

Light induced motion manipulation of micro-sized colloids in external flow - aggregation, fractioning and analysis

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All in one: What connects the interfacial sensitive fractioning and population-based interface analysis of micro-sized colloids between the mimicking blood clogging with inorganic particles. All those features are possible by light induced motion manipulation of micro-sized colloids in external fluid flows via microfluidics. This is possible with a photosensitive azobenzene containing surfactant, where the more surface-active *trans*-isomer adsorbs stronger on interfaces in comparison to the less surface active *cis*-isomer counterpart. This yields under illumination into dynamic exchange between *trans*- and *cis*-isomers making interfaces upon illumination phoretically/osmotically active – its phoretic strength is proportional to illumination parameters and more important also to interfacial properties.

In this sense sedimented colloids can be brought into a vertical displacement from solid-liquid interfaces. We show that this control of light induced chemical activity yields either to model representative of controlled blood coagulation for inorganic particles[1] or into simple and versatile population-based analysis[2] and fractioning of equally sized particles[3,4] varying only by their interfacial properties (porosity, material, surface coating or surface charge).

Keywords:

active colloids, photosensitive surfactants, interface based fractionation, population based porosity analysis, aggregation, microfluidics

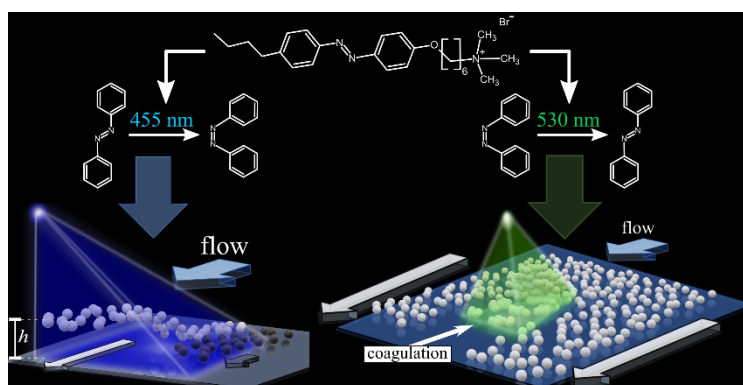


Figure 1. Light-induced control of microscale colloids with azobenzene surfactants. Blue light induces particle levitation and altered motion, enabling fractionation[3,4] and population-based porosity analysis [2]. Green light induces aggregation, causing localized, remotely controlled clogging.

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