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Filaments in curved streamlines: rapid formation of *Staphylococcus aureus* biofilm streamers

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**Abstract**

Biofilms are surface-associated conglomerates of bacteria that are highly resistant to antibiotics. These bacterial communities can cause chronic infections in humans by colonizing, for example, medical implants, heart valves, or lungs. *Staphylococcus aureus*, a notorious human pathogen, causes some of the most common biofilm-related infections. Despite the clinical importance of *S. aureus* biofilms, it remains mostly unknown how physical effects, in particular flow, and surface structure influence biofilm dynamics. Here we use model microfluidic systems to investigate how environmental factors, such as surface geometry, surface chemistry, and fluid flow affect biofilm development of *S. aureus*. We discovered that *S. aureus* rapidly forms flow-induced, filamentous biofilm streamers, and furthermore if surfaces are coated with human blood plasma, streamers appear within minutes and clog the channels more rapidly than if the channels are uncoated. To understand how biofilm streamer filaments reorient in flows with curved streamlines to bridge the distances between corners, we developed a mathematical model based on resistive force theory of slender filaments. Understanding physical aspects of biofilm formation of *S. aureus* may lead to new approaches for interrupting biofilm formation of this pathogen.
Keywords: Staphylococcus aureus, biofilm, biofilm streamers, flow, microfluidics, microbiology

1. Introduction

Staphylococcus aureus is a human pathogen notorious for causing hospital-acquired infections as well as fatal infections that occur outside of health care settings [1–3]. Methicillin-resistant S. aureus (MRSA), in particular, is a major concern due to its potent virulence coupled with resistance to many antibiotics [4–6]. MRSA is the most widespread cause of hospital-associated infections in the United States and Europe with a high mortality rate [7–10]. S. aureus and MRSA cause a variety of infections ranging from minor skin and soft tissue infections to serious illnesses such as infections of indwelling medical devices, osteomyelitis, endocarditis, sepsis, and toxic shock syndrome [11].

S. aureus infections that are associated with abiotic materials, such as intravenous catheters and implants, are of primary concern because S. aureus colonizes such medical devices and forms biofilms [12–17]. Biofilms are surface-associated three-dimensional conglomerates of bacteria that are enclosed by self-produced extracellular polymeric substances (EPS) [18–20]. Once biofilms have developed, their removal is challenging because, compared to planktonic organisms, cells in biofilms display enhanced resistance to antimicrobial treatments and host immune defenses [21–28]. Consequently, biofilms are often responsible for chronic infections [29], leading to high morbidity and significant healthcare costs [30, 31]. Although biofilm-associated infections have spurred intense research efforts to understand biofilm formation by S. aureus [32–34], how physical aspects of the microenvironments of medical devices impinge on biofilm dynamics remain poorly understood.

The microenvironment affecting biofilm formation on indwelling medical devices can be characterized by complex local geometries, mechanical shear forces due to flows, the local chemical milieu and surface chemistry. Typically, inserted medical devices that come into contact with human blood are coated immediately with blood plasma proteins. Plasma proteins are readily adsorbed on abiotic surfaces [35, 36], and thus, they generate a unique local surface chemistry. Previous studies show that S. aureus attachment to surfaces is enhanced by adsorbed blood plasma proteins, such as the matrix proteins fibrinogen and fibronectin [37, 38]. S. aureus cell surfaces are decorated with components that recognize adhesive matrix proteins and mediate binding to the molecules. These components include fibronectin-binding proteins A and B and clumping factors A and B [39]. However, it is unclear how these components function under realistic physical conditions that include fluid motion, the associated flow regimes and surface geometries. For example, previous studies of the role of these components in S. aureus biofilms primarily considered biofilm development on smooth surfaces with no flow or constant flow of nutrient-containing medium across the biofilm [40–42]. Recent reports have demonstrated that, in the presence of flow and complex surface geometries, biofilms of a bacterial pathogen, Pseudomonas aeruginosa, deform into flexible three-dimensional filaments, known as ‘streamers’ [43–45], which connect the gaps between corners [46, 47], causing
clogging in flow systems [48]. Whether and how rapidly *S. aureus* can form biofilm streamers and how factors related to surface chemistry affect *S. aureus* biofilms are unknown.

To study *S. aureus* biofilm formation in physical environments that mimic *in vivo* conditions, we used model microfluidic systems that include curvy channels, as well as networks of multiple channels. The wall shear stresses that we used range from \(~0.02\) Pa to \(~1\) Pa, which are comparable to physiological wall shear stresses present in capillaries, venules, or catheters (between \(0.02\) Pa to \(4\) Pa) [49, 50]. We examined several types of *S. aureus* strains, which differed in their quorum sensing system. For *S. aureus*, one of the quorum sensing system is also known as the accessory gene regulator (*agr*) system that regulates biofilm development and the expression of virulence factors [51–53]. Natural isolates of *S. aureus* can be grouped into four different *agr* classes, which correlate with different types of infection [54, 55]. We generated a realistic surface chemistry in our model microfluidic channels by coating the channels with human blood plasma. We discovered that the blood plasma triggered especially rapid cell attachment to the channels, resulting in the formation of biofilm streamers within minutes. The streamers led to clogging dynamics that are considerably more rapid than for other bacteria, such as *P. aeruginosa* [48]. To understand how streamers initiate and bridge the gaps between corners, we modeled elastic filaments in flows with curved streamlines, and qualitatively compare this model with our experiments.

### 2. Results and discussions

#### 2.1. Rapid biofilm streamer formation of *S. aureus*

In microfluidic channels containing corners and flow (figure 1(a)), all four *agr* groups of *S. aureus* strains that we examined rapidly form biofilm streamers (figure 1(b) and supplement figure S1, available from stacks.iop.org/NJP/16/065024/mmedia), with slight differences in morphology. However, for the detailed experiments on biofilm streamer initiation and clogging that are described below, we restrict ourselves to study only one representative strain, *S. aureus* RN6734, which possesses the *agr* I system.

To visualize the morphology of the biofilm streamers, we stained the *S. aureus* cells with a fluorescent dye. This method revealed a three-dimensional structure suspended between the corners of the channel (figure 1(b)). High-resolution confocal images of these streamers show that they initially consist of porous networks that, over time, become more rigid and less porous. *S. aureus* forms biofilm streamers more rapidly than *P. aeruginosa* [48, 56] under equivalent conditions. *S. aureus* biofilms initially form on the walls of the channel (figure 2(a)), and are subsequently deformed by flow into thicker clusters on the corners of the channel (figure 2(b)). Finally, due to flow, small filaments are pulled out from these clusters. We visualized the entire time sequence as the filaments elongate and attach to the channel walls of the closest downstream corner to form a biofilm bridge between adjacent corners (figure 2(c)). At this stage, streamers are thin (<10 \(\mu\)m), flexible, and they vibrate in the flow. While the streamers remain flexible, they occasionally detach and reattach elsewhere downstream (figure 2(d)) [57, 58]. It is also apparent that the filamentous biofilms form independently at different corners. After biofilm-corner-to-corner-bridge formation occurs, the width of the streamer increases due to accumulation of EPS and/or trapping of cells that flow past. As the streamers thicken, they also appear to become more rigid as we observe that they cease to vibrate in the flow (figure 2(e)).
In these models of physiological systems, flow is the major contributor to the shape of the biofilm structures, whereas bacterial motility is less significant since non-motile staphylococci exhibit a similar biofilm structure to that of their motile Gram-negative counterparts [48]. This feature suggests that streamer formation may not depend on bacterial motility as flow itself can transport cells or biomass to the streamer site [48, 59]. We conclude that, in this context, flow plays a major role in actively shaping the three-dimensional structure of the biofilms [46, 48, 50, 60–64], in addition to affecting nutrient gradients and social behaviors in biofilms [65–68]. By contrast, in environments lacking flow, motility seems to be important for accumulation of cells near surfaces and subsequent biofilm formation [69–71].

2.2. Blood plasma coating expedites biofilm streamer formation and clogging of the channel

To examine the effect of realistic surface chemistry on S. aureus biofilm formation in our microfluidic channels with complex geometries, we coated the channels with human blood plasma [72]. In static conditions, S. aureus biofilm formation is absent, or strongly reduced, when the surfaces are not coated with human blood plasma [73, 74]. However, in our microfluidic flow channels, S. aureus biofilms also form in the absence of blood plasma surface.
Figure 2. Initiation of the *S. aureus* biofilm streamer and its rapid expansion. The displayed images are representative snapshots from movie 1, available from stacks.iop.org/NJP/16/065024/mmedia, acquired at 30 fps using bright field illumination. The images were taken at a focal plane that is perpendicular to gravity and in the middle of the channel, which is 100 μm from the bottom of the channel. (a) Initial image of the channel, which is coated with human blood plasma proteins. (b) A thin streamer originates from the corner on which surface-attached biomass has accumulated. (c) A streamer has bridged the gap between adjacent corners. At this stage, the streamer is flexible and vibrates in the flow. (d) Occasionally streamers become dislodged and they can reattach elsewhere downstream. (e) Stable streamers have formed on all corners and accumulated additional biomass, making them less flexible. The suspension we flowed through the channel was at a cell density that corresponds to OD<sub>600</sub> = 1.2. The scale bar represents 200 μm.
coating. Nonetheless, biofilm formation is strongly affected by the surface coating in our system. Indeed, after initiation of flow in microfluidic channels that were coated with human plasma proteins, streamers robustly bridge the distance between corners at all 36 bends of our microfluidic channel within a few minutes, following rapid attachment of cells to the surface. By contrast, in the absence of blood plasma surface coating, it takes several hours to form streamers between all the bends in our channel. After streamer formation between the bends of the channel, rapid biomass accumulation of streamers occurs in a way that is independent of the size of the channels (figure 3). Furthermore, detachment of streamers from the surface occurs less frequently in the blood-plasma-coated channel, indicating that the cells were more strongly attached to the blood-plasma-coated chambers than to the non-coated chambers. Enhanced surface-attachment is consistent with previous reports of binding of bacterial membrane-associated proteins to immobilized blood plasma proteins [38]. In addition, the presence of blood plasma may also induce changes in S. aureus gene expression that promote attachment or EPS production to further accelerate biofilm streamer formation. In summary, S. aureus biofilm streamers initiate via a rapid mechanism that relies on binding of S. aureus cells to immobilized blood plasma proteins on the walls of the channels. For P. aeruginosa, not only the initial surface-attachment of a few cells is important, but also cell-growth and division is required for the cells to form confluent surface-attached biofilms over tens of hours. Thus, in the case of P. aeruginosa, the process takes tens of hours [48]. By contrast, the time scale for S. aureus biofilm streamer initiation is not sufficient for substantial cell growth, particularly when the surfaces are coated with human blood plasma (figure 3(a)), which indicates that cell growth is not essential for S. aureus streamer formation.

Expansion of biofilm streamers increases the hydrodynamic resistance of the channel, which ultimately leads to clogging of the microfluidic device, if the pressure difference that drives the flow remains constant. To investigate clogging dynamics caused by S. aureus biofilms, we applied a constant pressure difference across the microfluidic system containing

Figure 3. Coating the channels with human blood plasma affects S. aureus biofilm streamer formation. (a) We measured the time, $T_{1/2}$, until biofilm streamers have grown in situ to occupy half of the channel width at a fixed flow rate of $1.5 \pm 0.05 \mu \text{L min}^{-1}$ and at a fixed bacterial cell concentration (optical density at 600 nm = $0.46 \pm 0.05$). We used three different channels with square cross-sections of different side lengths: 100 μm, 150 μm and 200 μm. Coating the microfluidic channels with human plasma prior to bacterial inoculation leads to rapid accumulation of biomass compared with non-coated channels, independent of the size of the channels. The error bars indicate the range of values from three independent replicates. (b) A typical image used to measure the biofilm thickness.
Figure 4. Dynamics of biofilm-induced clogging for *S. aureus* under constant pressure drop flow conditions. (a) Schematic drawing of the experimental apparatus: early or mid exponential phase *S. aureus* cells are loaded into a reservoir. Using wide-bore tubing, this reservoir is connected to the microfluidic channel. The effluent collection dish is placed on an analytical balance. The height difference, $\Delta h$, between the reservoir suspension and the effluent collection dish is proportional to the applied pressure difference. (b) The weight of the effluent suspension as a function of time is converted into the flow rate, $Q_0$, as a function of time using equation (A.1). Using this flow rate measurement, the time to clogging, $T$, and the duration of the clogging transition, $\tau$, were measured for each channel. (c), (d) $T$ and $\tau$ for different flow rates, wall shear stress, mean flow speeds and different channel sizes. All channels had a square cross-section, with a width and height equal to $d = 100 \mu m$, $150 \mu m$, or $200 \mu m$. Wall shear stress and mean flow speed were calculated as $\mu_{\text{water}} \times \frac{Q_0}{d^3}$ and $\frac{Q_0}{d^2}$ where $\mu_{\text{water}}$ is the viscosity of water ($\sim 0.001$ Pa s). $T$ and $\tau$ appear to be independent of channel size, flow rate, wall shear stress, and mean flow speeds for the flow rates we investigated. The suspension we flowed through the channel in these experiments was at a cell density that corresponds to $\text{OD}_{600} = 0.06 \pm 0.01$. (e), (f) $T$ and $\tau$ depend strongly on the bacterial cell concentration of the suspension flowed through the channel.
channels coated with blood plasma (figure 4(a)), which drives a constant flow rate $Q_0$ prior to the clogging transition. We measured the clogging time, $T$, at which the flow rate was reduced to $Q_0/2$ (figure 4(b); materials and methods). In addition, we measured the duration of the clogging transition, $\tau$, which we defined as the time at which the flow rate decreases from 75% to 25% of $Q_0$ (figure 4(b); materials and methods). We found that $T$ and $\tau$ are approximately independent of flow rates, flow speed, shear stress, and channel sizes, for the flow rates we investigated (figures 4(c) and (d)). However, increasing the bacterial cell density that flows through the channel from $\text{OD}_{600} \sim 0.05$ to $\text{OD}_{600} \sim 0.2$ ($\text{OD}_{600}$ is the optical density at 600 nm, which is a common measure of cell density in suspensions) significantly reduced both the clogging time and the clogging transition duration (figures 4(e) and (f)), yet increasing the cell density further had no strong effects on $T$ and $\tau$. As the onset of the clogging transition is highly correlated with the formation of biofilm streamers, and the clogging duration is highly correlated with the streamer expansion, figures 4(e) and (f) reveal that streamer initiation and expansion have a strong dependence on cell density at low cell densities, and a weak dependence on cell density at high cell densities. The effect of cell density in initiation and clogging is consistent with the interpretation that biofilm streamers initiate by blood-plasma-protein-mediated attachment of cells to the wall. In addition, increased production of surface-attachment-mediating proteins by $S. aureus$ during the exponential growth phase should also increase the probability of attachment to surfaces and consequently stimulate biofilm formation [75–77], ultimately leading to more rapid clogging.

2.3. $S. aureus$ streamers form in complex geometries

We demonstrate that $S. aureus$ biofilm streamers are ubiquitous in complex geometries (figure 5). In particular, we found that biofilm streamers develop rapidly following inoculation and they span the gaps in a complex, branched channel. This example indicates that biofilm streamers might be universal in porous media in environmental and medical environments [48, 78–80], where they could cause rapid clogging.
Figure 6. (a) Image of a biofilm streamer during initial formation in a microfluidic channel where a bacterial cell concentration corresponding to OD$_{600}=0.5$ was flowed through the channel by a syringe pump, which was set to a flow rate of 1.5 ± 0.05 μL min$^{-1}$. The red arrows indicate the flow direction: (i) The streamer grows in length.
2.4. Modeling biofilm streamers as hinged or clamped flexible filaments in curved flow

To understand how biofilm streamers that are bound on one corner can attach to the adjacent downstream corner, we examined the deformation of streamers in the flow as the streamers grow longer. The streamer stems from the left corner, growing in length with a reorientation in the clockwise direction towards the right corner (figures 6(a)–(i)). The curved shapes of the streamer filaments show that they are highly flexible. Once a biofilm bridge has formed between two corners, the biofilm bridge expands in radius and becomes straighter (figures 6(a)–(ii)) likely due to a reduction in flexibility of thicker biofilm bridges.

To characterize this system mathematically, we consider an idealized model flow problem capturing some essential features of the current scenario, based on a flexible filament that is anchored at one end and expands in length in the presence of curved streamlines. Arguably the simplest flow with curved streamlines is the steady rigid body rotation flow with velocity \( \nu = \hat{\gamma} - \vec{e}_y \vec{e}_x \), where \( \hat{\gamma} \) is the rotation rate. This flow results in circular streamlines, as shown in figure 6(b). We model the biofilm streamer as a flexible slender filament of radius \( r \), length \( L \), and bending stiffness \( B \). The position vector of the filament is denoted as \( \vec{x} (s, t) = x (s, t) \hat{e}_i + y (s, t) \hat{e}_j \), where \( s \in [0, L] \) is the arc length. One end of the filament is tethered at the position \( \vec{x}_0 = -L_0 \hat{e}_x \), a distance \( L_0 \) from the origin, where this geometry is motivated by a streamer tethered at the surface of the experimental channel.

To model the dynamics of the filament, we exploit the slenderness of the streamer and use resistive force theory [81–83] to characterize the viscous force per unit length \( \vec{f}_{\text{viscous}} \) on the streamer:

\[
\vec{f}_{\text{viscous}} = \left[ \zeta_\perp (I - \vec{x}\vec{x}) + \zeta_\parallel \vec{x}\vec{x} \right] (\vec{\nu} - \vec{x}).
\]
where $\zeta_{\parallel} = 2\pi\mu/\log (L/r)$ and $\zeta_{\perp} = 2\zeta_{\parallel}$ are the drag coefficients for a slender filament moving, respectively, parallel and perpendicular to its axis. We denote derivatives on $\bar{x}$ with respect to arclength $s$ or time $t$ with subscripts. The streamer flexibility is described by the Euler–Bernoulli beam theory [84], which gives rise to an elastic force per unit length

$$\vec{f}_{\text{elastic}} = -B\ddot{x}_{\text{xss}} + \{T\ddot{x}_y\},$$

where $B = EI$ is the constant bending stiffness of the filament, i.e. the product of the Young’s modulus $E$ and the area moment of inertia $I$, and $T(s, t)$ can be interpreted as the tension along the streamer to enforce its inextensibility. The force balance $\vec{f}_{\text{viscous}} + \vec{f}_{\text{elastic}} = 0$ results in an elastohydrodynamic equation governing the deformation of a flexible streamer with position vector $\bar{x}(s, t)$ in the rotating flow (see Materials and Methods for details).

A dimensionless parameter for comparing viscous and elastic forces is given by $\eta_s = \zeta_s U_0 L^3/B$, where the background flow $U_0$ scales with the distance from the origin as $U_0 \sim \dot{\gamma} L_0$. A larger value of $\eta_s$ represents effectively a more flexible filament. The dimensionless parameter can be rewritten as $\eta_s = (L/L_0)^3\eta$, where $\eta = \zeta_s\dot{\gamma} L_0^3/B$ is independent of the filament length $L$. Here we consider a fixed distance from the origin $L_0$ (hence a fixed $\eta$), and we vary the effective flexibility $\eta_s$ by changing the dimensionless filament length $L/L_0$, so that the effective flexibility increases with filament length. To understand how the clockwise reorientation of the streamer occurs, we consider filaments that are hinged and those that are clamped at the tethering location. Hinged filaments are free to rotate at the tethering location, whereas clamped filaments have a fixed tangent at the tethering location.

For a vertically clamped filament (figure 6(c)), flexibility is essential for reorientation, since the filament would be vertical if it were completely rigid ($\eta = 0$). For flexible filaments ($\eta > 0$), increasing the dimensionless filament length $L/L_0$ leads to higher effective flexibility $\eta_s$, which allows the filament to bend more in the clockwise direction. Filaments that are more flexible ($\eta = 100$, solid lines in figure 6(c)) can deform more strongly in flow than less flexible filaments ($\eta = 1$, dotted lines).

For a hinged filament, the physical picture is quite different from the clamped filaments (figure 6(d)), in spite of the apparent similarity of the filament shape in the two cases. We again consider flexible filaments ($\eta = 100$) with increasing length $L/L_0$ (solid lines). However, for hinged filaments, flexibility does not play an essential role in the clockwise reorientation. Relatively rigid filaments ($\eta = 1$, dotted lines) turn more towards the clockwise direction than very flexible filaments ($\eta = 100$, solid lines). It is also found that the filament adopts similar shapes for large values of $\eta$. Unlike a clamped tether, a hinge cannot support any torques, and the filament is therefore free to rotate at the tethered end. The reorientation mechanism in the hinged case is primarily due to the change of filament length, which alters the torque balance at the hinge. In the case of completely rigid hinged rods ($\eta = 0$) in the simple rotating flow considered here, we can derive an analytical formula for the steady-state tangent vector of the straight filament

$$\bar{x}_{\text{x rigid}} = 2L/\left(3L_0\right)\hat{z} + \sqrt{1-4L^2/\left(9L_0^2\right)} \hat{y},$$

(3)
which was obtained by a torque balance about the hinge. Equation (3) illustrates that increasing the filament length $L/L_0$ reorients the filaments in the clockwise direction, as a consequence of the altered torque balance. The limit $L/L_0 = 3/2$ corresponds to a rigid rod oriented horizontally in the rotating flow, and any further increase in $L/L_0$ does not support a steady-state configuration so that the rod rotates about the hinge.

Biofilm streamers are likely to behave more like hinged filaments than clamped filaments, because there are no geometric constraints that prevent the streamers from freely rotating at the tether. This model predicts that, as biofilm streamers become longer, their orientation changes towards the clockwise direction (figures 6(a)(i) and (d)). We also note that the model also predicts that as hinged filaments become less flexible, the stiffening induces further clockwise reorientation (figures 6(a)(ii) and compare black solid and dotted lines in figure 6(d) for a streamer with the same length $L/L_0$ but different flexibility $\eta$). By contrast, if the streamer is clamped at one end, the stiffening streamer should reorient in the counter-clockwise direction.

Our mathematical model can also be used to estimate the tension in the biofilm streamer, which is important for determining under which conditions streamers become dislodged. For vertically clamped filaments, the tension varies non-monotonically along the filament with maxima present near the tethered end (figure 6(e)). The location of the maximum tension is at the point at which the filament will break if it cannot withstand the tension. By contrast, tension decreases monotonically along hinged filaments, and the maxima always occur at the hinged end (figure 6(f)). Experimentally, we also observe that biofilm streamers become dislodged close to the tethering location.

In this section we illustrated how flexibility and tethering conditions of biofilm filaments affect their orientation in the curved flow fields with a simple rotating flow. Although the model captures the fundamental feature of curved streamlines, quantitative comparisons with the experiments are, however, beyond the reach of the current work because the background flow employed in the model is a highly idealized rotating flow, compared with the actual flow in the microfluidic channel. The idealized flow model here is intended to uncover qualitatively interesting physical features associated with curved streamlines. In future work, we will consider more realistic flow models and explore the idea of estimating the elasticity of biofilm streamers based on their deformed shape in more realistic flows.

3. Conclusion

We used model microfluidic systems to investigate how environmental factors, such as surface geometry, surface chemistry, and fluid flow affect biofilm formation of _S. aureus_. We discovered that _S. aureus_ forms flow-induced, filamentous biofilm streamers in curvy channels of different sizes more rapidly than _P. aeruginosa_. When surfaces were coated with human blood plasma, streamers appeared within minutes and grew to clog the channels on a rapid timescale. We found that _S. aureus_ biofilm streamers also formed in different flow geometries, which led us to hypothesize that _S. aureus_ biofilm streamers are ubiquitous in natural, and possibly infectious, environments. Using mathematical modeling, we showed how flexibility and tethering conditions of biofilm filaments affect their orientation in the flow fields with curved streamlines. Whether species other than _P. aeruginosa_ and _S. aureus_ form biofilm streamers, and which rheological features of the biofilm matrix are required for streamer formation are open questions that we are currently pursuing. Studying biofilm dynamics in environments that mimic physical...
and chemical conditions of natural habitats, such as inserted medical devices, holds the potential for identifying new biofilm structures, and perhaps such studies will reveal novel methods to prevent biofilm-related diseases.

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**Appendix: Materials and methods**

**Bacterial strains and culture conditions**

All *S. aureus* strains used in this study were generous gifts from Dr Tom W Muir’s group (Princeton University), and were originally obtained from Dr Richard Novick’s group (New York University). *S. aureus* RN6734 is the standard *agr* group I laboratory strain [55]. *S. aureus* RN6607 was isolated from a healthy human, and it represents *agr* group II [55, 85]. *S. aureus* RN8465 is the *agr* group III prototype, also known as menstrual toxic shock syndrome strain [86, 87]. *S. aureus* RN4850 is the *agr* IV prototype, and it was isolated from a patient with scalded skin syndrome [55, 88]. All *S. aureus* strains were grown overnight in tryptic soy broth (TSB) supplemented with 0.5% glucose and 3% NaCl at 32 °C. Overnight cultures of *S. aureus* were back-diluted 1:100 in TSB, and grown to early or mid-exponential phase at 37 °C with shaking.

**Microfluidic experiments and microscopy**

Microfluidic channels were prepared from polydimethylsiloxane (Sylgard 184, Dow Corning) using conventional soft lithography techniques. Each channel was sealed to a glass microscope slide (75 × 50 × 1.0 mm, Corning) after air plasma treatment (Harrick Scientific, NY). The microfluidic devices consist of a single channel containing many corners and have a square cross-section. We used three different channels with different cross-sections, which had side lengths of 100 μm, 150 μm or 200 μm (figures (1)–(4), and (6)). Microfluidic devices consisting of a network of channels were 1600 μm wide and 30 μm high (figure 5). Some channels were coated with human blood plasma, according to the following procedure: human blood plasma was purchased from Biological Specialty Corporation, PA, and was diluted with H₂O to a 20% human blood plasma solution, which was flown into the microfluidic channels. These channels were then stored in a humid environment at 4 °C for 24 h prior to bacterial inoculation [72] to ensure that the channels stay wetted. Bacterial cultures were introduced to the channels using either a constant flow driven by a syringe pump (Harvard Apparatus, MA) (figure 1(a)), or a constant pressure (details are provided below). Images of the biofilms were acquired using a Nikon Ti-Eclipse microscope under epifluorescence illumination and a CCD camera (iXon, Andor), or using phase contrast illumination on a Leica DM IRB microscope. Images were analyzed and processed using Matlab (Mathworks).
Flow driven by a constant pressure

When bacterial cultures were introduced into microfluidic channels at constant pressure, 100 mL of *S. aureus* culture was used to fill a reservoir connected via Tygon tubing (inner diameter 2.4 mm) and Luer connectors (Cole–Parmer) into an inlet of a microfluidic channel (figure 4(a)). The outlet was also connected to Tygon tubing, and the effluent culture was collected in a dish placed on an analytical balance capable of weighing to 0.1 mg precision. The height of the culture reservoir above the effluent culture dish determined the applied pressure difference $\Delta p$, and therefore the flow rate through the channel.

Analysis of flow rate time series

The weight of the effluent culture was measured every 4 s on an analytical balance (GD503, Sartorius), controlled via Labview (National Instruments). To obtain the flow rate time series $Q(t)$ from the effluent weight time series $w(t)$, we computed

$$Q(t) = \frac{w(t + 30 \text{ s}) - w(t - 30 \text{ s})}{60 \text{ s}} \frac{1}{\text{density}}$$

where the density was assumed to be the density of water, 1 g mL$^{-1}$. The time to clogging, $T$, and the duration of the clogging transition, $\tau$, were calculated by fitting the function $Q_0 / \left[ 1 + \exp \left( \frac{t - T}{\tau/2} \right) \right]$ to the measured flow rate $Q(t)$, where $Q_0$ is the flow rate prior to the clogging transition. The time $\tau$ is therefore defined as the time period in which the fitted flow rate decreases from 75% to 25% of $Q_0$, while $T$ is the time at which the fitted flow rate is $0.5Q_0$ [48].

Staining of biofilms in situ

To visualize the three-dimensional structures of streamers, we fluorescently stained the biofilms produced by *S. aureus* (figure 1(b)). For this assay, we very carefully and slowly removed the tubing carrying the bacterial culture from the inlet of the microfluidic channel after a streamer had formed. We then connected a syringe containing 5 µM SYTO 9 stain (Invitrogen) in phosphate buffered saline to the channel inlet, and flowed this solution through the channel to stain nucleic acids. To avoid the formation of bubbles in the channels, the liquid in the tubing that contains the staining solution has to make contact with the liquid in the channel before slowly pushing the tubing all the way into the channel inlet. The staining solution was flowed through the channel slowly to avoid disrupting the structure of the streamer.

Elastohydrodynamic model for the biofilm streamer

Using the resistive force theory described in the main text [81–83], the viscous force per unit length is given by

$$\bar{f}_{\text{viscous}} = \left[ \zeta (I - \vec{x}_s \vec{x}_s) + \zeta_\parallel \vec{x}_s \vec{x}_s \right] (\vec{v} - \vec{x}_s)$$

$$= \zeta (I - \vec{x}_s \vec{x}_s/2) (\vec{v} - \vec{x}_s)$$

where $\zeta = 2\zeta_\parallel$, $\vec{x}_s$ is the position vector, $\vec{x}_s$ is the unit tangent vector, $\vec{x}_s$ is the local filament velocity, and $\vec{v}$ is the background flow velocity. The elastic force per unit length is modeled by
the Euler–Bernoulli beam theory as
\[ \vec{f}_{\text{elastic}} = -B \vec{x}_{sx} + (T \vec{x}_s), \tag{A.3} \]
where \( T(s) \) is a Lagrange multiplier enforcing inextensibility of the filament; it may be interpreted as the tension along the filament. A force balance \( \vec{f}_{\text{viscous}} + \vec{f}_{\text{elastic}} = \vec{0} \) leads to the elastohydrodynamic equation
\[ \zeta (I - \vec{x}_s / 2) ( \vec{v} - \vec{x}_s ) = -B \vec{x}_{sx} + (T \vec{x}_s), \tag{A.4} \]
which can be inverted to obtain the equation governing the evolution of the filament shape \( \vec{x}(s, t) \)
\[ \zeta ( \vec{v} - \vec{x}_s ) = B \left[ \vec{x}_{sx} + \vec{x}_s (\vec{x}_s \cdot \vec{x}_{sx}) \right] - 2T \vec{x}_s - T \vec{x}_{ss}. \tag{A.5} \]

We normalize lengths by \( L_0 \), velocities by \( \gamma L_0 \), time by \( 1/\gamma \), and tension by \( \zeta \gamma L_0^2 \), resulting in the following dimensionless equation
\[ \eta ( \vec{v} - \vec{x}_s ) = \vec{x}_{sx} + \vec{x}_s (\vec{x}_s \cdot \vec{x}_{sx}) - 2 \eta T_{ss} \vec{x}_s, \tag{A.6} \]
where the stars represent dimensionless variables, \( \eta = \zeta \gamma L_0^4 / B, s^* \in [0, L^*], \) and \( L^* = L / L_0 \). We omit the stars hereafter for simplicity and refer only to dimensionless variables unless otherwise stated.

The inextensibility condition prevents any local stretching, requiring \( \vec{x}_s \cdot \vec{x}_s = 1 \) for all time, which implies \( \partial (\vec{x}_s \cdot \vec{x}_s) / \partial t = 0 \Rightarrow \vec{x}_s \cdot \vec{x}_s = 0 \). An equation for tension \( T(s) \) can be obtained from the elastohydrodynamic equation (equation (A.6)) with the aid of this inextensibility condition as follows:
\[ 2 \eta T_{s} - \eta (\vec{x}_s \cdot \vec{x}_s) = -\eta \vec{x}_s \cdot \vec{v}_s - 7 \vec{x}_{xs} \cdot \vec{x}_{sx} - 6 \vec{x}_{xss} \cdot \vec{x}_{sx}, \tag{A.7} \]
where the following identities (obtained by repeated differentiation of the identity \( \vec{x}_s \cdot \vec{x}_s = 1 \)) have been used for simplification: \( \vec{x}_s \cdot \vec{x}_s = 0, \vec{x}_s \cdot \vec{x}_{sx} = -\vec{x}_s \cdot \vec{x}_s, \vec{x}_s \cdot \vec{x}_{sx} = -3 \vec{x}_s \cdot \vec{x}_{ss}, \) and \( \vec{x}_s \cdot \vec{x}_{xss} = -4 \vec{x}_s \cdot \vec{x}_{xss} - 3 \vec{x}_s \cdot \vec{x}_{ss}. \) The coupled equations (equations (A.6) and (A.7)) can be solved numerically using the method outlined in Tornberg and Shelley [89], when appropriate boundary conditions are supplied.

For a vertically clamped filament, the dimensionless boundary conditions at the tethered end are given by \( \vec{x}(s = 0, t) = -\hat{c}_x, \vec{x}_s(s = 0, t) = \hat{c}_s \). At the free end, we have the force-free and torque-free conditions \( \vec{x}_{sx}(s = L, t) = \vec{0} \) and \( \vec{x}_s(s = L, t) = \vec{0} \), respectively. For a hinged filament, in place of the geometrical constraint on the slope \( \vec{x}_s(s = 0, t) = \hat{c}_s \) in the clamped case, we have the torque free condition \( \vec{x}_{sx}(s = 0, t) = \vec{0} \) at the tethered end, and all other boundary conditions remain identical. Note that the steady-state filament shape is independent of the initial condition.

The following boundary conditions of the tension equation apply to both the clamped and hinged cases. At the free end, we simply require the tension to vanish \( T(s = L) = 0 \). The boundary condition for tension at the tethered end is less discussed in the literature and requires
more consideration. We obtain the boundary condition by taking the inner product of the local tangent vector at the tethered end $\vec{x}(s = 0, t)$ with the elastohydrodynamic equation (equation (A.6)) evaluated at the tethered end, where $\vec{x}(s = 0, t) = 0$. This treatment leads to the following boundary condition for the tension equation at the tethered end:

$$T_s(s = 0, t) = -\frac{\eta}{2} \nu(s = 0, t) \cdot \vec{x}(s = 0, t) - 3\vec{x}_{ss}(s = 0, t) \cdot \vec{x}_{ss}(s = 0, t).$$  

(A.8)

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