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# Electric Double Layer Composed of an Antagonistic Salt in an Aqueous Mixture: Local Charge Separation and Surface Phase Transition

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**Venue: S4-625**

**Time: 12:00-13:00**

**Abstract:** We examine an electric double layer containing an antagonistic salt in an aqueous mixture, where the cations are small and hydrophilic but the anions are large and hydrophobic. In this situation, a strong coupling arises between the charge density and the solvent composition. As a result, the anions are trapped in an oil-rich adsorption layer on a hydrophobic wall. We then vary the surface charge density  $\sigma$  on the wall. For  $\sigma > 0$  the anions remain accumulated, but for  $\sigma < 0$  the cations are attracted to the wall with increasing  $|\sigma|$ . Furthermore, the electric potential drop  $\Psi(\sigma)$  is nonmonotonic when the solvent interaction parameter  $\chi(T)$  exceeds a critical value  $\chi_c$  determined by the composition and the ion density in the bulk. This leads to a first-order phase transition between two kinds of electric double layers with different  $\sigma$  and common  $\Psi$ . In equilibrium such two-layer regions can coexist. The steric effect due to finite ion sizes is crucial in these phenomena.