarities to other previously reported ricin-like or R-type lectin-

sugar complexes. Although SSA displays only a single carbohydrate-binding site at site α , the mode of binding to Gal resembles that observed in other galactose-binding lectins. Structural comparison of SSA and the galactose-binding lectin EW29 (earthworm 29-kDa lectin)²⁵ from the earthworm *Lumbricus terrestris*, which contains two carbohydrate-binding sites at sites α and γ (Fig. 2), reveals a

Table 3. SSA–Gal β 1,3GalNAc interactions

Ligand atom ^a	Direct contact (Å)	Interaction partner
Gal O4	2.8	Asn22 OD1
	2.8	Glu25 N
Gal O3	2.9	Asn22 ND2
	3.0	Asn46 ND2
	2.7	Wat25
Gal O2	2.8	Wat45
	Gal C6–O6	3.3–3.5
Gal C3–C4	3.6–3.8	Tyr37 phenol ring
GalNAc O4	2.7	Glu25 O
	2.8	Wat98
GalNAc O7	3.0	Wat46
GalNAc O1	2.7	Wat130
	2.7	Wat118

Contacts observed in a single subunit within a 3.2 and 4.0 Å distance for hydrogen bonds and hydrophobic carbon–carbon interactions, respectively.

^a Hydrophobic interactions are indicated in boldface.

similar orientation of the terminal Gal moiety in EW29, with an outward 1.6 Å displacement of the sugar ring compared to its position in SSA along with a shift of the β 1– β 2 loop. Despite the close structural similarities between SSA and the β -trefoil domain of MOA (rmsd of 1.24 Å for 125 C α atoms), the Gal moiety adopts a distinct position and orientation in the two structures. In MOA, the non-reducing terminal Gal moiety of the blood group B trisaccharide is shifted by \sim 1.5 Å and rotated by \sim 30° compared to its position in SSA to favor interactions between the O4 and O6 hydroxyl groups of Gal and MOA (Fig. 2).

A novel dimeric interface

The SSA crystal structure identifies a novel dimer interface for an R-type fungal lectin, consistent with data from gel-filtration experiments (Supplementary Fig. S2). The overall architecture of the SSA dimer consists of a compact assembly with dimensions of 60 Å \times 30 Å \times 25 Å that buries a flat surface on each β -trefoil monomer. The dimer interface is formed by the tight association of the β 4 and β 5 strands from

each monomer with the participation of two peripheral loop regions (β 5– β 6 and β 7– β 8) (Fig. 2). In turn, the dimeric assembly of SSA differs from the heterodimeric assembly of the lectin domain of ricins.²⁸ In fact, a structural overlay between SSA and the ricin B chain from the castor bean plant²⁹ reveals that a nine-residue insertion in the β 4– β 5 loop of SSA prevents formation of a dimeric assembly as seen in the ricin R-type lectin domain. In SSA, the calculated buried surface area to a 1.4 Å probe radius encompasses \sim 950 Å² on each subunit and involves 16 residues that are dominantly apolar. The topology of the β -strands recruited in this novel dimer interface is conserved in sequence family members of SSA (see below) and residues involved in this assembly are well conserved, suggesting that this type of dimeric association should be preserved in SSA homologs.

Diversity in the oligomeric assembly of various lectins that share a similar fold has been already described and may account for their multivalent carbohydrate-binding specificities.³⁰ In the SSA dimer, the two carbohydrate-binding sites, which are located on the same face of the dimer, are separated by 38 Å, a distance preventing the formation of cross-linking interactions between different antennae of a single N-glycan. Although the nature and biological significance of cross-linking interactions remain to be investigated, the remote location of the two carbohydrate-binding sites in the SSA dimer may favor interaction with distinct glycan chains harboring terminal galactosyl residues.

BLAST searches using the SSA sequence as a template identified close homologs in the genome of the three plant pathogenic fungi, *B. fuckeliana* (*Botrytis cinerea*), *P. tritici-repentis* and *C. heterostrophus*, the former two belonging to the same *Pleosporaceae* family. These three putative SSA homologs share approximately 77%, 27% and 28% sequence identity, respectively, and may belong to the same family of fungal lectins, as key residues at the carbohydrate-binding site are conserved in site α along with most of side chains involved in the dimeric assembly (Fig. 3).

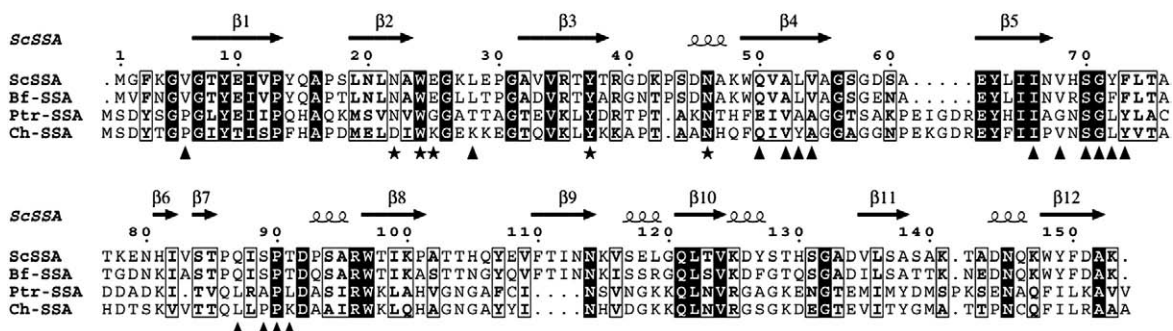


Fig. 3. Sequence alignment of SSA with homologs from *B. fuckeliana* (Bf-SSA, accession EDN28997.1), *P. tritici-repentis* (Ptr-SSA, accession EDU42915.1) and *C. heterostrophus* (Ch-SSA, accession FK679805.1), as putative representative members of this new class of fungal lectins. Identical residues are indicated with a black background while similar residues are boxed. Amino acids involved in carbohydrate binding and dimer assembly of SSA are indicated by ★ and ▲, respectively. Secondary-structure elements are indicated above the sequence.

In summary, the crystal structure of SSA reveals that the specific binding to terminal Gal/GalNAc sugars with 1,3 linkages is achieved by both direct and water-mediated hydrogen bonds to the O4 and O3 hydroxyls of Gal at the non-reducing end. The novel dimeric assembly demonstrates that different quaternary structures exist within the R-type lectin family, resulting from a different association of structurally similar domains. Whereas other types of quaternary arrangements may occur within the extended group of fungal lectins, the restricted number of functional binding sites per β -trefoil domain in SSA could dictate the type of quaternary structures within this new family of fungal lectins and affect the ability to establish multiple interactions and cross-link glycan receptors. By analogy with the dual storage–defense role attributed to a large group of abundant plant lectins, the *Sclerotiniaceae* lectins might also combine a likely function as storage protein in the development and morphogenesis of the fungus, with a possible defense role against predating organisms. Further studies to elucidate/investigate the biological role of this emerging class of fungal lectins are required.

Materials and Methods

Materials

SSA was isolated from mature *S. sclerotiorum* sclerotes and purified as described previously.¹¹

Crystallization and data collection

For crystallization, ammonium sulfate-precipitated SSA was extensively dialyzed against 10 mM HEPES, pH 7.5, and 150 mM NaCl and concentrated to 7.3 mg/ml. Crystallization experiments were performed by vapor diffusion with Greiner plates (Greiner-BioOne) using a Freedom (Tecan) and Honeybee (Cartesian) robot. Initial crystals of apo SSA were obtained at 20 °C from a condition containing 2.0 M sodium chloride and 0.1 M sodium acetate, pH 4.6, of the Structure Screen 2 (Molecular Dimension Limited). After manual optimization in Limbro plates, larger crystals appeared within 10 days from typically 1 μ l of protein solution and 0.5 μ l of reservoir containing 2.1 M sodium chloride and 0.1 M sodium acetate, pH 4.6. The SSA–Gal β 1,3GalNAc complex was made by mixing the protein solution with 2.5 mM Gal β 1,3GalNAc followed by preincubation overnight at 4 °C prior to crystallization experiments. Crystals of the complex appeared at 20 °C from a condition containing 1.26 M ammonium sulfate, 0.2 M lithium sulfate and 0.1 M Tris, pH 8.5. Crystals selected for data collection were rapidly transferred in the reservoir solution supplemented with 25% glycerol (apo) or 30% polyethylene glycol 400 (complex), flash cooled at 100 K in a nitrogen gas stream, and stored in liquid nitrogen. Data were collected on the European Synchrotron Radiation Facility beamlines ID23-1 and ID23-2 (Grenoble, France). Oscillation images were integrated with MOSFLM³¹ and scaled and merged with SCALA.³²

Structure determination and refinement

Attempts to solve the structure using a Se-labeled D-Gal- β -SePh derivative were unsuccessful. Initial phases were obtained by molecular replacement with the Phaser program³³ using as search model a structural template generated with the protein homology recognition engine Phyre,³⁴ which uses profile–profile matching algorithms for the detection of homologs of known three-dimensional structure, the so-called template-based homology modeling or fold recognition. The model generated by the Phyre server was based on the crystal structure of *C. botulinum* non-toxin hemagglutinin HA33/A¹³ (Protein Data Bank accession number 1YBI) and led to an unambiguous solution in Phaser with a Z-score value of 51 in the 15- to 2.8 Å resolution range. The resulting model, comprising four molecules of SSA in the asymmetric unit, was rebuilt in an automatic fashion with the program ARP/wARP.³⁵ Final refinement was then performed with REFMAC³⁶ using data up to 1.6 Å resolution, and the resulting σ_A -weighted $2mF_o - DF_c$ and $mF_o - DF_c$ electron density maps were used to correct the model with the graphics program COOT.³⁷ The structure of the SSA disaccharide complex was solved by molecular replacement with Phaser,³³ using apo SSA as a search model. The structure of the complex was subsequently refined and manually adjusted with the programs REFMAC³⁶ and Coot,³⁷ respectively.

The final apo model comprises residues Gly2 to Lys153 for the four SSA molecules and two glycerol molecules arising from the cryo buffer; that of the disaccharide complex comprises residues Gly2 to Lys153 for a single SSA molecule, a Gal β 1,3GalNAc disaccharide, four sulfate ions arising from the crystallization buffer, and two small polyethylene glycol molecules arising from the cryo buffer. In the two structures, the backbone region of residues Gly58 to Ser60 adopts two alternate conformations while, in the apo structure, the surface loop region Thr103–Glu108 is disordered. The rmsd between the apo and sugar-bound form is 0.64 Å for 144 C α atoms. In the apo form, the β 2– β 3 surface loop region containing the residue pair Trp24–Glu25 at site α undergoes large conformational changes in one out of four molecules, but these movements are mediated by the crystal packing environment. The stereochemistry of the models was analyzed with MolProbity;²¹ no residues were found in the disallowed regions of the Ramachandran plot. Data collection and refinement statistics are reported in Table 1. Structural sequence alignment was performed with the RAPIDO server.³⁸ Figures 1 and 2 were generated with PyMOL.³⁹ Figure 3 was prepared with ESPript.⁴⁰

Glycan array screening

The glycan binding specificity of SSA was analyzed by the Consortium for Functional Glycomics using micro-arrays v3.1.²³ Full details of protocols used and the ~380 glycans included in the array are presented on the Consortium's web site† and described by Paulson *et al.*²³ Lyophilized SSA was dissolved in phosphate-buffered saline at 1 mg/ml and labeled with tetrafluorophenyl-Alexa Fluor 488 using the Invitrogen protein labeling kit following the manufacturer's instructions. Labeled SSA was diluted to 0.25 μ g/ml in Tris-buffered saline (20 mM Tris, 150 mM NaCl, 2 mM CaCl₂, and 2 mM MgCl₂,

† <http://www.functionalglycomics.org/static/consortium/resources/resourcecoreh8.shtml>

pH 7.4) containing 1% bovine serum albumin and 0.05% Tween 20. An aliquot (70 μ l at 0.25 μ g/ml) of the labeled lectin solution was applied to separate microarray slides and incubated under a coverslip for 60 min in a dark, humidified chamber at room temperature. After incubation, the coverslips were gently transferred to a solution of Tris-buffered saline containing 0.05% Tween 20 and successively washed four times by gently dipping the slides in Tris-buffered saline containing 0.05% Tween 20 and deionized water. After the last wash, the slides were spun in a slide centrifuge for approximately 15 s to dry and immediately scanned in a ProScanArray MicroArray Scanner (PerkinElmer) using an excitation wavelength of 488 nm and ImaGene software (BioDiscovery, Inc., El Segundo, CA) to quantify fluorescence. Data are expressed as average relative fluorescence units for binding each glycan calculated by averaging four values after removing the highest and lowest values; STDEV is the standard deviation, SEM is the standard error measurement, and CV is the coefficient of variation (SD/mean) calculated as percentage. Analysis of the carbohydrate specificity of SSA on the glycan array was repeated at lectin concentrations of 0.50 and 1.0 μ g/ml.

Molecular docking

α -GalNAc was selected as a candidate ligand for automated docking to SSA using Autodock Vina.²⁷ SSA and the carbohydrate ligands were treated as a rigid protein and flexible molecules, respectively. The grid of the docking simulation was defined by a 20 $\text{Å} \times 20 \text{Å} \times 20 \text{Å}$ cube centered on the SSA α site. The docking simulation was performed using the default parameters. For each ligand, the nine top-ranked generations based on the predicted binding affinity (in kilocalories per mole) were analyzed.

Accession numbers

Coordinates and structure factors have been deposited in the Protein Data Bank with accession numbers 2X2S and 2X2T.

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Supplementary Data

Supplementary data associated with this article can be found, in the online version, at [doi:10.1016/j.jmb.2010.05.038](https://doi.org/10.1016/j.jmb.2010.05.038)

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