

Optically trapped plasmonic nanoparticles for manipulating model lipid bilayers and biological membranes

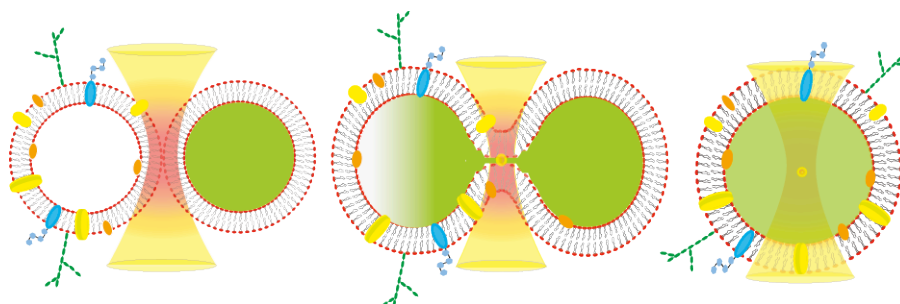
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Abstract

We use optically trapped plasmonic nano-heaters to facilitate efficient and controlled local heating of both model membranes and membranes of living cells. The associated effects on membranes phase behavior, membrane permeability and membrane morphology are visualized by confocal microscopy. Local heating near a gel phase membrane generates a phase transition together with an associated increased permeability of the membrane¹⁻⁴. Trapping a plasmonic nanoparticle at an interface between two Giant Unilamellar Vesicles (GUVs) results in immediate full fusion of the apposing membranes which we verify by measuring lipid and content mixing⁵. Fusion also occurs when trapping a plasmonic nanoparticle between two cells to transform the cells into a single viable syncytium⁶. Other fusion products realized with this method include a GUV fused to a cell or a GUV fused to a giant plasma membrane vesicle (GPMV). We show how this assay can be used in biophysical research; Membrane proteins encapsulated in an inert vesicle are transferred to a vesicle composed of negative lipids by optically inducing fusion. Mixing of the two membranes results in a fused vesicle with a high affinity for the protein and we observe immediate membrane tubulation due to the tubulating activity of the protein^{5,7}. Fusion of distinct membrane compartments has interesting applications in small scale chemistry for realizing pico-liter reactions and offers many exciting applications within biology which are discussed here.



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