Separation of Microscale Chiral Objects by Shear Flow

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We show that plane parabolic flow in a microfluidic channel causes nonmotile, helically shaped bacteria to drift perpendicular to the shear plane. Net drift results from the preferential alignment of helices with streamlines, with a direction that depends on the chirality of the helix and the sign of the shear rate. The drift is in good agreement with a model based on resistive force theory, and separation is efficient (>80%) and fast (<2 s). We estimate the effect of Brownian rotational diffusion on chiral separation and show how this method can be extended to separate chiral molecules.

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Many biochemically active molecules are naturally chiral and can only bind to target chiral molecules of a specific handedness [1]. The other enantiomer (i.e., the molecule having opposite handedness) may be inactive or cause undesirable effects. Chemical synthesis of chiral molecules usually produces a racemic mixture, with equal amounts of both enantiomers, and their separation based on chirality is of importance in fields ranging from agriculture to food and pharmaceutical industries. Currently favored approaches rely on gas, liquid, or capillary electromigration chromatography [2], requiring costly chiral media. Thus, simpler, alternative approaches to chiral separation are desirable.

Several alternative proposals for chiral separation exploit hydrodynamic forces. Some of these, yet untested experimentally, rely on the presence of a surface [3] or array of microvortices [4], while there has been successful chiral separation of centimeter-sized crystals in a rotating drum [5]. Other methods [6,7] stem from the prediction that a chiral particle in a simple shear flow experiences a lateral drift [8]. However, the feasibility of this approach has remained questionable, as measurements in Couette cells reported that the drift of millimeter-sized chiral objects [9] and the forces on centimeter-sized ones [10] differ from predictions by 2 orders of magnitude [9] or even in sign [10].

Here we report that microscale chiral objects, 3 orders of magnitude smaller than previous studies [9,10], experience a lateral drift in a microfluidic shear flow, and that the magnitude of the drift is in agreement with our theory. Previous work has demonstrated the ability of microfluidics to separate and sort colloids by size [11], spermatozoa by motility [12], and microbes by their preference for dissolved chemicals [13]. Our method uses microchannels to sort particles by chirality. We show that an enantiomer drifts with direction determined by the local shear, demonstrate the feasibility of this method for chiral separation, and indicate how the high shear rates achievable in micro-

channels ($>10^6 \text{ s}^{-1}$ [14]) allow our method to be extended to smaller scales (<40 nm).

The origin of chirality-dependent drift at low Reynolds number can be simply understood for the case of a helix. In a shear flow, objects undergo periodic rotations known as Jeffery orbits [15]: a sphere rotates with constant angular velocity, whereas for an elongated body, such as a helix, the velocity depends on orientation. The more elongated a body, the longer its residence time when aligned with streamlines. In addition to rotating in a Jeffery orbit, a helix drifts across streamlines. To see why, consider a right-handed helix aligned with a simple shear flow (Fig. 1), and decompose the velocity at a segment of the helix into components perpendicular and parallel to the segment. Drag on a thin rod in low Reynolds number flow is anisotropic, with a greater resistance when oriented

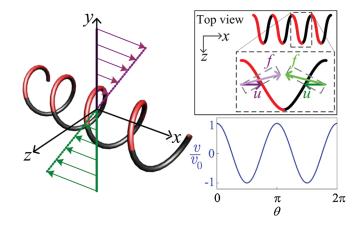


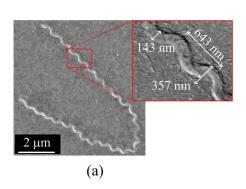
FIG. 1 (color). Schematic of a right-handed helix in simple shear flow. Red shading and black shading show the top and bottom halves of the helix. Upper inset: The net force acting on one pitch of the helix is along $-\hat{\mathbf{z}}$. Lower inset: Predicted normalized drift velocity v/v_0 versus helix orientation θ . θ is the angle between a helix in the x-y plane and the flow, and v_0 is the drift velocity of a helix aligned with the flow ($\theta = 0$).

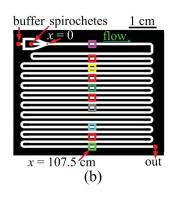
perpendicular rather than parallel to the flow [16]. Since the flows at the top (red) and bottom (black) halves of the helix are in opposite directions, both halves have a drag component along $-\hat{\mathbf{z}}$ (Fig. 1, top inset). Thus, a righthanded helix aligned with the shear flow drifts in the $-\hat{z}$ direction. Reversing the chirality of the helix or the sign of the shear produces a drift along $+\hat{\mathbf{z}}$. Furthermore, the drift depends on the orientation of the helix. A righthanded helix aligned with the y axis drifts along $+\hat{\mathbf{z}}$. In general, the drift velocity v of a helix lying in the x-y plane is a function of the angle θ between the helix and the flow (Fig. 1, bottom inset). Hence, the mean drift \bar{v} depends on the distribution of the orientation of the helix relative to the flow. When the helix is preferentially aligned with streamlines, for example, due to Jeffery orbits, it has a net drift. When all orientations are equally likely, for example, due to strong Brownian rotational diffusion, the net drift is zero [7]. Here we use microscale objects to demonstrate the chirality-induced drift and determine the size of the smallest helices that can be separated using this principle.

We use a nonmotile, right-handed, helically shaped bacterium, Leptospira biflexa flaB mutant [17] [Fig. 2(a)], as the microscale chiral object. These spirochetes have an average length of 16 μ m, thickness of 150 nm, and diameter of 200 nm (Fig. 2 and [18]). To expose spirochetes to a shear flow, we fabricated a 110 cm long, serpentine-shaped microfluidic channel via soft lithography [19] [Figs. 2(b) and 2(c)]. The channel has rectangular cross section of depth $H = 90 \mu m$ and width W = 1 mm, resulting in a parabolic flow, $u(y) = (3U/2)[1 - (2y/H)^2]$, with mean speed $U = 3.09 \text{ mm s}^{-1}$. Because of the high aspect ratio W/H, the velocity u is uniform in z, except for a 150 μ m layer adjacent to the two sidewalls [20], inconsequential for these experiments. The shear rate $\dot{\gamma}(y) = du/dy =$ $-12yU/H^2$ varies linearly along y (between $\pm 206 \text{ s}^{-1}$), with $\dot{\gamma} < 0$ for y > 0, and $\dot{\gamma} > 0$ for y < 0 [Fig. 2(c)]. The microchannel is equipped with two inlets, one for spirochetes and one for buffer solution (EMJH liquid medium [17]). Spirochetes were introduced through a microinjector, forming a band of width $w \approx 100 \ \mu \text{m}$ at the center of the channel, uniform over the depth, with buffer streams on either side. As $\dot{\gamma}$ varies with y, one expects the drift to vary with vertical position, and spirochetes in the top and bottom halves of the channel to drift in opposite directions.

We imaged the spirochete population using phase contrast microscopy with a 40× objective and a CCD camera. At ten locations along the channel [Fig. 2(b)], sets of 50 images were recorded at 1.5 s intervals at three depths: y =-H/4 ("bottom"), 0 ("mid"), and H/4 ("top"). Experiments revealed a clear and reproducible drift, with direction determined by the sign of the local shear rate. A superposition of 50 images acquired at x = 107.5 cm shows drift of comparable magnitude and opposite direction in the top and bottom halves, and no drift at midplane, where $\dot{\gamma} = 0$ [Fig. 3(c) and supplemental movie [21]]. Using image analysis to obtain the z position of individual spirochetes, we determined the probability density function of the across-channel distribution of cells, their mean position \bar{z} , and the one-standard-deviation spread σ , for each imaging location (x, y). The drift of the top and bottom populations increases monotonically with x or, equivalently, with time. The apparent spread of the spirochete distribution with increasing x [Figs. 3(b) and 3(c)] is a result of the variation of drift with depth combined with the objective's finite depth of focus [21]. Brownian motion was negligible ($\sigma \approx 6 \mu \text{m}$ over x =107.5 cm [21]).

Since drift (\bar{z}) increases linearly with distance x along the channel [Fig. 3(d)], the mean drift velocity is constant $(\bar{v} = 0.41 \ \mu \text{m s}^{-1})$, resulting in a linearly increasing separation distance Δz between the top and bottom populations [Fig. 3(e)]. Control experiments with neutrally buoyant 1 μ m spheres revealed no drift, irrespective of the vertical position in the channel [Fig. 3(d), empty squares], and thus no separation [Fig. 3(e), empty squares], supporting the conclusion that drift is a result of chirality. We further verified that the 180° turns in the channel are inconsequential for these observations [21].





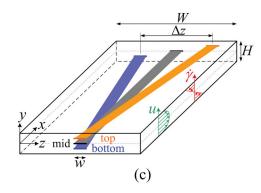


FIG. 2 (color). (a) Scanning electron micrograph (SEM) of *L. biflexa flaB* mutant, with typical dimensions (inset). The bent configuration is a result of SEM preparation and live organisms are nearly always straight. (b) Microchannel design with separate inlets for spirochetes and buffer. The color-coded squares refer to the locations of data collection [Fig. 3(b)]. (c) Schematic of the separation process (for one enantiomer) in the microchannel (W=1 mm, H=90 μ m, $w\approx 100$ μ m). The lateral drift direction depends on the sign of the shear $\dot{\gamma}$, resulting in divergence of top and bottom streams.

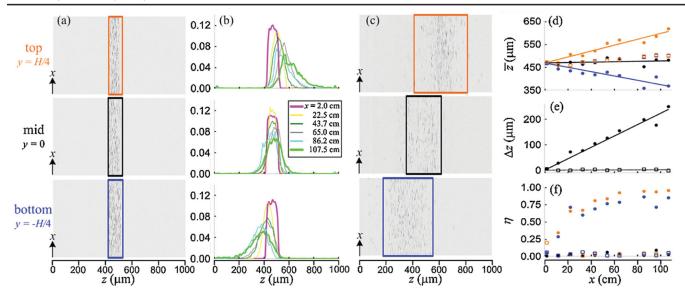


FIG. 3 (color). The observed spirochete distribution across the channel at (a) x = 2 cm and (c) x = 107.5 cm. Colors represent populations at different depths y: top (orange), mid (black), and bottom (blue) [see Fig. 2(c)]. The midpoint of each rectangle corresponds to the mean position (\bar{z}) of the distribution, while the half-width is 2 standard deviations (2σ). (b) Probability density function of spirochete distribution at various distances x along the channel [see Fig. 2(b)]. The thick pink and green profiles correspond to (a) and (c), respectively. (d)–(f) Quantification of lateral drift and separation along x. (d) Mean position of the population at three depths. (e) Divergence between the top and bottom populations. (f) Separation efficiency. In all panels, full circles refer to spirochete data, lines are linear fits, and empty squares are experiments with spherical beads. Over 10 000 spirochetes were imaged and located at each position (x, y).

To quantify how the bottom and top streams diverge, we define a separation efficiency $\eta = |N_L - N_R|/(N_L + N_R)$, where N_L and N_R are the numbers of spirochetes to the left and right of the mean position \bar{z} of the mid population (y=0), and η ranges from zero for no separation to one for perfect separation. We found η to increase with x for spirochetes, reaching a plateau at $\eta \approx 0.8$ after x=70 cm, while $\eta < 0.2$ for spheres [Fig. 3(f), empty squares]. This separation was achieved in 227 s, and experiments with $U=600 \text{ mm s}^{-1}$ ($\dot{\gamma}$ between $\pm 4 \times 10^4 \text{ s}^{-1}$) further accelerated separation (<2 s). While we have separated a single enantiomer based on the sign of the shear, by symmetry our results also apply to separation of a racemic mixture in shear of a given sign.

To confirm that the observed lateral drift is due to the coupling of shear and chirality, we use resistive force theory [16] and model a spirochete as a 25-turn helix with dimensions shown in Fig. 2(a). The drag force per unit length is $\mathbf{f} = \zeta_{\parallel} \mathbf{v}_{\parallel}^r + \zeta_{\perp} \mathbf{v}_{\perp}^r$, where $\zeta_{\parallel}, \zeta_{\perp}$ and $\mathbf{v}_{\parallel}^r, \mathbf{v}_{\perp}^r$ are the resistive coefficients and relative velocities parallel and perpendicular to a given segment of the helix, respectively. The drag anisotropy $\zeta_{\perp}/\zeta_{\parallel}$ depends on the thickness of the spirochete body and varies between 1.4 and 1.7 [16]. In Stokes flow, the linear and angular velocities of the helix are determined by total force and total moment balance. This yields the drift velocity v of a spirochete for an arbitrary orientation relative to the flow. Since we observe that the spirochetes lie in the x-y plane to within 10° , as expected on theoretical grounds [21], we adopt a twodimensional model with orientations limited to the x-y plane. The mean drift velocity then depends on the probability distribution $c(\theta)$ of orientations of the helix relative to the flow. This distribution obeys $\partial_t c = D_R \partial_\theta^2 c - \partial_\theta(\dot{\theta}c)$, where $D_R = 2.3 \times 10^{-3} \text{ s}^{-1}$ is the rotational diffusivity of an $L = 16 \mu \text{m}$ long spirochete [21] and $\dot{\theta}$ is the rotation rate caused by the shear flow. We approximate $\dot{\theta}$ as

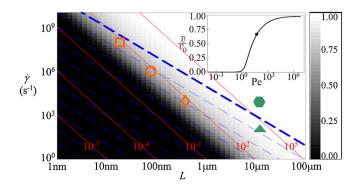


FIG. 4 (color online). Size limit for separation of isometric helices of length L and equivalent aspect ratio r=70. Dashed contours show constant Re (spaced by a factor of 10) and the thick contour is Re=0.1. Solid contours show constant Pe. The gray scale shows \bar{v}/v_0 , where v_0 is the drift velocity of a helix aligned with the flow. The smallest helices which can be separated for a given shear rate, determined by $\bar{v}/v_0=0.66$ (Pe = 50, marked by circle on inset) and Re<0.1, have $L\approx20$ (square), 80 (circle), and 400 nm (diamond) for $\dot{\gamma}\approx10^8$, 10^6 , and 10^4 s⁻¹, respectively. Full symbols mark the parameter regimes of our experiments: Re=0.03 (triangle) and Re=5 (hexagon). Inset: \bar{v}/v_0 vs Pe.

the rotation rate of an ellipsoid with aspect ratio r, $\dot{\theta} = -\dot{\gamma}(\cos^2\theta + r^2\sin^2\theta)/(1+r^2)$ [15], with r=70, as computed from the ratio of the rotation rates of the spirochetes at $\theta=0$ and $\theta=\pi/2$ obtained from resistive force theory. The steady-state solution ($\partial_t c=0$) for $c(\theta)$ was obtained numerically and the mean drift velocity computed as $\bar{v}=\int_0^{2\pi}v(\theta)c(\theta)d\theta$. This resulted in $\bar{v}=0.81$ –1.22 μ m s⁻¹, which is of the same order of the measured value (0.41 μ m s⁻¹). The residual discrepancy might be associated with the flexibility of the spirochetes and irregularities in their geometry.

For chirality-induced separation to take place, preferential alignment of helices with the flow is required. In our experiments, spirochetes were aligned hydrodynamically via Jeffery orbits. For smaller particles, rotational Brownian motion is more effective at randomizing orientation. To determine the size of the smallest helical particles that can be separated in this manner, we find the steady distribution for helices with r = 70 and length L. The mean drift velocity \bar{v} is uniquely determined by the Peclet number $Pe = \dot{\gamma}/D_R$ (Fig. 4, inset). For $Pe \gg 1$, alignment by shear dwarfs rotational diffusion and $\bar{v} \approx$ v_0 , where v_0 is the drift velocity of a helix aligned with the flow. Our experiments are in this limit, with Pe $\approx 4 \times 10^4$ $(\dot{\gamma} \approx 103 \text{ s}^{-1} \text{ at } y = H/4)$. In contrast, for Pe < 10, diffusion destroys the alignment and $\bar{v} \rightarrow 0$. Assuming isometrically scaled helices of length L, D_R scales as L^{-3} , decreasing Pe for smaller helices. This can be partially counteracted by increasing $\dot{\gamma}$, but the particle Reynolds number Re = $\dot{\gamma}L^2/\nu$ [11] must remain small (Re \ll 1) to operate in the Stokes regime. Hence, as L decreases, $\dot{\gamma}$ can be increased as L^{-2} . The competition between Pe and Re determines the smallest helices that can be separated for a given shear rate, which we estimate as $L \approx 20$, 80, and 400 nm for $\dot{\gamma} \approx 10^8$, 10^6 , and $10^4 \, {\rm s}^{-1}$, respectively (Fig. 4, open symbols). Shear rates as high as 10^6 s⁻¹ have been reported [14] in microchannels, and current technology allows pressure heads of 2×10^8 Pa (Microfluidics Corp., Newton, MA), such that a 10 μ m deep, 5 cm long channel could generate shear rates of 10⁷ s⁻¹, sufficient to separate 40 nm particles.

While we have demonstrated the lateral drift of chiral objects, one is ultimately interested in separating a mixture of enantiomers. Because the top and bottom halves of the channel produce drift in opposite directions, a racemic mixture would separate into four quadrants, with opposite quadrants containing the same enantiomer. We are currently implementing three-dimensional microfabrication to separately collect particles from the four quadrants. While we believe our technique represents a significant step forward in enabling reliable chiral separation by shear, it remains to be seen whether it can extend to the molecular scale (<10 nm). To this end, in addition to increasing shear rates via improved manufacturing techniques, it will be

interesting to explore the feasibility of using alternative alignment mechanisms to counteract rotational diffusion. One could operate at higher Reynolds number: preliminary experiments show successful separation at Re=5, but further research is required to elucidate the role of inertia on separation and any constraints on microchannel operation, such as Dean vortices and performance under high pressure. Finally, not all chiral particles are helical, so it would be useful to understand what geometries lead to the best separation.

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