I. INTRODUCTION

Soft deformable objects, such as liquid droplets, vesicles, cells, and synthetic capsules exhibit a complex behavior under flows. For example, in capillary flow, fluid vesicles [1], red blood cells (RBCs) [2–8], and synthetic capsules [9] deform to parachute shapes, and RBCs also deform to slipper shapes [4–7]. Shape transitions of fluid vesicles occur in simple shear flow [10–12]. Membrane wrinkling appears for fluid vesicles after inversion of an elongational flow [13,14] and for synthetic capsules in simple shear flow [15,16]. Among these soft objects, RBC have received a great deal of attention since they are important for both fundamental research and medical applications. In microcirculation, the deformation of RBCs reduces the flow resistance of microvessels. In patients with diseases such as diabetes mellitus and sickle cell anemia, the RBCs have a reduced deformability and often block the microvascular flow [2,17–20].

In a simple shear flow with flow velocity \( \mathbf{v} = \gamma \mathbf{e}_y \), fluid vesicles and RBCs show a transition from a tank-treading (TT) mode with a constant inclination angle \( \theta \) to a tumbling (TB) mode with increasing viscosity of the internal fluid \( \eta_m \) [21–24] or membrane viscosity \( \eta_{mb} \) [10,11]. This transition is described well by the theory of Keller and Skalak (KS) [21], which assumes a fixed ellipsoidal vesicle shape. Experimentally, synthetic capsules and RBCs show the oscillation of their lengths and \( \theta \), called swinging (SW) [15,25,26], during TT motion, and RBCs also transit from TB to TT with increasing \( \dot{\gamma} \) [26,27]. Recently, this dynamics was explained by the KS theory with the addition of an energy barrier for the TT rotation caused by the membrane shear elasticity [26,28]. More recently, this transition was also obtained by simulations [29,30]. However, the detailed dynamics has not yet been investigated.

For fluid vesicles in high shear flow, shape transitions [10–12] occur, and a swinging phase [24,31–34], where the shape and \( \theta \) oscillate around \( \theta = 0 \), appears between the TT and TB phases. This SW mode is also called trembling [24,33,34] or vacillating breathing [32]; it is explained by the KS theory extended to a deformable ellipsoidal vesicle [31] and the perturbation theory for a quasispherical vesicle [32–34]. Shape deformation plays an essential role in the SW of fluid vesicles. The deformation is not necessary to explain the SW of elastic capsules [15,25,26,28–30,35,36] but is required for quantitative analysis. In this paper, we extend the theory in Ref. [28] to include the shape deformation of RBCs and investigate the dynamics of deformable RBCs.

The internal fluid of RBCs behaves as a Newtonian fluid since RBCs do not have a nucleus and other intracellular organelles. The RBC membrane consists of a lipid bilayer with an attached spectrin network as cytoskeleton. The lipid bilayer is an area-incompressible fluid membrane. The shear elasticity of the composite membrane is induced by the spectrin network. Under physiological conditions, an RBC has a constant volume of \( V = 94 \mu \text{m}^3 \), surface area of \( S = 135 \mu \text{m}^2 \), \( \eta_m = 0.01 \text{Pa} \cdot \text{s} \), \( \eta_{mb} = 10^{-7} – 10^{-6} \text{Ns/m} \), membrane shear elasticity of \( \mu = 6 \times 10^{-6} \text{N/m} \), and bending rigidity of \( \kappa = 2 \times 10^{-19} \text{J} \) [3,12,17,37,38].

The models and results are presented with dimensionless quantities (denoted by a superscript \( \ast \)). The lengths and energies are normalized by \( R_0 = \sqrt{S/4\pi} \) and \( \mu R_0^2 \), respectively. For RBCs, they are \( R_0 = 3.3 \mu \text{m} \) and \( \mu R_0^2 = 6.5 \times 10^{-17} \text{J} \). There are two intrinsic time units: the shape relaxation time \( \tau = \eta_0 R_0/\mu \) by the shear elasticity \( \mu \) and the time of shear flow \( 1/\dot{\gamma} \); the reduced shear rate is defined as \( \dot{\gamma}_\ast = \dot{\gamma} \eta_0 \). The relative viscosities are \( \eta_m^\ast = \eta_m/\eta_0 \) and \( \eta_{mb}^\ast = \eta_{mb}/\eta_0 R_0 \), where \( \eta_0 \) is the viscosity of the outside fluid. In typical experimental conditions, the Reynolds number is low, \( Re < 1 \); hence, the effects of the inertia are neglected.

In Sec. II, we describe the extended KS theory [28] for an elastic capsule with a fixed ellipsoidal shape and the phase behavior of the capsule. In Sec. III, we introduce the shape equation for deformable RBCs and present the dynamics of deformed RBCs. The dependence of the function shape of the RBC free-energy potential is described in Sec. IV. Discussion and summary are given in Secs. V and VI, respectively. The comparison with experimental results is presented in Sec. V.

Swinging and synchronized rotations of red blood cells in simple shear flow

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The dynamics of red blood cells (RBCs) in simple shear flow was studied using a theoretical approach based on three variables: a shape parameter, the inclination angle \( \theta \), and phase angle \( \phi \) of the membrane rotation. At high shear rate and low viscosity contrast of internal fluid, RBCs exhibit tank-treading motion, where \( \phi \) rotates with swinging oscillation of shape and \( \theta \). At low shear rate, tumbling motion occurs and \( \theta \) rotates. In the middle region between these two phases, it is found that synchronized rotation of \( \phi \) and \( \theta \) with integer ratios of the frequencies occurs in addition to intermittent rotation. These dynamics are robust to the modification of the potential of the RBC shape and membrane rotation. Our results agree well with recent experiments.

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II. DYNAMICS OF ELASTIC CAPSULES WITH FIXED SHAPE

A. Models

1. Keller-Skalak theory

Keller and Skalak [21] analytically derived the equation of the motion of vesicles or capsules based on Jeffery’s theory [39]. In the KS theory, the vesicles are assumed to have a fixed ellipsoidal shape,

$$\left(\frac{x_1}{a_1}\right)^2 + \left(\frac{x_2}{a_2}\right)^2 + \left(\frac{x_3}{a_3}\right)^2 = 1,$$

(1)

where $a_i$ are the semiaxes of the ellipsoid, and the coordinate axes $x_i$ point along its principal directions. The $x_1$ and $x_2$ axes, with $a_1 > a_2$, are on the vorticity $(xy)$ plane and the $x_3$ axis is in the vorticity $(z)$ direction. The maximum lengths in three directions are $L_1 = 2a_1$, $L_2 = 2a_2$, and $L_3 = 2a_3$. The velocity field on the membrane is assumed to be

$$\mathbf{v}^m = \omega\hat{n}^m = \omega \left(\frac{a_1}{a_2} \frac{a_3}{a_1} \frac{a_2}{a_1}, 0 \right).$$

(2)

The energy $W_{ex}$ supplied from the external fluid has to be balanced with the energy dissipated in the vesicle, $W_{ex} = D_{in} + D_{mb}$ where $D_{in}$ and $D_{mb}$ are the energies dissipated inside the vesicle and on the membrane, respectively. The motion of the vesicle is derived from this energy balance. Then the motion of the inclination angle $\theta$ is given by

$$\frac{d\theta}{dt} = \frac{\gamma}{2} \left(-1 + f_0 f_1 \cos(2\theta)\right) - f_0 \omega = \frac{\gamma}{2} \left(-1 + B \cos(2\theta)\right),$$

(3)

$$B = f_0 \left(f_1 + f_2^1 \frac{f_1^1}{1 + f_2^1 (\eta_{mb}^1 - 1) + f_2^3 \eta_{mb}^3}\right),$$

(4)

$$\omega = -\frac{\gamma \cos(2\theta)}{2f_1 (1 + f_2^1 (\eta_{mb}^1 - 1) + f_2^3 \eta_{mb}^3)}. $$

(5)

The membrane-viscosity term was derived by Tran-Son-Tay et al. [17]. The factors appearing in Eqs. (3)–(5) are given by

$$f_0 = \frac{2}{(a_1/2 + a_2/a_1)},$$

$$f_1 = 0.5(a_1/a_2 - a_2/a_1),$$

$$f_2 = 0.5g(a_1^2 + a_2^2),$$

$$f_3 = 0.5E_3 R_0/\sqrt{f_1 V},$$

$$g = \int_0^\infty (a_1^2 + s)^{-3/2}(a_2^2 + s)^{-3/2}(a_3^2 + s)^{-1/2} ds,$$

$$\alpha_i = \frac{a_i}{(a_1 a_2 a_3)^{1/3}},$$

$$E_3 = \oint \mathbf{e}_i \cdot \mathbf{e}_j ds,$$

where $E_3$ is an integral over the membrane surface and $\mathbf{n}$ is the normal vector of the surface.

For $B > 1$, a stable fixed point $\theta = 0.5 \arccos(1/B)$ exists, and TT motion occurs, while for $B < 1$, there is no fixed point, and the angle $\theta$ periodically rotates (TB). As $\eta_{mb}$ or $\eta_{mb}$ increases, the transition from TT to TB motion occurs, where $B$ decreases from $B > 1$ to $B < 1$. The membrane viscosity $\eta_{mb}$ and the internal viscosity $\eta_{mb}$ have a similar effect; hence, an effective internal viscosity can be defined as $\eta_{eff} = \eta_{mb}^1 + f_3 \eta_{mb}^3$. The factor $f_3$ in $\eta_{eff}$ depends on the vesicle shape and can give different dynamics for deformable vesicles, in particular, for shape transformations between prolate and oblate vesicles [10,11].

The KS theory quantitatively predicts the TT-TB transition with increasing $\eta_{eff}$. However, it cannot explain the TB-TT transition with increasing $\gamma$. In the KS theory, vesicle motion does not depend on $\gamma$ except that the TT or TB rotation velocity increases linearly with $\gamma$.

2. KS Theory with an energy barrier

Skötheim and Secomb [28] extended the KS theory to take into account an energy barrier during TT membrane rotation. For RBCs and synthetic capsules with non-spherical rest shape, their membranes are locally deformed during the TT rotation. Fischer experimentally demonstrated that the RBC membrane rotates back to the original position when the shear flow is switched off [40]. To describe the energy barrier, a phase angle $\phi$ and free-energy potential $F(\phi)$ are introduced; see inset of Fig. 1. The potential is periodic, $F(\phi + \pi n) = F(\phi)$, and $\phi = 0$ at the rest shape. Thus, the motions of the inclination angle $\theta$ and phase angle $\phi$ are given by

$$\frac{d\theta}{\gamma dt} = \frac{1}{2} \left(-1 + f_0 f_1 \cos(2\theta)\right) - \frac{f_0 d\phi}{\gamma dt},$$

(6)

$$\frac{d\phi}{\gamma dt} = \frac{c_0 (\gamma' V')}{2f_1 (1 + f_2^1 (\eta_{mb}^1 - 1) + f_2^3 \eta_{mb}^3)} F' + \cos(2\theta),$$

(7)

where $c_0 = f_3 / 8 \pi f_1$. The equations of the original KS theory are recovered in the absence of barriers of the free energy $F$, i.e., $\partial F'/\partial \phi = 0$, where $\omega = d\phi/dt$ is independent of $\phi$.

Skötheim and Secomb used a simple potential $F_0(\phi) = E_0 \sin^3(\phi)$ and a reduced energy $U_e = f_3 E_0 f_3^1 / 2 f_1 \eta \gamma V = E_0^0 \gamma V \gamma' V'$. We employ the potential $F_0$ in this section and describe the dependence on the potential shape in Sec. IV. Equations (6) and (7) are numerically integrated using the fourth-order Runge-Kutta method. An oblate capsule with $L_2/L_1 = 0.25$ and $L_3/L_1 = 1$ is used as a model RBC.
Figures 1–4 show the dynamics of elastic capsules with fixed shapes. At low shear rate \( \gamma \) (large \( U_e \)), the capsules show TB motion, since the free-energy barrier locks the phase angle at \( \phi = 0 \). In TB, \( \theta \) rotates but \( \phi \) oscillates; see the trajectory at \( U_e = 0.81 \) in Fig. 3. At higher \( \gamma \) (smaller \( U_e \)) and low \( \eta_{rot} \), TT motion occurs. In TT, \( \phi \) oscillates (swing), and \( \phi \) rotates instead of \( \theta \); see the trajectory at \( U_e = 0.76 \) in Fig. 3. The oscillation of \( \phi \) or \( \theta \) in TB or TT occurs with the rotation frequency \( f_{rot}^\phi \) or \( f_{rot}^\theta \), respectively. Here, an angle change of \( \pi \) is counted as one rotation. Skotheim and Secomb [28] reported an intermittent phase between the TT and TB phases, where both rotations of \( \theta \) and \( \phi \) occur. The \( \phi \) (TT) rotation is intermittently interrupted by the \( \theta \) (TB) rotation slightly above the maximum energy barrier \( U_c^\phi \) of the TT phase. However, we found that the phases of synchronized rotation of \( \theta \) and \( \phi \) also exist in this middle range between the TB and TT phases. An infinite number of synchronization phases with integer ratios of \( f_{rot}^\phi \) and \( f_{rot}^\theta \) exist; see Fig. 2. This type of synchronization is called the Devil’s staircase [41].

The trajectories of the synchronized rotations with \( f_{rot}^\phi : f_{rot}^\theta = 1:1 \) and \( 2:1 \) are shown in Fig. 3. The former has the widest \( U_e \) range. The approach to synchronization is explained by the return map \( \phi_{rot}(\phi_0) \) at \( \theta = \pi/2 \) in Fig. 4. The curve \( \phi_{rot}(\phi_0) \) shifts to the left with increasing \( U_e \) and has two (stable and unstable) crossing points with the line

\[ \theta = 1:1 \]

\[ \phi = 1:1 \]

\[ \phi = 2:1 \]

\[ \phi = 3:1 \]

\[ \phi = 5:2 \]

\[ \phi = 3:2 \]

\[ \phi = 5:1 \]

Fig. 1. (Color online) Dynamic phase diagram of elastic capsules with fixed shapes calculated from Eqs. (6) and (7) with the potential \( F_0 = E_0 \sin^2(\phi) \). (a) Viscosity \( \eta_{rot} \) dependence for the RBC-like oblate shape with \( L_2/L_1 = 0.25 \) and \( L_1/L_1 = 1 \). (b) Aspect ratio \( L_2/L_1 \) dependence at the TT-TB transition viscosity \( \eta_{crit}^\theta = \eta_{crit}^\phi \) of the KS theory (\( U_e = 0 \)). The boundary lines of the TB (brown), TT (violet), and synchronization regions \([f_{rot}^\phi : f_{rot}^\theta = 1:1 \text{ red}, 2:1 \text{ blue}, \text{ and } 3:1 \text{ green}] \) are shown.

B. Results

Fig. 2. (Color online) Rotation frequency \( f_{rot}^\phi \) of the inclination angle \( \theta \) and phase angle \( \phi \) for the fixed oblate shape with \( L_2/L_1 = 0.25 \) at \( \eta_{crit}^\theta = 1 \). Numbers represent \( f_{rot}^\phi : f_{rot}^\theta \).

Fig. 3. (Color online) Dynamics of oblate capsules for the fixed shape. Top panel: time development of angles \( \theta \) and \( \phi \) at \( U_e = 0.78 \). Middle and bottom panels: trajectories on the phase space \((\theta, \phi)\). The capsules exhibit TT and TB rotations at \( U_e = 0.76 \) and 0.81, respectively. Synchronized rotations with \( f_{rot}^\phi : f_{rot}^\theta = 1:1 \) and \( 2:1 \) and intermittent rotation are observed at \( U_e = 0.78, 0.793, \) and 0.7782, respectively. The other parameters are the same as in Fig. 2.
Here, we have adapted it to RBC dynamics. The shape parameter $\tilde{f}_{rot} = f_{rot}/f_{rot}^{1:1}$ occurs at $0.77823 < U_c < 0.78722$. A dashed line represents $\phi_{\alpha=1} = \phi_{\alpha}$. The other parameters are the same as in Fig. 2.

$\phi_{\alpha=1} = \phi_{\alpha}$ at $0.77823 < U_c < 0.78722$, where the capsules approach the limit cycle with $\phi_{\alpha=1} = \phi_{\alpha}$. At slightly below or above the synchronization region ($0 < 0.77823 - U_c < 1$ or $0 < U_c - 0.78722 < 1$), intermittent rotation appears, like near the TT and TB regions [28], see the trajectory at $U_c = 0.7782$ in Fig. 3.

The qualitative behavior of capsules does not depend on the aspect ratios $L_2/L_1$ and $L_3/L_1$. However, the ranges of the synchronized rotations are narrower for more spherical capsules, as shown in Fig. 1(b). Thus, it would be difficult to observe synchronized rotations in quasispherical capsules. Note that the dynamics is independent of $L_3/L_1$ at $\eta_{eff} = \eta_{\star}(B=1)$.

A fluid vesicle has no membrane shear elasticity ($U_c = 0$) and shows no synchronization between $\theta$ and $\phi$ since the potential does not depend on $\ddot{\phi}$; see Fig. 1(a). At $0 < \eta_{eff} - \eta_{\star} < 1$, intermittent $\theta$ rotation also appears for the fluid vesicle. The tumbling frequency is given by $\tilde{f}_{rot} = \gamma \sqrt{1 - B^2/2 \pi \eta_{eff} - \eta_{\star}}$, in the KS theory [21]. A similar frequency dependence on $U_c$ is obtained in the extended KS theory: $f_{rot} = \sqrt{U_c - U_0^2}$ slightly above $U_0^2$ [28].

III. DYNAMICS OF DEFORMABLE RBCS

A. Shape equation

Previously, we extended the KS theory to include the shape deformation of fluid vesicles, on the basis of the perturbation theory [32,34,42] of quasispherical vesicles [31]. It showed very good agreement with experimental data [24].

Here, we have adapted it to RBC dynamics. The shape parameter $\alpha_{12} = (L_1 - L_2)/(L_1 + L_2)$ does not increase monotonically for elongation because of RBC dimples. Therefore, the shape parameter $\alpha_{13} = (L_1 - L_3)/(L_1 + L_3)$ is employed, where $d\alpha_{13}/d\alpha_{12} = 2$ for an oblate ellipsoid ($\alpha_{13} = 0$). The equation of the shape evolution is given by

$$\frac{d\alpha_{13}}{dt} = \left( 1 - \left( \frac{\alpha_{13}^{max}}{\alpha_{13}} \right)^2 \right) \frac{A_0}{\gamma^2 \alpha_{13}} \frac{\partial F^*}{\partial \alpha_{13}} + A_1 \sin(2\theta),$$

where $A_0 = 45/2 \pi (32 + 23 \eta_{mb}^* + 16 \eta_{mb}^2) V^0$ and $A_1 = 60/(32 + 23 \eta_{mb}^* + 16 \eta_{mb}^2)$. Here, the terms of $\eta_{mb}^*$ are added in $A_0$ and $A_1$ based on the theory in Refs. [33,34]. This revision improves the $\eta_{eff}$ dependence of fluid vesicles in Ref. [31]; the phase diagram for $\eta_{mb}^*$ becomes similar to that for $\eta_{eff}$.

The free energy $F(\alpha_{13}, \phi)$ is estimated by the simulation of a model RBC with $\phi = 0$, $\pi/2$ elongated by mechanical forces, where the RBC membrane is modeled as a triangular network [12]. In the simulation, 578 vertices are connected by a bond potential $U_{bond} = (k_{1}/2)(r - r_0)^2 [1 + (k_{2}/2)(r - r_0)^2]$, with $\mu = (\sqrt{3}/4)k_0 = 6 \times 10^{-14}$ N/m, $\kappa = 2 \times 10^{-19}$ J, and $k_2 = 1$. The area and volume of the RBC are kept constant by harmonic potentials. Our simulation reproduces the force-length curves of the optical-tweezer experiment [43] and other simulations [38,44,45]; see Fig. 5 in Ref. [12]. The effective force $F(\alpha_{13}, \phi)/\partial \phi$ is estimated from these force-length curves; see Fig. 5. The model RBC at $\phi = \pi/2$ has $9 \times 10^{-18}$ J higher energy than at $\phi = 0$ with $\alpha_{13} = 0.1$ in the absence of external forces. This height of the energy barrier agrees with the value $E_0 = 10^{-17}$ J in Ref. [28], which was estimated from Fischer’s experiments [40]. Abkarian et al. [26] estimated the height as $E_0 = \mu S(\alpha_{13} - \alpha_{13}/2)/2 \sim 3 \times 7 \times 10^{-10} [n^2 / \mu] \times \mu$ based on the velocity field of the KS theory [Eq. (2)]. However, it gives much higher barrier $E_0 \sim 1 \times 10^{-15}$ J for $\mu = 6 \times 10^{-6}$ N/m or smaller shear modules $\mu \sim 10^{-8}$ N/m for $E_0 = 10^{-17}$ J. Since the KS velocity field does not satisfy the local area conservation of membrane, it may give more stress on the membrane than the area-conserving velocity field [46], and the barrier height may be overestimated.

In this section, we employ the free-energy potential $F(\alpha_{13}, \phi) = F_1(\alpha_{13}) + F_2(\alpha_{13})\sin^2(\phi)$. The dependence on the potential function is discussed in Sec. IV. Instead of an interpolation [10,11,31], we used fit functions to obtain smooth functions for the numerical calculations; the normalized potentials $F_1(\alpha_{13}) = 5 \alpha_{13}^2 + (40/3)\alpha_{13}^3 + (230/4)\alpha_{13}^4$ and $F_2(\alpha_{13}) = 0.2 + 0.8 \alpha_{13}$; the shape parameter $\alpha_{12} = 0.56 + 0.35 \alpha_{13} - 0.23 \alpha_{13}^2 + 0.034 \alpha_{13}^3$; the coefficients $f_2 = 0.6018 + 0.064 \alpha_{13} - 0.19 \alpha_{13}^2 - 0.42 \alpha_{13}^3$ and $f_3 = 0.734 + 0.54 \alpha_{13} + 0.91 \alpha_{13}^2 + 3.2 \alpha_{13}^3$; and $\alpha_{13}^max = 0.7$. Equations (6)–(8) are numerically integrated using the fourth-order Runge-Kutta method.

In this model, the viscosity ratio of the membrane and inner fluid and the reduced volume are fixed at $\eta_{mb}^* / \eta_m^* = 3.1$ and $V^* = V/(4\pi R_0^3/3) = 0.64$. Experimentally, the viscosity $\eta_0^*$ of outside fluid is typically varied and $\eta_{mb}^*$ and $\eta_m^*$ are...
fixed at physiological values. Thus, $\eta_{in}=1$ corresponds to $\eta=0.01 \text{ Pa s}$. The reduced shear rate $\dot{\gamma}=1$ corresponds to $\dot{\gamma}=90$ or $180 \text{ s}^{-1}$ at $\eta_{in}=0.5$ or 1, respectively.

B. Results

The phase diagram and rotation frequencies of the deformable RBC in simple flow are shown in Fig. 6. The shape deformation does not qualitatively change the phase diagram. The tumbling phase appears in the bottom of Fig. 6(a) since $\dot{\gamma}$ is used for the vertical axis instead of $U_z \approx 1/\dot{\gamma}$.

In the TT phase, the shape parameter $\alpha_{13}$ and $\theta$ oscillate with the frequency $f_{rot}$; see Figs. 7(b) and 8(a). As $\gamma^*$ increases, the time-average $\langle \alpha_{13} \rangle$ increases, and the SW oscillation amplitudes of $\alpha_{13}$ and $\theta$ decrease; see Fig. 9(a). The peak-to-peak amplitude $\theta_{amp}$ is inversely proportional to $\gamma^*$ for both the deformable and fixed-shape RBCs. For the fixed-shape RBCs, the mean angle $\langle \theta \rangle$ increases with increasing $\gamma^*$ and reaches the angle of the KS theory at $U_z=0 (\dot{\gamma} \rightarrow \infty)$. For the deformable RBCs, $\langle \theta \rangle$ has a maximum and then decreases because of the elongation of the RBCs; see Figs. 9(b).

In the TT phase, $\alpha_{13}$ and $\theta$ oscillate with a fixed phase difference; see Fig. 8. The phase difference is calculated from the time difference $t_{max}^\alpha$ between the maximum values of $\alpha_{13}$ and $\theta$ and from $t_{min}^\alpha$ between the minimum values. The difference between $t_{max}^\alpha/T_\alpha$ and $t_{min}^\alpha/T_\alpha$ represents the asymmetry of the oscillation functions. In a sinusoidal function, $t_{max}^\alpha=t_{min}^\alpha$. It is found that $\alpha_{13}$ and $\theta$ show in-phase oscillation ($t_{\alpha}/T_\alpha=0$) at small $\gamma^*$, and the phase difference approaches $\pi/2$ with increasing $\gamma^*$. Walter et al. experimentally observed a phase difference of $\pi/4$ for synthetic capsules; see Fig. 7 in Ref. [15]. Our results agree with their experiments.

IV. DEPENDENCE ON POTENTIAL FUNCTION

We compared the dynamics of RBCs with the fixed and deformable shapes in the previous section. The fixation of the RBC shape can be interpreted as the bending rigidity $\kappa \rightarrow \infty$ or $F(\alpha_{13}, \phi)=k \alpha_{13}^4+F_\phi(\phi)$ with $k \rightarrow \infty$. This difference of the potential functions in $\alpha_{13}$ does not change the dynamics greatly. In this section, we investigate the dependence on the potential functions in $\phi$. For fixed-shape capsules, we compare the dynamics with three potential functions: $F_0=E_0 \sin^2(\dot{\phi})$, $F_4=E_0 \sin^4(\dot{\phi})$, and $F_8=E_0 \sin^8(\dot{\phi})$; see Fig. 11(a). The angle $\varphi$ is the rotational angle $\varphi=\arctan(x_i/y_i)$, where $r_i=(x_i, y_i, z_i)$ is the position of a tracer on the membrane. The phase angle $\phi$ is defined as $\phi=\arctan(x_i L_2/y_i L_1)$. The angle $\varphi$ can be defined without assuming an ellipsoidal shape. The potential $F_4$ has the sharpest peak.

As the peak of potentials ($F_0, F_4, F_8$) sharpens, the transition shear rates $\dot{\gamma}_0^\alpha$ and $\dot{\gamma}_0^\phi$ increase ($U_z$ decreases) since the maximum forces $\partial F/\partial \phi$ increases at constant $E_0$. Two limit cycles of the synchronized rotations can coexist for $F_4$ and $F_8$, while no coexistence is observed for $F_0$. The capsules approach different limit cycles with increasing and decreasing $\dot{\gamma}^*$. For the synchronized rotations with $f_{rot}^0/f_{rot}^\alpha=1:1$, a steeper but continuous change appears in the $f_{rot}$ curve for $F_4$.
FIG. 8. (Color online) Dynamics of the deformable RBCs at \( \eta^*_{in}=0.5 \). (a) Time evolution of \( \alpha_{13}, \theta \), and \( \phi \) at \( \dot{\gamma}=0.05 \). (b) Trajectories in the phase space \((\alpha_{13}, \theta)\) at \( \dot{\gamma}=0.02, 0.05 \), and 0.1. (c) Shear rate \( \dot{\gamma} \) dependence of the phase difference between \( \alpha_{13} \) and \( \theta \) oscillations at \( \eta^*_{in}=0.1 \). Solid and dashed lines represent \( T_{tt}^{\max}/T_{tt} \) and \( T_{tt}^{\min}/T_{tt} \), respectively, where \( T_{tt} \) is the TT rotation period, and \( T_{tt}^{\max}(T_{tt}^{\min}) \) are the time difference between the maximum (minimum) of \( \alpha_{13} \) and \( \theta \); see the explanation in (a).

than for \( F_0 \), and a discrete change with hysteresis appears for \( F_a \); compare Figs. 2(a) with Figs. 11(b) and 11(c).

The coexistence of two synchronized rotations occurs for the deformable RBCs with the free energy \( F_a(\alpha_{13}, \phi) = F_1(\alpha_{13}, \phi) + F_2(\alpha_{13}) \sin^2(\phi) \). The shape parameter \( \alpha_{13} \) and \( \theta \) steeply change at \( \phi = \pm 0.5 \pi \) during TT motion, as shown in Fig. 12. The oscillations of \( \alpha_{13} \) and \( \theta \) do not approach sinusoidal curves, unlike the case for \( F_0 \); compare Figs. 8(c) and 12(b). In the TT and TB phases, the averages and amplitudes of the angles \( \theta \) and \( \phi \) do not show significant differences between the potential shapes in \( \phi \) (\( F_0, F_a \), and \( F_F \)) and in \( \alpha_{13} \) (fixed or deformable shape); see Figs. 9 and 10.

V. DISCUSSION

Let us compare our results with the experimental results [26]. The TT-TB transition is observed at \( \dot{\gamma}=0.01 \sim 0.04 \) for

FIG. 9. (Color online) Time averages and peak-to-peak amplitudes of the (a) \( \alpha_{13} \) and (b) \( \theta \) oscillations during TT motion. Solid lines represent the deformable RBCs with the potential \( F_0 \) at \( \eta^*_{in}=0.1, 0.5 \), and 0.8. Dashed-dotted lines represent the deformable RBCs with the potential \( F_a \) at \( \eta^*_{in}=0.5 \). Dashed lines in (b) and (c) represent the RBCs with the fixed shape at \( \eta^*_{in}=0.5 \) and 2. The shear rate \( \dot{\gamma} \) is normalized by the minimum shear rate \( \dot{\gamma}_{in} \) of the TT phase.

\( \eta^*_{in}=0.45 \) in Ref. [26]. It is in good agreement with our results, \( \gamma^*_{in}=0.0158455 \) and \( \gamma^*_{in}=0.0173455 \) with \( F_0 \) and \( \eta^*_{in}=0.0272 \) and \( \eta^*_{in}=0.028 \) with \( F_a \). Akbarian et al. also reported the dependencies in the TT phase, \( f_{rot} \propto \dot{\gamma}(\theta_{eff} \propto 1/\dot{\gamma}) \), and the SW amplitude \( \theta_{amp} \propto 1/\dot{\gamma} \). We also obtained these dependencies for all types of potentials; see Figs. 7(a) and 9(c). Most of the dynamics in our calculations are qualitatively independent of the potential shapes, and the quantitative differences are smaller than the distribution widths of the experimental data. In the experiments, the transition shear rate and the other quantities have wide distributions because of the polydispersity of RBCs. RBCs become smaller and more viscous with age [17,18]. The viscoelasticity of the RBC membrane is changed by some diseases such as diabetes mellitus [19]. Further experimental and simulation studies are needed to tune up the RBC potential in our theoretical model.

In experiments, it is difficult to distinguish the intermittent rotations from transient rotations [26]. The intermittency has not yet been obtained by numerical simulations [29,30]. The synchronized rotations, in particular with \( f_{rot}^{\theta} \propto f_{rot}^{\phi}=1:1 \), would be much easier to observe in experiments and simu-
FIG. 10. (Color online) Time averages and peak-to-peak amplitudes of the (a) $\alpha_{13}$ and (b) $\phi$ oscillations during TB motion. Solid lines represent the deformable RBCs with the potential $F_a$ at $\eta_{\text{in}} = 0.1$, 0.5, and 0.8. Dashed-dotted lines represent the deformable RBCs with the fixed shape at $\eta_{\text{in}} = 0.5$. Dashed lines in (b) represent the RBCs with the fixed shape at $\eta_{\text{in}} = 0.5$ and 2. The shear rate $\dot{\gamma}$ is normalized by the maximum shear rate $\dot{\gamma}_{tb}$ of the TB phase.

FIG. 11. (Color online) (a) Three potentials $F_0 = E_0 \sin^2(\phi)$, $F_4 = E_0 \sin^4(\phi)$, and $F_8 = E_0 \sin^8(\phi)$. The rotation frequencies $f_{\text{rot}}$ of $\theta$ and $\phi$ for fixed-shape capsules with (b) $F_4$ and (c) $F_8$ at $\eta_{\text{in}} = 1$. Dashed or solid lines in (b) represent data obtained with increasing or decreasing $\dot{\gamma}$, respectively. The other parameters are the same as in Fig. 2.

FIG. 12. (Color online) Dynamics of the deformable RBCs with the potential $F_{a, c}$. (a) Time evolution of $\alpha_{13}$, $\theta$, and $\phi$ at $\dot{\gamma}$ = 1 and $\eta_{\text{in}}^* = 0.5$. (b) Solid and dashed lines represent $t_{\text{max}}^*/T_{\text{tt}}$ and $t_{\text{min}}^*/T_{\text{tt}}$, respectively.

In summary, we described the dynamics of RBCs in simple shear flow using a simple theory. The phase diagram...
of RBCs is divided into three regions: tank-treading, tumbling, and intermediate regions. In the intermediate regions, RBCs exhibit intermittent or synchronized rotations of the inclination angle $\theta$ and phase angle $\phi$. Synchronized rotations, in particular with $f_{\text{rot}}/f_{\text{rot}} = 1:1$, would be much easier to experimentally observe than intermittent rotations. In the TT (TB) phase, the shape and $\mathcal{H}(\phi)$ oscillate with the frequency of $\phi(\theta)$ rotation. The coexistence of two synchronized rotations can appear when the potential function of $\phi$ has a sharp peak. The other dynamic properties are not sensitive to the function shape of the free-energy potential. We focused on the dynamics of RBCs in this paper, but the resulting dynamics would be generally applicable to other elastic capsules.

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