

Diffusion in colloidal monolayers: bridging the gap between two and three spatial dimensions

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It is well established that, unlike for a three-dimensional fluid, particle interactions prevent the hydrodynamic transport coefficients from being defined for a two-dimensional fluid due to the notorious “long-time tail” feature of the velocity autocorrelation.

A colloidal monolayer formed at a fluid interface builds a bridge between these two limiting cases, and it provides insight on the transition from three down to two spatial dimensions: the positions of the colloidal particles are constrained to a plane and the colloid thus resembles a two-dimensional fluid. But the exchange of particle momentum takes place in three-dimensional space because it is mediated by the ambient fluid in the form of hydrodynamic interactions.

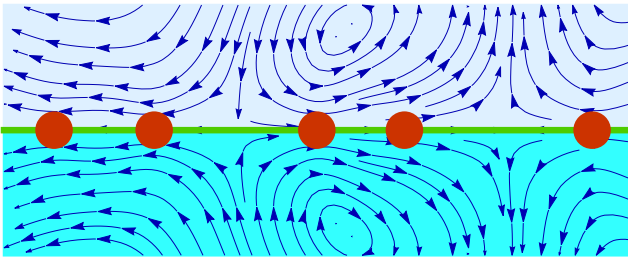


Fig. 1. Side view of a colloidal monolayer formed at the interface between two fluids, and sketch of the three-dimensional flows responsible for the hydrodynamic interactions.

Here we study the behavior of the colloidal diffusivity, which is the only hydrodynamic transport coefficient for the

two-dimensional colloidal fluid. The starting point is the Smoluchowski equation, i.e., the Fokker-Planck equation for the colloidal particles in the overdamped regime with due account of the hydrodynamic interactions. We show that the diffusivity exhibits an intermediate behavior between purely two-dimensional and fully three-dimensional fluid: on the one hand, Fick’s law, which pertains to *collective diffusion*, breaks down altogether [1, 2, 3], as confirmed experimentally [4]. On the other hand, the coefficient of *self-diffusion* (or single-particle diffusion) is finite [5], but the transitional nature of the monolayer shows up in a non-analytic dependence on the colloidal packing fraction [6], at odds with the case of a fully three-dimensional colloid.

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