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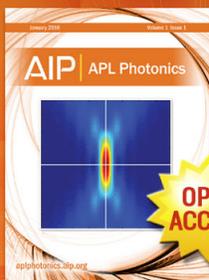
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Perspective: Coulomb fluids—Weak coupling, strong coupling, in between and beyond

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We present a personal view on the current state of statistical mechanics of Coulomb fluids with special emphasis on the interactions between macromolecular surfaces, concentrating on the *weak* and the *strong* coupling limits. Both are introduced for a (primitive) counterion-only system in the presence of macroscopic, uniformly charged boundaries, where they can be derived systematically. Later we show how this formalism can be generalized to the cases with additional characteristic length scales that introduce new coupling parameters into the problem. These cases most notably include asymmetric ionic mixtures with mono- and multivalent ions that couple differently to charged surfaces, ions with internal charge (multipolar) structure and finite static polarizability, where weak and strong coupling limits can be constructed by analogy with the counterion-only case and lead to important new insights into their properties that cannot be derived by any other means. © 2013 AIP Publishing LLC. [<http://dx.doi.org/10.1063/1.4824681>]

I. INTRODUCTION

A *Coulomb fluid* refers specifically to any mobile (thermalized) collection of charges, which interact via Coulomb interactions. These charges may be small ions (such as Na⁺ and Ca²⁺ with sizes around 0.1 nm), ions with internal structure (such as the rod-like spermine and spermidine ions with sizes around a nanometer) as well as charged nano-particles and macromolecules (such as proteins, colloids, polymers, and membranes), which are typically immersed in an aqueous solvent, as is the case in soft- and bio-matter context.^{1–8} Macromolecular surfaces get charged mainly due to the dissociation of their surface chemical groups in the solvent; this process releases small ions (referred to as co- and counterions depending on the sign of their charge with respect to that of the surface) into the solution. The type, magnitude, and the particular distribution of surface charges and those of the resulting mobile ions depend also on the specific chemistry of the surface as well as a number of other key factors including, among other things, the pH, temperature, dielectric properties or, in general, the specific molecular properties of the solvent. Once dissolved, these mobile charges mediate the interactions between macromolecular surfaces.

The presence of Coulomb fluids can significantly alter electrostatic interactions of charged (bounding) surfaces and modify the behavior of charged macromolecular dispersions.^{1–6} The role of Coulomb fluids can be so drastic that, as evidenced by numerous experimental and theoretical investigations over the last several years,^{9–123} it can challenge our understanding of electrostatic effects as demonstrated by non-conventional phenomena such as electrostatic attraction between like-charged surfaces, especially when multi-

valent ions (or counterions) are present in the system. Like-charge attraction is manifested in many experimental examples; a few notable cases include formation of large aggregates of like-charged polymers such as microtubules¹¹⁵ and F-actin^{18,21} as well as formation of large condensates of DNA in the bulk^{12,116–118} and in the DNA packaging inside viral shells that are observed in the presence of multivalent cations.^{119–123} Similarly, numerous numerical simulations have already demonstrated the emergence of like-charge attraction and investigated its underlying mechanism in many examples including charged membranes, colloids, and polymers.^{23–61}

All major theoretical proposals that aim to explain the phenomenon of like-charge attraction go beyond the standard mean-field or Poisson-Boltzmann (PB) theories, which have been studied since the early years of the last century,^{1–3} by including the effects of electrostatic fluctuations and correlations that are neglected in the description of Coulomb fluids on the mean-field level. These proposals include integral-equation methods (see, e.g., Refs. 29, 62, and 63), perturbative improvement of the mean-field theory including loop expansions and other Gaussian-fluctuations approximations (see, e.g., Refs. 64–78), variational methods (see, e.g., Refs. 79–82), and local density functional theory (see, e.g., Refs. 83 and 84). These approaches turn out to be applicable mostly at large separations between charged surfaces or, generally, for relatively small coupling (and/or electrostatic correlation) strengths. This regime is known as the weak-coupling (WC) or “high-temperature” regime. A complementary strong-coupling (SC) or “low-temperature” approach was pioneered by Rouzina and Bloomfield⁸⁶ based on the observation that counterions tend to form two-dimensional strongly

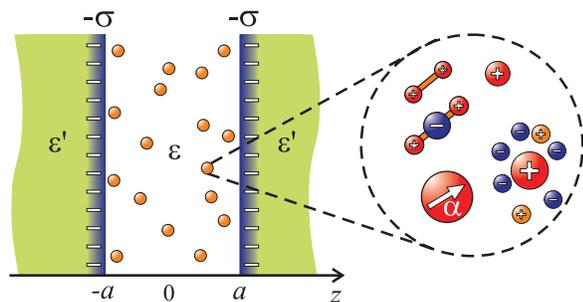


FIG. 1. Schematic representation of some of possible generalizations of the primitive “counterion-only” model as considered in this paper. It shows a system of large (mobile) multivalent ions (shown by red spheres), which may possess an internal structure (e.g., a rod-like shape or a static polarizability) and may be dispersed in a bathing solution of monovalent anions and cations in the background (shown by blue and orange spheres, respectively).

correlated layers at oppositely charged bounding surfaces when the coupling parameter is large. Such structural correlations can lead to dominant attractive forces of mainly energetic origin between like-charged surfaces that have been studied by means of several different theoretical approaches in recent years (see, e.g., Refs. 45–52, 58–60, and 88–100).

For a Coulomb fluid consisting of only a single charge species next to oppositely charged boundaries (the so-called “counterion-only” model), the WC and SC coupling theories were shown to follow systematically as two limiting laws of a single unified formalism,⁹⁴ a view that was completely corroborated by extensive numerical simulations (see, e.g., Refs. 42–59) or available exact solutions.^{124,125}

In what follows, we provide a guided personalized tour of recent advances in the theory of Coulomb fluids by reviewing various aspects of the WC-SC paradigm within the primitive counterion-only model. We then discuss how this framework can be generalized to derive powerful limiting laws for more complicated, but also more realistic models of Coulomb fluids with additional characteristic length scales (coupling parameters) than envisioned in the original framework of the WC-SC dichotomy; most notably, asymmetric ionic mixtures with mono- and multivalent ions, ions with internal multipolar structure, and finite static polarizability next to charged bounding surfaces (see Fig. 1).

II. COULOMB FLUIDS: GENERAL CONSIDERATIONS

A. The primitive model

The stated features of Coulomb fluids make a full understanding of the equilibrium properties of charged macromolecular systems quite difficult and many aspects of real systems have been either neglected (depending on the particular case or the application under investigation) or heavily approximated, often though with reasonable justification. Perhaps the most simple and yet efficient idealization is the so-called *primitive model* where solvent is treated as a featureless continuum of a fixed dielectric constant ϵ stemming from the degrees of freedom associated with the solvent molecules; small mobile ions are taken to be featureless and characterized only by their charge, without any internal structure and/or

polarizability. Furthermore, macromolecules or other macroscopic surfaces are often treated as objects with a fixed uniform charge distribution (see Fig. 1).

A great deal of theoretical effort has been devoted recently to improve upon these simplifying assumptions, in particular regarding the solvent structure (see, e.g., Refs. 1, 6, 126–130, and references therein) and the polarizability and internal structure of mobile ions as conceived by Debye¹³¹ (see, e.g., Refs. 132–137 and references therein), etc. Unfortunately, many of these interesting developments such as ionic liquids,^{138–140} surface ion-adsorption effects,^{141,142} the discreteness, and/or heterogeneity of the surface charge distribution (see, e.g., Refs. 143–156 and reference therein), pH-controlled charge regulation,^{157–161} many-body interactions,^{162,163} or Bjerrum pair formation^{164–166} must remain outside the domain of this review. Even on the level of the above simplified model assumptions, the underlying physics of the Coulomb fluids exhibits conceptual challenges and intriguing results, which have only recently been corroborated by computer simulations and, less often, experimental evidence.

B. Physical scales and parameters

Of the plethora of possible length scales (associated with electric charges, van der Waals (vdW) or chemical bonding energies, ion sizes, solvent molecular size, or spacing between discrete surface charges, etc., see Ref. 7), one can argue that within the primitive model discussed above only a few will be important.

The thermal energy $k_B T = \beta^{-1}$ can be compared with inter-ionic Coulomb interactions (e.g., for two elementary charges e_0), giving rise to the Bjerrum length $\ell_B = e_0^2 / (4\pi\epsilon\epsilon_0 k_B T)$, with gas-like behavior for small and liquid- or solid-like behavior for large ℓ_B . For an ionic fluid consisting of ions (counterions) of charge valency q at a charged interface of uniform surface charge density $-\sigma e_0$, the Gouy-Chapman length $\mu = 1 / (2\pi q \ell_B |\sigma|)$ determines the strength of the thermal energy with respect to the electrostatic interaction (attraction) with the surface (we assume that $q > 0$ and $-\sigma < 0$ with no loss of generality). The neutralizing counterions next to a charged surface are thus expected to form a diffuse gas-like phase when the *electrostatic coupling parameter*, defined as the ratio

$$\Xi = q^2 \ell_B / \mu = 2\pi q^3 \ell_B^2 |\sigma|, \quad (1)$$

is small, i.e., $\Xi \ll 1$. In the opposite situation with large $\Xi \gg 1$, one observes a very different behavior,^{44–46,50–52,90–95,99,100} see Fig. 2, where the ionic cloud is reduced from a three-dimensional (3D) layer to a quasi-two-dimensional (2D) sheet. For $\Xi \gg 1$, the mean separation between counterions, which follows from the local electroneutrality condition as $a_\perp \sim \sqrt{q/|\sigma|} \sim \mu\sqrt{\Xi}$, is much bigger than the thickness of the surface (counterion) layer given by μ . Conjointly with the surface-counterion interaction, the inter-counterion repulsions, $q^2 \ell_B / a_\perp \sim \sqrt{\Xi}$, also dominate the thermal energy. Therefore, increasing Ξ engenders a strongly correlated liquid, or even a crystalline phase of counterions, in the limit $\Xi \rightarrow \infty$.

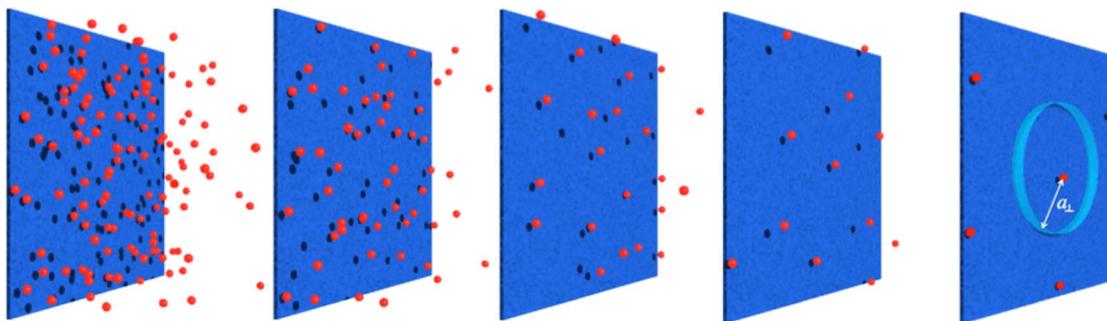


FIG. 2. An extended 3D counterion layer in the WC regime, $\Xi \ll 1$, reduces to a quasi-2D layer in the SC regime, $\Xi \gg 1$, upon increasing the coupling parameter (from left to right). In the limit of $\Xi \rightarrow \infty$, a single-particle description becomes relevant because of large correlation holes, $a_{\perp} \rightarrow \infty$, forming around individual counterions.

This picture follows from extensive Monte-Carlo (MC) simulations,^{44–47,50,51} which show that the *crossover* from the *WC regime* ($\Xi \ll 1$) to the *SC regime* ($\Xi \gg 1$) is associated with a hump in the heat capacity of the system and development of short-range correlations between counterions in the range $10 < \Xi < 100$. The transition to a Wigner crystalline phase (characterized by a diverging heat capacity) then occurs at a very large value of the coupling parameter around $\Xi \simeq 3 \times 10^4$.^{8,44–46}

The WC-SC paradigm was derived from a functional-integral (field-theoretic) representation of the partition function directly and systematically only in the case of the primitive counterion-only model.⁹⁴ The presence of additional mobile charge components, surface charge heterogeneity or mobile ion multipolar structure, and/or polarizability introduces new length scales and thus new coupling parameters into the problem. Nevertheless, one can always identify a principal coupling parameter, in analogy to the counterion-only case, that implies a SC-like fixed point. The thusly defined SC fixed point then displays a fine structure and can bifurcate into other strongly coupled states governed by these additional length scales and the ensuing coupling parameters. In what follows, we will discuss a few illuminating examples (see Fig. 1) displaying the salient features of this approach and its main results.

III. FIELD THEORY AND THE WC-SC PARADIGM IN THE COUNTERION-ONLY MODEL

The preceding observations suggest that the WC and the SC regimes should follow as complementary descriptions from a single, unified formalism. This clearly transpires from the field-theoretic approach to Coulomb fluids,^{66,67,78,94,167} wherein the WC and the SC limits can indeed be derived as *asymptotic* theories from a single *ansatz* for $\Xi \rightarrow 0$ and $\Xi \rightarrow \infty$, respectively.⁹⁴

In general, the Hubbard-Stratonovich transformation allows the partition function of the primitive model to be mapped *exactly* to a functional integral over a fluctuating (electrostatic) potential, $\phi(\mathbf{r})$,^{66,67,78,94,167} as

$$\mathcal{Z} = \int \mathcal{D}\phi e^{-\beta S[\phi]}, \quad (2)$$

where the effective “field-action” can be written (up to a “self-energy” prefactor, which we ignore here, see, e.g.,

Refs. 53–55, 66, 67, 78, and 94) as

$$S[\phi] = \frac{1}{2} \iint d\mathbf{r} d\mathbf{r}' \phi(\mathbf{r}) u^{-1}(\mathbf{r}, \mathbf{r}') \phi(\mathbf{r}') + i \int d\mathbf{r} \rho_0(\mathbf{r}) \phi(\mathbf{r}) - \int d\mathbf{r} \mathcal{U}(\phi(\mathbf{r})) = S_0[\phi(\mathbf{r})] - \int d\mathbf{r} \mathcal{U}(\phi(\mathbf{r})), \quad (3)$$

where the Coulomb interaction is $u(\mathbf{r}, \mathbf{r}') = 1/(4\pi\epsilon\epsilon_0 |\mathbf{r} - \mathbf{r}'|)$ and its kernel (operator inverse) $u^{-1}(\mathbf{r}, \mathbf{r}') = -\epsilon_0 \nabla \cdot \epsilon(\mathbf{r}) \nabla \delta(\mathbf{r} - \mathbf{r}')$. The fixed external (macromolecular) charge distribution is described by $\rho_0(\mathbf{r})$ and is assumed, for further specification, to be distributed on planar surfaces with either one plate (placed at $z = 0$) or two plane-parallel surfaces (placed at $z = -a$ and $z = +a$, i.e., at separation distance $D = 2a$ along the z axis) with a uniform surface charge density of $-\sigma e_0$. In addition, the system can exhibit a non-trivial spatial dielectric constant profile $\epsilon(\mathbf{r})$. The model of the Coulomb fluid is embodied in the generally non-linear “field self-interaction” term $\mathcal{U}(\phi(\mathbf{r}))$. The presence of these nonlinear field self-interactions renders an exact evaluation of the partition function difficult, except in a few special cases (for such exact solutions, see, e.g., Refs. 124 and 125 and references therein). Nonetheless, remarkable analytical progress has been made in analyzing the behavior of such systems, especially in the cases that go beyond the usual WC-SC framework (see, e.g., Refs. 53–55, 94, 133, and 134).

In the *counterion-only model*, one assumes that only counterions of charge valency q are present in the system and exactly neutralize the fixed charges. Although this model may be inapplicable in most real situations, it has nevertheless served as a useful paradigm, elucidating fundamental aspects of the WC-SC dichotomy. In this context, the field self-interaction is obtained in the grand-canonical ensemble as^{66,67,94}

$$\mathcal{U}(\phi(\mathbf{r})) = \lambda \Omega(\mathbf{r}) e^{-i\beta q e_0 \phi} \equiv \lambda \tilde{\mathcal{U}}(\phi(\mathbf{r})). \quad (4)$$

Here, λ is the fugacity of counterions and the “blip” function $\Omega(\mathbf{r})$ defines the region of space allowed to the mobile counterions (i.e., it is equal to one in the region where counterions are allowed to be present and zero elsewhere). The functional integral can be cast in a dimensionless form that depends on only one parameter, i.e., Ξ .⁹⁴ The WC-SC limits then follow directly from appropriate evaluations of the partition function.⁹⁴

A. WC theory: Mean-field theory and weak fluctuations

The WC limit, $\Xi \rightarrow 0$, for this model is obtained by solving the saddle-point equation of the field-action,^{66,67,94}

$$\left. \frac{\delta S[\phi]}{\delta \phi} \right|_{i\phi=\psi_{\text{PB}}} = 0, \quad (5)$$

which governs the mean (real-valued) electrostatic potential $\psi_{\text{PB}}(\mathbf{r}) = i(\phi(\mathbf{r}))$ and leads to the standard PB equation

$$-\varepsilon_0 \nabla \cdot [\varepsilon(\mathbf{r}) \nabla \psi_{\text{PB}}(\mathbf{r})] = \rho_0(\mathbf{r}) + qe_0 \lambda \Omega(\mathbf{r}) e^{-\beta qe_0 \psi_{\text{PB}}(\mathbf{r})}, \quad (6)$$

with Neumann boundary conditions at the charged interfaces. For one or two planar surfaces, the dielectric discontinuity at the bounding surfaces plays no role and the above equation leads to the standard Gouy-Chapman theory of electrical double layers¹⁻⁶ with $\psi_{\text{PB}}(\mathbf{r}) = \psi_{\text{PB}}(z)$. The corresponding expressions for the counterion density profile $n_{\text{PB}}(z)$ and the disjoining pressure P_{PB} imparted by the counterions on the bounding surfaces are well known,^{4,5} with the latter obtained from the contact-value theorem¹⁶⁸⁻¹⁷⁰ through the counterion density at the mid-plane $z = 0$. This leads to the following asymptotic form for large $D = 2a$:

$$P_{\text{PB}}(D) = k_B T n_{\text{PB}}(z; D) \Big|_{z=0} \simeq \frac{k_B T}{q^2 \ell_B D^2} \Big|_{D \rightarrow \infty}, \quad (7)$$

which clearly shows that the PB pressure at large separations is independent of σ , has an entropic origin, and is thus repulsive. Quite generally, it can be proven that the disjoining pressure within the PB theory is always repulsive, regardless of the shape of the charged surfaces as long as the boundary conditions are symmetric.¹⁷¹⁻¹⁷³

The PB equation is exact in the strict limit of $\Xi \rightarrow 0$ and has been applied successfully to study weakly charged systems.¹⁻⁵ When Ξ is finite but small, one expects subdominant Gaussian fluctuations to occur around the PB solution, $\phi = -i\psi_{\text{PB}} + \varphi$. These fluctuations are described by the field-action

$$S[\phi] \simeq S[-i\psi_{\text{PB}}] + \frac{1}{2} \int d\mathbf{r} d\mathbf{r}' \varphi(\mathbf{r}) \varphi(\mathbf{r}') \left. \frac{\delta^2 S[\phi]}{\delta \phi(\mathbf{r}) \delta \phi(\mathbf{r}')} \right|_{\phi=-i\psi_{\text{PB}}} + \mathcal{O}(\varphi^3), \quad (8)$$

and lead to small deviations from the PB pressure. The total WC pressure can then be written as

$$P_{\text{WC}}(D) = P_{\text{PB}}(D) + \Xi P_1(D) + \mathcal{O}(\Xi^2), \quad (9)$$

where P_1 is the Gaussian (one-loop) correction around the mean-field solution, analogous to vdW-type interactions^{66,67,78,94,174} and is thus attractive, $P_1 < 0$. For two like-charged surfaces,

$$\Xi P_1(D) \sim -k_B T \left. \frac{\ln D}{D^3} \right|_{D \rightarrow \infty}. \quad (10)$$

which has a similar algebraic dependence on D as in the case of the classical vdW force between neutral dielectrics, generated by the thermal fluctuations of the zero-frequency Mat-

subara modes of the electromagnetic field,¹⁷⁴ in the present case, however, the attractive correction is caused by the Gaussian fluctuations around a non-uniform background defined by the PB solution that produces also a logarithmic D -dependent correction. It should be noted that in the WC regime the fluctuation-induced attraction cannot overcome the repulsive leading-order PB pressure, the total pressure thus remaining repulsive.

In certain models that assume surface condensation or adsorption of counterions on (fixed) charged boundaries,⁷⁰⁻⁷⁷ the repulsive mean-field effects are strongly suppressed and the total pressure can eventually turn out to be attractive. Such condensation behavior cannot occur within the primitive model with purely Coulombic interactions and may occur only if one takes into account non-electrostatic surface effects going beyond the models considered here.

B. SC theory

As shown by Netz,⁹⁴ a systematic description for the SC limit follows from the same field-theoretic formalism that yields the PB equation in the limit $\Xi \rightarrow 0$, provided that one takes the limit of $\Xi \rightarrow \infty$. In this case, one can perform a fugacity (virial) expansion combined with a $1/\Xi$ expansion, the leading order of which was shown to be finite and given by single-particle contributions only. Physically, since in this limit the counterions are strongly attracted to the surface and are isolated in large correlation holes (of size $\sim a_{\perp}$) surrounding them, the partition function of the system is expected to be dominated by single-particle contributions from the interaction between counterions and charged surfaces. To the lowest order, one finds

$$\mathcal{Z}_{\text{SC}} = \mathcal{Z}_{\text{SC}}^{(0)} + \lambda \mathcal{Z}_{\text{SC}}^{(1)} + \mathcal{O}(\lambda^2), \quad (11)$$

where the first term involves only $S_0[\phi(\mathbf{r})]$ of (3) due to external charges, while

$$\mathcal{Z}_{\text{SC}}^{(1)} / \mathcal{Z}_{\text{SC}}^{(0)} = \int d\mathbf{r} \Omega(\mathbf{r}) e^{-\beta e_0 q \int d\mathbf{r}' u(\mathbf{r}, \mathbf{r}') \rho_0(\mathbf{r}')}, \quad (12)$$

is the single-ion partition function in the field of external charges (note the grand-canonical formalism, which can be transformed back to a canonical description involving N particles by a Legendre transform;⁹⁴ the number of particles in this case is fixed by the electroneutrality condition). As discussed in Sec. II B, counterion-counterion interactions are also very strong in the limit $\Xi \rightarrow \infty$, but appear as subleading if compared with counterion-surface contributions.

In the SC limit and for two like-charged bounding surfaces, the disjoining pressure can be expressed to the leading-order as⁹⁴

$$P_{\text{SC}}(D) = -\frac{\sigma^2}{2\varepsilon\varepsilon_0} + \left(\frac{2|\sigma|}{q} \right) \frac{k_B T}{D}. \quad (13)$$

It is easy to see that the first term follows from the electrostatic energy of individual counterions sandwiched between two equally charged surfaces, leading to an attractive pressure, while the repulsive second term is clearly nothing but the ideal-gas entropy of the counterions confined to the slit between the surfaces. In fact, as noted above in the SC limit,

the counterions are isolated in correlation holes of size a_{\perp} so that $|\sigma|a_{\perp}^2 \sim q$, and hence the energetic repulsion between the two like-charged surfaces is overcompensated by the attraction between them and the individual counterion in between in such a way that it changes the sign of the pressure!^{50,51,94}

The above result holds for small separations $D < a_{\perp}$ or equivalently, $D/\mu < \sqrt{\Xi}$. Within the regime of validity of the SC limit, one recovers both the repulsive regime for $D < D_*$ and the attractive regime for $D > D_*$, where $D_* = 2\mu$ is the separation at which the total force acting on the surfaces vanishes, as confirmed by MC simulations^{44–46} (this argument needs to be amended in the case where dielectric inhomogeneities are present in the system, see, e.g., Refs. 44–46, 54, and 55 and references therein).

C. Crossover from WC to SC limit

For separation distances larger than the correlation hole size or at smaller coupling parameters, one needs to account for the higher order terms in the SC expansion. Even though the single-particle SC limit appears to work very well for coupling parameters down to $\Xi \sim 100$ (corresponding, e.g., to DNA with trivalent counterions), the perturbative expansion turns out to be very inefficient and can hardly be extended into the so-called *crossover regime*, i.e., $10 < \Xi < 100$.^{50,51} This regime remains accessible *fully* only through simulations. Despite the challenging nature of correlations in this regime, Santangelo,⁹⁷ Weeks *et al.*,^{58,59} and others^{60,98} have shown that approximations based on the *decomposition method* used within the liquid-state theories can be applied with reasonable success. Also a test-charge theory developed by Burak *et al.*⁹⁶ seems to capture the onset of correlations and provide very useful insights into the crossover behavior of the counterion-only system.

Another approach to study counterion-only systems in the limit of large couplings has been proposed by considering the ground-state structure of the system in the limit $T \rightarrow 0$.^{86,91} It should be noted that this latter limit does not in general coincide with the SC limit, which is based on a combined virial and $1/\Xi$ expansion. The virial approach leads to finite entropic (temperature) corrections at the leading order (e.g., the second term in Eq. (13)) and deviations have been reported for the subleading corrections calculated from ground-state considerations and those obtained from the virial approach.^{99,100}

We should also note that *modified PB equations* are sometimes invoked as *ad hoc* generalizations of the standard PB theory that include the ion excluded-volume term and the so-called fluctuation potential.^{175,176} While these kinds of theories certainly describe some features of steric and correlation interactions inaccessible to the standard PB theory, they cannot be viewed as a limiting law in the sense that it follows systematically from a single unified formalism valid in both the weak and the strong coupling limit.

IV. GENERALIZATIONS OF THE WC-SC PARADIGM

In the original form,^{46,50,51,94} the WC and SC dichotomy applies only to a counterion-only system confined by uni-

formly charged surfaces, a situation of profound theoretical significance but seldom encountered in the real world. While the WC limit, through its correspondence with the saddle-point equation, can be defined for any field-action, the existence of additional length scales, besides the Bjerrum and the Gouy–Chapman ones, precludes a direct introduction of a unique electrostatic coupling parameter and consequently a systematic derivation of the SC limit with its virial expansion.

In this case, however, one can generalize not the SC theory itself, but the single-particle aspect of the SC limit, which can be then applied to any system that contains highly charged species, irrespective of the number of governing length scales (see, e.g., Refs. 42, 43, 48, 49, 53, 54, 103, 133, 134, and 149). The approach is reasonable because the highly charged ions are usually present only in small concentrations and thus a virial expansion, of which the single-particle limit is the lowest order, makes sense. In addition, the single-particle SC limit has been tested against detailed simulations and its regime of applicability is determined for any particular case.

In quite general context, for any field-action, one could thus derive a saddle-point theory as a substitute for the WC limit and a single-particle theory as a substitute for the SC limit. This is a powerful consequence of the field-theoretic approach and can be applied even to the cases where other approaches (such as those based on the ground-state methods^{99,100}) are not necessarily applicable (e.g., asymmetric ionic mixtures containing large amounts of monovalent salt). This is what we will illustrate for a few interesting cases detailed below.

A. Asymmetric ionic mixtures: Dressed multivalent-ion approach

A particularly relevant case is that of a mixture of *multivalent* ions in a bathing solution of *monovalent* ions (Figs. 3(a) and 4). This is a typical situation in the formation of liquid crystalline mesophases of semiflexible biopolymers,^{21,177,178} multivalent-ion driven condensation of DNA in the bulk,^{12,116–120} or in viruses, where multivalent ions are believed to play a key role in the stability of the viral capsid and/or packaging of its genome.^{121–123}

For an ionic mixture consisting of a single species of multivalent counterions (c) in a neutralizing background of monovalent anions ($-$) and cations ($+$), we have

$$\mathcal{U}(\phi(\mathbf{r})) = \Omega(\mathbf{r})(\lambda_c e^{-i\beta q e_0 \phi} + \lambda_+ e^{-i\beta e_0 \phi} + \lambda_- e^{i\beta e_0 \phi}), \quad (14)$$

which follows directly from the Hubbard–Stratonovich transformation of the microscopic (Coulomb) Hamiltonian of the system.^{53,66,67,94} Here, λ_c and λ_{\pm} are the respective ionic fugacities, and the “blip” function $\Omega(\mathbf{r})$ is assumed to be the same for all mobile species.

Obviously, in an asymmetric mixture, multivalent counterions and monovalent ions are coupled to macromolecular charges quite differently: multivalents *strongly*, while monovalents only *weakly*, as evident from their respective electrostatic coupling parameters. This presents a challenging problem as no single limit, neither WC nor SC will apply to all of the components of the system.⁵³ The saving grace in this situation is the fact that usually multivalent ions are

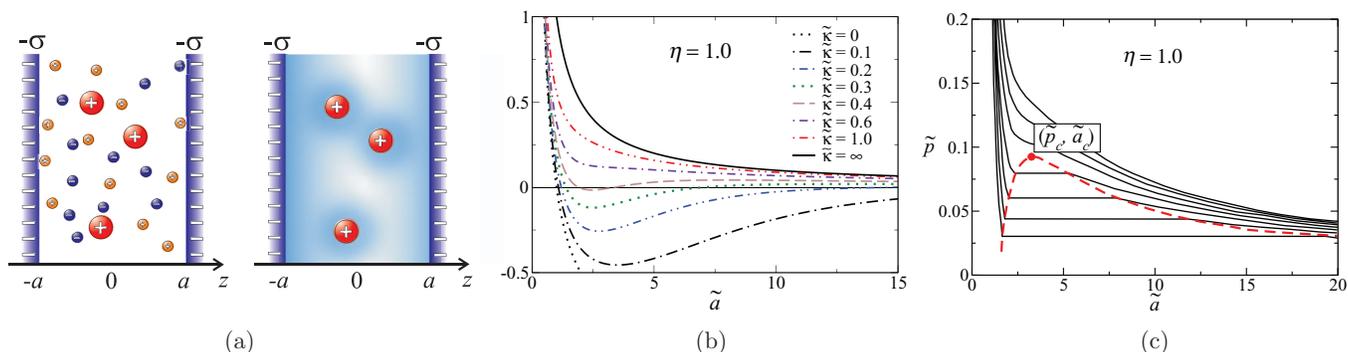


FIG. 3. (a) A mixture of multivalent counterions (red spheres) in a bathing solution of monovalent salt ions (blue and orange spheres); the former are strongly and the latter are weakly coupled to external (surface) charges. The monovalent ions can be integrated out from the explicit microscopic model (left) in the case of a highly asymmetric mixture (with highly charged multivalent ions), giving rise to the effective dressed multivalent-ion model (right). (b) The rescaled interaction pressure, $\tilde{p} = \beta P_{SC}/2\pi\ell_B\sigma^2$, as predicted by the dressed multivalent-ion theory, Eq. (18), is shown as a function of the rescaled half-distance between the surfaces, $\tilde{a} = a/\mu$. It bridges between the pure osmotic regime of multivalent counterions (large κ) where the electrostatic interactions are screened out and the SC theory for a counterion-only system (small κ), which exhibits attractive interaction between the like-charged surfaces for a wide range of parameters (here shown for $\eta = 1$, i.e., when the total charge due to multivalent counterions exactly compensates the surface charges). (c) In the canonical dressed multivalent-ion ensemble, there appears a non-monotonic behavior for the interaction pressure showing vdW-like loops, suggesting a coexistence regime between two different phases. The phase coexistence region actually ends with a critical point (p_c, a_c) . Adapted from Ref. 53.

present at very small concentrations, e.g., around just a few mM (see, e.g., Refs. 116, 119, 120, 177, and 178), and thus their behavior is expected to fit naturally within the virial scheme based on an expansion in terms of their fugacity (bulk concentration).

Additionally, in highly asymmetric solutions with $q \gg 1$ (e.g., with tri- and tetravalent counterions), the problem can be furthermore greatly simplified by employing the following approximation:⁵³

$$\mathcal{U}(\phi(\mathbf{r})) \simeq \lambda_c e^{-i\beta q e_0 \phi} - n_b (\beta e_0 \phi)^2 / 2 + \mathcal{O}(\phi^3) \quad (15)$$

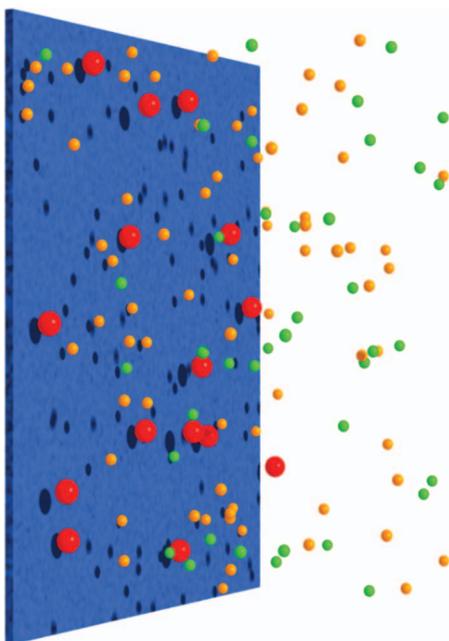


FIG. 4. Snapshot of an explicit MC simulation of a single (negatively) charged surface with (positive) tetravalent counterions (red spheres) and monovalent salt anions and cations (green and orange spheres); while the former are strongly attracted toward the surface, the latter form an extended DH-like atmosphere. (In this case, there is also a dielectric discontinuity at the charged surface corresponding to that of the hydrocarbon/water interface.)

(in the region where the ions are present $\Omega(\mathbf{r}) = 1$), where $n_b = 2n_0 + qc_0$, assuming that the multivalent counterions result from a $q:1$ salt of bulk concentration c_0 , which is mixed with a 1:1 salt of bulk concentration n_0 ; hence, $\lambda_c = c_0$, $\lambda_+ = n_0$, and $\lambda_- = n_0 + qc_0$ (note that the multivalent salt contributes an additional amount of monovalent ions with concentration qc_0).

From this form, it is clear that the degrees of freedom due to monovalent ions can be integrated out, resulting in an effective formalism including only screened interactions between the remaining “dressed” multivalent ions and fixed macromolecular charges.^{53–55} This follows from the first term of the field-action $S_0[\phi(\mathbf{r})]$, Eq. (3), and the quadratic term in the above expansion. In other words, the effective field-action of the system is given by

$$S[\phi] \simeq S_{DH}[\phi] + \lambda_c \int d\mathbf{r} \Omega(\mathbf{r}) e^{-i\beta q e_0 \phi}, \quad (16)$$

where $S_{DH}[\phi]$ has the same form as $S_0[\phi]$ except that $u^{-1}(\mathbf{r}, \mathbf{r}')$ is replaced by the screened Debye-Hückel (DH) kernel $u_{DH}^{-1}(\mathbf{r}, \mathbf{r}') = -\epsilon_0[\nabla \cdot \epsilon(\mathbf{r})\nabla - \epsilon(\mathbf{r})\kappa^2]\delta(\mathbf{r} - \mathbf{r}')$, and the inverse Debye length is introduced as $\kappa^2 = 4\pi\ell_B n_b$. Obviously, the multivalent ion concentration c_0 introduces a new parameter and new physics that goes beyond the simple DH screening picture. The multivalent ion effects can be codified by a new length scale defined as $\chi^2 = 8\pi q^2 \ell_B c_0$,^{53–55} which is analogous to the Debye length in the case of monovalent salt.

We refer to the theory based on the field-action (16) as the *dressed multivalent-ion theory*. The key point here is thus that the above approach can be applied only to highly asymmetric ionic mixture with $q \gg 1$.⁵³ The regime of validity of this approach can be checked against explicit-ion MC simulations, where all ions, including the monovalent ones, are explicitly simulated (see Fig. 4), which show that this approach can indeed give quantitatively accurate results in a wide range of realistic parameter values.^{54,55} This may not seem obvious at first since the above approach treats the monovalent ions

on the implicit DH level and one might thus expect strong deviations to occur when multivalent ions are present in the solution due, e.g., to nonlinear charge renormalization and/or Bjerrum pairing effects,^{164–166} these effects, however, turn out to be absent or negligible in the regime of parameters that is of concern to our discussion.^{53–55}

On the saddle-point level of the dressed multivalent-ion field-action, the PB equation, Eq. (5), reads

$$-\varepsilon_0[\nabla \cdot \varepsilon(\mathbf{r})\nabla - \varepsilon(\mathbf{r})\kappa^2]\psi_{\text{PB}} = \rho_0 + qe_0c_0 \Omega(\mathbf{r}) e^{-\beta qe_0\psi_{\text{PB}}}. \quad (17)$$

The corresponding disjoining pressure is again consistently repulsive for symmetric boundary conditions. The underlying assumption here is that electrostatic correlations are zero and thus all ions remain weakly coupled to each other and the external fixed charges.

When $c_0 < n_0/q^2$, one can furthermore develop a simple theory based on a virial expansion, which can capture the behavior of such asymmetric solutions in the limit corresponding to a generalized SC limit, where multivalent counterions are coupled strongly (to each other and the external charges), while monovalent ions still remain weakly coupled to other charge species. By inspection of the field-action (16), the single-particle limit has exactly the same form as Eq. (11), except that the screened DH potential replaces the Coulomb interaction, i.e., $u(\mathbf{r}, \mathbf{r}') \rightarrow u_{\text{DH}}(\mathbf{r}, \mathbf{r}')$. For a single charged surface, the dressed multivalent-ion approach can successfully predict the density profile of ions next to the surface as well as the charge inversion induced by multivalent counterions.⁵⁴

For two apposed like-charged planar surfaces,⁵³ the interaction pressure behaves very differently if the dressed multivalent counterions are treated on the canonical (they are not in chemical equilibrium with a bulk reservoir) or grand-canonical level (in chemical equilibrium with the bulk).⁵³ The corresponding interaction pressure (within the canonical ensemble for dressed multivalent counterions and assuming, for the sake of simplicity, that the system is dielectrically homogeneous and that the monovalent salt ions are present both inside and outside the slit) is then transformed from Eq. (13) to

$$P_{\text{SC}}(D) = \frac{(e_0\sigma)^2}{2\varepsilon\varepsilon_0} [e^{-\kappa D} + 2\mu\eta I'(D)/I(D)], \quad (18)$$

where $\eta = (Nq)/2|\sigma|S$ is the amount of multivalent counterions in the slit between the surfaces relative to the total surface charge and $I(D = 2a) = \int_{-a}^a \exp((2/\kappa\mu)e^{-\kappa a} \cosh \kappa z) dz$. The case $\eta = 0$ represents a system with salt only and the DH theory is recovered, whereas $\eta = 1$ is the case when the total charge due to counterions exactly compensates the surface charges; the counterion-only SC theory discussed above can be recovered within this latter case by letting $\kappa \rightarrow 0$, see Fig. 3(b). This again illustrates that the dressed multivalent-ion theory can bridge between the standard WC and SC limits. For multivalent dressed counterions within the canonical description, the interaction pressure can even become non-monotonic, displaying repulsive branches at small and large inter-surface separations,⁵⁴ see Figs. 3(b) and 3(c). In fact,

for certain values of the parameters, the interaction pressure shows vdW-like loops, which could suggest a coexistence regime between two different phases, see Fig. 3(c). The binodal or the coexistence curve actually ends with a critical point corresponding to the largest amount of salt in the system that still leads to a non-monotonic dependence of the pressure on the inter-surface separation.

The dressed multivalent-ion approach can also give an insight into attractive image-induced depletion forces between two neutral dielectric walls.⁵⁵ These forces arise as a result of expulsion of multivalent ions from the vicinity of the walls in the slit region in the situation where the two dielectric walls have a dielectric constants smaller than that of water in the slit, as is typically the case. In this case, the dielectric image charges of individual ions have the same sign as the ions themselves and so are repelled from one another, pushing the mobile ions from the slit region back to the bulk solution. Subsequently, this effect leads to a lower osmotic pressure acting on the walls from the ionic solution in the slit, and hence eventually a net inter-surface attraction between the walls due to the bulk osmotic pressure that pushes the surfaces together. Such depletion-induced forces become important at small inter-surface separations, where they can even be comparable to or stronger than the usual vdW forces as confirmed also by explicit-ion simulations.⁵⁵ Similar attractive interactions can appear in the absence of dielectric discontinuities, albeit they are smaller; namely, in the close proximity of the walls, the multivalent ions are stripped off their “counterion clouds,” leading to an increase of the excess chemical potential, and thus a decrease of the ideal part (i.e., the density of multivalent ions).

The polarization depletion mechanism,^{179,180} such as the one described above, shows a generic similarity with the depletion-induced effects in polymer solutions.¹⁸¹ In the present case, however, one should note that the depletion is a consequence of strong correlations between individual ions and their image charges, which turn out to be the dominant leading-order effect in the case of neutral surfaces with multivalent ions, and thus, as such, can be described analytically within the dressed multivalent-ion approach (see Ref. 55 for more details).

B. Ions with internal charge structure

Multivalent mobile ions are not always describable by simple point-like monopolar charge structure.¹³³ Some structured counterions can be described with a dipolar^{182,183} or quadrupolar^{184,185} charge distribution, which introduce additional features in electrostatic interactions that are quite distinct from the standard PB framework. The rigid internal structure of a single counterion can be described by a charge distribution $\hat{\rho}(\mathbf{r}; \mathbf{R}, \omega)$, where \mathbf{R} gives the location of the counterion and ω are the orientational variables specifying the angular dependence of the counterion charge distribution. Here, we shall focus on the case of uniaxial counterions corresponding, e.g., to mobile charged particles of rod-like structure such as spermine or spermidine. In this case, the charge distribution $\hat{\rho}(\mathbf{r}; \mathbf{R}, \omega)$ can be expanded in a

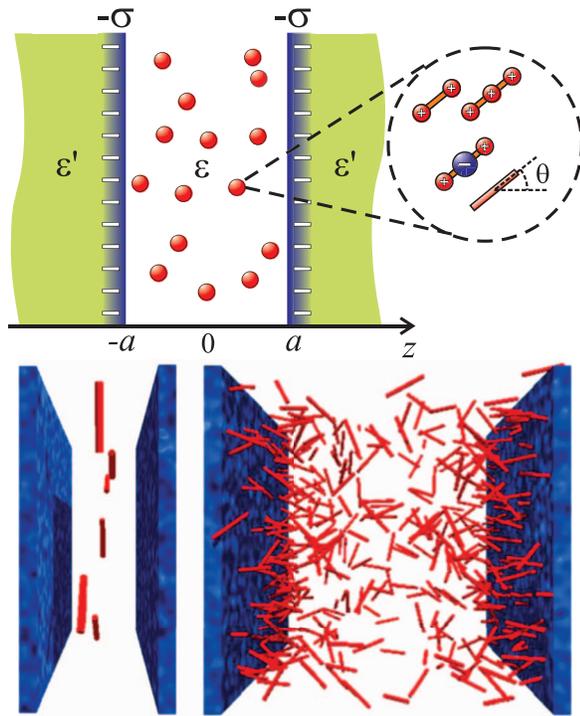


FIG. 5. Counterions can have an internal structure, such as a rod-like quadrupolar charge distribution (top). For rod-like counterions with non-zero monopolar and quadrupolar moments, the orientational order parameter in the WC limit (bottom, right) indicates preferred orientation perpendicular to the bounding surfaces, whereas in the SC limit (bottom, left) the counterions are preferably aligned parallel to the bounding surfaces.

standard multipolar series

$$\hat{\rho}(\mathbf{r}; \mathbf{R}, \boldsymbol{\omega}) = e_0 q \delta(\mathbf{r} - \mathbf{R}) - p_0 (\mathbf{n} \cdot \nabla) \delta(\mathbf{r} - \mathbf{R}) + t_0 (\mathbf{n} \cdot \nabla)^2 \delta(\mathbf{r} - \mathbf{R}) + \dots, \quad (19)$$

where $e_0 q$ is the monopolar moment of each counterion, $\mathbf{p} = p_0 \mathbf{n}$ is its dipolar moment, and $\mathbf{Q} = t_0 \mathbf{n} \otimes \mathbf{n}$ its uniaxial quadrupolar moment with director unit vector \mathbf{n} . One possible example where only the monopolar and quadrupolar moments are non-zero is a uniformly charged rod with total charge $e_0 q$ and length l for which $t_0 = e_0 q l^2 / 24$. Another possible case is a rod-like configuration with a negative charge ($-e_2$) in the center and two positive charges ($+e_1$) located at both ends of the rod, giving $e_0 q = 2e_1 - e_2$ and $t_0 = e_1 l^2$, respectively (see Fig. 5, top).

Assuming for simplicity that the system is again composed of counterions only and that these mobile counterions have a complicated internal charge distribution $\hat{\rho}(\mathbf{r}; \mathbf{R}, \boldsymbol{\omega})$, the field self-interaction in the partition function (see Eq. (3)) can be derived as¹³³

$$\mathcal{U}(\phi(\mathbf{r})) = \lambda \int d\boldsymbol{\omega} \Omega(\mathbf{r}) \exp \left(i\beta \int d\mathbf{r}' \hat{\rho}(\mathbf{r}'; \mathbf{r}, \boldsymbol{\omega}) \phi(\mathbf{r}') \right), \quad (20)$$

where λ is the fugacity of counterions. The WC limit is obtained from the saddle point approximation and the SC limit by the lowest order virial expansion corresponding to a single-particle limit. Though in this case the field-action contains more length scales than one, corresponding to the internal

structure of the counterions with in general q , p_0 , and t_0 all having distinct values, one can identify a principal coupling parameter, in analogy to the primitive counterion-only model, that implies a SC-like fixed point with a fine structure as a consequence of these additional length scales.

The WC limit for a system of rod-like counterions bounded by two like-charged plane-parallel surfaces is then given by a generalization of the PB equation that can be written in dimensionless form as¹³³

$$\psi'' = -\frac{1}{2} \int_{-1}^{+1} dx \Omega(x, z) (u(z) - pxu'(z) + tx^2u''(z)), \quad (21)$$

where ψ is the dimensionless potential (i.e., actual potential multiplied by $\beta e_0 q$), $u(z) = C \exp(-\psi - px\psi' - tx^2\psi'')$ and dimensionless multipolar moments $p = p_0/e_0 q \mu$ and $t = t_0/e_0 q \mu^2$. Here, $u(z)$ is proportional to the local orientationally dependent number density of the counterions. Since we are not interested in steric effects, we can set the blip function to $\Omega(x, z) = 1$ everywhere inside the slit between the two surfaces. Furthermore, $x = \cos \theta$ is an orientational variable, where θ corresponds to the angle between the z -axis and the director \mathbf{n} . The integral over this variable gives the orientational average. The constant C is set by the boundary conditions on the two bounding surfaces.

Numerical solutions of Eq. (21) show that increase of the quadrupolar moment leads to a *higher* concentration of counterions at the surfaces.¹³³ This can be explained by invoking the potential energy of every quadrupolar particle in electrostatic potential, proportional to the second derivative of the mean-field potential, which in the symmetric case is a concave function of the coordinate z . Thus the quadrupolar force acts away from the center toward both surfaces. Furthermore, the orientational order parameter, $S_2 = \frac{1}{2}(3\langle x^2 \rangle - 1)$, increases with increasing quadrupolar strength and is larger at both surfaces than at the center, indicating that counterion axes are preferentially aligned parallel to the z -axis and thus *perpendicular* to the bounding surfaces in the WC limit (see Fig. 5, bottom, right).

The SC limit is obtained by a virial expansion up to the term linear in λ , corresponding to a single-particle partition function.¹³³ One can show that $\mathcal{Z}_{SC}^{(0)}$ is again given by the first term in Eq. (11) involving only electrostatic interactions between both charged surfaces, so that the multipolar nature of the mobile ions is not important for this lowest order term. It becomes, however, important at the next order, linear in λ , which equals to

$$\begin{aligned} \mathcal{Z}_{SC}^{(1)} / \mathcal{Z}_{SC}^{(0)} &= \iint d\mathbf{R} d\boldsymbol{\omega} \Omega(\mathbf{r}) \\ &\times \exp \left[-\beta \iint d\mathbf{r} d\mathbf{r}' \hat{\rho}(\mathbf{r}; \mathbf{R}, \boldsymbol{\omega}) u(\mathbf{r}, \mathbf{r}') \rho_0(\mathbf{r}') \right. \\ &\left. - \frac{1}{2} \beta \iint d\mathbf{r} d\mathbf{r}' \hat{\rho}(\mathbf{r}; \mathbf{R}, \boldsymbol{\omega}) u(\mathbf{r}, \mathbf{r}') \hat{\rho}(\mathbf{r}'; \mathbf{R}, \boldsymbol{\omega}) \right]. \end{aligned} \quad (22)$$

The interaction potential in this part of the partition function is in general composed of the direct and image electrostatic interactions, i.e., $u(\mathbf{r}, \mathbf{r}') = u_0(\mathbf{r}, \mathbf{r}') + u_{\text{im}}(\mathbf{r}, \mathbf{r}')$, if the bounding surfaces have a different dielectric permittivity from the solvent. The second term in the exponent of Eq. (22) gives the image self-interactions among monopolar, dipolar, and quadrupolar moments (9 terms), whose explicit forms have been calculated in Ref. 133. Note that, in general, the self-energy contributions cannot be simply renormalized away into the rescaled fugacity.^{78,186}

In the case of two charged plane-parallel surfaces with uniform surface charge density, the electrostatic potential does not depend on the z coordinate, is thus spatially homogeneous and given by $\sigma e_0 a / (\epsilon \epsilon_0)$. Since all the terms in the density operator (19), except the first one, depend on spatial gradients (derivatives), the counterion energy in a homogenous external electrostatic potential depends only on the monopolar (first) term. The corresponding energy of a counterion in this electrostatic potential, i.e., the first term in the exponent of (22) is then given by $\beta q \sigma e_0^2 a / (\epsilon \epsilon_0)$. As for the self-energy term (second term in the exponent of (22)), it only picks up contributions from the z -dependent parts of the image self-interaction, u_{im} . Note that if there is no dielectric discontinuity, so that $u_{\text{im}}(\mathbf{r}, \mathbf{r}') = 0$, this term is identically equal to zero!

It then turns out that multipolar contributions (beyond monopoles) in the SC free energy, $\beta \mathcal{F}_{\text{SC}} = -\ln \mathcal{Z}_{\text{SC}}$, enter only via self-image interactions (see Ref. 133). Therefore, if there is no dielectric discontinuity, the multipolar effects in this limit vanish! Note that we have again assumed that the blip function Ω does not depend on the coordinate z , thus disregarding the possible entropic effects due to the finite size and anisotropy in the shape of counterions,^{184,185} which would be relevant only for small inter-surface separation on the order of the counterion size, where other competing interactions, besides electrostatic, come into play as well.^{128,187}

As for the inter-surface interaction pressure in the SC limit, the quadrupolar contribution can become negative (attractive) due to dielectric image contributions. On increase of the quadrupolar moment, the interaction becomes smaller and is eventually overwhelmed by repulsive contributions. The

orientational order parameter in the SC limit indicates that the counterions are preferably aligned perpendicular to the z -axis, *parallel* with the surfaces (see Fig. 5, bottom, left), which is contrary to the WC case. This is caused by strong quadrupole-quadrupole image repulsion when there is a dielectric discontinuity at the surfaces; otherwise, quadrupolar effects are nil as discussed above. Therefore, in the plane-parallel geometry, higher order multipoles in the internal structure of mobile ions can play an important role, but *only* for a dielectrically inhomogeneous system, i.e., when the bounding surfaces and the slit region possess different dielectric constants. Although, we should also emphasize that multipolar effects can be present if the charged surfaces are curved (and thus generate spatially varying external potentials) *even* without dielectric discontinuities, but these effects remain largely unexplored.

C. Polarizable ions—A two-parameter model

Apart from possible multipolar moments in addition to the monopolar charge, we have been up to now dealing with a *one-parameter model* of mobile (counter-)ions, as by assumption they differ only in the amount of charge they bear. Equally charged ions, such as Na^+ and Li^+ , or Ca^{2+} and Mn^{2+} , are thus indistinguishable and have no chemical identity. One possible generalization of this model is to additionally characterize the ion with its excess static ionic polarizability^{127,128,188–190} (see Fig. 6(a)), proportional to the volume of the cavity created by the ion in the solvent. Static excess ionic polarizability is then a second parameter that differentiates between different, but equally charged ionic species, and thus obviously introduces ionic specificity into the theory. Excess ionic polarizability studies go all the way back to the classical work by Debye on polar molecules.¹³¹

For a system composed of polarizable monopolar counterions, the field self-interaction can be derived¹³⁴ in the form of Eq. (3), but with

$$\mathcal{U}(\phi(\mathbf{r})) = \lambda \Omega(\mathbf{r}) \exp \left[-\beta \frac{\alpha}{2} (\nabla \phi(\mathbf{r}))^2 - i \beta q e_0 \phi(\mathbf{r}) \right], \quad (23)$$

where α is the excess polarizability of the counterions, while other details of the model are the same as for a

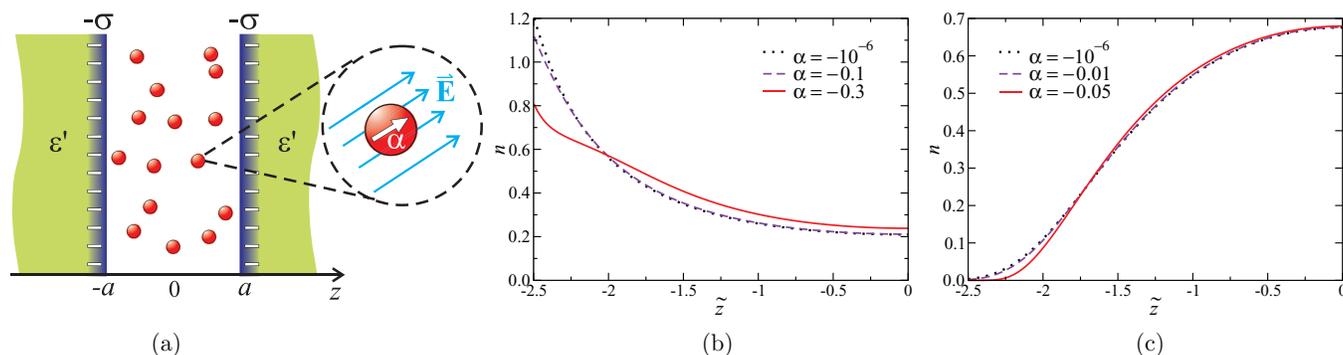


FIG. 6. (a) Counterion polarizability, α , introduces *effective interactions* between the counterions and the bounding surfaces. For negative excess polarizability, these additional interactions seem to be strongly repulsive and long ranged. (b) The WC counterions density profile (in units of $2\pi \ell_B \sigma^2$) as a function of the rescaled position, $\tilde{z} = z/\mu$, in the slit is shown next to the left surface for rescaled inter-surface half-distance $\tilde{a} = a/\mu = 2.5$ and different polarizability values α as indicated on the graph; here, the contribution from fluctuations around the mean-field solution is taken into account as well. (c) Same as (b), but here we show the SC density profile for different α . A clear effective repulsion is seen. Adapted from Ref. 134.

standard counterion-only system. The excess polarizability is defined precisely as the difference between the aqueous solvent polarizability and the proper ionic polarizability, and may thus be negative.^{128,131}

One should note here that the partition function depends on two parameters: the coupling constant Ξ as well as the polarizability α , i.e., it is a *two-parameter function*. These two parameters can be introduced in the following way: we first rewrite the electrostatic coupling constant (Eq. (1)) in the form $\Xi = q^2 \ell_B / \mu = 2\pi q^3 \ell_B^2 |\sigma| \equiv q^3 \Xi_0$, where we specifically decomposed the coupling parameter into its q and σ dependence. Then we introduce a rescaled polarizability that represents an additional independent parameter of the theory and is defined as $\tilde{\alpha} = \alpha(\beta/(q e_0 \mu)^2)$. Also, instead of using the excess polarizability, we can use the *dielectric decrement* $\tilde{\beta}$ in units of inverse mole per liter (denoted by M^{-1}),¹²⁷ i.e., $\alpha = \epsilon_0 \tilde{\beta}$. The dielectric decrement for various salts is typically negative and of the order of $10 M^{-1}$ in magnitude.

Again, we introduce the corresponding WC-SC proxies as the saddle-point and the single-particle theory, based on the lowest order virial expansion. Contrary to the cases described before, there exist at present no pertinent simulations that could help in assessing the accuracy of these approximations (however, see Ref. 191).

The saddle-point level, corresponding to the WC limit, can be derived in the form of a generalized PB equation governing the mean (real-valued) potential $\psi_{MF} = i\phi$ as^{128,188}

$$-\epsilon_0 \nabla \cdot [(\epsilon + \alpha n_{MF}(\mathbf{r})) \nabla \psi_{MF}(\mathbf{r})] = \rho_0(\mathbf{r}) + n_{MF}(\mathbf{r}), \quad (24)$$

where the mean-field counterion density is given by

$$n_{MF}(\mathbf{r}) = \lambda \exp \left[\beta \frac{\alpha}{2} (\nabla \psi_{MF}(\mathbf{r}))^2 - \beta q e_0 \psi_{MF}(\mathbf{r}) \right]. \quad (25)$$

Comparing this equation with Eq. (6), it is clear that the dielectric response function has a term proportional to the concentration of the ions, stemming from their dielectric decrement. This was in fact first noted by Bikerman in 1942.^{101,132} The solutions of this equation have been investigated in great detail in Refs. 128 and 188 and in general depend on the magnitude and sign of the excess polarizability.

From numerical solutions of the above PB equation, it appears that the counterion polarizability introduces *effective interactions* between the ions and the bounding surfaces. For negative excess polarizability, these additional interactions seem to be strongly repulsive and long ranged. They again lead to a depletion of the ions in the vicinity of the surface,¹⁷⁹ see Fig. 6(b). In the opposite case, the interactions are also repulsive, but weak, and the perturbations introduced by polarization are negligible. This yields a good measure for ion specificity, as the ions can be differentiated according to the sign of their polarizability.

On the single-particle level, corresponding to the SC limit, $\mathcal{Z}_{SC}^{(0)}$ is again given by the first term in Eq. (11), involving only electrostatic interactions between fixed surface charges. The polarizability of the mobile ions appears in the first-virial-order partition function, which is again formally of

a single-particle type and can be cast into the form¹³⁴

$$\mathcal{Z}_{SC}^{(1)} = \int d\mathbf{r}_0 z_{SC}^{(1)}(\mathbf{r}_0), \quad (26)$$

where we have introduced

$$\begin{aligned} z_{SC}^{(1)}(\mathbf{r}_0) / \mathcal{Z}_{SC}^{(0)} &= \det(1 + \alpha \nabla_i \nabla'_j u(\mathbf{r}_0, \mathbf{r}_0))^{-1/2} \\ &\times \exp \left(\frac{1}{2} \nabla_i C'(\mathbf{r}_0)^T \left(\frac{1}{\alpha} + \nabla_i \nabla'_j u(\mathbf{r}_0, \mathbf{r}_0) \right)^{-1} \right. \\ &\left. \times \nabla_i C'(\mathbf{r}_0) - \frac{C'(\mathbf{r}_0)}{2} \right), \end{aligned} \quad (27)$$

with $C'(\mathbf{r}_0) = u(\mathbf{r}_0, \mathbf{r}_0) - \int \rho_0(\mathbf{r}) u(\mathbf{r}_0, \mathbf{r}) d\mathbf{r}$. Here, $u(\mathbf{r}, \mathbf{r}')$ is again just the Coulomb interaction potential and ∇ as well as ∇' denote the gradients with respect to the first and the second variables.

Though the above form of the single-particle partition function appears to be quite complicated, it can be seen straightforwardly that the first term in Eq. (27) describes the thermal Casimir or zero-frequency vdW interaction between a single polarizable particle and the dielectric interfaces in the system. It can be written as

$$\frac{1}{2} \text{Tr} \ln(1 + \alpha \nabla_i \nabla'_j u(\mathbf{r}_0, \mathbf{r}_0)) \simeq \frac{1}{2} \alpha \text{Tr}[\nabla_i \nabla'_j u(\mathbf{r}_0, \mathbf{r}_0)]. \quad (28)$$

For large $|\mathbf{r}_0|$, we obtain the scaling form $|\mathbf{r}_0|^{-3}$ corresponding to the zero-frequency vdW interaction between the polarizable particle and a single dielectric discontinuity.^{174,192}

The corresponding single-particle-level counterion density profile, Fig. 6(c), exhibits strong image-repulsive interactions that deplete the vicinal space next to the bounding surfaces to an extent much larger than in the case of the saddle-point limit in Fig. 6(b).¹⁷⁹ Furthermore, on the single-particle level, the effective permittivity around the ion may turn overall negative, leading to a field instability that shows up in the partition function. Other nonlinear solvation-related effects¹⁸⁷ that have not been taken into account would then take over and stabilize the system.

In summary, for polarizable ions, the validity of the SC vs. WC description no longer depends on a single coupling parameter, but actually on two parameters. The parameter space is thus quite complicated and the validity of the WC-SC dichotomy is difficult to assess in general. The general conclusion would be that the contribution of polarizable counterions to the total partition function is highly non-additive at the weak coupling level, whereas it can sometimes be reduced to an additive contribution in the free energy at the strong coupling level, but only if the polarizability is large enough. Simply adding a vdW ion-polarizability-dependent contribution to the electrostatic potential of mean force is questionable.^{192,193}

V. CONCLUSIONS

We have provided a *guided personalized tour* of recent advances in Coulomb fluids based on the functional integral representation of the partition function via a field-action, as pioneered in the fundamental work of Edwards and Lenard.¹⁶⁷ This representation was taken later as a point of departure

for the introduction of the WC-SC dichotomy in the description of a (primitive) counterion-only system bounded by two charged surfaces. The WC limit was shown to stem from the saddle-point description of the field-action, while the SC description was based on the lowest order virial expansion. In the case of the primitive counterion-only model, the two limits can be derived directly and systematically. For more complicated, but also more realistic cases that imply a multiplicity of different length scales, one can identify the principal coupling parameter and introduce the pertaining saddle-point and the single-particle virial description as the proxies for the proper WC and SC limits. The saddle-point and the single-particle virial form of the partition function can then be invoked for any system, irrespective of the number of parameters or coupling constants describing it, and thus present a convenient point of departure to derive powerful limiting laws even for much more complicated systems than envisioned in the original framework of the WC-SC dichotomy. We tried to make a clear case that this philosophy can provide a solid foundation and a fairly accurate guide for an approximate and sometimes even analytical treatment of Coulomb systems that formally do not easily yield themselves to a simple single-coupling-parameter description.

ACKNOWLEDGMENTS

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- ¹J. Israelachvili, *Intermolecular and Surface Forces* (Academic Press, London, 1991).
- ²E. J. Verwey and J. T. G. Overbeek, *Theory of the Stability of Lyophobic Colloids* (Elsevier, Amsterdam, 1948).
- ³R. J. Hunter, *Foundations of Colloidal Science*, 2nd ed. (Oxford University Press, New York, 2001).
- ⁴S. A. Safran, *Statistical Thermodynamics of Surfaces, Interfaces, and Membranes* (Addison-Wesley Publishing Co., Reading, MA, 1994).
- ⁵*Soft Condensed Matter Physics in Molecular and Cell Biology*, edited by W. C. K. Poon and D. Andelman (Taylor & Francis, New York, London, 2006).
- ⁶*Electrostatic Effects in Soft Matter and Biophysics*, edited by C. Holm, P. Kekicheff, and R. Podgornik (Kluwer Academic, Dordrecht, 2001).
- ⁷R. French *et al.*, *Rev. Mod. Phys.* **82**, 1887 (2010).
- ⁸M. Baus and J. Hansen, *Phys. Rep.* **59**, 1 (1980).
- ⁹A. Khan, B. Jönsson, and H. Wennerström, *J. Phys. Chem.* **89**, 5180 (1985).
- ¹⁰R. Kjellander, S. Marčelja, and J. P. Quirk, *J. Colloid Interface Sci.* **126**, 194 (1988).
- ¹¹V. A. Bloomfield, *Biopolymers* **31**, 1471 (1991).
- ¹²V. A. Bloomfield, *Curr. Opin. Struct. Biol.* **6**, 334 (1996).
- ¹³P. Kékicheff, S. Marčelja, T. J. Senden, and V. E. Shubin, *J. Chem. Phys.* **99**, 6098 (1993).
- ¹⁴P. Gonzalez-Monzuelos and M. Olvera de la Cruz, *J. Chem. Phys.* **103**, 3145 (1995).
- ¹⁵M. Olvera de la Cruz, L. Belloni, M. Delsanti, J. P. Dalbiez, O. Spalla, and M. Drifford, *J. Chem. Phys.* **103**, 5781 (1995).
- ¹⁶M. Dubois, T. Zemb, N. Fuller, R. P. Rand, and V. A. Parsegian, *J. Chem. Phys.* **108**, 7855 (1998).
- ¹⁷H. H. Strey, R. Podgornik, D. C. Rau, and V. A. Parsegian, *Curr. Opin. Struct. Biol.* **8**, 309 (1998).
- ¹⁸J. X. Tang, T. Ito, T. Tao, P. Traub, and P. A. Janmey, *Biochemistry* **36**, 12600 (1997).
- ¹⁹G. C. L. Wong, A. Lin, J. X. Tang, Y. Li, P. A. Janmey, and C. R. Safinya, *Phys. Rev. Lett.* **91**, 018103 (2003).
- ²⁰J. C. Butler, T. Angelini, J. X. Tang, and G. C. L. Wong, *Phys. Rev. Lett.* **91**, 028301 (2003).
- ²¹T. E. Angelini, H. Liang, W. Wriggers, and G. C. L. Wong, *Proc. Natl. Acad. Sci. U.S.A.* **100**, 8634 (2003).
- ²²G. C. L. Wong and L. Pollack, *Annu. Rev. Phys. Chem.* **61**, 171 (2010).
- ²³L. Guldbrand, B. Jönsson, H. Wennerström, and P. Linse, *J. Chem. Phys.* **80**, 2221 (1984).
- ²⁴B. Svensson and B. Jönsson, *Chem. Phys. Lett.* **108**, 580 (1984).
- ²⁵D. Bratko, B. Jönsson, and H. Wennerström, *Chem. Phys. Lett.* **128**, 449 (1986).
- ²⁶L. Guldbrand, L. G. Nilsson, and L. Nordenskiöld, *J. Chem. Phys.* **85**, 6686 (1986).
- ²⁷C. E. Woodward, B. Jönsson, and T. Åkesson, *J. Chem. Phys.* **89**, 5145 (1988).
- ²⁸J. P. Valleau, R. Ivkov, and G. M. Torrie, *J. Chem. Phys.* **95**, 520 (1991).
- ²⁹R. Kjellander, T. Åkesson, B. Jönsson, and S. Marčelja, *J. Chem. Phys.* **97**, 1424 (1992).
- ³⁰A. P. Lyubartsev and L. Nordenskiöld, *J. Phys. Chem.* **99**, 10373 (1995).
- ³¹N. Grønbech-Jensen, R. J. Mashl, R. F. Bruinsma, and W. M. Gelbart, *Phys. Rev. Lett.* **78**, 2477 (1997).
- ³²N. Grønbech-Jensen, K. M. Beardmore, and P. Pincus, *Physica A* **261**, 74 (1998).
- ³³J. Wu, D. Bratko, and J. M. Prausnitz, *Proc. Natl. Acad. Sci. U.S.A.* **95**, 15169 (1998).
- ³⁴E. Allahyarov, I. D'Amico, and H. Löwen, *Phys. Rev. Lett.* **81**, 1334 (1998).
- ³⁵M. J. Stevens, *Phys. Rev. Lett.* **82**, 101 (1999).
- ³⁶P. Linse and V. Lobaskin, *Phys. Rev. Lett.* **83**, 4208 (1999).
- ³⁷P. Linse and V. Lobaskin, *J. Chem. Phys.* **112**, 3917 (2000).
- ³⁸B. Hribar and V. Vlady, *Biophys. J.* **78**, 694 (2000).
- ³⁹R. Messina, C. Holm, and K. Kremer, *Phys. Rev. Lett.* **85**, 872 (2000).
- ⁴⁰M. Deserno, A. Arnold, and C. Holm, *Macromolecules* **36**, 249 (2003).
- ⁴¹K.-C. Lee, I. Borukhov, W. M. Gelbart, A. J. Liu, and M. J. Stevens, *Phys. Rev. Lett.* **93**, 128101 (2004).
- ⁴²A. Naji and R. R. Netz, *Phys. Rev. Lett.* **95**, 185703 (2005).
- ⁴³A. Naji and R. R. Netz, *Phys. Rev. E* **73**, 056105 (2006).
- ⁴⁴A. G. Moreira and R. R. Netz, *Europhys. Lett.* **52**, 705 (2000).
- ⁴⁵A. G. Moreira and R. R. Netz, *Phys. Rev. Lett.* **87**, 078301 (2001).
- ⁴⁶A. G. Moreira and R. R. Netz, *Eur. Phys. J. E* **8**, 33 (2002).
- ⁴⁷A. G. Moreira and R. R. Netz, *Europhys. Lett.* **57**, 911 (2002).
- ⁴⁸A. Naji, A. Arnold, C. Holm, and R. R. Netz, *Europhys. Lett.* **67**, 130 (2004).
- ⁴⁹A. Naji and R. R. Netz, *Eur. Phys. J. E* **13**, 43 (2004).
- ⁵⁰H. Boroudjerdi, Y. W. Kim, A. Naji, R. R. Netz, X. Schlagberger, and A. Serr, *Phys. Rep.* **416**, 129 (2005).
- ⁵¹A. Naji, S. Jungblut, A. G. Moreira, and R. R. Netz, *Physica A* **352**, 131 (2005).
- ⁵²A. Naji, M. Kanduč, R. R. Netz, and R. Podgornik, in *Understanding Soft Condensed Matter via Modeling and Computation*, edited by W.-B. Hu and A.-C. Shi, Series in Soft Condensed Matter Vol. 3, edited by D. Andelman and G. Reiter (World Scientific, Singapore, 2010), Chap. 9; e-print [arXiv:1008.0357](https://arxiv.org/abs/1008.0357).
- ⁵³M. Kanduč, A. Naji, J. Forsman, and R. Podgornik, *J. Chem. Phys.* **132**, 124701 (2010).
- ⁵⁴M. Kanduč, A. Naji, J. Forsman, and R. Podgornik, *Phys. Rev. E* **84**, 011502 (2011).
- ⁵⁵M. Kanduč, A. Naji, J. Forsman, and R. Podgornik, *J. Chem. Phys.* **137**, 174704 (2012).
- ⁵⁶Y. S. Jho, M. Kanduč, A. Naji, R. Podgornik, M. W. Kim, and P. A. Pincus, *Phys. Rev. Lett.* **101**, 188101 (2008).
- ⁵⁷M. Kanduč, M. Trulsson, A. Naji, Y. Burak, J. Forsman, and R. Podgornik, *Phys. Rev. E* **78**, 061105 (2008).
- ⁵⁸Y.-G. Chen and J. D. Weeks, *Proc. Natl. Acad. Sci. U.S.A.* **103**, 7560 (2006).
- ⁵⁹J. M. Rodgers, C. Kaur, Y.-G. Chen, and J. D. Weeks, *Phys. Rev. Lett.* **97**, 097801 (2006).
- ⁶⁰J. Forsman, *J. Phys. Chem. B* **108**, 9236 (2004).
- ⁶¹M. Trulsson, B. Jönsson, T. Åkesson, J. Forsman, and C. Labbez, *Phys. Rev. Lett.* **97**, 068302 (2006).
- ⁶²R. Kjellander and S. Marčelja, *Chem. Phys. Lett.* **112**, 49 (1984).
- ⁶³R. Kjellander and S. Marčelja, *J. Chem. Phys.* **82**, 2122 (1985).
- ⁶⁴P. Attard, R. Kjellander, and D. J. Mitchell, *Chem. Phys. Lett.* **139**, 219 (1987).

- ⁶⁵P. Attard, D. J. Mitchell, and B. W. Ninham, *J. Chem. Phys.* **88**, 4987 (1988).
- ⁶⁶R. Podgornik and B. Žekš, *J. Chem. Soc., Faraday Trans. 2* **84**, 611 (1988).
- ⁶⁷R. Podgornik, *J. Phys. A: Math. Gen.* **23**, 275 (1990).
- ⁶⁸R. Podgornik and V. A. Parsegian, *Phys. Rev. Lett.* **80**, 1560 (1998).
- ⁶⁹J.-L. Barrat and J.-F. Joanny, *Adv. Chem. Phys.* **XCIV**, 1 (1996).
- ⁷⁰P. A. Pincus and S. A. Safran, *Europhys. Lett.* **42**, 103 (1998).
- ⁷¹B.-Y. Ha and A. J. Liu, *Phys. Rev. Lett.* **79**, 1289 (1997).
- ⁷²B.-Y. Ha and A. J. Liu, *Phys. Rev. Lett.* **81**, 1011 (1998).
- ⁷³D. B. Lukatsky and S. A. Safran, *Phys. Rev. E* **60**, 5848 (1999).
- ⁷⁴A. W. C. Lau, D. B. Lukatsky, P. Pincus, and S. A. Safran, *Phys. Rev. E* **65**, 051502 (2002).
- ⁷⁵M. Kardar and R. Golestanian, *Rev. Mod. Phys.* **71**, 1233 (1999).
- ⁷⁶B.-Y. Ha, *Phys. Rev. E* **64**, 031507 (2001).
- ⁷⁷A. W. C. Lau and P. Pincus, *Phys. Rev. E* **66**, 041501 (2002).
- ⁷⁸R. R. Netz and H. Orland, *Eur. Phys. J. E* **1**, 203 (2000).
- ⁷⁹R. R. Netz and H. Orland, *Eur. Phys. J. E* **11**, 301 (2003).
- ⁸⁰S. Buyukdagli, M. Manghi, and J. Palmeri, *Phys. Rev. E* **81**, 041601 (2010).
- ⁸¹S. Buyukdagli, M. Manghi, and J. Palmeri, *Phys. Rev. Lett.* **105**, 158103 (2010).
- ⁸²S. Buyukdagli, M. Manghi, and J. Palmeri, *J. Chem. Phys.* **134**, 074706 (2011).
- ⁸³M. J. Stevens and M. O. Robbins, *Europhys. Lett.* **12**, 81 (1990).
- ⁸⁴A. Diehl, M. N. Tamashiro, M. C. Barbosa, and Y. Levin, *Physica A* **274**, 433 (1999).
- ⁸⁵O. Gonzalez-Amezcuca, M. Hernandez-Contreras, and P. Pincus, *Phys. Rev. E* **64**, 041603 (2001).
- ⁸⁶I. Rouzina and V. A. Bloomfield, *J. Phys. Chem.* **100**, 9977 (1996).
- ⁸⁷A. A. Kornyshev and S. Leikin, *J. Chem. Phys.* **107**, 3656 (1997).
- ⁸⁸J. J. Arenzon, J. F. Stilck, and Y. Levin, *Eur. Phys. J. B* **12**, 79 (1999).
- ⁸⁹Y. Levin, J. J. Arenzon, and J. F. Stilck, *Phys. Rev. Lett.* **83**, 2680 (1999).
- ⁹⁰B. I. Shklovskii, *Phys. Rev. Lett.* **82**, 3268 (1999).
- ⁹¹A. Yu. Grosberg, T. T. Nguyen, and B. I. Shklovskii, *Rev. Mod. Phys.* **74**, 329 (2002).
- ⁹²A. Diehl, H. A. Carmona, and Y. Levin, *Phys. Rev. E* **64**, 011804 (2001).
- ⁹³A. W. C. Lau, D. Levine, and P. Pincus, *Phys. Rev. Lett.* **84**, 4116 (2000).
- ⁹⁴R. R. Netz, *Eur. Phys. J. E* **5**, 557 (2001).
- ⁹⁵Y. Levin, *Rep. Prog. Phys.* **65**, 1577 (2002).
- ⁹⁶Y. Burak, D. Andelman, and H. Orland, *Phys. Rev. E* **70**, 016102 (2004).
- ⁹⁷C. D. Santangelo, *Phys. Rev. E* **73**, 041512 (2006).
- ⁹⁸M. M. Hatlo and L. Lue, *Soft Matter* **5**, 125 (2009).
- ⁹⁹L. Šamaj and E. Trizac, *Phys. Rev. Lett.* **106**, 078301 (2011).
- ¹⁰⁰L. Šamaj and E. Trizac, *Phys. Rev. E* **84**, 041401 (2011).
- ¹⁰¹P. Duncan, M. M. Hatlo, and L. Lue, in *Electrostatics of Soft and Disordered Matter*, edited by D. S. Dean, J. Dobnikar, A. Naji, and R. Podgornik (Pan Stanford Publishing, Singapore, in press), Chap. 2.
- ¹⁰²M. Kanduč, J. Dobnikar, and R. Podgornik, *Soft Matter* **5**, 868 (2009).
- ¹⁰³M. Kanduč, A. Naji, and R. Podgornik, *J. Chem. Phys.* **132**, 224703 (2010).
- ¹⁰⁴M. Kanduč and R. Podgornik, *Eur. Phys. J. E* **23**, 265 (2007).
- ¹⁰⁵I. Rouzina and V. A. Bloomfield, *Biophys. J.* **74**, 3152 (1998).
- ¹⁰⁶R. Golestanian, M. Kardar, and T. B. Liverpool, *Phys. Rev. Lett.* **82**, 4456 (1999).
- ¹⁰⁷Y. Levin, *Physica A* **265**, 432 (1999).
- ¹⁰⁸T. T. Nguyen, I. Rouzina, and B. I. Shklovskii, *J. Chem. Phys.* **112**, 2562 (2000).
- ¹⁰⁹O. Punkkinen, A. Naji, R. Podgornik, I. Vattulainen, and P.-L. Hansen, *Europhys. Lett.* **82**, 48001 (2008).
- ¹¹⁰R. Messina, *J. Phys.: Condens. Matter* **21**, 113102 (2009).
- ¹¹¹F. Oosawa, *Biopolymers* **6**, 1633 (1968).
- ¹¹²F. Oosawa, *Polyelectrolytes* (Marcel Dekker, New York, 1971).
- ¹¹³G. N. Patey, *J. Chem. Phys.* **72**, 5763 (1980).
- ¹¹⁴J. Ray and G. S. Manning, *Macromolecules* **30**, 5739 (1997).
- ¹¹⁵D. J. Needleman, M. A. Ojeda-Lopez, U. Raviv, H. P. Miller, L. Wilson, and C. R. Safinya, *Proc. Natl. Acad. Sci. U.S.A.* **101**, 16099 (2004).
- ¹¹⁶J. Pelta, D. Durand, J. Doucet, and F. Livolant, *Biophys. J.* **71**, 48 (1996); J. Pelta, F. Livolant, and J.-L. Sikorav, *J. Biol. Chem.* **271**, 5656 (1996).
- ¹¹⁷K. Yoshikawa, *Adv. Drug Deliv. Rev.* **52**, 235 (2001).
- ¹¹⁸M. Takahashi, K. Yoshikawa, V. V. Vasilevskaya, and A. R. Khokhlov, *J. Phys. Chem. B* **101**, 9396 (1997).
- ¹¹⁹G. E. Plum and V. A. Bloomfield, *Biopolymers* **27**, 1045 (1988).
- ¹²⁰E. Raspaud, I. Chaperon, A. Leforestier, and F. Livolant, *Biophys. J.* **77**, 1547 (1999).
- ¹²¹H. S. Savithri, S. K. Munshi, S. Suryanarayana, S. Divakar, and M. R. N. Murthy, *J. Gen. Virol.* **68**, 1533 (1987).
- ¹²²M. de Frutos, S. Brasiles, P. Tavares, and E. Raspaud, *Eur. Phys. J. E* **17**, 429 (2005).
- ¹²³A. Leforestier, A. Siber, F. Livolant, and R. Podgornik, *Biophys. J.* **100**, 2209 (2011).
- ¹²⁴D. S. Dean, R. R. Horgan, and D. Sentenac, *J. Stat. Phys.* **90**, 899 (1998).
- ¹²⁵D. S. Dean, R. R. Horgan, A. Naji, and R. Podgornik, *J. Chem. Phys.* **130**, 094504 (2009).
- ¹²⁶D. Ben-Yaakov, D. Andelman, D. Harries, and R. Podgornik, *J. Phys.: Condens. Matter* **21**, 424106 (2009).
- ¹²⁷D. Ben-Yaakov, D. Andelman, and R. Podgornik, *J. Chem. Phys.* **134**, 074705 (2011).
- ¹²⁸D. Ben-Yaakov, D. Andelman, R. Podgornik, and D. Harries, *Curr. Opin. Colloid Interface Sci.* **16**, 542 (2011).
- ¹²⁹Y. Burak and D. Andelman, *Phys. Rev. E* **62**, 5296 (2000).
- ¹³⁰Y. Burak and D. Andelman, *J. Chem. Phys.* **114**, 3271 (2001).
- ¹³¹P. Debye, *Polar Molecules* (Dover, New York, 1929).
- ¹³²J. J. Bikerman, *Philos. Mag.* **33**, 384 (1942).
- ¹³³M. Kanduč, A. Naji, Y. S. Jho, P. A. Pincus, and R. Podgornik, *J. Phys.: Condens. Matter* **21**, 424103 (2009).
- ¹³⁴V. Démery, D. S. Dean, and R. Podgornik, *J. Chem. Phys.* **137**, 174903 (2012).
- ¹³⁵M. M. Hatlo, R. van Roij, and L. Lue, *Europhys. Lett.* **97**, 28010 (2012).
- ¹³⁶R. R. Netz, *Eur. Phys. J. E* **5**, 189 (2001).
- ¹³⁷D. Horinek and R. R. Netz, *Phys. Rev. Lett.* **99**, 226104 (2007).
- ¹³⁸A. A. Kornyshev, *J. Phys. Chem. B* **111**, 5545 (2007).
- ¹³⁹V. Démery, D. S. Dean, T. C. Hammant, R. R. Horgan, and R. Podgornik, *Europhys. Lett.* **97**, 28004 (2012).
- ¹⁴⁰V. Démery, D. S. Dean, T. C. Hammant, R. R. Horgan, and R. Podgornik, *J. Chem. Phys.* **137**, 064901 (2012).
- ¹⁴¹O. Spalla and L. Belloni, *Phys. Rev. Lett.* **74**, 2515 (1995).
- ¹⁴²J. Forsman, *Langmuir* **22**, 2975 (2006).
- ¹⁴³I. Borukhov, D. Andelman, and H. Orland, *Eur. Phys. J. B* **5**, 869 (1998).
- ¹⁴⁴Y. Kantor and M. Kardar, *Europhys. Lett.* **28**, 169 (1994).
- ¹⁴⁵E. E. Meyer, Q. Lin, T. Hassenkam, E. Oroudjev, and J. N. Israelachvili, *Proc. Natl. Acad. Sci. U.S.A.* **102**, 6839 (2005).
- ¹⁴⁶S. Perkin, N. Kampf, and J. Klein, *Phys. Rev. Lett.* **96**, 038301 (2006).
- ¹⁴⁷D. Ben-Yaakov, D. Andelman, and H. Diamant, *Phys. Rev. E* **87**, 022402 (2013).
- ¹⁴⁸C. Fleck and R. R. Netz, *Europhys. Lett.* **70**, 341 (2005).
- ¹⁴⁹A. Naji and R. Podgornik, *Phys. Rev. E* **72**, 041402 (2005).
- ¹⁵⁰R. Podgornik and A. Naji, *Europhys. Lett.* **74**, 712 (2006).
- ¹⁵¹Y. Sh. Mamasakhlisov, A. Naji, and R. Podgornik, *J. Stat. Phys.* **133**, 659 (2008).
- ¹⁵²A. Naji, D. S. Dean, J. Sarabadani, R. Horgan, and R. Podgornik, *Phys. Rev. Lett.* **104**, 060601 (2010).
- ¹⁵³J. Sarabadani, A. Naji, D. S. Dean, R. R. Horgan, and R. Podgornik, *J. Chem. Phys.* **133**, 174702 (2010).
- ¹⁵⁴D. S. Dean, A. Naji, and R. Podgornik, *Phys. Rev. E* **83**, 011102 (2011).
- ¹⁵⁵A. Naji, J. Sarabadani, D. S. Dean, and R. Podgornik, *Eur. Phys. J. E* **35**, 24 (2012).
- ¹⁵⁶V. Rezvani, J. Sarabadani, A. Naji, and R. Podgornik, *J. Chem. Phys.* **137**, 114704 (2012).
- ¹⁵⁷B. W. Ninham and V. A. Parsegian, *J. Theor. Biol.* **31**, 405 (1971).
- ¹⁵⁸D. Chan, J. W. Perram, L. R. White, and T. H. Healy, *J. Chem. Soc., Faraday Trans. 1* **71**, 1046 (1975).
- ¹⁵⁹H. H. von Grünberg, *J. Colloid Interface Sci.* **219**, 339 (1999).
- ¹⁶⁰G. S. Longo, M. Olvera de la Cruz, and I. Szleifer, *Soft Matter* **8**, 1344 (2012).
- ¹⁶¹G. S. Longo, M. Olvera de la Cruz, and I. Szleifer, *ACS Nano* **7**, 2693 (2013).
- ¹⁶²J. Dobnikar, M. Brunner, H. H. von Grünberg, and C. Bechinger, *Phys. Rev. E* **69**, 031402 (2004).
- ¹⁶³J. Dobnikar, Y. Chen, R. Rzehak, and H. H. Grünberg, *J. Phys.: Condens. Matter* **15**, S263 (2003).
- ¹⁶⁴N. Bjerrum, *Kgl. Dan. Videnskab. Selsk. Mat. Fys. Medd.* **7**, 1 (1926).
- ¹⁶⁵J. Zwanikken and R. van Roij, *J. Phys.: Condens. Matter* **21**, 424102 (2009).
- ¹⁶⁶M. E. Fisher and Y. Levin, *Phys. Rev. Lett.* **71**, 3826 (1993).
- ¹⁶⁷S. F. Edwards and A. Lenard, *J. Math. Phys.* **3**, 778 (1962).
- ¹⁶⁸D. Henderson and L. Blum, *J. Chem. Phys.* **75**, 2025 (1981).
- ¹⁶⁹H. Wennerström, B. Jönsson, and P. Linse, *J. Chem. Phys.* **76**, 4665 (1982).

- ¹⁷⁰D. S. Dean and R. R. Horgan, *Phys. Rev. E* **68**, 061106 (2003).
- ¹⁷¹J. C. Neu, *Phys. Rev. Lett.* **82**, 1072 (1999).
- ¹⁷²J. E. Sader and D. Y. C. Chan, *J. Colloid Interface Sci.* **213**, 268 (1999).
- ¹⁷³J. E. Sader and D. Y. C. Chan, *Langmuir* **16**, 324 (2000).
- ¹⁷⁴V. A. Parsegian, *Van der Waals Forces: A Handbook for Biologists, Chemists, Engineers, and Physicists* (Cambridge University Press, 2005).
- ¹⁷⁵V. Vlachy, *Annu. Rev. Phys. Chem.* **50**, 145 (1999).
- ¹⁷⁶G. Lamm, *Reviews in Computational Chemistry*, edited by K. B. Lipkowitz, R. Larter, and T. R. Cundari (Wiley-VCH, 2003), Vol. 19, pp. 147–365.
- ¹⁷⁷D. C. Rau and V. A. Parsegian, *Biophys. J.* **61**, 246 (1992).
- ¹⁷⁸D. C. Rau and V. A. Parsegian, *Biophys. J.* **61**, 260 (1992).
- ¹⁷⁹J. W. Zwanikken and M. Olvera de la Cruz, *Proc. Natl. Acad. Sci. U.S.A.* **110**, 5301 (2013).
- ¹⁸⁰S. Buyukdagli and T. Ala-Nissila, *Europhys. Lett.* **98**, 60003 (2012).
- ¹⁸¹H. N. W. Lekkerkerker and R. Tuinier, *Colloids and the Depletion Interaction* (Springer, New York, 2011).
- ¹⁸²A. Abrashkin, D. Andelman, and H. Orland, *Phys. Rev. Lett.* **99**, 077801 (2007).
- ¹⁸³S. Maset and K. Bohinc, *J. Phys. A: Math. Theor.* **40**, 11815 (2007).
- ¹⁸⁴Y. W. Kim, J. Yi, and P. A. Pincus, *Phys. Rev. Lett.* **101**, 208305 (2008).
- ¹⁸⁵S. Maset, J. Reščič, S. May, J. I. Pavlič, and K. Bohinc, *J. Phys. A: Math. Theor.* **42**, 105401 (2009).
- ¹⁸⁶Z.-G. Wang, *Phys. Rev. E* **81**, 021501 (2010).
- ¹⁸⁷B. E. Conway, *Ionic Hydration in Chemistry and Biophysics* (Elsevier, New York, 1981).
- ¹⁸⁸D. Frydel, *J. Chem. Phys.* **134**, 234704 (2011).
- ¹⁸⁹S. Buyukdagli and T. Ala-Nissila, *Phys. Rev. E* **87**, 063201 (2013).
- ¹⁹⁰S. Buyukdagli and T. Ala-Nissila, *J. Chem. Phys.* **139**, 044907 (2013).
- ¹⁹¹J. R. Bordin and C. Holm, “Influence of static polarization in counterions distribution near charged surfaces: A coarse-grained molecular dynamics study” (unpublished).
- ¹⁹²B. W. Ninham and V. Yaminsky, *Langmuir* **13**, 2097 (1997).
- ¹⁹³P. Lo Nostro and B. Ninham, *Chem. Rev.* **112**, 2286 (2012).