Staphylococcus aureus Fibronectin-Binding Protein A Mediates Cell-Cell Adhesion through Low-Affinity Homophilic Bonds

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ABSTRACT Staphylococcus aureus is an important opportunistic pathogen which is a leading cause of biofilm-associated infections on indwelling medical devices. The cell surface-located fibronectin-binding protein A (FnBPA) plays an important role in the accumulation phase of biofilm formation by methicillin-resistant S. aureus (MRSA), but the underlying molecular interactions are not yet established. Here, we use single-cell and single-molecule atomic force microscopy to unravel the mechanism by which FnBPA mediates intercellular adhesion. We show that FnBPA is responsible for specific cell-cell interactions that involve the FnBPA A domain and cause microscale cell aggregation. We demonstrate that the strength of FnBPA-mediated adhesion originates from multiple low-affinity homophilic interactions between FnBPA A domains on neighboring cells. Low-affinity binding by means of FnBPA may be important for biofilm dynamics. These results provide a molecular basis for the ability of FnBPA to promote cell accumulation during S. aureus biofilm formation. We speculate that homophilic interactions may represent a generic strategy among staphylococcal cell surface proteins for guiding intercellular adhesion. As biofilm formation by MRSA strains depends on proteins rather than polysaccharides, our approach offers exciting prospects for the design of drugs or vaccines to inhibit protein-dependent intercellular interactions in MRSA biofilms.

IMPORTANCE Staphylococcus aureus is a human pathogen that forms biofilms on indwelling medical devices, such as central venous catheters and prosthetic joints. This leads to biofilm infections that are difficult to treat with antibiotics because many cells within the biofilm matrix are dormant. The fibronectin-binding proteins (FnBPs) FnBPA and FnBPB promote biofilm formation by clinically relevant methicillin-resistant S. aureus (MRSA) strains, but the molecular mechanisms involved remain poorly understood. We used atomic force microscopy techniques to demonstrate that FnBPA mediates cell-cell adhesion via multiple, low-affinity homophilic bonds between FnBPA A domains on adjacent cells. Therefore, FnBP-mediated homophilic interactions represent an interesting target to prevent MRSA biofilms. We propose that such homophilic mechanisms may be widespread among staphylococcal cell surface proteins, providing a means to guide intercellular adhesion and biofilm accumulation.

Staphylococcus aureus is a human commensal and opportunistic pathogen that causes both superficial and invasive infections (1, 2). This species is a major cause of infections associated with indwelling medical devices such as central venous catheters and prosthetic joints (1, 2). The ability to form biofilms on implanted devices results in infections that are difficult to treat with antibiotics because many bacteria within the biofilm matrix are dormant. This is compounded by the prevalence of strains that are resistant to multiple antibiotics (methicillin-resistant S. aureus [MRSA]) (3, 4). Consequently, understanding the molecular mechanisms leading to the formation of staphylococcal biofilms may contribute to the development of novel therapeutic approaches for combating biofilm-related infections.

Until recently, the accumulation phase of S. aureus biofilms was attributed solely to the elaboration of polysaccharide intercellular adhesin (PIA), also known as poly-N-acetylglucosamine (PNAG) (1, 2, 5). However, it is now clear that proteins that are covalently anchored to the cell wall by sortase (cell wall-anchored proteins) can also promote biofilm accumulation. In the cases of SraP, SdrC, and SasG, cell-cell interactions have been shown to be due to specific homophilic binding between protein molecules located on adjacent cells (2, 5).

The fibronectin (Fn)-binding proteins (FnBPs) FnBPA and FnBPB promote biofilm formation by clinically relevant MRSA strains, both community associated and hospital associated (6–8). Both FnBPA and FnBPB have N-terminal A domains that are structurally and functionally related to clumping factor A (ClfA) and the Staphylococcus epidermidis SdrG protein and bind to fibrinogen by a variation of the dock, lock, and latch (DLL) mechanism whereby conformational changes in subdomains N2 and N3 within the A region result in highly stabilized complexes (9–11). The C-terminal fibronectin-binding domain comprises tan-
dem repeats that are intrinsically disordered, resulting in an extended flexible “stalk” that projects the A domain from the cell surface (Fig. 1A). The biofilm-forming region of FnBPA was localized to subdomains N2 and N3 of the N-terminal A region, but accumulation was shown not to involve a DLL mechanism (6, 7). FnBP-promoted biofilms could involve direct homophilic interactions or binding of the proteins to surface-located receptors on adjacent cells (Fig. 1B) (2).

Atomic force microscopy (AFM) has provided valuable insights into the molecular basis of staphylococcal adhesion. Force spectroscopy with biospecific probes has been used to probe the localization and binding strength of adhesins, including FnBPs, down to the single-molecule level (12–16). Furthermore, the use of bacterial cell probes has enabled the quantification of cell-substrate and cell-cell adhesive forces at the whole-cell level (17–19). In this study, we explore the molecular mechanism of FnBPA-dependent cell-cell adhesion using these AFM techniques (20, 21).

Specifically, we address the following questions: how strong are intercellular bonds, how many FnBPA proteins do they involve, and is FnBPA-mediated intercellular adhesion achieved by means of homophilic interactions or ligand binding? We analyze the binding mechanism of full-length FnBPA expressed from a plasmid in S. aureus strain SH10020 defective in clumping factors (Clfs) A and B, and in FnBPA and FnBPB (here S. aureus FnBPA+ cells), as well as of the recombinant FnBPA A domain immobilized on model surfaces. The results demonstrate that FnBPA mediates specific cell-cell adhesion via multiple, low-affinity homophilic bonds that depend on Zn2+ ions and involve the A domain.

RESULTS

FnBPA is involved in fibronectin binding and in cell aggregation. We first confirmed that FnBPA adhesins were expressed on FnBPA+ bacteria by analyzing their ability to adhere to Fn-coated substrates at the microscopic scale. Optical microscopy showed that FnBPA+ cells adhered strongly to Fn-coated surfaces, unlike FnBPA− cells, which hardly adhered at all (Fig. 2). Consistent with published data, this shows that FnBPA promotes bacterial adhesion to Fn and that the adhesin is expressed appropriately on FnBPA+ cells described here.

We then studied the involvement of FnBPA in cell-cell adhesion (Fig. 3). Figure 3A to C shows that FnBPA+ cells resuspended in buffer were isolated, without any evidence for aggregation. Addition of 1 mM Zn2+ induced the formation of large aggregates, 5 μm to 5 mm in size (Fig. 3D to F). Cell aggregates were disrupted in the presence of 1 mM EDTA (Fig. 3G to I) but restored upon further addition of Zn2+ (Fig. 3J to L). Aggregation was much less pronounced in S. aureus cells expressing no FnBPA (FnBPA− cells; Fig. 3M to O). These results show that FnBPA mediates cell-cell adhesion via Zn2+-dependent interactions.

Force spectroscopy of FnBPA interactions. AFM-based force spectroscopy was applied to living bacteria and to purified proteins to investigate the forces driving FnBPA-mediated cell-cell adhesion (Fig. 4). We used single-cell force spectroscopy (SCFS) (22, 23) to quantify the adhesion forces between single S. aureus cells. Single FnBPA+ cells were attached on colloidal cantilevers coated with polydopamine, allowing us to record force-distance curves between the cellular probes and the edges of small cell aggregates adhering on solid substrates (Fig. 4A). Single-molecule force spectroscopy (SMFS) was employed to probe single FnBPA bonds on live cells (Fig. 4B). Recent work showed that the region required for biofilm formation by FnBPA localizes to residues 166 to 498 of the A domain (7). Recombinant full-length FnBPA A domain was covalently bound to AFM tips in a random orientation, and the modified tips were used to record force-distance curves on FnBPA+ cells immobilized in porous membranes. To probe single FnBPA-FnBPA bonds in the absence of any other interactions, forces were also measured between tips and substrates both functionalized with fully oriented recombinant A domains (Fig. 4C).

FnBPA promotes specific cell-cell adhesion forces. We measured the adhesion forces between two individual S. aureus cells. Adhesion forces, rupture lengths, and typical force signatures obtained for three representative pairs of FnBPA+ cells are shown in Fig. 5A and B (see also Fig. S1 in the supplemental material for data obtained on more cells). Many curves featured large adhesion

FIG 1  FnBPA-dependent biofilm formation. (A) Schematic representation of the S. aureus FnBPA protein. S, secretory signal sequence; the A region comprising N1, N2, and N3 subdomains involved in fibrinogen and elastin binding and cell-cell aggregation during biofilm formation; R, tandem repeats of fibronectin-binding domains; W, proline-rich cell wall-spanning region; SS, sorting signal comprising the LPXTG motif, membrane-spanning domain, and positively charged tail. (B) Role of FnBPA in biofilm formation. In a first stage, FnBPA proteins promote attachment to host plasma proteins on biomaterial surfaces (left part of the left cartoon). Then, FnBPA mediates cell aggregation and biofilm accumulation (right part of the left cartoon). Whether this is achieved by homophilic protein-protein interactions or by binding to other ligands on adjacent cells is not yet established (right cartoons).

FIG 2  FnBPA-mediated adhesion to fibronectin. Optical (differential interference contrast) images showing the microscopic adhesion behavior of FnBPA+ cells (A) and FnBPA− cells (B), after a 2-h incubation on Fn-coated surfaces. Insets are representative images from a duplicate experiment.

Herman-Bausier et al.
Intriguingly, the same effect was observed for the interaction between FnBPA+ and FnBPA− cells (Fig. 5G and H; see also Fig. S2). This suggests that interaction between two FnBPA+ cells involves FnBPA proteins located on the two cell surfaces. So, our single-cell experiments show that FnBPA proteins on the S. aureus cell surface mediate specific, zinc-dependent, cell-cell interactions.

Binding strength of single FnBPA proteins on living bacteria.

How strong is a single FnBPA bond? To address this question, we analyzed the forces between recombinant FnBPA A domains attached to AFM tips and full-length FnBPA proteins on FnBPA− cells (Fig. 6; see also Fig. S3 in the supplemental material). Figure 6A to C (see Fig. S3 for more cells) shows that a large fraction (~50%) of the force curves recorded across three different cells featured single adhesion peaks with a moderate force of 125 ± 65 pN (mean ± SD from a total of 3,072 curves obtained on three cells). We believe that these forces originate from single FnBPA bonds as (i) adhesion was strongly decreased upon addition of EDTA (Fig. 6D to F; see also Fig. S3), when using S. aureus FnBPA− cells (Fig. 6G to I); or when using bare tips (see Fig. S3); (ii) the A domains were attached at low density on the tip; and (iii) the measured forces are in the range of the binding force typically reported for single-cell adhesion proteins (24, 25). Presumably, the larger adhesion forces sometimes detected (200 to 300 pN) are due to multiple FnBPA bonds. How does the measured bond strength compare with that reported for Fn-FnBPA bonds? Our ~125-pN force is stronger than the ~60-pN force measured for single Fn-FnBPA bonds at a similar loading rate (12, 15), suggesting that different mechanisms are involved. Strikingly, several groups have reported much larger adhesion forces for Fn-FnBP interactions, up to 6 nN depending on the strain investigated (13, 14, 16). However, the authors attached Fn at high density on the tip, meaning that many Fn-FnBP bonds were probed in parallel (up to 80). This is an important difference from the present work, in which FnBPA A domains were covalently attached at low density on the tip to favor single-bond detection.

Adhesion force maps indicate that the adhesins were largely exposed on the cell surface. Assuming that each adhesion event in the maps represents the detection of a single FnBPA, we find that the protein is exposed at a surface density of ~2,000 proteins/μm². In earlier work, the distribution of adhesins was mapped on the surface of staphylococcal cells. Immunogold electron microscopy was used to study the distribution of SsaF and UabB adhesins on Staphylococcus saprophyticus (26, 27). While the authors claimed that there was abundant labeling, the density was lower than that here, an effect that may reflect actual differences between species or variations in the sensitivity of the techniques. AFM force mapping was also used to map FnBPs on staphylococcal cells (15, 28). Lower et al. (15) revealed the spatial localization of putative FnBPs on S. aureus bacteria deposited on different substrates. Results suggested that the production and localization of FnBPP proteins may be induced by an external stimulus, such as the presence of Ftn on a surface.

Do the observed rupture lengths, 98 ± 44 nm (mean ± SD; 3 different cells), compare with the lengths of the probed molecules? As the full-length FnBPA protein and its His-tagged A domain are 948 and 481 amino acids in length (6, 7), respectively, and assuming that each amino acid contributes 0.36 nm to the contour length of a fully extended polypeptide chain (29), the lengths of fully extended proteins and A domains are expected to be 341 nm and 173 nm, respectively. Full extension of the A domain-FnBPA peptide is consistent with the notion that FnBPA interactions require Zn²⁺ (Fig. 5C and D; also see Fig. S2 in the supplemental material), that EDTA dramatically decreased the adhesion probability, suggesting that multiple bonds were involved. However, adhesion steps were often seen before rupture of the main adhesion peak, as well as small variations in cell-cell contact area. Discrete rupture may be induced by an external stimulus, such as the presence of Fn on a surface.

Do cell adhesion forces involve FnBPA proteins? We found that EDTA dramatically decreased the adhesion probability (Fig. 5C and D; also see Fig. S2 in the supplemental material), consistent with the notion that FnBPA interactions require Zn²⁺ ions (7). As can be seen in Fig. 5E and F (see also Fig. S2), we also measured the forces between S. aureus FnBPA− cells that lack FnBPA. Most adhesion events were abolished, indicating that the large adhesion forces on FnBPA− cells involve FnBPA proteins.
complex should therefore lead to a length of about 500 nm, which is much longer than what we observed. This means that the bond ruptures before complete unfolding of the proteins and, hence, that the latter are mechanically stable. This observation agrees well with earlier single-molecule results showing that SdrG, which shares strong structural similarities with FnBP, is not completely unfolded when subjected to large forces (30). We note that longer extensions were observed for cell-cell bonds (Fig. 5), supporting the notion that they involve pairs of full-length FnBPA proteins.

How many FnBPA bonds are involved in a cell-cell bond? As a rough estimate, this number may be inferred by comparing our single-molecule (Fig. 6) and single-cell (Fig. 5) forces and by considering the interaction area between two cells. The contact zone of two deformable spheres pressed on each other may be described by the Hertz model (31):

$$A = \frac{3FR}{4E^*}^{1/3},$$

in which $A$ is the radius of the contact area, $R$ is the effective radius ($1/R = 2/r$, where $r$ is the cell radius), $F$ is the applied load (here, 250 pN), and $E^*$ is the effective Young modulus $[1/E^* = (2 - 2\nu^2)/E]$, in which $E$ is the elastic moduli and $\nu$ is the Poisson ratio associated with the cell. Assuming that the $S. aureus$ cell radius $r$ is 0.5 μm, the Young modulus is 1.8 MPa (32), and the Poisson ratio is 0.3, we found an area radius of ~36 nm, thus yielding a contact area of ~0.004 μm². What is the number of interacting molecules in this area? Considering a surface density of ~2,000 FnBPA proteins/μm², as determined above from the force maps, the adhesion forces measured between two cells would involve about 8 FnBPA proteins in parallel on each cell. Given the variability of the data and the assumptions made in the model, this is in the range of the value obtained by comparing the adhesion forces measured for single molecules and single cells (~125 versus 800 to 2,000 pN). This leads us to believe that FnBPA-dependent cell-cell adhesion involves about 10 cumulative bonds, as also indicated by the sharp peak ruptures.

**FnBPA mediates low-affinity homophilic bonds.** To explain how FnBPA mediates cell-cell adhesion, two possible mechanisms were recently postulated, i.e., homophilic interactions or receptor-ligand binding (7) (Fig. 1B). Although our single-cell results are in favor of homophilic bonds, a direct demonstration for these was still lacking. We therefore measured the forces and dynamics of the interaction between purified FnBPA A domains in the absence of any other cell wall components (Fig. 4C). Force measurements between A domains revealed adhesion events with a mean adhesion force of 182 ± 78 pN (Fig. 7A, inset), which we attribute to single FnBPA-FnBPA bonds as A domains were uniformly oriented at low density on the two interacting surfaces. The measured strength is slightly larger than that between the A domain and full-length FnBPA on cells, an effect that could reflect differences in the orientation and accessibility of the molecules. The ~180-pN force is much weaker than the 2-nN force measured for SdrG-Fg bonds (30), indicating that FnBPA-mediated cell-cell adhesion does not involve a DLL binding mechanism. FnBPA-FnBPA bonds showed extensions (~50 nm) that were much shorter than those on live cells, suggesting that the A domains were hardly unfolded.

We then explored the dynamics of the FnBPA-FnBPA interaction, with the aim to assess the affinity of the bond. The forces needed to rupture homophilic bonds have been shown to depend on the loading rate, i.e., the rate at which the force is applied to the complex (24, 25, 33, 34). In agreement with this, we found that the
mean adhesion force ($F$) between two A domains increases linearly with the logarithm of the loading rate ($r$), as illustrated in Fig. 7A. The length scale of the energy barrier, $x_{p}$, was assessed from the slope $f_{p}$ of the $F$ versus $\ln(r)$ plot and found to be $0.2$ nm, i.e., in the range of values ($0.2$ to $1$ nm) typically measured by single-molecule AFM. Extrapolation to zero forces yielded the kinetic off-rate constant of dissociation at zero force:

$$k_{off} = \frac{x_{p}}{k_{T}} = 0.11 \text{s}^{-1}$$

This high dissociation rate, similar to that of homophilic bonds in cadherins (33) and in bacterial trimeric autotransporter adhesins (24), suggests that FnBPA homophilic bonds are highly dynamic. This short duration of adhesion may contribute to biofilm dissemination, by helping the bacteria to rapidly detach and colonize new sites.

We also studied how the adhesion frequency (i.e., number of curves with adhesion events) varies with interaction time, while keeping the loading rate constant (Fig. 7B). The binding frequency increased to reach a plateau corresponding to almost $100\%$ binding probability after only $0.5$ s, indicating that bond formation is fast. Similar fast bond formation was also observed with cadherins (33). Considering the interaction time needed for half-maximal probability of binding, $t_{0.5} = 168$ ms, we assessed the association rate constant, $k_{on} = \frac{t_{0.5}}{N_{A}} V_{eff} = 7.5$ $M^{-1} \text{s}^{-1}$, where $V_{eff}$ is the effective volume explored by the tip-tethered protein (approximated here to a half-sphere of $1$-nm radius) (35). We then estimated the equilibrium dissociation constant: $K_{D} = k_{off}/k_{on} = 15$ $mM$. The obtained $K_{D}$ value is much higher than that for FnBPA binding to fibrinogen ($\sim 1$ $\mu M$) (11) but in the range of that of homophilic interactions by trimeric autotransporter adhesins (24), thus indicating that homophilic FnBPA bonds have low affinity. This finding may have important biological implications. Low-affinity binding by means of FnBPA may represent the primary step in biofilm accumulation, enabling dynamic cell behaviors to occur, while subsequent higher-affinity binding would lead to firm cell-cell adhesion. In particular, the positively charged PIA polysaccharide, also known as poly-$N$-acetylglucosamine (PNAG), may enhance intercellular interactions by high-affinity multivalent binding with the negatively charged cell surfaces.

**DISCUSSION**

There is now considerable evidence that FnBPs participate in biofilm accumulation by *S. aureus* (6–8, 36), but the underlying molecular mechanisms are poorly understood. This study demonstrates that FnBPA mediates strong cell-cell adhesion via multiple, low-affinity homophilic bonds between A domains on adjacent cells. We speculate that homophilic interactions may be widespread among staphylococcal cell surface proteins, providing a means to promote intercellular adhesion and biofilm accumulation. Our methodology offers exciting prospects for the design of drugs or vaccines to inhibit protein-dependent intercellular interactions in MRSA biofilms.

The N2 and N3 subdomains of SdrC (37) have also been shown to promote cell-cell interactions, suggesting that homophilic interactions by these domains could be a general mechanism to promote the accumulation phase in *S. aureus* biofilms. Although the
FIG 6  Single-molecule force spectroscopy of FnBPA bonds on living bacteria. (A to C) Adhesion force maps (500 nm by 500 nm; gray scale, 500 pN) (A), adhesion force histograms (B), and rupture length histograms (C) together with representative force curves obtained by recording force curves in TBS with 1 mM Zn²⁺ across the surface of three *S. aureus* FnBPA⁺ cells using tips labeled with the FnBPA A domain. Shown in the insets in panel A are deflection images of the cells. (D to I) Force data obtained for an FnBPA⁺ cell (cell 3) in the presence of 1 mM EDTA (D to F) and for an *S. aureus* FnBPA⁻ cell (G to I). All curves were obtained using a contact time of 100 ms, a maximum applied force of 250 pN, and approach and retraction speeds of 1,000 nm · s⁻¹.
The significance of the Zn2⁺-dependent cell-cell interactions promoting homophilic cellular interactions, thus providing a driving force for biofilm formation. The biological importance of these interactions is supported by the observation that they are required for bacterial attachment to surfaces. The occurrence of FnBPA homophilic bonds correlated with the cell aggregation behavior, leading us to believe that these interactions represent an important driving force for biofilm formation. Unlike the very strong and stable DLL bonds, homophilic bonds show moderate strength and fast dissociation, a trait which may be important for biofilm dissemination. Several factors can lead to biofilm detachment, including mechanical stress like fluid flow, and detachment agents like enzymes or surfactants (1, 38). Together with these factors, the fast dissociation of the FnBPA bonds may contribute to cell detachment (isolated cells or cell clusters), therefore favoring colonization of new sites.

Our finding that zinc is required to form homophilic bonds is consistent with earlier reports showing that FnBPA (7) but also other staphylococcal adhesins, like SasG (39) and Aap (40, 41), promote zinc-dependent biofilm accumulation. It is therefore tempting to speculate that S. aureus has evolved these subdomains to promote homophilic cellular interactions, thus providing a general mechanism to favor biofilm accumulation. The biological significance of the Zn2⁺-dependent cell-cell interactions promoted by FnBPA can be called into question if the cation is limiting in vivo. The mammalian host restricts access to cations such as Zn and Mn that bacteria need for growth and proliferation in vivo, a phenomenon called nutritional immunity (42). An important host factor that contributes to this phenomenon is calprotectin, a Zn2⁺-binding protein that can reach high levels in infected tissue (43). However, successful pathogens such as S. aureus produce dedicated uptake machinery for cations (42). It should be noted that Zn2⁺ is present in the cytosol of mammalian cells and bacteria (44). S. aureus lyases host cells by secreting cytolytic toxins which release cytoplasmic contents. In addition, during biofilm development, some of the bacterial cells undergo autolysis to release DNA, which is an important component of the biofilm matrix (45). This altruistic action will also release bacterial cytoplasmic contents, including Zn2⁺. More than 3% of Escherichia coli proteins contain Zn2⁺ (46). The extracellular zinc-dependent metalloprotease aureolysin of S. aureus contributes to virulence in mice, indicating that it is active in vivo and presumably acquires its Zn2⁺ cofactor following secretion (47). We thus argue that the local concentration of Zn2⁺ at the early stages of biofilm development will be sufficient to support FnBPA-mediated aggregation. Finally, the expression of FnBPs has been shown to support biofilm formation on subcutaneous catheters during experimental infections of mice, arguing that adequate Zn2⁺ is likely to be present in vivo (36).

**Materials and Methods**

**Bacterial strains and growth conditions.** The S. aureus FnBPA⁺ strain (strain SH1000 clfA clfB fnbA fnbB) is defective in clumping factors A and B and fibronectin-binding proteins A and B (6). FnBPA⁻ cells were grown overnight in tryptic soy broth (TSB), washed once with TSB, subcultured into TSB at a 1:100 dilution, and allowed to grow to an optical density at 600 nm (OD600) of 0.4. The S. aureus FnBPA⁺ strain is a derivative of strain SH1000 clfA clfB fnbA fnbB carrying plasmid pFNBA4 expressing fibronectin-binding protein A from strain 8325-4 (7). For expression of strain FnBPA, FnBPA⁺ cells were grown overnight in TSB with chloramphenicol (10 µg/ml), washed once in TSB, subcultured into TSB at a 1:100 dilution, and allowed to grow to an OD600 of 0.4 in TSB plus chloramphenicol.

**Recombinant proteins.** Plasmid pQE30::FnBPA37–511 (10) was used as the template for inverse PCR with the phosphorylated primers 5’-TCA GAAACAAAGCAACTACAG-3’ and 5’-GGATCCTCCCTGATGC-3’ to eliminate DNA encoding the N-terminal His tag. The PCR product was treated with DpnI to eliminate template DNA, and following blunt-end ligation, the plasmid was transformed into E. coli XL1-Blue. The plasmid was extracted from E. coli and used as template for inverse PCR with phosphorylated primers 5’-CATCACCACCATCCTCACTA CCTAAAGTCGACCTGCAGCCAAG-3’ and 5’-TTAATTTTTCTCATTTC GGTACG-3’ to introduce an in-frame fusion with DNA encoding a C-terminal 6xHis tag. This PCR product was treated with DpnI, blunt-end ligation was carried out, and the plasmid was transformed into E. coli XL1-Blue. The C-terminally hexahistidine-tagged FnBPA A domain protein (residues 37 to 511) was expressed and purified by Ni²⁺ affinity chromatography as previously described (39).

**Adhesion assay.** To assess the adhesion phenotype of the bacterial strains, bacteria were incubated with Fn-coated substrates prepared as follows. Glass cover slips coated with a thin layer of gold were immersed...
overnight in an ethanol solution containing 1 mM 10% 16-
mercaptoundecanethiol acid–90% 1-mercapto-1-undecanediol (Sigma),
rinsed with ethanol, and dried with N2. Substrates were then immersed for
30 min in a solution containing 10 mg · ml⁻¹ N-hydroxysuccinimide
(NHS) and 25 mg · ml⁻¹ 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDC) (Sigma), rinsed 5 times with Ultrapure water (ELGA LabWater), incubated with 0.1 mg · ml⁻¹ of Fn from bovine plasma
(Sigma) for 1 h, and rinsed further with phosphate-buffered saline (PBS)
buffer. Fn substrates were incubated at 37°C in 200-μl bacterial suspensions
adjusted in Tris-buffered saline (TBS) buffer supplemented with 1 mM
ZnCl₂ to an OD₆00 of 0.3 to 0.4. After 2 h, the substrates were gently
rinsed by 3 consecutive washes in TBS buffer supplemented with 1 mM
ZnCl₂ and directly imaged using an inverted optical microscope (Zeiss
Axio Observer Z1) equipped with a Hamamatsu Ci0600 camera.

Aggregation assays. Aggregation phenotypes were directly observed
after cell resuspension in TBS buffer (pH 7.4) or TBS buffer supplemented
with 1 mM ZnCl₂, addition of 1 mM EDTA, and further addition of 1 mM
ZnCl₂. Aggregation levels were observed in test tubes, by optical micros-
copy at low magnification (Zeiss Stevi DV4 stereomicroscope;
Oberkochen, Germany) and at high magnification (Zeiss Axio Observer
Z1 equipped with a Hamamatsu Ci0600 camera; Oberkochen, Germany).

Cell-cell force spectroscopy. To probe bacterial aggregates with
single-cell probes, hydrophobic substrates were prepared by coating glass
coverslips with a thin layer of gold, immersing them overnight in a solution
of 1 mM 1-dodecanethiol (Sigma-Aldrich), rinsing them with etha-
nol, and drying them under N₂. Cells resuspended in TBS buffer plus
1 mM ZnCl₂ were deposited and allowed to adhere on hydrophobic sub-
strates for 2 h. Nonadhering cells were removed by gentle rinsing, and the
cell-coated substrates were attached, while avoiding dewetting, to the bot-
tom of a glass petri dish. Bacterial cell probes were obtained as previously
described (22, 23). Briefly, colloidal probes were prepared by attaching a
single silica microsphere (6.1-μm diameter; Bangs Laboratories) with a
thin layer of UV-curable glue (NOA 63; Norland Edmund Optics) on
triangular tipless cantilevers (NP-O10; Microlevers, Veeco Metrology
Group) and using a Nanoscope VIII Multimode AFM (Bruker Corpora-
tion, Santa Barbara, CA). Recombinant FnBPA A domain with a
C-terminal His tag was immobilized onto cantilevers and substrates as
follows. Silicon substrates were coated by thermal evaporation with a thin
layer of Cr (5 nm) followed by a thin layer of gold (30 nm). Gold substrates
and gold cantilevers (see above) were rinsed in ethanol, cleaned for 10 min
by UV-ozone treatment, rinsed in ethanol, and dried with N₂. They were
immersed overnight in a 0.1 mM solution of 100% HS-C₁₁⁻(EG₉⁻)₂-
OH thiols (ProChimia) and 1% HS-C₁₁⁻(EG₉⁻)₂-NTA thiols (ProChimia),
rinsed with ethanol, dried with N₂, and immersed in a 40 mM aqueous
solution of NiSO₄ (pH 7.2) for 30 min. Cantilevers and substrates were
then incubated in a 200-μl droplet of a 200-μg/ml solution of FnBPA A
domains for 1 h and rinsed and stored in PBS. Unless stated otherwise,
multiple force curves were recorded at 100-ms contact time, with a max-
imum applied force of 250 pN, and using a constant approach and retraction
speed of 1,000 nm/s.

Single-molecule force spectroscopy on model surfaces. SMFS mea-
surements using tips and substrates functionalized with FnBPA A do-

Supplemental material for this article may be found at
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Figure S1, PDF file, 1.3 MB.
Figure S2, PDF file, 1.3 MB.
Figure S3, PDF file, 1.6 MB.

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Homophilic Adhesion by FnBPA

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