

Phase diagram for a copolymer in a micro-emulsion

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Abstract

In this paper we study a model describing a copolymer in a micro-emulsion. The copolymer consists of a random concatenation of hydrophobic and hydrophilic monomers, the micro-emulsion consists of large blocks of oil and water arranged in a percolation-type fashion. The interaction Hamiltonian assigns energy $-\alpha$ to hydrophobic monomers in oil and energy $-\beta$ to hydrophilic monomers in water, where α, β are parameters that without loss of generality are taken to lie in the cone $\{(\alpha, \beta) \in \mathbb{R}^2: \alpha \geq |\beta|\}$. Depending on the values of these parameters, the copolymer either stays close to the oil-water interface (localization) or wanders off into the oil and/or the water (delocalization). We derive two variational formulas for the quenched free energy per monomer, one that is ‘‘column-based’’ and one that is ‘‘slope-based’’. Using these variational formulas we identify the phase diagram in the (α, β) -cone. There are two regimes: *supercritical* (the oil blocks percolate) and *subcritical* (the oil blocks do not percolate). The supercritical and the subcritical phase diagram each have two localized phases and two delocalized phases, separated by four critical curves meeting at a quadruple critical point. The different phases correspond to the different ways in which the copolymer can move through the micro-emulsion.

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Key words and phrases. Random copolymer, random micro-emulsion, free energy, percolation, variational formula, large deviations, concentration of measure.

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Remark: The part of this paper dealing with the ‘‘column-based’’ variational formula for the free energy has appeared as a preprint on the mathematics archive: arXiv:1204.1234.

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