



Cite this: *Soft Matter*, 2024, 20, 3845

Received 22nd January 2024,  
 Accepted 8th April 2024

DOI: 10.1039/d4sm00095a

[rsc.li/soft-matter-journal](http://rsc.li/soft-matter-journal)

## Spontaneous oscillation of an active filament under viscosity gradients†

Zhaorong Liu,<sup>a</sup> Youchuang Chao,<sup>b</sup> Zhijun Zheng<sup>c</sup> and Lailai Zhu<sup>\*a</sup>

We investigate the effects of uniform viscosity gradients on the spontaneous oscillations of an elastic, active filament in viscous fluids. Combining numerical simulations and linear stability analysis, we demonstrate that a viscosity gradient increasing from the filament's base to tip destabilises the system, facilitating its self-oscillation. This effect is elucidated through a reduced-order model, highlighting the delicate balance between destabilising active forces and stabilising viscous forces. Additionally, we reveal that while a perpendicular viscosity gradient to the filament's orientation minimally affects instability, it induces asymmetric ciliary beating, thus generating a net flow along the gradient. Our findings offer new insights into the complex behaviours of biological and artificial filaments in complex fluid environments, contributing to the broader understanding of filament dynamics in heterogeneous viscous media.

### 1 Introduction

Motile cilia, or interchangeably, flagella, are hair-like, flexible filaments protruding from a wide range of cells. Their oscillatory motion facilitates various biological processes, such as propulsion of flagellated algae that can capture CO<sub>2</sub><sup>1,2</sup> and that of planktonic ciliates important in aquatic ecosystems,<sup>3</sup> and feeding of marine invertebrates. In mammals,<sup>4</sup> flagellar movement is essential for sperm locomotion, while ciliary beating in the airways helps expel mucus and is crucial for transporting oocytes and embryos in the oviduct. Ciliary dysfunction in humans causes up to 35 diseases known as ciliopathies<sup>5</sup> including male infertility owing to weakened sperm motility, female infertility due to impaired oviductal cilia, and chronic bronchitis resulting from defective clearance of respiratory mucus. Besides its biological significance, the behaviour of ciliary beating has inspired the design of active filamentous structures for applications in adaptive materials, microfluidics, and drug delivery.<sup>6–9</sup>

Motivated by its fundamental importance and broad application prospects, a substantial amount of research has been dedicated to examining ciliary oscillation in diverse contexts.<sup>10</sup> It is established that a biological filament oscillates spontaneously,

typically driven by its internal structure—the central axoneme comprising microtubule doublets and dynein molecular motors. A variety of models have been proposed to elucidate the mechanisms of how dynein motors power spontaneous ciliary oscillation.<sup>11–22</sup> Despite these efforts to achieve mechanistic understanding, the proposed models have also facilitated the investigation on synchronisation of two active filaments<sup>23–25</sup> and the emergence of metachronal waves generated by multiple filaments,<sup>26,27</sup> focusing on inter-filament interactions. More recently, motivated by the typically non-Newtonian bodily fluids hosting cilia and flagella, research has been conducted on a spontaneously oscillating filament in viscoelastic<sup>28</sup> or shear-thinning/thickening fluids.<sup>29</sup>

Besides these recognised rheological complexities, heterogeneity is also a notable characteristic in bodily fluids, often manifesting as spatially varying viscosity. A key question is how will the viscosity gradient affect ciliary oscillation? In this work, we will address this question, but before that, let us mention the biological relevance of this scenario. It is relevant to the beating of pulmonary cilia in the airway surface liquid (ASL) that is essential for mucociliary clearance. Overlaying the airway epithelium, the ASL comprises two distinct layers: a gel-like mucus layer above a periciliary layer (PCL), as depicted in Fig. 1.

Both layers contain mucins—large macromolecules forming networks<sup>31</sup>—but differ in their types and, consequently, in fluid viscosities; the PCL is less viscous than the superficial gel layer. In addition to such a viscosity patchiness in the ASL, the PCL—typically as deep as cilia, presents its own viscosity heterogeneity. Specifically, the cilia immersed in the PCL are rich in transmembrane mucins, producing a glycan meshwork with a density that increases towards the airway surface.<sup>30</sup> This density gradient, in turn, results in a viscosity variation parallel to the ciliary orientation.

<sup>a</sup> Department of Mechanical Engineering, National University of Singapore, 117575, Singapore. E-mail: [lailai\\_zhu@nus.edu.sg](mailto:lailai_zhu@nus.edu.sg)

<sup>b</sup> School of Energy Science and Engineering, Harbin Institute of Technology, Harbin, 150001, China

<sup>c</sup> CAS Key Laboratory of Mechanical Behavior and Design of Materials, Department of Modern Mechanics, University of Science and Technology of China, Hefei, 230027, China

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4sm00095a>





**Fig. 1** Schematic of pulmonary cilia in the airway surface liquid adapted from ref. 30 with permission from Elsevier. This liquid consists of a more viscous gel layer on top of a less viscous periciliary layer (PCL). The PCL approximately contains the cilia and features a growing density of glycan from top to bottom, thus leading to a viscosity gradient.

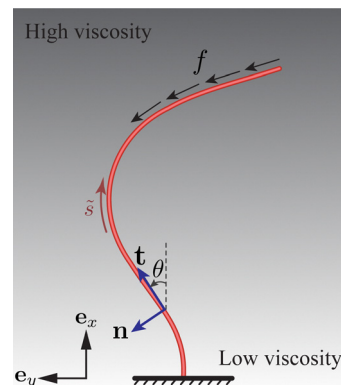
Inspired by the picture depicted above, this study explores the spontaneous oscillation of an active elastic filament in viscous fluids exhibiting viscosity heterogeneity. Simulations are conducted in the creeping flow regime to examine the influence of uniform viscosity gradients on the emergence of oscillatory instability. These numerical findings are corroborated by stability analysis. We then employ a reduced-order model to further clarify the underlying mechanisms by which viscosity gradients affect the instability.

## 2 Problem setup

We consider a slender, inextensible filament undergoing spontaneous oscillation in a quiescent creeping flow of Newtonian fluid with spatially varying viscosity. The filament of length  $L$  features a circular cross-section with diameter  $b \ll L$ , leading to a vanishing slenderness  $\beta = b/L \ll 1$ . Its profile is described by the centerline coordinates  $\tilde{\mathbf{r}}(\tilde{s}, \tilde{t})$  as functions of arc length  $\tilde{s} \in [0, L]$  and time  $\tilde{t}$ . From hereinafter, dimensional variables (but not constant values) are indicated by a tilde.

The filament is vertically clamped at its base ( $\tilde{s} = 0$ ) and, in its undeformed, relaxed state, aligns along the vertical ( $\mathbf{e}_x$ ) direction (see Fig. 2). To emulate the active force exerted by the molecular motors, we employ the follower force approach,<sup>17,32</sup> applying a compressive force density uniformly distributed along the filament centerline, aligned with its tangent everywhere. For a related configuration featuring a point follower force of constant magnitude at the tip ( $\tilde{s} = L$ ), the results, exhibiting trends similar to those for the distributed force, are given in the ESL.† A sufficiently strong follower force or force density can induce the filament to self-oscillate, simultaneously driving the surrounding flow.<sup>20,32,33</sup>

The motion of the filament is restricted to the  $xy$  plane, see Fig. 2. Consequently, its profile remains planar, and is determined by the angle  $\theta(\tilde{s}, \tilde{t})$  between the tangent vector



**Fig. 2** Schematic of an elastic filament submerged in a fluid with a vertical viscosity gradient. The filament is anchored at one end and free at its tip. Its shape is characterised by the tangent angle  $\theta(\tilde{s}, \tilde{t})$ , where  $\tilde{s}$  represents the arc length. The local tangent and normal vectors are denoted as  $\mathbf{t}(\tilde{s}, \tilde{t})$  and  $\mathbf{n}(\tilde{s}, \tilde{t})$ , respectively. The filament experiences a distributed active force with density  $-\mathbf{f}\mathbf{t}$  along its centerline.

$\mathbf{t}(\tilde{s}, \tilde{t}) = \cos \theta \mathbf{e}_x + \sin \theta \mathbf{e}_y$  and the  $\mathbf{e}_x$  axis. The associated normal vector is  $\mathbf{n}(\tilde{s}, \tilde{t}) = -\sin \theta \mathbf{e}_x + \cos \theta \mathbf{e}_y$ .

To probe the influence of the viscosity gradient, we prescribe a viscosity  $\tilde{\mu}(\tilde{x}, \tilde{y})$  linearly changing in one direction—either vertically ( $\mathbf{e}_x$ ) or horizontally ( $\mathbf{e}_y$ ). Choosing a reference viscosity  $\mu_0$  measured at the middle  $(L/2, 0)_{\tilde{x}\tilde{y}}$  of the undeformed filament, we express the viscosity as  $\tilde{\mu}(\tilde{x}, \tilde{y}) = \mu_0 g(\tilde{x}, \tilde{y})$ . Hence,  $g(\tilde{x}, \tilde{y})$  represents a scaled viscosity profile, which features two forms depending on the viscosity gradient direction:

$$g(\tilde{x}, \tilde{y}) = \begin{cases} 1 + \gamma(2\tilde{x}/L - 1) & \text{(vertical gradient),} \\ 1 + 2\gamma\tilde{y}/L & \text{(horizontal gradient),} \end{cases} \quad (1)$$

where  $\gamma$  signifies a dimensionless viscosity gradient magnitude.

## 3 Elasto-hydrodynamics at a low Reynolds number

Here, we describe the low-Reynolds-number elasto-hydrodynamics in the Stokes flow limit. Instantaneous balance between the active force, elastic force, and hydrodynamic force holds everywhere on the filament centerline. The active force density (per unit length)  $-\mathbf{f}\mathbf{t}$ , imposed anti-parallel to the tangent vector  $\mathbf{t}$ , possesses a constant magnitude  $f$ . The internal elastic force density  $\tilde{\mathbf{f}}_e = \tilde{\tau}\mathbf{t} + \tilde{N}\mathbf{n}$  consists of a tangential tension component  $\tilde{\tau}$  and a normal component  $\tilde{N} = -A\theta_{\tilde{s}\tilde{s}}$ , where  $A$  is the bending modulus of the filament (see the ESL,† Section S1) and the subscript  $\tilde{s}$  denotes differentiation with respect to the arc-length.

The hydrodynamic force on slender filaments in Stokesian fluids with a constant viscosity is typically addressed using the resistive force theory—the leading order slender-body theory.<sup>34–38</sup> Kamal and Lauga<sup>39</sup> recently extended the classical theories to incorporate a constant viscosity gradient in fluids. At the leading order of filament slenderness  $\beta$ , the hydrodynamic force aligns with the classical theories for fluids with



a uniform viscosity, except that the viscosity everywhere takes its locally non-constant value. At the next order of  $\beta$ , non-local terms emerge. Here, we address solely the leading-order effects, and consequently, the hydrodynamic force density reads

$$\tilde{\mathbf{f}}_h = -\left(\tilde{\zeta}_{\parallel}(\tilde{\mathbf{r}})\mathbf{t}\mathbf{t} + \tilde{\zeta}_{\perp}(\tilde{\mathbf{r}})\mathbf{n}\mathbf{n}\right) \cdot \frac{\partial \tilde{\mathbf{r}}}{\partial \tilde{t}} \quad (2)$$

where  $\tilde{\zeta}_{\perp}(\tilde{\mathbf{r}})$  and  $\tilde{\zeta}_{\parallel}(\tilde{\mathbf{r}})$  denote the viscous resistance coefficients that linearly scale with the local viscosity  $\tilde{\mu}(\tilde{\mathbf{r}})$  measured at  $\tilde{\mathbf{r}}(\tilde{s})$ . Based on this linear relationship, we obtain  $\tilde{\zeta}_{\perp}(\tilde{x}, \tilde{y}) = g(\tilde{x}, \tilde{y})\tilde{\zeta}_{\perp,0}$  and  $\tilde{\zeta}_{\parallel}(\tilde{x}, \tilde{y}) = g(\tilde{x}, \tilde{y})\tilde{\zeta}_{\parallel,0}$ . Here,  $\tilde{\zeta}_{\perp,0} = 4\pi\mu_0/(-\ln\beta + 1/2)$  indicates the resistance coefficient defined at the reference position  $(L/2, 0)_{\tilde{x}\tilde{y}}$ , and  $\tilde{\zeta}_{\perp,0}/\tilde{\zeta}_{\parallel,0} = 2$  as we take the limit of  $\beta \rightarrow 0$  throughout this study. Accordingly,  $\tilde{\zeta}_{\perp}(\tilde{x}, \tilde{y})/\tilde{\zeta}_{\parallel}(\tilde{x}, \tilde{y}) = 2$ .

## 4 Governing equations and nondimensionalisation

The equations governing the dynamics of the filament can be derived by imposing force and torque balances along the filament, as elaborated in Section S1 of the ESI.†

Scaling forces with  $A/L^2$ , lengths with  $L$ , and time with  $\xi_{\perp,0}L^4/A$ , we obtain the following dimensionless governing equations:

$$\frac{\partial \theta}{\partial t} + \theta_{ssss} - (\tau\theta_s)_s - 2\theta_s(\tau_s + \theta_s\theta_{ss} - \sigma) + G(\tau\theta_s - \theta_{sss}) = 0, \quad (3a)$$

$$\tau_{ss} + (\theta_s\theta_{ss})_s - \frac{1}{2}\theta_s(\tau\theta_s - \theta_{sss}) - G(\tau_s + \theta_s\theta_{ss} - \sigma) = 0, \quad (3b)$$

where  $G(x, y) = g_s/g$  indicates the relative change of viscosity over the filament arc length, and  $\sigma = fL^3/A$  represents the dimensionless active force scaled by elastic forces. In the absence of viscosity gradients,  $G = 0$ , eqn (3) recovers to the governing equations in ref. 33. Imposing a vertical viscosity gradient defined in eqn (1),

$$G(x, y) = \frac{1}{g} \frac{\partial g}{\partial x} \frac{\partial x}{\partial s} = \frac{2\gamma}{1 + \gamma(2x - 1)} \cos \theta, \quad (4)$$

where we have employed  $x(s) = \int_0^s \cos \theta(s') ds'$  to compute  $\partial x/\partial s$ . Similarly, for a horizontal viscosity gradient, we derive

$$G(x, y) = \frac{2\gamma}{1 + 2\gamma y} \sin \theta. \quad (5)$$

Eqn (3) is solved with boundary conditions (BCs). At the filament tip ( $s = 1$ ), force- and torque-free conditions hold, corresponding to:

$$s = 1: \quad \tau = \theta_s = \theta_{ss} = 0. \quad (6)$$

At the clamped end ( $s = 0$ ), the BCs are:

$$s = 0: \quad \tau\theta_s - \theta_{sss} = \tau_s + \theta_s\theta_{ss} - \sigma = \frac{\partial \theta}{\partial t} = 0. \quad (7)$$

We adopt a Chebyshev spectral method to discretize the governing equations in space, combining an implicit–explicit temporal scheme, as detailed in Section S2 of the ESI.†

## 5 Linearisation of the governing equations

Subject to a sufficient small active force, the filament maintains a stationary and straight equilibrium state. To investigate its stability, we linearise the governing equations eqn (3) around this equilibrium. Assuming small filament deformation, we adopt a linear expansion:  $\theta = \theta^{(0)} + \varepsilon\theta^{(1)}(s, t)$  and  $\tau = \tau^{(0)}(s) + \varepsilon\tau^{(1)}(s, t)$ , where  $(\theta^{(0)}, \tau^{(0)})$  and  $(\theta^{(1)}, \tau^{(1)})$  denote the zero-th and first order solutions, respectively, with  $\varepsilon \ll 1$  being a small parameter.

Upon linearisation, we obtain  $\theta^{(0)} = 0$  and

$$\tau_{ss}^{(0)} - G^{(0)}(\tau_s^{(0)} - \sigma) = 0, \quad (8)$$

with

$$G^{(0)} = \begin{cases} 2\gamma/[1 + \gamma(2x - 1)] & \text{(vertical gradient),} \\ 0 & \text{(horizontal gradient).} \end{cases} \quad (9)$$

From eqn (8) and the BCs, we obtain  $\tau^{(0)} = \sigma(s - 1)$ .

Knowing the base state, we derive the governing equation for  $\theta^{(1)}$ ,

$$\frac{\partial \theta^{(1)}}{\partial t} + \theta_{ssss}^{(1)} - \sigma\theta_s^{(1)} - \sigma(s - 1)\theta_{ss}^{(1)} + G^{(0)}[\sigma(s - 1)\theta_s^{(1)} - \theta_{sss}^{(1)}] = 0, \quad (10)$$

where  $\theta^{(1)}$  and  $\tau^{(1)}$  have been decoupled. Considering eqn (9) and (10) together, we recognise that the horizontal viscosity gradient does not affect the linear stability because of the vanishing  $G^{(0)}$ , but only the vertical gradient does.

The linearised BCs for  $\theta^{(1)}$  are:

$$s = 0: \quad \sigma\theta_s^{(1)} + \theta_{sss}^{(1)} = \frac{\partial \theta^{(1)}}{\partial t} = 0, \quad (11a)$$

$$s = 1: \quad \theta_s^{(1)} = \theta_{ss}^{(1)} = 0. \quad (11b)$$

Employing the normal form  $\theta^{(1)}(s, t) = \hat{\theta}(s)\exp(\omega t)$  with  $\omega$  the complex growth rate, we arrive at a classical eigenvalue problem,

$$\omega\hat{\theta} + \hat{\theta}_{ssss} - \sigma\hat{\theta}_s - \sigma(s - 1)\hat{\theta}_{ss} + G^{(0)}[\sigma(s - 1)\hat{\theta}_s - \hat{\theta}_{sss}] = 0. \quad (12)$$

The leading (most unstable) eigenvalue and its associated eigenmode can be obtained by numerically solving eqn (12) using the same spectral discretization as for solving eqn (3), see the ESI,† Section S2.

## 6 Results: nonlinear simulations and linear stability analysis

We numerically solve eqn (3) to examine the effect of viscosity gradient on the spontaneous filament oscillation, particularly focusing on the instability threshold. These nonlinear simulations are corroborated by linear stability analysis (LSA). We then employ



a reduced-order bead-spring model to seek deeper insights into the instability mechanism modulated by the viscosity gradient. The effects of vertical and horizontal viscosity gradients will be demonstrated separately below, starting from the former.

### 6.1 Filament in a vertical viscosity gradient

Fig. 3(a) shows how the complex growth rate  $\omega$  of perturbation depends on the dimensionless active force  $\sigma$  and the vertical viscosity gradient  $\gamma \in [-0.6, 0.6]$ . The vertical dashed lines represent the critical active force  $\sigma_c$ , beyond which spontaneous filament oscillation emerges due to instability. Close to this threshold, the values of  $\sigma$  predicted by LSA and nonlinear simulations show excellent agreement.

More importantly, a negative correlation between the instability threshold  $\sigma_c$  and the gradient  $\gamma$  is clearly observed. This trend is

further revealed in Fig. 3(b), showing that both  $\sigma_c$  and the frequency of the corresponding most unstable eigenmode  $\text{Im}(\omega_c)/2\pi$  at the onset decrease with  $\gamma$  monotonically. This suggests that a viscosity profile increasing from the base to the tip of the filament destabilises the system, thereby promoting its self-oscillation.

### 6.2 Filament in a horizontal viscosity gradient

The scenario of linear instability is independent of the horizontal viscosity gradient, as indicated by eqn (9). This independence near the onset of instability is confirmed by the numerically computed  $\omega$  versus  $\sigma$  (see the ESI,† Section S4). Notably, even when  $\sigma$  substantially exceeds the threshold, the influence of the viscosity gradient on  $\omega$  remains marginal.

The horizontal viscosity gradient breaks the left-right symmetry in a forced fashion; hence, the filament exhibits asymmetric beating as expected. However, this asymmetry remains mild even at a substantial gradient level of  $\gamma = 0.6$ , as illustrated by the bottom panel of Fig. 4(a). Intuitively, the asymmetric beating resembling the recover-and-stroke pattern of a natural cilium would pump the fluid horizontally. To characterise this pumping effect, we calculate the average volume flow rate over one oscillation period  $T = 2\pi/\text{Im}(\omega)$ :<sup>40,41</sup>

$$Q = \frac{\xi_{\perp,0}L}{\pi\mu_0A} \times \frac{1}{T} \int_0^T \int_0^1 \frac{x(\mathbf{f}_h \cdot \mathbf{e}_y)}{g} ds dt. \quad (13)$$

As evidenced in Fig. 4(b), the flow rate  $Q$  exhibits an almost linear increase with the dimensionless active force  $\sigma$  above the threshold. The fluid is propagated along the viscosity gradient. Moreover, the net flow rate grows with the viscosity gradient  $\gamma$ , following an approximately linear relationship, as illustrated in Fig. 4(c).

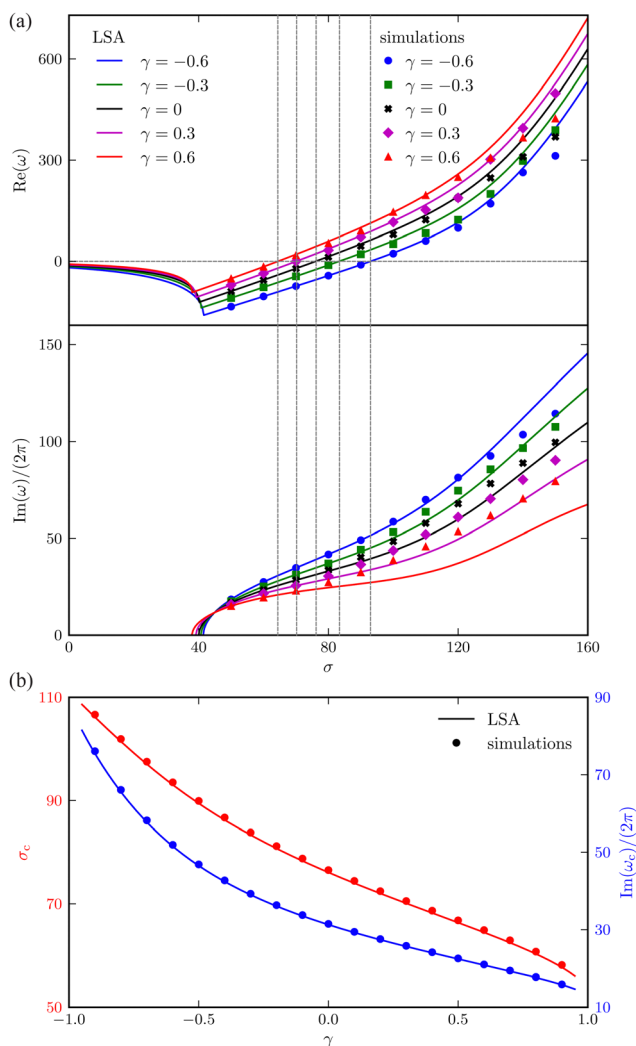
## 7 Reduced-order model

In the preceding section, we show the significant effect of vertical viscosity gradient on the onset of instability. In this section, we examine this effect further based on a reduced-order minimal model. Similar models have been employed by other studies on the behaviour of flexible structures in viscous fluids.<sup>32,42–50</sup>

As shown in Fig. 5, this model comprises two beads numbered 1 and 2 sequentially connected to two links of length  $L/2$ . Akin to the full model, only planar motion restricted to the  $xy$  plane is considered. To incorporate elasticity, the links are joined by a torsional spring of stiffness  $k$ . Besides, the lower link is attached to the base using a similar spring. The  $i$ -th ( $i = 1, 2$ ) bead is centred at  $\tilde{\mathbf{r}}_i$ . The angles  $\theta_1$  and  $\theta_2$  represent the orientation of the lower and upper links, respectively, relative to the  $\mathbf{e}_x$  axis.

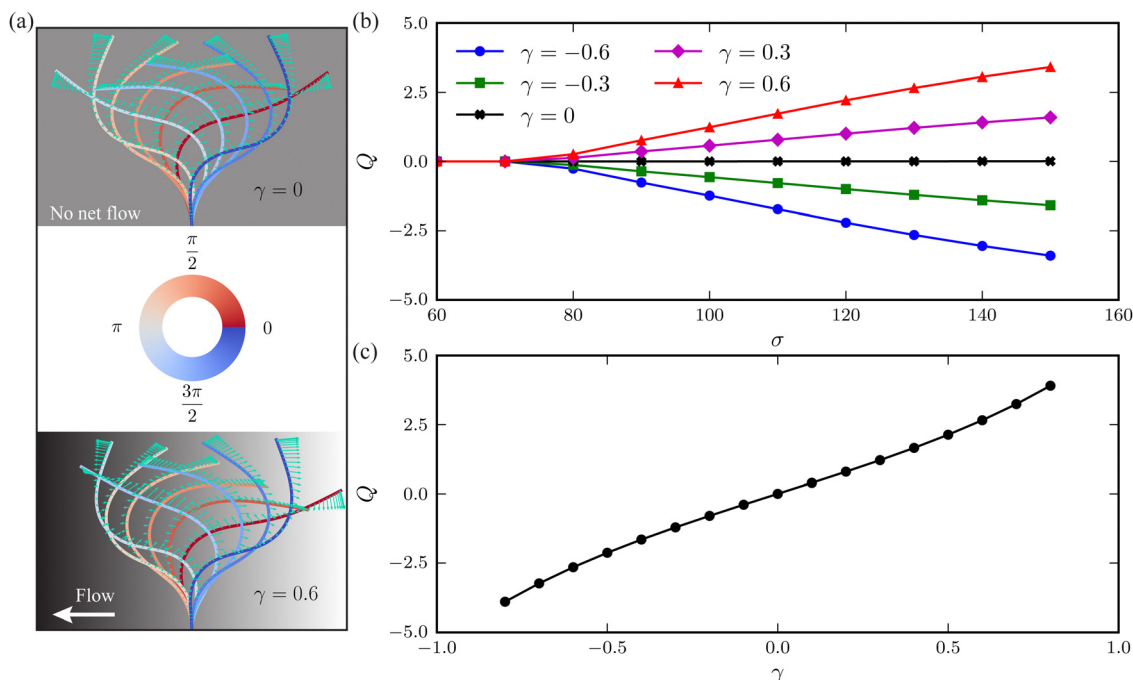
A follower force  $\tilde{\mathbf{F}}_a = -F_a(\cos\theta_2, \sin\theta_2)$  is imposed on bead 2 to represent the active force, persistently opposite to the tangent of the upper link. The hydrodynamic force exerted on bead  $i$  takes the form  $\tilde{\mathbf{F}}_{h,i} = -\tilde{\zeta}_i \partial \tilde{\mathbf{r}}_i / \partial t$ , with  $\tilde{\zeta}_i$  indicating the drag coefficient of bead  $i$ . Considering the vertical viscosity gradient,  $\tilde{\zeta}_i/\zeta_0 = 1 + \gamma(2\tilde{\mathbf{r}}_i \cdot \mathbf{e}_x/L - 1)$ . Here,  $\zeta_0$  represents the reference drag coefficient.

Employing the principle of virtual work, or the Euler-Lagrangian equations detailed in the Appendix, we derive the dimensionless



**Fig. 3** Comparison between the LSA (lines) and numerical results (symbols) for an active filament. (a) The real part (growth rate of the perturbation) of the most unstable eigenvalues, and corresponding frequency of oscillation (imaginary part divided by  $2\pi$ ), versus the dimensionless active force  $\sigma$  for different  $\gamma$ . The vertical dashed lines indicate the critical  $\sigma_c$  corresponding to the onset of instability. (b) Critical  $\sigma_c$  and oscillation frequency  $\text{Im}(\omega_c)/(2\pi)$  as a function of  $\gamma$ .



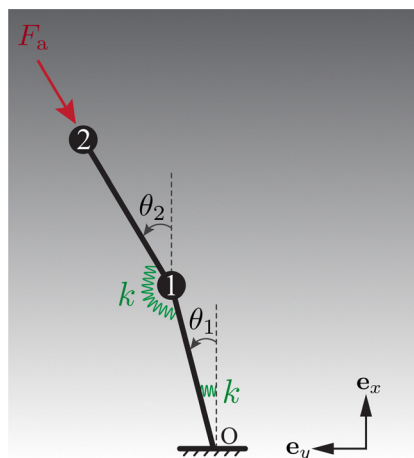


**Fig. 4** (a) Time-lapse of filament waveforms over one beating cycle for two viscosity gradients  $\gamma = 0$  (top panel) and  $\gamma = 0.6$  (bottom panel) at a dimensionless active force  $\sigma = 130$ . The black and white background suggests the fluid viscosity, where black corresponds to a higher viscosity. Arrows indicate the instantaneous velocities of the filament's centerline. (b) Average volume flow rate  $Q$  versus  $\sigma$  at varying viscosity gradients  $\gamma$ . (c) Dependence of  $Q$  on  $\gamma$  when  $\sigma = 130$ .

governing equations for the reduced-order model:

$$\begin{aligned} \Sigma \sin(\theta_1 - \theta_2) - 2\theta_1 + \theta_2 - (1 - \gamma + \gamma \cos \theta_1) \dot{\theta}_1 \\ - [1 - \gamma + \gamma(\cos \theta_1 + \cos \theta_2)] [\dot{\theta}_1 + \dot{\theta}_2 \cos(\theta_2 - \theta_1)] = 0, \end{aligned} \quad (14a)$$

$$\theta_1 - \theta_2 - [1 - \gamma + \gamma(\cos \theta_1 + \cos \theta_2)] [\dot{\theta}_2 + \dot{\theta}_1 \cos(\theta_2 - \theta_1)] = 0, \quad (14b)$$



**Fig. 5** The reduced-order model comprises two torsional springs and two beads, encapsulating the minimal physical ingredients of elasto-hydrodynamics: elastic torques generated by the springs and hydrodynamic forces exerted on the moving beads.

where  $\Sigma = F_a L / 2k$  represents the dimensionless active force. Here, we have chosen  $\zeta_0 L^2 / 4k$  as the characteristic time scale, and the overdot denotes taking a derivative with respect to time. Linearising eqn (14) about the equilibrium configuration  $\theta_1 = \theta_2 = 0$ , we examine the stability. Using the normal mode approach, we substitute  $\theta_{1,2} = \hat{\theta}_{1,2} \exp(\omega t)$  in the linearised equation and obtain,

$$\Sigma(\hat{\theta}_1 - \hat{\theta}_2) - 2\hat{\theta}_1 + \hat{\theta}_2 - \omega[(2 + \gamma)\hat{\theta}_1 + (1 + \gamma)\hat{\theta}_2] = 0, \quad (15a)$$

$$\hat{\theta}_1 - \hat{\theta}_2 - \omega(1 + \gamma)(\hat{\theta}_1 + \hat{\theta}_2) = 0. \quad (15b)$$

Non-trivial solutions to eqn (15) exist when the determinant of the system is zero, *i.e.*,  $1 + [6 + \gamma(5 - 2\Sigma) - 2\Sigma]\omega + (1 + \gamma)\omega^2 = 0$ , leading to the complex growth rate:

$$\omega_{\pm} = \Sigma - \frac{5}{2} - \frac{1}{2}(1 + \gamma)^{-1} \pm \sqrt{\left[\Sigma - \frac{5}{2} - \frac{1}{2}(1 + \gamma)^{-1}\right]^2 - (1 + \gamma)^{-1}}. \quad (16)$$

The complex growth rate  $\omega$  implies five stability regimes depending on  $\Sigma$  and  $\gamma$ :

1. Stable, when  $\Sigma \leq \frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1} - (1 + \gamma)^{-\frac{1}{2}}$ :  
 $\text{Re}(\omega_{\pm}) < 0$  and  $\text{Im}(\omega_{\pm}) = 0$ , corresponding to a stable state.
2. Decaying oscillations, when  $\frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1} - (1 + \gamma)^{-\frac{1}{2}} < \Sigma < \frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1}$ :  
 $\text{Re}(\omega_{\pm}) < 0$  and  $\text{Im}(\omega_{\pm}) \neq 0$ , indicating decaying oscillations.
3. Stable periodic oscillations, when  $\Sigma = \frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1}$ :



$\text{Re}(\omega_{\pm}) = 0$  and  $\text{Im}(\omega_{\pm}) = \pm(1 + \gamma)^{-\frac{1}{2}}$ , and the system oscillates with a constant amplitude. Besides, increasing  $\gamma$  decreases  $\Sigma_c$  and the oscillation frequency.

4. Exponentially growing oscillations, when  $\frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1} < \Sigma < \frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1} + (1 + \gamma)^{-\frac{1}{2}}$ :

$\text{Re}(\omega_{\pm}) > 0$  and  $\text{Im}(\omega_{\pm}) \neq 0$ , indicating exponentially growing oscillations.

5. Unstable, when  $\Sigma \geq \frac{5}{2} + \frac{1}{2}(1 + \gamma)^{-1} + (1 + \gamma)^{-\frac{1}{2}}$ :

$\text{Re}(\omega_{\pm}) > 0$  and  $\text{Im}(\omega_{\pm}) = 0$ , the amplitude grows without bound, indicating an unstable system that diverges.

As illustrated in Fig. 6, the complex growth rate  $\omega$  versus  $\Sigma$  and  $\gamma$  reveals the first four regimes discussed above. Moreover, the trend qualitatively agrees with that of the continuum filament model shown in Fig. 3(a). Based on the capacity of the reduced-order model to capture the characteristic behaviours of the full model, we will further exploit the model to understand the physical mechanisms underlying the destabilisation of the vertical viscosity gradient.

### 7.1 Physical mechanisms of destabilising vertical viscosity gradient

Upon applying a vertical viscosity gradient  $\gamma > 0$ , the fluid surrounding bead 2 (upper) becomes more viscous, whereas the viscosity around bead 1 (lower) remains unchanged, because bead 1 stays near its equilibrium state close to the reference position  $\tilde{y} = L/2$ . Naturally, we may wonder whether the stabilisation results from an overall viscosity enhancement. Intriguingly, owing to the independence of  $\Sigma$  on the viscosity, increasing the viscosity homogeneously by the same factor (thus eliminating the gradient) does not alter the onset  $\Sigma_c$  of instability. A uniform viscosity enhancement merely affects the oscillation frequency. Consequently, we deduce that it is the gradient of viscosity, rather than its overall magnitude, that affects the stability, as we will explain below based on an energetic point of view.

In the Stokes flow regime, the energy of the bead-spring system comprises the elastic energy,  $E$ , of the springs, because the kinetic energy of the beads vanishes. The temporal growth rate of  $E$  is powered by the active force and simultaneously dissipated by the viscous force, namely,

$$\dot{E} = P_a - P_v, \quad (17)$$

where  $P_a$  denotes the power of active force, and  $P_v$  the viscous dissipation rate. They take the form

$$P_a = \Sigma \sin(\theta_1 - \theta_2) \dot{\theta}_1, \quad (18a)$$

$$P_v = 2\mathcal{F}, \quad (18b)$$

where  $\mathcal{F}$  is the dissipation function derived in the Appendix. Near the equilibrium where  $\theta_1 \approx \theta_2 \ll 1$ , they can be approximated by

$$P_a \approx (1 + \gamma)\Sigma(\dot{\theta}_1 + \dot{\theta}_2)\dot{\theta}_1, \quad (19a)$$

$$P_v \approx \dot{\theta}_1^2 + (1 + \gamma)(\dot{\theta}_1 + \dot{\theta}_2)^2. \quad (19b)$$

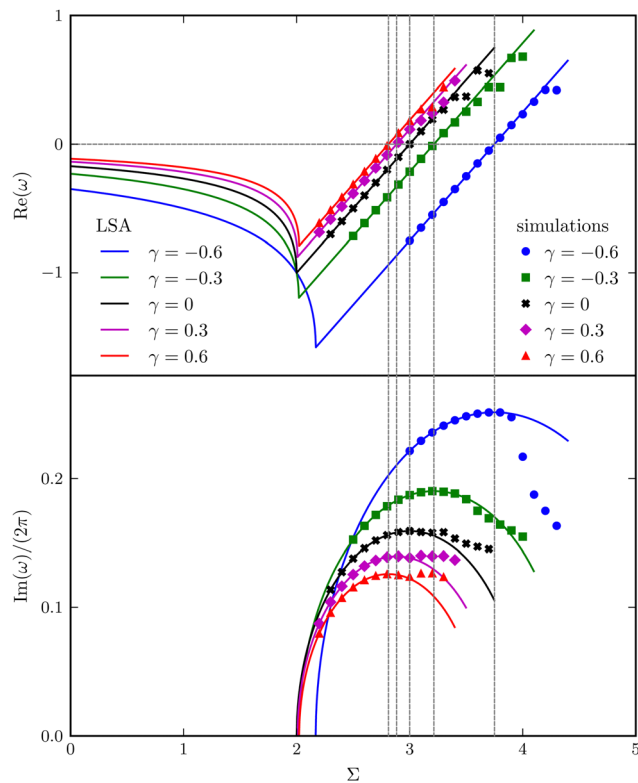


Fig. 6 Real and imaginary parts of the complex growth rate  $\omega$  versus the dimensionless active force  $\Sigma$  with varying viscosity gradients  $\gamma$ , obtained from LSA (symbols) and simulations (lines) for the bead-spring model.

Eqn (19) allows us to analyse the linear stability from the energetic perspective. Evidently,  $P_v$  is always positive, indicating the consistent stabilisation due to the viscous dissipation. On the other hand, the active power  $P_a$  can generally switch sign, with  $P_a > 0$  corresponding to the destabilisation triggered by the active force. Furthermore, eqn (19) implies that the active force is the only destabiliser in this system, as indeed aligning with the physical intuition.

The viscosity gradient  $\gamma$  affects the stability by modulating the  $(1 + \gamma)$ -dependent  $P_a$  and  $P_v$ . In both cases,  $(1 + \gamma)$  appears as a prefactor of active force strength  $\Sigma$  in the former, and of a portion of the viscous power in the latter. Hence, a positive gradient  $\gamma$  amplifies both  $P_a$  and  $P_v$ , whereas a negative  $\gamma$  decreases both.

In the limit of  $\gamma \rightarrow -1$ , the active power reduces to zero whereas the viscous dissipation rate remains finite. This leads to a stable system, irrespective of the active force strength. This picture delineates the stabilising effect of a decreasing  $\gamma$ , particularly when it approaches its lower limit. However, beyond this limit, eqn (19) fails to explicitly elucidate the consequences of altering  $\gamma$  on the system's power balance and stability. To address this gap, we analyse the numerical data for  $\Sigma = 3$  at  $\gamma = -0.01, 0$ , and  $0.01$ . Our findings, presented in Fig. 7, depict the impact of varying  $\gamma$  on the temporal evolution of  $P_a$  and  $-P_v$ , as well as their summation  $\dot{E}$ . Notably, increasing  $\gamma$  leads to a more pronounced amplification of the active power  $P_a$  compared to the negative viscous dissipation rate  $-P_v$ , and



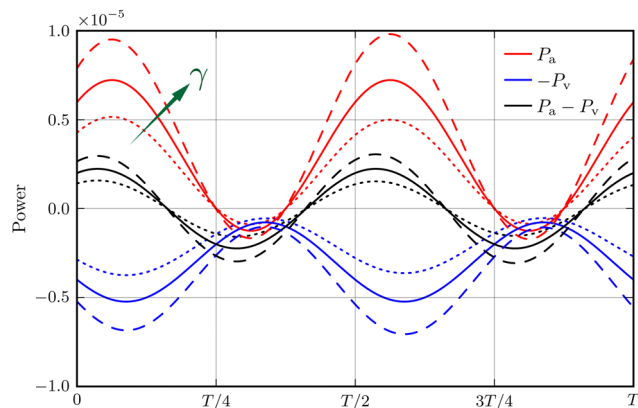


Fig. 7 Power balance within an oscillation period  $T$  of the bead-spring system, when the viscosity gradient  $\gamma = 0.01$  (dashed lines),  $\gamma = 0$  (solid lines), and  $\gamma = -0.01$  (dotted lines) at  $\Sigma = 3$ . Here,  $P_a$  (red) and  $P_v$  (blue) represent the active power and viscous dissipation rate, respectively.

concomitantly more intense oscillations in  $\dot{E}$ . This observation reflects, semi-phenomenologically, the destabilising effect of a larger  $\gamma$ .

## 8 Conclusions and discussion

In this work, we have numerically studied the spontaneous oscillation of an anchored active filament in viscous fluids with a linearly varying viscosity. The activity of the filament is modelled by a point follower force or a uniformly distributed force density. In both scenarios, our simulations reveal that a positive viscosity gradient, extending from the filament's base to tip, facilitates its self-oscillation, namely, decreasing the minimum activity required to trigger the elasto-hydrodynamic instability. On the other hand, the viscosity gradient orthogonal to the orientation of the filament does not alter the onset of instability. However, by introducing forced symmetry breaking, this orthogonal gradient enables asymmetric beating of the filament, consequently generating a net fluid pumping along the gradient.

The numerically demonstrated effects of viscosity gradients on the self-oscillatory stability of the active filament are supported by our LSA. Numerical results and predictions of LSA agree excellently near the instability threshold. To better understand how viscosity gradients affect the instability, we have used a reduced-order model. This model reproduces the various stability regimes identified in the original filament model and provides analytical insights into the interplay between the destabilising active force and the stabilising viscous dissipation. Specifically, by increasing the vertical viscosity gradient, we observe an enhanced temporal oscillation in both the active power and the viscous dissipation rate. This enhancement is more pronounced for the former than for the latter, resulting in a more intense oscillation of the elastic energy's growth.

While our study is inspired by respiratory cilia in heterogeneous bodily fluids, it does not delve into the biological implications within this specific context. As a highly complex

fluid, ASL is not only heterogeneous and layered,<sup>51</sup> but also features viscoelasticity<sup>52</sup> and potentially shear-thinning properties.<sup>53</sup> The rheological behaviour of ASL is substantially more complex compared to the Newtonian fluid model with uniform viscosity gradient used here. Notably, a similar Newtonian model with spatial viscosity variation has been employed in the investigation of mucociliary clearance enabled by beating cilia.<sup>53</sup>

To investigate the effects of viscosity gradients on ciliary oscillation, we have employed the simplified follower force approach, which may lack realism. For a deeper, biologically pertinent understanding, future work should explore models of ciliary oscillation driven by dynein motors.<sup>11–22</sup>

It is worth noting that this study resonates with the recent investigations on microswimmers' self-propelling in viscous fluids with viscosity gradients.<sup>54–61</sup> Consequently, a pertinent inquiry is the behaviour of a spontaneously beating flagellum, with or without a cell body, in environments featuring continuous or sharp viscosity gradients. In the absence of the gradient, a single flagellum is expected to swim along a spontaneously emerging direction depending on its initial orientation. We hypothesise that introducing an external viscosity gradient could direct the flagellum to move along a preferential direction. This hypothesis will be tested in our future studies.

## Conflicts of interest

There are no conflicts to declare.

## Appendix

### Dynamics of the bead-spring system

The governing equations of the bead-spring model are derived based on the Euler–Lagrangian framework:<sup>62</sup>

$$\frac{d}{dt} \left( \frac{\partial \tilde{\mathcal{L}}}{\partial \dot{\theta}_i} \right) - \frac{\partial \tilde{\mathcal{L}}}{\partial \theta_i} + \frac{\partial \tilde{\mathcal{F}}}{\partial \theta_i} = \tilde{Q}_i, \quad (i = 1, 2). \quad (20)$$

Here,  $\tilde{\mathcal{L}}$  represents the Lagrangian of the system  $\tilde{\mathcal{L}} = \tilde{T}_{\text{kin}} - \tilde{V}$ , where  $\tilde{T}_{\text{kin}}$  and  $\tilde{V}$  denote the kinetic and potential energies, respectively. The generalised forces  $\tilde{Q}_i$  are expressed as:

$$\tilde{Q}_i \equiv \tilde{\mathbf{F}}_a \cdot \frac{\partial \tilde{\mathbf{r}}_2}{\partial \theta_i}, \quad (i = 1, 2), \quad (21)$$

where the follower force  $\tilde{\mathbf{F}}_a$  can be regarded as an external force of the system.

The dissipation function  $\tilde{\mathcal{F}}$  corresponding to eqn (20) is defined as<sup>62</sup>

$$\begin{aligned} \tilde{\mathcal{F}} &\equiv \frac{1}{2} \tilde{\zeta}_1 \left( \frac{d\tilde{\mathbf{r}}_1}{dt} \right)^2 + \frac{1}{2} \tilde{\zeta}_2 \left( \frac{d\tilde{\mathbf{r}}_2}{dt} \right)^2 \\ &= \frac{L^2}{8} \tilde{\zeta}_1 \dot{\theta}_1^2 + \frac{L^2}{8} \tilde{\zeta}_2 \left[ \dot{\theta}_1^2 + \dot{\theta}_2^2 + 2\dot{\theta}_1 \dot{\theta}_2 \cos(\theta_2 - \theta_1) \right]. \end{aligned} \quad (22)$$

This function remains non-negative, namely,  $\tilde{\mathcal{F}} \geq 0$ , offers two physical interpretations. Firstly, it represents the velocity-



dependent potential, wherein the hydrodynamic force can be computed as

$$\tilde{\mathbf{F}}_{h,i} = -\frac{\partial \tilde{\mathcal{F}}}{\partial \dot{\theta}_i}, \quad (i = 1, 2). \quad (23)$$

Secondly, it characterises the rate of total energy dissipation, specifically, the viscous dissipation rate  $\tilde{P}_v$  in this system,

$$\tilde{P}_v = 2\tilde{\mathcal{F}}. \quad (24)$$

Because the bead mass is neglected, the total kinetic energy of the system is zero,  $\tilde{T}_{\text{kin}} = 0$ . Hence,  $\tilde{\mathcal{L}} = -\tilde{V}$ , with the potential energy  $\tilde{V}$  written as:

$$\tilde{V} = \frac{1}{2}k\theta_1^2 + \frac{1}{2}k(\theta_1 - \theta_2)^2. \quad (25)$$

Besides, the two generalised forces corresponding to  $\theta_1$  and  $\theta_2$  are

$$\tilde{Q}_1 = -F_a \frac{L}{2} \sin(\theta_2 - \theta_1), \quad (26a)$$

$$\tilde{Q}_2 = 0. \quad (26b)$$

In addition, the drag coefficients of both beads under the viscosity gradient are:

$$\frac{\tilde{\zeta}_1}{\tilde{\zeta}_0} = 1 - \gamma + \gamma \cos \theta_1, \quad (27a)$$

$$\frac{\tilde{\zeta}_2}{\tilde{\zeta}_0} = 1 - \gamma + \gamma(\cos \theta_1 + \cos \theta_2). \quad (27b)$$

Choosing  $k$ ,  $2k/L$ , and  $\zeta_0 L^2/4k$  as the characteristic energy, force, and time, respectively, the dimensionless potential energy, generalised force, and dissipation function are given as follows:

$$V = \frac{1}{2}(2\theta_1^2 + \theta_2^2 - 2\theta_1\theta_2), \quad (28a)$$

$$Q_1 = \Sigma \sin(\theta_1 - \theta_2), \quad (28b)$$

$$\begin{aligned} \mathcal{F} = & \frac{1}{2}(1 - \gamma + \gamma \cos \theta_1)\dot{\theta}_1^2 + \frac{1}{2}[1 - \gamma + \gamma(\cos \theta_1 + \cos \theta_2)] \\ & \times [\dot{\theta}_1^2 + \dot{\theta}_2^2 + 2\dot{\theta}_1\dot{\theta}_2 \cos(\theta_2 - \theta_1)]. \end{aligned} \quad (28c)$$

By substituting eqn (28) into the dimensionless equation of eqn (20), we obtain eqn (14). The rate of work done by the follower force is

$$P_a = \mathbf{F}_a \cdot \dot{\mathbf{r}}_2 = Q_1 \dot{\theta}_1 = \Sigma \sin(\theta_1 - \theta_2) \dot{\theta}_1. \quad (29)$$

## Acknowledgements

The work is funded by the Singapore Ministry of Education Academic Research Fund Tier 2 grant (MOE-T2EP50122-0015).

L. Z. acknowledges the partial support from the Paris-NUS joint research grant (ANR-18-IDEX-0001 & A-0009528-01-00). The National Natural Science Foundation of China (Project no. 12202438) is also acknowledged. The computation of the work was performed on resources of the National Supercomputing Centre, Singapore (<https://www.nscc.sg>).

## Notes and references

- 1 D. Moreira and J. C. M. Pires, *Bioresour. Technol.*, 2016, **215**, 371–379.
- 2 C. M. Beal, I. Archibald, M. E. Huntley, C. H. Greene and Z. I. Johnson, *Earths Future*, 2018, **6**, 524–542.
- 3 R. W. Pierce and J. T. Turner, *Rev. Aquat. Sci.*, 1992, **6**, 139–181.
- 4 R. R. Strathmann, T. L. Jahn and J. R. Fonseca, *Biol. Bull.*, 1972, **142**, 505–519.
- 5 F. Hildebrandt, T. Benzing and N. Katsanis, *N. Engl. J. Med.*, 2011, **364**, 1533–1543.
- 6 R. Dreyfus, J. Baudry, M. L. Roper, M. Fermigier, H. A. Stone and J. Bibette, *Nature*, 2005, **437**, 862–865.
- 7 J. M. den Toonder and P. R. Onck, *Trends Biotechnol.*, 2013, **31**, 85–91.
- 8 W. Wang, Q. Liu, I. Tanasijevic, M. F. Reynolds, A. J. Cortese, M. Z. Miskin, M. C. Cao, D. A. Muller, A. C. Molnar, E. Lauga, P. L. McEuen and I. Cohen, *Nature*, 2022, **605**, 681–686.
- 9 S. Park, G. Choi, M. Kang, W. Kim, J. Kim and H. E. Jeong, *Microsyst. Nanoeng.*, 2023, **9**, 153.
- 10 W. Gilpin, M. S. Bull and M. Prakash, *Nat. Rev. Phys.*, 2020, **2**, 74–88.
- 11 C. J. Brokaw, *J. Exp. Biol.*, 1971, **55**, 289–304.
- 12 M. Hines and J. J. Blum, *Biophys. J.*, 1978, **23**, 41–57.
- 13 C. B. Lindemann, *J. Theor. Biol.*, 1994, **168**, 175–189.
- 14 S. Camalet and F. Jülicher, *New J. Phys.*, 2000, **2**, 324.
- 15 I. H. Riedel-Kruse, A. Hilfinger, J. Howard and F. Jülicher, *HFSP J.*, 2007, **1**, 192–208.
- 16 R. H. Dillon and L. J. Fauci, *J. Theor. Biol.*, 2000, **207**, 415–430.
- 17 P. V. Bayly and S. K. Dutcher, *J. R. Soc., Interface*, 2016, **13**, 20160523.
- 18 P. Sartori, V. F. Geyer, A. Scholich, F. Jülicher and J. Howard, *eLife*, 2016, **5**, e13258.
- 19 D. Oriola, H. Gadêlha and J. Casademunt, *R. Soc. Open Sci.*, 2017, **4**, 160698.
- 20 L. G. Woodhams, Y. Shen and P. V. Bayly, *J. R. Soc., Interface*, 2022, **19**, 20220264.
- 21 J. F. Cass and H. Bloomfield-Gadêlha, *Nat. Commun.*, 2023, **14**, 5638.
- 22 M. T. Gallagher, J. C. Kirkman-Brown and D. J. Smith, *PNAS Nexus*, 2023, **2**, pgad072.
- 23 B. Chakrabarti and D. Saintillan, *Phys. Rev. Lett.*, 2019, **123**, 208101.
- 24 Y. Man and E. Kanso, *Phys. Rev. Lett.*, 2020, **125**, 148101.
- 25 S. Maretvadakethope, Y. Hwang and E. E. Keaveny, *Phys. Rev. Fluids*, 2022, **7**, 053101.
- 26 T. A. Westwood and E. E. Keaveny, *Phys. Rev. Fluids*, 2021, **6**, L121101.





- 27 B. Chakrabarti, S. Fürthauer and M. J. Shelley, *Proc. Natl. Acad. Sci. U. S. A.*, 2022, **119**, e2113539119.
- 28 K. G. Link, R. D. Guy, B. Thomases and P. E. Arratia, *arXiv*, 2023, preprint, arXiv:2306.03178, DOI: [10.48550/arXiv.2306.03178](https://doi.org/10.48550/arXiv.2306.03178).
- 29 C. Wang, S. Gsell, U. D'Ortona and J. Favier, *J. Fluid Mech.*, 2023, **965**, A6.
- 30 L. E. Wallace, M. Liu, F. J. van Kuppeveld, E. de Vries and C. A. de Haan, *Trends Microbiol.*, 2021, **29**, 983–992.
- 31 M. Zanin, P. Baviskar, R. Webster and R. Webby, *Cell Host Microbe*, 2016, **19**, 159–168.
- 32 G. De Canio, E. Lauga and R. E. Goldstein, *J. R. Soc., Interface*, 2017, **14**, 20170491.
- 33 Y. Fily, P. Subramanian, T. M. Schneider, R. Chelakkot and A. Gopinath, *J. R. Soc., Interface*, 2020, **17**, 20190794.
- 34 E. O. Tuck, *J. Fluid Mech.*, 1964, **18**, 619–635.
- 35 G. K. Batchelor, *J. Fluid Mech.*, 1970, **44**, 419–440.
- 36 R. G. Cox, *J. Fluid Mech.*, 1970, **44**, 791–810.
- 37 R. G. Cox, *J. Fluid Mech.*, 1971, **45**, 625–657.
- 38 J. P. K. Tillett, *J. Fluid Mech.*, 1970, **44**, 401–417.
- 39 C. Kamal and E. Lauga, *J. Fluid Mech.*, 2023, **963**, A24.
- 40 N. Liron, *J. Fluid Mech.*, 1978, **86**, 705–726.
- 41 D. Smith, J. Blake and E. Gaffney, *J. R. Soc., Interface*, 2008, **5**, 567–573.
- 42 Z. Peng, G. J. Elfring and O. S. Pak, *Soft Matter*, 2017, **13**, 2339–2347.
- 43 F. Ling, H. Guo and E. Kanso, *J. R. Soc., Interface*, 2018, **15**, 20180594.
- 44 Y. Man and E. Kanso, *Soft Matter*, 2019, **15**, 5163–5173.
- 45 H. A. Stone and L. Zhu, *J. Fluid Mech.*, 2020, **888**, A31.
- 46 Z. Liu, F. Qin and L. Zhu, *Phys. Rev. Fluids*, 2020, **5**, 124101.
- 47 K. Qin, Z. Peng, Y. Chen, H. Nganguia, L. Zhu and O. S. Pak, *Soft Matter*, 2021, **17**, 3829–3839.
- 48 W.-Z. Fang, F. Viola, S. Camarri, C. Yang and L. Zhu, *Non-linear Dyn.*, 2021, **106**, 2005–2019.
- 49 S. Hu, J. Zhang and M. J. Shelley, *Soft Matter*, 2022, **18**, 3605–3612.
- 50 S. Hu and F. Meng, *J. Fluid Mech.*, 2023, **966**, A23.
- 51 B. Button, L.-H. Cai, C. Ehre, M. Kesimer, D. B. Hill, J. K. Sheehan, R. C. Boucher and M. Rubinstein, *Science*, 2012, **337**, 937–941.
- 52 Z. Chen, M. Zhong, Y. Luo, L. Deng, Z. Hu and Y. Song, *Respir. Res.*, 2019, **20**, 274.
- 53 R. Chatelin and P. Poncet, *J. Biomech.*, 2016, **49**, 1772–1780.
- 54 B. Liebchen, P. Monderkamp, B. ten Hagen and H. Löwen, *Phys. Rev. Lett.*, 2018, **120**, 208002.
- 55 C. Datt and G. J. Elfring, *Phys. Rev. Lett.*, 2019, **123**, 158006.
- 56 R. Dandekar and A. M. Ardekani, *J. Fluid Mech.*, 2020, **895**, R2.
- 57 M. R. Stehnach, N. Waisbord, D. M. Walkama and J. S. Guasto, *Nat. Phys.*, 2021, **17**, 926–930.
- 58 C. E. López, J. Gonzalez-Gutierrez, F. Solorio-Ordaz, E. Lauga and R. Zenit, *Phys. Rev. Fluids*, 2021, **6**, 083102.
- 59 R. V. More and A. M. Ardekani, *Annu. Rev. Fluid Mech.*, 2023, **55**, 157–192.
- 60 J. Gong, V. A. Shaik and G. J. Elfring, *Sci. Rep.*, 2023, **13**, 596.
- 61 C. Feng, J. J. Molina, M. S. Turner and R. Yamamoto, *Phys. Fluids*, 2023, **35**, 051903.
- 62 H. Goldstein, C. Poole and J. Safko, *Classical Mechanics*, Addison Wesley, 3rd edn, 2002.

