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Direct Numerical Simulations of Electrophoresis of Charged Colloids

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We propose a numerical method to simulate electrohydrodynamic phenomena in charged colloidal dispersions. This method enables us to compute the time evolutions of colloidal particles, ions, and host fluids simultaneously by solving Newton, advection-diffusion, and Navier-Stokes equations so that the electrohydrodynamic couplings can be fully taken into account. The electrophoretic mobilities of charged spherical particles are calculated in several situations. The comparisons with approximation theories show quantitative agreements for dilute dispersions without any empirical parameters; however, our simulation predicts notable deviations in the case of dense dispersions.

Electrohydrodynamic phenomena are of great importance in physical, chemical, and biological science, and also in several engineering fields [1]. In the case of electrophoresis of charged particles, for example, the particles start to move on the application of external electric fields. The electric double layer, i.e., the cloud of counterions around charged particles, tends to be deformed and its distribution becomes anisotropic because of the applied external field and also of the friction between ions and fluids. The electrophoretic mobility of a single colloidal particle is then determined by the balance between the electrostatic driving force and the hydrodynamic frictional force acting on the particle. In this situation, the time evolutions of the colloidal particles, the ions, and the host fluids are described by coupled equations of hydrodynamics (Navier-Stokes) and electrostatics (Poisson) with proper boundary conditions imposed on the surfaces of the colloidal particles. However, the usual numerical techniques of partial differential equations are hopeless in dealing with the dynamical evolution of many-particle systems since the moving particle-fluid boundary condition must be treated at every discrete time step.

In recent years, efforts to resolve hydrodynamic interactions in colloidal dispersions have attracted a lot of attention. Various advanced methods have been proposed such as the Stokesian dynamics (SD) [2], the finite element method (FEM) [3], the lattice Boltzmann method (LBM) [4], the stochastic rotation dynamics [5], the fluid particle dynamics (FPD) [6], and yet another method which treats solid-fluid interaction efficiently [7]. Pioneering approaches have been proposed also to simulate charged colloidal dispersions without hydrodynamics [8–12]. Although extensions have been done to take into account the hydrodynamics by using SD [13], FEM [14], LBM [15–19], and FPD [20], quantitatively reliable simulations have not yet been performed successfully for many-particle dispersions due to the complexity of the system.

Recently, we developed a reliable and efficient numerical method, called the smoothed profile (SP) method [21,22], to resolve the hydrodynamic interactions acting on solid particles immersed in Newtonian host fluids. In the SP method, the original sharp boundaries between colloids and host fluids are replaced with diffuse interfaces with finite thickness $\xi$. This simple modification greatly improves the performance of numerical computations since it enables us to use the fixed Cartesian grid even for the problems with moving boundary conditions.

The SP method is not only applicable to the dispersions in Newtonian fluids, but particularly suitable for the particle dispersions in complex fluids. It has already been applied successfully to liquid crystal colloidal dispersions [23,24] and charged colloidal dispersions [25]. Field-particle hybrid simulations were performed where the average direction of the liquid crystal molecules and the density of ions were treated as coarse-grained continuum objects while colloids were treated explicitly as particles. The interaction between fields and particles was taken through the diffuse interface. The above methods for the dispersions in complex fluids are, however, not yet appropriate for simulating dynamical phenomena since hydrodynamic effects are completely neglected. The purpose of the present study is to establish an efficient and reliable simulation method applicable for electrohydrodynamic phenomena such as electrophoresis by combining our SP methods for hydrodynamic [21,22] and electrostatic [25] interactions.

In the present Letter, we briefly outline our numerical modeling for charged colloidal dispersions and then demonstrate the reliability of the combined SP method by comparing our numerical results with classical approximation theories [26–29]. Finally, comparisons are made for the electrophoretic mobilities of dense dispersions, where the simulation results show notable deviations from a mean-field–type theory according to the cell model [30,31].

Let us consider $N$ spherical particles with radius $a$, the mass $M_p$, and the inertia tensor $I_p$ in a host fluid consisting of multicomponent ions of species $\alpha$ with charges $Z_{\alpha}e$, where $e$ is the unit charge. The local number density of $\alpha$ ions is $C_\alpha(\mathbf{r},t)$ at a time $t$. The total charge on a colloidal particle at time $t$ is $Q_p(t) = \int d^3r C_\alpha(\mathbf{r},t) Z_{\alpha}e$. The total force acting on the particle is $F_p = \sum_{\alpha} \int d^3r \mathbf{E}_\alpha(\mathbf{r},t) Z_{\alpha}e$. The force on the particle is then given by $F_p = \sum_{\alpha} \int d^3r \mathbf{E}_\alpha(\mathbf{r},t) Z_{\alpha}e - \mathbf{F}_h - \mathbf{F}_d - \mathbf{F}_s$, where $\mathbf{F}_h$, $\mathbf{F}_d$, and $\mathbf{F}_s$ are hydrodynamic, electrohydrodynamic, and short-range forces, respectively. The force on the particle is then given by $F_p = \sum_{\alpha} \int d^3r \mathbf{E}_\alpha(\mathbf{r},t) Z_{\alpha}e - \mathbf{F}_h - \mathbf{F}_d - \mathbf{F}_s$.
potential due to the constant electric field
Then the charge neutrality
Hereafter, soft-core potential of the truncated Lennard-
distribution of the surface charge
represented smoothly everywhere in the system by
\[ C_a = \tilde{C}_a \exp[-\frac{1}{\sigma_a} (\Psi + \Psi_{ex})/k_BT]. \]
The bulk salt concentration \( \tilde{C}_a \) is related to the Debye
screening length \( \kappa^{-1} = 1/\sqrt{4\pi \lambda_B \sum Z_i^2 C_a} \), where \( \lambda_B = e^2/4\pi \varepsilon k_BT \) is the Bjerrum length, which is approximately
0.72 nm in water at 25 °C.
Simulations have been performed in a three-dimensional
cubic box with the periodic boundary conditions. The
linear dimension is \( L/\Delta = 64 \), where \( \Delta \) is the lattice spacing
chosen as a unit length, which is taken related to the
Bjerrum length as \( \Delta = 4\pi \lambda_B \). We use the particle radius
\( a = 5 \) and the thickness parameter \( \xi = 2 \) throughout the
present simulations. The host fluid contains 1:1 electrolyte
composed of monovalent counterions (\( \alpha = + \)) and coions
(\( \alpha = - \)). The unit of the energy and the electrostatic
potential is \( k_BT \) and \( k_BT/e \), respectively. The later corre-
spends to 25.7 mV at 25 °C. The nondimensional parameter
\( m_a = 2e k_BT f/a \omega/e^2 \) is set to \( m_a = m_\sigma = 0.184 \),
which corresponds to the value of KCl solution at 25 °C.
Our unit of time \( \tau = \Delta^2 f_a/k_BT \) corresponds to 0.44 μs.
We first consider a single charged particle moving with
the drift velocity \( \tilde{V} = (-\tilde{V}, 0, 0) \) in a constant electric field
\( \tilde{E} = (E, 0, 0) \). The electrophoretic mobility \( V/E \) is related
to the zeta potential \( \zeta \), which is defined as the electrostatic
potential on a slipping plane as
\[ V/E = \xi/e \]
when \( \zeta \) is small [1]. The Smoluchowski equation \( f = 1 \)
is valid in the limiting case \( \kappa a \to \infty \) [27], while the Hückel
equation \( f = 2/3 \) is valid in the opposite case \( \kappa a \to 0 \)
[28]. Henry derived an expression \( f = f_h (\kappa a) \) which is
valid for a general value of \( \kappa a \) [29]. These equations
indicate that the mobility is proportional to \( \zeta \); however,
this relation tends to fail for larger \( \zeta \) where the relaxation
effect due to deformations of electric double layer becomes
notable. O’Brien and White proposed an approximation
theory which is valid also for larger \( \zeta \) [26].
We have performed simulations for electrophoresis of a
single particle in linear response regimes \( E \approx 0.15 \) and
compared them with the O’Brien-White theory. A constant
uniform electric field \( E = 0.1 \), which corresponds to
2.85 × 10^5 V/cm, was applied. The terminal \( V \) was calcu-
lated for \( 50 \leq -Z \leq 750 \) with \( \kappa^{-1} = 10 \), corresponding
to the salt concentration 11 μmol/l at 25 °C in water. We
chose \( \nu = \eta/\rho = 5 \), so the Reynolds number \( Re = aV/\nu \)
remains small. In both the O’Brien-White theory and our
simulations, the zeta potential is assumed to be the electro-
static potential at the particle surface, \( \zeta = \Psi_{surface} \). In our
simulations, the surface charges were chosen as \( Z = -50, -100, -200, -300, -400, -500, \) and \( -750 \), correspond-
ing to \( y = 0.525, 1.044, 2.035, 2.927, 3.692, 4.332, \) and

\[ \mu_a = k_BT \ln C_a + Z_a e(\Psi + \Psi_{ex}) \]
5.510, respectively. Here the dimensionless zeta potential $y \equiv e\zeta/\kappa T$ is introduced. A relationship between the surface charge $|Z|e$ and the dimensionless zeta potential $y$ is shown in Fig. 1(a), where our numerical results are plotted together with an analytic solution of the nonlinear Poisson-Boltzmann (PB) equation [34] and the Debye-Hückel linearized solution $\zeta = |Z|e/4\pi a^2\varepsilon\kappa(1 + \kappa a^{-1})$. We see that our numerical results are consistent with the nonlinear PB theory. In Fig. 1(b), the dimensionless mobility $E_m \equiv 3\varepsilon\eta V/2e\kappa T E$ is plotted as a function of the dimensionless zeta potential with $\kappa a = 0.5$. It is clearly demonstrated that our method reproduced the O’Brien-White theory almost perfectly including the nonlinear regime $y \geq 2$ within only a few percent error. We emphasize that such a precise agreement with the theory has never been obtained by any simulation methods so far proposed. The distributions of charge density due to counterions and coions are shown in Fig. 1(c) for $y = 1.044$ and (d) for $y = 3.692$. One can see that the electric double layer is deformed considerably in the nonlinear regime (d), while it is almost isotropic in the linear regime (c). As mentioned before, the relaxation effect due to the deformed double layer causes the nonlinearity in $E_m$.

Our simulation method is easily applicable to dense dispersions consisting of many particles. We thus examined the effect of the particle concentration on the electrophoretic mobility. The linearized theory for a single particle Eq. (6) is still valid for dense dispersions when $E$ is small; however, $f$ is now depending on both $\kappa a$ and $\phi$. Simulations were carried out with $Z = -100$ and $E = 0.1$ for various particle volume fractions $\phi \equiv 4\pi a^3 N/3L^3$ to calculate $f(\kappa a, \phi) = \eta V/\varepsilon E$. The Debye length $\kappa^{-1}$ is taken to be 5 and 10, which correspond to $\kappa a = 1$ and 0.5, respectively. The corresponding salt concentration is 44 $\mu$mol/l for $\kappa^{-1} = 5$ and 11 $\mu$mol/l for $\kappa^{-1} = 10$, respectively. Figure 2 shows typical snapshots of the systems with (a) fcc, (b) bcc, and (c) random configurations [35]. The horizontal color map represents the charge density for $\kappa a = 1$ on a cross section perpendicular to the $z$ axis. In the cases of fcc and bcc, $E$ was applied perpendicular to the (1,0,0) and (1,1,1) faces, but we obtained very small differences within only 1%.

The mobility coefficient $f(\kappa a, \phi)$ for $\kappa a = 1$ and 0.5 is plotted as a function of $\phi$ in Figs. 3(a) and 3(b), respectively. We found that $f$ decreases rapidly with increasing $\phi$. Furthermore, the overall behavior looks almost independent of the particle configurations. A theoretical model has been proposed by Levine and Neale for the electrophoretic mobility of dense dispersions by using the cell model [30]. They assumed the situation that a spherical particle with radius $a$ is located at the center of a spherical container (cell) with radius $b$ and calculated $V$ as a function of $\kappa a$ and $\phi = (a/b)^3$. Ohshima proposed a simpler expression for the mobility coefficient $f$ according to the cell model [31]. Ohshima’s prediction is shown in Figs. 3(a) and 3(b) together with our numerical results. The overall agreement between our simulation and Ohshima’s theory is better in (a) with a smaller Debye length $\kappa^{-1} = 5 = a$ than in (b) with a larger one $\kappa^{-1} = 10$. In both (a) and (b), the simulation results tend to be larger than the theory as $\phi$ increases. We expect that the deviation is attributable to the occurrence of overlapping of the electric double layers for larger $\phi$ because such an effect is totally neglected in the theory. To this end, we estimated the effective radius $a + \kappa^{-1}$ of the ionically dressed particles and defined the effective volume fraction $\phi_{\text{ef}} \equiv 4\pi(a + \kappa^{-1})^3 N/3L^3 = [1 + (\kappa a)^{-1}]^3 \phi$. As is clearly seen in Figs. 3(a) and 3(b), our results agree well with Ohshima’s theory for $\phi_{\text{ef}} \leq 1$ where the effect of

**FIG. 1 (color).** Relationship between surface charge $|Z|e$ and dimensionless zeta potential $y$ (a). Our numerical data follow nicely on the analytic solution of the nonlinear PB equation [34] but deviates notably from the Debye-Hückel linearized theory. Dimensionless mobility $E_m$ of a single particle is plotted in (b) as a function of dimensionless zeta potential $y$. For comparison, results of Smoluchowski, Henry, and O’Brien-White for $\kappa a = 0.5$ are plotted. The color contours in (c) and (d) represent the total ionic charge density $\sum e Z C_a$ around a single particle for (c) $Z = -100$ and (d) $Z = -500$. The electric field is applied in the horizontal (+x) direction.

**FIG. 2 (color).** Snapshots of the electrophoresis of dense dispersions with (a) fcc, (b) bcc, and (c) random particle configurations. The color map represents the total ionic charge density $\sum e Z C_a$ in a plane perpendicular to the $z$ axis. The electric field is applied in the +x direction normal to (1,0,0) face for fcc and bcc. See movies in Ref. [35].
lapping is small. However, for \( \varphi_{\text{eff}} > 1 \) where the overlapping of the electric double layers becomes large, deviations between our simulations and the theory become notable. We emphasize that the present study is the first successful simulations which provide quantitative data consistent with recent studies which also take into account the effects of double layer overlapping [19,36,37].

In summary, we have developed a unique numerical method for simulating electrohydrodynamic phenomena in colloidal dispersions. The method was first applied to simulate electrophoresis of a single spherical particle, and we found that our method can reproduce quantitatively the reliable analytical theory proposed by O'Brien and White. Simulations were then performed for electrophoresis of colloids in dense dispersions, and we compared them with the theoretical analysis based on the cell model. We found that the cell model is reliable when overlapping of electric double layers is small but less reliable as overlapping becomes larger.

![FIG. 3 (color online). The mobility coefficient \( f(\kappa a, \varphi) \) as a function of the volume fraction \( \varphi \) in (a) \( \kappa a = 1 \) and (b) \( \kappa a = 0.5 \). The solid lines represent the approximation theory proposed by Ohshima [31]. The theory is confirmed to be accurate for \( \varphi_{\text{eff}} \leq 1 \), but tends to deviate from our numerical results for \( \varphi_{\text{eff}} > 1 \) where overlapping of the electric double layers becomes notable.](image-url)