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Purification, crystallization and preliminary X-ray diffraction analysis of the carbohydrate-binding region of the *Streptococcus gordonii* adhesin GspB

The carbohydrate-binding region of the bacterial adhesin GspB from *Streptococcus gordonii* strain M99 (GspB_{BR}) was expressed in *Escherichia coli* and purified using affinity and size-exclusion chromatography. Separate sparse-matrix screening of GspB_{BR} buffered in either 20 mM Tris pH 7.4 or 20 mM HEPES pH 7.5 resulted in different crystallographic behavior such that different precipitants, salts and additives supported crystallization of GspB_{BR} in each buffer. While both sets of conditions supported crystal growth in space group $P2_12_12_1$, the crystals had distinct unit-cell parameters of $a = 33.3$, $b = 86.7$, $c = 117.9$ Å for crystal form 1 and $a = 34.6$, $b = 98.3$, $c = 99.0$ Å for crystal form 2. Additive screening improved the crystals grown in both conditions such that diffraction extended to beyond 2 Å resolution. A complete data set has been collected to 1.3 Å resolution with an overall R_{merge} value of 0.04 and an R_{merge} value of 0.33 in the highest resolution shell.

1. Introduction

GspB is a 3072-residue cell-wall-anchored glycoprotein from *Streptococcus gordonii* that mediates the binding of this bacterium to salivary glycoproteins and human platelets (Bensing & Sullam, 2002). The former interaction is important for colonization of the oral cavity, while the latter is a key event in the pathogenesis of infective endocarditis (Xiong *et al.*, 2008). This unusual adhesin is a member of an expanding family of serine-rich glycoproteins that are found in most pathogenic streptococci and staphylococci (Takahashi *et al.*, 2006; Siboo *et al.*, 2005; Rose *et al.*, 2008; Seifert *et al.*, 2006) and are believed to mediate pathogen host attachment and promote bacterial biofilm formation (Sanchez *et al.*, 2010; Wu *et al.*, 2007). GspB contains an atypical 90-amino-acid N-terminal signal peptide, a serine-rich region, a basic region, a second serine-rich region and a C-terminal cell-wall-anchoring domain (Fig. 1). The basic region (GspB_{BR}) mediates the binding of GspB to sialylated carbohydrate moieties on platelet glycoprotein Ib α (Takamatsu *et al.*, 2005), salivary mucin MG2 and salivary agglutinin (Takamatsu *et al.*, 2006) through its high-affinity interaction with NeuAc α (2–3)Gal β (1–3)GalNAc (sialyl-T antigen). Of note, the binding properties of GspB homologs vary considerably, with some homologs (*e.g.* Hsa of *S. gordonii* strain Challis and SrpA of *S. sanguinis*) binding to a broader or different range of carbohydrate motifs, while others (such as PsrP of *S. pneumoniae*) have no apparent lectin-like activity (Shivshankar *et al.*, 2009; Takamatsu *et al.*, 2005; Yajima *et al.*, 2005).

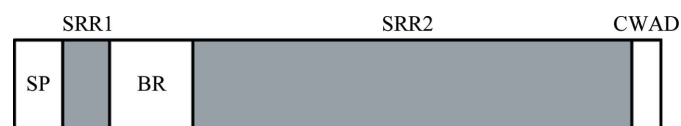


Figure 1
Overall architecture of GspB. GspB is comprised of a signal peptide (SP), a short serine-rich region (SRR1), a unique basic region (BR) that is responsible for carbohydrate binding, a second, longer, serine-rich region (SRR2) and a cell-wall-anchoring domain (CWAD). A recently published structural study of the Fap1 adhesin from *S. parasanguinis* identified the structural elements of both the serine-rich repeats and the unique region in that protein (Ramboarina *et al.*, 2010). The repeat region of GspB is likely to form a super-helical fibril like that observed in the repeat region of Fap1.

These studies will ultimately identify the molecular details of carbohydrate selectivity by GspB and related serine-rich repeat adhesins. This is a critical starting point for understanding how carbohydrate binding by GspB and related lectins affects pathogen infectivity. Furthermore, a crystal structure may allow the design of small-molecular inhibitors to disrupt carbohydrate binding, which offers a new route for the design of therapeutics. Here, we report the expression, purification and crystallization of the carbohydrate-binding domain GspB_{BR} from *S. gordonii* strain M99.

2. Materials and methods

2.1. Expression and purification

GspB_{BR} comprises residues 233–615 of GspB and retains lectin behavior (Takamatsu *et al.*, 2005). The gene encoding GspB_{BR} was amplified from a patient isolate of *S. gordonii* strain M99 and was cloned into the pGEX plasmid encoding an N-terminal GST fusion tag as previously described (Takamatsu *et al.*, 2005). This plasmid was transformed into electrocompetent *Escherichia coli* BL21 Gold (DE3) cells (Stratagene) and grown in LB medium with 100 µg ml⁻¹ ampicillin at 310 K until the optical density at 600 nm (OD₆₀₀) was

between 0.4 and 0.5. The cells were then cold-shocked by incubation in an ice-water bath for 20 min. GspB_{BR} expression was induced by the addition of isopropyl β-D-1-thiogalactopyranoside (IPTG) to a final concentration of 0.1 mM and the cells were incubated with shaking at 291 K overnight. The cells were harvested by centrifugation at 4000g for 15 min at 277 K and the pellets were washed with ice-cold phosphate-buffered saline (PBS) and spun down at 4500g for 15 min at 277 K.

The cell pellets were resuspended in ice-cold PBS supplemented with 1 mM dithiothreitol, 1 µg ml⁻¹ leupeptin, 1 µg ml⁻¹ pepstatin A, 0.17 mg ml⁻¹ phenylmethylsulfonyl fluoride, 1 µg ml⁻¹ DNaseI and 10 µg ml⁻¹ RNase. The cells were disrupted by sonication for 20 min with 0.7 s pulses at 277 K. The lysate was clarified by centrifugation at 27 000g for 30 min. The supernatant was passed through a 0.45 µm filter and loaded onto a 5 ml GSTrap column (GE Healthcare) at 1 ml min⁻¹. The column was washed with PBS supplemented with 1 mM dithiothreitol until the absorbance at 280 nm (A₂₈₀) returned to 0 and was then eluted with buffer containing 50 mM Tris pH 8.0 and 10 mM L-glutathione. The eluted protein was concentrated using a 50 kDa molecular-weight cutoff spin concentrator (Millipore). The GST tag was cleaved using 1 U factor Xa (NEB) in a buffer consisting of 100 mM NaCl, 2 mM CaCl₂ and 50 mM Tris pH 8.0 for 36 h at room temperature (296 K). The cleavage reaction was passed through a 0.2 µm filter (Costar, Spin-X) and separated at 0.3 ml min⁻¹ on a 24 ml Hi-Load Superdex 200 10/300 size-exclusion column (GE Healthcare) equilibrated in 20 mM Tris pH 7.4 or 20 mM HEPES pH 7.5. Fractions were analyzed using SDS-PAGE (Fig. 2) and concentrated using a 30 kDa molecular-weight cutoff concentrator (Millipore). The protein concentration was determined using the Quickstart Bradford Assay Kit (Bio-Rad).

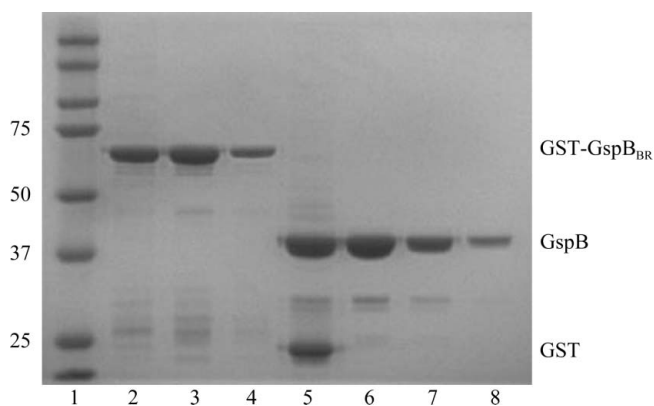


Figure 2 Purification of GspB_{BR} for crystallization. The gel shows two separate concentrations of the GST-GspB_{BR} fusion protein after affinity purification and two concentrations of purified isolated GspB_{BR}(246–601) after size-exclusion chromatography. Lane 1, Kaleidoscope Ladder (Bio-Rad; labeled in kDa); lanes 2 and 3, 4 µg GST-GspB_{BR}; lane 4, 1 µg GST-GspB_{BR}; lane 5, 4 µg of GST and GspB_{BR} after factor Xa cleavage; lane 6, 4 µg GspB_{BR} after size-exclusion chromatography; lane 7, 1 µg GspB_{BR} after size-exclusion chromatography; lane 8, dissolved crystals grown in 26% Jeffamine-ED 2001 and 0.05 M HEPES pH 7.5.

2.2. Crystallization

GspB_{BR} formed diffraction-quality crystals (Fig. 3) in two chemically distinct sets of conditions that resulted from sparse-matrix screening of the protein in two different buffers: 20 mM Tris pH 7.4 and 20 mM HEPES pH 7.5. Sparse-matrix screening of GspB_{BR} in either buffer was performed using a Mosquito crystallization robot (TTP LabTech) and sampled conditions from Crystal Screen, Crystal Screen 2 and Index Screen (Hampton Research) and Wizard I and II (Emerald BioSystems).

The first crystal form was grown using the hanging-drop vapor-diffusion method with drops consisting of 1 µl 10 mg ml⁻¹ GspB_{BR} in 20 mM HEPES pH 7.5 mixed with 1 µl reservoir solution (25% PEG

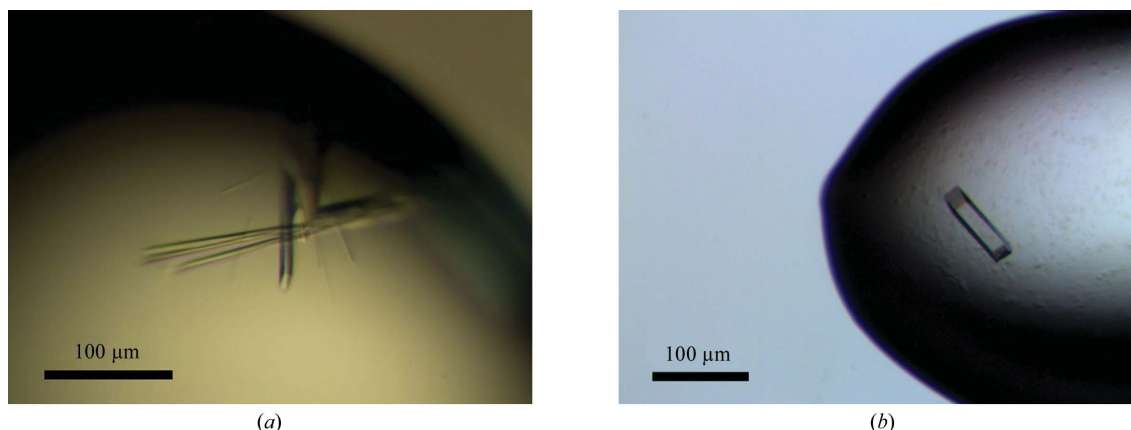


Figure 3 Crystals of GspB_{BR} from *S. gordonii* strain M99 grown using the hanging-drop vapor-diffusion method. (a) Crystals of GspB_{BR} grown from 25% PEG 3350, 0.15 M ammonium acetate, 0.1 M HEPES pH 7.5 and 10 mM spermidine. (b) Crystals of GspB_{BR} grown from 34% Jeffamine ED-2001, 0.15 M KCl and 0.1 M HEPES pH 7.5.

3350, 0.15 M ammonium acetate, 0.1 M HEPES pH 7.5) and equilibrated against 1 ml reservoir solution. Crystals grew within 3 d and were flash-cooled in liquid nitrogen without the addition of an additional cryopreservation agent. The initial crystals diffracted to 3 Å resolution. Optimization was performed using grid screening together with the Additive Screen kit from Hampton Research. The inclusion of 10 mM spermidine in the crystallization conditions increased the crystal size and improved the diffraction limit to 1.9 Å resolution (Fig. 4a).

A second set of chemically distinct crystallization conditions was identified for GspB_{BR} after repeating sparse-matrix screening with 6 mg ml⁻¹ GspB_{BR} in 20 mM Tris pH 7.4. Crystals were grown using the hanging-drop vapor-diffusion method with drops consisting of 1 µl GspB_{BR} and 1 µl reservoir solution [33% Jeffamine ED-2001 (Hampton Research) and 0.1 M HEPES pH 7.5] equilibrated against 1 ml reservoir solution. Crystals appeared within 2 d, but required three weeks to grow to full size. Prior to flash-cooling in liquid nitrogen, crystals were cryoprotected in a solution containing all of the components of the reservoir solution with the addition of 15% glycerol. Additive screening identified that the inclusion of 0.15 M KCl improved the diffraction quality of these crystals from 1.8 to 1.3 Å resolution (Fig. 4b).

2.3. Data collection and processing

Crystal quality was assessed on Stanford Synchrotron Radiation Lightsource (SSRL) beamlines 9-2, 11-1 and 12-2 and the Life Sciences Collaborative Access Team (LS-CAT) ID-21-D/F/G beamlines. Although the two crystallization conditions are chemically distinct, both resulted in the formation of orthorhombic crystals in space group $P2_12_12_1$ (Table 1). Diffraction data for the crystals obtained using PEG 3350 as the precipitant were collected on beamline 11-1 at SSRL using a MAR 325 CCD detector with a distance of 250 mm and a wavelength of 1.03034 Å. A data set consisting of 90 frames was collected with a rotation angle of 90° and an exposure time of 20 s per frame. Diffraction data for the crystals

Table 1

X-ray data-collection statistics for crystals scaled in space group $P2_12_12_1$.

Values in parentheses are for the highest resolution shell.

	Crystals obtained using	
	PEG 3350	Jeffamine ED-2001
Space group	$P2_12_12_1$	$P2_12_12_1$
Wavelength (Å)	1.03034	0.97856
Beamline	SSRL 11-1	LS-CAT ID-21-G
Resolution (Å)	50–1.90	50–1.29
Unit-cell parameters (Å, °)	$a = 33.3, b = 86.7,$ $c = 117.9,$ $\alpha = \beta = \gamma = 90$	$a = 34.6, b = 98.3,$ $c = 99.0,$ $\alpha = \beta = \gamma = 90$
No. of measured reflections	84235	380613
No. of unique reflections	26193	83230
Multiplicity	3.2 (2.9)	4.6 (2.8)
$\langle I/\sigma(I) \rangle$	16.8 (3.9)	21.3 (2.6)
Completeness (%)	93.8 (92.4)	96.7 (73.6)
R_{merge}^\dagger (%)	6.8 (35.3)	4.0 (33.2)
No. of molecules in asymmetric unit	1	1
Matthews coefficient (Å ³ Da ⁻¹)	2.2	2.1
Solvent content (%)	43.2	42.5

$^\dagger R_{\text{merge}} = \frac{\sum_{hkl} \sum_i |I_i(hkl) - \langle I(hkl) \rangle|}{\sum_{hkl} \sum_i I_i(hkl)}$, where $I_i(hkl)$ is the i th instance of the intensity at position hkl and $\langle I(hkl) \rangle$ is the average of all instances of the reflections at position hkl .

obtained using Jeffamine ED-2001 as the precipitant were collected on beamline ID-21-G at the Advanced Photon Source using a MAR 325 CCD detector with a distance of 165 mm and a wavelength of 0.97856 Å. A data set consisting of 240 frames was collected with a rotation angle of 90° and an exposure time of 2 s per frame. All data were processed and scaled using the *HKL-2000* program package (Otwinowski & Minor, 1997).

3. Results and discussion

3.1. Crystallization and data collection

Sparse-matrix screening of the GspB_{BR} protein in different buffers resulted in the identification of dramatically different crystallization conditions (Fig. 3). This suggests that parallel sparse-matrix screening

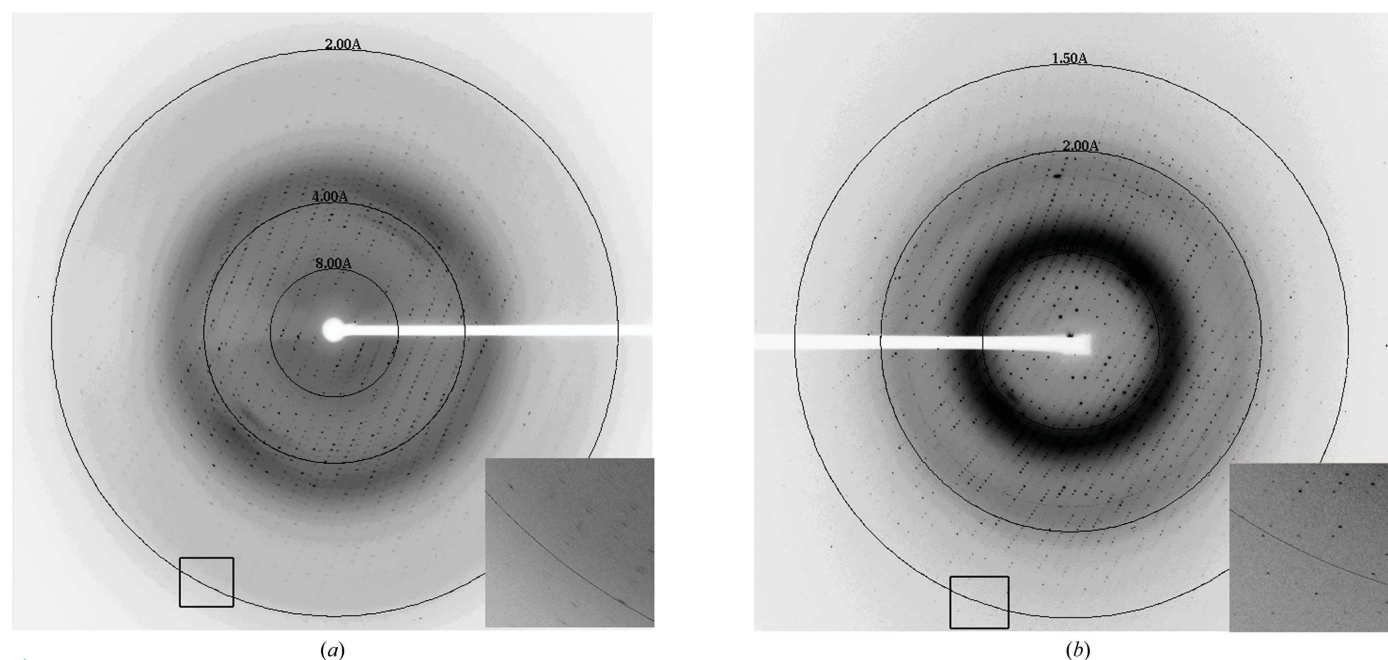


Figure 4

Typical diffraction for each of the crystal forms. (a) Diffraction image for crystals grown using PEG 3350 as the precipitant. This diffraction image was collected on the SSRL 11-1 beamline and diffraction to 1.8 Å resolution can be observed. (b) Typical diffraction image for crystals grown using Jeffamine ED-2001 as the precipitant. This diffraction image was collected on the LS-CAT ID-21-G beamline and diffraction to 1.3 Å resolution can be observed.

Table 2

 Data-collection statistics for the DyCl₃ and HoCl₃ MAD data sets.

Values in parentheses are for the highest resolution shell.

	DyCl ₃			HoCl ₃		
	Peak	Inflection	Remote	Peak	Inflection	Remote
Wavelength (Å)	1.59083	1.59122	1.02463	1.53523	1.53583	1.01623
Beamline	SSRL 9-2	SSRL 9-2	SSRL 9-2	SSRL 9-2	SSRL 9-2	SSRL 9-2
Resolution (Å)	50–1.98	50–1.98	50–1.55	50–2.90	50–2.80	50–2.75
Unit-cell parameters (Å, °)	$a = 35.5, b = 98.5,$ $c = 99.1,$ $\alpha = \beta = \gamma = 90$	$a = 34.5, b = 98.5,$ $c = 99.1,$ $\alpha = \beta = \gamma = 90$	$a = 34.5, b = 98.4,$ $c = 99.0,$ $\alpha = \beta = \gamma = 90$	$a = 34.6, b = 99.0,$ $c = 98.3,$ $\alpha = \beta = \gamma = 90$	$a = 34.6, b = 99.0,$ $c = 99.0,$ $\alpha = \beta = \gamma = 90$	$a = 34.7, b = 99.1,$ $c = 98.8,$ $\alpha = \beta = \gamma = 90$
No. of measured reflections	203951	206064	431338	45488	48993	53679
No. of unique reflections	23918	23929	48691	6172	6842	6889
Multiplicity	8.5 (8.1)	8.6 (8.1)	8.9 (8.3)	7.4 (6.0)	7.2 (5.1)	7.8 (6.6)
$I/\sigma(I)$	31.2 (19.3)	31.2 (18.4)	27.2 (6.2)	12.4 (3.6)	11.1 (2.7)	10.4 (3.1)
Completeness (%)	98.0 (96.2)	98.0 (95.6)	97.2 (99.6)	77.1 (80.6)	76.9 (75.6)	73.1 (77.7)
$R_{\text{merge}}^{\dagger}$ (%)	8.7 (17.9)	8.0 (17.7)	6.1 (35.9)	14.5 (32.6)	13.9 (32.0)	12.8 (36.2)

$\dagger R_{\text{merge}} = \sum_{hkl} \sum_i |I_i(hkl) - \langle I(hkl) \rangle| / \sum_{hkl} \sum_i I_i(hkl)$, where $I_i(hkl)$ is the i th instance of the intensity at position hkl and $\langle I(hkl) \rangle$ is the average of all instances of the reflection at position hkl .

of proteins in different buffers may be a general method to increase the probability of growing diffraction-quality crystals of a protein. In this case, orthorhombic crystals of GspB_{BR} with 20 mM HEPES pH 7.5 as the buffer formed from conditions that used PEG 3350 as the precipitant, while crystals of GspB_{BR} with 20 mM Tris pH 7.4 as the buffer formed from conditions using Jeffamine ED-2001 as the precipitant. Crystals of GspB_{BR} grown using Jeffamine ED-2001 as the precipitant displayed superior diffraction quality (Fig. 4) and reproducibility. As a result, this crystal form was used exclusively for structure determination.

The unit-cell parameters for the crystals grown from the conditions containing PEG 3350 were $a = 33.3, b = 86.7, c = 117.9$ Å, $\alpha = \beta = \gamma = 90^\circ$, while the unit-cell parameters for the crystals grown from the conditions containing Jeffamine ED-2001 as the precipitant were $a = 34.6, b = 98.3, c = 99.0$ Å, $\alpha = \beta = \gamma = 90^\circ$ (Table 1). Since the b and c unit-cell parameters are similar in length for the crystals grown from Jeffamine ED-2001, the crystals were originally assumed to be tetragonal; however, scaling in *SCALEPACK* (Otwinowski & Minor, 1997) resulted in unreasonable R_{merge} values and an unreasonable number of rejected reflections. This strongly suggested that the crystals were orthorhombic. Specific volume calculations (Matthews, 1968) suggested the presence of one molecule of GspB_{BR} per asymmetric unit and a solvent content of 43% for both crystal forms (Table 1).

3.2. Identification of heavy-atom derivatives

GspB_{BR} does not exhibit significant sequence similarity to any protein of known structure; it contains only two methionines and no cysteines. As a result, heavy-atom derivatives were prepared for phasing. Two heavy-atom derivatives, Dy³⁺ and Ho³⁺, were prepared by soaking GspB_{BR} crystals with either 1 mM DyCl₃ for 3 d or 10 mM HoCl₃ for 3 d. Data were collected using the wavelengths and beamlines listed in Table 2 and were processed and scaled using the *HKL* (Otwinowski & Minor, 1997) and *CCP4* (Collaborative Computational Project, Number 4, 1994) suites of programs. The location of the positions of both Dy³⁺ and Ho³⁺ were independently identified as the same site using the *SHELXD* (Sheldrick, 2008) subroutine in the program *SHARP* (de La Fortelle & Bricogne, 1997). The extreme non-isomorphism between all data sets suggested that the inclusion of multiple crystals in the phasing calculation would be detrimental. As a result, three-wavelength multiple-wavelength anomalous dispersion data sets for both the Dy³⁺ and Ho³⁺ derivatives were carefully collected and used for phasing in the absence of a

native reference data set. Phases were calculated using *SHARP* (de La Fortelle & Bricogne, 1997) and were further improved by solvent flattening using *DM* (Collaborative Computational Project, Number 4, 1994; Cowtan, 1994). Details of the structure determination and analysis will be published elsewhere.

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