Forces between single pairs of charged colloids in aqueous salt solutions

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Forces between single pairs of negatively charged micrometer-sized colloids in aqueous solutions of monovalent, divalent, or trivalent counter-ions at varying concentrations have been measured by employing optical tweezers. The experimental data have been analyzed by using the Derjaguin-Landau-Verwey-Overbeek (DLVO) theory and a numerical solution of the Poisson-Boltzmann (PB) equation. With monovalent counter-ions, the data are well described by the DLVO and PB theories, suggesting that the DLVO theory is adequate to describe the colloidal forces at these conditions. At higher counter-ion valence, the approximations within the two theories become evident.

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I. INTRODUCTION

Colloidal dispersions possess a large number of interesting and important properties and hence find applications as coatings, aerosols, and ceramics and for separation or filter processes. The size and charge of the colloids can be varied extensively, and thus the range and strength of the intercolloidal interactions can be adjusted. This makes colloidal dispersions an excellent model system to investigate fundamental issues in condensed matter physics [1]. A long-standing issue is the electrostatic interaction among charged colloids. For instance, the screening of charged colloids by counter-ions is central for the understanding of liquid suspensions, the stability of colloidal dispersions, and particle aggregation. Colloidal crystals are often considered as a model system for atomic crystals, with the advantage of tunable repulsion by controlling the salt concentration [2,3]. Another important subject is the behavior of vesicles and compartment stability in life science [4–8]. The latter ones play also a role in problems of drug delivery [9] and in microfluidic devices [6–8]. Also membrane fusion is strongly controlled by electrostatic interactions, by regulating the distance between the two opposite intermediate membrane “buds,” which initiate the fusion [4].

Optical tweezers [10] are ideal tools to carry out experiments with micrometer-sized objects in materials research [11–13], biological sciences [14–16], and soft matter research [17,18]. The idea behind optical tweezers is using strongly focused laser beams to trap a small dielectric particle in its focal point. Several considerations—e.g., laser power, ratio of the refractive index of the trapped particle and of its surrounding medium, trapped bead size, and bead position in the trap—have to be considered to optimize the use of optical tweezers. Their unique ability to hold and manipulate a single particle in a suitable medium without mechanical contact enables exciting new experiments in micro rheology [19,20]. Optical tweezers allow studying in detail the interactions between single pairs of colloids with a force resolution on the order of 0.3 pN and nanometer positioning accuracy.

The interaction potential between two colloids in solutions is intensely discussed in the literature [21–32] and remains a challenge for both experiment and theory. Despite its limitations, the Derjaguin-Landau-Verwey-Overbeek (DLVO) theory [21,22] with a repulsive electrostatic and an attractive van der Waals interaction still constitutes the cornerstone in describing the interaction between charged colloids in aqueous salt solutions since more than 50 years. The analytic expression of the repulsive part of the DLVO potential can be obtained by applying approximations to the solution of the Poisson-Boltzmann (PB) equation in planar geometry and employing the Derjaguin approximation (see Sec. III and the Appendix). The applicability of the DLVO theory deteriorates at increasing magnitude of the surface potential and charge due to the linearization and superposition approximation and at increasing counter-ion valence due to the neglect of small-ion correlations.

In the present contribution, we report on measurements of forces between single pairs of negatively charged colloids in aqueous salt solutions using optical tweezers. Moreover, our observations are analyzed by using the DLVO theory and solving the Poisson-Boltzmann equation numerically.

II. METHODS AND MATERIAL

Figure 1(a) illustrates the experimental setup, in which one colloid is fixed at the tip of a micropipette and the other one in an optical trap. An inverted microscope (Axiovert S 100 TV, Carl Zeiss, Jena, Germany) was used, and the optical trap was realized with a diode-pumped Nd:YAG laser (1064 nm, 1 W, LCS-DTL 322; Laser 2000, Wessling, Germany), power stabilized to achieve long-term stability. Additionally, the profile of the laser beam was monitored to ensure optimal trapping conditions. After passing an optical isolator, a quarter-wave plate was used to produce circularly polarized light to exclude effects due to reflection differences of the mirrors between the p and s polarization of the laser light. The beam was expanded and coupled into the back aperture of the microscope objective (Plan-NeoFluor 100 × 1.30 Oil, Carl Zeiss, Jena, Germany). Video imaging and the optical position detection were accomplished by a digital camera (KPF 120, Hitachi, Düsseldorf, Germany). The opti-
which was determined with an accuracy of ±0.3 pN.

The force versus separation dependence \( F(D) \) of the interaction between one single pair of negatively charged colloids was obtained by image analysis [Fig. 1, panels (b) and (c)]. From the digital images, the displacement of the colloid in the optical trap from the equilibrium position and the separation between the two colloids were determined using a custom-made algorithm with a special edge detection routine in a LabView environment [35,36]. A 60° section, where the two diffraction images overlap [Fig. 1(c)], was removed, and the pathway of the remaining inner edge (given by maximum intensity contrast) was detected and fitted to a circle. The center of the circle defined the position of the center of the colloid. This tracking procedure enabled a reduction of the effects of artifacts in \( x \) and \( y \) positions to below 2 nm.

Spherical polystyrene beads (microParticles, Berlin, Germany), possessing a negative charge due to surface sulfate groups with \( pK_a = 1.92 \), have been used as colloids. To exclude effects arising from differences among the colloids (variations in diameter, surface roughness, surface charge, etc.), the experiments were performed with single pairs of colloids with the surrounding medium changing by flushing the sample cell with salt solutions of different concentration and valency. With this procedure an experiment with a single pair of colloids lasted typically 4–6 h. A measurement sequence comprised 0.3–30 mM NaCl, 0.15–3 mM CaCl\(_2\), and 0.003–0.1 mM LaCl\(_3\) salt solutions at \( pH \approx 7.5 \). The colloid concentration is below 10\(^{-18}\) M, and hence their effect on the ionic strength is negligible. After finalizing the measurement of each valence, the reproducibility of the experiment as a whole was ensured by remeasuring the force-separation dependence in the initial 0.3-M NaCl solution. At the end of a measurement the average diameter of the two colloids was determined by measuring the force-separation dependence in a 1-M NaCl solution, where the force is well described by a hard-sphere potential (data not shown). For all colloids a diameter of 2.26±0.02 μm was obtained, which was in accordance with light scattering experiments.

### III. Theory

The experimental force-distance data have been analyzed using (i) a size-corrected screened Coulomb interaction and (ii) a numerical approach based on the PB equation in planar geometry augmented with the Derjaguin approximation. The first approach originates from the linearized PB equation in spherical geometry (also referred to as the Debye-Hückel theory). The size-corrected screened Coulomb force is often taken as the electrostatic contribution to the DLVO force acting between two charged colloids (see, e.g., Ref. [1]), but this expression generally differs from the original one given by DLVO [21,22]. In the Appendix we show that the two expressions become the same in a certain limit that approximately holds at the present experimental conditions. The equation proposed by DLVO is only applicable for colloids in symmetric electrolytes; hence, it cannot be used here.

In the screened Coulomb formalism, the electrostatic interaction energy \( U \) operating between two equal colloids
possessing the charge $Z$ and radius $R$ at the center-to-center distance $r$ is given by\[1\]

$$U(r) = \frac{(eZ)^2}{4\pi\varepsilon_0\varepsilon_r} \exp[-\kappa(r - 2R)] \left(1 + \kappa R\right)^2 r,$$  

where $e$ is the elementary charge, $\varepsilon_0$ the permittivity of vacuum, $\varepsilon_r$ the relative permittivity of the solution, and

$$\kappa = \left[\frac{1}{\varepsilon_0\varepsilon_r kT} \sum (e_z i)^2 c_{i,\text{bulk}}\right]^{1/2}$$

the inverse Debye screening length, with $k$ being Boltzmann constant, $T$ the temperature, $z_i$ the valence of species $i$, and $c_{i,\text{bulk}}$ the bulk concentration of species $i$. The interaction energy and the force at surface-to-surface separation $D = r - 2R$ become

$$U(D) = \frac{(eZ)^2}{4\pi\varepsilon_0\varepsilon_r} \exp(-\kappa D) \left(1 + \kappa R\right)^2 (D + 2R),$$

and

$$F(D) = \frac{(eZ)^2}{4\pi\varepsilon_0\varepsilon_r} \exp(-\kappa D) \left[1 + \kappa (D + 2R)\right] \left(1 + \kappa R\right)^2 (D + 2R)^2.$$

Since we do not have means to characterize the two colloids individually, we assume their charge and their radius to be equal. The van der Waals attraction also entering in the DLVO interaction can here be neglected, since it contributes only at high salt concentrations ($\geq 1$ M NaCl) and at surface separations smaller than 10 nm [37–39], in agreement with results from atomic force microscopy [40].

Our numerical PB approach starts with two planar surfaces with the surface charge density $\sigma$, which are separated by the distance $D$ and have their normals in the $z$ direction. The intervening medium is in equilibrium with a bulk electrolyte solution. Because of symmetry, only one-half of the system needs to be considered, $0 \leq z \leq b, b = D/2$. The amount of ions and their distribution between the two surfaces can be approximated by the PB equation according to

$$\varepsilon_0\varepsilon_r \frac{d^2 \phi(z)}{dz^2} = -e \sum c_{i}(z),$$

where

$$c_{i}(z) = c_{i,\text{bulk}}(z) \exp[-e_z i \phi(z)/kT],$$

with the boundary conditions

$$\left.\frac{d\phi(z)}{dz}\right|_{z=0} = -\frac{\sigma}{\varepsilon_0\varepsilon_r},$$

and

$$\left.\frac{d\phi(z)}{dz}\right|_{z=b} = 0.$$

In Eqs. (5)–(8), $\phi(z)$ is the electrostatic potential at position $z$ relative to the bulk, $c_{i}(z)$ the bulk concentration of species $i$ at position $z$, $c_{i,\text{bulk}}$ the concentration of species $i$ in the bulk, and $z_i$ the valence of species $i$. The numerical solution of Eq. (5)–(8) gives us the potential profile $\phi(z)$ and the

FIG. 2. (Color online) Force $F$ vs separation $D$ for one single pair of colloids in aqueous solution of varying salt and salt concentrations from experiments (symbols) and the DLVO theory with fitted values of the parameters $Z$ and $R$ (curves). (a) NaCl at 0.3 mM (black squares), 0.55 mM (red circles), 1 mM (green triangles), 2 mM (blue nablás), 4 mM (cyan diamonds), 10 mM (magenta left triangles), and 30 mM (orange right triangles). To ensure a full reproducibility of the exchange of the medium and to exclude hysteresis effects due to possible adsorption effects on the colloids the sample cell was flushed again with 0.3 mM NaCl (gold diamonds). (b) CaCl$_2$ at 0.15 mM (black open squares), 0.3 mM (red open circles), 0.5 mM (green open triangles), 1 mM (blue open nablás), 1.5 mM (cyan open diamonds), 3 mM (magenta open left triangles), 0.15 mM (orange open right triangles), and finally 0.3 mM NaCl (gold open diamonds). (c) LaCl$_3$ at 3 $\mu$M (black red solid square), 10 $\mu$M (black green solid circle), 30 $\mu$M (black blue solid triangle), 100 $\mu$M (black cyan solid nabla), 3 $\mu$M (black magenta solid diamonds), and 0.3 mM NaCl solution (black gold solid diamonds). Some indicative error crosses are given.
The force between two spherical and charged colloids $F(D)$ at a surface-to-surface separation $D$ is obtained from $U_{\text{plane}}(D)$ by the Derjaguin approximation [21,37] according to

$$F(D) = \pi R \frac{U_{\text{plane}}(D)}{A},$$

where $R$ is the radius of the colloids. The Derjaguin approximation is valid as long as $D \ll R$, which holds for our colloidal system. Finally, the relation between the surface charge density $\sigma$ and the colloidal charge $Z$ is given through

$$\sigma = \frac{eZ}{4\pi R^2}.$$  

Following the experimental work, $T=298$ K and $\varepsilon_r = 78.3$ are used throughout in the model calculations.

**IV. RESULTS AND DISCUSSION**

The experimental force-separation dependences in aqueous salt solution with monovalent (NaCl), divalent (CaCl$_2$), or trivalent (LaCl$_3$) counter-ions at varying concentration are displayed in Fig. 2 (symbols). At sufficiently long separation the force is essentially zero, whereas at decreasing separation the force is monotonously repulsive. The onset of the repulsion appears at shorter separation (i) at increasing salt concentration for a given counter-ion and (ii) at increasing counter-ion valence at a given salt concentration.

Using the DLVO theory we have made global fits of the colloidal charge $Z$ and radius $R$ for three different pairs of colloids using all force curves for a given type of salt by minimizing $\sigma^2 = \left(\frac{1}{N}\sum_{i=1}^{N} \left(F_{\text{calc}}^{i} - F_{\text{exp}}^{i}\right)^2\right)^{1/2}$, where $N$ is the number of data points, $F_{\text{calc}}^{i}$ the calculated force using Eq. (4), and $F_{\text{exp}}^{i}$ the experimental force. In the case of trivalent counter-ions, the influence of CO$_2$ in air has to be accounted for by assuming 34 $\mu$M HCO$_3^-$.

<table>
<thead>
<tr>
<th>Colloid pair</th>
<th>Salt</th>
<th>No. of data points</th>
<th>$Z$/1000</th>
<th>$R$ [nm]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>NaCl</td>
<td>245</td>
<td>218 [156, 267]</td>
<td>1134.6 [1133.0, 1135.4]</td>
</tr>
<tr>
<td></td>
<td>CaCl$_2$</td>
<td>108</td>
<td>137 [110, 161]</td>
<td>1131.9 [1130.5, 1132.8]</td>
</tr>
<tr>
<td></td>
<td>LaCl$_3$</td>
<td>160</td>
<td>69 [58.9, 79.4]</td>
<td>1135.4 [1131.5, 1138.0]</td>
</tr>
<tr>
<td>2</td>
<td>NaCl</td>
<td>266</td>
<td>441 [318, 536]</td>
<td>1135.9 [1133.9, 1136.8]</td>
</tr>
<tr>
<td></td>
<td>CaCl$_2$</td>
<td>180</td>
<td>257 [156, 326]</td>
<td>1135.7 [1132.9, 1136.8]</td>
</tr>
<tr>
<td></td>
<td>LaCl$_3$</td>
<td>231</td>
<td>66 [42, 82]</td>
<td>1136.9 [1132.6, 1145.0]</td>
</tr>
<tr>
<td>3</td>
<td>NaCl</td>
<td>231</td>
<td>283 [203, 346]</td>
<td>1137.1 [1129.0, 1143.3]</td>
</tr>
<tr>
<td></td>
<td>CaCl$_2$</td>
<td>174</td>
<td>220 [163, 265]</td>
<td>1138.2 [1136.3, 1139.2]</td>
</tr>
<tr>
<td></td>
<td>LaCl$_3$</td>
<td>76</td>
<td>56 [49, 63]</td>
<td>1139.0 [1136.0, 1141.2]</td>
</tr>
</tbody>
</table>
forces between single pairs of charged colloids in aqueous solution of varying salt and monovalent counterions at $c_{\text{salt}}=0.15, 0.3, 0.5, 1, 2, 4, 10,$ and $30 \text{ mM}$ (right to left) with $Z=218\,000$ and $R=1134.6 \text{ nm}$ and (b) divalent counterions at $c_{\text{salt}}=0.15, 0.3, 0.5, 1, 1.5,$ and $3 \text{ mM}$ (right to left) with $Z=137\,000$ and $R=1131.9 \text{ nm}$ using the DLVO theory (solid curves) and the PB theory (dotted curves).

Now, (i) the reasonable agreement between the DLVO theory and PB approaches and (ii) the accumulated experience that the small-ion correlations neglected in both approaches are of minor importance in aqueous solutions with monovalent small ions [39] imply that, except for the lowest salt concentrations, the DLVO theory satisfactorily represents the electrostatic interaction between the colloids in the presence of monovalent salt. Second, a comparison of the fitted $Z$ and $R$ among the three colloidal pairs with monovalent counterions shows that the radii are virtually the same, whereas the fitted charges ranges from $Z=200\,000$ to $450\,000$. Third, for a given pair of colloids, at increasing counter-ion valence the fitted radius $R$ remains essentially constant, whereas the fitted charge $Z$ decreases (Table I). We attribute the latter effect to the neglect of the small-ion correlations in the DLVO theory, which gain importance at increasing counterion valence. The neglect of those leads to an underestimation of screening of the colloids, while the functional force-separation dependence is essentially preserved.

Finally, Fig. 5 displays the surface separation versus the (a) NaCl, (b) CaCl$_2$, and (c) LaCl$_3$ concentrations at the specified forces 2 and 4 pN for the three different pairs of colloids as obtained from the experiments and the predictions of the DLVO theory. A similar degree of agreement between the experimental (symbols) and the calculated (lines) as for the force versus distance curves are obtained.

V. CONCLUSION

To conclude, the force-separation dependence for single pairs of charged colloids in aqueous solution of monovalent, divalent, or trivalent counterions of varying concentration has been measured using optical tweezers. The results are quantitatively described by the DLVO theory using
concentration-independent effective colloidal charge and stochiometric Debye screening lengths. However, the resulting effective colloidal charge decreases with increasing counterion valence, which is attributed to the approximations of the DLVO theory.

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**APPENDIX**

Starting from the PB equation for planar geometry, we will (i) briefly account for the derivation of the electrostatic force according to the DLVO theory and (ii) relate this force with the size-corrected screened Coulomb force [Eq. (4)]. The first part is given in more detail in textbooks; see, e.g., Refs. [37,41,42].

Equation (9) provides the net force between two charged planes immersed in an electrolyte solution. Insertion of Eq. (6) into Eq. (9) and the assumptions of weakly interacting colloids, enabling an expansion of the exponential in Eq. (6) to the second order, give

$$\frac{F_{\text{plane}}}{A} = \frac{1}{2} \varepsilon_{0} \varepsilon_{r} \kappa^{2} \left[ \phi(b) \right]^{2}. \quad (A1)$$

The next step is to apply the superposition approximation where the electrostatic potential at the mid-plane $\phi(b)$ is approximated according to

$$\phi(b) \approx 2 \phi^{\text{GC}}(b), \quad (A2)$$

where $\phi^{\text{GC}}(b)$ denotes the electrostatic potential appearing at the distance $b$ outside a single charged surface in contact with a semi-infinite electrolyte solution. The latter system possesses an analytic solution for symmetric electrolytes (also referred to as the Gouy-Chapman theory). For a $z \varepsilon z$ electrolyte and sufficiently far from the charged surface such that the electrostatic potential is small, we have

$$\phi^{\text{GC}}(b) = \frac{4kT}{\varepsilon z} \Gamma_{0} \exp(-\kappa b), \quad (A3)$$

where here and in the following $z$ denotes the valence of the $z \varepsilon z$ electrolyte and $\Gamma_{0}$ is related to the surface charge density $\sigma$ through the surface potential $\phi_{0}$ according to

$$\Gamma_{0} = \tanh \left( \frac{\varepsilon z \phi_{0}}{4kT} \right) \quad (A4)$$

and

$$\sigma = (8kTc_{\text{bulk}}\varepsilon_{0}\varepsilon_{r})^{1/2} \sinh \left( \frac{\varepsilon z \phi_{0}}{2kT} \right). \quad (A5)$$

Substitution of Eq. (A3) into Eq. (A2) and Eq. (A2) into Eq. (A1) gives

$$\frac{F_{\text{plane}}(D)}{A} = 32\varepsilon_{0}\varepsilon_{r}\kappa^{2} \left( \frac{kT}{\varepsilon z} \right)^{2} \Gamma_{0}^{2} \exp(-\kappa D)$$

$$= 64kTc_{\text{bulk}}\Gamma_{0}^{2} \exp(-\kappa D), \quad (A6)$$

where Eq. (2) has been used. Finally, after integration with respect to $D$ and applying the Derjaguin approximation, we have
\[ F(D) = 64\pi R k T c_{\text{bulk}} \Gamma_0^2 \exp(-\kappa D)/\kappa, \quad (A7) \]

which is the electrostatic force acting between two like-charged colloids of the original DLVO theory [21,22,37,41,42].

Our aim is now to examine under conditions at which Eqs. (4) and (A7) are identical. Assume now that the colloids are weakly charged such that Eqs. (A4) and (A5) can be linearized. Thereafter, an elimination of the surface potential \( \phi_0 \) gives the relation

\[ \Gamma_0 = \frac{\sigma}{(32 k T c_{\text{bulk}} e_0 e_r)^{3/2}}, \quad (A8) \]

which inserted into Eq. (A7) gives

\[ F(D) = 2\pi R \frac{\sigma^2}{e_0 e_r \kappa} \exp(-\kappa D)\kappa. \quad (A9) \]

The replacement of the surface charge density \( \sigma \) with the colloid charge \( Z \) using Eq. (12) results in

\[ F(D) = \frac{1}{24\pi e_0 e_r} \frac{(eZ)^2}{\kappa R^3} \exp(-\kappa D). \quad (A10) \]

Finally, when the colloid separation is much smaller than their size \( D \ll R \) and the Debye screening length is much smaller than the colloids \( \kappa^{-1} \ll R \), Eqs. (4) and (A7) are equivalent. At the present experimental conditions, this is practically the case.

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