



We study soft-matter physics and biophysics theoretically and numerically. Our main target is physics of biomembrane and cells under various conditions. There are many interesting shape transitions and dynamic behaviors. We develop membrane models and hydrodynamic simulation methods.

Dynamics of red blood cells and lipid vesicles in flow

Red blood cell in capillary flow



Solvent-free bilayer

coarse-graining

no explicit solvent -> effective attractive potential between hydrophobic segments

We found several pathways

of membrane fusion and fission.

membrane model

THEFT





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nerecity

Red blood cells in glass capillary tsukada et al. Microvasc. Res.61, 231 (2001).

fission

0.5 Small shear Small shear

Lipid vesicle in simple shear flow

time t/τ Stomatocyte to prolate shape. Shear also induces elongational and shrinking transitions between discocyte and prolate.

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Slow flow: discocyte Fast flow: parachute

The transition velocity linearly depends on the membrane bending rigidity and shear elasticity. On the other hand, lipid vesicles elongate to prolate shape.

simulations by MPC-SR with dynamically-triangulated membrane model

Membrane fusion and fission

one hydrophilic

segment

two hydrophobic segments



Fusion mediated by particle adhesion



New pathway; Fusion occurs via pore-opening beside stalk intermediate

• Meshless membrane model
• Disparticle represents, appl(ρ) + ε (Σ Upp(ρ(ρ) + Σ Upp(ρ(ρ))))
• α_p(ρ): deviation from plane
• Bending rigidity and line tension
• Stepsembly
• Output of the tension
• Output of tension
• Output

Hydrodynamic simulation methods



We proposed the intermediate models (DPD-MT and MPC-LD) and clarify the relations between DPD (dissipative particle dynamics) and MPC (multiparticle collision dynamics). We also clarified artifacts when the angular momentum is not conserved.