Photo-triggered assemblies of amphi philic copolymers with and with no surfactants: experimental studies and model predicting the response to light Chair: Prof. Françoise Winnik(MANA Satellite PI)



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Amphiphilic copolymers form in aqueous solutions co-assemblies, for instance with micelles of surfactants, lipids. They also form selfassemblies upon transferring a fraction of their hydrophobic groups from water into apolar clusters. The energy balance between attractive hydrophobic binding and intra/inter chain repulsions controls these assemblies, that are used since decades to tailor properties of complex fluids, for instance emulsions, physical gels, dispersion of cargoe-particles for drug delivery, etc... In practice, the hydrophobic drive toward (self-)assembly can be modulated at the microscopic level upon variation of control parameters (pH, temperature, or exposure to stimuli: e.g. light), which in turn provide remote control of macroscopic properties.

We tailor azobenzene-modified copolymers to achieve marked photo-switch in such assemblies. Experimental illustrations include mixed micellization with surfactants, and photo-triggered permeabilization of lipid and plasma membrane of mammal cells. To analyze experimental data on the degree of hydrophobic binding of polymer chains, we propose a model based solely on one parameter: a critical hydrophobicity of a (bound) polymer segment. This model is shown to reproduce the measured fractions of micelle-bound hydrophobes (here azobenzene) as a function of density of hydrophobic groups per chain. Non-monotonic variation of LCST are also predicted and experimentally evidenced. The model enables us to discuss a bottom up approach to the design of systems having optimal responses to light.

## Venue: Seminar Room #431, 4F, MANA Bldg., Date: May 7<sup>th</sup>, Monday Time: 15:00-15:45

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