Depletion forces often dominate the effective forces among (colloidal) particles and their interactions with walls when suspended in fluids containing additional depletion agents, e.g., a background of smaller particles, nonadsorbing polymers, micelles, or vesicles [1–3]. At sufficiently small colloid-colloid or colloid-wall distances, excluded volume interactions between the particles and their interactions with walls (the pressure) [5–7], until now such a direct connection between the nonequilibrium steady-state distribution of APs and the active depletion forces has not been reported in experiments.

In general, the mechanical force exerted on a probe particle is given by

$$F = \int_{\partial A} \mathbf{d} s \mathbf{n} \cdot \mathbf{\sigma},$$

where $\mathbf{\sigma}(r)$ is the local stress in the bath due to the probe and walls and $\partial A$ is an arbitrary boundary with the normal vector $\mathbf{n}$ enclosing the probe. In passive baths, the stress is directly related to the equilibrium distribution of depletants. For hard particles, highly accurate approximations have been constructed, but this approach also extends to other interactions [3,24].

Here we apply the theoretical framework of an “active stress” [25–27] to an experimental system to obtain the force exerted by APs onto a probe particle within a circular cavity from the measured AP distribution around the probe. The results are in excellent agreement with independent force measurements using optical tweezers and are also corroborated by numerical simulations. Notably, our approach does not require knowledge of the specific interactions between APs and the probe, thereby facilitating the application of this concept to many other synthetic or living active systems.

Active colloidal particles are made by evaporating 20 nm thin carbon caps onto silica spheres with diameter $2r_a \approx 2 \mu m$. They are suspended in a mixture of water and propylene glycol $n$-propyl ether (PnP) (0.4 mass fraction of PnP). Under laser illumination ($\lambda = 532$ nm, $I \approx 7 \mu W/\mu m^2$), the fluid near the light-absorbing caps locally demixes, leading to compositional surface flows and thus to self-propulsion of the particles [28,29]. Owing to gravity and hydrodynamic interactions with the lower sample plate, the translational and rotational motion is effectively restricted to two dimensions (2D) [28]. A suspension of APs with area fraction $0.31 \pm 0.02$ is laterally confined to a lithographically fabricated circular cavity with a 100 $\mu m$ diameter. For the slow propulsion velocity $v = 0.40 \pm 0.08 \mu m/s$ used in this study, no motility-induced phase separation, i.e., formation of particle clusters—is observed [30,31]. Under such conditions, the demixing zones of the binary fluid remain rather close to the AP surfaces, which largely reduces phoretic interactions between the particles. From the APs’ rotational diffusion time $\tau_r \approx 23.7 s$, one obtains their persistence length $\ell = v \tau_r \approx 9.5 \mu m$ (see the Supplemental Material [32]). For the measurement of active depletion forces, specifically designed probe particles were fabricated via a

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Δ symmetric around whether green laser illumination is off or on, whether the colloidal particles are passive or active (i.e., the probe is near the center of the cavity. Regardless of the two-photon laser writing process [41]. They consist of disks with diameter 2rp = 15 μm and thickness 3.5 μm with four legs underneath to minimize the friction with the substrate [Fig. 1(a)]. On top of the disks, we designed hemispheres with diameter 11 μm to guarantee a harmonic optical trapping potential when a vertically incident laser beam (λ = 1064 nm) is focused onto the probe. The chosen intensity and focus diameter of the laser yield an optical trap stiffness k = 0.50 ± 0.02 pN/μm (see the Supplemental Material [32]). The force exerted on the probe is then obtained from the probe’s mean displacement relative to the trap center Δx = xp − xtr [Fig. 1(b)]. Prior to each force measurement, we allow the system to equilibrate for ≈15 minutes with the AP’s activity turned off. After illuminating the sample, we wait ≈10 minutes to reach a steady state before recording the position of the probe and APs for 2000 s using a center-of-mass tracking algorithm at a frame rate of 2 Hz.

Figure 2(a) shows a typical snapshot of the experiment. From the time-averaged configurations we calculate the spatial density ρ(x, y) of APs, which is plotted in Figs. 2(b)–2(f) for different probe-wall distances normalized by the AP diameter, i.e., d′ = d/(2rp). In agreement with previous studies, as a result of the APs’ finite reorientation dynamics, their density is increased at the surfaces of the cavity wall and the probe [42]. In particular, at small d′, APs strongly accumulate in the wedges formed by the probe and the wall [10]. This asymmetry in the AP distribution leads to d′-dependent changes of the probe’s fluctuations relative to the optical trap center as shown by the corresponding probability distributions P(Δx). Figure 3(a) corresponds to the situation where the probe is near the center of the cavity. Regardless of whether the colloidal particles are passive or active (i.e., whether green laser illumination is off or on), P(Δx) is symmetric around Δx = 0, indicating the absence of an effective force (solid lines). However, compared to passive particles where P(Δx) agrees well with a Gaussian distribution, in the case of an active background P(Δx) becomes broader and exhibits non-Gaussian tails. Such tails are caused by the finite persistence length of APs and are in good agreement with other studies using active baths [43,44]. When the probe is positioned closer to the cavity wall, however, the maximum of the distributions shifts to the right relative to Δx = 0, and the distributions become asymmetric [Fig. 3(b)], both indicating an additional force acting on the probe.

From the probe’s mean displacement relative to the optical trap’s center in the x and y directions, we obtain the force \( F = (F_x, F_y) \) acting on the probe, which is shown with symbols in Fig. 4(a) as a function of d′. In particular, at short distances, \( F_x(d′) \) displays a strong nonmonotonic

FIG. 1. (a) Electron scanning microscope image (top) and schematic picture (bottom) of the probe particle used. The scale bars are 3 μm each. (b) Schematic illustration of the experimental situation. The probe particle at \( x_p \) and surface-to-surface distance to the circular wall \( d \) is trapped by an optical tweezers (\( x_{tr} \) denotes the center of the trap). The smaller spheres with the dark caps correspond to the APs.

FIG. 2. (a) Experimental snapshot showing APs (dark points) and the optically trapped probe particle inside the circular cavity. Measured AP density distributions are displayed for different probe-wall distances: (b) d′ = 0.46, (c) 1.02, (d) 1.59, (e) 2.78, (f) 6.30. Scale bars are 10 μm each.

FIG. 3. (a) Distribution \( P(Δx) \) at the center of the circular confinement in the presence of active (squares) and passive (circle) particles. The solid lines are Gaussian fits. (b) Distribution \( P(Δx) \) with the probe at \( d′ = 1.02 \) (stars), 1.59 (squares), and 6.30 (circles). The solid lines correspond to fits with a Gaussian distribution. The error bars show the standard deviation of experimental data calculated over five independent measurements.
The view expressed in Eq. (2) relates the nonvanishing inhomogeneity around the probe causes a large repulsive active depletion force \( F_x(d') \) is essentially zero and only fluctuates slightly around zero.

A qualitative understanding of the nonmonotonic and repulsive behavior of \( F_x(d') \) is immediately obtained from the density distributions shown in Figs. 2(b)–2(f). At a probe-wall distance \( d' = 0.46 \), APs are unable to pass the probe-wall spacing, which leads to a strong increase of \( \rho \) in the wedges formed by the probe and the wall. This density inhomogeneity around the probe causes a large repulsive active depletion force \( F_x \) acting on the probe [Fig. 2(b)], in stark contrast to attractive forces generated by passive depletion agents at such small distances. Upon slightly increasing \( d' \), APs are able to “squeeze” through the space between the probe and the wall, thereby further pushing the probe away from the wall. When \( d' \) becomes comparable to the AP diameter, this squeezing-induced effect disappears and the AP density enrichment near the wall is much less pronounced [Fig. 2(c)]. This explains the appearance of the first maximum of \( F_x \) [Fig. 4(a)]. Because APs generally tend to accumulate near walls, for \( d' > 1 \), on average, a dense monolayer forms at the circular cavity wall, and the above discussed accumulation of APs at the wedges will repeat in a similar fashion at larger distances. This is shown for \( d' = 1.59 \), where the density of APs between the probe and the wall is again increased [Fig. 2(d)], which rationalizes the second peak in \( F_x \) [Fig. 4(a)]. For larger distances, the density around the probe becomes more and more homogeneous, which eventually leads to the disappearance of an active depletion force [Figs. 2(e) and 2(f)].

As a first step towards a quantitative relationship between the AP distribution and the active depletion force, we determine from \( \rho(x,y) \) the angle-resolved AP density distribution \( \rho(\varphi) \) around the probe particle, with \( \varphi \) defined in Fig. 2(b) [for details regarding the determination of \( \rho(\varphi) \), see the Supplemental Material [32]]. Figure 4(b) shows \( \rho(\varphi) \) normalized by the total particle density \( \rho_0 \) for several values of \( d' \) [marked as colored symbols in Fig. 4(a)]. Clearly, the nonmonotonic behavior in \( F_x(d') \) is also reflected in the corresponding \( \rho(\varphi,d') \), suggesting a close link between the two quantities.

To yield a direct relationship between the active depletion force and the AP distribution, we write the force on the probe as \( \mathbf{F} = \int_\mathcal{A} d^3r \rho \nabla u \) with local density \( \rho(r) \) of active bath particles, and with \( u(r) \) as the pair potential characterizing the bath-probe interaction as a function of their (center-to-center) distance \( r \). The force thus depends on the averaged AP configuration given by their density profile with respect to the probe center [cf. Fig. 4(b)]. In contrast to a dilute passive bath in thermal equilibrium, where \( \rho(r) \propto e^{-u(r)/(k_B T)} \), such a relationship is no longer valid in the case of an active bath [46] and thus fails to predict the force \( \mathbf{F} \) [13].

Under steady-state conditions and regardless of whether the system is in thermal equilibrium or not, force balance dictates the validity of the hydrostatic equation \( \nabla \cdot \sigma = \rho \nabla u = 0 \) (in the absence of particle currents [26]). Exploiting the divergence theorem leads to Eq. (1). Importantly, if we choose a contour \( \partial \mathcal{A} \) outside the range of \( u(r) \), the interactions \( u(r) \) do not show up explicitly in the expression for the local stress [25]

\[
\sigma = -\rho k_B T \mathbf{I} - k_B T v_{\text{col}} \left[ \frac{\hat{v}}{2D_0} \rho \mathbf{I} - (\nabla \rho)^{ST} \right],
\]

(2)
even though they do shape the distribution of APs around the probe. Here, the local polarization field \( \mathbf{p}(r) \) of APs enters. Moreover, the local propulsion speed \( \hat{v}(r) \) is reduced compared to the bare propulsion speed \( v \) due to interactions with neighboring particles. The symmetric and traceless derivative is \( (\nabla \mathbf{p})^{ST} = \partial_i p_j + \partial_j p_i - \langle \nabla \cdot \mathbf{p} \rangle \delta_{ij} \). The view expressed in Eq. (2) relates the nonvanishing...
polarization due to the aggregation at the probe and confinement to an active stress tensor, but alternative views have been proposed in which the polarization is identified with an external one-body force [47]. However, this distinction changes neither the force balance nor the mechanical force $\mathbf{F}$ on the probe.

We now consider a circular contour $\partial \mathcal{A}$ with radius $r_b > r_p + r_a$ and normal vector $\mathbf{n} = \mathbf{e}_r$ given by the radial unit vector. We choose $r_b$ as close to the probe as possible but outside the range of the pair potential, $u(r > r_b) \approx 0$. Plugging Eq. (2) into Eq. (1), we obtain (for details, see the Supplemental Material [32])

$$
\mathbf{F} = -k_B T \left( 1 + \frac{v_x r_{v_{\text{eff}}}}{2D_0} \right) \int_0^{2\pi} d\varphi r_b \mathbf{e}_r \rho(r_b, \varphi) \\
+ k_B T v_x r_b \left[ \frac{\partial}{\partial r} \int_0^{2\pi} d\varphi \mathbf{p}(r, \varphi) \right]_{r=r_b}.
$$

The second integral involving the polarization can be eliminated by setting $\hat{\mathbf{p}} = D_0 \nabla \rho$, exploiting the fact that our symmetric probe does not generate a particle current. Moreover, symmetry dictates that $F_y = 0$ [cf. Fig. 4(a)]. If we further use the fact that the density decays exponentially away from the probe, we obtain the rather simple result [32]

$$
F_x = k_B T_{\text{eff}} C_x
$$

for the force, whereby the integral $C_x(r_b) = -\int_0^{2\pi} d\varphi r_b \cos \varphi \rho(r_b, \varphi)$ only depends on the density of APs outside the probe. While Eq. (4) resembles the passive result with an elevated effective temperature $T_{\text{eff}}(r_b)$, the distribution of APs around the probe markedly deviates from the equilibrium profile. The temperature $T_{\text{eff}}$ simply encodes the elevated density fluctuations due to the activity of the particles.

The solid line in the inset of Fig. 4(a) shows the predicted force $F_x$ calculated from the experimentally measured density profiles with $r_b = r_p + 2.2 r_a$. We observe excellent agreement with the independently measured forces, with an effective temperature $T_{\text{eff}} \approx 17 T$ for all distances $d'$. Note that for distances $d' < 1$, the integration contour would intersect the wall, which does not allow the evaluation of forces in this range. We remark that $k_B T_{\text{eff}}$ and $C_x$ individually depend on $r_b$, but $F_x$ is independent of the precise choice of the integration contour.

We corroborate our results with numerical simulations employing a minimal model of active Brownian particles. Within this model, hydrodynamic interactions are neglected, and only the excluded volume of APs and their self-propulsion with constant speed $v$ is considered. All model parameters are fixed by the experimental values. Further details are provided in the Supplemental Material [32]. Figure 5(a) shows the force $\mathbf{F}$ as a function of $d'$ obtained from simulations. As in the experiments [cf. Fig. 4(a)], the force experienced by the probe is always repulsive and oscillates before eventually decaying to zero. In addition, the positions of the maxima agree very well with experiments. The inset of Fig. 5(a) shows the measured force $F_x$ (symbols) together with the prediction of Eq. (4), which also for this model system shows excellent agreement with a single elevated temperature $T_{\text{eff}} \approx 16.25 T$ independent of the probe-wall separation.

The only difference lies in the magnitude of $\mathbf{F}$ and the angular densities $\rho(\varphi)$ [Fig. 5(b)], which are consistently smaller in simulations. To understand those differences in more detail, we have determined the distribution of contact times, i.e., the times APs remain near the lateral confinement and the probe particle, respectively. We find that using an entirely repulsive model underestimates the width of the experimentally determined distribution of contact times; see the Supplemental Material [32]. Adding an attraction to the model that effectively accounts for, e.g., phoretic attractions and the slowing down of AP velocities due to surface roughness or hydrodynamic effects yields better agreement with the independently measured forces.
consistency with the experimental data and concurrently increases the force magnitude, reaching good quantitative agreement [32].

In summary, we have demonstrated with experiments and simulations that the forces exerted by an active bath onto a probe particle can be obtained from the AP density around the probe by employing the recently proposed concept of active stress. Remarkably, this approach does not require explicit knowledge of system-specific AP-probe interactions; therefore, it is applicable to many other active systems. In addition to understanding the motion of Brownian objects in baths of synthetic active particles, our work may also elucidate the role of active depletion forces in living systems which are believed to contribute to the organization of intracellular structures [48,49].

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[1] S. Asakura and F. Oosawa, On interaction between two bodies immersed in a solution of macromolecules, J. Chem. Phys. 22, 1255 (1954).
[2] Y. Mao, M. Cates, and H. Lekkerkerker, Depletion force in colloidal systems, Physica (Amsterdam) 222A, 10 (1995).
[3] B. Götzelmann, R. Evans, and S. Dietrich, Depletion forces in fluids, Phys. Rev. E 57, 6785 (1998).
[4] C. Bechinger, R. Di Leonardo, H. Löwen, C. Reichhardt, G. Volpe, and G. Volpe, Active particles in complex and crowded environments, Rev. Mod. Phys. 88, 045006 (2016).
[5] L. Angelani, C. Maggi, M. L. Bernardini, A. Rizzo, and R. Di Leonardo, Effective Interactions Between Colloidal Particles Suspended in a Bath of Swimming Cells, Phys. Rev. Lett. 107, 138302 (2011).
[6] D. Ray, C. Reichhardt, and C. J. Olson Reichhardt, Casimir effect in active matter systems, Phys. Rev. E 90, 013019 (2014).
[7] R. Ni, M. A. Cohen Stuart, and P. G. Bolhuis, Tunable Long Range Forces Mediated by Self-Propelled Colloidal Hard Spheres, Phys. Rev. Lett. 114, 018302 (2015).
[8] W. Yan and J. F. Brady, The force on a boundary in active matter, J. Fluid Mech. 785, R1 (2015).
[9] A. P. Solon, Y. Fily, A. Baskaran, M. E. Cates, Y. Kafri, M. Kardar, and J. Tailleur, Pressure is not a state function for generic active fluids, Nat. Phys. 11, 673 (2015).
[10] L. R. Leite, D. Lucena, F. Q. Potiguar, and W. P. Ferreira, Depletion forces on circular and elliptical obstacles induced by active matter, Phys. Rev. E 94, 062602 (2016).
[11] G. Junot, G. Briand, R. Ledesma-Alonso, and O. Dauchot, Active Versus Passive Hard Disks Against a Membrane: Mechanical Pressure and Instability, Phys. Rev. Lett. 119, 028002 (2017).
[12] M. Z. Yamchi and A. Naji, Effective interactions between inclusions in an active bath, J. Chem. Phys. 147, 194901 (2017).
[13] P. Liu, S. Ye, F. Ye, K. Chen, and M. Yang, Constraint Dependence of Active Depletion Forces on Passive Particles, Phys. Rev. Lett. 124, 158001 (2020).
[14] F. Feng, T. Lei, and N. Zhao, Tunable depletion force in active and crowded environments, Phys. Rev. E 103, 022604 (2021).
[15] X.-L. Wu and A. Libchaber, Particle Diffusion in a Quasi-Two-Dimensional Bacterial Bath, Phys. Rev. Lett. 84, 3017 (2000).
[16] L. Ortlieb, S. Rafai, P. Peyla, C. Wagner, and T. John, Statistics of Colloidal Suspensions Stirred by Microswimmers, Phys. Rev. Lett. 122, 148101 (2019).
[17] L. Angelani, R. Di Leonardo, and G. Ruocco, Self-Starting Micromotors in a Bacterial Bath, Phys. Rev. Lett. 102, 048104 (2009).
[18] R. Di Leonardo, L. Angelani, D. Dell’Arciprete, G. Ruocco, V. Iebba, S. Schippa, M. P. Conte, F. Mecarini, F. De Angelis, and E. Di Fabrizio, Bacterial ratchet motors, Proc. Natl. Acad. Sci. U.S.A. 107, 9541 (2010).
[19] A. Sokolov, M. M. Apodaca, B. A. Grzybowski, and I. S. Aranson, Swimming bacteria power microscopic gears, Proc. Natl. Acad. Sci. U.S.A. 107, 969 (2010).
[20] S. A. Mallory, C. Valeriani, and A. Cacciuto, Curvature-induced activation of a passive tracer in an active bath, Phys. Rev. E 90, 032309 (2014).
[21] C. P. Royall, D. G. A. L. Aarts, and H. Tanaka, Fluid structure in colloid–polymer mixtures: The competition between electrostatics and depletion, J. Phys. Condens. Matter 17, S3401 (2005).
[22] D. Kleshchanok, R. Tuinier, and P. R. Lang, Direct measurements of polymer-induced forces, J. Phys. Condens. Matter 20, 073101 (2008).
[23] H. N. W. Lekkerkerker and R. Tuinier, Depletion interaction, in Colloids and the Depletion Interaction (Springer Netherlands, Dordrecht, 2011), pp. 57–108.
[24] P. Attard, Spherically inhomogeneous fluids: II. Hard-sphere solute in a hard-sphere solvent, J. Chem. Phys. 91, 3083 (1989).
[25] T. Speck, Collective forces in scalar active matter, Soft Matter 16, 2652 (2020).
[26] T. Speck and A. Jayaram, Vorticity Determines the Force on Bodies Immersed in Active Fluids, Phys. Rev. Lett. 126, 138002 (2021).
[27] T. Speck, Coexistence of active Brownian disks: Van der Waals theory and analytical results, Phys. Rev. E 103, 012607 (2021).
[28] J. R. Gomez-Solano, S. Samin, C. Lozano, P. Ruedas-Batuecas, R. van Roij, and C. Bechinger, Tuning the motility and directionality of self-propelled colloids, Sci. Rep. 7, 14891 (2017).
[29] J. R. Gomez-Solano, S. Roy, T. Araki, S. Dietrach, and A. Maciolek, Transient coarsening and the motility of optically heated Janus colloids in a binary liquid mixture, Soft Matter 16, 8359 (2020).
[30] I. Buttutoni, J. Bialké, F. Kümmel, H. Löwen, C. Bechinger, and T. Speck, Dynamical Clustering and Phase Separation
in Suspensions of Self-Propelled Colloidal Particles, Phys. Rev. Lett. 110, 238301 (2013).

[31] J. Palacci, S. Sacanna, A. P. Steinberg, D. J. Pine, and P. M. Chaikin, Living crystals of light-activated colloidal surfers, Science 339, 936 (2013).

[32] See Supplemental Material at http://link.aps.org/supplemental/10.1103/PhysRevLett.129.058001 for further details and a discussion of extended models citing Refs. [25,26,33–41].

[33] S. Das, A. Garg, A. I. Campbell, J. Howse, A. Sen, D. Velegol, R. Golestanian, and S. J. Ebbens, Boundaries can steer active Janus spheres, Nat. Commun. 6, 1 (2015).

[34] M. Schmid, D. Ludescher, and H. Giessen, Optical properties of photoresists for femtosecond 3D printing: Refractive index, extinction, luminescence-dose dependence, aging, heat treatment and comparison between 1-photon and 2-photon exposure, Opt. Mater. Express 9, 4564 (2019).

[35] J. R. Howse, R. A. L. Jones, A. J. Ryan, T. Gough, R. Vafabakhsh, and R. Golestanian, Self-Motile Colloidal Particles: From Directed Propulsion to Random Walk, Phys. Rev. Lett. 99, 048102 (2007).

[36] S. Bera, S. Paul, R. Singh, D. Ghosh, A. Kundu, A. Banerjee, and A. Adhikari, Fast Bayesian inference of optical trap stiffness and particle diffusion, Sci. Rep. 7, 1 (2017).

[37] J. D. Weeks, D. Chandler, and H. C. Andersen, Role of repulsive forces in determining the equilibrium structure of simple liquids, J. Chem. Phys. 54, 5237 (1971).

[38] B. Dünweg and W. Paul, Brownian dynamics simulations without Gaussian random numbers, Int. J. Mod. Phys. C 02, 817 (1991).

[39] G. Volpe, I. Buttinoni, D. Vogt, H.-J. Kümmcr, and C. Bechinger, Microswimmers in patterned environments, Soft Matter 7, 8810 (2011).

[40] X. Wang, S. Ramírez-Hinestrosa, J. Dobnikar, and D. Frenkel, The Lennard-Jones potential: When (not) to use it, Phys. Chem. Chem. Phys. 22, 10624 (2020).

[41] N. Anscombe, Direct laser writing, Nat. Photonics 4, 22 (2010).

[42] K. Schaar, A. Zöttl, and H. Stark, Detention Times of Microswimmers Close to Surfaces: Influence of Hydrodynamic Interactions and Noise, Phys. Rev. Lett. 115, 038101 (2015).

[43] C. Maggi, M. Paoluzzi, N. Pellicciotta, A. Lepore, L. Angelani, and R. Di Leonardo, Generalized Energy Equipartition in Harmonic Oscillators Driven by Active Baths, Phys. Rev. Lett. 113, 238303 (2014).

[44] A. Argun, A.-R. Moradi, E. Pince, G. B. Bagci, A. Imparato, and G. Volpe, Non-Boltzmann stationary distributions and nonequilibrium relations in active baths, Phys. Rev. E 94, 062150 (2016).

[45] A. Duzgun and J. V. Selinger, Active Brownian particles near straight or curved walls: Pressure and boundary layers, Phys. Rev. E 97, 032606 (2018).

[46] F. Ginot, I. Theurkauff, D. Levis, C. Ybert, L. Bocquet, L. Berthier, and C. Cottin-Bizonne, Nonequilibrium Equation of State in Suspensions of Active Colloids, Phys. Rev. X 5, 011004 (2015).

[47] A. K. Omar, Z.-G. Wang, and J. F. Brady, Microscopic origins of the swim pressure and the anomalous surface tension of active matter, Phys. Rev. E 101, 012604 (2020).

[48] M. Almonacid, W. W. Ahmed, M. Bussonnier, P. Mailly, T. Betz, R. Voituriez, N. S. Gov, and M.-H. Verlhac, Active diffusion positions the nucleus in mouse oocytes, Nat. Cell Biol. 17, 470 (2015).

[49] N. Razin, R. Voituriez, J. Elgeti, and N. S. Gov, Generalized Archimedes’ principle in active fluids, Phys. Rev. E 96, 032606 (2017).