Localization and characterization of the ligand-binding domain of the fibrinogen-binding protein (FgBP) of *Streptococcus equi* subsp. *equi*

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The group C streptococcus *Streptococcus equi* subsp. *equi* possesses a 498-residue major cell-wall-associated protein (FgBP) which binds horse fibrinogen (Fg), reacts with convalescent horse serum and protects against lethal *S. equi* challenge in a small animal model. In the present study, analysis of a panel of 17 purified N- and C-terminal FgBP truncates by ligand affinity blotting and SDS-PAGE revealed that the region required for maximum binding of Fg extended over the first half of the mature protein. The C-terminal two-thirds of this domain is predicted to be α-helical coiled-coil and the N-terminal one-third to possess non-coiled-coil single strands. Residues at the extreme N-terminus and within the coiled-coil region are both required for ligand binding. A high incidence of α-helical coiled-coil structure also seems to be responsible in part for the aberrant mobility of FgBP on SDS gels. The efficiency with which FgBP binds Fg from different animal species decreases in the order horse > mouse, pig > rat > sheep, dog, bovine, human. Binding to horse Fg is inversely related to temperature over the range 45–4°C and is independent of Ca2+ ions. MS analysis provided corroborative evidence that FgBP is covalently linked to the cell wall peptidoglycan.

**Keywords:** *Streptococcus equi* subsp. *equi*, fibrinogen-binding protein, ligand-binding domain

**INTRODUCTION**

Fibrinogen (Fg) is a major component of plasma, and the ability to bind Fg is common amongst cell wall proteins of pathogenic Gram-positive bacteria (Kehoe, 1994; Patti et al., 1994). Fg consists of three pairs of non-identical chains (α, β, γ) linked by disulphide bonds and has an M₀ of 340000. For the host, Fg has two main functions: polymerization into fibrin and mediation of platelet aggregation. Together, these result in the formation of platelet–fibrin thrombi (Doolittle, 1994). In Gram-positive bacteria, Fg binds to the variable N-terminal/central regions of many cell-wall-associated proteins, an interaction which appears to serve several different functions. For example, although the precise mechanism by which streptococci resist phagocytosis remains controversial, Fg binding to M protein does appear to play a role in resistance to phagocytosis for several strains of *Streptococcus pyogenes* (Courtney et al., 1997; Kehoe, 1994; Navarre & Schneewind, 1999; Poirier et al., 1989; Whitnack & Beachey, 1982, 1985). On the other hand, Fg binding to *Staphylococcus aureus* has been shown to be important in the initiation of infections of implanted medical devices and surgical wounds by promoting bacterial attachment (Herrmann et al., 1993; Moreillon et al., 1995; Vaudaux et al., 1989, 1995).

*Streptococcus equi* subsp. *equi*, a group C streptococcus, is the causative agent of strangles, a highly contagious and debilitating upper respiratory tract disease of the family *Equidae*. The disease is initially characterized by nasal discharge and fever followed by swelling and abscess formation in local lymph nodes (Timoney, 1993). In a recent publication, we described the puri-
 purification of a fibrinogen-binding protein (termed FgBP) from the cell wall of *S. equi* subsp. *equi*. FgBP was shown to have an apparent Mr 220,000 when analysed by SDS-PAGE, reacted with convalescent horse serum and was protective in a small animal model against lethal *S. equi* challenge (Meehan *et al*., 1998a). The sequence of the corresponding gene (*fbp*) was determined and was shown to encode a protein of 534 amino acids (Mr 38,344) which possessed some structural and sequence similarities to other streptococcal cell wall proteins. Of note were characteristic signal sequence and cell wall/membrane-spanning domains, two blocks (A and B) of degenerate repeated sequences and a high probability of α-helical coiled-coil structure over about 60% of the molecule. The sequence was identical, with the exception of six amino acids, to the M-like protein (SeM) described by Timoney *et al.* (1997) for a different isolate of *S. equi* subsp. *equi*. There is evidence to suggest that, like streptococcal M proteins, FgBP (SeM) is anti-phagocytic by limiting C3b deposition on the bacterial cell surface, that antiserum to the protein is opsonic and that Fg binding may enhance the ability of *S. equi* subsp. *equi* to resist killing by equine neutrophils (Boschwitz & Timoney, 1994a, b; Timoney *et al*., 1997).

Previously, we described the overexpression and purification of a recombinant FgBP truncate which lacked the signal sequence and C-terminal cell wall/membrane domain and showed that this recombinant protein behaved in an analogous fashion to wild-type FgBP in terms of Fg binding, seroreactivity and protective immunogenic potential (Meehan *et al*., 1998a, b). In the present communication, we describe the construction of a panel of FgBP truncates, show that the Fg-binding domain is localized in the N-terminal region of mature FgBP, and that the efficiency of Fg binding is dependent on temperature and source of Fg. We also provide evidence from MS analysis of recombinant and wild-type FgBP that the native protein may be covalently linked at its C-terminus to the streptococcal cell wall.

**METHODS**

**Bacterial strains, plasmids and growth conditions.** *Escherichia coli* XL-1 Blue (Strattagene) and plasmid pQE30 (Qiagen) were used for expression of the recombinant FgBP proteins. The virulent isolate, strain TW, of *S. equi* subsp. *equi* was obtained from the late Dr Paul Storm of International BV, The Netherlands (Meehan *et al*., 1998a). *E. coli* strains were grown at 37 °C in either LB medium or 2×YT medium (Lech & Brent, 1997). Where appropriate, media were supplemented with ampicillin (100 µg ml⁻¹) and IPTG (1 mM). *S. equi* subsp. *equi* was grown at 37 °C on Columbia base agar supplemented with 5% (v/v) horse blood or statically overnight in Todd–Hewitt broth supplemented with 0.2% (w/v) yeast extract.

**Recombinant DNA techniques.** Genomic DNA was isolated from *S. equi* TW by a modification of the method of Yu & Ferretti (1989) as described previously (Meehan *et al*., 1998a). Plasmid DNA was purified from *E. coli* by a modified alkaline lysis method (Felicello & Chinali, 1993). DNA restriction digestions, ligations and transformations were carried out by standard methods (Sambrook *et al*., 1989). Biochemical procedures. SDS-PAGE was performed using either 12.5% (w/v) or 7.5% (w/v) polyacrylamide separating gels and a 4.5% (w/v) polyacrylamide stacking gel (Laemmli, 1970). Samples were routinely heated for 3 min at 100 °C in Laemmli sample buffer (Laemmli, 1970) prior to electrophoresis. Proteins were detected by staining with Coomassie brilliant blue. Mr values were determined from the relative mobilities of 15 standard marker proteins (Gibco-BRL; BenchMark protein ladder).

Protein concentration was estimated by a modification (Dulley & Grieve, 1975) of the Lowry method using BSA as standard. Matrix-assisted laser desorption/ionization time-of-flight (MALDI-TOF) MS was performed as follows by Dr Len C. Packman, Dept Biochemistry, Cambridge, UK. FgBP samples (0.5 µl) dissolved in dilute (10–50 mM) sodium phosphate buffer (pH 7.2) or 0.1% (v/v) trifluoroacetic acid/50% (v/v) acetonitrile were mixed with 0.5 µl matrix solution [sinapinic acid (10 mg ml⁻¹) in 50% (v/v) acetonitrile containing 0.1% (v/v) trifluoroacetic acid] and dried onto sample slides at 25–30 °C in a fan-ventilated drying box. Samples were washed twice with 4 µl water to remove salts, then dried, and a further 0.5 µl matrix was added. As needed to improve the signal, samples were rewashed and further matrix solution was applied. An empirical dilution series of the protein sample (5–50-fold) was tested to ascertain the sample conditions necessary to give the optimal mass spectral signal. Mass analysis was carried out on a Kratos MALDI 4 time-of-flight mass spectrometer following the manufacturer’s recommendations. Masses were determined using rabbit aldolase (Mr 39,211) and horse myoglobin (Mr 16,951) as internal calibrants. Errors were minimized by averaging at least five mass determinations.

**Immunological and affinity procedures.** Western immunoblotting was performed as described previously (Caffrey *et al*., 1988) using 5% (w/v) dried skimmed milk as blocking reagent and rabbit anti-FgBP (1:10,000 dilution; Meehan *et al*., 1998a). Affinity-purified horseradish-peroxidase-labelled goat anti-rabbit IgG H + L (ICN) was used as localizing antibody and 4-chloro-1-naphthol as developing reagent. The procedure for FgBP immunoblotting after electrophoresis of SDS-PAGE gels was as described by Meehan *et al.* (1998a). For dot Fg-affinity blots, solutions containing FgBP truncates were subjected to doubling dilutions and 150 µl aliquots (containing 100–0.045 pmol) were transferred onto nitrocellulose using the Bio-Rad Bio-Dot apparatus. After the blots had dried, they were blocked, incubated with labelled horse Fg and developed as described for Fg-affinity electroblots (Meehan *et al*., 1998a). ELISA tests were carried out using standard procedures (Newell *et al*., 1988). For analysis of FgBP binding to Fgs from different animal species, wells of microtitre plates were coated overnight with 50 µl Fg solutions, ranging in concentration from 10 to 0.0195 µg per ml of 1 M sodium carbonate buffer (pH 9.6). Wells were blocked with 4% (w/v) dried skimmed milk, incubated with FgBP (1.25 µg ml⁻¹), followed by rabbit anti-FgBP antibodies (1:5,000 dilution) and peroxidase-labelled anti-rabbit IgG (1:5,000 dilution; ICN). For analysis of Fg binding at different temperatures and to determine the concentration of ligand resulting in 50% saturation of receptor, wells were coated with 25 ng FgBP1, blocked with 1% (w/v) BSA and probed with doubling dilutions of horseradish-peroxidase-labelled horse/human Fg as appropriate. For these experiments, blocking was performed in two stages: (1) incubation at 22 °C for 30 min followed by (2) incubation for 30 min at one of several temperatures (4, 22, 30, 37 or 45 °C). Subsequent washing and incubation steps were carried out at the chosen temperature. All ELISA tests...
were developed at room temperature using 3,3'5,5' tetramethylbenzidine as substrate and the A \textsubscript{\text{450}} was measured.

**Purification of FgBP.** FgBP was purified from *S. equi* as described previously (Meehan et al., 1998a). Briefly, this involved lysis of *S. equi* cells by passage through a French pressure cell (221 MPa) and harvesting of cell envelopes by centrifugation (45 000 g, 1 h, 4 °C). Cell envelopes were then digested overnight with mutanolysin (800 U ml\textsuperscript{-1}) in the presence of a cocktail of protease inhibitors. Following centrifugation (45 000 g, 1 h, 4 °C), proteins, including FgBP, which were released from the cell wall were recovered in the supernatant fraction. FgBP was then purified from this preparation by horse Fg-affinity chromatography (Meehan et al., 1998a).

**Expression and purification of FgBP truncates.** Oligonucleotide primers (Genosys) complementary to selected *fbp* sequence were used to amplify *fbp* fragments. Restriction enzyme cleavage sites (BamHI or XmaI) were engineered into the 5' end of all primers. A stop codon was also engineered directly after the restriction enzyme cleavage site at the 5' end of reverse primers. One hundred microlitre PCR reactions were performed in cloned pfu polymerase buffer (Stratagene) containing 5 U cloned pfu polymerase, 250 ng forward and reverse primers, 250 µM dNTPs and 100 ng pFP21 (recombinant pQE30 which expresses truncate FgBP1) or 500 ng *S. equi* genomic DNA. Amplification conditions consisted of 30 cycles of denaturation at 94 °C for 1 min, annealing at 55 °C for 1 min and extension at 72 °C for 2 min/1 kb of product followed by a final extension at 72 °C for 10 min. DNA was then purified using Wizard PCR preps (Promega), cleaved with appropriate restriction enzymes and cloned into pQE30. Recombinant plasmids were transformed into *E. coli* XL-1 Blue.

Hexahistidyl-tagged FgBP truncates were purified by metal chelate affinity chromatography using a modification of the method described by Meehan et al. (1998a). *E. coli* containing the recombinant pQE30 plasmids was grown, harvested and lysed by passage through a French pressure cell (221 MPa) as previously described. The soluble cytoplasmic/periplasmic fraction was applied to a nickel-iminodiacetic acid column, which was then washed extensively with binding buffer containing 40 mM imidazole until the A \textsubscript{\text{405}} of the effluent reached zero. Hexahistidyl-tagged FgBP truncates were recovered using an 80–400 mM gradient of imidazole in binding buffer. Most of the purified proteins were then extensively dialysed against several changes of 50 mM sodium phosphate (pH 7.2) containing decreasing concentrations (0–5, 0–2 and 0–1 M) of NaCl. Some of the smaller truncates tended to precipitate at lower NaCl concentrations. Accordingly, these were ultimately dialysed against either 0–2 M NaCl (FgBP6–8) or 0–5 M NaCl (FgBP9). Where necessary, purified proteins were concentrated by ultrafiltration using an Amicon PM10 filter.
RESULTS
Localization of the Fg-binding domain
In order to localize the Fg-binding domain of FgBP, DNA fragments of fbp were amplified by PCR and cloned in-frame into the expression vector pQE30. The resultant recombinant FgBP proteins (FgBP1–17; Fig. 1) were purified by metal chelate chromatography in the presence of protease inhibitors and analysed by SDS-PAGE (Fig. 2). The panel of recombinant molecules so constructed covered truncates lacking the wall/membrane anchor domain (FgBP1), ones lacking the wall/membrane anchor domain and increasing amounts of the putative coiled-coil domain (which includes the A and B repeat regions; FgBP2–9), together with ones lacking the wall/membrane anchor domain and an increasing number of N-terminal residues (FgBP10–17). All purified truncates gave a single dominant band following SDS-PAGE, with the exception of FgBP3, which routinely purified as two components (FgBP3a

![Fig. 2. SDS-PAGE of purified hexahistidyl-tagged FgBP truncates. Lanes 1–17, FgBP truncate numbers 1–17, respectively (see Fig. 1). Proteins have been stained with Coomassie brilliant blue, and the positions and \( M_r \) values of marker proteins are indicated to the left.]

<table>
<thead>
<tr>
<th>FgBP truncate</th>
<th>Loss of coiled-coil structure (%)</th>
<th>( M_r ) predicted from sequence</th>
<th>( M_r ) estimated by SDS-PAGE</th>
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<tr>
<td>Native FgBP</td>
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<tr>
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<td>40</td>
<td>23110</td>
<td>60000</td>
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</table>

* Estimated by MALDI-TOF MS.
and FgBP3b; see Fig. 2 and Table 1). MS, nucleotide sequencing and affinity binding data confirmed that the lower-\( M_t \) species (FgBP3b) was a degradation product lacking a further 30 amino acids from its C-terminus.

All purified truncates (including both FgBP3a and 3b) reacted with anti-FgBP antibodies as assessed by both Western and dot immunoblotting (data not shown). However, only the native protein and FgBP truncates 1, 2, 3a, 3b, 4, 5 and 10 bound detectable levels of Fg in corresponding Western affinity blotting experiments (see Fig. 3a). To place these observations on a more quantitative basis and to obviate demonstrable problems associated with SDS-induced reduction in Fg binding during SDS-PAGE and with less efficient electrotransfer of more tightly focused high-\( M_t \) truncates, dot-blotting experiments were performed using undenatured FgBPs (Fig. 3b). These semi-quantitative experiments confirmed the general trend observed during Western affinity blotting and revealed that in the absence of SDS (a) wild-type FgBP, FgBP1 (which lacks the wall/membrane anchor domain) and FgBP2–3a/3b (missing 113 and 182/212 amino acids from their C-termini, respectively) all bound similar levels of Fg to a first approximation, (b) FgBP4–6 (missing 255, 309 and 330 C-terminal residues, respectively) bound approximately 16-, 32- and 128-fold less Fg than FgBP1, (c) N-terminal truncates FgBP10–12 (which have 8, 11 and 19 residues, respectively, deleted from their N-termini) bound about 4-, 16- and 128-fold less Fg than their FgBP2 control and (d) the remaining truncates, i.e. FgBP7–9 and FgBP13–17 (see Fig. 1), bound no detectable Fg (Fig. 3b).

**Effect of Ca\(^{2+}\), temperature and Fg type on Fg binding**

Ca\(^{2+}\) has recently been shown to play a regulatory role in the interaction of Fg with several staphyloccocal Fg-binding proteins, e.g. ClfA, ClfB and Efb (Ni Eidhin et al., 1998; O’Connell et al., 1998; Palma et al., 1998). However, this does not appear to be the case for *S. equi* subsp. *equi* since preincubation of FgBP with a range of concentrations (in twofold dilutions) of Ca\(^{2+}\) (50–0·24 mM), EDTA (20–0·01 mM) or EGTA (20–0·01 mM) did not alter Fg binding to immobilized FgBP in ELISA tests (data not shown). Nor did the sequence of FgBP show the presence of a potential cation-binding EF-hand-like motif reportedly implicated in the binding of Fg to staphyloccocal ClfA and ClfB proteins (Meehan et al., 1998a; Ni Eidhin et al., 1998; O’Connell et al., 1998). In contrast, temperature did affect Fg binding to FgBP1 (and to other smaller Fg-binding FgBP truncates), the binding being significantly more efficient at 4°C than at 37°C (see Fig. 4a). Other assays performed with Fgs from a number of different mammals (see Fig. 4b) showed that FgBP bound the various molecules with the following approximate efficiencies with regard to horse Fg: viz. mouse and pig Fg, 50%; rat Fg, 25%; sheep, dog, bovine and human Fg, 3–8%. Additional ELISA experiments (not shown) revealed that (a) the concentrations of horse and human Fg resulting in 50%...
saturation of receptor at 4 °C were 40 ng ml−1 and 320 ng ml−1, respectively, and (b) at 45 °C, an approximately ninefold higher concentration of horse Fg was required to cause 50% saturation of receptor than at 4 °C.

**Aberrent mobility during SDS-PAGE**

Previous studies had shown that the $M_r$ values of native FgBP and recombinant FgBP1 as estimated by SDS-PAGE (220000 and 200000, respectively) were greatly at variance with those ($M_r$ 54597 and 49541, respectively) predicted from consideration of gene sequence (Meehan et al., 1998a). SDS-PAGE analysis of the current panel of FgBP truncates provides evidence that a high incidence of $\alpha$-helical coiled-coil structure may be responsible in part for this phenomenon (Table 1). Thus removal of residues from the N-terminus of the protein where there is a low probability of $\alpha$-helical coiled-coil structure and/or removal from the C-terminus of residues accounting for approximately 18% of the total coiled-coil structure generated truncates (FgBP1, 2, 10–15) in which the dominant protein species showed an apparent:predicted $M_r$ ratio (4–4.3) almost identical to that observed for wild-type FgBP. Removal of a further 12–22% of predicted coiled-coil structure from the C-terminus caused resultant truncates (FgBP16–17) to migrate with an apparent $M_r$ about 2-6–2-8 times that predicted, whereas removal of 39% or more of predicted coiled-coil structure from the C-terminus caused truncates (FgBP3–9) to migrate on SDS gels with an estimated $M_r$ close to that predicted from a consideration of sequence (see Table 1). Thus it would appear that (a) various FgBP truncates have the ability to migrate as apparent tetramers, trimers/dimers or monomers during SDS-PAGE and (b) this phenomenon can be correlated, at least in part, with the extent of predicted coiled-coil structure in the molecule. Whether these observations have any bearing on the macromolecular structure of FgBP in situ remains to be determined (see Meehan et al., 1998a, for further discussion of this issue).

**Wall association of FgBP**

Several lines of evidence strongly support the view that FgBP is covalently linked to peptidoglycan in situ. Firstly, it can only be released from the purified cell walls following digestion of the latter with the muralytic enzyme mutanolysin. Secondly, it possesses an LPSTG cell-wall sorting sequence (Meehan et al., 1998a). In staphylococci, there is formal evidence that such an amide bond is formed between Thr of the sorting sequence and the uncross-linked peptide crossbridge of peptidoglycan (Navarre & Schneewind, 1999; Ton-That et al., 1997). Finally and importantly, MALDI-TOF MS analysis of wild-type FgBP (purified following mutanolysin digestion) supports this general scenario. Thus the $M_r$ of FgBP1 estimated by mass spectrometry ($M_r$ 49540 ± 22) was in excellent agreement with that predicted from consideration of sequence (49541). However, the estimated $M_r$ of wild-type FgBP purified following mutanolysin digestion of the cell wall was determined to be 52440 ± 90. This is at considerable variance with that predicted for the primary translation product after N-terminal processing of the signal sequence at N37 ($M_r$ 54597; Meehan et al., 1998a) or after additional C-terminal processing at T382 in the cell wall sorting motif ($M_r$ 51467). On the other hand, it is consistent with a FgBP molecule (predicted $M_r$ 52486) which had been processed at both its N- and C-termini (i.e. N37–T382) and amide-linked at T383 to the di-alanyl crossbridge of the putative murein subunit AAK(A)QAMurNAcGlcNAc (Schleifer & Kandler, 1972). However, additional experimentation is required to formally prove this point.

**DISCUSSION**

Analysis of overlapping truncates has succeeded in defining for the first time the region within S. equi FgBP required for Fg binding. Maximum ligand binding requires a considerable proportion (about 52–68%) of the native (wall-associated) protein, and involves a region extending from the vicinity of the N-terminus (residue 37) to beyond residue 279, possibly as far as residue 352. Within this domain, individual residues at the N-terminus appear to be more crucial to Fg-binding activity than those positioned at the C-terminus. Thus removal of as few as 19 residues from the N-terminus results in dramatic (over 128-fold) loss of binding activity, whereas similar loss of Fg binding is only evidenced after the removal of about 150 residues from the C-terminal aspect of this (316-residue) domain. The presence in the various truncates of the N-terminal hexahistidyl affinity tag did not itself appear to compromise ligand binding as evidenced by the similarity in Fg-binding efficiencies of wild-type FgBP and recombinant derivatives such as FgBP1–3.

An N-terminal localization of Fg binding for FgBP is reminiscent of the reported binding of Fg to other streptococcal proteins such as the M proteins of serotypes 1, 3 and 5, and the FAI protein. However, sequence comparison of the relevant domains of these and other similar proteins has failed to reveal any regions of significant homology or recognized motif structures. It seems that Fg is capable of binding to some of the most variable regions of these proteins (Åkesson et al., 1994; Kehoe, 1994; Reichardt et al., 1997; Talay et al., 1996). In these respects, the mechanism of Fg binding to S. equi FgBP is clearly distinct from that displayed by the staphylococcal Fg-binding proteins ClfA, ClfB and Efb. These latter proteins have been shown to bind preferentially to one or more of the three Fg chains in a manner which probably involves Ca$^{2+}$ ions and a cation-binding EF-hand-like motif (McDevitt et al., 1997; Meehan et al., 1998a; Ni Eidhin et al., 1998; O’Connell et al., 1998; Palma et al., 1998).

Of probable relevance to the mechanism of Fg binding
are computer predictions of structure. These suggest that (a) residues 37–45 at the extreme N-terminus of wild-type FgBP (and FgBP1–9) exist in a random coil structure, whilst the balance of the molecule is largely (over 84%) \(\alpha\)-helical; and (b) 60% of the mature protein (from residues 166 to 475 of the precursor protein) possesses an extremely high probability (95–100%) of \(\alpha\)-helical coiled-coil structure, with zero to very low probability of such structures in the first 110 residues (residues 37–147; Meehan et al., 1998a). Ongoing structural studies of key FgBP truncates by circular dichroism (CD) have confirmed many of these predictions. Thus FgBP1 shows almost 100% \(\alpha\)-helical content by CD using the CONTIN secondary structure estimation procedure (Provencher & Glockner, 1981), and shows a molar ellipticity \(\langle[\theta_{222}]\rangle\rangle=106\) similar to that observed for other coiled coils (M. Meehan and others, unpublished data). Based on these observations and the ligand-binding profiles detailed above, it is tempting to envisage critical Fg-binding domain(s) located in the less structured non-coiled-coil N-terminal aspect of the FgBP dimer, with \(\alpha\)-helical coiled-coil sequences to the flanking (C-terminal) side providing, in part, a stabilizing structure which anchors and places constraints on the relative positioning of the non-coiled-coil single strands. In this model, excessive C-terminal truncation might serve to alter the gross structure of the FgBP molecule, either prior to or following interaction with its macromolecular ligand, leading to loss of effective interaction with Fg. Alternatively, Fg might interact with both the N-terminal and coiled-coil aspects of FgBP. Indeed, the coiled-coil region apparently necessary for full Fg binding (approx. 27 heptads, or 52 helical turns) is substantially greater than the minimum length generally considered necessary for the formation and stabilization of coiled-coil structures (three to four heptads; Su et al., 1994; Lau et al., 1984). Perhaps, as has been suggested for other systems, the coiled-coil structure of FgBP interacts in zipper-like fashion with the coiled-coil structure of Fg (Åkesson et al., 1994; Kehoe, 1994; Reichardt et al., 1997). This might serve to facilitate/stabilize interactions at the distal N-terminal site (or vice versa) and enhance binding efficiency. Both A and B repeated sequences of \(S.\) equi FgBP are present in the region of predicted coiled-coil structure (see Fig. 1). Of these, the degenerate B repeats are unlikely to be involved in binding since they are absent in truncate FgBP3, which displays wild-type Fg-binding efficiency. In contrast, the A repeats are clearly positioned within the region of the molecule required for full Fg binding. However, the A repeats are neither essential nor sufficient for Fg binding since they are absent in FgBP5, which still retains some, albeit weak, binding activity and are present in truncates (FgBP13–16) which are devoid of detectable activity. Whatever the precise mechanisms involved, it would appear that sites at the immediate N-terminus of FgBP and an extended region within the coiled-coil section of the molecule are both required for optimum Fg-binding activity. Biophysical studies should allow more critical appraisal of the role of coiled-coil structure in binding. The observation that FgBP binds ligand more efficiently at lower temperatures has also been noted for some class C M proteins of group A streptococci. This in turn has been correlated with the stability of the \(\alpha\)-helical coiled-coil structure and a tendency for these proteins to unfold to random coil monomers at higher temperatures (Åkerström et al., 1992; Nilson et al., 1995; Cedervall et al., 1997). A similar explanation seems likely for FgBP since ongoing CD analysis has shown FgBP1 to become progressively more unfolded in response to increased temperature over the range 8–80 °C and to be about 0, 16 and 30% unfolded at 8, 37 and 45 °C, respectively (M. Meehan and others, unpublished data).

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REFERENCES


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