



Institute of Physical Chemistry, BAS
Sofia, Bulgaria

Study of Foam Film Drainage via micro-interferometric techniques

Plamen Tchoukov, Elena Mileva, Dotchi Exerowa

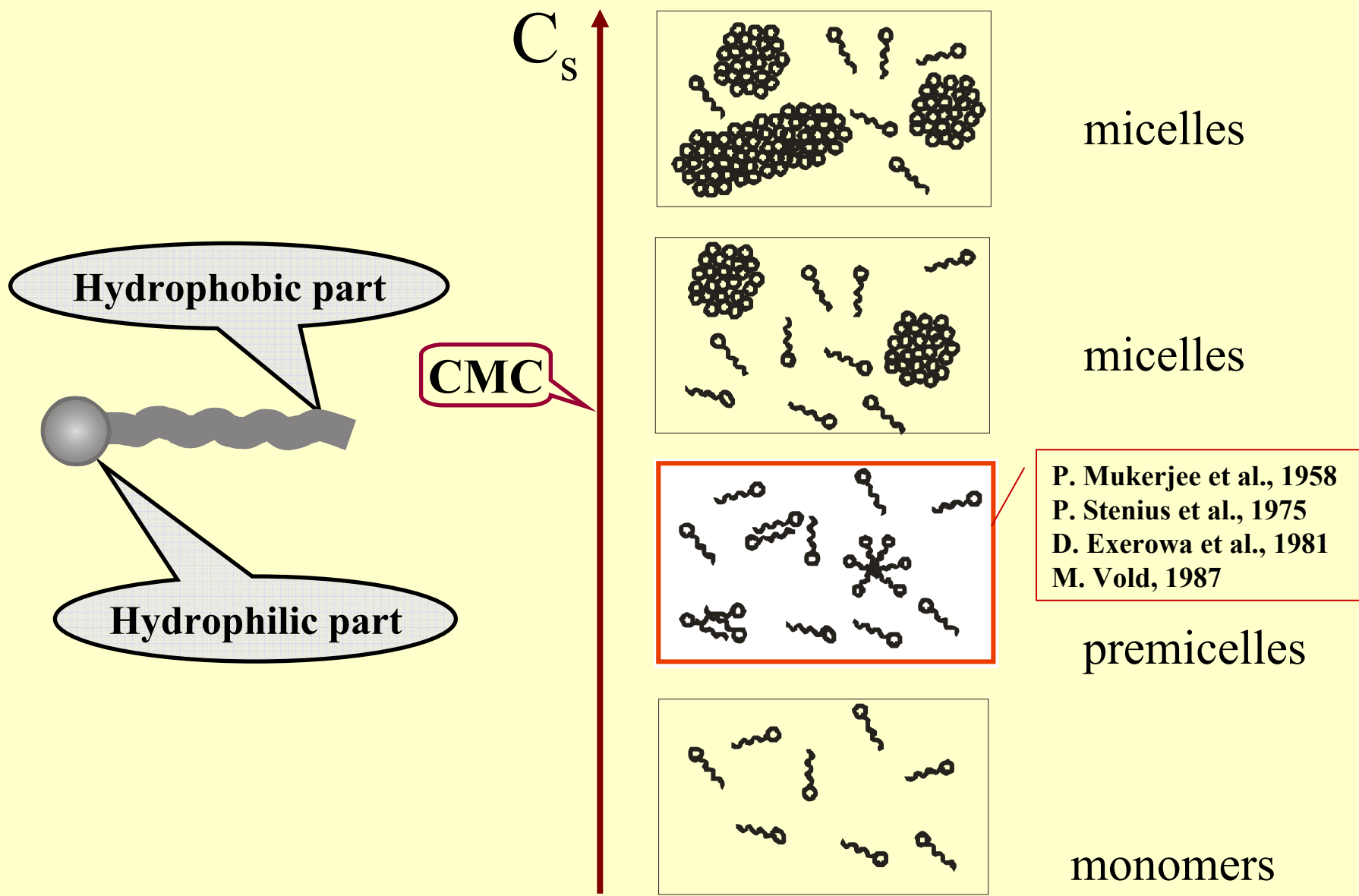
Partially supported by:

Contract with EC "Nanoscale Phenomena and Structures in Bulk and Surface Phases" (NANOPHEN, № INCO-CT-2005-016696)

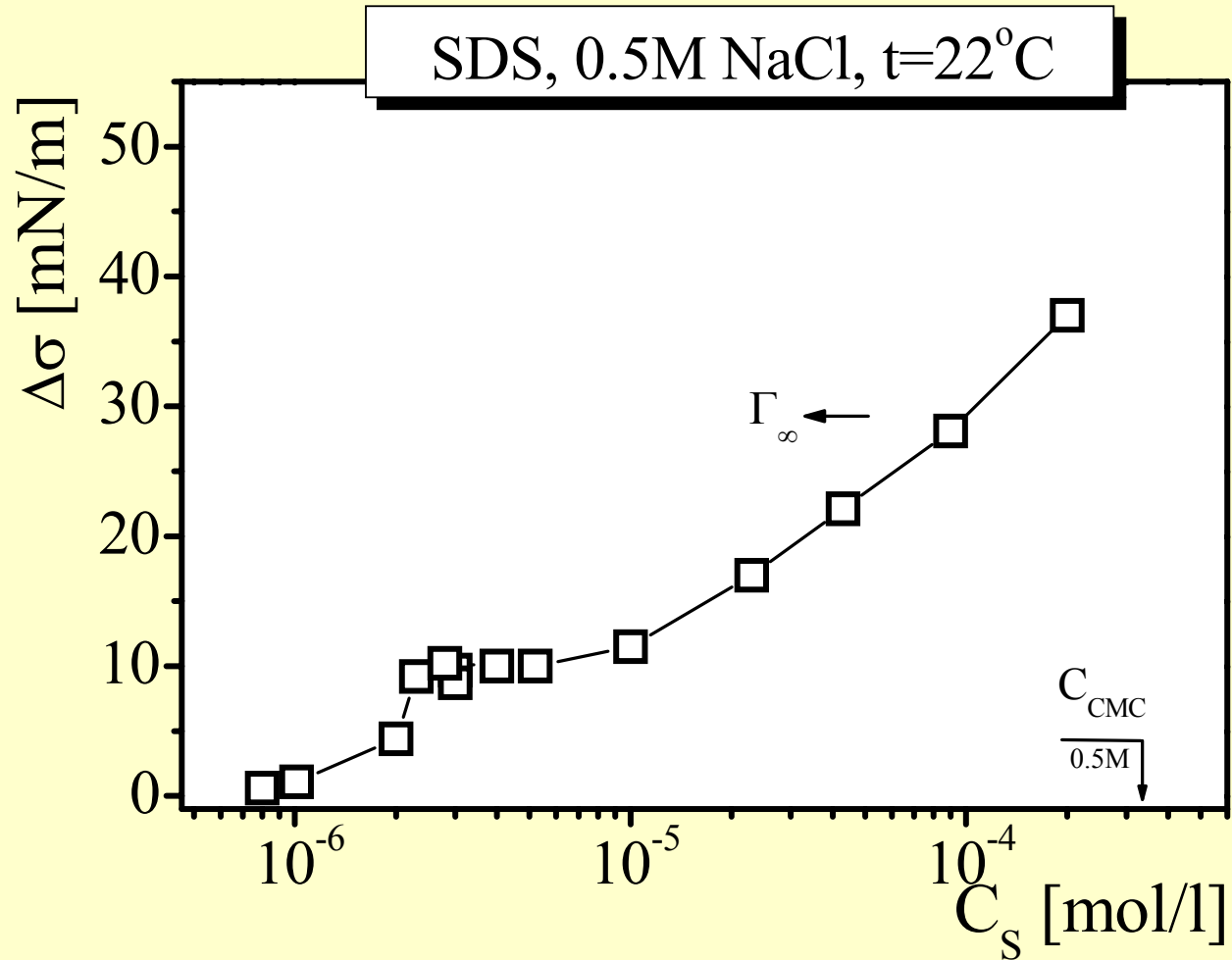
Contract with NSF „Colloid Aspects Of Nano-Science: Nanostructures At Interfaces, In Bulk And In Three-Phase-Contact Zone”, module “Amphiphilic nanostructures in fluid media”

Aim

*To outline the impact of
amphiphilic self-assembled
structures on the drainage of
microscopic foam films from
aqueous surfactant solutions*

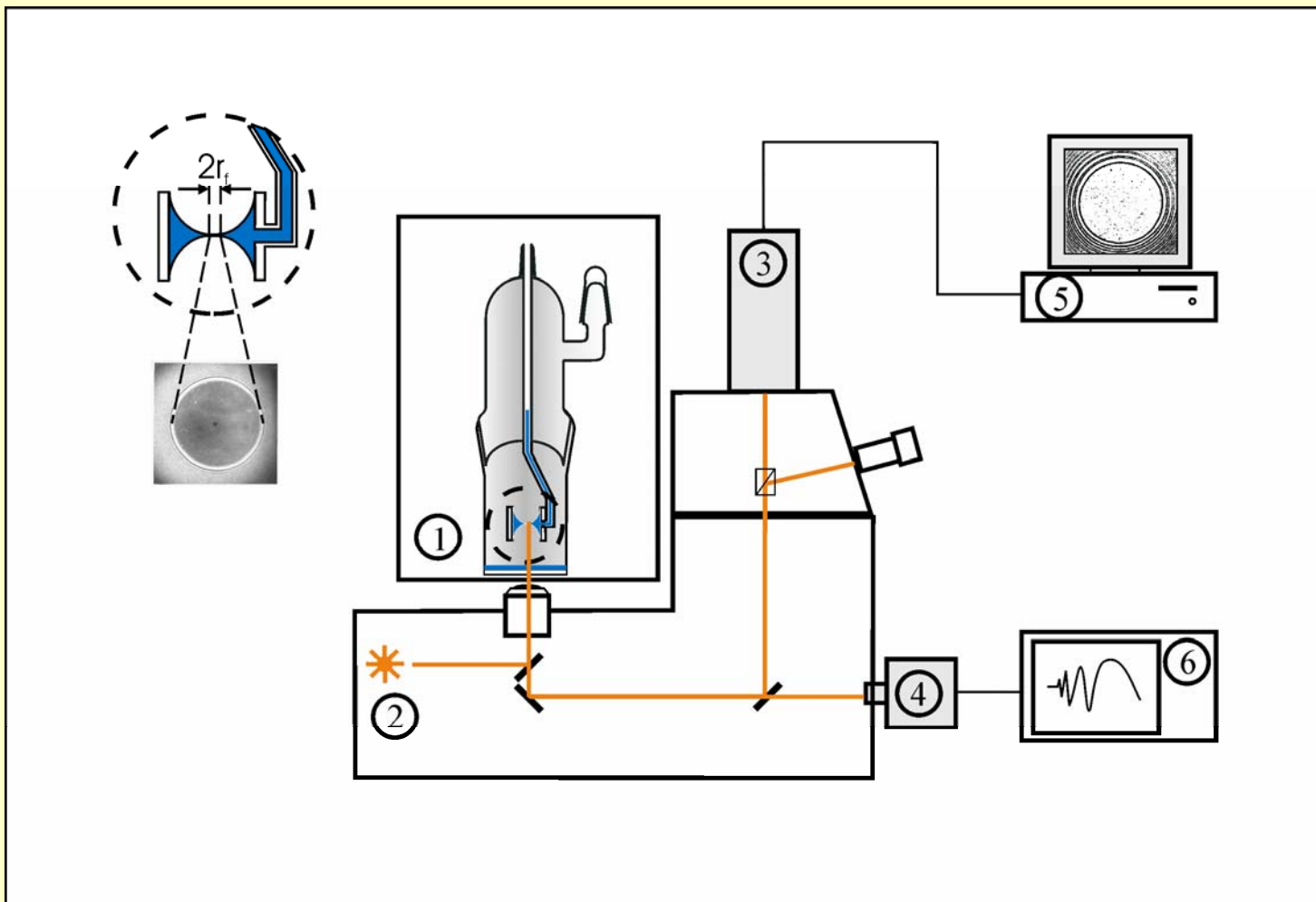


Motivation



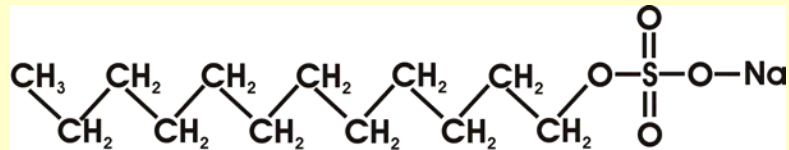
Exerowa et al., *J.Coll.Interf.Sci.*,1981,81.

Experimental setup



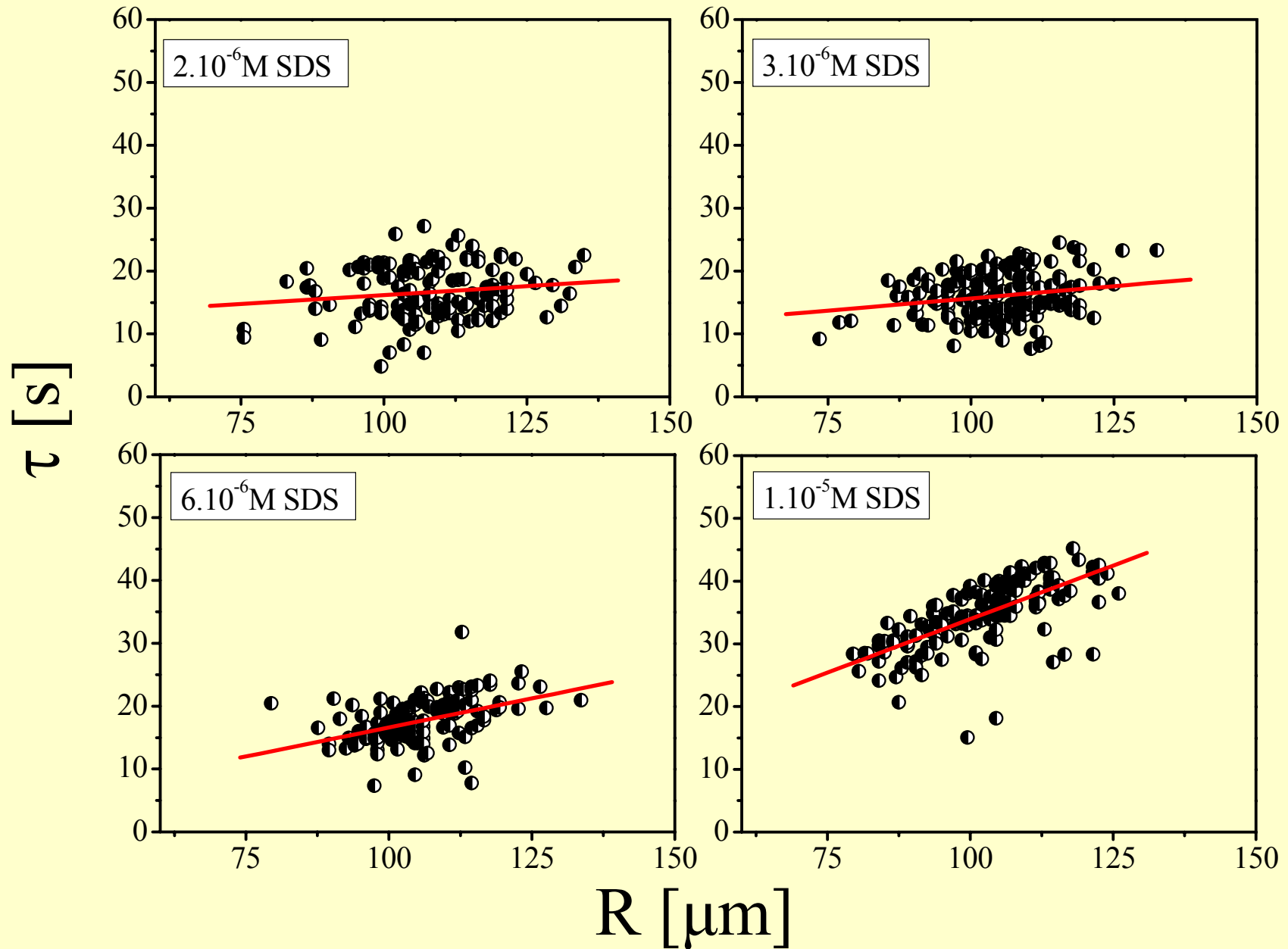
Experimental conditions

- Anionic surfactant SDS

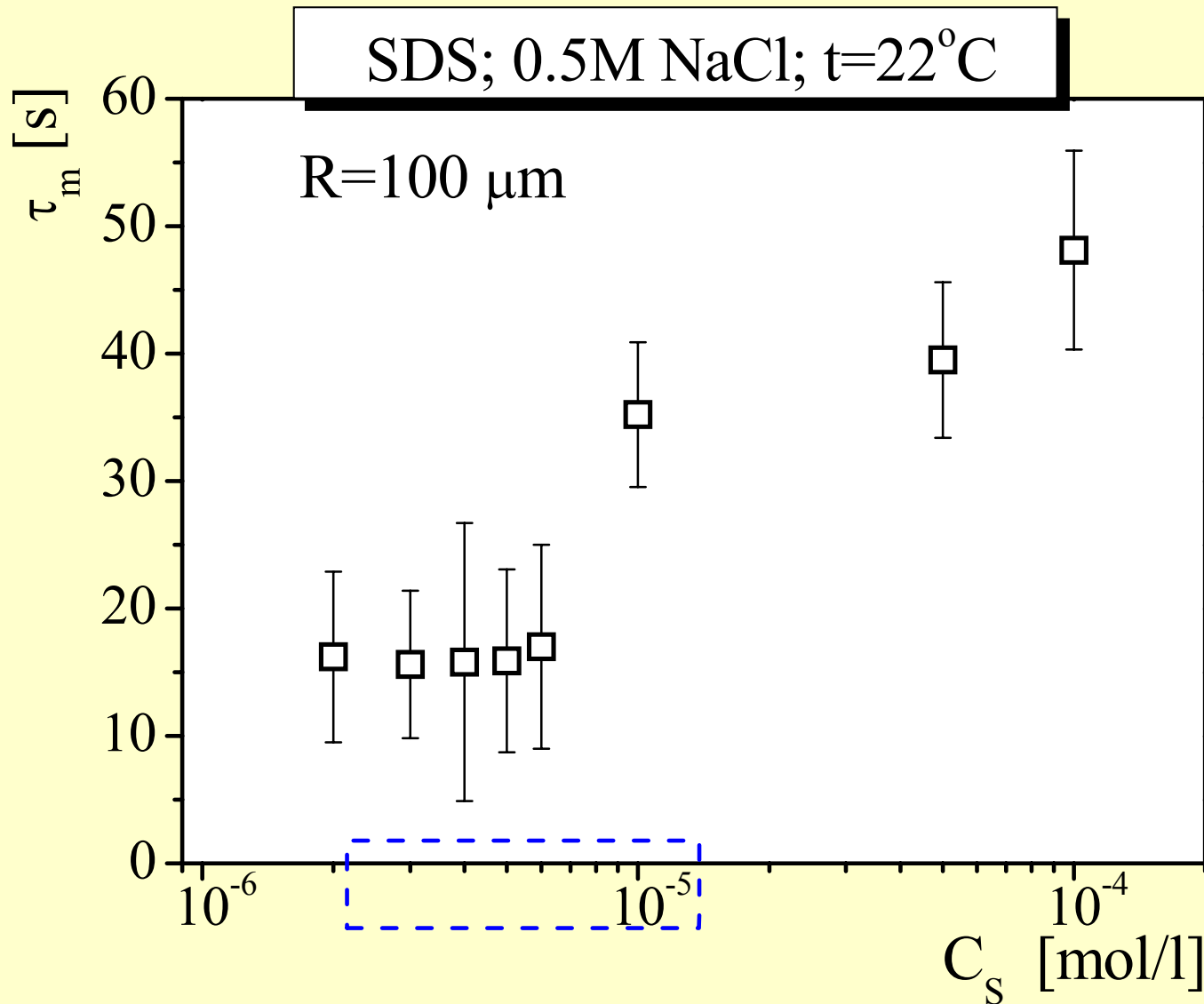


- Surfactant Concentration:
 - ✓ $2 \times 10^{-6} \div 10^{-4}$ mol/l (0.5M NaCl)
- Electrolyte concentration :
 - ✓ $0.5\text{M NaCl} > C_{\text{el,cr}}$

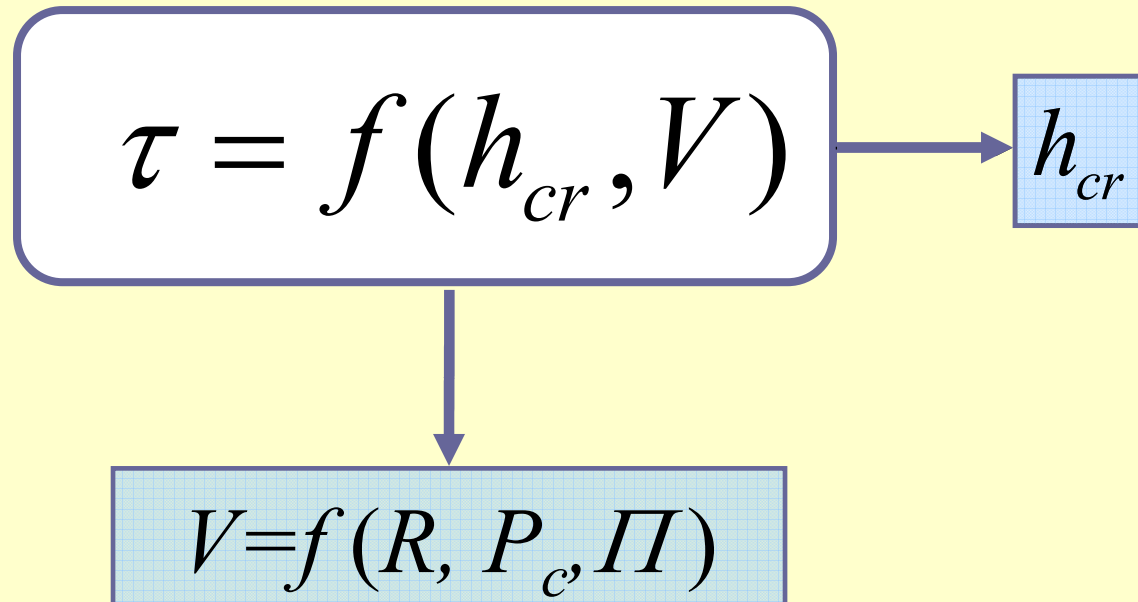
SDS, 0.5M NaCl, $t=22^{\circ}\text{C}$



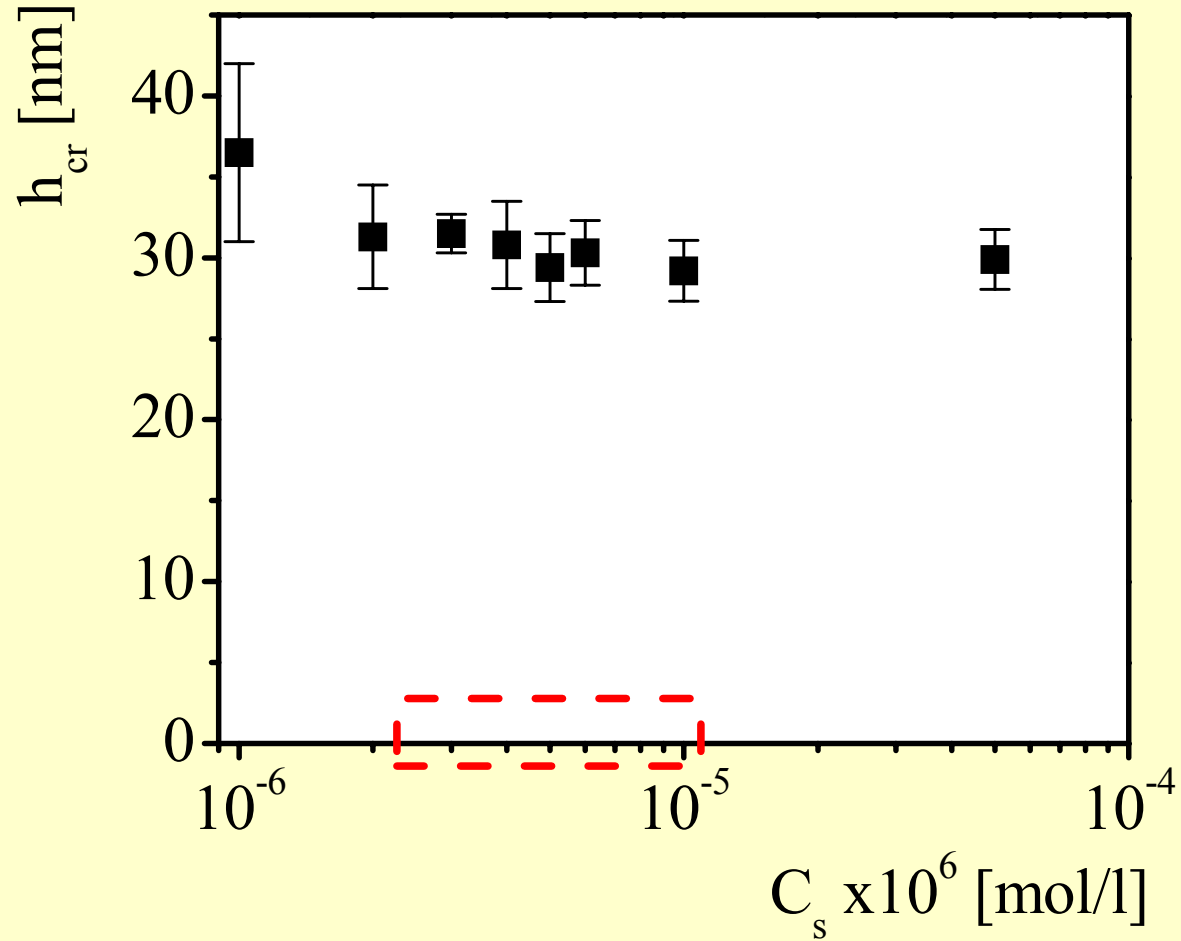
Foam film kinetics: drainage time



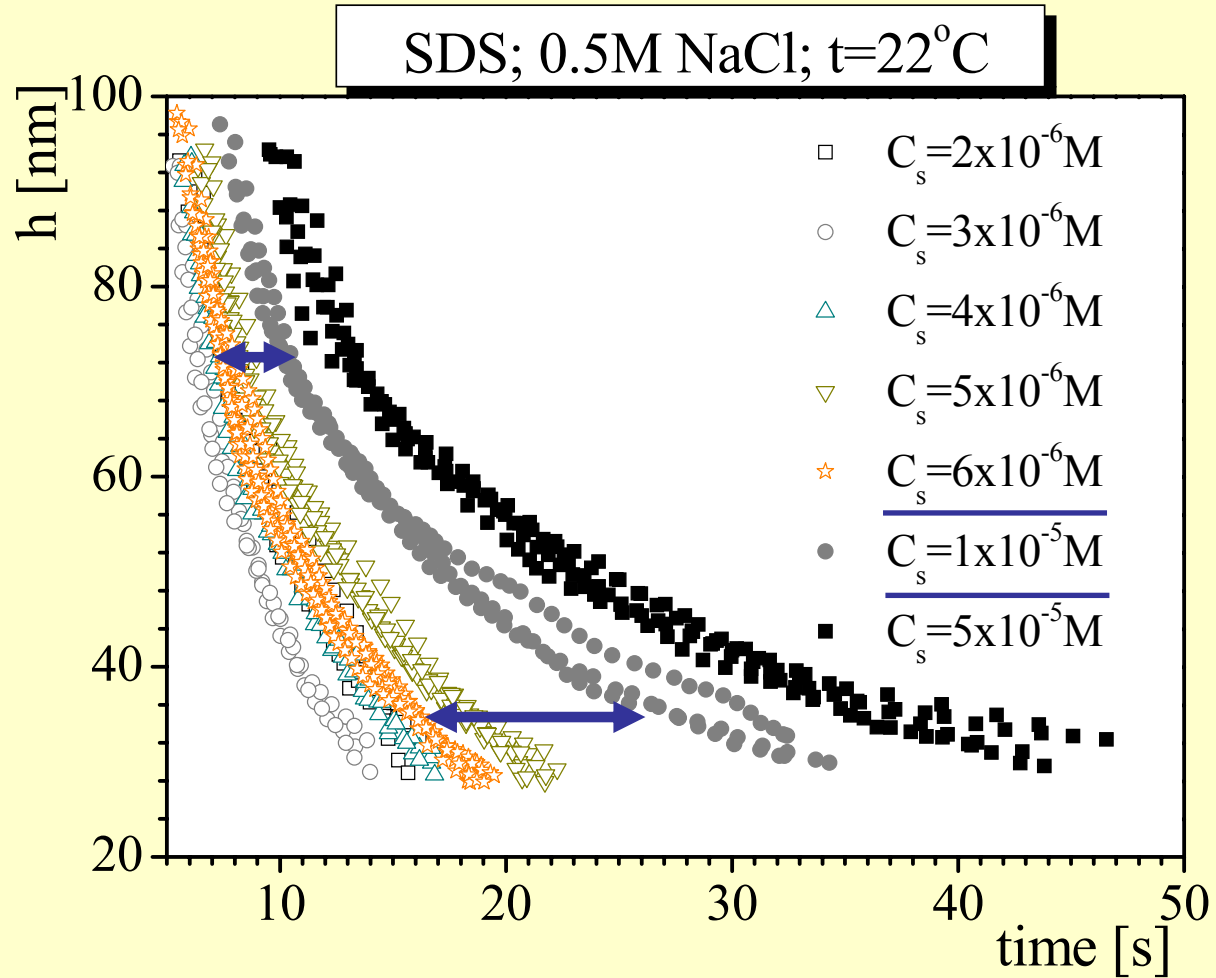
Time? What does it depend on?



What does time depend on? - critical film thickness



Foam film kinetics: $h(t)$



Sheludko,
Adv.Coll.&Interface Sci., 1, 391,1967

$$V_{Re} = \frac{2h^3 \Delta P}{3\eta R^2}$$

$$\Delta P = P_c - \Pi \quad \eta = 1.0 \times 10^{-3} \text{ (Pa.s)}$$

Radoev, Dimitrov, Ivanov,
Coll&polymer Sci., 252, 50, 1974

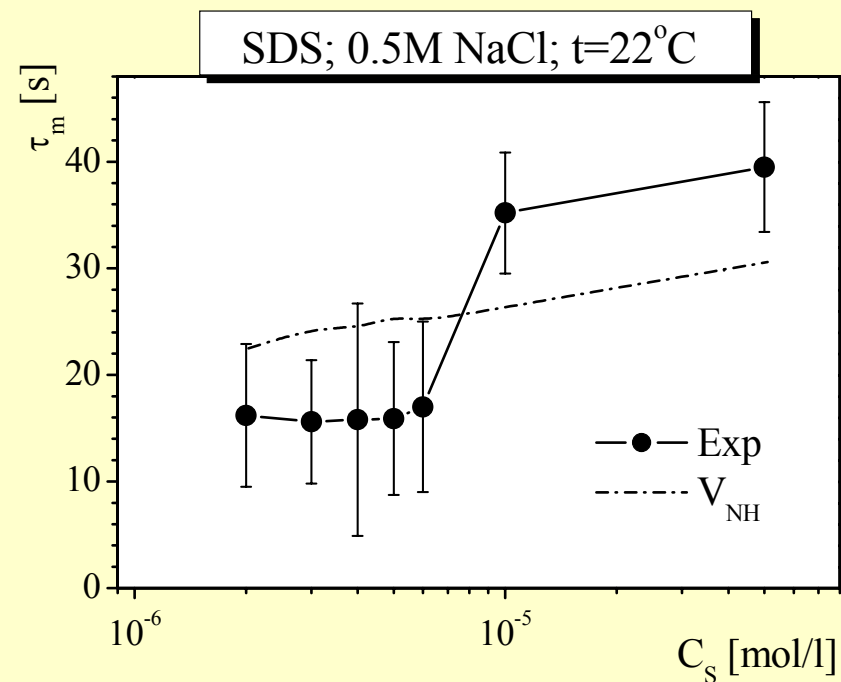
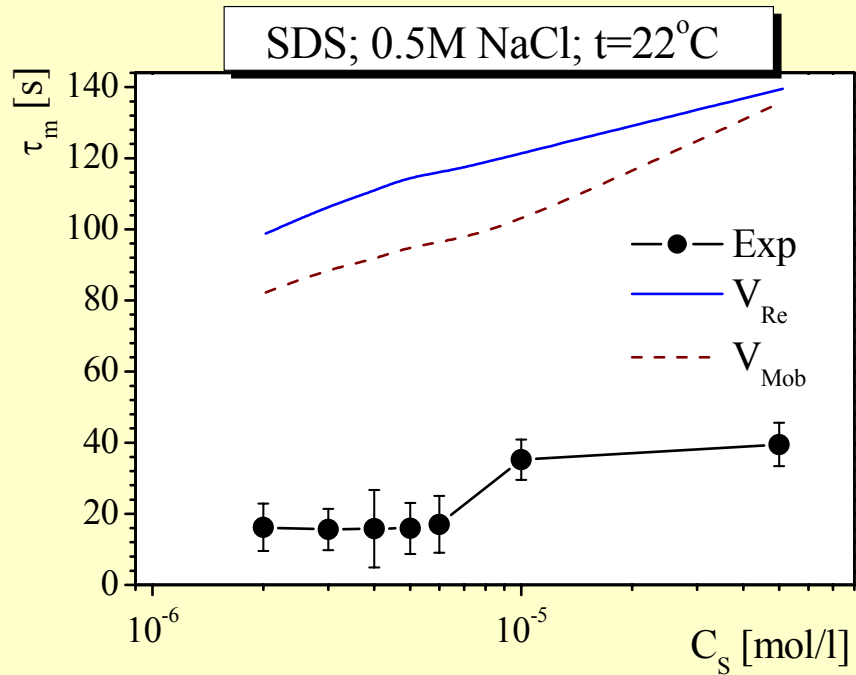
$$\tau = \int_{h_0}^{h_{cr}} \frac{dh}{V}$$

$$\frac{V_{Mob}}{V_{Re}} = 1 + b + \frac{h_s}{h}$$

$$b = -\frac{3\eta D}{\Gamma_0 \left(\frac{\partial \sigma_0}{\partial c_0} \right)} \quad h_s = -\frac{6\eta D_s \left(\frac{\partial \Gamma_0}{\partial c_0} \right)}{\Gamma_0 \left(\frac{\partial \sigma_0}{\partial c_0} \right)}$$

$$V_{NH} = \frac{1}{6\eta} \sqrt[5]{\frac{h^{12} \Delta P^8}{4\sigma^3 R^4}}$$

Manev, Tsekov, Radoev,
J.Disp. Sci.&Technology, 18, 769, 1997



Our interpretation

- Amphiphilic structures (premicelles) exist in initial surfactant solutions
- Under the action of negative (van der Waals) disjoining pressure, the surfactant aggregates disintegrate, and the size distribution curve is shifted to the monomers.

Mileva E, Exerowa D, Colloids Surf. A 149, 207, (1999)

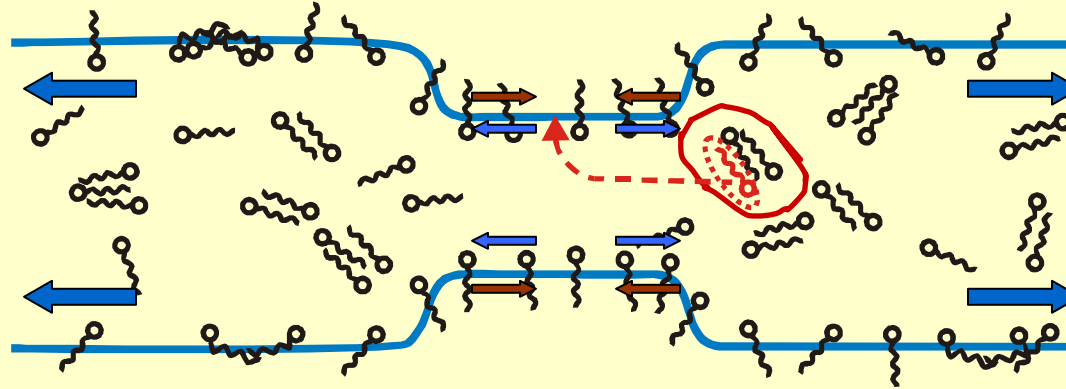
$$\Pi(h) = \Pi_{vw}(h)$$



$$X_n(h < h_d) < X_n(h_d)$$

- Films drain in a regime of high interfacial mobility and thickness inhomogeneities

Our interpretation



$$P_{n\tau} = \nabla_{\tau} \sigma \quad \Rightarrow \quad \mu^f \frac{\Delta U}{H^f} \sim \left| \frac{\partial \sigma}{\partial c^f} \right| \frac{\Delta c_{\tau}^f}{R^f}$$

$$\nabla_{\tau} (\Gamma v_{\tau}^f) = j_n^f \quad \Rightarrow \quad j_n^f \sim D^f \frac{\Delta c_n^f}{H^f} \sim \epsilon j_{\tau}^f$$

$$\Delta c_n^f = c^f - c_s^f, \quad \Delta c_{\tau}^f = c^m - c_s^f$$

$$\mu \frac{\Delta U}{H} \sim \left| \frac{\partial \sigma}{\partial c} \right| \frac{\Delta c_{\tau} + c_{agg}}{R}$$

Mileva, Radoev, (1993)

$$c_{new}^m = c^m + c_{agg}$$

To summarize:

- ✓ Concentration coincidence of surface tension and the film drainage results is due to the presence of **amphiphilic structures** as coupled with the **local hydrodynamics** and **mass transfer of surfactant** molecules.