The skin-colonizing commensal bacterium *Staphylococcus epidermidis* is a leading cause of hospital-acquired and device-related infections. Its pathogenicity in humans is largely due to its propensity to form biofilms, surface-adherent bacterial accumulations that are remarkably resistant to chemical and physical stresses. Accumulation-associated protein (Aap) from *S. epidermidis* has been shown to be necessary and sufficient for mature biofilm formation and catheter infection. Aap contains up to 17 tandem B-repeat domains, capable of zinc-dependent assembly into twisted, rope-like intercellular filaments in the biofilm. Using microscopic and biophysical techniques, we show here that Aap B-repeat constructs assemble further into zinc-dependent functional amyloid fibrils. We observed such amyloid fibrils by confocal microscopy during both early and late stages of *S. epidermidis* biofilm formation, and we confirmed that extracellular fibrils from these biofilms contain Aap. Unlike what has been observed for amyloidogenic biofilm proteins from other bacteria, which typically use chaperones or initiator proteins to initiate amyloid assembly, our findings indicate that Aap from *S. epidermidis* requires Zn$^{2+}$ as a catalyst that drives amyloid fiber formation, similar to many mammalian amyloid-forming proteins that require metals for assembly. This work provides detailed insights into *S. epidermidis* biofilm formation and architecture that improve our understanding of persistent staphylococcal infections.

*Staphylococcus epidermidis* is a critical component of the normal human flora that helps control the colonization and invasion of potentially dangerous microbial pathogens. However, *S. epidermidis* has emerged as a leading opportunistic pathogen due to its high prevalence on epithelial surfaces and ability to colonize prosthetic medical devices (1). *S. epidermidis* specifically is the leading cause of nosocomial infections and device-related infections (2, 3) and is, along with its other coagulase-negative relatives, the leading cause of bacteremia (4, 5). Although *S. epidermidis* infections are typically nonaggressive, they are extremely resistant to antibiotic therapy (6). Therefore, staphylococcal infections often require invasive treatment methods and frequently lead to chronic morbidity, mortality, and high healthcare costs (5, 7, 8). *S. epidermidis* pathogenesis and chronic persistence is primarily associated with its ability to form a biofilm (9, 10), a multilayered bacterial aggregation surrounded by an extracellular matrix (11). Biofilm formation occurs in three general stages: primary adherence to a surface by individual staphylococcal cells; accumulation into multicellular colonies through intercellular adhesion events; and formation of a mature biofilm through cycles of remodeling that create characteristic cellular towers separated by channels that allow access to nutrients. The biofilm is typically surrounded by a secreted extracellular matrix composed of macromolecular components that serve to anchor the staphylococcal cells together, including the polysaccharide poly-GlcNAc (PNAG)$^4$ as well as proteins, extracellular DNA, or teichoic acid, depending on the strain and growth conditions (12–14). Biofilm formation can occur through both polysaccharide-dependent and independent pathways, the latter mediated by protein–protein interactions (15). The protein Aap (accumulation-associated protein) is primarily responsible for protein-dependent intercellular accumulation of *S. epidermidis* biofilm formation (16), and it is also required for the polysaccharide-dependent mechanism (17). Rohde et al. (18) and Corrigan et al. (19) described a protein-based mechanism for staphylococcal biofilm formation.

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$^4$The abbreviations used are: PNAG, poly-N-acetylglucosamine; Aap, accumulation-associated protein; AUC, analytical ultracentrifugation; TEM, transmission electron microscopy; CFM, confocal microscopy; SasG, *Staphylococcus aureus* surface protein G; MBP, maltose-binding protein; HMBP, His-tagged MBP; ThT, thioflavin T; FA, formic acid; DTPA, 2-[bis(2-hydroxyethyl)amino]ethy]amino]acetic acid; Ni-NTA, nickel-nitritolriacetic acid; TSA, tryptic soy agar; ThS, thioflavin S; TSB, tryptic soy broth; IPTG, isopropyl 1-thio-β-D-galactopyranoside; SSR, summed square of the residual; Brpt, B-repeat.

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This article contains Figs. S1–S8 and Tables S1–S4.

1 Both authors contributed equally to this work.

2 Present address: Enable Injections, Inc., Cincinnati, OH 45241.

3 To whom correspondence should be addressed. Tel.: 513-803-7490; Fax: 513-636-5355; E-mail: andrew.herr@cchmc.org.

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tion that is independent of PNAG; in several strains of *S. epidermidis* or *Staphylococcus aureus*, the protein Aap or its ortholog SasG can mediate biofilm formation in the absence of polysaccharide secretion (18, 19). Indeed, 39% of biofilm-positive clinical isolates are PNAG-negative, whereas 90% are positive for Aap (15, 20, 21). Recent work by Schaeffer et al. (22) demonstrated the critical role of Aap in *S. epidermidis* infection in vivo. Under fluid shear, Aap-deficient strains formed significantly less biofilm than those expressing normal levels of Aap. A rat catheter model was then used to evaluate the biological implication of these results, which revealed that Aap, but not PNAG, was required for infection (22).

Aap is a multidomain protein consisting of an N-terminal export signal followed by the A-repeat region (11 partially-conserved 16-residue repeats), a lectin domain, the B-repeat region containing 5–17 conserved repeats of a 128-amino acid sequence, a proline/glycine-rich (P/G-rich) region that forms a highly-extended stalk that is resistant to compaction (23), and an LPXTG cell wall anchor motif, as shown in Fig. 1A. The N-terminal portion of Aap containing the A-repeat region and lectin domain (collectively called the A-domain) can be proteolytically cleaved to expose the B-repeat region, which can then initiate bacterial accumulation into microcolonies (18). The staphylococcal metalloprotease SepA is responsible for this proteolytic cleavage event, switching the role of Aap from A-domain–mediated surface adhesion to its namesake role in biofilm accumulation (24). Each B-repeat (Brpt) contains a 78-amino acid G5 domain and a 50-amino acid spacer domain (also called an E domain); the B-repeat sequences in Aap are highly conserved, with 83–100% sequence identity (25). The final repeat in the B-repeat region is composed of a single G5 domain without the spacer motif (Fig. 1A); this C-terminal half-repeat “cap” plays a role in stabilizing the protein (25). We have previously demonstrated that a single B-repeat domain with the half-repeat cap (Brpt1.5) will self-associate in the presence of Zn²⁺ to form an anti-parallel dimer, leading to a model for Zn²⁺-mediated protein-dependent intercellular accumulation between staphylococci in a nascent biofilm (25, 26). Subsequent work demonstrated that Brpt1.5 and longer B-repeat constructs can self-assemble in the presence of Zn²⁺ or Cu²⁺, whereas other metals (Mn²⁺, Co²⁺, and Ni²⁺) can bind to B-repeats but do not induce assembly (27). Recent work has shown similar Zn²⁺-dependent self-association behavior for the B-repeat region of SasG, the *S. aureus* ortholog of Aap (28, 29).

Full-length Aap contains 5–17 of these nearly identical B-repeats; Corrigan et al. (19) have demonstrated that at least five tandem B-repeats were required for *S. aureus* biofilm formation, suggesting that the biological function of SasG and, presumably, Aap relies on longer stretches of at least five consecutive B-repeats. Therefore, the focus of this study is to characterize a biologically-relevant construct of Aap consisting of five consecutive B-repeats and the C-terminal cap (called Brpt5.5; see Fig. 1A) and to determine the role of tandem B-repeats in *S. epidermidis* biofilm formation. In the presence of Zn²⁺, tandem B-repeats assembled into a range of oligomeric states, including highly-elongated fibers. We show that Zn²⁺-induced B-repeat fibers are functional amyloid fibers that assemble in a temperature- and time-dependent fashion. Importantly, fibers formed in vitro by recombinant, tandem B-repeats show at least partial resistance to Zn²⁺ chelation and acidification, suggesting that these highly stable amyloid fibers may play a critical part in rendering the biofilm resistant to physical or chemical stresses. We apply a newly-developed analytical approach to deconvolute the early-stage assembly of fibril intermediates using analytical ultracentrifugation. Finally, we show that amyloid fibers form during early and late stages of *S. epidermidis* biofilm growth, and we demonstrate that fibrils isolated from *S. epidermidis* biofilms are primarily composed of Aap. This is the first report demonstrating that Aap, an essential protein for *S. epidermidis* infectivity, forms amyloid fibrils; these findings reveal important mechanistic details regarding *S. epidermidis* infections responsible for extensive morbidity and mortality.

Results

Solution characterization of tandem B-repeats from Aap

To determine the functional relationship between tandem copies of the Aap B-repeat region and *S. epidermidis* biofilm formation, we have generated a construct containing the C-terminal five intact B-repeats, along with the C-terminal half-repeat (called Brpt5.5) (Fig. 1A). The design of this construct was based on the minimum number of B-repeats previously shown to support biofilm formation by SasG, an Aap ortholog from *S. aureus* (19). In addition, we utilized a Brpt3.5 construct, both in isolation and as an N-terminally His₆-tagged maltose-binding protein (MBP) fusion protein (called HMBP–Brpt3.5). The HMBP–Brpt3.5 construct showed more efficient expression and was more stable in solution, so it was predominantly used for initial experiments.

Far-UV circular dichroism (CD) was used to verify proper folding of the B-repeats in these constructs (Fig. 1B). Based on our previous studies of related Brpt1.5 constructs (25, 26) and the high sequence identity between B-repeats (Fig. S1) (25, 30), it was expected that the Brpt3.5 and Brpt5.5 constructs would have similar secondary structure content. Indeed, these constructs all contain high β-strand and coil content. The increased negativity around 200 nm observed for the Brpt3.5 and Brpt5.5 constructs may be due to greater random coil contribution from the increased proportion of B-repeat spacer motifs compared with G5 domains in Brpt3.5 and Brpt5.5 (i.e. Brpt5.5 contains five spacer motifs versus six G5 domains (~45% spacer), whereas Brpt1.5 contains one spacer motif versus two G5 domains (~33% spacer) (see Fig. 1A)). The CD spectrum for uncleaved HMBP–Brpt3.5 (Fig. S2) revealed a combination of α-helical, β-strand, and coil secondary structure, consistent with the α-helical content of MBP (31).

Sedimentation velocity analytical ultracentrifugation (AUC) was used to characterize each construct in solution under native conditions (Fig. 1C). Brpt3.5 and Brpt5.5 sedimented as monomers with very high frictional ratios (2.58 and 3.55, respectively; the frictional ratio *f*ₐ₀ is the frictional coefficient of the sedimenting species compared with that of an ideal sphere of identical volume). Although these constructs have similar sedimentation coefficients, the difference in frictional ratios is an important indication of the increased mass of
Brpt5.5 compared with Brpt3.5. Such high frictional ratios are indicative of highly-elongated global conformations, such as what might be expected based on our prior characterization of Brpt1.5 (25, 26). Furthermore, such an extended conformation for tandem B-repeats, in conjunction with the extended Pro/Gly-rich stalk (23), makes Aap well-suited to project itself from the S. epidermidis surface in order to more easily interact with adjacent cells and surfaces and to avoid steric hindrance from other cell wall–anchored proteins. The HMBP–Brpt3.5 construct also sedimented as a monomer but had a significantly increased sedimentation coefficient due to the additional 42-kDa mass contributed by the more globular HMBP fusion tag (Fig. 1C).

**Tandem B-repeats assemble into multiple higher-order species in the presence of Zn$^{2+}$**

We previously reported that shorter Aap B-repeat constructs, Brpt1.5 and Brpt2.5, specifically self-associate to form dimers in the presence of Zn$^{2+}$ (25–27, 30). These data indicated that tandem B-repeats self-associate in a modular fashion, with each B-repeat capable of forming an adhesive contact with another B-repeat in the presence of approximately two Zn$^{2+}$ ions, and that longer B-repeats constructs can dimerize at lower free Zn$^{2+}$ concentrations (25, 32). This phenomenon is known as the chelation effect, in which the binding of Zn$^{2+}$ and the resulting self-association at the first site in each of two multisite protromers reduces the entropic penalty for neighboring sites to self-assemble in the presence of Zn$^{2+}$. Thus, the effective free Zn$^{2+}$ concentration required for assembly of the entire multirepeat protein becomes progressively lower as the number of repeats increases. Prior biophysical work on B-repeat constructs has focused on shorter constructs that were suitable for detailed thermodynamic analyses; these short constructs were limited to monomer–dimer equilibria (25–27, 30). For the longer Brpt3.5 and Brpt5.5 constructs described here, self-association in the presence of Zn$^{2+}$ could be significantly more complicated than a simple monomer–dimer equilibrium, due to the potential for multiple adhesive interactions within a stretch of three to five intact B-repeats. Indeed, we observed a dramatic change in the sedimentation coefficient distribution for HMBP–Brpt3.5 in the presence of Zn$^{2+}$ compared with monomeric apo-HMBP–Brpt3.5 (Fig. 2, A and B), consistent with the formation of a wide range of very large oligomeric states (Fig. 2C). In addition to the reaction boundaries visible in the 0–40 s$^*$ range, the increasing trend at 40 s$^*$ indicates the presence of even larger aggregated species. This aggregation behavior is quite distinct from previously observed oligomerization of B-repeat constructs in the presence of Zn$^{2+}$ (Fig. S3). Although the HMBP fusion tag alone is able to assemble in the presence of Zn$^{2+}$ (Fig. 3), the formation of such enormous aggregates of HMBP–Brpt3.5 in the presence of Zn$^{2+}$ is unexpected and suggests that a distinct mode of assembly or aggregation is occurring.

**2D size–and–shape sedimentation analysis indicates formation of fiber-like species**

A recently developed AUC analysis method was used to better resolve the multiple sedimentation boundaries observed for HMBP–Brpt3.5 in the presence of Zn$^{2+}$ and to provide additional information on the size and shape of the sedimenting species. We utilized the 2D size–and–shape $c(s,ff)$ analysis in SEDFIT that characterizes each sedimenting species in terms of both sedimentation coefficient and frictional ratio (33). This approach is particularly useful when analyzing co-sedimenting species that differ greatly in shape and therefore experience very different degrees of drag (34). For these experiments, we used samples of 5 μM HMBP–Brpt3.5 alone or in the presence of 500 μM or 1 mM ZnCl$_2$. The samples were analyzed at both 25
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To characterize the nature of these extremely large, fiber-like B-repeat species, samples of HMBP–Brpt3.5 incubated with Zn$^{2+}$ for 2, 4, or 7 days were visualized by transmission EM (TEM) (Fig. 4A). Negative-stained TEM images revealed large assemblies of protein fibers. One possibility raised by the TEM images is that HMBP–Brpt3.5 can form amyloid fibers in the presence of Zn$^{2+}$. Amyloid fibers are highly stable protein fibers that have a characteristic β-strand–based fibril architecture (35). Formation of amyloid fibers was initially linked to protein misfolding implicated in disease states; however, a number of “functional” amyloid proteins required for specific cell processes have recently been characterized in organisms ranging from archaea (36) and bacteria to humans (37). Functional amyloid proteins from bacterial species, such as Escherichia coli, Bacillus subtilis, Enterobacteriaceae, and Pseudomonas spp., have recently been implicated in biofilm formation; specifically, these functional amyloid fibers are integral to the overall stability of the biofilm (38–41). To confirm that these HMBP–Brpt3.5 protein fibers were amyloid, samples were incubated with thioflavin T (ThT) and characterized by fluorescence spectroscopy. Thioflavin T is a small-molecule fluorophore that can intercalate within amyloid fibers, which restricts rotation about an internal bond and results in a dramatic increase in quantum yield and characteristic fluorescence emission at 482 nm (42, 43).

To ensure that the ThT fluorescence produced was truly from Zn$^{2+}$-dependent amyloid fibers, HMBP–Brpt3.5 was first pretreated with 10% formic acid (FA) to depolymerize any fibers that may have spontaneously formed (44). As seen with other amyloid proteins, HMBP–Brpt3.5/Zn$^{2+}$ produced minimal ThT fluorescence in the presence of FA (Fig. 4B). This effect was not due to FA-induced structural changes in the HMBP–Brpt3.5 monomer, because the secondary structure of the protein was unchanged upon addition of FA (Fig. S2). Upon removal of FA and incubation for 24 h, HMBP–Brpt3.5/Zn$^{2+}$ protein fibers demonstrated strong ThT fluorescence (Fig. 4B). However, after removal and incubation, HMBP–Brpt3.5 alone also produced a low level of ThT fluorescence, suggesting that

Figure 2. Sedimentation behavior of tandem B-repeats in the presence of Zn$^{2+}$. A, raw sedimentation velocity data for 5 µM HMBP–Brpt3.5 in the absence of Zn$^{2+}$, and B, in the presence of 3 mM ZnCl$_2$, both at 36,000 rpm and 20 °C. In both panels, scans 1–100 (5-h elapsed time) were loaded, with every 7th scan plotted. Note the dramatic increase in spacing between scans in B compared with A; the much faster-moving sedimentation boundaries of HMBP–Brpt3.5 in the presence of Zn$^{2+}$ compared with HMBP–Brpt3.5 alone indicate the sedimentation of very large species. C, sedimentation coefficient distribution of HMBP–Brpt3.5 alone (gray line) and HMBP–Brpt3.5 in the presence of 3 mM ZnCl$_2$ (blue line). The broad peak distribution in the presence of Zn$^{2+}$ indicates that HMBP–Brpt3.5 sediments as a mixture of assembled oligomers; however, resolution of individual species is difficult due to overlapping sedimentation profiles. The inset of C shows the same data analyzed by Wide Distribution Analysis in SEDANAL; note the x axis is on the natural log scale, and the y scale has been normalized by area. Here, the sample containing 3 mM ZnCl$_2$ shows some material around the 5–10 s$^{-1}$ range, with the majority of material sedimenting between 10 and 20 s$^{-1}$, in good agreement with the c(s) distribution calculated by SEDFIT. AU, absorbance units.
some fibers could potentially form in the absence of Zn\(^{2+}\). Based on quantification of monomeric HMBP–Brpt3.5 in the Zn\(^{2+}\)-free AUC data, the proportion of protein that forms fibers in the absence of Zn\(^{2+}\) is 3% or less (Figs. 1 and 2). A separate construct was additionally tested to better replicate the natural setting of Aap tethered in dense tufts to the cell wall of S. epidermidis. This construct was Brpt3.5 (cleaved from HMBP) with a C-terminal Cys residue added (Brpt3.5–Cys). Under nonreducing conditions, a disulfide bond would link two Brpt3.5 molecules in a parallel fashion, similar to their orientation on the cell surface, while raising the local B-repeat concentration, a well-known factor in amyloidogenesis. In Fig. 4C, there is an increase in ThT fluorescence corresponding to overall protein concentration. Additionally, there is a dependence on local B-repeat concentration, with disulfide-linked samples (“D” for dimer) yielding higher ThT fluorescence than reduced samples (“M” for monomer).

To ensure that amyloid formation was not due to the presence of the His\(_6\)–MBP tag (or the C-terminal Cys tag), native Brpt5.5 (tag-free) was incubated with Zn\(^{2+}\) and then visualized by TEM. Brpt5.5 formed fibers (Fig. 4D) primarily of the “branched” morphology seen with HMBP–Brpt3.5 (Fig. 4A). This morphology resembles one often observed with light-chain amyloid (45) but that is also observable with Aβ peptide (46). Analyzing these Brpt5.5 fibers spectroscopically (Fig. 4E, green bars) revealed that incubation with amyloid-binding dyes...
led to increases in both ThT fluorescence (47) and Congo Red absorbance at 540 nm (Fig. 4E, red bars) (48). Because amyloid fibers have characteristic structures rich in β-strand, we examined the far-UV CD spectrum under conditions that promote these amyloid-like aggregates. It should be noted that based on X-ray crystallography and NMR studies (26, 30, 49), along with CD data presented here and elsewhere (25, 30), natively-folded B-repeats contain β-sheet and random coil. Importantly, how-

Figure 4. Amyloid properties of tandem B-repeat constructs in the presence of Zn^{2+}. A, negative-stained TEM image of HMBP–Brpt3.5 protein fibers generated 7 days postincubation with 500 μM ZnCl_{2}. B, fluorescence emission spectra of 10 μM ThT alone (black) or ThT in the presence of HMBP–Brpt3.5 with 500 μM ZnCl_{2} (red), HMBP–Brpt3.5 alone (dark blue), and HMBP–Brpt3.5 with 10% formic acid (cyan) after excitation at 432 nm. HMBP–Brpt3.5 without ThT (green) did not produce fluorescence. The HMBP–Brpt3.5 alone and HMBP–Brpt3.5 + Zn^{2+} samples were pretreated with 10% FA to remove aggregates, followed by dialysis back into 50 mM MOPS, pH 7.2, 50 mM NaCl, and addition of ZnCl_{2}. C, ThT fluorescence of cleaved Brpt3.5 tagged with Cys added to the C-terminus (Brpt3.5–Cys) at 1 mM ZnCl_{2}, there is increased fluorescence of samples according to overall protein concentration. Furthermore, disulfide-linked Brpt3.5–Cys dimers (0.5–5.0 mg/ml; D) show even higher fluorescence. These results demonstrate the dependence of amyloid formation on local B-repeat concentration. Triplicate samples were examined, with empty circles representing individual measurements and averages shown as filled bars with a standard deviation shown with error bars. Statistically significant increases in the ThT fluorescence of a ZnCl_{2}-containing sample over the sample without ZnCl_{2} at the same protein concentration is denoted by an asterisk, and a pound sign indicates a significant increase of Brpt3.5–Cys without ZnCl_{2} over buffer.

G, Brpt5.5 samples were tested for recognition by the anti-amyloid antibody OC using a dot-blot assay. Antibody binding was observed for samples incubated at temperatures and Zn^{2+} concentrations conducive of the CD spectral change to a minimum near 225 nm. Natively-folded Brpt5.5 (without Zn^{2+}) also showed binding, but this was lost when Brpt5.5 was unfolded. Two regions from the same dot-blot assay are shown in G, both after a 5-min exposure. The solid border indicates regions of continuity on the blot. RFU, relative fluorescence units.
ever, Fig. 4F demonstrates that when Brpt5.5 is incubated with 5 mM Zn$^{2+}$ at near-physiological temperature (40 °C), there is a significant change in the CD spectrum, resulting in a strong, broad minimum near 225 nm. This is similar to CD spectra observed with insulin fibrils (50) and glucagon fibrils (51). Together, these spectroscopic results and TEM observations are consistent with amyloid-like fibril formation by native, tandem B-repeats.

A number of antibodies have been raised against amyloid-forming peptides or proteins. Kayed et al. (52) reported an antibody (OC antibody), which could specifically recognize $\beta_{42}$ fibers, but not monomers or oligomers. Furthermore, it appeared to recognize generic amyloid fiber conformation(s) irrespective of protein sequence, as it was able to also detect amyloid fibers from islet amyloid polypeptide and $\alpha$-synuclein (52). Thus, we chose this antibody to test against Brpt5.5 amyloid fibers (Fig. 4G). By dot-blot assays, OC antibody recognized Brpt5.5 that had been incubated with Zn$^{2+}$ at temperatures that resulted in a significant change in the CD signal in Fig. 4F. Interestingly, OC antibody recognized native, monomeric Brpt5.5 as well. This suggests that conformations present in the amyloid fiber are also present in the natively-folded protein. Incubating Brpt5.5 in the absence of Zn$^{2+}$ and at a temperature high enough to unfold the protein resulted in loss of OC antibody binding. Collectively, Brpt5.5 and HMBP–Brpt3.5 both share features characteristic of amyloid fibers.

### B-repeat fiber assembly is time- and temperature-dependent

Metal ions, namely Zn$^{2+}$ and Cu$^{2+}$, play complex roles in the aggregation and amyloidogenesis of $\beta$-amyloid peptide important in Alzheimer’s disease (53–56). However, functional amyloid-forming proteins from bacteria assemble into amyloid fibers without a known requirement for zinc ions or other triggering molecules. Rather, these amyloid proteins often form as a result of the coordinated action of accessory proteins, including nucleator proteins and chaperones that prevent aberrant polymerization in the cytoplasm (38, 41, 57, 58). In other cases, amyloidogenesis can require proteolytic activity to release the amyloidogenic regions of the protein, such as with Bap (59). Thus, the mechanism of Zn$^{2+}$-dependent amyloid formation by the B-repeat region of Aap is of significant interest for comparison with other systems. We have therefore used a combination of HPLC, TEM, confocal microscopy, and AUC to analyze the assembly of Aap B-repeat amyloid formation as a function of time, temperature, and solution conditions.

To assess the effect of temperature on the rate and morphology of amyloid fiber formation, we used the HPLC quantification assay previously shown to effectively differentiate between monomer and oligomer or soluble fiber species (60). A relative time course for HMBP–Brpt3.5 fiber formation was established by incubating 10 $\mu$M HMBP–Brpt3.5 with 500 $\mu$M ZnCl$_2$ for 0, 2, and 6 h and 1, 4, and 30 days at both 20 °C (Fig. S5) and 37 °C (Fig. 5A). Samples were first centrifuged for 1 h at 13,000 rpm to remove any insoluble aggregates prior to separation by a C4 HPLC column; therefore, the elution profile only reports on monomer, oligomer, and/or soluble amyloid fiber species. The HPLC elution profile of HMBP–Brpt3.5 alone (Fig. 5A, black trace) gives a peak eluting at 28.5 ml, which corresponds to a single monomeric species, as shown by its sedimentation coefficient distribution (Fig. 1B). Upon addition of Zn$^{2+}$ to HMBP–Brpt3.5, samples gradually showed a shift in the distribution of peaks toward higher-elution volumes that represent higher-order oligomer or fiber species. As time progressed, the monomer peak decreased and shifted to the right as the oligomer peak increased. The transition from monomer to putative fiber species was more pronounced after 1, 4, or 30 days for samples incubated at 37 °C (Fig. 5A) compared with the samples incubated at 20 °C (Fig. S5), indicating accelerated HMBP–Brpt3.5 self-assembly at higher temperatures.

We verified that our observations were not an artifact of the His$_6$–MBP tag, using an orthogonal approach. Because the HPLC assay is limited to only oligomers and soluble fiber species, we used a turbidity assay that follows light scattering by large assemblies and aggregates, including amyloid fibers (61). With the Brpt5.5 construct, we observed significant increases in the turbidity of the samples containing Zn$^{2+}$ when incubated at 37 °C (Fig. 5C). These data demonstrate the Zn$^{2+}$, time, and temperature dependence of B-repeat fiber assembly using native, untagged B-repeats.

### B-repeat fibers are resistant to acid and chelator treatment

Our previous work on Brpt1.5 revealed that this shorter construct formed a reversible Zn$^{2+}$-dependent dimer that was sensitive to removal of Zn$^{2+}$ (via chelators such as DTPA) or to even a modest decrease in pH (from 7.4 to 6.0) (25). To assess the stability of mature amyloid fibers, we allowed HMBP–Brpt3.5 fibers to form in the presence of Zn$^{2+}$ over a period of 24 h or 4 days at 20 or 37 °C. We then treated the samples by addition of a sufficient volume of dilute HCl to lower the pH to 5, addition of 2 mM DTPA, or addition of buffer as a control. The samples were incubated for 2 h before loading onto a C4 reverse-phase HPLC column, without spinning out the fibrous aggregates. The elution profile for the 20 °C sample at 4 days resembled the 24-h samples (Fig. S5) in terms of the monomer versus oligomer distribution and showed evidence of limited remodeling upon incubation with DTPA or HCl. In contrast, the elution profile for the 37 °C sample (Fig. 5B) showed that the oligomer/fiber peak predominated and that it was more resistant to the action of DTPA or HCl. TEM was performed on each sample to confirm that fiber structure was maintained (Fig. S6). Furthermore, although some conformational rearrangement of the fibers may occur at lower temperatures and early time points (62), mature fibers at 37 °C are highly resistant to these environmental conditions.

The initial Zn$^{2+}$-dependent assembly of untagged Brpt5.5 was reversible with the addition of DTPA or HCl, as shown by sedimentation velocity AUC data in Fig. S7. These results coincide with the reversible assembly of Brp1.5 (25), suggesting tandem B-repeats still undergo an initial phase of reversible assembly. Brpt5.5 aggregates from Fig. 5C were examined by dynamic light scattering to determine their resistance against DTPA. Although a decrease in the average hydrodynamic radius of all particles present was observed within 15 min, the average particle size was still about 10 times that of initial, reversible
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**Figure 5.** HPLC and turbidity assays to monitor time and temperature dependence of amyloidogenesis. A, C4 reverse-phase HPLC elution profiles of 10 μM HMBP–Brpt3.5 alone (black) or incubated with 500 μM ZnCl₂ for 1 (red), 4 (blue), or 30 (green) days at 37 °C. Samples were centrifuged at 13,000 rpm for 1 h prior to HPLC separation to remove insoluble aggregates. B, samples of 10 μM HMBP–Brpt3.5 with 500 μM ZnCl₂ were incubated for 4 days at 37 °C prior to addition of Tris-buffered saline (red trace); sufficient diluted HCl to lower the pH to 5 (blue trace); or 2 mM DTPA (green trace). Separation by HPLC revealed that oligomer/soluble fiber species were maintained in the presence of both acid (blue) and DTPA (green) when compared with the buffer-treated control samples (red) at both temperatures. TEM images of these samples are shown in Fig. S5. C, samples of 5 μM Brpt5.5 incubated at 37 °C and followed by absorbance at 400 nm, showing turbidity in the presence of ZnCl₂ with maximal turbidity reached by 24 h. Buffer controls for each condition were also monitored. For clarity, only the buffer + 8 mM ZnCl₂ data are shown, which had the highest turbidity of all buffer conditions. D, investigation of the DTPA resistance of Brpt5.5 aggregates in 5 mM ZnCl₂ from C by dynamic light scattering. Very large particle sizes were observed in the two samples examined. After addition of 5 mM DTPA, measurements were taken at 15 min and 1 h. In each case, particles of very large size were still present. One sample was examined at 4 h, with an average hydrodynamic radius of 620 Å. Natively-folded Brpt5.5 samples (n measurements were taken at 15 min and 1 h. In each case, particles of very large size were still present. One sample was examined at 4 h, with an average hydrodynamic radius of 620 Å.

Zn²⁺-induced complexes (Fig. 5D, inset). Even at 1 and 4 h, the average particle size of the aggregates exceeded that of natively-folded Brpt5.5. This result suggests that whereas some higher-order oligomers or nonamyloid aggregate may have been reversed or solubilized by DTPA, there is a fraction of material that remained in an aggregate form, as indicated by the remaining turbidity. It is possible the DTPA-sensitive material had not yet developed into mature amyloid fibers or that a higher local concentration of the tandem B-repeats (e.g. as seen with HMBP–Brpt3.5 or with native cell wall-anchored Aap) is required for complete DTPA resistance. DTPA resistance is an important feature of Aap functional amyloid, in that we previously reported the addition of Zn²⁺-chelator could prevent biofilm formation but could not disrupt mature biofilms (25), consistent with the presence of highly-stable Aap-derived amyloid fibers in the biofilm.

**Amyloid fibers are structural components in S. epidermidis biofilms**

Although it has been well-established that Aap is critical for *S. epidermidis* biofilm formation (15, 16, 22, 25), the mechanism by which Aap promotes stable, long-term intercellular adhesion is not well-understood. Others have shown that bundles of Aap fibrils extend outward from the cell wall on planktonic *S. epidermidis* cells (63, 64), but these were presumably free-standing proteins attached to the cell wall rather than amyloid fibrils. We sought to explore the significance of our findings in the context of *S. epidermidis* biofilms. *S. epidermidis* strain RP62A biofilms were grown in tryptic soy broth (TSB) supplemented without additional Zn²⁺ (Fig. 6A) or with additional Zn²⁺ (Fig. 6B) and visualized by TEM. Large networks of extracellular fibrils were observed around clusters of cells in the biofilm. This morphology closely resembles the fiber morphology observed by native Brpt5.5 when examined at the same magnification (see Fig. 6G for Brpt5.5 incubated at 37 °C with 5 mM ZnCl₂, compared with Fig. 6H for RP62A biofilm + 20 μM ZnCl₂). These extracellular fibrils were observed in biofilms grown in TSB with or without addition of Zn²⁺, although the presence of Zn²⁺ increased the prevalence of the fibers. TSB contains Zn²⁺, which allows for Aap-based, Zn²⁺-dependent accumulation and biofilm formation (25, 26, 65). Addition of the Zn²⁺ chelator, DTPA, at the start of growth inhibits biofilm formation (25); by TEM, these *S. epidermidis* cells showed no evidence of the extracellular fibrous networks.
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seen in the mature biofilms (Fig. 6C). The addition of DTPA at the start of growth would prevent the initial reversible self-assembly of Aap B-repeat regions (Fig. S7), presumably preventing subsequent nucleation of amyloid. However, once a mature biofilm is formed, DTPA can be added with no resulting effect (Fig. 6D) (25), similar to the observation that adding DTPA to mature amyloid fibers formed in vitro does not result in depolymerization (Fig. 5B). The highly-resistant nature of these amyloid fibers would contribute to the strength and stability of the mature biofilm, as shown previously for other systems, including TasA from B. subtilis (40).

In Fig. 4B, we demonstrated that B-repeat fibers formed in vitro are sensitive to FA. Based on this result, Aap fibers in biofilms should also show sensitivity to FA. Indeed, addition of FA to mature biofilms demonstrated the ability to disrupt the majority of the extracellular fibers in the biofilm (Fig. 6E). Furthermore, we examined the effect of pH on biofilm stability. As with DTPA, we previously observed that mildly-acidic pH disrupts the reversible assembly of B-repeat proteins (25, 32), but low pH has no effect on pre-formed B-repeat amyloid fibrils (Fig. 5B). Consistent with these in vitro results, we observed that lowering the pH of an established biofilm to 5.0 had no effect on the extracellular amyloid fiber network (Fig. 6F).

A biofilm formation assay (Fig. 6I) was performed to complement the TEM observations and evaluate the role of Aap functional amyloid fibers in biofilm formation in a more quantitative way. We once again observed the ability of DTPA to inhibit biofilm formation, but only when present before biofilm formation; when DTPA or pH 5.0 buffer was added after biofilm maturation (t = 24 h), neither was able to disrupt the biofilm, likely due to the resistance of the amyloid fibers (Fig. 6D, F, and I). Furthermore, addition of FA to mature biofilm was able to significantly disrupt the biofilm, as expected due to its ability to depolymerize the functional amyloid (Fig. 6E). The correlation between depolymerization of amyloid fibrils and weakening of the biofilm structure is a key observation that supports the idea of functional
amyloid fibrils contributing strength and stability to the biofilms.

**S. epidermidis amyloid fibers are composed of processed Aap**

As a final characterization of the extracellular amyloid fibers we observed in *S. epidermidis* biofilms, we determined the composition of these fibers by a combination of SDS-PAGE and MS. Biofilms were collected as they were for TEM characterization (Fig. 6B). The biofilm mixtures were then centrifuged to separate the bacteria from extracellular material and media. After removing the supernatant, we used lysozyme to digest the polyglycine cross-links of the cell wall peptidoglycan. The soluble proteins released from the cell wall were examined by SDS-PAGE. We observed SDS-insoluble aggregate trapped in the well and stacking gel (Fig. 6). This aggregated material contained primarily Aap when examined by nanoLC-MS/MS after in-gel tryptic digestion (Tables S1 and S2). In addition to Aap, peptides from several cytoplasmic proteins were identified, but with sparse coverage, suggesting these proteins are present in small quantities and are likely irrelevant to the amyloid fibers. We observed this result in three separate biological samples, one of which was not exposed to lysozyme and thus contained only extracellular material and not cell wall-anchored material (Fig. S8 and Table S3 and S4).

Peptides were observed from the B-repeats and the lectin region of Aap, with one peptide including Leu-601, which is one of two SepA cleavage sites (the other being Leu-335) (24), suggesting the lectin domain is still attached to Aap. The Aap A-repeats also contain tryptic sites, so the fact that no A-repeat peptides were detected by MS/MS is highly suggestive that SepA cleavage occurred at Leu-335, downstream of the A-repeat region. This observation is in agreement with Rohde et al. (18), who demonstrated that proteolytic processing of the N terminus of Aap is required for biofilm formation, and with Paharik et al. (24), who observed SepA-processed Aap molecules that retained the lectin domain but not the A-repeat region in the context of biofilms. This suggests the lectin does not need to be removed for biofilm formation (accumulation and amyloidogenesis specifically) to occur, but that it is the A-repeat region that inhibits biofilm formation.

**Amyloid fibers form early in biofilm formation and correlate with DTPA resistance of biofilms**

To further explore the importance of Aap amyloidogenesis in the context of developing biofilms, we followed biofilm formation by *S. epidermidis* strain RP62A at distinct time points throughout the first 24 h of biofilm formation. The addition of ThT to the media during the initial inoculation of the bacteria allowed us to follow amyloid formation by CFM (Fig. 7A). In parallel, biofilms were stained with LIVE/DEAD fluorescent dye to ensure ThT was not affecting cell growth or biofilm formation (Fig. 7B). As early as 2 h post-inoculation, punctate ThT fluorescence was visible on planktonic cells. This suggests that fibers start to form to a limited degree prior to cellular accumulation or biofilm assembly (Fig. 7B). At 6 h, the formation of microcolonies, and therefore intercellular accumulation, begins to occur. At this stage, ThT fluorescence is located at the boundaries between associating cells, which is consistent with the role of Aap as the critical factor for intercellular adhesion. ThT fluorescence increases throughout the biofilm over time, with a predominance of fluorescence in the core of the biofilm with the highest cellular densities. Although ThT is regularly used to identify amyloid fibers, there are also other molecules that can interact with ThT (43, 66–69).

To provide further evidence that the increased ThT fluorescence observed by CFM is indicative of Aap-dependent fibers forming in the developing biofilms, we took advantage of the DTPA resistance of the amyloid fibers observed *in vitro* and in mature biofilms. Specifically, we tested the ability of DTPA to inhibit biofilm formation when added at distinct time points along a similar time frame. Addition of DTPA during the 1st h of incubation was able to prevent biofilm formation from occurring, and by 2 h, DTPA was significantly less effective. The DTPA resistance of the biofilm correlates with the time frame of amyloid formation in growing biofilms (Fig. 7A). To ensure that the lack of biofilm formation at early time points was not due to mixing of the cultures upon adding DTPA, a duplicate experiment (Fig. 7D) was performed where DTPA was not added but wells were mixed in the same way as in Fig. 7C. Each of the conditions was still able to grow very strong biofilms, similar to the untreated control ("RP62A"), indicating that the mixing does not significantly affect biofilm formation. Therefore, these results support our hypothesis that Aap is important both for intercellular accumulation, by the formation of reversible assemblies and subsequently amyloid fibers between bacteria, and for stabilizing mature biofilms due to the remarkable resistance of amyloid fibers to physical and chemical insults.

**Discussion**

Our previous work established that the B-repeat region of Aap can undergo Zn\(^{2+}\)-mediated self-association to form proteinaceous rope-like filaments between staphylococcal cells in the biofilm and that Zn\(^{2+}\) chelation was able to inhibit biofilm formation by both *S. epidermidis* and *S. aureus* (25–27, 30). This initial work was carried out with the short B-repeat constructs BrpT1.5 and BrpT2.5, which showed reversible self-association that could be inhibited upon addition of chelator or moderate reduction in pH. Given that staphylococcal biofilms typically undergo acidification over time, it was unclear how this Zn\(^{2+}\)-mediated self-assembly mechanism could be maintained within the biofilm *in vivo*.

The results presented here using a more biologically relevant construct (BrpT5.5) and a BrpT3.5 fusion protein (HM1B–BrpT3.5) demonstrate that the B-repeat region of Aap is capable of two distinct Zn\(^{2+}\)-dependent assembly processes, forming both reversible oligomers and functional amyloid fibers within biofilms. Although the formation of amyloid fibers in biofilms has been established in several bacterial species, the mechanism of Aap amyloid fiber assembly displays some unique features among this group of bacterial biofilm proteins. Unlike the amyloidogenic biofilm proteins curli (*E. coli* and *Salmonella* spp.) (57, 70), TasA (*B. subtilis*) (40, 58), or FapC (*Pseudomonas* spp.) (41) that use additional chaperones or initiator proteins to help control and initiate amyloid fiber assembly, our data suggest that Aap utilizes Zn\(^{2+}\) as a catalyst to drive amyloid fiber formation.
formation. This mechanism for metal-dependent amyloid nucleation is instead reminiscent of several mammalian amyloid-forming proteins, including $\beta$-amyloid (54, 71, 72), prions (73, 74), and $\beta_2$-microglobulin (75, 76). Our proposed mechanism of Aap amyloid assembly shares a few similarities to that reported for *S. aureus* Bap (59). The N-terminal region of this cell wall–anchored protein is cleaved and released into the local environment, where it can self-assemble into amyloid-like structures in the presence of low Ca$^{2+}$ concentrations or at acidic pH. Bap, like Aap, can therefore act as both a sensor and a scaffold protein (59). Aap requires cleavage of the N-terminal A-repeat region by staphylococcal proteases or human proteases to unmask the B-repeat region before intercellular contact and amyloidogenesis occur in the presence of Zn$^{2+}$ (18, 24).

Bap has a similar arrangement of tandemly repeated domains to Aap and SasG (but the domains differ from the A- and B-repeats found in Aap and SasG), and it seems to be critical for infection in a mouse catheter model (59). However, the *bap* gene has not been found in *S. aureus* or *S. epidermidis* human clinical isolates (59). A Bap homolog, Bhp, is present in human isolates of *S. epidermidis* (77), but no experimental evidence exists regarding its role in biofilm formation or ability to form amyloid fibers, with the exception of a study by Lembre et al. (78), who showed a six-residue peptide from Bhp was able to form amyloid fibers in vitro. The *bhp* gene was found in less than half of *S. epidermidis* strains isolated from prosthetic knee joint and hip infections, whereas *aap* was found in 89% of isolates (15).

A recently identified *S. epidermidis* protein, Sbp, was shown to form amyloid fibers both in vitro and when expressed in intracellular inclusions in *E. coli* (79). Wang et al. (79) also showed that an *sbp* knockout of *S. epidermidis* strain 1457 did not show thioflavin S (ThS) fluorescence by confocal microscopy nor did it form a biofilm, an observation consistent with the initial study by Decker et al. (80). The WT strain 1457 did show biofilm formation, as expected, along with ThS fluorescence indicative of amyloid-like fibers. However, the authors did not identify Sbp as a component of purified amyloid fibers derived from biofilms, so there is uncertainty whether Sbp directly forms amyloid fibers in the biofilm or whether Sbp plays an indirect role in the nucleation of Aap amyloid fibers. If the latter case were true, then genetic knockout of either *aap* or *sbp* would abrogate biofilms as well as eliminate ThT fluorescence.

**Figure 7.** Formation of amyloid fibers is well-correlated to DTPA resistance in *S. epidermidis* biofilms. A, biofilm formation by *S. epidermidis* was characterized 2, 6, 12, 18, and 24 h post-inoculation by confocal microscopy, staining with 10 $\mu$M ThT (cyan) over a bright field. In the 2-hr panel, a zoomed inset is shown with increased contrast and brightness to highlight punctate ThT fluorescence visible around planktonic *S. epidermidis* cells. ThT fluorescence is seen primarily at cell–cell junctions at the 6-hour time point. Mature biofilms (24 h) show ThT throughout the biofilm as a major structural component. B, *S. epidermidis* biofilms at 2, 6, 12, 18, and 24 h post-inoculation were stained in parallel with LIVE/DEAD stain (green/red) as a control (scale bar, 10 $\mu$m). Images were generated in Zen 2009 Light Edition. C, biofilm formation assays show DTPA resistance develops within 2–6 h, with full DTPA resistance reached within 24 h. D demonstrates these observations are, in fact, due to addition of DTPA and not mixing. The asterisk denotes statistically significant difference ($p < 0.05$) to DTPA added at $t = 0$ h, and the pound sign indicates no significant difference to RP62A without treatment, as determined by a two-tailed Student’s t-test. Three separate experiments were performed for data in C and D, each with four replicate wells. The averages from the three separate experiments are plotted as empty circles, with the average of those plotted as gray bars with error bars showing ± S.D. Images of the crystal violet–stained wells (after washing and drying) of two representative samples are also shown.
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cence. Until there are well-defined mutations identified in either Aap or Sbp that specifically eliminate interactions, it will be difficult to establish the exact role played by Sbp. For this reason, we utilized MS to confirm that the fibers we observed were indeed composed of Aap.

The molecular characteristics of diverse types of amyloid fiber architecture have been well-described in multiple studies using a combination of structural and biophysical techniques (81–89). AUC approaches have also been used to characterize mature amyloid fibers in solution (90, 91). However, there have been few studies about the assembly of early oligomers and proto-fibril intermediates that produce mature amyloid fibers (92–94), particularly in terms of resolving discrete intermediate species. For example, pre-amyloid oligomerization of transthyretin has been characterized by AUC but without resolution of individual species (95). We have applied the $c(s_{m0})$ analysis to sedimentation velocity AUC data to resolve the broad array of species present at early stages of Aap amyloidogenesis, including the oligomeric complexes and nascent fibers. This is one of the first uses of this 2D size—shape analysis approach to an amyloid-forming protein, and its ability to separate many species with differing sizes and shapes in solution holds promise for resolving assembly intermediates in other amyloid systems as well. A similar approach, based on the van Holde-Weischet method (96) and implemented in Ultrascan (97) (RRID:SCR_018126), designed to resolve size and shape information from sedimentation velocity data has been applied to mature amyloid fibers formed by Aβ peptides (98, 99), but here we examined amyloidogenesis of a much larger protein. The $c(s_{m0})$ analysis model is capable of distinguishing species of different size and shape that are sedimenting at the same rate (34), which can occur in amyloid assembly systems due to the complicated nature of the assembly process and the number of species to be resolved. This analysis works particularly well in systems such as this one with slow kinetics, which allows resolution of discrete assembly intermediates rather than simply showing broad reaction boundaries representing multiple species in rapid exchange. The $c(s_{m0})$ analysis of Aap presented here clearly discriminates between compact oligomers and fibers that show similar sedimentation coefficients due to the increased drag and resulting slower sedimentation of the highly-elongated fibers. This analytical approach could prove useful in providing mechanistic details for how amyloid nucleation occurs by this or other proteins.

Biofilm formation is the primary characteristic responsible for pathogenicity, and it contributes to antibiotic resistance in chronic infections caused by *S. epidermidis*. One of the biggest challenges with recurrent infections caused by biofilms is their highly adhesive and cohesive nature and their resistance to chemical and physical insults. Our data indicate that intercellular amyloid fibers appear early during the accumulation phase of nascent biofilms, and they continue to increase until they are ubiquitous throughout the mature biofilm. Using highly-purified tandem B-repeats in solution, we show that mature amyloid fibers are highly resistant to chemical stresses such as $\text{Zn}^{2+}$ chelation or acidic pH, suggesting that the amyloid fibers forming the intercellular network are one reason *S. epidermidis* biofilms are so resistant to harsh environmental conditions. We have previously shown (25) that addition of $\text{Zn}^{2+}$ chelators to staphylococcal cultures prevents biofilm formation. However, addition of these same chelators to pre-formed mature staphylococcal biofilms does not appreciably disperse the biofilm, which is now explained by the presence of resistant amyloid fibers between cells. Furthermore, we predict that homologous surface proteins containing tandem B-repeats in *S. aureus* (i.e. SasG and Pls) and other Gram-positive bacteria will form similar $\text{Zn}^{2+}$-dependent amyloid fibers between cells in biofilms. The data presented here provide new insights into the mechanism of *S. epidermidis* biofilm formation and infection.

Materials and methods

**Bacterial strains and media**

*S. epidermidis* strain RP62A (ATCC 35984) was purchased directly from ATCC as a glycerol stock and was cultured in TSB.

**Expression construct generation**

The Brpt3.5 construct (amino acids 1761–2223) of Aap (NCBI accession no. AAW53239.1 (10)) was PCR-amplified from RP62A genomic DNA and inserted into the expression vector pHisMBP–DEST (kindly provided by Dr. Artem Evdokimov) using Gateway technology, which adds an N-terminal hexahistidine–maltose-binding protein (His–MBP) fusion to the Brpt3.5 construct, with an intervening tobacco etch virus protease site. The Brpt5.5 construct (amino acids 1505–2223) of Aap (NCBI accession no. AAW53239.1 (10)) was synthesized by Life Technologies, Inc., GeneArt®, also containing an intervening tobacco etch virus protease site for removal of the His–MBP tag. The plasmids were then transformed into the *E. coli* expression cell line BLR(DE3) (Novagen). The Brpt3.5–Cys mutant was synthesized by Life Technologies, Inc., GeneArt®, as well, containing amino acids 1761–2223 of Aap (NCBI accession no. AAW53239.1 (10)), followed by a single cysteine residue inserted before the stop codon.

**Protein expression and purification**

One-liter cultures were inoculated with 10 ml of His–MBP-Brpt3.5/BLR(DE3) culture at an $A_{600}$ of 0.6–0.8 and then allowed to incubate overnight with shaking at 37 °C. Protein expression was then induced using 250 μM IPTG for 6 h at 25 °C. The cells were then harvested, resuspended, frozen, and thawed prior to lysis by French press. The cell lysate was centrifuged, and the soluble fraction was decanted onto a nickel-NTA gravity column. The Brpt3.5 protein was eluted by imidazole step gradient at 300 mM imidazole. Fractions confirmed by SDS gel to contain Brpt3.5 were then further purified by anion-exchange using a 5-ml anion Q fast-flow column (GE Healthcare) followed by size-exclusion chromatography using a Superdex 200 column (GE Healthcare). The presence of the His–MBP-fusion tag improved solubility and behavior in solution and was therefore left attached to enhance protein stability, except for initial CD and AUC experiments (Fig. 1).

The Brpt5.5 construct was induced overnight at 20 °C with 200 μM IPTG, lysed by sonication, then run over a homemade 50-ml Ni-NTA column containing 50 ml of IMAC-Sepharose
Fast Flow resin (GE Healthcare) charged with Ni$^{2+}$ and housed inside a XK 16/40 column. After elution by an imidazole linear gradient from 0 to 300 mM imidazole, the fractions containing Brpt5.5 were dialyzed into 20 mM Tris, pH 7.4, 300 mM NaCl and cleaved by TEV for at least 6 h under reducing conditions. Another 50-ml Ni-NTA column was run, this time collecting the flow-through (cleaved Brpt5.5). Finally, a Superdex 200 (GE Healthcare) column was run to separate remaining contaminants and truncations of Brpt5.5.

**CD**

Brpt3.5 and HMBP–Brpt3.5 samples were dialyzed into 10 mM Tris, pH 7.4, and 150 mM NaF. Far-UV CD spectra were obtained using an Aviv 215 spectrometer (Aviv Biomedical, Inc., Lakewood, NJ). The concentration of cleaved Brpt3.5 (fusion tag removed) was determined using the molar extinction coefficient of 16,390 M$^{-1}$ cm$^{-1}$ calculated by ProtParam. An additional sample of HMBP–Brpt3.5 was treated with 10% FA to depolymerize amyloid fibers, followed by dialysis of the sample into 10 mM Tris, pH 7.4, and 150 mM NaF. The concentration of Brpt5.5 was based on the molar extinction coefficient of 16,390 M$^{-1}$ cm$^{-1}$, and the spectrum was collected in 10 mM Tris, pH 7.4, and 50 mM NaF. Data were analyzed using the CDSSTR program on the online Dichroweb server with reference set 4 (RRID:SCR_018125) (100). Data are plotted as mean residue ellipticity, [θ], which is in units of degrees cm$^2$ dmol$^{-1}$ residue$^{-1}$.

**Analytical ultracentrifugation**

Experiments were performed with a Beckman XL-I analytical ultracentrifuge (Beckman Coulter, Indianapolis, IN) using absorbance optics at 280 nm. Sedimentation velocity experiments were performed at 36,000 rpm at 20 °C with and without 3 mM ZnCl$_2$ (Fig. 2) or at 25 and 37 °C with or without 500 μM or 1 mM ZnCl$_2$ as indicated (Fig. 3). Data were analyzed using Sedit software (101) and the c(s) (Figs. 1 and 2) or c(s,ff$_c$) models (Fig. 3). The c(s,ff$_c$) model describes the sedimentation behavior of the species in solution as a two-dimensional distribution based on both sedimentation coefficient and frictional ratio, which allows resolution of multiple species of widely-differing size and shape sedimenting at the same sedimentation coefficients (33, 34). Parameters of buffer density and viscosity and the partial specific volume of constructs were calculated using SEDNTERP (102) at all relevant experimental temperatures. The data described in Fig. 3, B and C, were analyzed using several different models in SEDFIT; the c(s,ff$_c$) model yielded fits with the lowest value for the summed square of the residuals (SSR). The standard c(s) model fit gave an SSR value of 0.5140; the best-fit worm-like chain model gave an SSR value of 0.5102; the c(s,ff$_c$) model gave an SSR value of 0.5029 when setting the frictional ratio limits from 1 to 5. To show reversibility of Brpt5.5 initial assembly (Fig. S7), 6 μM Brpt5.5 was dialyzed overnight into 50 mM MOPS, pH 7.2, 50 mM NaCl, 3.5 mM ZnCl$_2$. A sample was analyzed by sedimentation velocity AUC with the addition of buffer (Brpt5.5 + Zn), 7 mM DTPA (+Zn + DTPA), or enough HCl to reach pH near 6 (+Zn + HCl). As a control, Brpt5.5 was also dialyzed into 20 mM MES, pH 5.0, 50 mM NaCl, 3.5 mM ZnCl$_2$. This sample sedimented as a monomer near 2 S (Fig. S7). The absorbance at 280 nm was used for the c(s) distribution analysis.

**Transmission EM**

5-μl samples of 10 μM HMBP–Brpt3.5 incubated with 500 μM ZnCl$_2$ were applied to 200 mesh Formvar carbon/copper grids for 2 min, washed with distilled H$_2$O, stained by 1% uranyl acetate dropwise for 30 s, and washed a second time with distilled H$_2$O. Samples were then dried for 1 h prior to viewing on a Hitachi 7600 transmission electron microscope at an accelerating voltage of 80 kV. Images were captured using an AMT 2k CCD camera. Brpt5.5 was incubated at 20 μM with 5 mM Zn$^{2+}$ and incubated at 37 °C for 3 weeks. Samples were stained using the same protocol as HMBP–Brpt3.5, but with 2% uranyl acetate (Electron Microscopy Sciences). To prepare biofilms for TEM, the protocol for “Harvesting biofilms” was followed up through the step of washing the biofilms off the dialysis tubing (see below). The collected washes were treated with DTPA, FA, or buffer and incubated for 1 h at room temperature while shaking. A 3-μl sample was added to the grids for negative staining as described above. For extracellular biofilm material, washes were centrifuged at 17,000 × g for 5 min before collecting the supernatant.

**Harvesting biofilms**

This protocol was based on an approach used by Sun et al. (17) to isolate extracellular and cell wall–associated proteins from S. epidermidis biofilms. RP62A was grown overnight on tryptic soy agar (TSA) with sheep blood (Thermo Fisher Scientific™ R01202). Colonies were scraped from the agar plate and suspended in TSB to an absorbance of 0.1. A piece of the 3500 MWCO dialysis tubing was cut down the edge and opened to be arranged over a TSA agar plate. A 2-ml aliquot (with the addition of ZnCl$_2$, DTPA, FA, or 100 mM MES, pH 5.0) was added to the dialysis tubing on top of the TSA agar. After 16 h of growth at 37 °C, the biofilm was washed off of the dialysis tubing by pipetting 1 ml of H$_2$O across the surface of the tubing. For lysostaphin digestion of the cell wall, mixtures were centrifuged for 10 min at 17,000 × g, and the pellet was resuspended in 50 mM Tris, pH 7.4, 150 mM NaCl, 30% raffinose. Samples were then incubated at 37 °C for 1 h with 1 mg/ml of lysostaphin (Sigma L3876).

**Thioflavin T protein fluorescence assay**

10 μM HMBP–Brpt3.5 and 20 μM Brpt5.5 samples were treated with 10% FA to depolymerize amyloid fibers (44), followed by dialysis into 50 mM MOPS, pH 7.2, 50 mM NaCl. The FA-treated protein was then incubated with or without 500 μM ZnCl$_2$ for 24 h at 20 °C prior to adding thioflavin T (MilliporeSigma) to a final concentration of 10 μM. In parallel, one sample of HMBP–Brpt3.5 was treated with 10% FA and was tested for ThT fluorescence without removing the FA (labeled FA + HMBP–Brpt3.5 + ThT in Fig. 4B). Fluorescence was measured using a PerkinElmer Life Sciences LS50B luminescence spectrophotometer or a Biotek Synergy at an excitation of 434
or 440 nm and collecting the complete emission spectrum between 450 and 600 nm. Congo Red absorbance was measured from 400 to 600 nm, and the absorbance at 540 nm was plotted in Fig. 4E. ThT emission was plotted at 482 nm (Fig. 4, C and E).

**HPLC assays**

For the HPLC fiber/oligomer quantification assay, 10 μM HMBP–Brpt3.5 samples were incubated with or without 500 μM ZnCl₂ for 1, 4, or 30 days at 20 or 37 °C. Samples were prepared based on a protocol previously described by O’Nuallain et al. (60). Briefly, samples were centrifuged for 1 h at 13,500 rpm and then resuspended in 250 μL of 5% acetonitrile. Samples were loaded on a Sepax Bio-C4 reverse-phase column (Sepax Technologies, Newark, DE) with a 5-μm particle size and 300 Å pore size and run on an Akta purifier in a running buffer of 5% acetonitrile and 0.1% trifluoroacetate at a flow rate of 0.75 ml/min. A linear gradient of 0–100% eluent (95% acetonitrile and 0.1% trifluoroacetate) from 13.33 to 66.67 min was used to elute the proteins. Peaks were detected by absorbance at 280 nm and integrated using the Unicorn software, normalizing the peak area to a standardized elution volume for monomer and oligomer peaks.

To determine the stability of the Zn²⁺-induced amyloid fibers, samples of 10 μM HMBP–Brpt3.5 were incubated with 500 μM ZnCl₂ for 1 or 4 days at 20 or 37 °C, followed by the addition of an equal volume of the following: 1) buffer (20 mM Tris, pH 7.4, 150 mM NaCl); 2) dilute hydrochloric acid sufficient to lower the pH 5.0; or 3) 2 mM Na₂-DTPA. Samples were further incubated for 2 h, mixed with equal volume of 5% acetonitrile, and then separated on a C4 column.

**Turbidity assays**

5 μM Brpt5.5 in 2 ml of 20 mM MOPS, pH 7.2, 50 mM NaCl was incubated with 3, 5, or 8 mM ZnCl₂ at 37 °C in a shaking incubator. The sample was transferred to a cuvette, and the absorbance measured at 400 nm (Fig. 5). Significant difference was identified when p < 0.05.

**Dynamic light scattering**

A Malvern Analytical Zetasizer Nano S was used to perform particle size measurements at 20 °C on 5 μM Brpt5.5 samples incubated with 5 mM ZnCl₂ for at least 30 days at 37 °C in a quartz microcuvette. The Z-average of at least three series of measurements was reported. Native Brpt5.5 samples were measured in the same way but were not incubated with ZnCl₂ prior to the experiment.

**Confocal microscopy**

For analysis of Brpt5.5 protein fibers, 10 μM samples were incubated with 500 μM ZnCl₂ for 2, 6, 12, 18, or 24 h prior to addition of thioflavin T (10 μM final concentration). Samples were then added to 8-well borosilicate glass slides (Nunc) and viewed using a Zeiss LSM 710 inverted microscope using an Apochromat ×63/1.40 oil differential interference contrast M27 objective for bright field and a laser set to 458 nm with emission filters at 469–580 nm to measure thioflavin T fluorescence. For analysis of amyloid formations in biofilms, overnight cultures of *S. epidermidis* RP62A were diluted 1:200, and 400-μL aliquots were added to 8-well borosilicate glass slides. The slides were incubated for 18 h at 37 °C without shaking, followed by addition of thioflavin T to 10 μM as indicated. The media were aspirated, and each well was washed three times with distilled H₂O. The biofilms were viewed using a Zeiss LSM 510 inverted microscope as described above. Control biofilm samples were alternatively labeled with the LIVE/DEAD stain kit (Invitrogen), using 3 μL of stain diluted in 1 ml of sterile saline buffer.

**Biofilm formation assay**

To quantify the ability for RP62A to form biofilms under varying conditions, a crystal violet-based biofilm formation assay was utilized (25). An overnight culture grown in TSB was diluted 1:200 into TSB. To wells of a 96-well plate (Corning 351172), 200 μl of diluted culture were added, along with initial treatments of ZnCl₂ or DTPA (t = 0 h). DTPA, FA, or 100 mM MES, pH 5.0, was added at the specified time points and incubated for an additional 1 h at 37 °C. The liquid was then removed from the wells, followed by two washes of 200 μl of H₂O before being allowed to dry. Next, 100 μl of 0.1% crystal violet was added and allowed to stain the biofilms for 10 min at room temperature. The crystal violet solution was then removed, followed by two more washes with 200 μl of H₂O. The plates were allowed to dry before being scanned. Finally, 200 μl of 33% acetic acid was added, and plates were incubated at 4 °C for 30 min to remove crystal violet stain from the adherent biofilms. The solution in the wells was then transferred to new wells and then diluted 1:1 (100 μl of solution to 100 μl of H₂O) before scanning at 520 nm. Statistical analysis was performed in Microsoft Excel, using the two-tailed, two-sample unequal variance Student’s t test. Significant difference was identified when p < 0.05.

**Mass spectrometry**

Biofilms were prepared as described under “Harvesting biofilms,” with the exception that no H₂O was added to the dialysis membrane during collection. This kept the material in the biofilm at a reasonable concentration to be visualized by SDS-PAGE with Coomassie Brilliant Blue staining. The section outlined in Fig. 6 was reduced with DTT, alkylated with iodoacetamide, and digested with trypsin. The resultant peptides were recovered and dried in a speed vac, before analysis by nanoLC-MS/MS. We provide data from two additional samples in Fig. S8 and Table S3 and S4. These samples were collected in the same way as in Fig. 6, but one sample was not exposed to lysostaphin (details provided in supporting information).

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