

# Shape dependent phoretic propulsion of slender active particles<sup>a)</sup>

Y. Ibrahim,<sup>1</sup> R. Golestanian,<sup>1, b)</sup> and T. B. Liverpool<sup>1, 1, c)</sup>

(Dated: 30 May 2020)

We theoretically study the self-propulsion of a thin (slender) colloid driven by asymmetric chemical reactions on its surface at vanishing Reynolds number. Using the method of matched asymptotic expansions, we obtain the colloid self-propulsion velocity as a function of its shape and surface physico-chemical properties. The mechanics of self-phoresis for rod-like swimmers has a richer spectrum of behaviours than spherical swimmers due to the presence of two small length scales, the slenderness of the rod and the width of the slip layer. This leads to subtleties in taking the limit of vanishing slenderness. As a result, even for very thin rods, the distribution of curvature along the surface of the swimmer, namely its shape, plays a surprising role in determining the efficiency of propulsion. We find that thin cylindrical self-phoretic swimmers with *blunt* ends move faster than thin prolate spheroid shaped swimmers with the same aspect ratio.

PACS numbers: pacs

Keywords: keywords

## I. INTRODUCTION

In recent years there has been an explosion of interest in the area of soft active materials<sup>1–3</sup>. Active materials are condensed matter systems driven out of equilibrium by components that autonomously convert stored energy into motion. These materials are of interest partly because of their potential as a basis for engineered smart materials. They may in addition serve as model systems for understanding fundamental non-equilibrium phenomena in cellular biology. While one the major foci of the field involves the emergence of collective behaviour from active components<sup>1–3</sup>, it is of particular importance to first understand and control the behaviour of the individual components that drive the system out of equilibrium<sup>4–9</sup>. A commonly studied active component is a synthetic self-phoretic swimmer<sup>4–9</sup>. These are colloidal particles with an asymmetric catalytic coating over their surface suspended in a viscous fluid in which are dissolved chemically reactive species (fuel) whose consumption (or production) is accelerated by the catalyst<sup>10</sup>. The asymmetric catalytic coating leads to asymmetric adsorption and release of the reactants and products. This asymmetry of the distribution of reactants and/or products coupled with a short-range interaction of the chemical species with the colloid surface, leads to an interfacial flow and hence to motion of the colloid<sup>11,12</sup>. Experimentally, the two most studied self-phoretic swimmer shapes are asymmetrically coated spheres and rods. Among the earliest studied synthetic self-phoretic swimmers were cylindrical bi-metallic nano-rods of micron length and nm-scale diameter (with two-halves made of different metal catalysts)<sup>13,14</sup> which speed up the decomposition of hydrogen peroxide ( $\text{H}_2\text{O}_2$ ), in a  $\text{H}_2\text{O}_2$  solution. Other well studied system are spherical colloids partially coated with catalysts<sup>15,16</sup> also in hydrogen peroxide solutions. Unravelling the collective behaviour of these systems<sup>17,18</sup> in order to design and control their macroscopic behaviour<sup>16,19</sup> requires a detailed understanding of the mechanisms of swimmer propulsion at the single particle level. Given this there have been several theoretical studies aimed at isolating the physical mechanisms behind the observed propulsion and linking them to properties of the swimmers such as their shape and coating<sup>7,10,20–25</sup>. These are all based on generalising results for phoretic motion in external gradients<sup>11</sup> to the situation where the gradient is self-generated.

In this article we focus on calculating the propulsion speed of self-phoretic rods as a function of their slenderness ratio (defined as the ratio of the cross-sectional diameter to the rod length),  $\epsilon$ . For thin rods with vanishing aspect ratio,  $\epsilon$  there is some debate about the particular form of the dependence on the self-propulsion speed on the length and aspect ratio<sup>10,23,25</sup>. Here we show this is because there are two small length scales in the problem, (1) the diameter of the rod and (2) the thickness of the phoretic slip layer. By a careful asymptotic analysis, we identify some subtleties in taking the limits of both these length scales to zero which clarifies the debate. We point out that physically, there is a clear hierarchy of length scales which implies that the thickness of the slip layer (a molecular scale) is much smaller than rod diameter (a colloidal scale) which is much smaller than the rod length. When we take this hierarchy into account, we find that details of the shape of the rods becomes of paramount importance in determining the asymptotic propulsion speed (in the limit  $\epsilon \rightarrow 0$ ). It has been widely assumed that thin cylindrical self-phoretic rods studied experimentally can be approximated by an equivalent thin spheroid (ellipsoid of revolution) of the same slenderness ratio<sup>5,23,25</sup>. Here we examine this assumption by comparing slender active particles of fixed slenderness  $\epsilon$  whose shape are given by thin spheroids with those whose shape is given by cylindrical rods and obtain

asymptotic results in the limit  $\epsilon \rightarrow 0$ . One conclusion of our analysis is that in fact, for issues of self-propulsion, this assumption is in general, not a justified one.

To be concrete, we do this by using the method of matched asymptotic expansions for swimmers with a generic shape function that interpolates between needle-like spheroid and cylindrical rod-like geometries. We find that rod-like shapes are faster than the needle-like shapes with the same  $\epsilon$ . Our results thus suggest a that a strategy for better performance (faster speed) of self-phoretic swimmers would be to design blunt rod-like shapes rather than tapered needle-like shapes.

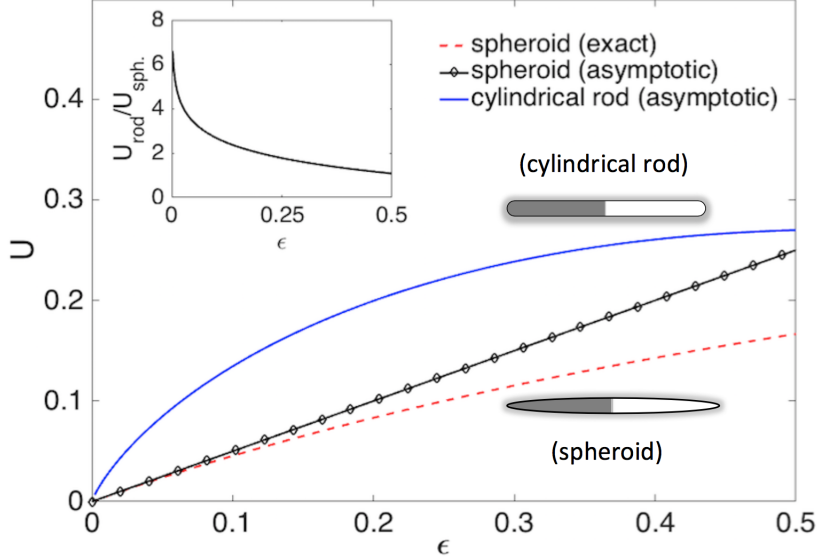


FIG. 1. Comparison of swimmer propulsion speeds. The propulsion speed,  $U$  of a half-coated rod as a function of the slenderness ratio,  $\epsilon$ . The speed,  $U$ , is expressed in units of  $|\mu^* \alpha^* / D|$ . The blue solid line is the asymptotic result ( $\epsilon \rightarrow 0$ ) for the cylindrical rod (see eqn. (44)), while the black diamonds indicate the asymptotic result for the spheroid (see eqn. (39) and ref.<sup>23</sup>). The dashed red line is the plot of the exact propulsion speed obtained in ref.<sup>25</sup> of a half-coated spheroid. (inset) Plot of the ratio of the asymptotic speeds of the cylindrical rod to the spheroid asymptotic against the slenderness ratio (see eqns. (39) and (44)).

## II. THE THIN DIFFUSIOPHORETIC SWIMMER

We consider a thin (slender) colloid which has a heterogeneous catalytic surface that produces (or consumes) molecules of a chemical species in a variable manner on its surface generating a local concentration gradient of the species in its vicinity. We then use the standard framework<sup>10,11</sup> for studying phoretic propulsion to calculate the steady-state speed of the colloid generated by this self-generated gradient. This is done by solving the coupled equations of motion of the local concentration of reactants/products and local momentum conservation taking account of the boundary conditions on the colloid surface.

### A. Equations of motion

$$\tilde{\nabla} \cdot \tilde{\mathbf{J}} = 0; \quad \tilde{\mathbf{J}} = -D \left( \tilde{\nabla} \tilde{c} + \frac{\tilde{c}}{k_B T} \tilde{\nabla} \tilde{\psi} \right), \quad (1)$$

in the zero Péclet number limit, where  $\tilde{\psi}(\tilde{\mathbf{r}} - aS(\tilde{\mathbf{z}}), \tilde{\mathbf{z}})$  is the short-range interaction energy of the reactants (products) with the colloid surface.  $S(\tilde{\mathbf{z}})$  is the colloid shape function, the surface of the colloid is given by the function  $\tilde{\mathbf{r}}(\tilde{\mathbf{z}}, \theta) = (aS(\tilde{\mathbf{z}}) \cos \theta, aS(\tilde{\mathbf{z}}) \sin \theta, \tilde{\mathbf{z}})$ . In the experimental systems motivating this work, the typical length of the self-phoretic colloids is  $2l \sim 2\mu\text{m}$  and they move with a speed  $U^* \sim 10\mu\text{ms}^{-1}$  while the diffusion coefficient of the

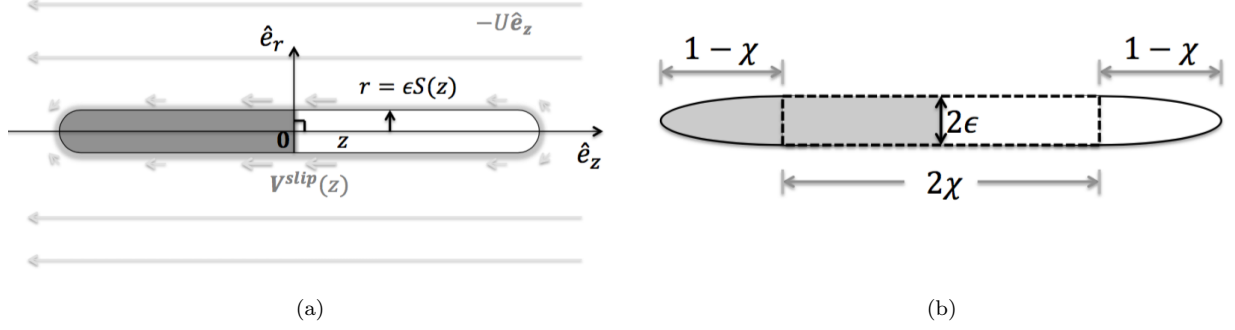


FIG. 2. (a) Sketch of a slender catalytic microswimmer showing the variation of catalytic coverage and shape along the long axis oriented in the  $z$  direction. The gray (shaded) half of the swimmer surface is made of a different catalyst than the unshaded half. The catalysts produce (or consume) molecules of the reactive species at different rates leading to an asymmetric distribution of molecules near the swimmer.  $\mathbf{V}^{\text{slip}}$  is the phoretic slip velocity induced by the coupling of the asymmetric distribution of the molecules and their short-range interaction with the swimmer surface.  $U\hat{e}_z$  is the axial translational velocity of the swimmer in the lab frame. (b) The generic thin rod with rounded ends which interpolates between a thin spheroid and a thin cylinder. The limit ( $\chi = 0$ ) is the spheroid, while ( $\chi = 1$ ) is the cylindrical rod.

reactants (products)  $D \sim 10^3 \mu\text{m}^2\text{s}^{-1}$ . Therefore, they are in the small Péclet number limit ( $\mathcal{P}e = U^*l/D \ll 1$ ).  $k_B$  is the Boltzmann constant and  $T$  is the temperature of the solution. The heterogeneous surface leads to a variable production/consumption rate,  $\tilde{\alpha}(\tilde{z})$  of molecules on the surface giving a flux boundary condition on the surface:

$$\hat{\mathbf{n}} \cdot \tilde{\mathbf{J}} = \tilde{\alpha}(z), \quad \text{at} \quad \tilde{r} = a S(\tilde{z}), \quad (2)$$

and the concentration attenuation decays far from the swimmer,  $\tilde{c} \rightarrow \tilde{c}_\infty$ , as  $\sqrt{\tilde{r}^2 + \tilde{z}^2} \rightarrow \infty$ .

We restrict our analysis to the fixed flux limit where the catalytic production/consumption rate,  $\alpha(z)$ , does not depend on the local concentration of the reactant (product) particles. In this regime, the particles flux is limited by the reaction rate rather than diffusion of the particles to the swimmer surface.

$$0 = \tilde{\nabla} \cdot \tilde{\mathbf{v}}, \quad (3)$$

$$\mathbf{0} = \eta \tilde{\nabla}^2 \tilde{\mathbf{v}} - \tilde{\nabla} \tilde{p} - \tilde{c} \tilde{\nabla} \tilde{\psi}, \quad (4)$$

where  $\eta$  is the fluid viscosity. This is justified as typical experimental swimming speeds of  $U^* \sim 10 \mu\text{ms}^{-1}$  and  $2l \sim 2 \mu\text{m}$  colloid length, and for typical solution kinematic viscosity  $\nu \sim 10^{-6} \text{m}^2\text{s}^{-1}$ , mean these colloids generate flows with very low Reynolds number ( $\mathcal{R}e = U^*l/\nu \ll 1$ ). The inhomogeneous term  $-\tilde{c} \tilde{\nabla} \tilde{\psi}$  arises due to the interactions of the reactant (product) particles with the colloid surface,  $\tilde{\psi}(r - aS(z), z)$ . We have no-slip boundary conditions on the colloid surface in otherwise uniform flow

$$\tilde{\mathbf{v}} = \mathbf{0}, \quad \text{at} \quad \tilde{r} = a S(\tilde{z}), \quad (5)$$

$$\tilde{\mathbf{v}} \rightarrow -\tilde{U} \hat{e}_z, \quad \sqrt{\tilde{r}^2 + \tilde{z}^2} \rightarrow \infty. \quad (6)$$

Note that we are concerned only with the axial propulsion of the swimmer (restricting the swimmer translation along  $\tilde{z}$ -axis). In addition, the colloid experiences no net body force (which uniquely determines  $\tilde{U}$ )

$$\oint_{\tilde{r}=aS(\tilde{z})} \tilde{\Pi} \cdot \hat{\mathbf{n}} d\tilde{S} - \iiint_{-\infty}^{\infty} \tilde{c} \tilde{\nabla} \tilde{\psi} d\tilde{V} = \mathbf{0} \quad (7)$$

where  $\tilde{\mathbf{\Pi}} = -\tilde{p}\mathbf{1} + \eta \left( \tilde{\nabla} \tilde{\mathbf{v}} + [\tilde{\nabla} \tilde{\mathbf{v}}]^T \right)$  is the fluid stress tensor and  $d\tilde{S} = a d\phi dz$ ,  $d\tilde{V} = r dr d\phi dz$  are the surface and volume elements in cylindrical coordinates respectively.

## B. Non-dimensionalization

We express the displacements  $(r, z)$  in terms of half the length of the swimmer,  $l$ . We consider swimmers with concentration gradients generated by the colloids on a scale  $c^*/a$ . Hence we express the flux  $\tilde{J}$  of reactant (product) particles in units of  $c^*D/a$ , the concentration field  $\tilde{c}$  in units of  $c^*$ , (where  $D$  is the diffusion coefficient of the reactant/product particles), the interaction energy  $\tilde{\Psi}$  in the units of thermal energy  $k_B T$ , ( $k_B$  the Boltzmann constant and  $T$  is the temperature of the solution), the flow field  $\tilde{\mathbf{v}}$  with  $U^* = \mu^* c^*/a$  (where  $\mu^* = k_B T L^{*2}/\eta$  is the characteristic phoretic mobility coefficient,  $\eta$  is the viscosity of the fluid and  $L^*$  is the short-range interaction lengthscale of the reactant (product) molecules with the swimmer surface). We scale the pressure field  $\tilde{p}$  with  $\eta U^*/l$ . We note that the problem has three length-scales; the swimmer characteristic length  $2l$ , its cross-sectional radius  $a$  and the range  $L^*$  of the interaction between the molecules and swimmer surface. Consequently, we have three asymptotic near-field regions and in addition the scale on which the ends of the rod are rounded.

We therefore define the dimensionless parameters,  $\epsilon = a/l$ , the slenderness ratio and  $\lambda = L^*/l$ , the interaction layer thickness to the swimmer largest lengthscale, with  $0 < \lambda \ll \epsilon \ll 1$ . In addition, we define dimensionless fields  $\mathbf{J} = \tilde{J}a/c^*D$ ,  $c = \tilde{c}/c^*$ ,  $\mathbf{v} = \tilde{\mathbf{v}}/U^*$ ,  $p = \tilde{p}/\eta U^*$ ,  $\psi = \tilde{\psi}/k_B T$ , and the dimensionless catalytic flux  $\alpha = \tilde{\alpha}a/c^*D$  on the swimmer surface.

$$\nabla \cdot \mathbf{J} = 0; \quad \mathbf{J} = -\epsilon (\nabla c + c \nabla \psi), \quad (8)$$

$$0 = \nabla \cdot \mathbf{v}, \quad (9)$$

$$\mathbf{0} = \nabla^2 \mathbf{v} - \nabla p - \epsilon \lambda^{-2} c \nabla \psi, \quad (10)$$

where  $\lambda = L^*/l$  is the ratio of the interaction length-scale to half the length of the swimmer,  $\epsilon = a/l$  is the swimmer slenderness ratio and  $\psi(r - \epsilon S(z), z)$  is the short-range interaction potential between reactant (product) molecules and surface.

$$\hat{\mathbf{n}} \cdot \mathbf{J} = \alpha(z), \quad \text{at } r = \epsilon S(z), \quad (11)$$

and the concentration decays to its value far from the swimmer,  $c \rightarrow c_\infty$ ,  $\sqrt{r^2 + z^2} \rightarrow \infty$ .

$$\mathbf{v} = \mathbf{0}, \quad \text{at } r = \epsilon S(z), \quad (12)$$

$$\mathbf{v} \rightarrow -U \hat{\mathbf{e}}_z, \quad \sqrt{r^2 + z^2} \rightarrow \infty. \quad (13)$$

The zero torque and force conditions are

$$\oint_{r=\epsilon S(z)} \mathbf{\Pi} \cdot \hat{\mathbf{n}} dS - \epsilon \lambda^{-2} \int_{-\infty}^{\infty} c \nabla \psi dV = \mathbf{0} \quad (14)$$

where  $\mathbf{\Pi} = -p\mathbf{1} + (\nabla \mathbf{v} + [\nabla \mathbf{v}]^T)$  is the dimensionless stress tensor and  $dS$ ,  $dV$  are the surface and volume elements in cylindrical coordinates respectively.

## C. The slender shape function

$$S(z; \chi) = \begin{cases} 1, & 0 \leq |z| \leq \chi; \\ \sqrt{1 - \left(\frac{|z| - \chi}{1 - \chi}\right)^2}, & \chi \leq |z| \leq 1, \end{cases} \quad (15)$$

where the parameter  $0 \leq |\chi| < 1$  is the additional lengthscale specifying the swimmer shape curvature. The limit  $\chi \rightarrow 0$  is the widely studied spheroid shape, while the limit  $\chi \rightarrow 1$  is the cylindrical rod shape.  $1 - \chi = \epsilon$  is a cylindrical rod with hemispherical caps.

$$\hat{\mathbf{t}} = \frac{\epsilon S_z(z) \hat{\mathbf{e}}_r + \hat{\mathbf{e}}_z}{\sqrt{1 + \epsilon^2 S_z^2(z)}}, \quad (16)$$

$$\hat{\mathbf{n}} = \frac{\hat{\mathbf{e}}_r - \epsilon S_z(z) \hat{\mathbf{e}}_z}{\sqrt{1 + \epsilon^2 S_z^2(z)}}, \quad (17)$$

where  $S_z(z) = dS(z)/dz$ .

### III. ANALYSIS

#### A. Overview

We use the method of matched asymptotics together with slender-body approximations to solve for the swimmer propulsion speed  $U$ . We shall first identify the different regions of the solution of the equations of motion, and define re-scaled coordinates for both the interaction layer  $r \sim \mathcal{O}(\lambda)$  and the slender body inner region  $r \sim \mathcal{O}(\epsilon)$  (see Figure 3). In the subsequent sections, we systematically find solutions to the diffusion and Stokes equations in these regions. Then we match these asymptotic results to obtain a consistent solution for the whole domain and hence find the propulsion speed  $U$  which is the goal of the analysis.

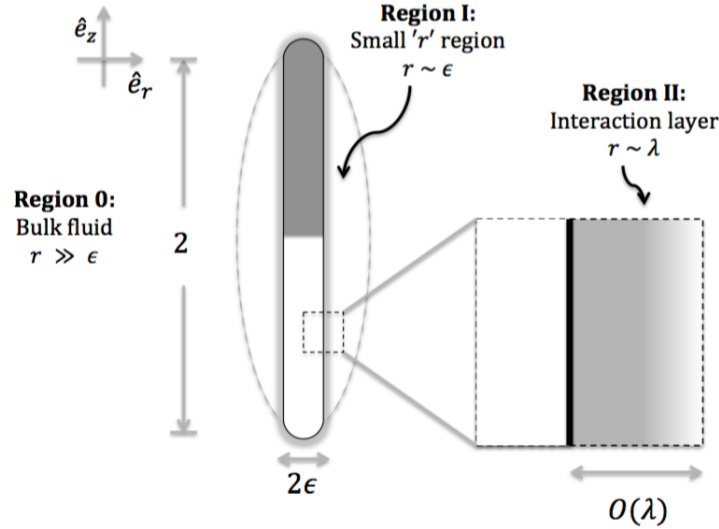


FIG. 3. Sketch of the slender swimmer showing the asymptotic regions.  $\lambda = L^*/l$  and  $\epsilon = a/l$  are dimensionless small parameters; where  $2l$  is the swimmer length,  $2a$  its diameter and  $L^*$  the range of the interaction between reactant (product) molecules and the swimmer surface (typically of the order of molecular size  $\sim \text{\AA}$ ).

We have three regions of solution for our matched asymptotics. In the outer layer, region 0,  $r \gg \epsilon, \lambda$ ; in the intermediate layer, region I,  $r \sim \epsilon$ ; and in the inner interaction layer, region II,  $r \sim \lambda$ . In regions 0 and I, the

hierarchy of scales  $\epsilon \gg \lambda$  mean that we can approximate the interaction layer as being of vanishing width: we can set  $\lambda$  to zero, and our analysis proceeds via an asymptotic expansion purely in  $\epsilon$ . However in region II, we must take account of both  $\lambda, \epsilon$  and need to perform a *double expansion* in both variables which we use for our asymptotic analysis. The details of our calculation are outlined in the appendix and we present only a summary of the main results here.

$$U + (\hat{e}_z \cdot \hat{\mathbf{t}}(z)) v_z(0, z) = 4q(z; \epsilon) \log\left(\frac{1}{\epsilon}\right) + 4q(z; \epsilon) \log\left(\frac{2\sqrt{1-z^2}}{S(z)}\right) - 2q(z; \epsilon) + 2 \int_{-1}^1 \frac{q(\xi; \epsilon) - q(z; \epsilon)}{|z - \xi|} d\xi, \quad (18)$$

where the slip velocity,  $\mathbf{v}(0, z) = \mathbf{V}^{\text{slip}}$  and force density along the swimmer centerline,  $q(z, \epsilon)$  are obtained by matching with the inner and intermediate regions (see Appendix ).

The propulsion speed  $U$  is determined by requiring that the solution of the integral equation above satisfies the constraint of zero total force on the swimmer, i.e requiring

$$\int_{-1}^1 q(z) dz = 0. \quad (19)$$

From region I and II, we obtain the slip velocity<sup>11</sup> as (see Appendices and )

$$\mathbf{V}^{\text{slip}}(z; \epsilon) = \epsilon \mu(z) \hat{\mathbf{t}} \cdot \nabla c(z; \epsilon) \hat{\mathbf{t}}, \quad (20)$$

where we have define the diffusiophoretic mobility<sup>11</sup>  $\mu(z) = \int_0^\infty \rho \left[1 - e^{-\Psi^{(0)}(\rho, z)}\right] d\rho$ , and the concentration gradient,  $\nabla c$  is obtained from region 0 and I.

The concentration,  $c(z; \epsilon)$  in region 0 and I can be calculated as (see Appendices and )

$$c(z; \epsilon) = C_I^{(-1)}(z) \log\left(\frac{1}{\epsilon}\right) + C_I^{(0)}(z) \quad (21)$$

where

$$C_I^{(-1)}(z) = \alpha(z) S(z), \quad (22)$$

$$C_I^{(0)}(z) = C_\infty + \alpha(z) S(z) \log\left(\frac{2}{S(z)}\right) - \frac{1}{2} \int_{-1}^1 \frac{d}{d\xi} \left[ \alpha(\xi) S(\xi) \right] \text{sgn}(z - \xi) \log|z - \xi| d\xi. \quad (23)$$

So we can calculate the propulsion speed of the swimmer by solving the coupled integral equations (18,19) for the unknowns  $q(z), U$ . The solutions will depend on the shape functions  $S(z)$  due to the explicit dependence on  $S(z)$  of equation (18) and the fact that  $\hat{\mathbf{t}}(z) = \hat{\mathbf{t}}(S(z))$ . In the next section, we will consider slender swimmers with both needle-like (or spheroid) and rod-like shapes and highlight the different asymptotic limits of their propulsion velocities.

## B. The swimmer propulsion velocity

In this section, we obtain an approximate value for the propulsion speed  $U$  of the swimmer and force density,  $q(z; \epsilon)$  by solving the coupled integral equations (18,19). There are various methods for solving this type of problem. For example, an approximate solution by a Legendre expansion of the force density<sup>30</sup>  $q(z; \epsilon)$  in the interval  $-1 \leq z \leq 1$  to obtain an infinite linear system of equations from which one obtains  $U$ . Other alternatives are perturbative methods<sup>28</sup> for solving integral equations where the force densities (and hence the propulsion speed  $U$ ) are found iteratively. However, since we are interested here in the slender limit, following Cox<sup>27</sup> and Tillett<sup>31</sup>, the presence of  $\log(\epsilon)$  in equation (18) suggests an asymptotic series expansion of  $q(z)$  and  $U$  with terms of type  $\epsilon^m (\log(\epsilon))^n$ , with  $n, m$  integers. Since the slip velocity vanishes as  $\epsilon \rightarrow 0$ , in the limit  $\epsilon \ll 1$ , the asymptotically leading terms in such

a series will be of the form  $\epsilon(\log \epsilon)^n$  where  $n \leq 1$ , which suggests an expansion<sup>23,27,31</sup> of the force density and speed in powers of  $(\log \epsilon)^{-1}$ .

$$V_z^{\text{slip}}(z; \epsilon) = V_{-1}^{\text{slip}}(z) \epsilon \log \left( \frac{1}{\epsilon} \right) (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) + V_0^{\text{slip}}(z) \epsilon (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) , \quad (24)$$

where

$$V_{-1}^{\text{slip}}(z) = \mu(z) \hat{\mathbf{t}} \cdot \nabla C_I^{(-1)}(z) , \quad V_0^{\text{slip}}(z) = \mu(z) \hat{\mathbf{t}} \cdot \nabla C_I^{(0)}(z) .$$

with  $C_I^{(-1)}, C_I^{(0)}$  defined in equations (22,23). The propulsion speed  $U$  and the force density  $q(z; \epsilon)$  are written as expansions in  $(\log \epsilon)^{-1}$

$$\frac{U}{\epsilon} = -U_{-1} \log(\epsilon) + U_0 - U_1 (\log(\epsilon))^{-1} + \dots , \quad (25)$$

$$\frac{q(z; \epsilon)}{\epsilon} = q_0(z) - q_1(z) (\log(\epsilon))^{-1} - q_2(z) (\log(\epsilon))^{-2} + \dots , \quad (26)$$

where the force-free condition and the validity of the expansion for all values of  $\epsilon$ , implies,  $\int_{-1}^1 q_n(z) dz = 0$ , for all  $n$ . Substituting equations (25,26) and equation (24) into equation (18) gives

$$U_{-1} + V_{-1}^{\text{slip}}(z) (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) = 4q_0(z) , \quad (27)$$

$$U_0 + V_0^{\text{slip}}(z) (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) = 4q_0(z; \epsilon) \log \left( \frac{2\sqrt{1-z^2}}{S(z)} \right) - 2q_0(z; \epsilon) + 2 \int_{-1}^1 \frac{q_0(\xi; \epsilon) - q_0(z; \epsilon)}{|z - \xi|} d\xi , \quad (28)$$

$$U_n = 4q_n(z; \epsilon) \log \left( \frac{2\sqrt{1-z^2}}{S(z)} \right) - 2q_n(z; \epsilon) + 2 \int_{-1}^1 \frac{q_n(\xi; \epsilon) - q_n(z; \epsilon)}{|z - \xi|} d\xi . \quad (29)$$

$$U_{-1} = -\frac{1}{2} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}(z)) V_{-1}^{\text{slip}}(z) dz , \quad (30)$$

and the next term (from

$$U_0 = -\frac{1}{2} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}(z)) V_0^{\text{slip}}(z) dz - 2 \int_{-1}^1 q_0(z) \log \left( \frac{2\sqrt{1-z^2}}{S(z)} \right) dz , \quad (31)$$

where

$$q_0(z) = \frac{1}{4} (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) V_{-1}^{\text{slip}}(z) - \frac{1}{8} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) V_{-1}^{\text{slip}}(z) dz . \quad (32)$$

Using equation (29) all the higher order contributions to  $U$  can in principle be evaluated. However, if one intends to go beyond asymptotic results iterative methods might converge faster than the expansion outlined above<sup>28</sup>. We note that for a spheroidal rod where  $S(z) = \sqrt{1-z^2}$ , we can explicitly evaluate the integral  $\int_{-1}^1 q_n(z) dz = 0$  in equation (29) and one finds that all  $U_n = 0, n \geq 1$ .

$$\alpha(z) = \frac{1}{2} (\alpha_+ + \alpha_-) + \frac{1}{2} (\alpha_+ - \alpha_-) \tanh \left( \frac{z}{\Delta} \right) , \quad (\Delta \rightarrow 0) . \quad (33)$$

Given the shape functions defined above (see equation (15)) and defining  $\delta = 1 - \chi$  (see Figure 4(a), 4(b), 4(c)), the tangent unit vector to the swimmer surface is

$$\hat{\mathbf{t}}(z) = \begin{cases} \hat{\mathbf{e}}_z, & z \in (-\chi, \chi); \\ \frac{-\epsilon \operatorname{sgn}(z)(|z| - \chi) \hat{\mathbf{e}}_r + \sqrt{\delta^4 - \delta^2 (|z| - \chi)^2} \hat{\mathbf{e}}_z}{\sqrt{\delta^4 + (\epsilon^2 - \delta^2)(|z| - \chi)^2}}, & z \in [-1, -\chi) \cup (\chi, 1], \end{cases} \quad (34)$$

where  $\operatorname{sgn}(x)$  is the sign function. By varying  $\chi$  between 0 and 1 we can interpolate between a tapered rod and a blunt rod.

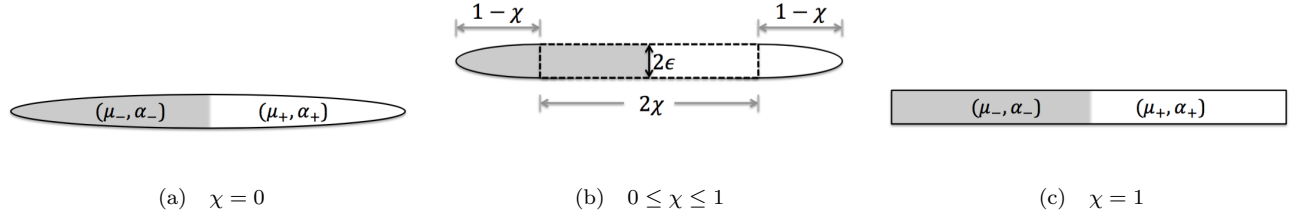


FIG. 4. Examples of slender self-phoretic swimmer shapes with different physico-chemical properties on their surfaces.

### 1. Needle-like shape (Tapered ends)

$$S(z) = S(z; \chi = 0) = \sqrt{1 - z^2}; \quad z \in [-1, 1]. \quad (35)$$

for which an exact solution exists<sup>21,25</sup>. For this shape, the unit tangent vector in equation (34) simplifies to

$$\hat{\mathbf{t}} = \frac{\sqrt{1 - z^2} \hat{\mathbf{e}}_z - \epsilon z \hat{\mathbf{e}}_r}{\sqrt{1 - z^2 + \epsilon^2 z^2}}; \quad z \in [-1, 1]. \quad (36)$$

For a half-coated ( $\alpha_+ = -1, \alpha_- = 0$ ) spheroid swimmer with uniform mobility ( $\mu(z) = \mu_+ = \mu_- = 1$ ), we obtain

$$U_{-1} = -\frac{1}{2} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) V_{-1}^{\text{slip}}(z) dz = -\frac{1}{2} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) \hat{\mathbf{t}} \cdot \nabla C_I^{(-1)}(z) dz = o(\epsilon), \quad (37)$$

$$U_0 = -\frac{1}{2} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) V_0^{\text{slip}}(z) dz = -\frac{1}{2} \int_{-1}^1 (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}) \hat{\mathbf{t}} \cdot \nabla C_I^{(0)}(z) dz = \frac{1}{2} + o(\epsilon), \quad (38)$$

where  $C_I^{(-1)}$  and  $C_I^{(0)}$  are defined in equations (A52,A53). From eqn. (29) all higher order terms vanish  $U_n = 0$  ( $n \geq 1$ ).

Therefore, the speed is asymptotically,

$$U = \frac{1}{2} \epsilon + o(\epsilon^2 \log(\epsilon)), \quad (39)$$



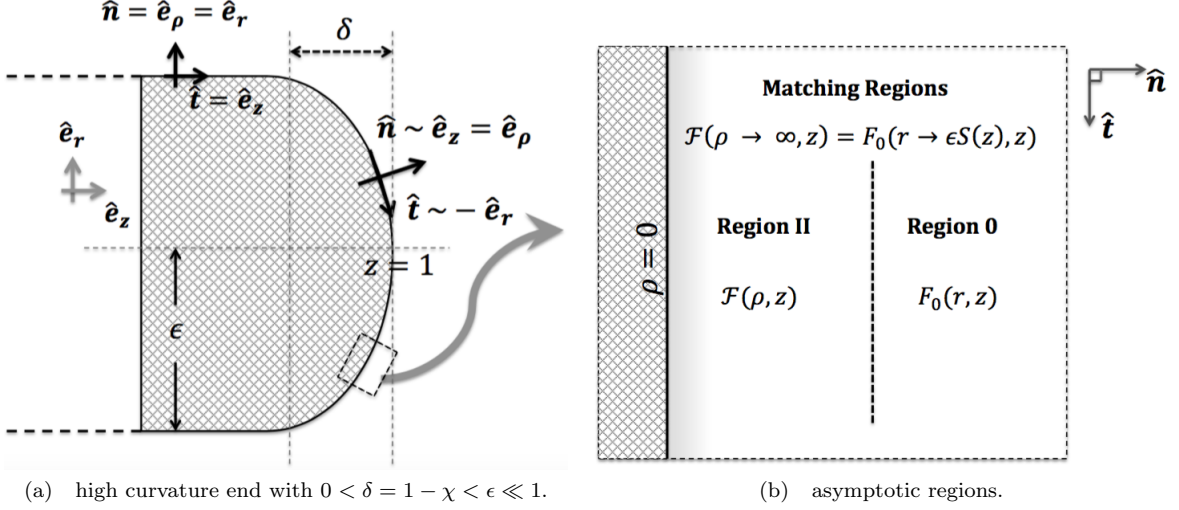


FIG. 5. Sketch of the swimmer *blunt* end - showing the matching regions.  $\delta = 1 - \chi$  is the width of the near end region where the shape function decays to zero. Note that the intermediate (region I) vanishes near the ends.

which agrees with the asymptotic limit of the exact solution<sup>25</sup> and the asymptotic result<sup>23</sup>.

## 2. Rod-like shape (Blunt ends)

We next consider a straight rod-like swimmer with blunt ends (see Figure 5). We define body shapes with *blunt* ends as shapes having a small but finite end width  $\delta = 1 - \chi$ , such that  $0 < \delta < \epsilon \ll 1$ , where the body shape drops rapidly but continuously to zero (see Fig. 5). Thus, the projection of the tangent vector along the axis is

$$\hat{e}_z \cdot \hat{t} = \begin{cases} 1, & z \in (-\chi, \chi) ; \\ o(\epsilon), & z \in [-1, -\chi) \cup (\chi, 1] , \end{cases} \quad (40)$$

since  $\text{maximum}\{\delta/\epsilon\} = 1$ . Hence, to the leading order, integrals involving the projection  $\hat{e}_z \cdot \hat{t}$  are evaluated as

$$\int_{-1}^1 (\hat{e}_z \cdot \hat{t}) \left( \cdot \right) dz = \int_{-\chi}^{\chi} \left( \cdot \right) dz + o(\epsilon), \quad (0 < 1 - \chi < \epsilon \ll 1). \quad (41)$$

This is crucial in determining the propulsion speed since for these shapes  $S(z = \pm\chi) = 1$  while  $S(z = \pm 1) = 0$ .

$$U_{-1} = -\frac{1}{2} \int_{-1}^1 (\hat{e}_z \cdot \hat{t}) V_{-1}^{\text{slip}} dz = -\frac{1}{2} \int_{-\chi}^{\chi} \frac{dC_I^{(-1)}}{dz} dz + o(\epsilon) = \frac{1}{2} + o(\epsilon), \quad (42)$$

$$\begin{aligned} U_0 &= -\frac{1}{2} \int_{-1}^1 (\hat{e}_z \cdot \hat{t}) V_0^{\text{slip}}(z; \epsilon) dz - 2 \int_{-1}^1 q_0(z) \log \left( \frac{2\sqrt{1-z^2}}{S(z)} \right) dz \\ &= -\frac{1}{2} \int_{-\chi}^{\chi} \frac{dC_I^{(0)}}{dz} dz - 2 \int_{-1}^1 q_0(z) \log \left( \frac{2\sqrt{1-z^2}}{S(z)} \right) dz + o(\epsilon) = \left( \log 2 - \frac{1}{2} \right) + o(\epsilon). \end{aligned} \quad (43)$$

Note that the integration limits are  $z = \pm\chi$  rather than  $z = \pm 1$  since the high curvature ends do not contribute to the propulsion velocity (see equation 41). Hence, we obtain a speed

$$U = \frac{1}{2} \epsilon \left[ \log \left( \frac{4}{\epsilon} \right) - 1 \right] + \mathcal{O} \left( \frac{\epsilon}{\log(\frac{1}{\epsilon})} \right). \quad (44)$$

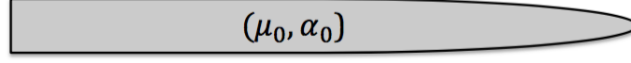


FIG. 6. Fore-aft asymmetric swimmer with uniform physico-chemical properties over its surface.

We note that with the same physico-chemical properties, this asymptotic speed is significantly faster than that of the spheroid shape (equation (39)).

### 3. Asymmetric shape (mixed ends)

$$S(z) = \begin{cases} S(z; 0 < 1 - \chi < \epsilon), & -1 \leq z \leq 0, \\ S(z; \chi = 0), & 0 \leq z \leq 1. \end{cases} \quad (45)$$

Hence we obtain the propulsion velocity at leading order in  $\epsilon$ ,

$$U = U_{-1} \epsilon \log\left(\frac{1}{\epsilon}\right) + U_0 \epsilon + \dots = \frac{1}{2} \epsilon \left[ \log\left(\frac{\sqrt{2}}{\epsilon}\right) - 1 \right] + \mathcal{O}\left(\frac{\epsilon}{\log(\frac{1}{\epsilon})}\right). \quad (46)$$

## IV. DISCUSSION

We have performed a detailed study of how the shape of slender diffusiophoretic self-propelled colloid (a phoretic swimmer) affects the speed at which the colloid 'swims'. By slender here, we mean that the aspect ratio of average diameter to length is much less than 1,  $\epsilon = a/l \ll 1$ . In our calculations, the thickness of the phoretic slip layer,  $L^*$  is taken as the smallest length in the problem since it is a molecular length-scale comparable to the interaction range of the reactant (product) molecules, while the diameter  $a$  is a much larger colloidal length-scale, so we have the hierarchy of length-scales  $l \gg a \gg L^*$ . Our approach is based on using simple axisymmetric shape functions which are able to interpolate between needle-like swimmers (spheroids) which have tapered ends and cylindrical rod-like swimmers with blunt ends. We note that cylindrical rod-like swimmers with blunt ends are easier to manufacture and hence have been the subject of many of the experimental studies to date. To obtain results in the limit  $\epsilon \rightarrow 0$ , we have used the method of matched asymptotic expansions.

Our main result is the observation that the swimmer propulsion speed is strongly shape-dependent and in particular depends on whether the swimmer has pointed or blunt ends, i.e has a needle-like or rod-like shape respectively. We show that swimmers with blunt ends (cylinders) have a higher propulsion speed than swimmers with tapered ends (needles) of the same aspect ratio,  $\epsilon$ .

A simple argument explaining the differences in swimming efficiency can be obtained by considering the competition of effects that lead to self-propulsion. Self-propulsion arises from the balance of the hydrodynamic stresses due to the slip velocity generated by the concentration gradients of reactants (products) and the associated hydrodynamic drag due to fluid velocity gradients near the surface of the swimmer. The constraint of force (torque) balance on the swimmer then selects a unique propulsion velocity. For a given shape, the higher the concentration gradient developed, the higher the propulsion velocity. The concentration gradients from the reaction occur on a length-scale which is set by the length of the swimmer and is mostly independent of the detailed shape of the swimmer, however the slip and drag they generate depend on the local surface area of the swimmer and are sensitive to the detailed shape of the swimmer. For a spheroid, the hydrodynamic stresses leading to drag are comparable to the stresses due to slip along the whole surface of the colloid, while for the blunt cylinder, the effects of phoretic slip dominate the drag at the highly curved ends<sup>25</sup>. At a fixed aspect ratio,  $\epsilon$ , the surface area of the spheroid,  $A_{sph}$  is larger than the surface area of the blunt cylinder,  $A_{cyl}$ . Hence the balance of stresses is skewed towards a higher propulsion velocity for blunt cylinders.

Our results clearly imply that care must be taken in designing theoretical models for self-phoretic slender swimmers and that slender self-phoretic cylindrical rods cannot be accurately modelled by slender self-phoretic spheroids. We note that there is a limit when such details of the swimmer shape do not matter - when the slip layer is larger than the diameter of the rods. However this limit, as discussed above, is unphysical.

## ACKNOWLEDGMENTS

This work was supported by EPSRC grant EP/G026440/1 (TBL, YI), and HFSP grant RGP0061/2013 (RG). YI acknowledges the support of University of Bristol. TBL acknowledges support of BrisSynBio, a BBSRC/EPSRC Advanced Synthetic Biology Research Centre (grant number BB/L01386X/1).

## Appendix A: Asymptotic analysis

### 1. Region 0: Outer region

$$c(r, z) = C_0(r, z) , \quad (\text{A1})$$

$$\mathbf{v}(r, z) = \mathbf{V}_0(r, z) . \quad (\text{A2})$$

Since,  $r \gg \epsilon$ , the short-range interaction potential of the reactive species with the swimmer surface vanishes in this region

$$\psi(r - \epsilon S(z), z) = \Psi_0(r - \epsilon S(z), z) = 0 . \quad (\text{A3})$$

We approximate the concentration by one generated by a line distribution of point sources<sup>23</sup>, along the swimmer centerline  $(r, z) = (0, \xi)$ , where  $-1 \leq \xi \leq 1$ .

$$C_0(r, z) = C_\infty + \int_{-1}^1 \frac{j(\xi)}{\sqrt{r^2 + (z - \xi)^2}} d\xi, \quad (\text{A4})$$

where  $j(z)$  is the source line density along the body centerline. It is to be determined by matching  $C_0(r, z)$  as  $r \rightarrow 0$  with the intermediate (region I) expansion of concentration field (see Fig. 3). We note that since  $\Psi_0(r - \epsilon S(z), z) = 0$  in this region, the concentration  $C_0(r, z)$ , is a solution of the steady state diffusion equation (8).

While the integral in equation (A4) is singular as  $r \rightarrow 0$  and  $|z| \leq 1$ , there are a number of ways to isolate the singularity and approximate the integral analytically as the observation point approaches the swimmer surface. A particularly simple method is subtracting the singular part of the integral which gives the approximate expression<sup>28</sup>

$$C_0(r, z) = C_\infty + 2j(z) \log\left(\frac{2}{r}\right) + \int_{-1}^1 \frac{j(\xi) - j(z)}{|z - \xi|} d\xi + 2j(z) \log\left(\sqrt{1 - z^2}\right) + \mathcal{O}(r^2) . \quad (\text{A5})$$

$$C_0(r, z) = C_\infty + 2j(z) \log\left(\frac{2}{r}\right) - \int_{-1}^1 \frac{d}{d\xi} j(\xi) \operatorname{sgn}(z - \xi) \log|z - \xi| d\xi + \mathcal{O}(r^2 \log(r)) . \quad (\text{A6})$$

Note that we assumed  $j(\pm 1) = 0$  in obtaining the above expression, which we confirm to be the case *a posteriori* for our generic shape function defined in equation (15).

We approximate the fluid velocity field by one generated by a line distribution of point force singularities along the swimmer centerline<sup>26</sup>

$$\mathbf{V}_0(r, z) = -U \hat{\mathbf{e}}_z + \int_{-1}^1 d\xi \left( \frac{\hat{\mathbf{e}}_z}{|\mathbf{x} - \mathbf{x}'|} + \frac{(z - \xi)(\mathbf{x} - \mathbf{x}')}{|\mathbf{x} - \mathbf{x}'|^3} \right) q(\xi; \epsilon) , \quad (\text{A7})$$

where  $\mathbf{x} - \mathbf{x}' = r \hat{\mathbf{e}}_r + (z - \xi) \hat{\mathbf{e}}_z$  and  $q(z; \epsilon)$  is a generic force line density along the body's centerline.  $q(z; \epsilon)$  will be determined by matching  $\mathbf{V}_0(r, z)$  as  $r \rightarrow 0$  with the intermediate (region I) velocity field (see Fig. 3). This velocity field is a solution of the Stokes equations (9,10) since the short-range interaction potential vanishes in this region. There is an associated hydrostatic pressure,

$$p(r, z) - p_\infty = P_0(r, z) = 2 \int_{-1}^1 d\xi \left( \frac{z - \xi}{|\mathbf{x} - \mathbf{x}'|^3} \right) q(\xi; \epsilon)$$

$$V_{0,z}(r, z) = -U + 4q(z; \epsilon) \log \left( \frac{2\sqrt{1 - z^2}}{r} \right) - 2q(z; \epsilon) - 2 \int_{-1}^1 \frac{q(\xi; \epsilon) - q(z; \epsilon)}{|z - \xi|} d\xi + \mathcal{O}(r^2). \quad (\text{A8})$$

In the next section, we will solve for the concentration and velocity fields in the intermediate region I (see Fig. 3).

## 2. Region I: Intermediate region

At intermediate length-scales, for which the radial distance is larger than the range of the short-ranged interaction but much less than the length of the swimmer, the swimmer is locally approximated by an infinite cylinder of thickness  $2\epsilon$ . Here, we re-scale the radial coordinate with the swimmer slenderness ratio ( $\epsilon = a/l$ );

$$R = \frac{r}{\epsilon} \quad . \quad (\text{A9})$$

$$c(r, z) = C_I(R, z) \quad , \quad (\text{A10})$$

$$\mathbf{v}(r, z) = \mathbf{V}_I(R, z) \quad . \quad (\text{A11})$$

and the short-range interaction between the molecules and the swimmer surface also vanishes in this region (we have set  $\lambda = 0$ )

$$\psi(r - \epsilon S(z), z) = \Psi_I(R - S(z), z) = 0 \quad . \quad (\text{A12})$$

This is of course because we have assumed  $\lambda \ll \epsilon$  (i.e the swimmer cross-sectional radius is much larger than the interaction range of the potential).

Using the definition of the unit vectors,  $\hat{\mathbf{t}}, \hat{\mathbf{n}}$  in equations (17) and (16), the local gradients (away from the ends) in the re-scaled coordinates<sup>33</sup> are

$$\hat{\mathbf{n}} \cdot \nabla = \frac{1}{\epsilon} \frac{\partial}{\partial R} + \mathcal{O}(\epsilon) \quad , \quad (\text{A13})$$

$$\hat{\mathbf{t}} \cdot \nabla = \frac{\partial}{\partial z} + S_z(z) \frac{\partial}{\partial R} + \mathcal{O}(\epsilon) \quad , \quad (\text{A14})$$

$$J_{I,R}(R, z) = -\frac{\partial C_I(R, z)}{\partial R} \quad ; \quad J_{I,z}(R, z) = -\frac{\partial C_I(R, z)}{\partial z} \quad . \quad (\text{A15})$$

Since the interaction vanishes, the equation of motion for the reactant (product) concentration is the steady state diffusion equation (8), to leading order has only radial contributions :

$$0 = \epsilon^{-1} \frac{1}{R} \frac{\partial}{\partial R} \left( R \frac{\partial C_I}{\partial R} \right) ; \quad \text{with boundary condition} \quad - \frac{\partial C_I}{\partial R} \Big|_{R=S(z)} = J_{I,R}(S(z), z) , \quad (\text{A16})$$

where in the boundary condition, the (unknown) radial flux  $J_{I,R}(S(z), z)$  is determined by matching  $C_I(R, z)$ ,  $J_{I,R}(R, z)$  with the inner interaction layer (region II) on the swimmer surface. The other boundary condition is obtained from matching with the outer region 0. Therefore, solving equation (A16) and imposing the boundary condition,

$$C_I(R, z; \epsilon) = A(z; \epsilon) + J_{I,R}(S(z), z) S(z) \log \left( \frac{1}{R} \right) , \quad (\text{A17})$$

where  $A(z; \epsilon)$  is an unknown function of  $z$  resulting from integrating over  $R$ , must be determined by matching with the outer (region 0) concentration,  $C_0(r, z)$  given in equation (A6). Using this result, equation (22) and the second term in equation (23) are determined by  $J_{I,R}$ , while the first and third terms in equation (23) are determined by  $A(z; \epsilon)$ . We note that while  $A$  is determined by the properties of the concentration as  $r, z \rightarrow \infty$ ,  $J_{I,R}$  depends on the concentration near the reactive colloidal surface.

$$0 = \frac{1}{R} \frac{\partial}{\partial R} \left( R \frac{\partial V_{I,z}}{\partial R} \right) ; \quad \text{with slip boundary condition} \quad V_{I,z}|_{R=S(z)} = \hat{\mathbf{e}}_z \cdot \mathbf{V}^{\text{slip}}(z; \epsilon) . \quad (\text{A18})$$

The hydrostatic pressure difference,  $p(R, z) - p_\infty = P_I(R, z)$  vanishes at this order. The slip velocity,  $\mathbf{V}^{\text{slip}}(z; \epsilon)$  is obtained by matching with the inner interaction layer (region II). Solving equation (A18) gives,

$$V_{I,z}(R, z; \epsilon) = \hat{\mathbf{e}}_z \cdot \mathbf{V}^{\text{slip}}(z; \epsilon) + \mathcal{W}(z; \epsilon) \log \left( \frac{R}{S(z)} \right) . \quad (\text{A19})$$

The unknown function,  $\mathcal{W}(z; \epsilon)$  resulting from integrating over  $R$  will be found by matching with the fluid velocity in region 0, far from the swimmer surface.

### 3. Region II: Inner interaction (boundary) layer

$$\rho = \frac{r - \epsilon S(z)}{\lambda} ; \quad \text{where} \quad \lambda = \frac{L^*}{l}, \quad \epsilon = \frac{a}{l}, \quad \lambda \ll \epsilon . \quad (\text{A20})$$

In this region, we must perform an expansion in both of the small parameters  $\lambda$  and  $\epsilon$  (noting our assumption  $L^* \ll a$  or equivalently  $\lambda \ll \epsilon$ ). We therefore express the fields (concentration, fluid velocity and pressure) w.l.g. as

$$c(r, z) = \mathcal{C}(\rho, z) = \mathcal{C}^{(0,0)}(\rho, z) + \epsilon \mathcal{C}^{(0,1)}(\rho, z) + \lambda \mathcal{C}^{(1,0)}(\rho, z) + \lambda \epsilon \mathcal{C}^{(1,1)}(\rho, z) + \dots \quad (\text{A21})$$

$$\mathbf{v}(r, z) = \mathbf{V}(\rho, z) = \mathbf{V}^{(0,0)}(\rho, z) + \epsilon \mathbf{V}^{(0,1)}(\rho, z) + \lambda \mathbf{V}^{(1,0)}(\rho, z) + \lambda \epsilon \mathbf{V}^{(1,1)}(\rho, z) + \dots \quad (\text{A22})$$

$$p(r, z) = \mathcal{P}(\rho, z) = \lambda^{-2} \mathcal{P}^{(-2,0)}(\rho, z) + \lambda^{-1} \mathcal{P}^{(-1,0)}(\rho, z) + \epsilon \lambda^{-2} \mathcal{P}^{(-2,1)}(\rho, z) + \epsilon \lambda^{-1} \mathcal{P}^{(-1,1)}(\rho, z) + \dots \quad (\text{A23})$$

The leading terms for the concentration are of order  $\mathcal{O}(\epsilon^0)$ , while the leading order terms of velocity and pressure are of  $\mathcal{O}(\epsilon)$ . Therefore for an asymptotic analysis, we can simplify the expansions above. We can set  $\mathcal{C}^{(0,1)}(\rho, z) = \mathcal{C}^{(1,1)}(\rho, z) = 0$  and rename  $\mathcal{C}^{(0,0)}(\rho, z) \equiv \mathcal{C}^{(0)}(\rho, z)$  and  $\mathcal{C}^{(1,0)}(\rho, z) \equiv \mathcal{C}^{(1)}(\rho, z)$ . Similarly we can set  $\mathbf{V}^{(0,0)}(\rho, z) = \mathbf{V}^{(1,0)}(\rho, z) = 0$ ,  $\mathcal{P}^{(-2,0)}(\rho, z) = \mathcal{P}^{(-1,0)}(\rho, z) = 0$  and rename  $\mathbf{V}^{(0,1)}(\rho, z) \equiv \mathbf{V}^{(0)}(\rho, z)$ ,  $\mathbf{V}^{(1,1)}(\rho, z) \equiv \mathbf{V}^{(1)}(\rho, z)$ ,  $\mathcal{P}^{(-2,1)}(\rho, z) \equiv \mathcal{P}^{(-2)}(\rho, z)$  and  $\mathcal{P}^{(-1,1)}(\rho, z) \equiv \mathcal{P}^{(-1)}(\rho, z)$ .

$$c(r, z) = \mathcal{C}(\rho, z) = \mathcal{C}^{(0)}(\rho, z) + \lambda \mathcal{C}^{(1)}(\rho, z) + \dots \quad (\text{A24})$$

$$\psi(r - \epsilon S(z), z) = \Psi(\rho, z) = \Psi^{(0)}(\rho, z) + \lambda \Psi^{(1)}(\rho, z) + \dots \quad (\text{A25})$$

with associated flux written in components (see equation (8)),  $\mathbf{J}(r, z) = \mathcal{J}(\rho, z) = \mathcal{J}_z \hat{e}_z + \mathcal{J}_\rho \hat{e}_r$ . Away from the ends  $\hat{\mathbf{t}} \simeq \hat{e}_z$ ,  $\hat{\mathbf{n}} \simeq \hat{e}_r$  and the leading contribution to the flux is radial,

$$\mathcal{J}_\rho(\rho, z) = \lambda^{-1} \mathcal{J}_\rho^{(-1)}(\rho, z) + \mathcal{J}_\rho^{(0)}(\rho, z) + \mathcal{O}(\lambda) . \quad (\text{A26})$$

where

$$\mathcal{J}_\rho^{(-1)} = -\epsilon \left[ \frac{\partial \mathcal{C}^{(0)}}{\partial \rho} + \mathcal{C}^{(0)} \frac{\partial \Psi^{(0)}}{\partial \rho} \right] \quad ; \quad \mathcal{J}_\rho^{(0)} = -\epsilon \left[ \frac{\partial \mathcal{C}^{(1)}}{\partial \rho} + \mathcal{C}^{(0)} \frac{\partial \Psi^{(1)}}{\partial \rho} + \mathcal{C}^{(1)} \frac{\partial \Psi^{(0)}}{\partial \rho} \right] . \quad (\text{A27})$$

Hence, equation (8) becomes

$$\lambda^{-2} \frac{\partial \mathcal{J}_\rho^{(-1)}}{\partial \rho} + \lambda^{-1} \left[ \frac{\mathcal{J}_\rho^{(-1)}}{(\lambda \rho + \epsilon S(z))} + \frac{\partial \mathcal{J}_\rho^{(0)}}{\partial \rho} \right] + \mathcal{O}(1) = 0 , \quad (\text{A28})$$

with the corresponding boundary condition

$$\left[ \lambda^{-1} \mathcal{J}_\rho^{(-1)} + \mathcal{J}_\rho^{(0)} + \mathcal{O}(\lambda) \right]_{\rho=0} = \alpha(z) . \quad (\text{A29})$$

Therefore at leading order in the flux, equation (8) becomes

$$\frac{\partial \mathcal{J}_\rho^{(-1)}}{\partial \rho} = 0 \quad ; \quad \text{with boundary condition} \quad \mathcal{J}_\rho^{(-1)} \Big|_{\rho=0} = 0 . \quad (\text{A30})$$

Integrating equation (A30) once gives  $\mathcal{J}_\rho^{(-1)} = 0$ . Integrating again, using the definition of the flux, equation (A25) and equation (A27), we obtain an expression for the concentration in the inner region<sup>12</sup>

$$\mathcal{C}^{(0)}(\rho, z; \epsilon) = C_s(z; \epsilon) \exp \left( -\Psi^{(0)}(\rho, z) \right) \quad ; \quad C_s(z; \epsilon) = \mathcal{C}^{(0)}(\rho \rightarrow \infty, z) . \quad (\text{A31})$$

The as yet undetermined integrating constant,  $C_s(z; \epsilon)$  will be found by matching with the concentration,  $C_I(R, z)$  in the intermediate region.

$$\frac{\partial \mathcal{J}_\rho^{(0)}}{\partial \rho} = 0 \quad ; \quad \text{with boundary condition} \quad \mathcal{J}_\rho^{(0)} \Big|_{\rho=0} = \alpha(z) . \quad (\text{A32})$$

It follows then that

$$\mathcal{J}_\rho^{(0)}(\rho = 0, z) = \lim_{\lambda \rightarrow 0, \rho \rightarrow \infty} \mathcal{J}_\rho^{(0)}(\rho, z) = \alpha(z) , \quad (\text{A33})$$

which will provide the required boundary condition for matching with the flux in the intermediate (region I).

(Fluid velocity):

$$\mathbf{v}(r, z) = \mathbf{V}(r, z) = \epsilon \mathbf{V}^{(0)}(\rho, z) + \epsilon \lambda \mathbf{V}^{(1)}(\rho, z) + \dots \quad (\text{A34})$$

$$p(r, z) = \mathcal{P}(r, z) = \epsilon \lambda^{-2} \mathcal{P}^{(-2)}(\rho, z) + \epsilon \lambda^{-1} \mathcal{P}^{(-1)}(\rho, z) + \dots . \quad (\text{A35})$$

It is convenient to express equations (10) in cylindrical coordinates for axisymmetric velocity fields,

$$\frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial}{\partial r} \mathbf{v} \right) + \left( \frac{\partial^2}{\partial z^2} \mathbf{v} \right) - (\nabla \mathcal{P} + \epsilon \lambda^{-2} \mathcal{C} \nabla \Psi) = \mathbf{0} \quad (\text{A36})$$

or alternatively

$$\frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial}{\partial r} \mathbf{v} \right) + \left( \frac{\partial^2}{\partial z^2} \mathbf{v} \right) - \hat{\mathbf{e}}_z \left( \frac{\partial \mathcal{P}}{\partial z} + \epsilon \lambda^{-2} \mathcal{C} \frac{\partial \Psi}{\partial z} \right) - \hat{\mathbf{e}}_r \left( \frac{\partial \mathcal{P}}{\partial r} + \epsilon \lambda^{-2} \mathcal{C} \frac{\partial \Psi}{\partial r} \right) = \mathbf{0} \quad (\text{A37})$$

Using equations (A20,A21,A22,A23) and projecting the equations onto the local inner coordinates  $(\hat{\mathbf{t}}(z), \hat{\mathbf{n}}(z))$  which are functions of  $z$  only,  $\mathbf{v}^{(0)} = \mathcal{V}_n^{(0)} \hat{\mathbf{n}} + \mathcal{V}_t^{(0)} \hat{\mathbf{t}}$  (see Fig. 5) :

$$\hat{\mathbf{n}} : \quad 0 = \epsilon \left( -\lambda^{-3} \hat{\mathbf{n}} \cdot \hat{\mathbf{e}}_r \left[ \frac{\partial \mathcal{P}^{(-2)}}{\partial \rho} + \mathcal{C}^{(0)} \frac{\partial \Psi^{(0)}}{\partial \rho} \right] + \lambda^{-2} \frac{\partial^2 \mathcal{V}_n^{(0)}}{\partial \rho^2} - \lambda^{-2} \hat{\mathbf{n}} \cdot \hat{\mathbf{e}}_z \left[ \frac{\partial \mathcal{P}^{(-2)}}{\partial z} + \mathcal{C}^{(0)} \frac{\partial \Psi^{(0)}}{\partial z} \right] \right. \quad (\text{A38})$$

$$\left. + \frac{\lambda^{-1}}{(\lambda \rho + \epsilon S(z))} \frac{\partial \mathcal{V}_n^{(0)}}{\partial \rho} + \mathcal{O}(\lambda^0) \right), \quad (\text{A39})$$

$$\hat{\mathbf{t}} : \quad 0 = \epsilon \left( \lambda^{-2} \left[ \frac{\partial^2}{\partial \rho^2} \hat{\mathbf{t}} \cdot \mathbf{v}^{(0)} - \hat{\mathbf{t}} \cdot \nabla \mathcal{P}^{(-2)} - \mathcal{C}^{(0)} \hat{\mathbf{t}} \cdot \nabla \Psi^{(0)} \right] + \frac{\lambda^{-1}}{(\lambda \rho + \epsilon S(z))} \frac{\partial}{\partial \rho} \mathcal{V}_t + \mathcal{O}(\lambda^0) \right), \quad (\text{A40})$$

while equation (9) becomes at leading order in  $\lambda$

$$\epsilon \left( \lambda^{-1} \frac{\partial \mathcal{V}_n^{(0)}}{\partial \rho} + \mathcal{O}(\lambda^0) \right) = 0. \quad (\text{A41})$$

The leading order,  $\mathcal{O}(\lambda^{-3})$  terms in equation (A39) imply that the stresses induced by the interaction of the reactant (product) molecules with the swimmer surface are balanced by a local radial pressure gradient,

$$\frac{\partial \mathcal{P}^{(-2)}}{\partial \rho} + \mathcal{C}^{(0)} \frac{\partial \Psi^{(0)}}{\partial \rho} = 0. \quad (\text{A42})$$

Solving equation (A42), and using the already calculated  $\mathcal{C}^{(0)}$  given in equation (A31), we obtain the hydrostatic pressure profile<sup>11,34</sup>

$$\mathcal{P}^{(-2)}(\rho, z; \epsilon) = C_I(S(z), z; \epsilon) \left[ e^{-\Psi^{(0)}(\rho, z)} - 1 \right]. \quad (\text{A43})$$

To obtain it we have matched with the pressure in the intermediate (region I) which requires that  $\lim_{\rho \rightarrow \infty} \mathcal{P}^{(-2)}(\rho, z) = 0$  and  $\lim_{\rho \rightarrow \infty} \Psi^{(0)}(\rho, z) = 0$ . In the tangential direction, at leading order, i.e.  $\mathcal{O}(\lambda^{-2})$ , in equation (A40), the viscous stresses balance the pressure gradient (due to the concentration gradient) and the tangential stresses induced by the interaction of the molecules of reactant (product) with the swimmer surface

$$\frac{\partial^2}{\partial \rho^2} \mathcal{V}_t - \hat{\mathbf{t}} \cdot \nabla \mathcal{P}^{(-2)} - \mathcal{C}^{(0)} \hat{\mathbf{t}} \cdot \nabla \Psi^{(0)} = 0. \quad (\text{A44})$$

$$\frac{\partial^2 \mathcal{V}_n^{(0)}}{\partial \rho^2} = 0, \quad (\text{A45})$$

since from equation (17) ,  $\hat{\mathbf{n}} \cdot \hat{\mathbf{e}}_z = O(\epsilon)$ . Note that  $\hat{\mathbf{t}} \sim \hat{\mathbf{e}}_z$  away from the ends and  $\hat{\mathbf{t}} \sim \hat{\mathbf{e}}_\rho$  at the ends (see Fig. 5). Now, using equation (A43) and equation (A31), we can integrate equation (A44) twice to obtain the tangential velocity and hence the limiting value

$$\lim_{\lambda \rightarrow 0; \rho \rightarrow \infty} \mathcal{V}_t = \epsilon \mu(z) \hat{\mathbf{t}} \cdot \nabla C_I(z; \epsilon) . \quad (\text{A46})$$

In addition, from equations (A45, A41), the normal component of the velocity vanishes,  $\mathcal{V}_n^{(0)} = 0$ . Thus, we obtain the slip velocity<sup>11</sup>

$$\mathbf{V}^{\text{slip}}(z; \epsilon) = \epsilon \mu(z) \hat{\mathbf{t}} \cdot \nabla C_I(z; \epsilon) \hat{\mathbf{t}} , \quad (\text{A47})$$

where we define the diffusiophoretic mobility<sup>11</sup>  $\mu(z) = \int_0^\infty \rho \left[ 1 - e^{-\Psi^{(0)}(\rho, z)} \right] d\rho$ .

In the next section, we match the solutions from the three regions to find the unknown functions  $J_{I,R}(R, z)$ ,  $A(z; \epsilon)$ ,  $j(z)$ ,  $\mathcal{W}(z; \epsilon)$  and  $q(z)$  arising from our integrations.

#### 4. Asymptotic matching

(*Concentration*): First, we match the  $\mathcal{O}(\epsilon^0, \lambda^0)$  fluxes between regions I and II;

$$(\text{II}) \quad \lim_{\lambda \rightarrow 0, \rho \rightarrow \infty} \mathcal{J}_\rho^{(0)}(\rho, z) = \alpha(z) = \lim_{R \rightarrow S(z)} J_{I,R}(R, z) \quad (\text{I}) , \quad (\text{A48})$$

using equation (A33). Next, we match the  $\mathcal{O}(\epsilon^0, \lambda^0)$  concentrations between intermediate (region I) with the outer (region 0). Using equations (A6 and A17)

$$(\text{I}) \quad C_I(R, z; \epsilon) = A(z; \epsilon) + \alpha(z) S(z) \log \left( \frac{1}{R} \right) = \lim_{r \rightarrow \epsilon R \ll 1} C_0(r, z) \quad (0) . \quad (\text{A49})$$

we obtain the source line density  $j(z)$ , introduced in equation (A4) and the unknown function  $A(z; \epsilon)$  introduced in equation (A17) :

$$j(z) = \frac{1}{2} \alpha(z) S(z) ; \quad A(z; \epsilon) = C_\infty + \alpha(z) S(z) \log \left( \frac{2}{\epsilon} \right) - \frac{1}{2} \int_{-1}^1 \frac{d}{d\xi} \left[ \alpha(\xi) S(\xi) \right] \text{sgn}(z - \xi) \log |z - \xi| d\xi . \quad (\text{A50})$$

Hence we obtain the leading order concentration in region I

$$C_I(R, z; \epsilon) = C_I^{(-1)}(z) \log \left( \frac{1}{\epsilon} \right) + C_I^{(0)}(z) + \alpha(z) S(z) \log \left( \frac{S(z)}{R} \right) , \quad (\text{A51})$$

where we have defined

$$C_I^{(-1)}(z) = \alpha(z) S(z) , \quad (\text{A52})$$

$$C_I^{(0)}(z) = C_\infty + \alpha(z) S(z) \log \left( \frac{2}{S(z)} \right) - \frac{1}{2} \int_{-1}^1 \frac{d}{d\xi} \left[ \alpha(\xi) S(\xi) \right] \text{sgn}(z - \xi) \log |z - \xi| d\xi . \quad (\text{A53})$$

(*Velocity*): We first match the  $\mathcal{O}(\epsilon, \lambda^0)$  velocities between the inner (region II) with the intermediate (region I)



$$(II) \quad \lim_{\lambda \rightarrow 0; \rho \rightarrow \infty} \epsilon \mathcal{V}^{(0)} = \mathbf{V}^{\text{slip}}(z; \epsilon) \quad (I) . \quad (A54)$$

Next we match the  $\mathcal{O}(\epsilon, \lambda^0)$  velocities between the outer (region 0) with the intermediate (region I)

$$(I) \quad V_{I,z}(R, z; \epsilon) = V_z^{\text{slip}} + \mathcal{W}(z; \epsilon) \log \left( \frac{R}{S(z)} \right) = \lim_{r \rightarrow \epsilon R \ll 1} V_{0,z}(r, z) \quad (0) , \quad (A55)$$

where  $V_z^{\text{slip}} = \mathbf{V}^{\text{slip}} \cdot \hat{\mathbf{e}}_z$ . From equations (A8) and (A19) we find the unknown  $\mathcal{W}(z; \epsilon)$  and  $q(z)$ , independent of the choice of  $R$ , as long as  $\epsilon R \ll 1^{26-28}$ . Thus,  $\mathcal{W}(z; \epsilon) = -4q(z; \epsilon)$  and

$$U + (\hat{\mathbf{e}}_z \cdot \hat{\mathbf{t}}(z)) V^{\text{slip}} = 4q(z; \epsilon) \log \left( \frac{1}{\epsilon} \right) + 4q(z; \epsilon) \log \left( \frac{2\sqrt{1-z^2}}{S(z)} \right) - 2q(z; \epsilon) + 2 \int_{-1}^1 \frac{q(\xi; \epsilon) - q(z; \epsilon)}{|z - \xi|} d\xi , \quad (A56)$$

where ,  $V^{\text{slip}} = |\mathbf{V}^{\text{slip}}|$  and  $\hat{\mathbf{t}}(z) = \hat{\mathbf{t}}(S(z))$ . The propulsion speed  $U$  is determined by requiring (see Section III B) that the solution of the integral equation above satisfies the constraint of zero total force on the swimmer, i.e requiring

$$\int_{-1}^1 q(z) dz = 0 .$$

<sup>a</sup>

<sup>b</sup>Electronic mail: ramin.golestanian@physics.oxford.ac.uk

<sup>c</sup>Electronic mail: t.liverpool@bristol.ac.uk

<sup>1</sup>M. C. Marchetti, J. F. Joanny, S. Ramaswamy, T. B. Liverpool, J. Prost, M. Rao, and R. A. Simha, “Hydrodynamics of soft active matter,” *Rev. Mod. Phys.* **85**, 1143–1189 (2013).

<sup>2</sup>S. Ramaswamy, “The Mechanics and Statistics of Active Matter,” *Annu. Rev. Condens. Matter Phys.* **1**, 323–345 (2010).

<sup>3</sup>F. Julicher, K. Kruse, J. Prost, and J. Joanny, “Active behavior of the Cytoskeleton,” *Phys. Rep.* **449**, 3–28 (2007).

<sup>4</sup>R. Golestanian, T. B. Liverpool, and A. Ajdari, “Propulsion of a molecular machine by asymmetric distribution of reaction products,” *Physical review letters* **94**, 220801 (2005).

<sup>5</sup>E. Yariv, “Electrokinetic self-propulsion by inhomogeneous surface kinetics,” in *Proceedings of the Royal Society of London A: Mathematical, Physical and Engineering Sciences* (The Royal Society, 2010) p. rspa20100503.

<sup>6</sup>S. Michelin and E. Lauga, “Phoretic self-propulsion at finite péclet numbers,” *Journal of Fluid Mechanics* **747**, 572–604 (2014).

<sup>7</sup>W. F. Paxton, A. Sen, and T. E. Mallouk, “Motility of catalytic nanoparticles through self-generated forces,” *Chemistry—A European Journal* **11**, 6462–6470 (2005).

<sup>8</sup>E. Yariv and O. Schnitzer, “The electrophoretic mobility of rod-like particles,” *Journal of Fluid Mechanics* **719**, R3 (2013).

<sup>9</sup>S. Ebbens, D. Gregory, G. Dunderdale, J. Howse, Y. Ibrahim, T. Liverpool, and R. Golestanian, “Electrokinetic effects in catalytic platinum-insulator janus swimmers,” *EPL (Europhysics Letters)* **106**, 58003 (2014).

<sup>10</sup>R. Golestanian, T. Liverpool, and A. Ajdari, “Designing phoretic micro-and nano-swimmers,” *New Journal of Physics* **9**, 126 (2007).

<sup>11</sup>J. L. Anderson, “Colloid transport by interfacial forces,” *Annual Reviews of Fluid Mechanics* **21**, 61–99 (1989).

<sup>12</sup>J. Anderson, M. Lowell, and D. Prieve, “Motion of a particle generated by chemical gradients part 1. non-electrolytes,” *Journal of Fluid Mechanics* **117**, 107–121 (1982).

<sup>13</sup>W. F. Paxton, K. C. Kistler, C. C. Olmeda, A. Sen, S. K. St. Angelo, Y. Cao, T. E. Mallouk, P. E. Lammert, and V. H. Crespi, “Catalytic nanomotors: autonomous movement of striped nanorods,” *Journal of the American Chemical Society* **126**, 13424–13431 (2004).

<sup>14</sup>J. L. Moran, P. M. Wheat, and J. D. Posner, “Locomotion of electrolytic nanomotors due to reaction induced charge autoelectrophoresis,” *Physical Review E* **81**, 065302 (2010).

<sup>15</sup>J. Howse, R. Jones, A. Ryan, T. Gough, R. Vafabakhsh, and R. Golestanian, “Self-Motile Colloidal Particles: From Directed Propulsion to Random Walk,” *Phys. Rev. Lett.* **99**, 048102 (2007).

<sup>16</sup>J. Palacci, S. Sacanna, A. P. Steinberg, D. J. Pine, and P. M. Chaikin, “Living crystals of light-activated colloidal surfers,” *Science* **339**, 936–940 (2013).

<sup>17</sup>N. Yoshinaga and T. B. Liverpool, “Hydrodynamic interactions in dense active suspensions : From polar order to dynamical clusters,” *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **96**, 020603(R) (2017).

<sup>18</sup>S. Saha, R. Golestanian, and S. Ramaswamy, “Clusters, asters, and collective oscillations in chemotactic colloids,” *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **89**, 1–7 (2014), arXiv:1309.4947.

<sup>19</sup>I. Theurkauff, C. Cottin-Bizonne, J. Palacci, C. Ybert, and L. Bocquet, “Dynamic clustering in active colloidal suspensions with chemical signaling,” *Physical review letters* **108**, 268303 (2012).

<sup>20</sup>J. L. Moran, P. M. Wheat, and J. D. Posner, “Locomotion of electrocatalytic nanomotors due to reaction induced charge autoelectrophoresis,” *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **81**, 1–4 (2010), arXiv:1001.5462.

<sup>21</sup>M. N. Popescu, S. Dietrich, M. Tasinkevych, and J. Ralston, “Phoretic motion of spheroidal particles due to self-generated solute gradients,” *The European Physical Journal E* **31**, 351–367 (2010).

<sup>22</sup>B. Sabass and U. Seifert, “Nonlinear, electrocatalytic swimming in the presence of salt,” *J. Chem. Phys.* **136** (2012), 10.1063/1.4719538, arXiv:1202.3797.

- <sup>23</sup>O. Schnitzer and E. Yariv, “Osmotic self-propulsion of slender particles,” *Physics of Fluids* **27**, 031701 (2015).
  - <sup>24</sup>E. L. Jewell, W. Wang, and T. E. Mallouk, “Catalytically driven assembly of trisegmented metallic nanorods and polystyrene tracer particles,” *Soft matter* **12**, 2501–2504 (2016).
  - <sup>25</sup>A. Nourhani and P. E. Lammert, “Geometrical performance of self-phoretic colloids and microswimmers,” *Physical review letters* **116**, 178302 (2016).
  - <sup>26</sup>G. K. Batchelor, “Slender-body theory for particles of arbitrary cross-section in stokes flow,” *Journal of Fluid Mechanics* **44**, 419–440 (1970).
  - <sup>27</sup>R. G. Cox, “The motion of long slender bodies in a viscous fluid part 1. general theory,” *Journal of Fluid mechanics* **44**, 791–810 (1970).
  - <sup>28</sup>J. B. Keller and S. I. Rubinow, “Slender-body theory for slow viscous flow,” *Journal of Fluid Mechanics* **75**, 705–714 (1976).
  - <sup>29</sup>J. Geer, “Stokes flow past a slender body of revolution,” *Journal of Fluid Mechanics* **78**, 577–600 (1976).
  - <sup>30</sup>E. Tuck, “Some methods for flows past blunt slender bodies,” *Journal of Fluid Mechanics* **18**, 619–635 (1964).
  - <sup>31</sup>J. P. K. Tillett, “Axial and transverse Stokes flow past slender axisymmetric bodies,” *J. Fluid Mech.* **44**, 401–417 (1970).
  - <sup>32</sup>S. Shklyaev, J. F. Brady, and U. M. Córdova-Figueroa, “Non-spherical osmotic motor: chemical sailing,” *Journal of Fluid Mechanics* **748**, 488–520 (2014).
  - <sup>33</sup>M. Van Dyke, *Perturbation methods in fluid mechanics* (Parabolic Press, Incorporated, 1975).
  - <sup>34</sup>E. Yariv, “Electrokinetic self-propulsion by inhomogeneous surface kinetics,” *Proc. R. Soc. A Math. Phys. Eng. Sci.* **467**, 1645–1664 (2011).
- a
- <sup>b</sup>Electronic mail: [ramin.golestanian@physics.oxford.ac.uk](mailto:ramin.golestanian@physics.oxford.ac.uk)
- <sup>c</sup>Electronic mail: [t.liverpool@bristol.ac.uk](mailto:t.liverpool@bristol.ac.uk)
- <sup>1</sup>M. C. Marchetti, J. F. Joanny, S. Ramaswamy, T. B. Liverpool, J. Prost, M. Rao, and R. A. Simha, “Hydrodynamics of soft active matter,” *Rev. Mod. Phys.* **85**, 1143–1189 (2013).
  - <sup>2</sup>S. Ramaswamy, “The Mechanics and Statistics of Active Matter,” *Annu. Rev. Condens. Matter Phys.* **1**, 323–345 (2010).
  - <sup>3</sup>F. Julicher, K. Kruse, J. Prost, and J. Joanny, “Active behavior of the Cytoskeleton,” *Phys. Rep.* **449**, 3–28 (2007).
  - <sup>4</sup>R. Golestanian, T. B. Liverpool, and A. Ajdari, “Propulsion of a molecular machine by asymmetric distribution of reaction products,” *Physical review letters* **94**, 220801 (2005).
  - <sup>5</sup>E. Yariv, “Electrokinetic self-propulsion by inhomogeneous surface kinetics,” in *Proceedings of the Royal Society of London A: Mathematical, Physical and Engineering Sciences* (The Royal Society, 2010) p. rspa20100503.
  - <sup>6</sup>S. Michelin and E. Lauga, “Phoretic self-propulsion at finite péclet numbers,” *Journal of Fluid Mechanics* **747**, 572–604 (2014).
  - <sup>7</sup>W. F. Paxton, A. Sen, and T. E. Mallouk, “Motility of catalytic nanoparticles through self-generated forces,” *Chemistry—A European Journal* **11**, 6462–6470 (2005).
  - <sup>8</sup>E. Yariv and O. Schnitzer, “The electrophoretic mobility of rod-like particles,” *Journal of Fluid Mechanics* **719**, R3 (2013).
  - <sup>9</sup>S. Ebbens, D. Gregory, G. Dunderdale, J. Howse, Y. Ibrahim, T. Liverpool, and R. Golestanian, “Electrokinetic effects in catalytic platinum-insulator janus swimmers,” *EPL (Europhysics Letters)* **106**, 58003 (2014).
  - <sup>10</sup>R. Golestanian, T. Liverpool, and A. Ajdari, “Designing phoretic micro-and nano-swimmers,” *New Journal of Physics* **9**, 126 (2007).
  - <sup>11</sup>J. L. Anderson, “Colloid transport by interfacial forces,” *Annual Reviews of Fluid Mechanics* **21**, 61–99 (1989).
  - <sup>12</sup>J. Anderson, M. Lowell, and D. Prieve, “Motion of a particle generated by chemical gradients part 1. non-electrolytes,” *Journal of Fluid Mechanics* **117**, 107–121 (1982).
  - <sup>13</sup>W. F. Paxton, K. C. Kistler, C. C. Olmeda, A. Sen, S. K. St. Angelo, Y. Cao, T. E. Mallouk, P. E. Lammert, and V. H. Crespi, “Catalytic nanomotors: autonomous movement of striped nanorods,” *Journal of the American Chemical Society* **126**, 13424–13431 (2004).
  - <sup>14</sup>J. L. Moran, P. M. Wheat, and J. D. Posner, “Locomotion of electrolytic nanomotors due to reaction induced charge autoelectrophoresis,” *Physical Review E* **81**, 065302 (2010).
  - <sup>15</sup>J. Howse, R. Jones, A. Ryan, T. Gough, R. Vafabakhsh, and R. Golestanian, “Self-Motile Colloidal Particles: From Directed Propulsion to Random Walk,” *Phys. Rev. Lett.* **99**, 048102 (2007).
  - <sup>16</sup>J. Palacci, S. Sacanna, A. P. Steinberg, D. J. Pine, and P. M. Chaikin, “Living crystals of light-activated colloidal surfers,” *Science* **339**, 936–940 (2013).
  - <sup>17</sup>N. Yoshinaga and T. B. Liverpool, “Hydrodynamic interactions in dense active suspensions : From polar order to dynamical clusters,” *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **96**, 020603(R) (2017).
  - <sup>18</sup>S. Saha, R. Golestanian, and S. Ramaswamy, “Clusters, asters, and collective oscillations in chemotactic colloids,” *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **89**, 1–7 (2014), arXiv:1309.4947.
  - <sup>19</sup>I. Theurkauff, C. Cottin-Bizonne, J. Palacci, C. Ybert, and L. Bocquet, “Dynamic clustering in active colloidal suspensions with chemical signaling,” *Physical review letters* **108**, 268303 (2012).
  - <sup>20</sup>J. L. Moran, P. M. Wheat, and J. D. Posner, “Locomotion of electrocatalytic nanomotors due to reaction induced charge autoelectrophoresis,” *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **81**, 1–4 (2010), arXiv:1001.5462.
  - <sup>21</sup>M. N. Popescu, S. Dietrich, M. Tasinkevych, and J. Ralston, “Phoretic motion of spheroidal particles due to self-generated solute gradients,” *The European Physical Journal E* **31**, 351–367 (2010).
  - <sup>22</sup>B. Sabass and U. Seifert, “Nonlinear, electrocatalytic swimming in the presence of salt,” *J. Chem. Phys.* **136** (2012), 10.1063/1.4719538, arXiv:1202.3797.
  - <sup>23</sup>O. Schnitzer and E. Yariv, “Osmotic self-propulsion of slender particles,” *Physics of Fluids* **27**, 031701 (2015).
  - <sup>24</sup>E. L. Jewell, W. Wang, and T. E. Mallouk, “Catalytically driven assembly of trisegmented metallic nanorods and polystyrene tracer particles,” *Soft matter* **12**, 2501–2504 (2016).
  - <sup>25</sup>A. Nourhani and P. E. Lammert, “Geometrical performance of self-phoretic colloids and microswimmers,” *Physical review letters* **116**, 178302 (2016).
  - <sup>26</sup>G. K. Batchelor, “Slender-body theory for particles of arbitrary cross-section in stokes flow,” *Journal of Fluid Mechanics* **44**, 419–440 (1970).
  - <sup>27</sup>R. G. Cox, “The motion of long slender bodies in a viscous fluid part 1. general theory,” *Journal of Fluid mechanics* **44**, 791–810 (1970).
  - <sup>28</sup>J. B. Keller and S. I. Rubinow, “Slender-body theory for slow viscous flow,” *Journal of Fluid Mechanics* **75**, 705–714 (1976).
  - <sup>29</sup>J. Geer, “Stokes flow past a slender body of revolution,” *Journal of Fluid Mechanics* **78**, 577–600 (1976).

- <sup>30</sup>E. Tuck, “Some methods for flows past blunt slender bodies,” *Journal of Fluid Mechanics* **18**, 619–635 (1964).
- <sup>31</sup>J. P. K. Tillett, “Axial and transverse Stokes flow past slender axisymmetric bodies,” *J. Fluid Mech.* **44**, 401–417 (1970).
- <sup>32</sup>S. Shklyaev, J. F. Brady, and U. M. Córdova-Figueroa, “Non-spherical osmotic motor: chemical sailing,” *Journal of Fluid Mechanics* **748**, 488–520 (2014).
- <sup>33</sup>M. Van Dyke, *Perturbation methods in fluid mechanics* (Parabolic Press, Incorporated, 1975).
- <sup>34</sup>E. Yariv, “Electrokinetic self-propulsion by inhomogeneous surface kinetics,” *Proc. R. Soc. A Math. Phys. Eng. Sci.* **467**, 1645–1664 (2011).