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Communication: Mean-field theory of water-water correlations in electrolyte solutions

David M. Wilkins,1,2,a) David E. Manolopoulos,3 Sylvie Roke,2 and Michele Ceriotti1
1Laboratory of Computational Science and Modeling, IMX, École Polytechnique Fédérale de Lausanne, 1015 Lausanne, Switzerland
2Laboratory for Fundamental BioPhotonics, Institutes of Bioengineering and Materials Science and Engineering, School of Engineering, and Lausanne Centre for Ultrafast Science, École Polytechnique Fédérale de Lausanne, CH-1015 Lausanne, Switzerland
3Physical and Theoretical Chemistry Laboratory, University of Oxford, South Parks Road, Oxford OX1 3QZ, United Kingdom

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Long-range ion induced water-water correlations were recently observed in femtosecond elastic second harmonic scattering experiments of electrolyte solutions. To further the qualitative understanding of these correlations, we derive an analytical expression that quantifies ion induced dipole-dipole correlations in a non-interacting gas of dipoles. This model is a logical extension of the Debye-Hückel theory that can be used to qualitatively understand how the combined electric field of the ions induces correlations in the orientational distributions of the water molecules in an aqueous solution. The model agrees with the results from molecular dynamics simulations and provides an important starting point for further theoretical work. Published by AIP Publishing. [http://dx.doi.org/10.1063/1.4983221]

The electric field of a solvated ion in water induces orientational ordering in the surrounding solvent molecules. However, the length scale over which this ordering persists has been a topic of significant debate, at least in part because the range at which correlations can be detected depends on the experimental probe.1 The results of neutron diffraction,2,3 X-ray scattering,2,3 dielectric relaxation,6 and femtosecond pump-probe experiments,7 as well as atomistic simulations of the reorientation time scales of water molecules8 and of the vibrational spectra of solutions,9,10 have suggested that the ordering of the surrounding water molecules by ions extends no further than about 3 solvation shells (around 0.8 nm) for sub-molar concentrations. On the other hand, infrared photodissociation experiments,11,12 and a study combining terahertz and femtosecond infrared spectroscopies,13 have found evidence for ordering extended to longer ranges. Molecular dynamics simulations looking directly at the orientational correlations between water molecules showed that the presence of ionic solutes has an effect on these correlations at distances of more than 1 nm.14

Femtosecond elastic second harmonic scattering (fs-ESHS)15,16 measurements have recently been used to probe the orientational order of water molecules in H2O and D2O electrolyte solutions,17 revealing intensity changes that are already detectable at micromolar concentrations, and which are identical for more than 20 different electrolytes. The non-specificity of the fs-ESHS response, its magnitude, and its onset at low concentration point to its long-range origin. This isotope exchange experiment, together with the recorded polarization combinations (in conjunction with the selection rules for nonlinear light scattering experiments16,18), shows that the recorded changes in the fs-ESHS response in the concentration range from 1 μM to 100 mM arise from water-water correlations that are induced by the ions (and not from the ions themselves). This effect shows intriguing correlations with changes in the surface tension of dilute electrolyte solutions, suggesting that the microscopic phenomenon underlying the second-harmonic signal can have an impact on macroscopic observables.

In this Communication, we derive an analytical expression for the correlations induced in a non-interacting gas of dipoles by the electric field of ions. This expression is a natural extension to a simple Debye-Hückel model, which has been shown to qualitatively capture the concentration-dependence of the second-harmonic response,17,19 and can be used to elucidate the nature, the range, and the energetics of the weak ion-induced ordering probed by fs-ESHS.20 The expression provides a benchmark for a fundamental understanding of the interplay of ion-dipole and ion-ion interactions. By comparison with classical molecular dynamics simulations of dilute NaCl solutions, we demonstrate that both of these factors are needed to characterize the ion-induced solvent correlations.

We begin by considering the water molecules in an ionic solution to be point dipoles that interact only with the solute and have no explicit dipole-dipole interactions. The orientational ordering of these dipoles is thus caused only by the electric field due to the ions. A similar philosophy has been applied in mean-field theories that have existed for many decades, in which water molecules are treated as point dipoles in a dielectric continuum.21 These theories have been used to understand several physical properties, including the local field factor that describes the reaction field on a molecule due to its polarization of the surrounding medium,22 the
dielectric constant of a liquid, and dynamical properties such as dielectric relaxation times. Although such a theory cannot report on short-ranged hydrogen-bonding and dipole-dipole interactions, the dipolar screening is implicitly included. We will also show later that dipole-dipole interactions can be included in a refined version of our model and have no impact on the long-range behavior.

Figs. 1(a) and 1(b) show how our model is built up: first, the ions are taken to be point charges in a dielectric continuum, with an appropriate spatial distribution, after which the system is filled with a uniform gas of independent dipoles, which will align with the local electric field. We then define the dipole correlation function for two solvent molecules separated by a distance \( r \) (that is, the average inner product of two dipoles as a function of their separation),

\[
\langle \cos \phi \rangle (r) = \frac{1}{V} \int_V \langle \hat{\mu}(R) \cdot \hat{\mu}(R + r) \rangle_{\text{ori}} d^3R,
\]

where \( \hat{\mu}(R) \) is the unit vector in the direction of the dipole moment of a molecule at \( R \), \( V \) is the volume of the system, and \( \text{“ori”} \) denotes an average over molecular orientations and ionic positions. Fig. 1(c) illustrates how the angle \( \phi \) is defined for two representative water molecules.

In the supplementary material, we show that by taking a Taylor expansion in the reciprocal temperature \( \beta = 1/k_B T \), we can make the approximation,

\[
\langle \cos \phi \rangle (r) \approx \frac{1}{V} \left( \frac{\beta \mu}{3} \right)^2 \int_V \langle E(R) \cdot E(R + r) \rangle_{\text{ori}} d^3R,
\]

where \( E(R) \) is the total electric field at position \( R \) due to all of the ions in the solution, and \( \mu \) is the permanent dipole moment of a water molecule. The subscript “ori” indicates that the average is taken over the positions of ions. For simplicity of notation, any angular brackets in the following work without a subscript are taken over ion positions only. Eq. (2) shows that in our model, the correlation between dipoles is proportional to the correlation between electric fields, which are taken to be the only source of ordering for the molecules.

The electric field \( E(R) \) at a given position \( R \) is the sum of electric fields due to all of the ions. This allows us to write,

\[
E(R) = \sum_{m \in \text{ions}} \frac{eZ_m f_0}{4\pi \epsilon_0 \epsilon_r} \mathbf{E}(R - r_m),
\]

with \( Z_m \) the charge of the \( m \)th ion in units of the electron charge \( e \) and \( r_m \) the position of this ion, \( f_0 \) the Onsager local field factor, \( \epsilon_0 \) the vacuum permittivity, \( \epsilon_r \) the solvent dielectric constant, and \( \mathbf{E}(r) \) the electric field associated with individual ions (most commonly the Coulomb field, \( r/r^3 \)). This gives

\[
\langle \cos \phi \rangle (r) \approx \frac{A}{V} \sum_{m,n} Z_m Z_n \int_V \langle \mathbf{E}(R) \cdot \mathbf{E}(R + r) \rangle \cdot \mathbf{E}(R + r - r_m) d^3R,
\]

in which we have defined \( A = \left( \frac{\beta \mu \epsilon_0}{12\pi \epsilon_r \epsilon_0} \right)^2 \).

In the thermodynamic \( (V \rightarrow \infty) \) limit, the integral in Eq. (4) is taken over all space and can be most conveniently expressed in reciprocal space,

\[
\langle \cos \phi \rangle (r) \approx \frac{A}{V} \int \mathbf{E} (K) \cdot \mathbf{E} (-K) \times \left( \sum_{m,n} Z_m Z_n e^{iK \cdot (r_m - r_n)} \right) e^{iK \cdot r} \frac{d^3K}{(2\pi)^3},
\]

where \( \mathbf{E}(K) \) is the Fourier transform of the field function \( \mathbf{E}(r) \). The term in angular brackets is proportional to the charge-charge structure factor \( S(K) \) of the ions. This gives the dipole correlation function in terms of the ion number density \( \rho \) as,

\[
\langle \cos \phi \rangle (r) = \frac{\rho}{(2\pi)^3} \left( \frac{\beta \mu f_0}{12\pi \epsilon_0 \epsilon_r} \right)^2 \times \int |\mathbf{E}(K)|^2 S(K) e^{iK \cdot r} d^3K.
\]

The most appropriate mean-field model can be obtained by taking the field function \( \mathbf{E}(r) \) to be the Coulomb field \( r/r^3 \), corresponding to \( \mathbf{E}(K) = -4\pi iK/K^2 \), and using the Debye-Hückel (DH) structure factor, \( S(K) = \frac{2K^2}{K^2 + \kappa^2} \), where \( \kappa = \left( \frac{2\epsilon_0 \epsilon_r \epsilon_r}{\epsilon_0} \right)^{1/2} \) is the inverse Debye length. This gives

\[
\langle \cos \phi \rangle_{\text{DH}} (r) = \frac{\rho}{2\pi} \left( \frac{\beta \mu f_0}{3\epsilon_0 \epsilon_r} \right)^2 e^{\kappa r}. \tag{7}
\]

The variation of \( \langle \cos \phi \rangle_{\text{DH}} (r) \) with ion concentration is instructive. As seen in Figure 2, for small \( \rho \), an increase in...
concentration leads to an increase in correlation between solvent dipoles, while for large $\rho$, the $e^{-\kappa r}$ factor dominates. Increasing the concentration results in ions being more screened and with a lesser propensity to orient solvent dipoles. It should be also noted that, at all of the concentrations shown in Fig. 2, the dipolar correlations at distances above 5 nm are very small. However, because the number of water molecules further than 5 nm away is very large, these correlations can be measured by fs-EHS experiments, a testimony to the exquisite sensitivity of the probe.

Eq. (6) allows us to investigate the interplay between the ion-ion spatial correlations (encoded in $S(K)$) and the ion-dipole orientational correlations (due to the electric field $E(r)$).

By changing the form of $S(K)$, one can estimate the response to an arbitrary distribution of ions: for instance, one could extend this model to investigate the correlations induced by charges on an interface. A particularly instructive example involves a completely uncorrelated arrangement of ions. This random-ion (RI) model is equivalent to setting $S(K) = 2$, which leads to dipole-dipole correlations corresponding to Eq. (7) with $\kappa = 0$, while the concentration $\rho$ is kept constant. At all concentrations, this RI model leads to increased dipole-dipole correlations because of the lack of screening of the Coulombic ion-dipole interaction by the correlated cloud of counterions. It is worth stressing that although it might be appealing to qualitatively discuss the dampening of correlations in terms of the exponentially screened DH field of an ion, this is not an appropriate model. Such a screened-field/random ions (SF-RI) model amounts to setting $L(r) = -\nabla \phi(r) = r \left( \frac{e^{-\kappa r}}{r^3} + \frac{e^{-\kappa r}}{r^3} \right)$ and $S(K) = 2$. The resulting functional form of the induced dipole-dipole correlations resembles that of the full DH model at short distances but then leads to unphysical anticorrelations at large distance (see the supplementary material).

Fig. 3 compares the predicted $\langle \cos \phi \rangle(r)$ using the full DH theory (Eq. (7)), the RI and the SF-RI models, and the correlations computed from a MD simulation using a ~ 20 nm cubic box with about 264 000 TIP4P/2005 water molecules. All curves correspond to a salt concentration of 8 mM and a temperature of 300 K. The other physical constants used are described in the supplementary material. Comparison with the MD results in Fig. 3 shows that only the full DH model captures the correct long-range behavior of the dipole-dipole correlations—although the short range structure is clearly absent. Neglecting ion-ion spatial correlations artificially increases the orientational correlations, since randomly distributed ions cannot efficiently screen the fields of other ions. A picture in which one interprets dipole-dipole correlations in terms of the screened electrostatic field of the ions, while providing a qualitative picture of the physics, is inconsistent with the linearized-Boltzmann structure of the mean-field model and fails to quantitatively reproduce the MD results. This comparison demonstrates that the long-ranged dipole-dipole correlations are most naturally interpreted as being due to the bare electric field of the ions, inducing (weak) ordering in a spherical domain surrounding each ion. The correlations are modulated by short range interactions (which are not included in this model), and by the presence of ion-ion spatial correlations, which result in a partial cancellation of the field and therefore to exponential damping of $\langle \cos \phi \rangle(r)$. This damping of the dipole-dipole correlations explains the saturation of the fs-EHS signal at high electrolyte concentrations.

We note that the mean-field model can be further improved to include more physical effects. $\langle \cos \phi \rangle_{\text{DH}}(r)$ diverges in the $r \to 0$ limit because of the singularity in the electric field at the ion positions. It is possible to remove this short-distance divergence by restricting the volume of space in which water molecules can be found; however, the fact that two water molecules have a distance of minimum approach, below which $\langle \cos \phi \rangle(r)$ is not meaningful, makes the divergence irrelevant. We can also estimate the impact of neglecting dipole-dipole interactions, by re-introducing them in a perturbative fashion. This can be done by following the procedure used to derive the approximation in Eq. (2), including also the dipole-dipole interaction. In doing so, we find (as described in the supplementary material) that up to order $\beta^4$, the correlation function can be decoupled into two terms, describing the correlations present in bulk water and the changes induced by the electric field of the ions. The lowest-order term in $\langle \cos \phi \rangle(r)$ that includes a coupling between the dipole-dipole and the ion-dipole interactions is proportional to $\beta^4 e^{-\kappa r}/r^7$. This term decays much more rapidly than does the model of Eq. (7) and makes essentially no contribution at long enough distances: above 0.33 nm, the magnitude of this correction is less than 1% of the magnitude of $\langle \cos \phi \rangle_{\text{DH}}(r)$, and less than $10^{-15}$ above 1 nm.

In order to elucidate the free energy scale associated with ion-induced long-range dipole-dipole correlations, we evaluate the total energetic contribution associated with the oriented dipoles at distances larger than a chosen cut-off distance $r_c$ from an ion, which reads (see the supplementary material),

$$U = 4\pi \rho_S \mu \int_{r_c}^{\infty} r^2 E(r) \mathcal{L}(\beta \mu E(r)) \, dr,$$

where $\mathcal{L}(x) = \coth(x) - 1/x$ is the Langevin function and $\rho_S$ is the solvent density. The mean electric field $E(r)$ around the ion is given by the Debye-Hückel theory. The integral can be computed by expanding the integrand as a Taylor series in $\beta$. 

![FIG. 3. Comparison of the solvent dipole correlation function for the full Debye-Hückel theory (black line), the random-ion approximation (red line) and the screened-field plus random-ion approximation (blue line), with a salt concentration of 8 mM at $T = 300$ K. We also show the correlation function calculated from MD at this concentration (green line). In all cases, we have subtracted the correlation function for pure water at the same temperature. Note that this correlation is zero for the mean-field model, and so $\langle \Delta \cos \phi \rangle(r) = \langle \cos \phi \rangle(r)$ for all curves but MD. Inset: the absolute correlation function $\langle \cos \phi \rangle(r)$ calculated from MD, showing considerable structure at short range.](image-url)
Fig. 4 shows the total energetic contribution of the dipoles oriented by an ion as a function of the electrolyte concentration and for different cut-off lengths. At mM concentrations, dipolar order beyond three solvation shells (∼0.8 nm) is associated with an energy scale of about $3 k_B T$, and even the tails beyond 4 nm correspond to a significant fraction of $k_B T$. Thus, even though each ion-dipole interaction is very small, the collective effect can be significant when considering the energetics of the ion that induces the orientation of the dipoles. Indeed, field-theoretical calculations on ionic solutions have shown that their surface tension can be written as a sum of two terms, one describing the interactions of ions with their image charges, and another which describes the energy stored in the dielectric medium surrounding these ions, which is analogous to the energy of dipole orientation described in Eq. (8). These arguments support the hypothesis that ion-induced dipole-dipole correlations underlie the concentration dependence of both fs-ESHS and surface tension measurements for dilute electrolyte solutions.

In conclusion, we have shown that long-range, non-specific electrolyte-induced correlations in water as recently observed in fs-ESHS experiments can be captured by a simple mean-field model that treats water molecules as non-interacting dipoles oriented by the electrostatic field of ions, which are themselves correlated following the Debye-Hückel theory. Although one can intuitively understand the orientational correlations as arising from the exponentially screened field of correlated ions, a more accurate picture, leading to quantitative predictions of MD simulations, regards them as arising from unscreened ion-dipole correlations that combine destructively when the physically relevant ion-ion correlations are included. This model is very useful to pinpoint what we think is the main physical origin of the electrolyte-induced change in the fs-ESHS intensity and to estimate the length and energy scale of the effect. It does not, however, explain the dramatic isotope effects that are seen in experiments, or the temperature dependence of the fs-ESHS signal. As such it is clearly only a first step in a complete description of the experimental data, which should also include a re-evaluation of the molecular hyperpolarizability tensor, particularly when probed by femtosecond laser pulses.

See supplementary material for more detailed derivations of the formulas used in the main text, as well as a list of the numerical values of physical constants used.

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