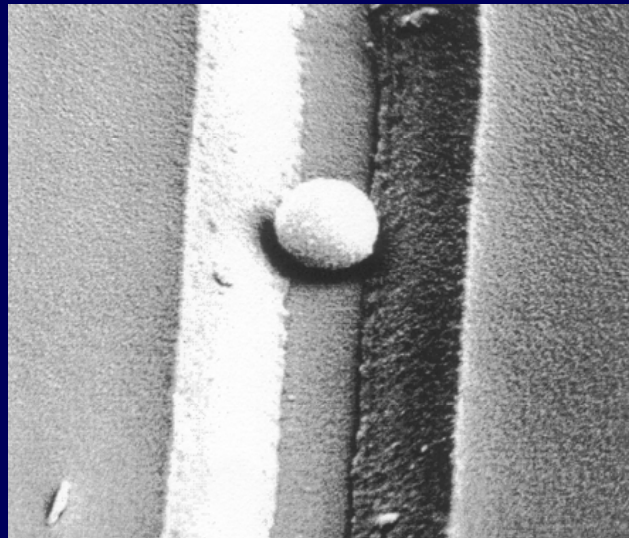
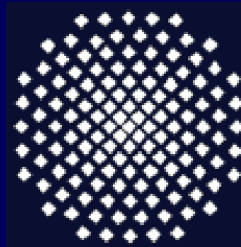


Diffusion in Reduced Dimensions

Clemens Bechinger

2. Physikalisches Institut, Universität Stuttgart

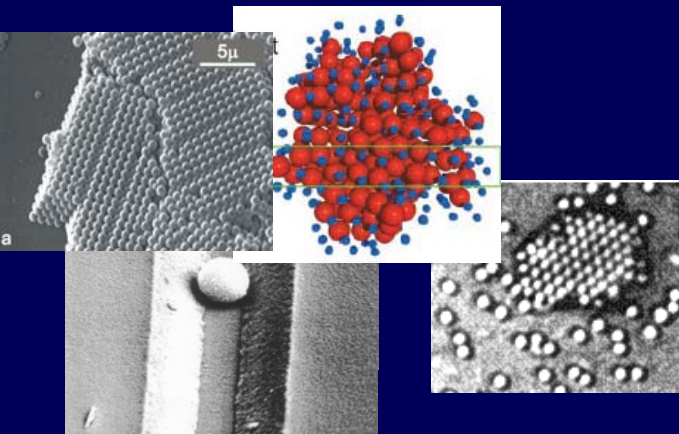
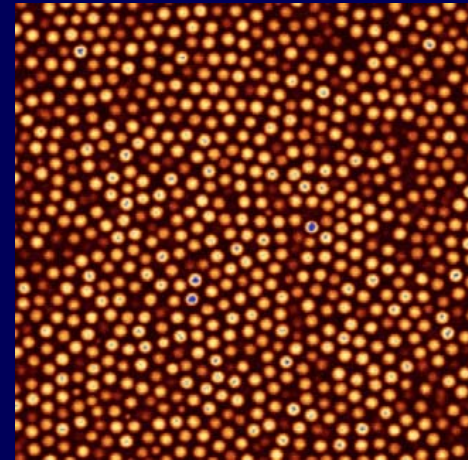


Colloids As Model Systems

Colloids: = Solid particles ranging between 10 nm and 10 μm dispersed in liquids

- overdamped motion: → **slow**
- very small elastic constants $\approx \frac{k_B T}{a^3}$ → **soft**
- typical length scales \approx visible light: → **seeable**

“... the same equations have the same solutions ...” (R. Feynman)



Atoms

$u(r)_{\text{atom}}$

\equiv

Colloids

$u(r)_{\text{colloid}}$

- Phase transitions
- Nucleation phenomena
- Glass formation
- Diffusion through pores and channels
- ...

Colloids are „giant“ atoms

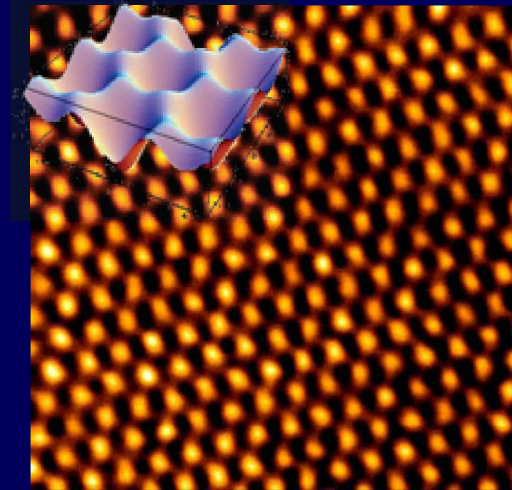
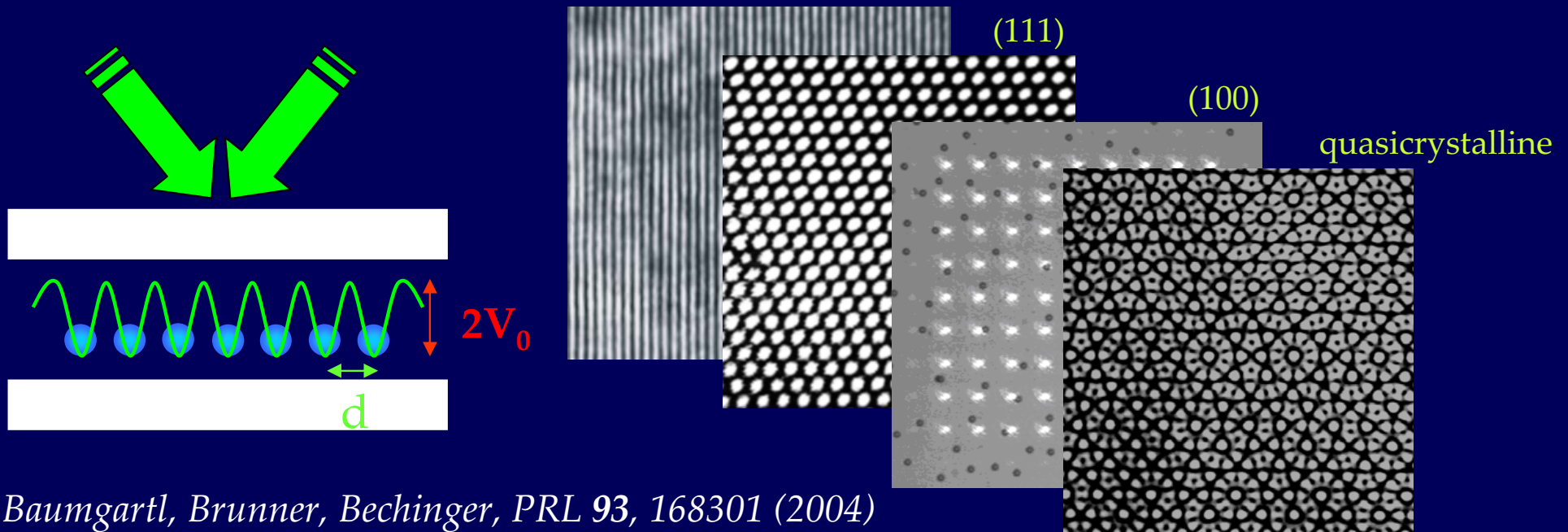
Epitaxial growth of monolayers

morphology depends on

- lattice mismatch
- substrate strength
- adsorbate-adsorbate interactions

parameters difficult to vary in atomic systems

→ colloids on light-induced substrate potentials

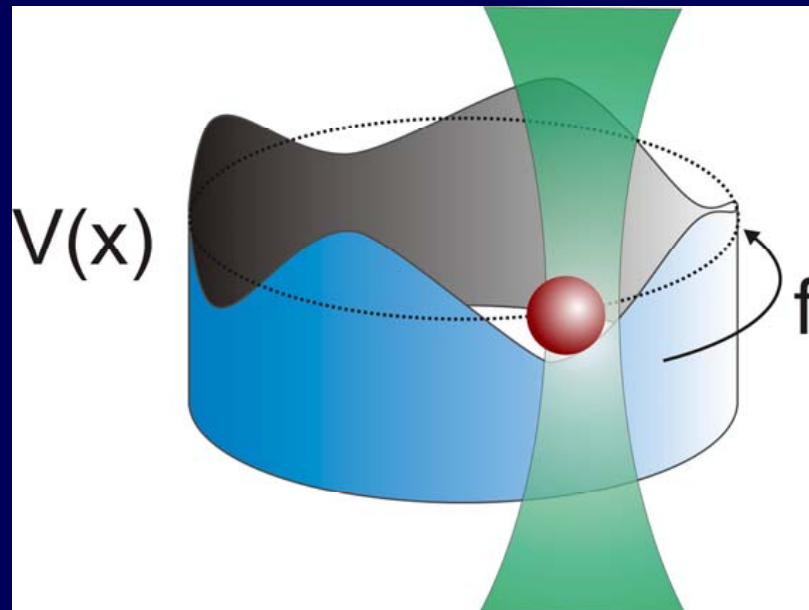


Stochastic Thermodynamics

Driven colloidal systems beyond linear response

Generalized Einstein relation

$$D = k_B T \mu + \int_0^{\infty} d\tau I(\tau)$$

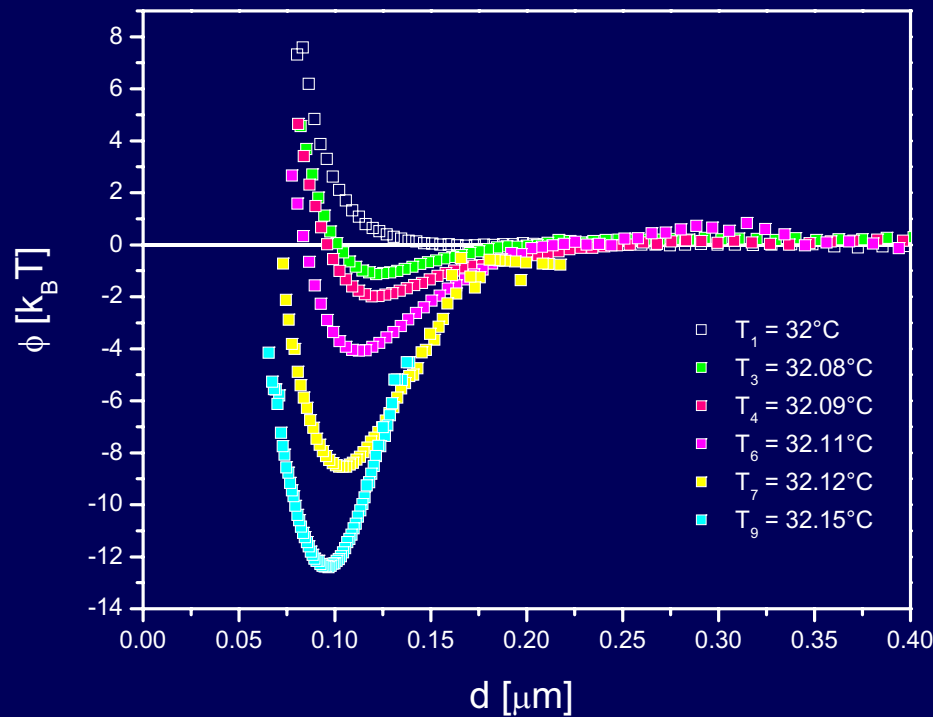
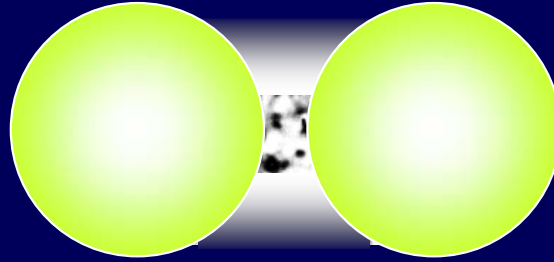


Blickle, Speck, Lutz, Seifert, Bechinger, PRL 98, 210601 (2007)

Critical Casimir Forces

Confinement of critical fluctuations in binary liquid mixtures near T_c

→ long-ranged forces



water/lutidin

$\eta_c = 29\%$

$T_c = 32^\circ\text{C}$

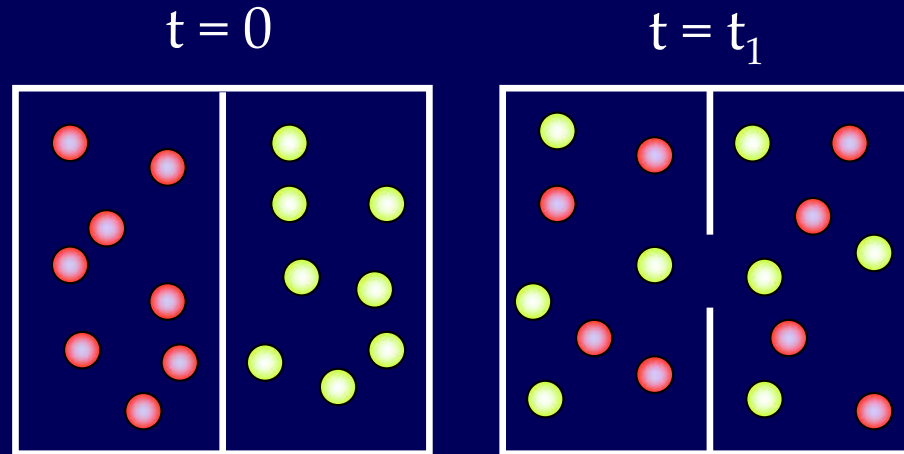
Single-file Diffusion of colloids

Experiments: *Q.-H. Wei, C. Lutz*

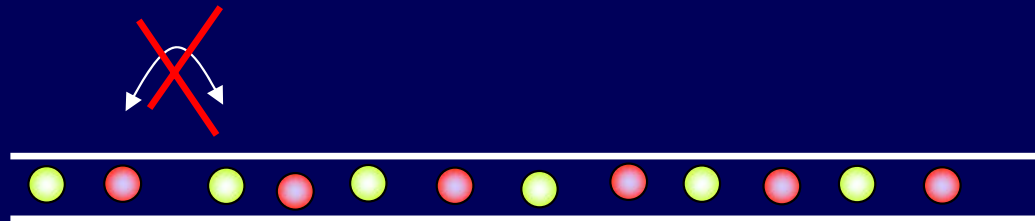
Theorie: *M. Kollmann*

Diffusion in narrow Channels

3D, 2D: mixing



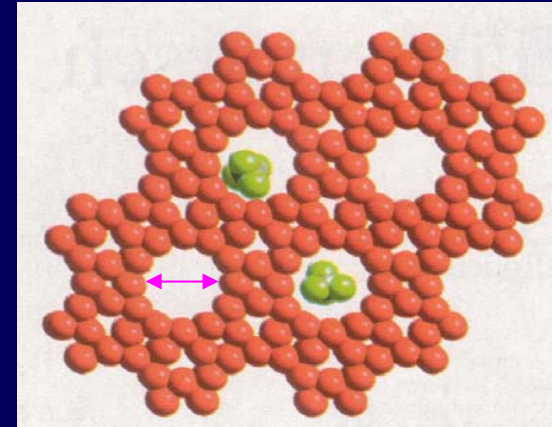
1D: sequence unchanged



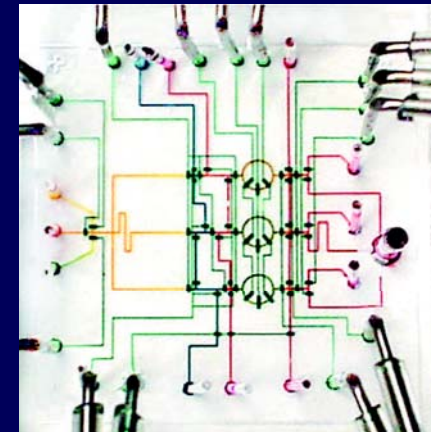
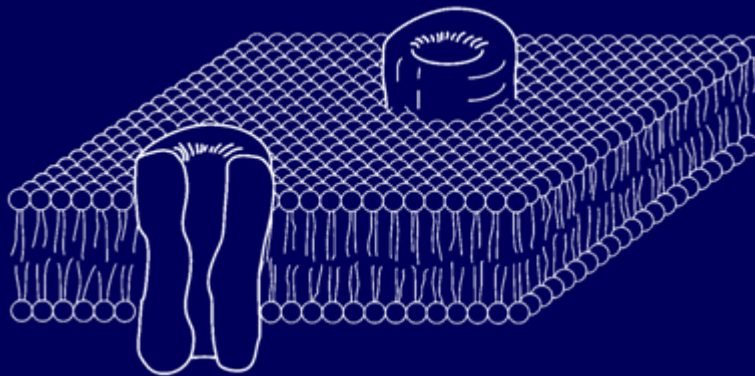
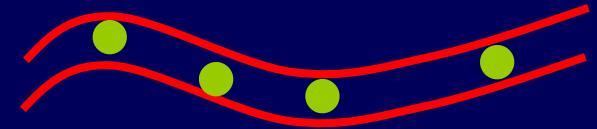
1D diffusion entirely different

Realization of SF conditions

- molecular sieves (zeolites)
- carbon nanotubes
- ionic transport through membranes
- reptation in polymer melts
- microfluidic devices
- ...



0.73nm



Single-File Diffusion

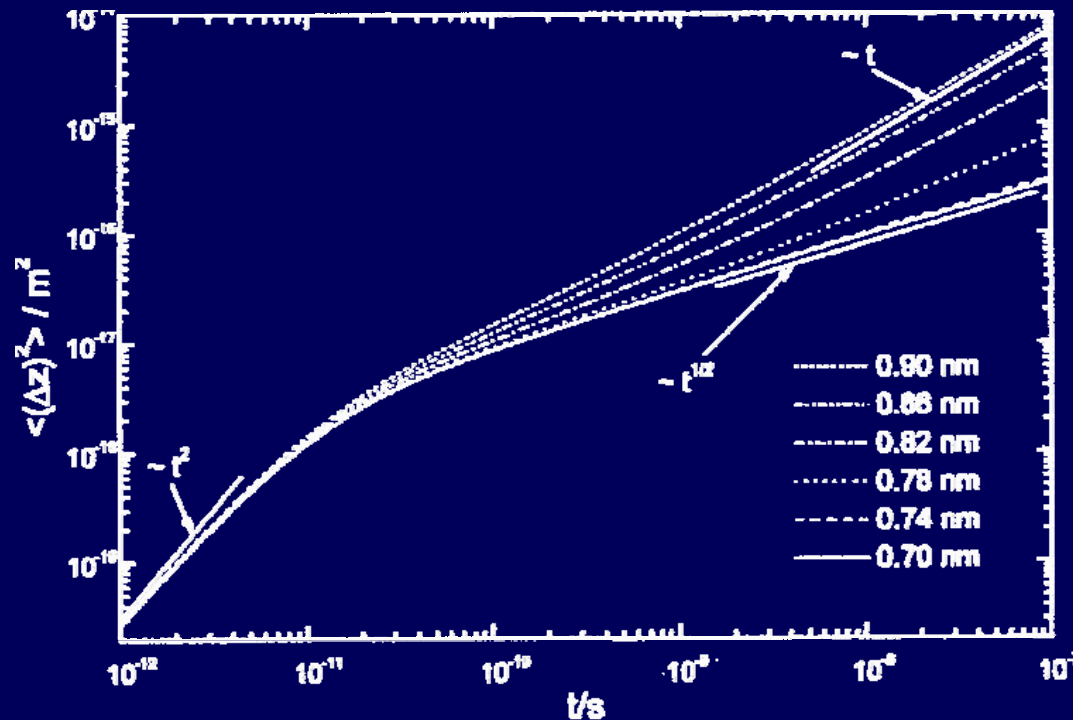
$$\lim_{t \rightarrow \infty} \langle \Delta x(t)^2 \rangle = 2F\sqrt{t}$$

Levitt, Phys. Rev. A 8, 3050 (1973)

Fedders, Phys. Rev. B 17, 40 (1978)

van Beijeren, Kehr, Kutner, Phys. Rev. B, 28, 5711 (1983)

Kärger, J. Phys. Rev. A 45, 4173 (1992)



$$\sigma_P = 0.383 \text{ nm } (\cong \text{CF}_4)$$

Hahn, Kärger, J. Phys. Chem. B, 102, 5766 (1998)

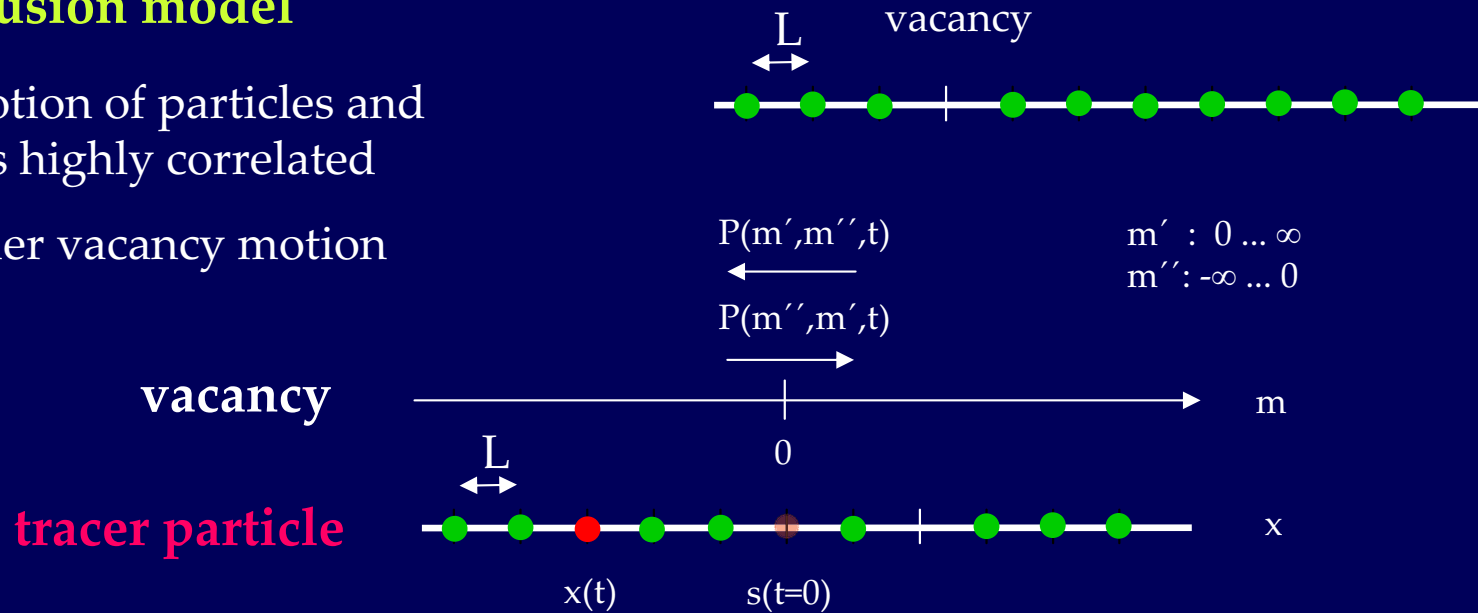
Kärger's Derivation of $t^{1/2}$ - Law

J. Kärger, Phys. Rev. A 45, 4173 (1992)

1D exclusion model

$\Theta \approx 1$: motion of particles and vacancies highly correlated

→ consider vacancy motion



$$\langle x^2(t) \rangle = L^2(1 - \Theta) \int_{m'=0}^{\infty} \int_{m''=-\infty}^0 [P(m', m'', t) + P(m'', m', t)] dm' dm''$$

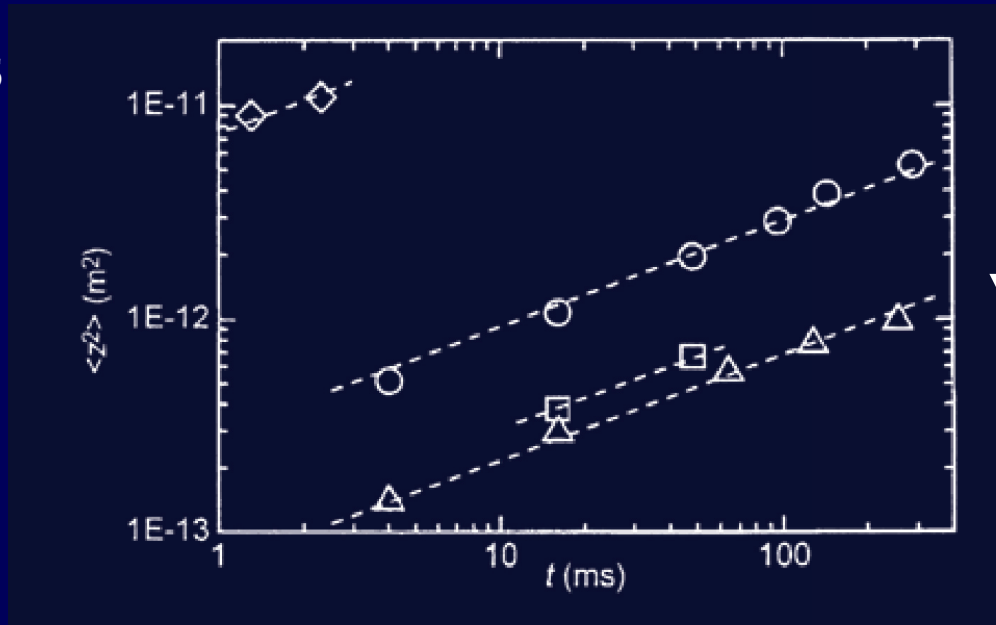
normal diffusion of vacancies

$$P(m', m'', t) = \left[\frac{L^2}{4\pi D_v t} \right]^{1/2} \exp \left[-(Lm' - Lm'')^2 / (4D_v t) \right]$$

$$\langle x^2(t) \rangle = \left[\frac{2}{\pi} \right]^{1/2} L^2 \frac{1 - \Theta}{\Theta} \left[\frac{t}{\tau} \right]^{1/2} \quad \checkmark$$

SFD in Zeolites

$\text{CF}_4 / \text{AlPO}_4\text{-5}$



SFD ✓

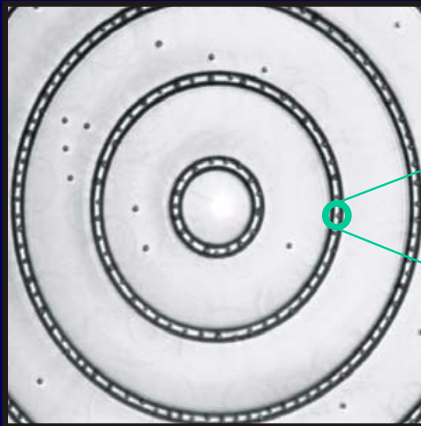
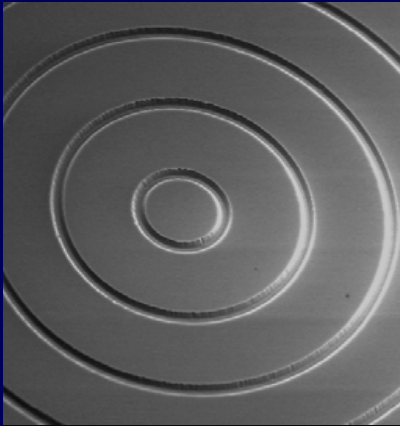
Hahn, Kärger, Kukla, Phys. Rev. Lett. 76, 2762 (1996)

However: controversial results for $\text{CH}_4 / \text{AlPO}_4\text{-5}$: SFD and ND

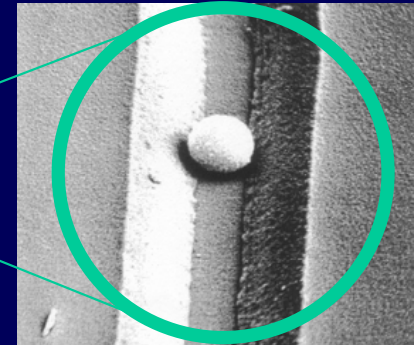
➔ no ideal pore structure?
interaction across adjacent pores ?

SFD in colloidal systems

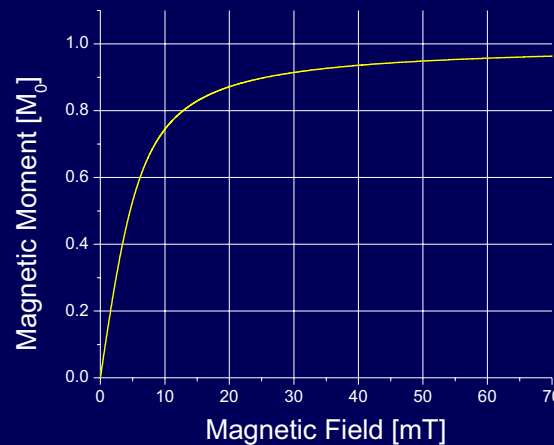
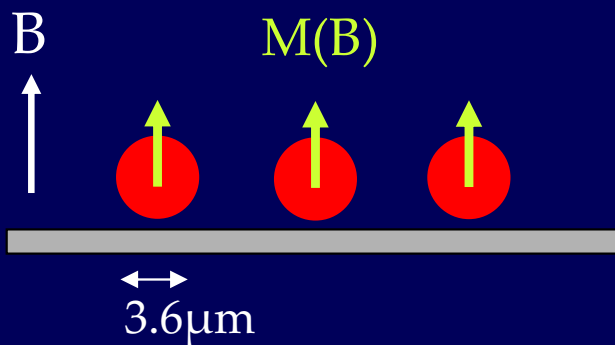
channel structures:



50 μm



particles:



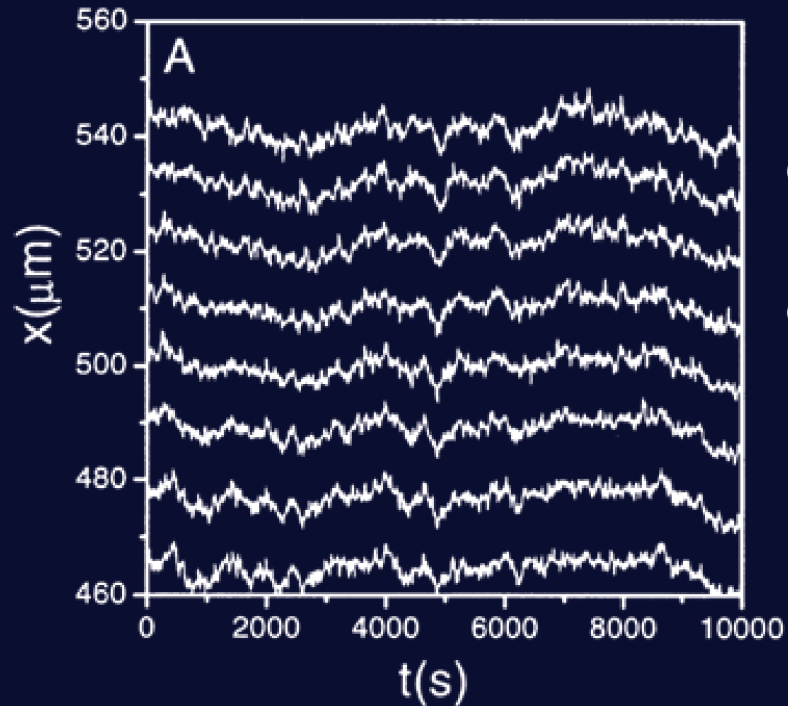
$$M = M_0 \cdot \left(\coth(\alpha B) - \frac{1}{\alpha B} \right)$$

$$M_0 = 6 \times 10^{-13} \text{ Am}^2$$

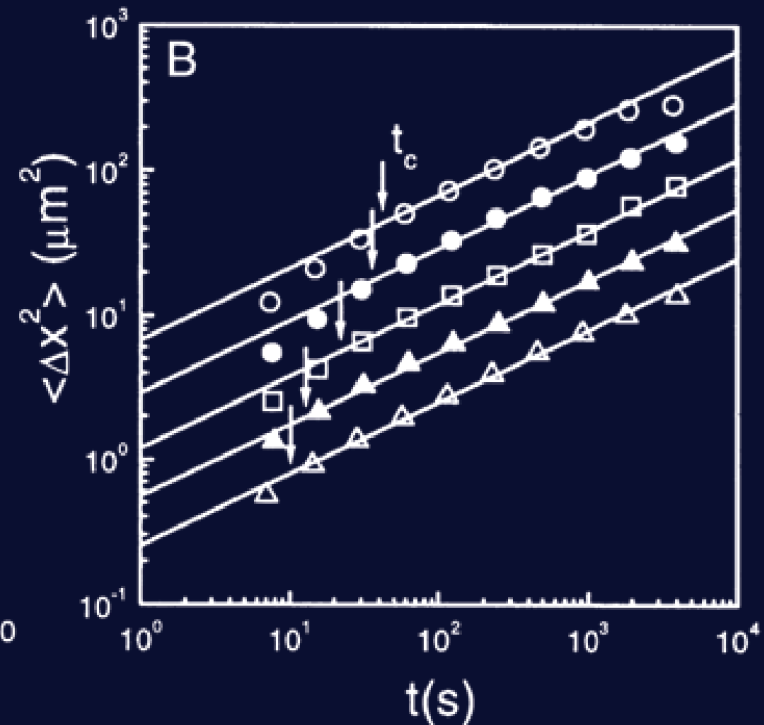
$$\alpha = \frac{\mu}{k_B T}$$

Direct Observation of SFD

single particle trajectories



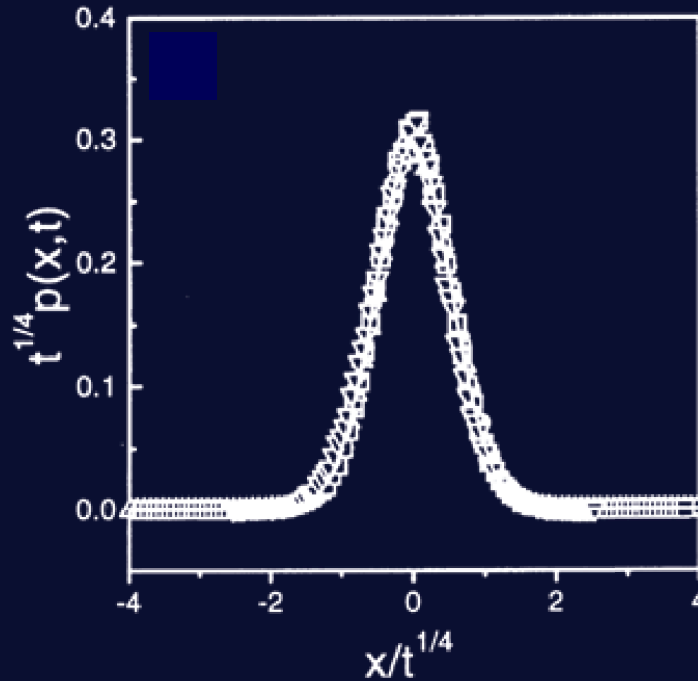
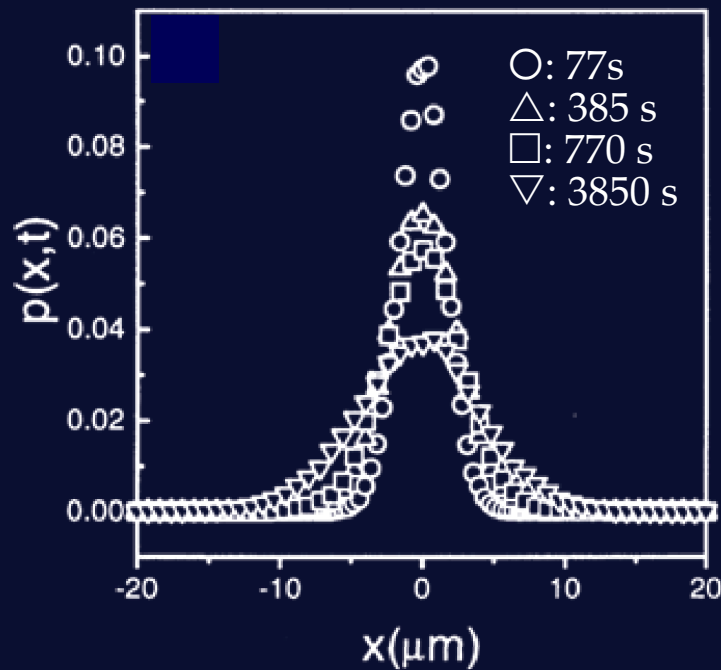
MSD



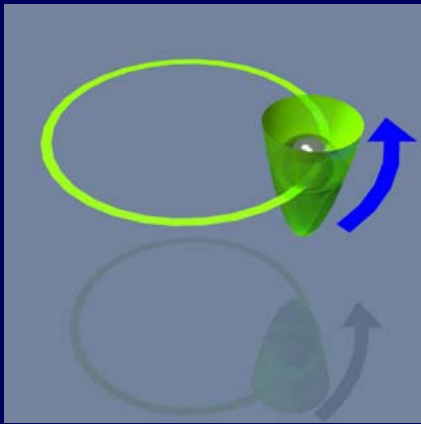
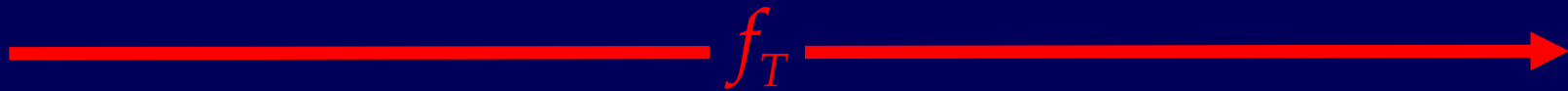
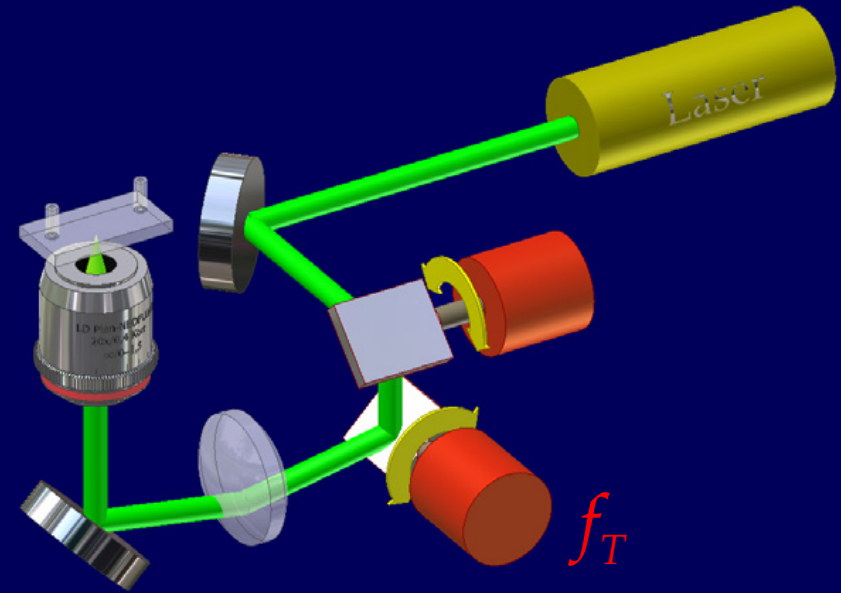
$$\langle \Delta x^2 \rangle = 2F\sqrt{t}$$

Propagator

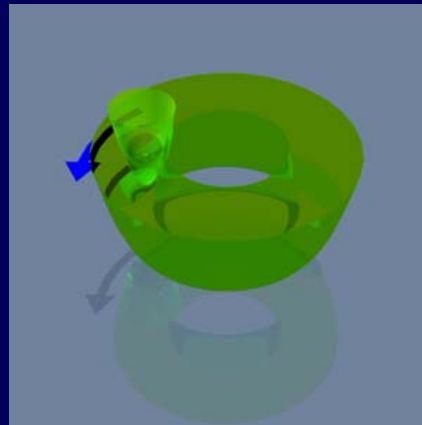
$$p(x,t)_{x=0,t=0} = \frac{1}{\sqrt{4\pi Ft^{1/2}}} \exp(-x^2/4Ft^{1/2}) \quad (\text{hard rods})$$



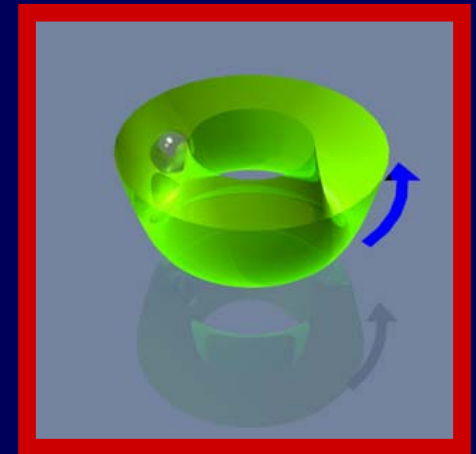
Channels Made by Optical Tweezers



Single moving trap

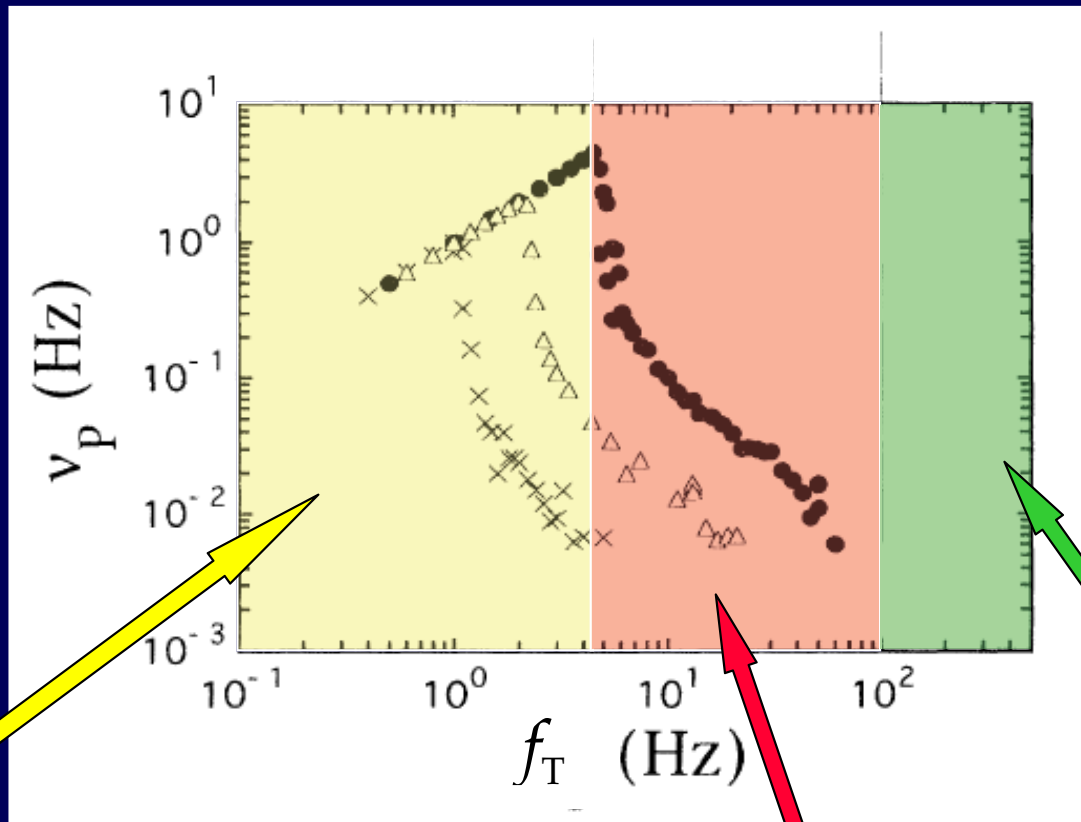


intermediate regime



quasi-static toroidal trap

Scanning Optical Fields



phase-locked

$$v_P = f_T$$

phase-slip

diffusive

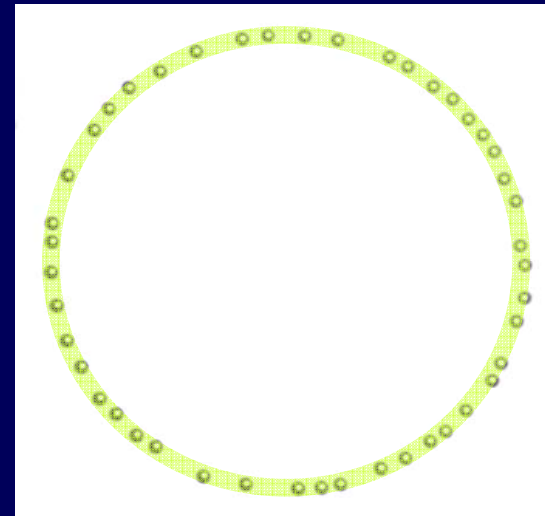
$$v_P = \frac{2}{(2\pi R)^2} \frac{1}{f_T} \frac{1}{w_0} \left[\frac{V_{foc}}{6\pi\eta a} \right]^2$$

Channels Made by Optical Tweezers

$$f_T \approx 300 \text{ Hz}$$

2.9 μm PS particles

$$\beta u(r) = (Z^*)^2 \lambda_B \left(\frac{\exp(\kappa\sigma)}{1 + \kappa\sigma} \right)^2 \frac{\exp(-\kappa r)}{r}$$

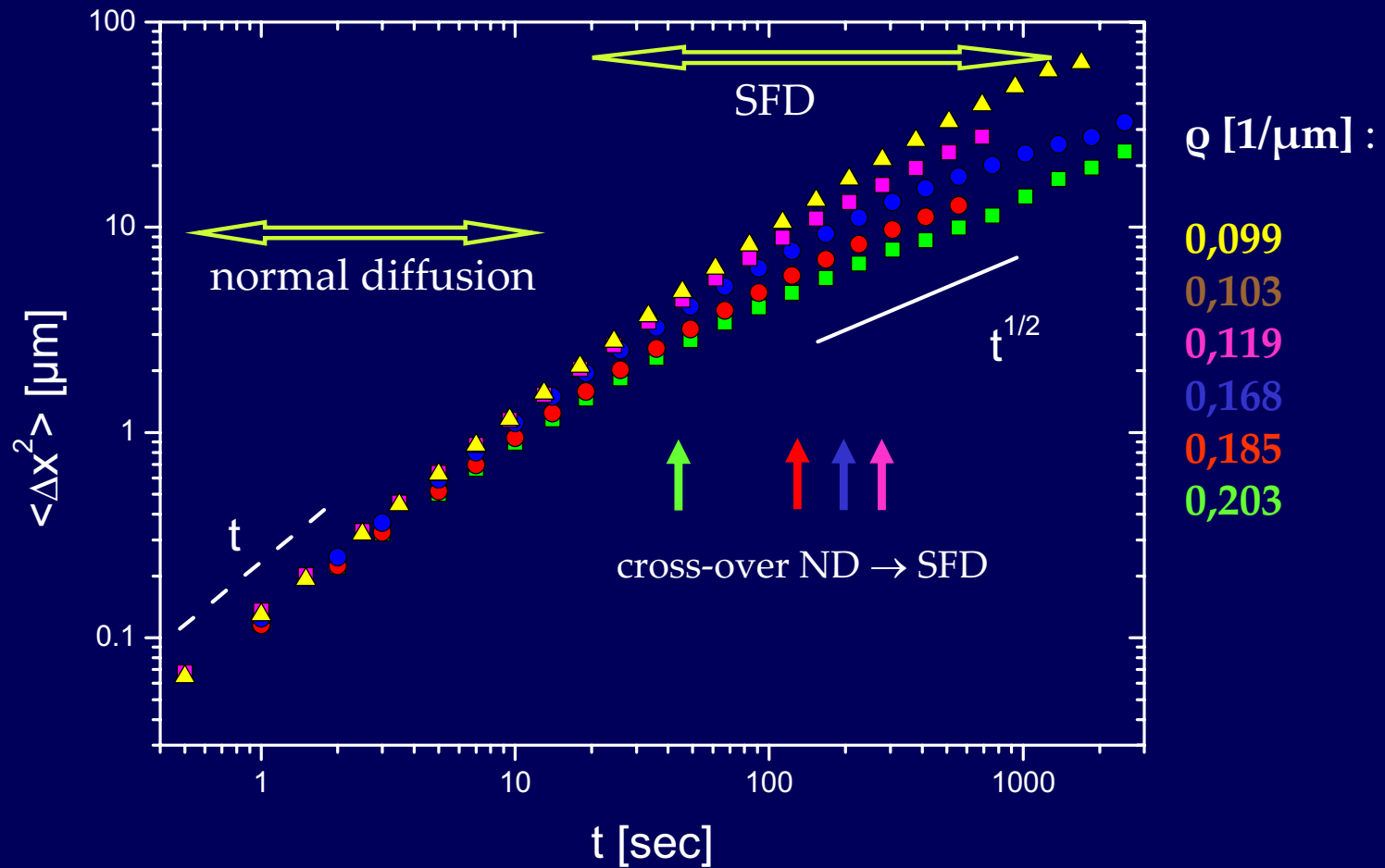


20 μm

Advantages

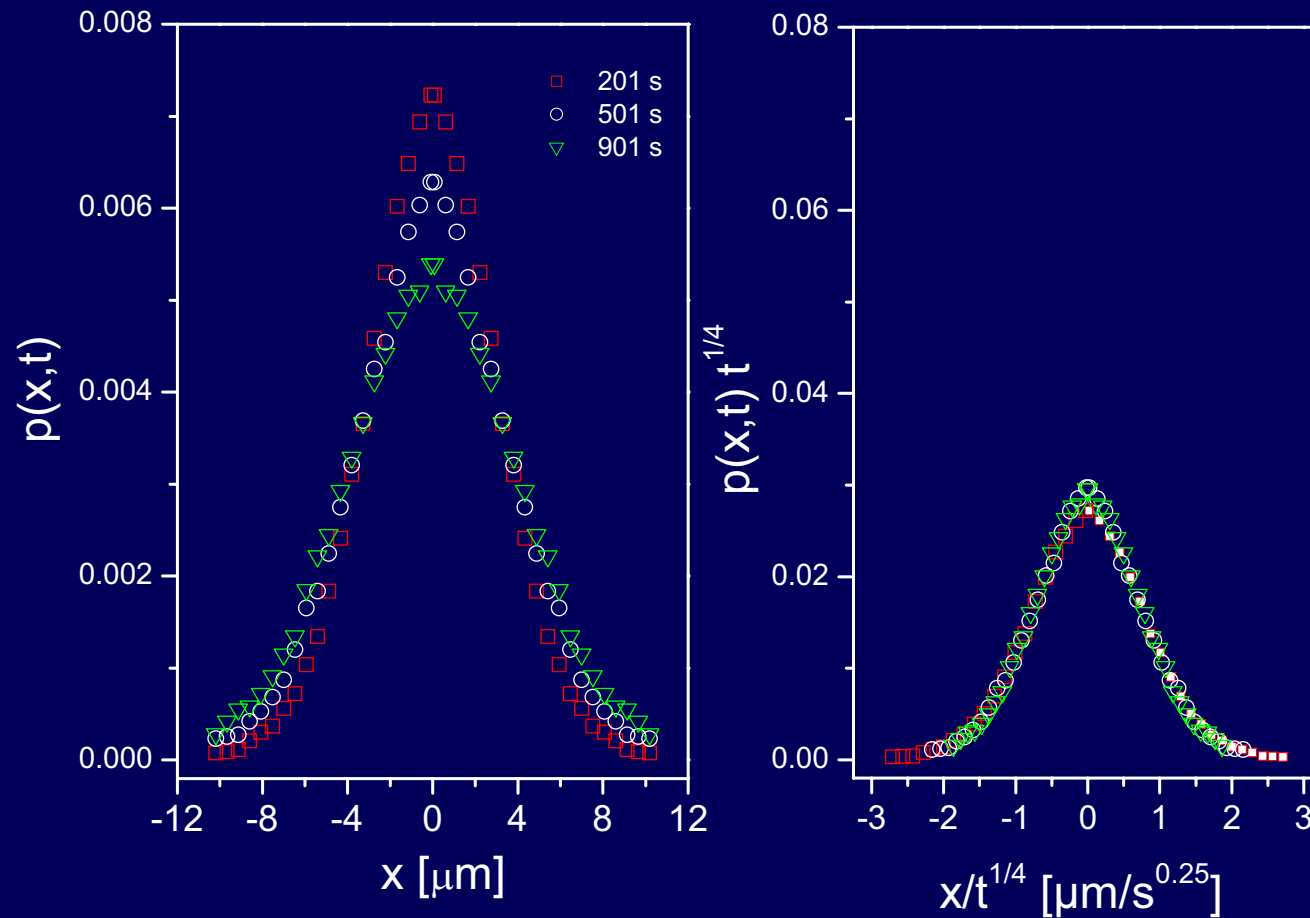
- In situ control of channel geometry and particle number density
- Higher particle mobility due to absence of sticking boundary conditions @ walls

Crossover: Normal Diffusion to SFD

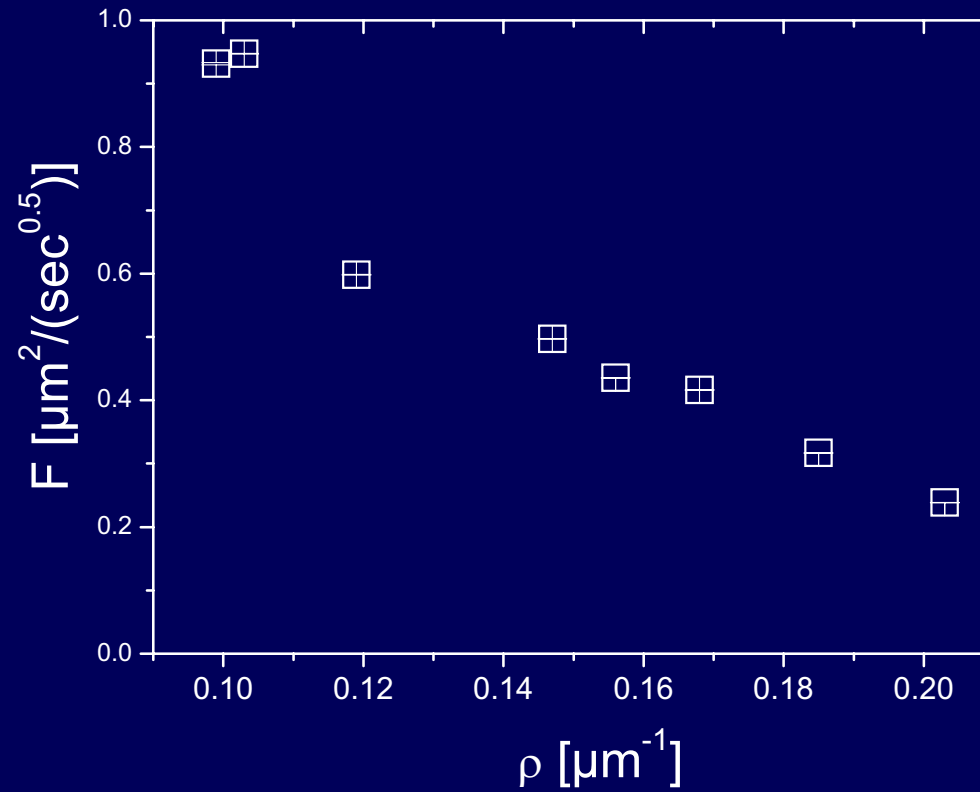


Propagator

$$p(x, t)_{x=0, t=0} = \frac{1}{\sqrt{4\pi Ft^{1/4}}} \exp(-x^2/4Ft^{1/2})$$



SF Mobility



F from Intrinsic System Properties

$$\lim_{t \rightarrow \infty} \langle \Delta x^2(t) \rangle = \underbrace{\frac{2S(q, t=0)}{\rho}}_{2F} \sqrt{\frac{D^{\text{eff}}(q)}{\pi}} \sqrt{t}$$

Kollmann PRL 90, 180602 (2003)

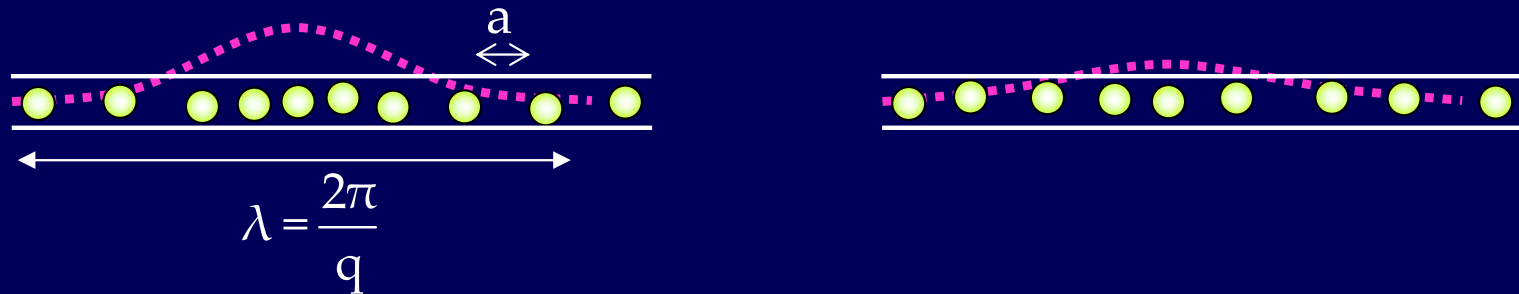
Valid for any pair interaction

HI treated pairwise additive

infinite system

long-wavelength limit ($q \ll a^{-1}$)

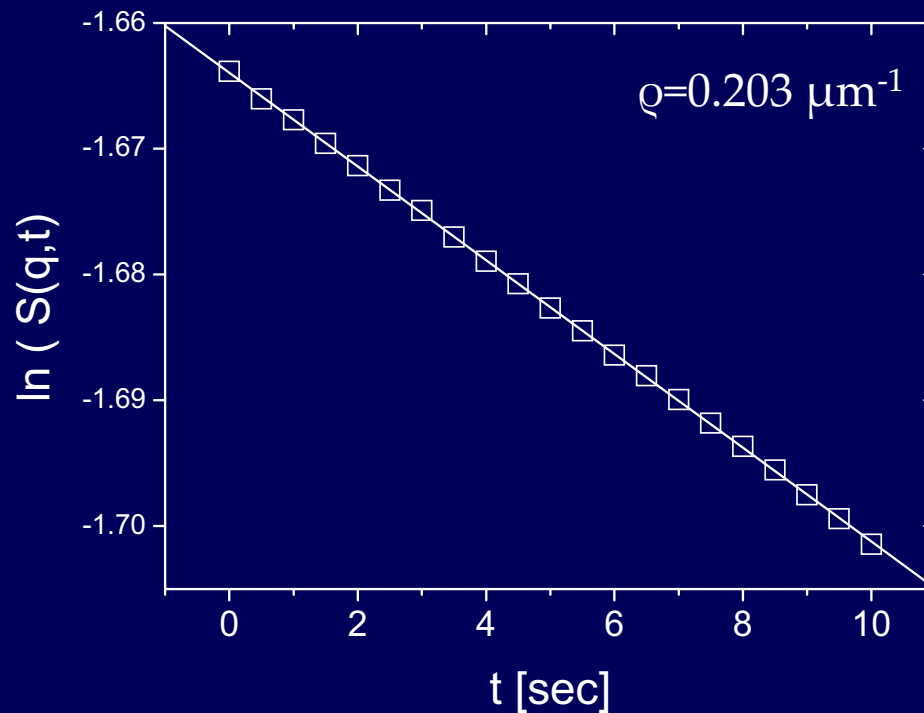
Why MSD is related to collective diffusion coefficient $D^{\text{eff}}(q)$?



1D: Decay of density mode \longleftrightarrow trajectory of every single particle

Dynamic Structure Factor

$$S(q, t) = \frac{1}{N} \left\langle \sum_{i,j} \exp(-iq[x_j(t + \tau) - x_i(\tau)]) \right\rangle_{\tau}$$



$$q = 4q_{\min}$$

$$\left(q_{\min} = \frac{2\pi}{\lambda_{\max}} = \frac{1}{R} \right)$$

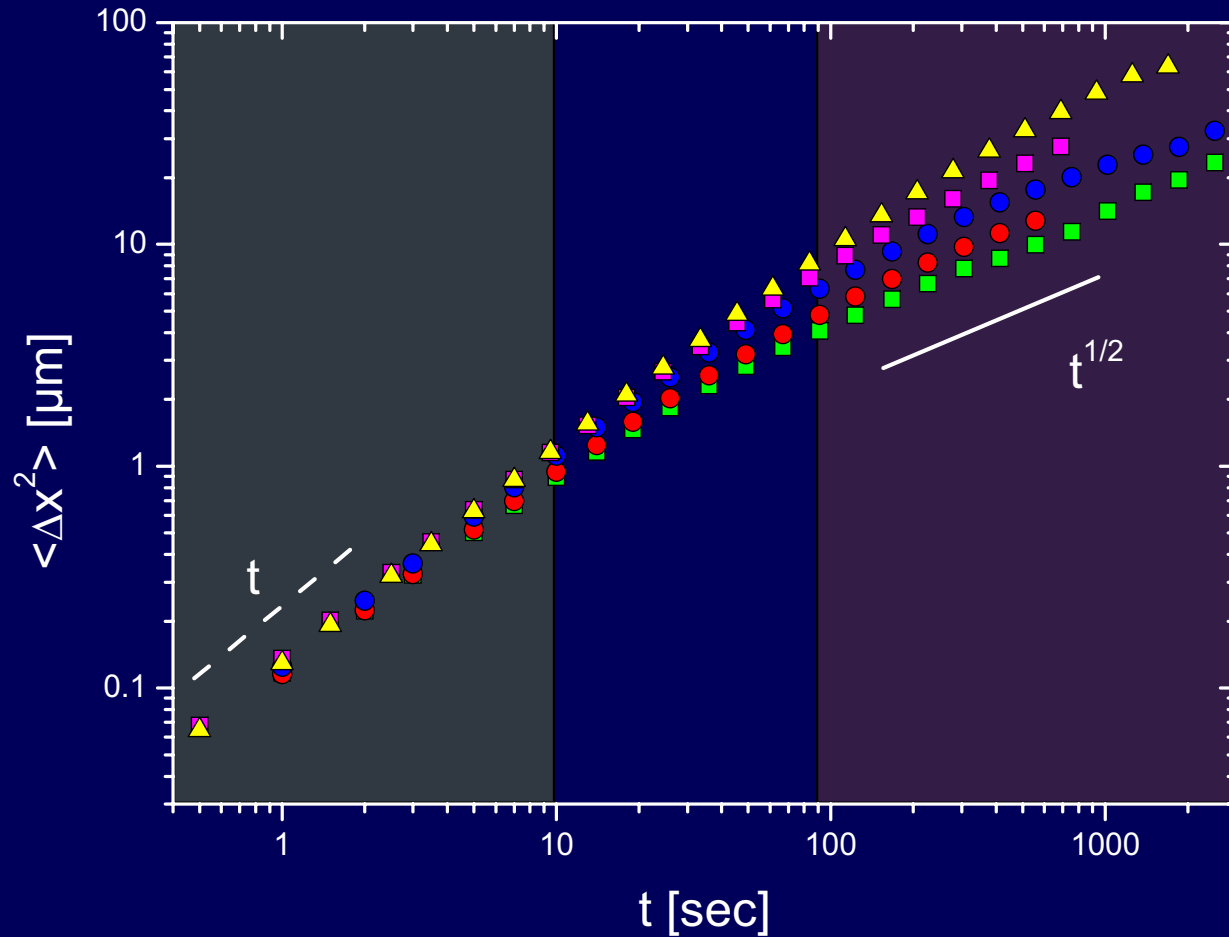
$q \ll a^{-1}$:

$$S(q, t) = S(q, 0) \exp(-q^2 D^{\text{eff}}(q)t)$$

Nägele, *Phys. Rep.* **272**, 215 (1996)

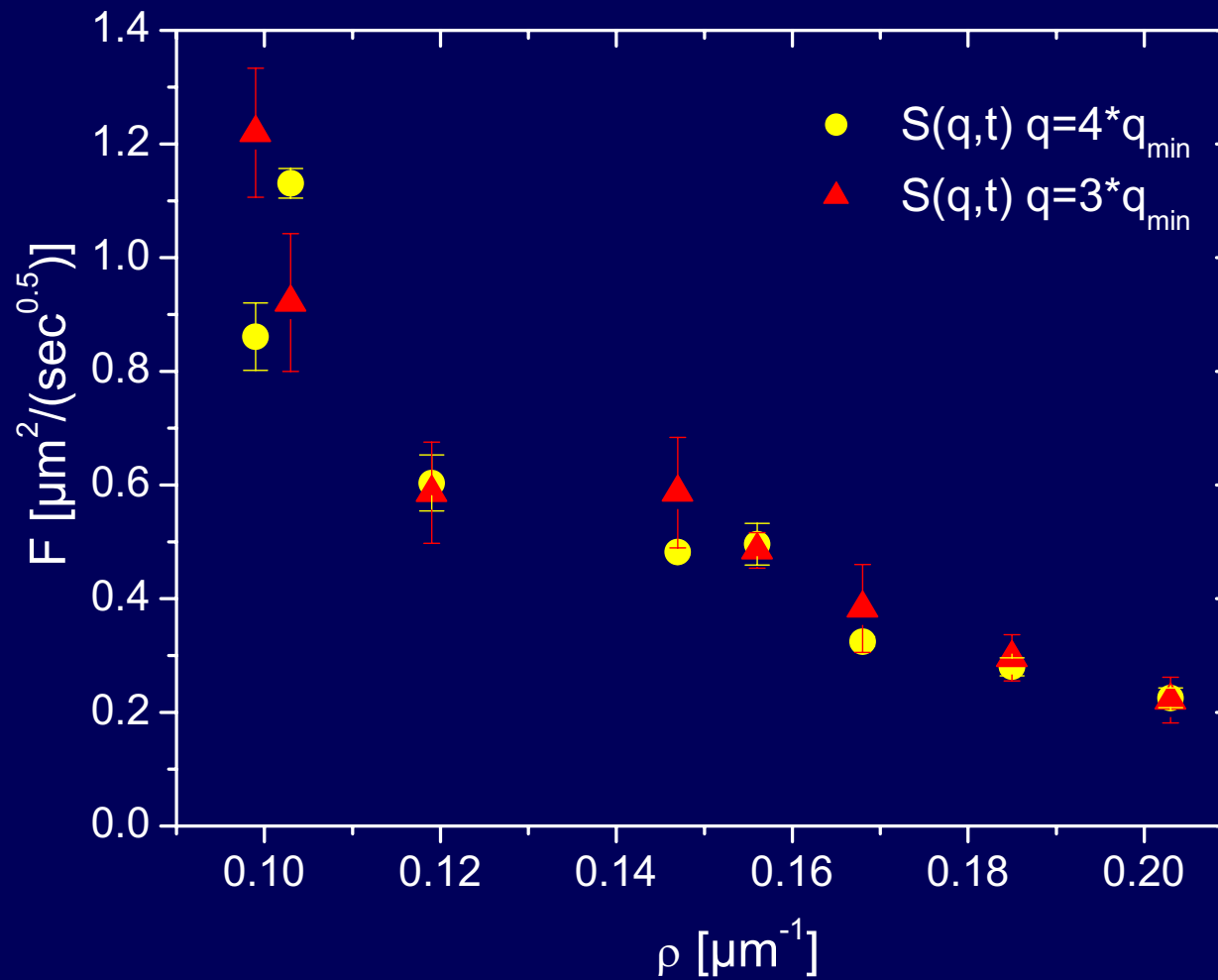
F can be obtained at short times ($t < t_c$) !!

F from Short-Time Behavior

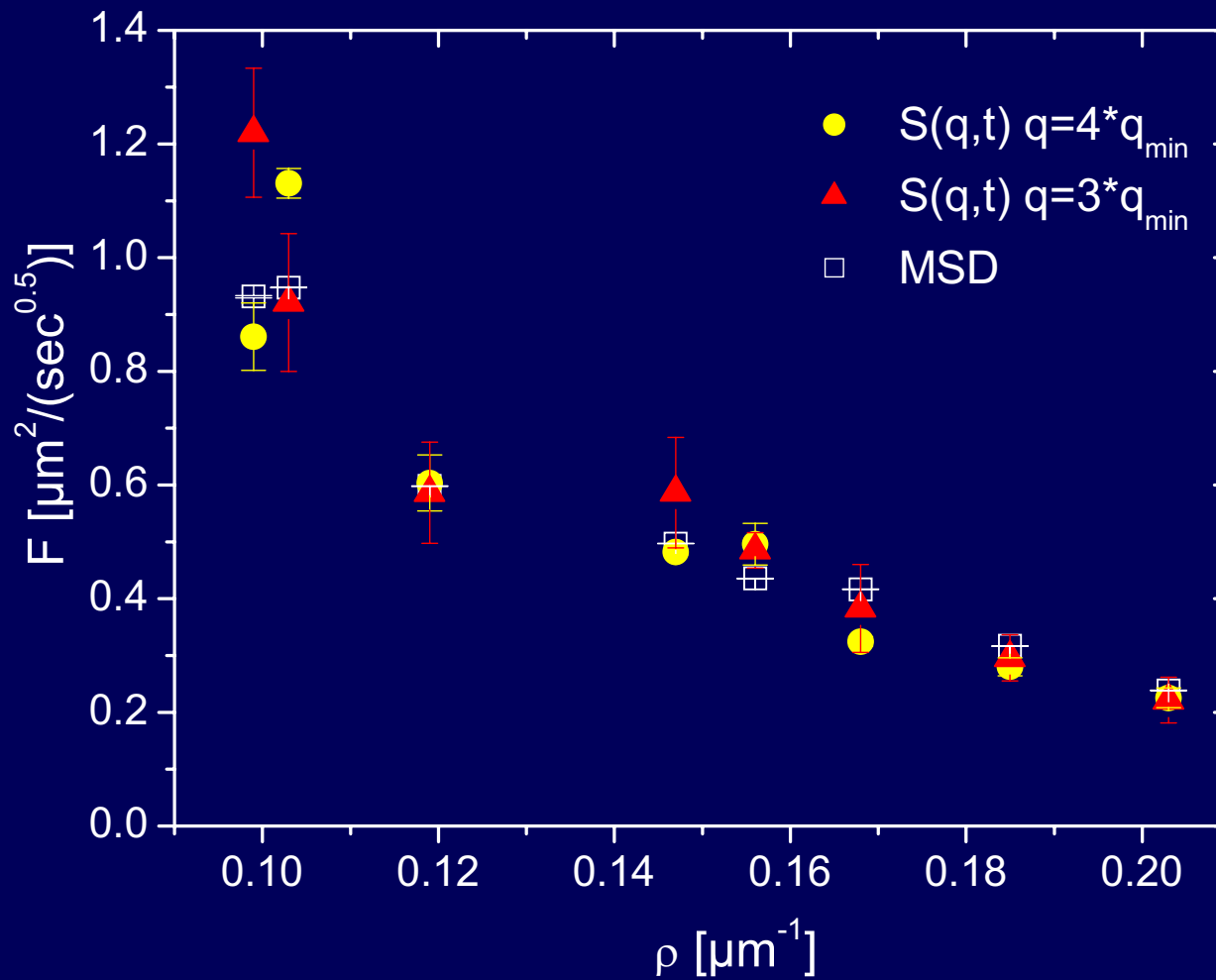


$$F = \frac{S(q, 0)}{\rho} \sqrt{\frac{D^{\text{eff}}(q)}{\pi}} \quad \Longrightarrow \quad \langle \Delta x^2 \rangle = 2F\sqrt{t}$$

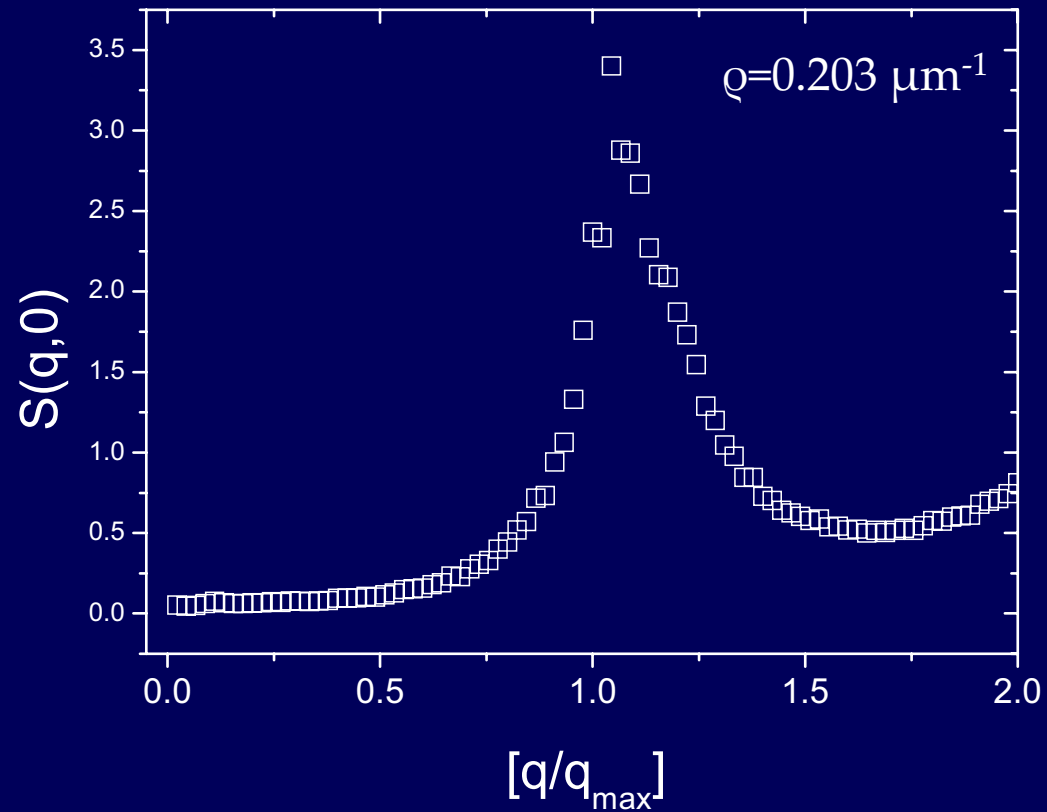
SFD - Mobility



SFD - Mobility



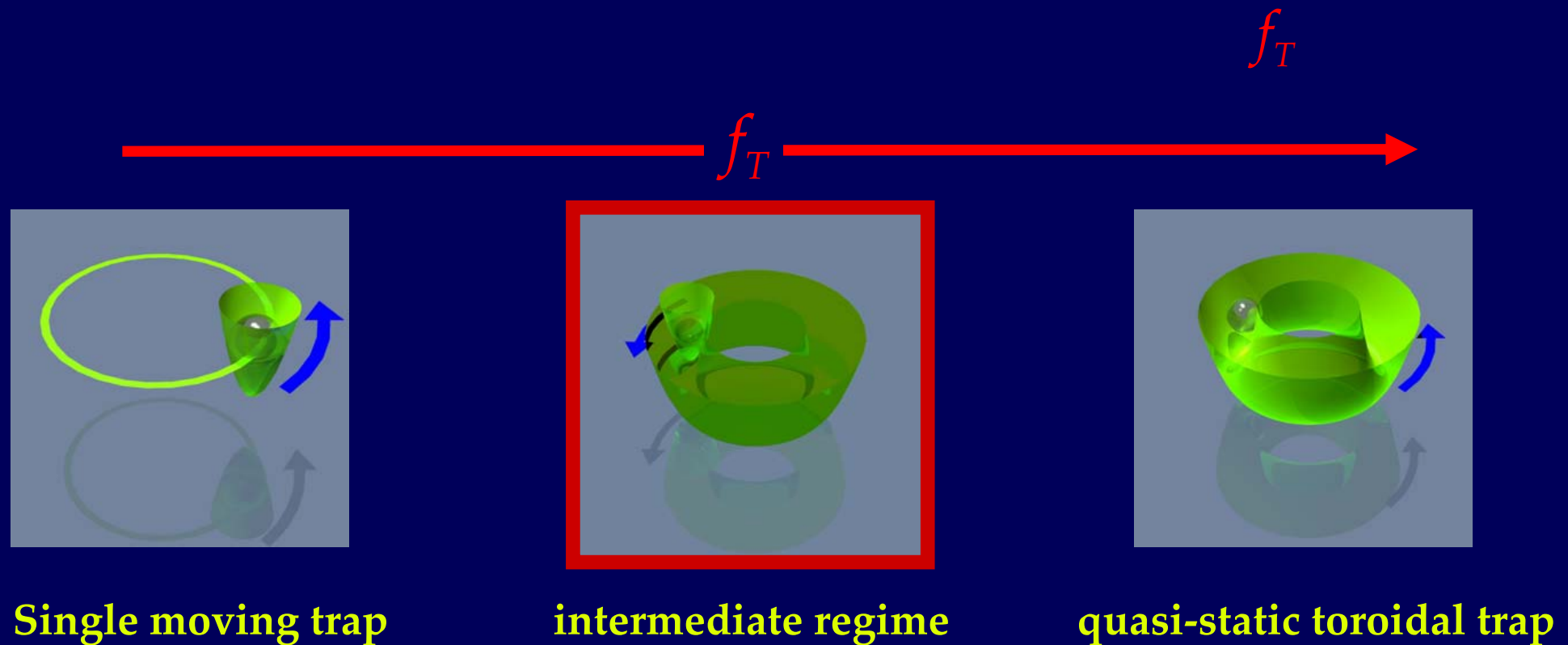
Finite Size Effects ?



$$q_{\max} = \frac{2\pi}{a}$$

$$\lim_{q \rightarrow 0} \frac{\partial S(q,0)}{\partial q} \approx 0$$

From Diffusive to Driven Motion

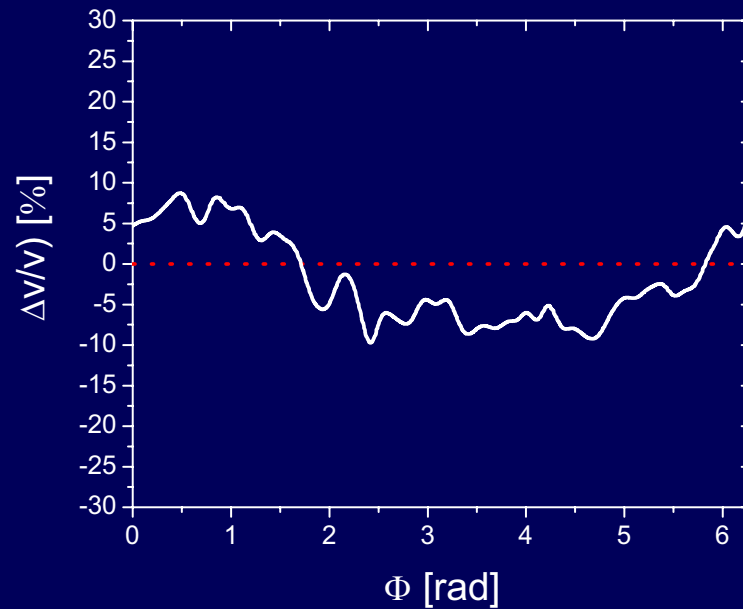


Lutz, Reichert, Stark, Bechinger, EPL 74, 719 (2006)

Phase-Sliper Regime



- silica particles, $\sigma = 3\mu\text{m}$
- ethanol (3D tweezing)
- $f_T = 76\text{Hz}$



constant, non-conservative force

Circling Particles in Toroidal Trap

- silica particles, $\sigma = 3\mu\text{m}$
- electrostatic interaction, $\kappa^{-1} \approx 300\text{nm}$
- ethanol (3D tweezing)
- $f_T = 76\text{Hz}$



mechanism:



particle pair
catches up with
isolated sphere

max. screening
from fluid flow
→ highest mobility

escape of the
two front
particles

particle pair
catches up with
isolated sphere

Summary

Colloids are versatile model systems for statistical physics

„Colloids are the computer simulator's dream“ (Daan Frenkel)

- **realization of SF-conditions in colloidal systems**
topographic structures, optical tweezers
- **transition from normal diffusion to SFD**
dependence of crossover from particle interaction and density
- **F obtained from collective system behavior**
asymptotic single-particle properties derived from short-time collective behavior