Host blood proteins as bridging ligand in bacterial aggregation as well as anchor point for adhesion in the molecular pathogenesis of *Staphylococcus aureus* infections

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Fibronectin (Fn) and fibrinogen (Fg) are major host proteins present in the extracellular matrix, blood, and coatings on indwelling medical devices. The ability of *Staphylococcus aureus* to cause infections in humans depends on favorable interactions with these host ligands. Closely related bacterial adhesins, fibronectin-binding proteins A and B (FnBPA, FnBPB) were evaluated for two key steps in pathogenesis: clumping and adhesion. Experiments utilized optical spectrophotometry, flow cytometry, and atomic force microscopy to probe FnBPA/B alone or in combination in seven different strains of *S. aureus* and *Lactococcus lactis*, a Gram-positive surrogate that naturally lacks adhesins to mammalian ligands. In the absence of soluble ligands, both FnBPA and FnBPB were capable of interacting with adjacent FnBPs from neighboring bacteria to mediate clumping. In the presence of soluble host ligands, clumping was enhanced particularly under shear stress and with Fn present in the media. FnBPB exhibited greater ability to clump compared to FnBPA. The strength of adhesion was similar for immobilized Fn to FnBPA and FnBPB. These findings suggest that these two distinct but closely related bacterial adhesins, have different functional capabilities to interact with host ligands in different settings (e.g., soluble vs. immobilized). Survival and persistence of *S. aureus* in a human host may depend on complementary roles of FnBPA and FnBPB as they interact with different conformations of Fn or Fg (compact in solution vs. extended on a surface) present in different physiological spaces.

1. Introduction

Fibronectin (Fn) and fibrinogen (Fg) are multidomain glycoproteins that are major protein components of blood plasma. Fn is part of the fibrous extracellular matrix supporting endothelial cells in an insoluble fibrillar form, and it circulates as a soluble form in blood plasma (Henderson et al., 2011; Mezzenga and Mitsi, 2019; Singh et al., 2010) at a concentration of 0.2 to 0.4 g/L (Mosher, 2006). Fg is the most abundant coagulation factor at a concentration of 1.5–4.5 g/L (Ariens, 2013) in the blood. Because Fn and Fg are found in blood, they also form coatings on devices implanted in humans (Herrmann et al., 1988; Vaudaux et al., 1993).

*Staphylococcus aureus* is commonly found living on the skin and anterior nares of humans (Krismer et al., 2014; Lowy, 1998). When it gains entry inside a human host, *S. aureus* can lead to serious diseases like bacteremia and infective endocarditis. The incidence of *S. aureus* infections is rising (Naber, 2008; Tong et al., 2015), and mortality can be as high as 15–50 % (van Hal et al., 2012). *S. aureus* is one of only 11 bacteria and fungi listed as a “Serious Threat” in the Antibiotic Resistant Threats Report by the Centers for Disease Control and Prevention (2019). Therefore, it is critical to understand the mechanisms underlying *S. aureus* virulence so that we can develop novel therapies for these infections. For example, clinical studies of bloodstream infections have recently found higher binding affinity for immobilized Fn in *S. aureus* strains collected form human patients with infected cardiovascular devices (Hos et al., 2015; Lower et al., 2011) and infected endocarditis...
Interaction with host proteins is a critical first step in pathogenesis of *S. aureus* in the body. Binding between *S. aureus* and Fn and Fg was first reported several decades ago (Kapral, 1966; Kuusela, 1978). Humans ligands, like Fn and Fg, often play a key role in bacterial infections (Henderson et al., 2011; Vaudaux et al., 1989; Vaudaux et al., 1993). When *S. aureus* first enter the blood, the bacterial cells may aggregate together. This clumping is mediated by Fn and Fg, two of the most abundant host plasma proteins. Past work has primarily focused on the role of Fg because it is present at 10x the blood concentration of Fn (1.5–4.5 g/L vs. 0.2–0.4 g/L, respectively) (Lowe et al., 2004; Mosher, 2006). Yet, Fn-mediated cell aggregation may become predominant in areas where recruitment of Fn occurs, for instance at sites of injury wounds (Henderson et al., 2011). Soluble Fn is also important because it mediates *S. aureus* internalization in host cells (Sinha et al., 1999). In addition to clumping and internalization, adherence of *S. aureus* also depends on cell-wall anchored adhesins of the MSCRAMM (microbial surface components recognizing adhesive matrix molecules) type, which bind to host proteins, particularly Fn and Fg (Foster et al., 2014; Herrmann et al., 1988). The fibronectin-binding proteins A and B (FnBPA and FnBPB) are two key members of the MSCRAMM family (Foster, 2016). FnBPA and FnBPB are multidomain adhesins meaning that they can bind several mammalian ligands common in the blood, particularly Fn and Fg.

FnBPA and FnBPB consist of ~1000 residues that contain an N-terminal signal sequence responsible for secretion, an A region comprising subdomains N1, N2, and N3, followed by the repeat region with 10–11 domains, and a C-terminal cell wall and membrane-spanning regions containing the cell wall anchoring motif LPETG (Fig. 1). The Fg-binding site is located in the A-region near the N-terminus (Foster, 2016); whereas the Fn-binding site is located in the C-terminal repeat regions of FnBPA and FnBPB (Meehan et al., 2007; Schwarz-Linek et al., 2003).

Fg-binding occurs through a variant of the dock-lock-latch mechanism of the N2 and N3 subdomains (Foster et al., 2014; Keane et al., 2007; Ponnapati et al., 2003, Wann et al., 2000). The A region has also been reported to mediate cell-cell aggregation of bacteria (Geoghegan et al., 2013; Herman-Bausier et al., 2015). Fn-binding takes place through a tandem β-zipper mechanism by forming anti-parallel strand along the type-I modules at the N-terminus of Fn (Bingham et al., 2008; Schwarz-Linek et al., 2003). Another Fn-binding site has also been identified within the N2 and N3 subdomains of the A region of FnBPA (Burke et al., 2011).

*S. aureus* interactions with Fn and Fg have been determined to be associated with infections in humans (Piroth et al., 2008; Que et al., 2005). For example, adhesive interactions between Fn and *S. aureus* have been linked to biofilm-based infections of the blood and circulatory system (Hos et al., 2015; Lower et al., 2011; Xiong et al., 2015). Fn and Fg have also been linked to other aspects of *S. aureus* pathogenesis such as aggregation (Heilmann et al., 2004; Henderson et al., 2011; McAdow et al., 2011). Cell aggregation or clumping between neighboring bacteria may occur through FnBPA and FnBPB, or other MSCRAMM surface adhesins such as clumping factor A and B (Crosby et al., 2016; Dastgheyb et al., 2015; Geoghegan et al., 2013; Herman-Bausier et al., 2015; McAdow et al., 2011). Immune evasion and antibiotic resistance are enhanced when *S. aureus* form cell aggregates in the bloodstream (Crosby et al., 2016). Host ligand proteins like Fn and Fg may even bind to *S. aureus* forming a protective shield around bacteria cells (Crosby et al., 2016; Thomas et al., 2019).

In this study, we examine aggregation and adhesion of *S. aureus* in the presence of Fg and Fn, present in either a free (i.e. solubile) or immobilized form. A number of complementary techniques were used including optical spectrophotometry, flow cytometry and atomic force microscopy (AFM). Cell clumping and adhesion were evaluated under both physiological levels of shear and static conditions. Full-length *S. aureus* mutant strains of 8325-4 as well as *Lactococcus lactis*, which is a non-virulent, Gram positive surrogate that lacks adhesins for mammalian proteins including Fn and Fg.

Overall, the results demonstrate that both FnBPA and FnBPB facilitate cell-to-cell clumping through interactions with neighboring bacteria. This aggregation is enhanced by the addition of soluble Fg, soluble zinc, and especially by soluble Fn under physiological levels of shear. Normalized for the density of cell wall proteins (molecules per nm²), FnBPB presented a greater ability to clump compared to FnBPA. In contrast, adhesion to immobilized ligand was similar for both bacterial adhesions under physical stress. This finding suggests that these two distinct but closely related bacterial adhesins, have different functional capabilities to interact with host ligands in different settings (e.g., solubile vs. immobilized). Further, these results reveal that the conformation of host ligand (compact in solution vs. extended on a surface) impacts the interactions with these bacterial adhesins. This would mean that planktonic bacteria in blood interact more favorably with circulating host ligands like Fn and Fg through FnBPB. Whereas both bacterial adhesions may play a role when interacting with host ligands immobilized on a surface (e.g. part of the extracellular matrix or coating on an implant). This apparent specialization of each Fn-binding adhesin could play complementary roles in the onset and progression of infection in the human body.

2. Materials and methods

2.1. Bacteria strains and growth conditions

Cryopreserved *S. aureus* strains were grown at 37°C in triptic soy broth (TSB) supplemented with 10 μg/mL erythromycin and 0.5 % dextrose. *L. lactis* strains were grown in M17 broth supplemented with 5 μg/mL erythromycin and 0.5 % dextrose at 30°C. Both bacteria were grown in the presence of antibiotics since these strains were constructed...
by insertion of DNA fragments encoding antibiotic resistance in their plasmids. (Greene et al., 1995; Que et al., 2000, 2001). For clumping assays, bacteria were grown in their respective broth and temperature conditions and then diluted to an OD600nm of 1.0 in sterile TSB or M17 (Pestrak et al., 2020). For the AFM studies, bacteria were grown to an exponential phase, harvested and then washed in PBS. AFM data were acquired within two hours after harvesting the cell to ensure cell viability (Boonar et al., 2001).

2.2. Clumping assay with soluble host ligands

For studies under shear conditions, host proteins (Fn or Fg) were added to tubes with broth to a final concentration of 1 μg/mL. Since Fn and Fg have different molecular weights (i.e., Fn is ~440 kDa; Fg is ~340 kDa), a larger number of Fg molecules were tested compared to Fn molecules. A control tube was included with only bacteria (no host protein added). Tubes were incubating under shaking conditions at 200 rpm inducing an estimated shear ~8 dyn/cm² (i.e., pulmonary blood flow), as estimated according to (Ley et al., 1989). Aggregation of cells caused sedimentation of the clumps. The amount of clumping was estimated by removing 700 μl aliquot from the top of the tube and measuring the OD at 600 nm according to Kwiecinski et al. (Kwiecinski et al., 2019). The percentage of clumping was calculated as the percentage decrease from the OD at time zero. Percentage difference of clumping relative to control conditions (ligand free) in the presence of soluble host proteins fibronectin, and fibrinogen was determined by subtracting the OD values from the ligand free broth minus protein-containing broth then dividing by their average and multiplying by 100. Results shown are the means ± standard deviation of at least three independent experiments. For every experiment, an independent, fresh preparation of each strain was used. p-values were calculated using three independent experiments. For every experiment, an independent, protein-containing broth then dividing by their average and multiplying.

2.3. Atomic force microscopy with immobilized ligand

Force measurements were acquired with a Bioscope AFM and NanoSCOPE IV controller (Veeco/Digital Instruments) as described in Buck et al. (Buck et al., 2010; Oestreich et al., 2012). The results presented herein focus on adhesion to immobilized Fn because we recently examined molecular bonding of these same S. aureus and L. lactis strains to immobilized Fg (Casillas-Ituarte et al., 2019). For the experiments in this manuscript, an attached inverted microscope (Axiovert 200 M; Zeiss) was used to position the AFM tip over bacteria cells. A total of 167 different S. aureus and L. lactis cells from 20 independent cell cultures were probed with Si3N4 probes with nominal tip radius of 20 nm. The spring constant for each AFM tip was estimated by thermal tuning method (average = 0.094 nN nm⁻¹). The AFM tips were coated with Fn according to published protocols (Casillas-Ituarte et al., 2012; Lower, 2011). Briefly, a clean AFM tip was coated with Fn by immersion in a 100 μg/mL Fn PBS solution for 45 min, and then rinsed in PBS. Fn was deposited through this non-specific method, to mimic the conditions in the human body where these blood proteins coat surfaces in the circulatory system.

A total of 23 different tips were used. AFM measurements were conducted in PBS, at a single retraction velocity of 5.4 μm/s generating over 100,000 force curves. From these, 5,120 and 5,522 force curves were obtained from S. aureus expressing FnBPA or FnBPB, respectively. A total of 8,332 curves were collected from S. aureus expressing both FnBPA and FnBPB. Force curves for FnBPA and FnBPB present in L. lactis were 5,823 and 6,194, respectively. A total of 29,823 force curves were obtained for non-specific interactions between Fn and the surface of S. aureus DU 5883 and L. lactis pIL253 (negative controls). Other control experiments (e.g., with uncoated AFM tips) generated >40, 000 force curves.

To ensure specificity, only the final binding peak was included in the analyses in all the studies. Specific interactions between Fn-FnBPA and Fn-FnBPB were confirmed by monitoring successive unbinding events. A peak-to-peak distance (ΔL) of ~30 nm was indicative of the unfolding distance of multiple F1 repeats in Fn (Meadows et al., 2003). These ΔL measurements were confirmed at the beginning and the end of each experiment. Each AFM tip was used on only a few cells (~8) before being discarded when the characteristic unfolding patterns (ΔL values) were no longer observed. Negative controls included S. aureus DU5883 (fnbA fnbB double mutant) and L. lactis cells with an empty plasmid (pIL253). Force strength (or adhesion) was plotted as a histogram of force frequency to see the distribution of force values. This force frequency or frequency of binding was reported as percentage of force curves observed in a force range divided by total number of curves with adhesion events multiplied by 100.

2.4. Western ligand blots

Surface expression of FnBPs in L. lactis and S. aureus were determined by ligand affinity blotting by incubation with pure Fn as described in detail by Que et al. and Bisognano et al. (Bisognano et al., 2000; Que et al., 2000) and summarized in Casillas et al. (Casillas-Ituarte et al., 2012, 2019).

3. Results

Bacterial cell aggregation and adhesion interactions were examined for full-length FnBPA and FnBPB expressed individually (and collectively) in S. aureus mutant strains and surrogate host L. lactis. These reference strains are described in Buck et al. (Buck et al., 2010). A total of four different S. aureus strains were tested for the experiments presented herein: (1) FnBPA+ FnBPB-, (2) FnBPA- FnBPB+, (3) wild type strain expressing both FnBPA and FnBPB and (4) FnBPA- FnBPB-, a fnbA fnbb double mutant called DU5883. Three strains were tested in the L. lactis envelop: (1) FnBPA+ FnBPB-, (2) FnBPA- FnBPB+ and (3) pIL253, an empty vector as negative control.

3.1. Aggregation with soluble ligand under physiological shear stress

Clumping experiments were performed with bacteria in moving solution to simulate physiological levels of shear stress (~8 dyn/cm²). In the absence of host ligands, S. aureus formed clumps (see ligand free experiment in Fig. 2A). Clumping was limited to only ~12 % in the DU5883 mutant strain of S. aureus, which does not express FnBPA nor FnBPB (see Fig. 2A). These observations demonstrate that cell-cell aggregation depends on the presence of FnBPA and FnBPB.

Clumping was indeed observed in the two mutant strains of S. aureus that produced only FnBPA or FnBPB (Fig. 2A) in the ligand free
conditions. These findings suggest that aggregation can occur as a result of FnBPA-FnBPA and FnBPB-FnBPB interactions. Aggregation with these two strains was significantly greater than the wild-type strain. The relatively smaller aggregation in the wild-type strain is attributed to a lower level of expression of both FnBPA/B (see Western blots in Fig. 3C). There was a slightly greater aggregation for S. aureus that produced only FnBPA compared to S. aureus that produced only FnBPB. This could be due to differences in the number of protein present in the surface of the bacteria and/or to variation in the binding affinities between FnBPA-FnBPA and FnBPB-FnBPB. Semi-quantification of these proteins with Western blots (Fig. 3C) shows slightly greater concentration of FnBPA (~30%) suggesting that number density could be the reason for greater clumping.

Clumping was also evaluated in the presence of host proteins Fn and Fg. Addition of Fn increased cell aggregation relative to the control experiments, which lacked host proteins (Fig. 2A and B). Relative to the ligand free conditions, the addition of soluble Fn increase clumping in ~30% in both the FnBPA and FnBPB strains whereas minimum increase was observed in the wild type and DU 5883 strain (~4%). In the presence of Fg, there was an increase in clumping in all the strains. Yet, this increase was more pronounced in the wild type and DU 5883 strains (~50% to 60%) (Fig. 2B). Clumping in the wild-type and DU 5883 cells was not substantially enhanced in the presence of Fn compared to the control. The exception was in the presence of Fg, where an increment in clumping was observed. This clumping increase is attributed to the presence of the other surface adhesins on S. aureus (e.g., clumping factor A and B), which are known to bind to Fg (Ganesh et al., 2008).

To address this confounding issue (i.e., S. aureus proteins other than FnBPA and FnBPB that participate in clumping), we tested aggregation in a L. lactis model. This surrogate is Gram positive like S. aureus but lacks all known mammalian adhesins (Que et al., 2000). Clumping in the model surrogate L. lactis expressing FnBPA or FnBPB is shown in Fig. 3.

Aggregation was significantly slower in L. lactis compared to that in S. aureus (Fig. 2A and B). Longer incubation times (150 vs. 90 min) were needed to observed clumping in both FnBPA and FnBPB cells of L. lactis. A smaller number of proteins expressed in L. lactis compared to S. aureus could be the reason for this slower rate (see results for adhesion studies below). In the absence of FnBPA and FnBPB, no clumping was observed in L. lactis.

Consistent with the S. aureus clumping experiments, FnBPA-FnBPA and FnBPB-FnBPB interactions under physiological levels of shear stress appear to promote clumping in the FnBPA L. lactis cells. Yet, clumping in the FnBPA variant was not detected under the incubation time shown here (150 min). The FnBPA-variant of L. lactis finally clumped after 24 h of incubation under shear conditions without the additional of a bridging ligand (data not shown).

Similar to S. aureus aggregation, L. lactis also demonstrate similar aggregation enhancement in the presence of Fn in both FnBPA and FnBPB. Clumping in the presence of Fg was similar to that of the control, but only for the FnBPB variant. In all cases, FnBPB presented a greater ability to clump compared to FnBPA, which is an interesting difference from the experiments with S. aureus (e.g., compare Fig. 2A with B). Western blots (Fig. 5D) showed comparable levels of expression of these bacterial proteins (<11% greater for FnBPB). These findings suggest higher FnBPB-mediated intercellular clumping compared to FnBPA molecules. L. lactis expressing only fnbB also showed greater clumping for soluble Fg compared to FnBPA (Fig. 3A).

3.2. Aggregation with soluble ligand under static conditions

Flow cytometry was used to assess FnBPA and FnBPB role in aggregate formation under static conditions in both S. aureus and L. lactis (Fig. 4A and B). Cell clumping was observed in the presence and absence of host ligands. In S. aureus, clumping was almost largely absent in the wild type due to the low levels of expression of FnBPA and FnBPB as
In the presence of Fn, *S. aureus* expressing exclusively FnBPA or FnBPB presented an increased in clumping at least 50 % (ligand-free vs. addition of Fn; see Fig. 4A). Clumping in the wild type and the negative control (DU 5883) cells was not significantly enhanced in the presence of Fn. These results are consistent with the studies conducted under shear stress (Fig. 2A and B), that is, aggregation is affected by the presence of FnBPA and FnBPB. Yet a clear difference between the results from the studies conducted under static and shear stress, is the remarkable clumping enhancement observed in the *S. aureus* variant that produced only FnBPB (~150 % difference relative to ligand free conditions). Addition of Fg to the different *S. aureus* variants, (Fig. 4A) produced a slight increase in aggregation in the FnBPA and wild type cells.

Clumping in *L. lactis*, was enhanced in the presence of soluble Fn in both FnBPA and FnBPB variants relative to free ligand control (Fig. 4B). Addition of soluble Fg addition did not play a significant contribution in cell aggregation (Fig. 4B).

3.3. Adhesion to immobilized Fn under physical stress

Atomic force microscopy (AFM) was used to measure adhesion or binding forces associated with bacterial adhesion to immobilized Fn. AFM data for immobilized Fg was the focus on a recent paper by Casillas et al. (Casillas-Ituarte et al., 2019), and will be presented in the Discussion section. For the AFM experiments presented here, full length FnBPA and FnBPB was expressed in the surface of *S. aureus* and *L. lactis*. AFM was performed as described in prior work (Buck et al., 2010; Casillas-Ituarte et al., 2019). Unlike traditional binding assays (static adhesion studies, e.g., microtiter), AFM allows direct measurement of bond strength on ligand-receptor pairs through a dynamic process of pushing and pulling the linkages. Example binding force spectra are shown as insets in Fig. 5A and B. Binding events are represented as a series of sawteeth (Evans, 2001) where the final sawtooth represents the rupture or unbinding force between Fn on the AFM tip and Fn-binding receptors on a bacterium.

Force histograms for Fn binding to FnBPA or FnBPB in *S. aureus* and *L. lactis* are shown in Fig. 5A and B, respectively. For comparison, a Fn force histogram for wild-type *S. aureus* expressing both FnBPA and FnBPB is also shown (bottom panel in Fig. 5A). Fn-FnBPA and Fn-FnBPB interactions exhibited a median of ~40 % binding frequency (i.e., retraction curves that exhibit an adhesion event since not all the molecular interactions result in the formation of a bond) in both *S. aureus* mutant strains; whereas the wild-type generated a frequency of binding of 22 %. There was a lower frequency of binding observed for *L. lactis* with median values of 16 % and 8 %, respectively for FnBPA and FnBPB. Non-specific binding between Fn and the surface of *S. aureus* DU 5883 and *L. lactis* pIL253 (negative controls) exhibited binding frequencies of <10 % and <4 %, respectively. Differences in binding frequency of Fn-BPA or FnBPB expressed in *S. aureus* and *L. lactis* can be attributed to the different levels of protein expression in each type of bacterium (compare Fig. 5C, D).

The force spectra obtained from the different bacterial systems and summarized in Fig. 5A and B, were further analyzed by the worm-like chain model to estimate the number of Fn-FnBP pairs according to prior work (Casillas-Ituarte et al., 2012). For *L. lactis*, three or fewer pairs were involved in the measured interactions. For AFM experiments with *S. aureus* ~10 pairs were estimated. This is consistent with the Western blot and binding frequency analyses described above. Western blots showed smaller amounts of FnBPs in *L. lactis*. Binding frequency (or frequency of observing curves with adhesion events) was also lower for *L. lactis*.

Fn-FnBPA and Fn-FnBPB interactions presented large adhesion force peaks from ~300 pN to ~4 nN in the *S. aureus* envelop. Binding forces for *S. aureus* mutants were centered around ~800 pN, and ~1.3 nN for the Fn-FnBPA, and Fn-FnBPB, respectively. Forces from the wild-type *S. aureus* (expressing both FnBPA and FnBPB) presented a bimodal distribution with one population centered at ~670 pN and other at ~2.5 nN. Interacting forces between FnBPA or FnBPB with Fn in *L. lactis* were centered around ~460 pN and ~410 pN, respectively. These forces were significantly weaker compared to those in *S. aureus*. This is attributed to a smaller number of proteins present in *L. lactis* as described above.

4. Discussion

Here, we examined how FnBPA and FnBPB on the outer cell wall of *S. aureus* impact clumping and adhesion, keys steps in molecular pathogenesis, as described above. Despite their name, FnBPs have sites that bind to Fn as well as Fg (see Fig. 1), which allowed us to test the roles of both human ligands. Experiments were performed with *S. aureus* as well as *L. lactis* to unravel potentially confounding attributes of FnBPA vs. FnBPB. Furthermore, the use of *L. lactis* allowed us to overcome the problem of redundancy since a single adhesin could be expressed alone in a surrogate gram-positive bacteria host lacking other receptors for mammalian ligands like Fn and Fg (Que et al., 2001, 2005). FnBPP-mediated intercellular adhesion or clumping was tested under both static and shear conditions. Table 1 summarizes the results presented in Figs. 2 through 4. One of the most striking observations is the influence of flow on cell aggregation. Under stagnant conditions, the maximum observed clumping was 11 % (*S. aureus* FnBPA with addition of Fn ligand). Under dynamic conditions mimicking 8 dyn/cm² of shear, cell aggregation was consistently higher reaching a maximum observed value of up to 78 % (*S. aureus* FnBPA with addition of Fn ligand). Shear stress clearly enhances clumping suggesting that conformational changes in Fn and/or FnBPA/B take place under these conditions. These mechanical deformations would result in an increased binding affinity and, hence clumping. That is, the forces created by the shear stress could partially unfold Fn and/or FnBPA/A exposing previously sequestered
regions and thus increase the likelihood of these molecules to interact productively to form a bond. Protein unfolding under shear stress and consequent enhanced aggregation have been described in other proteins (Dobson et al., 2017).

Homophilic aggregation between molecules from adjacent cells (ligand-free clumping between bacterial cells) of up to 56% was observed for both \textit{S. aureus} and \textit{L. lactis} expressing FnBP’s on their outer envelope. Lesser homo-aggregation for \textit{L. lactis} expressing FnBPs (Table 1) can be explained by the lower density of cell-wall proteins as confirmed by the Western blots (Fig. 5C vs D). This type of cell-cell clumping was also observed for \textit{S. aureus} in the absence of FnBPs (see data for DU5883 in Table 1), indicating that other cell wall proteins on \textit{S. aureus} (e.g., clumping factors A or B, ClfA/B; serine-aspartate repeat proteins D or E, SdrD/E; von Willebrand factor) may play a role in homo-aggregation. But, as shown in Table 1, FnBPA and/or FnBPB clearly play the major role in enhancing cell-cell aggregation, particularly under shear conditions. Cell-cell aggregation through FnBPA/B was also enhanced through the addition of soluble zinc; whereas removal of zinc with the divalent cation chelator EDTA decreased cell aggregation (see Supplemental Fig. 1). This finding is consistent with previous studies with zinc (Geoghegan et al., 2013; Herman-Bausier et al., 2015).

Homo-aggregation was also impacted by the different sequences of amino acids making up FnBPA vs. FnBPB. This was evident in the ligand-free experiments with \textit{L. lactis}. For instance, FnBPB mutants show enhanced clumping compared to FnBPA variants (Fig. 3A and 4B). FnBPA and FnBPB have a relatively low (~45%) sequence identity in the A region which has been previously identified as a possible site for cell-cell interactions (Geoghegan et al., 2013; Herman-Bausier et al., 2015; Jonsson et al., 1991). Our aggregation experiments suggest that the A-domain of FnBPB has a greater affinity for A-domains of FnBPB from adjacent cells compared to that observed for pairs of A-domains in FnBPA (see Fig. 3A). This might explain the reason clumping was
observed at a similar, low level (~3 %) for all \textit{L. lactis} under static conditions (Fig. 4A); whereas clumping was observed for only FnBPB variants of \textit{L. lactis} under shear conditions (Fig. 3A). It seems that interactions between adjacent A-domains on FnBPB molecules are more resilient than those between FnBPA molecules.

In terms of bridging host ligands, the addition of soluble Fn significantly enhanced clumping for both \textit{S. aureus} and \textit{L. lactis} strains expressing FnBPA and/or FnBPB (Table 1). Relative to ligand-free conditions, clumping increased by >100 % for \textit{L. lactis} expressing solely FnBPA or FnBPB on the cell wall (Fig. 3B). In \textit{S. aureus}, this increase was more modest (30 %; Fig. 2B) likely due to the confounding impact of other cell-wall adhesins able to bind to mammalian ligands like Fn.

As shown in Fig. 2B, the addition of soluble Fg significantly enhanced clumping for \textit{S. aureus} lacking FnBPs on their cell wall (DUS5883 mutant). Fg likely served as a bridging ligand between cell-wall MSCRAMMs like ClfA/B and SdrD/E, which are well-known to bind to this ligand (Foster et al., 2014). Fg-enhanced clumping was also observed for \textit{L. lactis} and \textit{S. aureus} expressing FnBPB/A/B (Table 1). This form of ligand-assisted clumping could be important in vivo since Fg is about ten times more abundant than Fn in the blood.

\textit{S. aureus} clumping in the presence of Fg likely involves interactions with the A region, which is the active Fg-binding site in FnBPA/B (Foster et al., 2014; Keane et al., 2007). Differences in the clumping affinity between FnBPA and FnBPB (see Figs. 2A, B, 3 A, 4 A and 4B) are attributed to the low sequence identity in this A region, as discussed previously for the cell-cell interactions. Studies of the diversity of the A domain of FnBPA and FnBPB from \textit{S. aureus} strains, have shown that there are at least seven distinct isoforms with 60–85 % sequence identity. Each distinct isoform binds to the same site in Fg although with a different affinity (Burke et al., 2016; Loughman et al., 2008).

Comparing the two host, blood proteins, significantly more clumping was observed in the presence of Fn compared to Fg (Table 1). This is likely due to the multivalent binding capacity towards Fn for both FnBPA and FnBPB. Each of these bacterial proteins are able to bind up to ten Fn molecules through the FnBR region (Bingham et al., 2008); whereas the A-region of FnBPA/B binds to a single Fg molecule (Foster et al., 2014). Therefore, Fn dimers could more readily act as a bridging molecule between FnBPs molecules on adjacent bacterium. It is also possible that a conformational change in Fn upon adhesion to one bacterial adhesin (Liang et al., 2016) could favor an attractive interaction with an adjacent adhesin.

Aggregation of bacterial cells is one of at least two key processes that govern the initiation of \textit{S. aureus} pathogenesis. Binding of \textit{S. aureus} to solid substrates such as internal tissue or implanted materials is the other key initiation step for infection. These binding reactions are often mediated through interactions between bacterial MSCRAMMs (e.g., FnBPA/B) and host ligands that are immobilized on surfaces.

### Table 1

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<td>78% (4%)</td>
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<tr>
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<td>FnBPB</td>
<td>44% (1%)</td>
<td>65% (11%)</td>
<td>55% (2%)</td>
</tr>
<tr>
<td></td>
<td>others</td>
<td>12% (1%)</td>
<td>12% (1%)</td>
<td>21% (1%)</td>
</tr>
<tr>
<td>\textit{L. lactis}</td>
<td>FnBPA</td>
<td>0% (3%)</td>
<td>25% (7%)</td>
<td>0% (3%)</td>
</tr>
<tr>
<td></td>
<td>FnBPB</td>
<td>16% (3%)</td>
<td>69% (8%)</td>
<td>8% (3%)</td>
</tr>
<tr>
<td></td>
<td>others</td>
<td>0% (2%)</td>
<td>0% (3%)</td>
<td>0% (3%)</td>
</tr>
</tbody>
</table>

*Clumping detected after 24 h.

Soluble Zn enhanced homo-aggregation under shear conditions.

Microtiter is commonly used to measure adhesion reactions involving bacterial cells, including \textit{S. aureus} binding to human ligands (Casillas-Ituarte et al., 2019; Peacock et al., 2000). While this is a straightforward technique it offers only an indirect measure of adhesion because it detects changes in the optical density of (dead) labelled cells on well plates. AFM, on the other hand, provides a means of directly probing biophysical forces and/or the mechanical stability of ligand-receptor bonds (on live cells). Furthermore, AFM is a more dynamic technique capable of pulling or tugging on ligand-receptor pairs. This is important for ligand interactions with MSCRAMMs as demonstrated above for the clumping experiments under static vs. shear conditions.

AFM adhesion data provided herein shows a bond strength centered at ~430 pN for immobilized Fn with FnBPA or FnBPB on \textit{L. lactis} (Fig. 5B). Strength of binding to Fg was slightly stronger for FnBPB compared to Fg-FnBPB; ~460 pN vs. ~410 pN, respectively. Because similar number of proteins contributed to the interaction, this difference in bond strength could be traced to the extra repeat of ~40 amino acids in FnBPA (see Fig. 1). Even single amino acid changes in FnBPA have been reported to change the binding affinity towards immobilized Fn in clinical isolates of \textit{S. aureus} (Hos et al., 2015; Lower et al., 2011).

The range of adhesion data shown in Fig. 5B is consistent with Fg-binding data that was reported in another publication for this same strain of \textit{L. lactis} expressing FnBPA with up to three amino acid substitutions in the repeat region (Casillas-Ituarte et al., 2019). Fig. 5B shows a narrow force distribution and small adhesion frequency (~20 %). This, along with an analysis with the worm-like chain model, indicate single ligand-receptor pairs for the AFM experiments with \textit{L. lactis}.

Stronger adhesion forces were acquired in \textit{S. aureus} (800–1300 pN, Fig. 5A). This is expected given the higher level of FnBPs in this species of bacteria (compare Fig. 5C vs. D). The magnitude of adhesion shown in Fig. 5A is consistent with previous AFM studies for immobilized Fn on \textit{S. aureus} expressing FnBPs (Buck et al., 2010). Stronger adhesion for \textit{S. aureus} likely originates through multivalent interactions (Casillas-Ituarte et al., 2012) with the repeat region (see Fig. 1) that can bind up to nine molecules of Fn (Bingham et al., 2008).

Focusing on the simpler, ligand-receptor interaction in the \textit{L. lactis} surrogate, the ~430 pN adhesion force on Fn is stronger than the 241 pN adhesion force (median value) for Fg binding to FnBPs on the same \textit{L. lactis} surrogate (Casillas-Ituarte et al., 2012). A different strength of binding for the two host ligands is not surprising since there is a different mode or mechanism of binding for each ligand (Fn vs. Fg). Binding to Fg is expected to take place through interactions with the A-domain of FnBPA/B (see Fig. 1).

Interestingly, these two host ligands respond quite differently to tensile loading on the ligand-FnBP bond. Under conditions comparable to physiological load, binding between single pairs of Fg and FnBPA...
reach strengths of greater than 1300 nN (Casillas-Ituarte et al., 2019; Millles et al., 2018). Furthermore, the bond strength between Fn-FnBP is found to be dependent on amino acid substitutions in the repeat region, a part of FnBP that does not directly interact with Fg. Casillas-Ituarte et al. (Casillas-Ituarte et al., 2019) attribute this to catch-bond behavior of Fg when it binds to FnBPs under high tensile force.

In summary, FnBP adhesins in S. aureus adhesins are capable of hydrophobic interactions with neighboring bacteria that leads to clumps. This aggregation is enhanced by soluble Fg, and particularly soluble Fn homophilic interactions with neighboring bacteria that leads to clumps. Under physiological levels of shear (Table 1). In general, FnBPB presents a greater ability to clump in the presence of solution host ligands compared to FnBP. Interestingly, when the host ligand was immobilized on a surface, both adhesins FnBPA and FnBPB presented similar strength of adhesion. This indicates a critical condition for the interaction of these bacterial adhesins is the conformation of the host ligand (e.g., soluble vs. immobilized). In previous studies, we found that small variations in the amino acid sequences of the bacterial receptor alter the strength of adhesion to immobilized Fn and Fg (Casillas-Ituarte et al., 2012, 2019; Lower et al., 2011). In this study, we demonstrate that host protein configuration also plays a role in the initial molecular pathogenesis of S. aureus.

5. Conclusions
S. aureus has evolved to interact with multiple components of the host to avoid immune response and to facilitate adhesion to surfaces of indwelling medical devices. We have shown that two closely related S. aureus adhesins, FnBPA and FnBPB promote clumping by intercellular adhesion, in addition to their well-known ability to adhere to Fg. FnBP-mediated clumping is affected by the different physiological conditions (static vs. shear) and by the presence of soluble host proteins, particularly Fn. We found that these bacterial adhesins have different capabilities to interact with soluble ligand. This might explain the reason that most clinical and reference strains of S. aureus express both of these two adhesins despite the fact that they bind to similar target ligands (Burke et al., 2011; Loughman et al., 2008). Perhaps FnBPB plays a role when S. aureus are in the bloodstream exposed to soluble ligand, whereas both FnBPs are important when S. aureus interact with immobilized ligands on a surface (e.g., extracellular matrix or foreign medical device). The specialization of FnBPB could also explain the reason that both adhesins cooperate in the induction of severe infections by S. aureus (Shinji et al., 2011). This could also mean that there are different regulatory mechanisms for these two genes allowing expression under different conditions.

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Declaration of Competing Interest
The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data
Supplementary material related to this article can be found, in the online version, at doi:https://doi.org/10.1016/j.micron.2021.103137.

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