

Probing the screening of the Casimir interaction with optical tweezers

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Quantum and Thermal Electrodynamical Fluctuations - KITP 2022



Instituto de Física

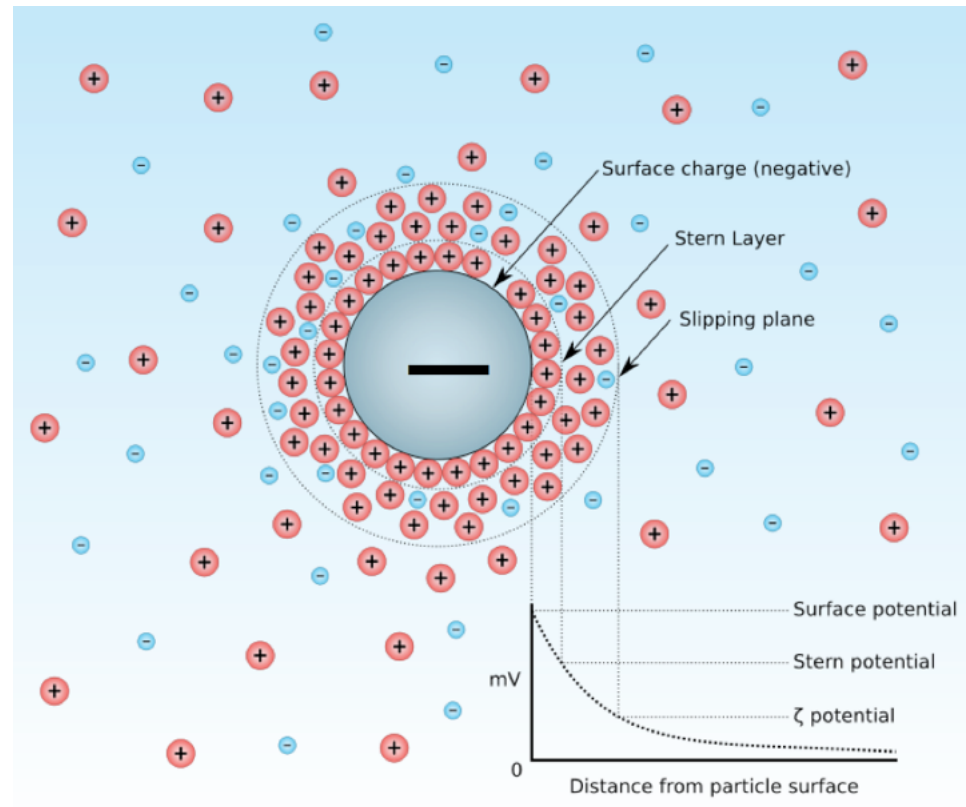
Universidade Federal do Rio de Janeiro



- ▶ Introduction
- ▶ Theory
- ▶ Optical tweezers
- ▶ Experimental results
- ▶ Conclusion

screening in electrolytes

- ions in solution form a diffuse layer around the charged particle
- characteristic length: **Debye screening length** λ_D
- electrostatic interaction between two particles exponentially suppressed for distances $> \lambda_D$
- Partial or total (?) screening of the zero-frequency contribution to the Casimir interaction



screening in electrolytes

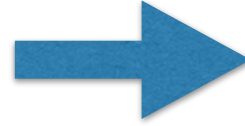
- Details of screening are essential for understanding the Casimir interaction between dielectric materials across a polar liquid
- Indeed, in this case zero-frequency contribution could be dominant already at distances > 100 nm as refractive indexes at positive Matsubara frequencies nearly match

- Example: aqueous solution; biological samples: $\lambda_D < 1$ nm
- Is there any interaction left at distances > 100 nm ?

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Nonlocal response in aqueous solution

movable ions in solution



Non-local response

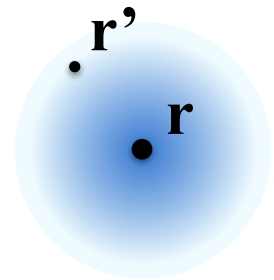
Bulk approximation: we ignore the presence of boundaries when deriving the constitutive equations (Ohm's law)

real space $\mathbf{J}(\mathbf{r}) = \int d^3r' \sigma(\mathbf{r} - \mathbf{r}') \cdot \mathbf{E}(\mathbf{r}')$

Fourier $\mathbf{J}(\mathbf{K}, \omega) = \sigma(\mathbf{K}, \omega) \cdot \mathbf{E}(\mathbf{K}, \omega)$



K dependence/spatial dispersion



Related problem: non-local response of metals and the Casimir effect

Kats, Barton, Contreras-Reyes, Esquivel-Sirvent, Mochan, Svetovoy, Villareal, Intravaia, Klimchitskaya, Mostepanenko, Henkel

vdW with aqueous solution: B. Davies and B. W. Ninham, J. Chem. Phys. (1972)

Electrodynamics in the aqueous solution: movable ions

$$\epsilon = \epsilon_b \mathbb{1} + \frac{i}{\omega} \sigma$$

Hydrodynamical model

Transverse permittivity $\epsilon_t(\omega) = \epsilon_b(\omega) - \frac{\omega_P^2}{\omega(\omega + i\gamma)}$

Longitudinal permittivity: nonlocal, Debye screening length λ_D

$$\epsilon_\ell(\mathbf{K}, \omega) = \epsilon_b(\omega) - \left(\frac{\omega(\omega + i\gamma)}{\omega_P^2} - \frac{\lambda_D^2}{\epsilon_b} K^2 \right)^{-1}$$

B. Davies and B. W. Ninham, J. Chem. Phys. (1972)

Screening of the Casimir force

Casimir interaction across an electrolyte

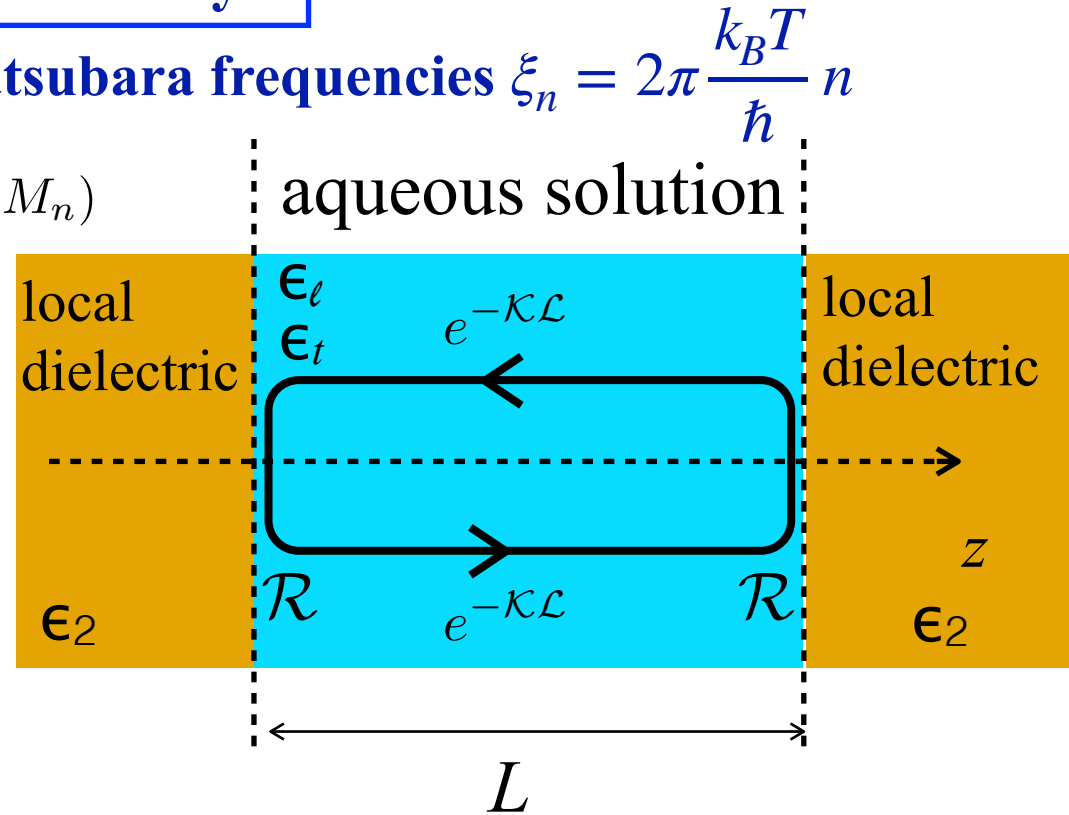
Scattering formula: sum over Matsubara frequencies $\xi_n = 2\pi \frac{k_B T}{\hbar} n$

$$\mathcal{F} = k_B T A \sum_{n=0}^{\infty}{}' \int \frac{d^2 k}{(2\pi)^2} \ln \det(1 - M_n)$$

$$M_n = \mathcal{R} e^{-\kappa L} \mathcal{R} e^{-\kappa L}$$

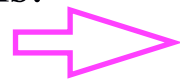
$$\mathcal{R} = \begin{pmatrix} r_{ss} & 0 & 0 \\ 0 & r_{pp} & r_{pl} \\ 0 & r_{lp} & r_{ll} \end{pmatrix}$$

describes the coupling between TM-polarized transverse waves (p) and longitudinal waves



plasma frequency for ions:

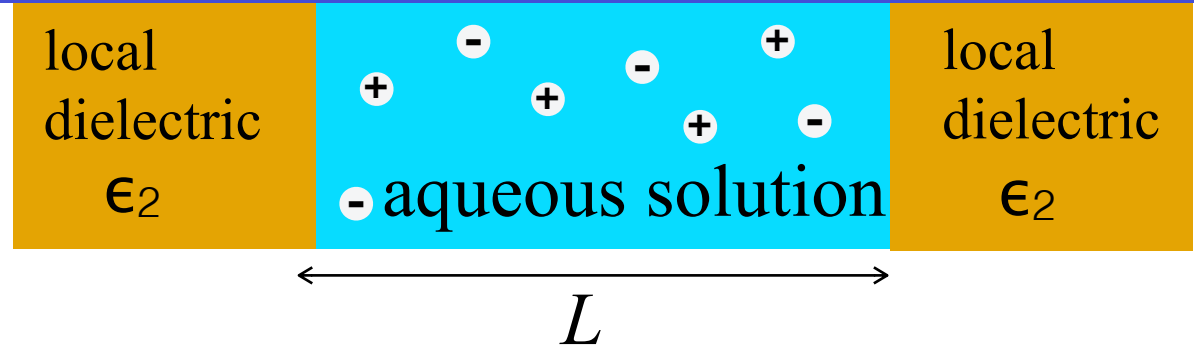
$$\omega_P \ll k_B T / \hbar$$



- only zero-frequency contribution is modified
- no coupling between TM and longitudinal channels

Screening of the Casimir force

Parallel planar interfaces



- zero-frequency limit of the longitudinal reflection amplitude:

$$r_{ll} = \frac{\epsilon_b \sqrt{k^2 + 1/\lambda_D^2} - \epsilon_2 k}{\epsilon_b \sqrt{k^2 + 1/\lambda_D^2} + \epsilon_2 k}$$

- Scattering approach: screened longitudinal contribution and non-screened transverse magnetic (TM) one [Apéry's constant $\zeta(3) \approx 1.20$]

$$\frac{\mathcal{F}_{n=0}}{A} = \frac{k_B T}{2} \left[-\frac{\zeta(3)}{8\pi L^2} + \int \frac{d^2 k}{(2\pi)^2} \ln \left(1 - r_{ll}^2 e^{-2\sqrt{k^2 + 1/\lambda_D^2} L} \right) \right]$$

- Previous result, from fluctuational electrostatics: zero-frequency completely screened. Mitchell & Richmond (1974), Mahanty & Ninham 1976, Parsegian 2006

$$\frac{\mathcal{F}_{n=0}^{\text{LPB}}}{A} = \frac{k_B T}{2} \int \frac{d^2 k}{(2\pi)^2} \ln \left(1 - r_{ll}^2 e^{-2\sqrt{k^2 + 1/\lambda_D^2} L} \right)$$

Screening of the Casimir force

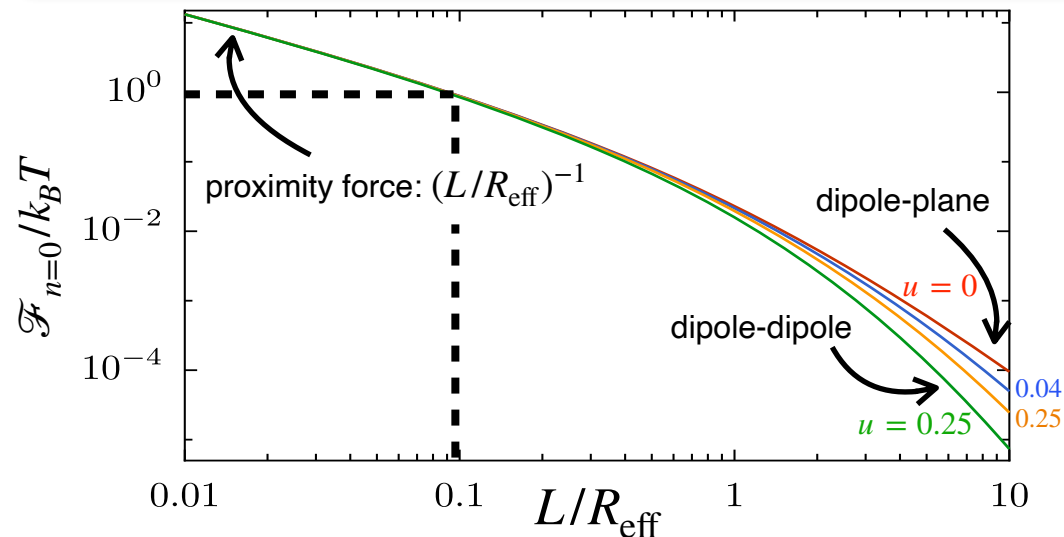
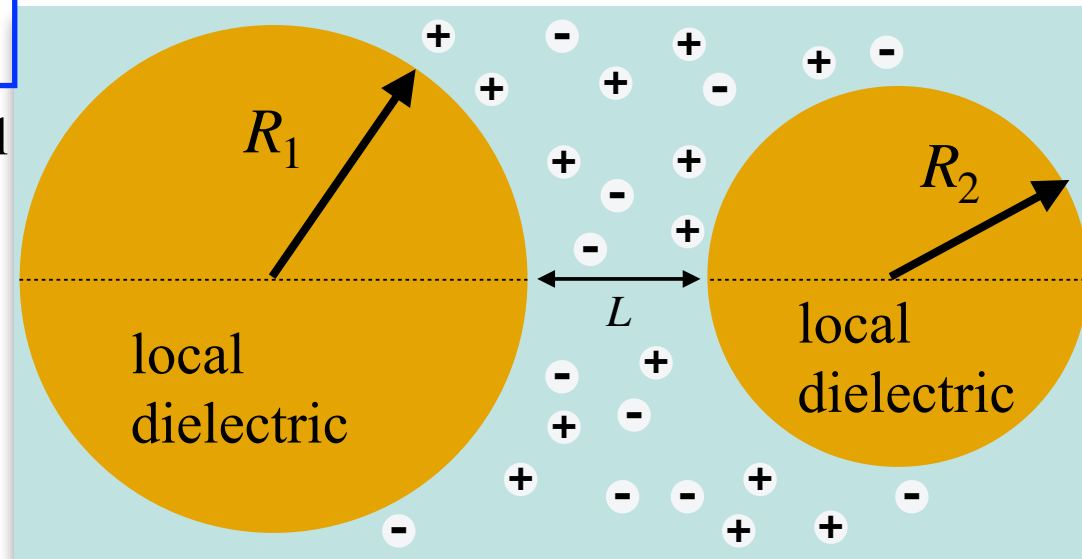
two dielectric spheres

- We assume $L \gg \lambda_D \Rightarrow$ longitudinal contribution completely suppressed by screening
- More general case: see Larissa Inacio's poster
- The only nonzero contribution modified by the ions is the zero-frequency transverse magnetic contribution

• effective radius $R_{\text{eff}} = \frac{R_1 R_2}{R_1 + R_2}$

- Free energy $\mathcal{F}_{n=0}/k_B T$ is an universal function of

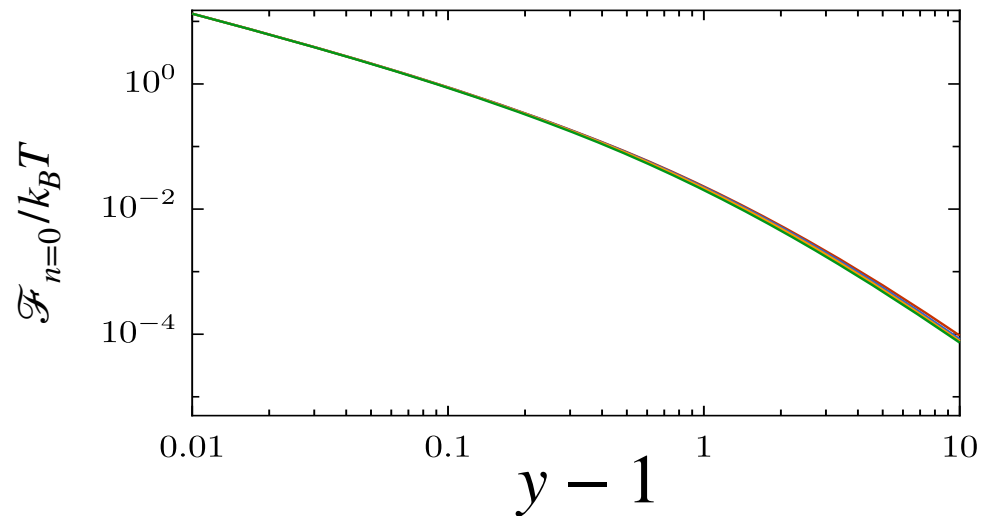
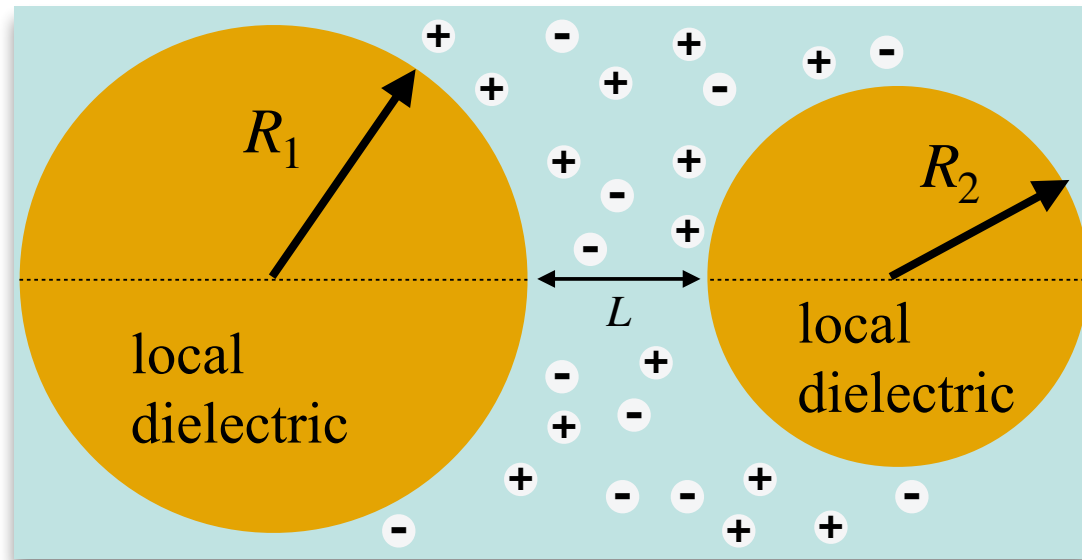
$$u = \frac{R_1 R_2}{(R_1 + R_2)^2} \quad \text{AND} \quad x = L/R_{\text{eff}}$$



Screening of the Casimir force

two dielectric spheres

- effective radius $R_{\text{eff}} = \frac{R_1 R_2}{R_1 + R_2}$
- Free energy $\mathcal{F}_{n=0}/k_B T$ is an universal function of
$$u = \frac{R_1 R_2}{(R_1 + R_2)^2} \quad \text{AND} \quad x = L/R_{\text{eff}}$$
- Better representation: conformally-invariant parameter (Fosco, Lombardo & Mazzitelli)
$$y = 1 + x + u \frac{x^2}{2}$$
- Similar universal function for Drude-vacuum-Drude configuration: Bimonte & Emig, Schoger & Ingold



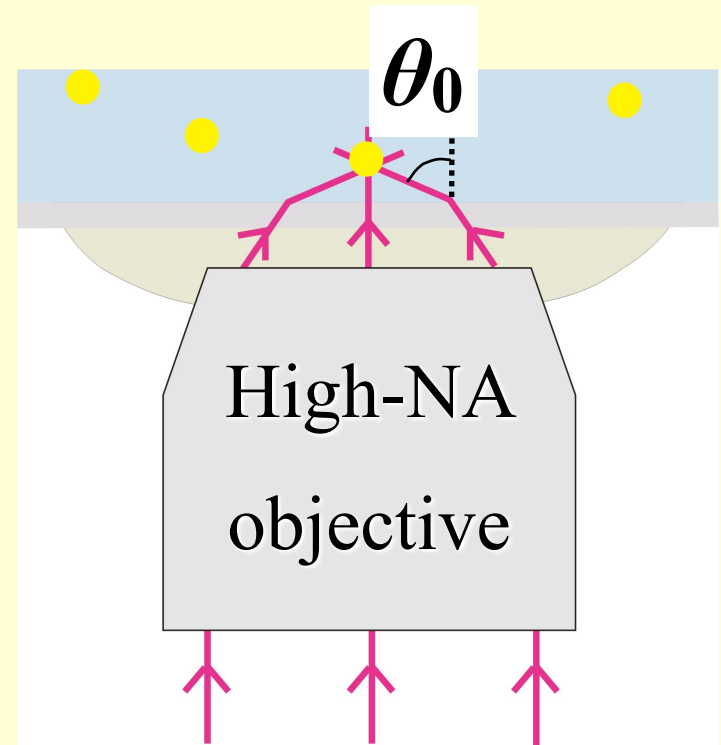
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Optical Tweezers

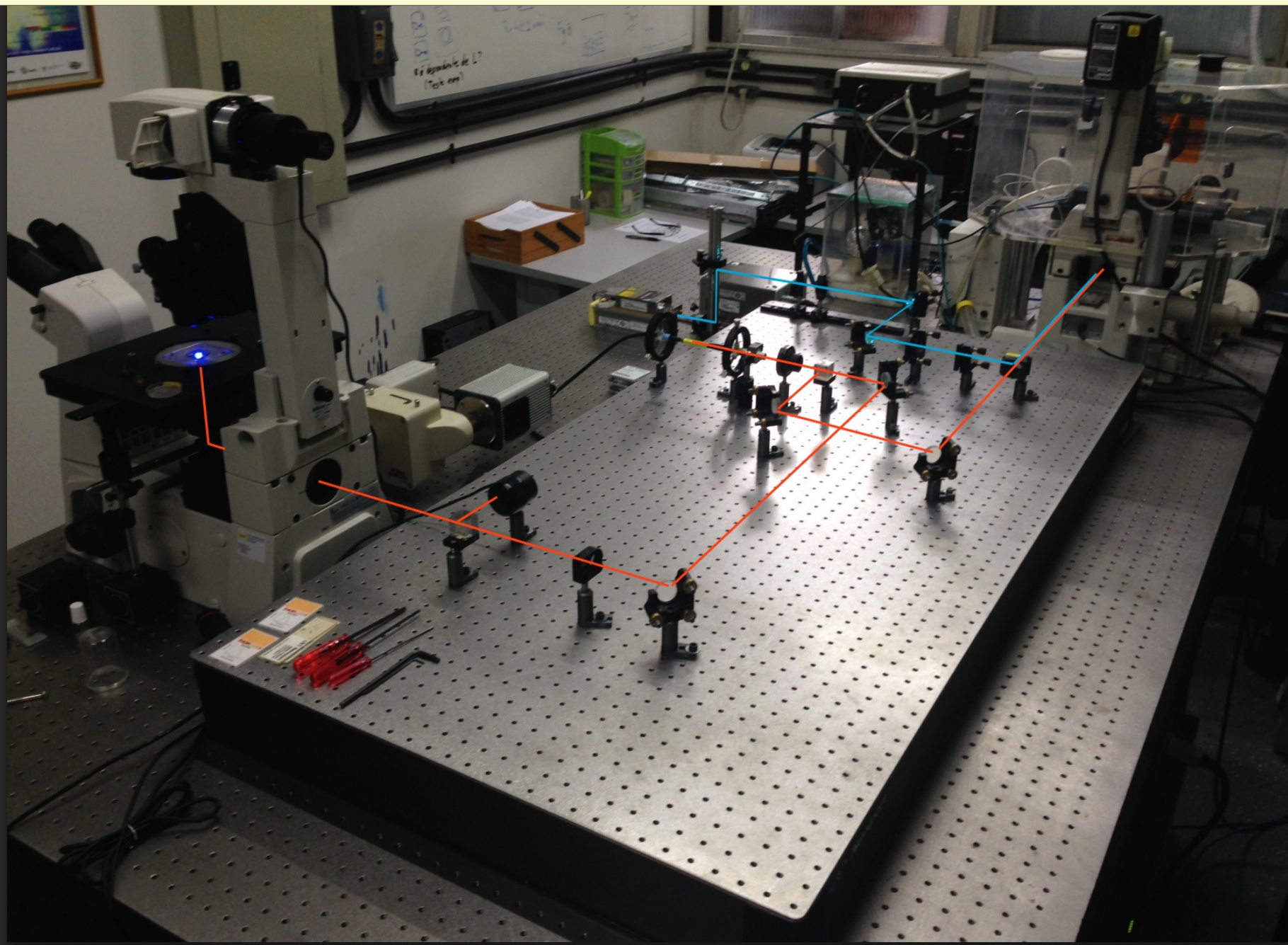
Optical trap with a single strongly focused laser beam...

→ "optical tweezers"



$$NA = n_{\text{water}} \sin(\theta_0)$$

Optical Tweezers Lab in Rio



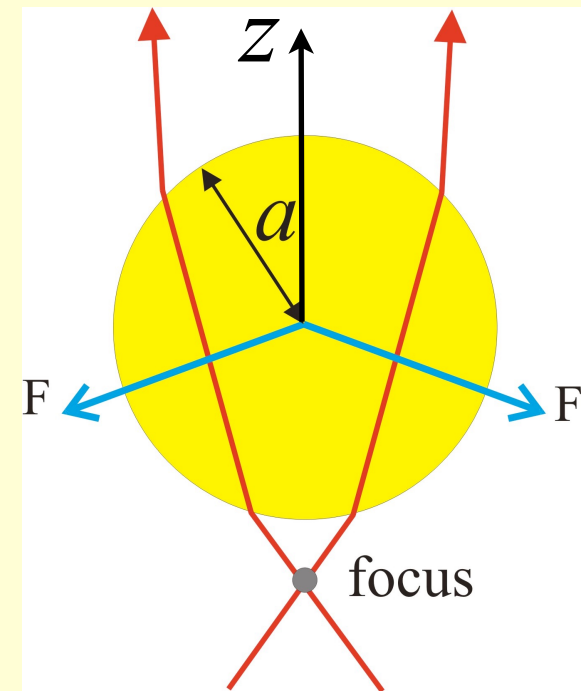
Optical Tweezers

Typically $a \sim \lambda$ \longrightarrow need many multipole terms (beyond dipole approximation) – **Mie scattering** regime

Physical picture in the ray optics regime $a \gg \lambda$

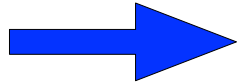
refracted rays: force **towards** the focus ('gradient force') ...

reflected rays: radiation pressure pushing along the propagation direction

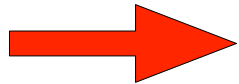


Screening of the Casimir interaction

Motivation: Casimir force experiments with Optical Tweezers



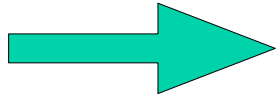
material of the trapped microsphere must have refractive index slightly higher than external medium @ laser wavelength - silica in water



materials allowing for trapping are such that positive Matsubara frequencies provide relatively small contribution

Measuring the Casimir interaction

Casimir (van der Waals) experiments with Optical Tweezers



trap stiffness $k \propto$ laser power P
can be tuned to very small values
 $k \sim$ fN/nm

measuring fN forces between silica microspheres in aqueous solution at distances ~ 100 nm

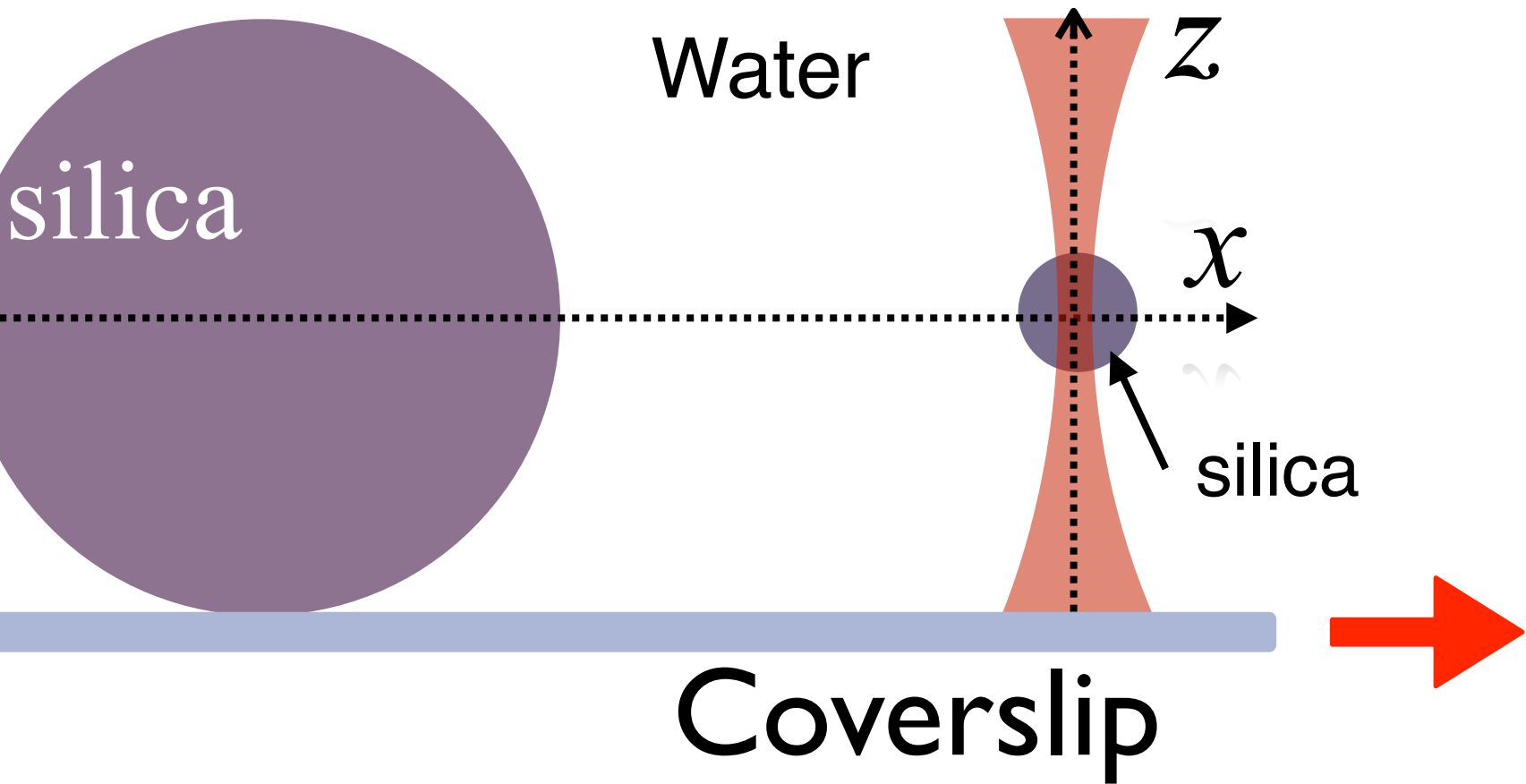
Casimir interaction: radiation pressure of the **quantum electromagnetic field modes in thermal equilibrium**

Typical (usually AFM) measurements of the van-der-Waals/Casimir interaction across an electrolyte (polar liquid)

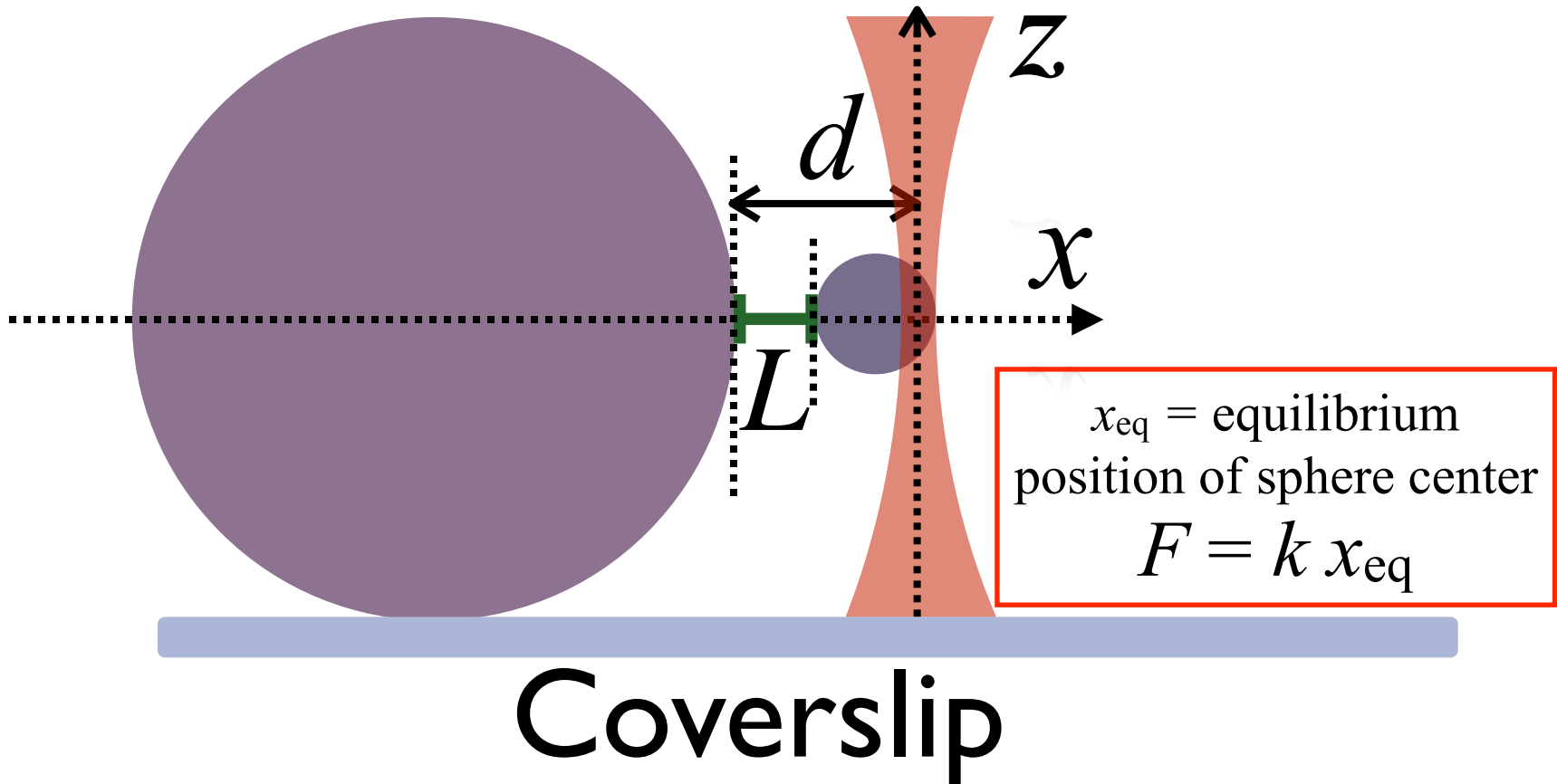
- **metallic surfaces:** distance ~ 100 nm (Munday, Palasantzas, Svetovoy, Ciliberto,...)
- **dielectric surfaces:** distance ~ 1 nm (Borkovec, Trefalt,...)

In both cases, Matsubara zero frequency contribution is usually negligible, and so is screening!

Measuring the Casimir force

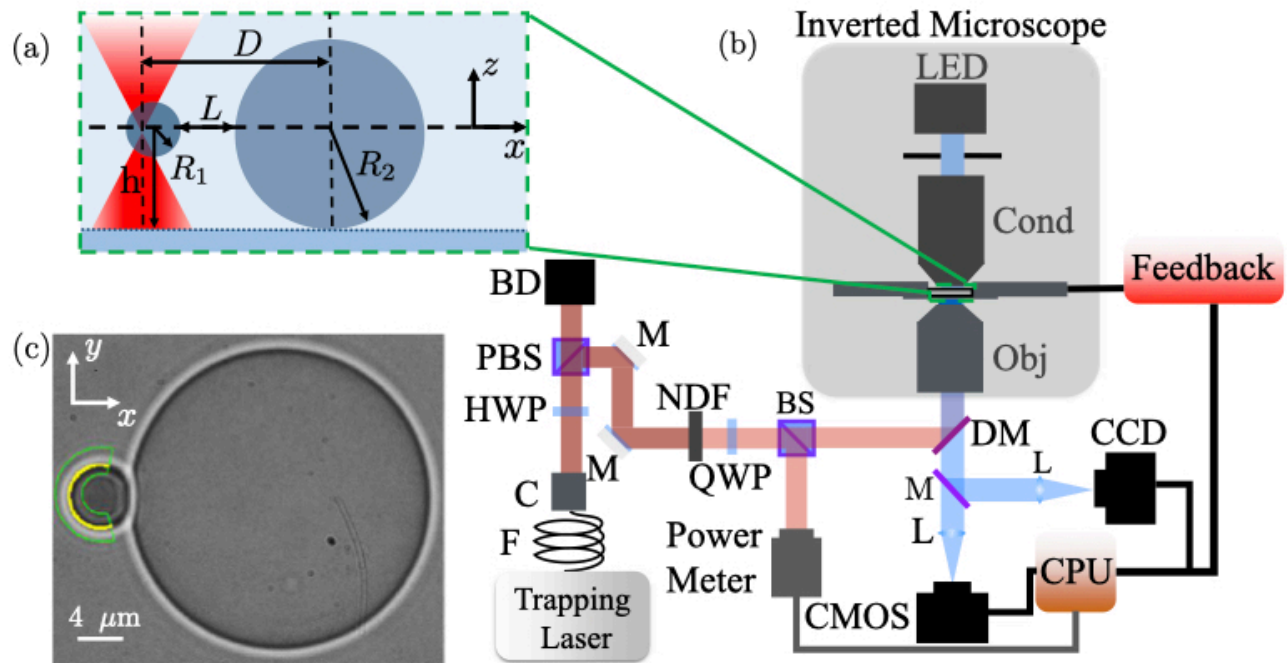
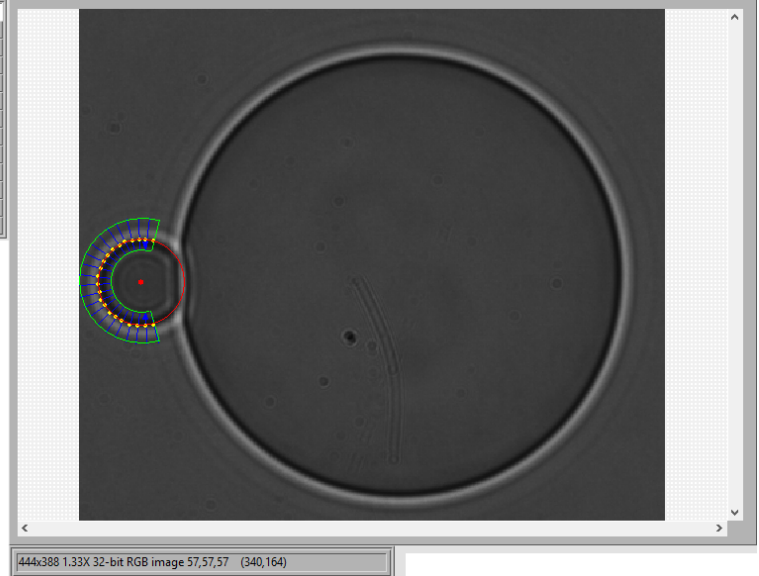
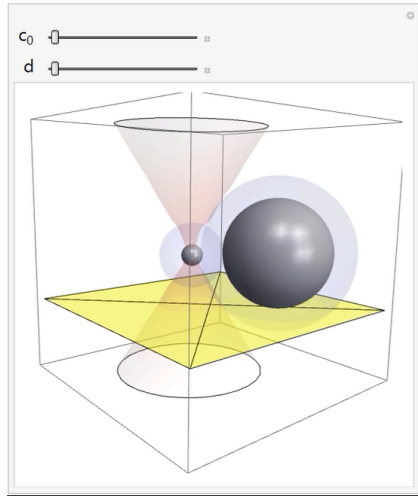


Measuring the Casimir force



measuring the interaction energy from fluctuations

Fluctuations of a trapped sphere near a bigger sphere attached to the glass slide

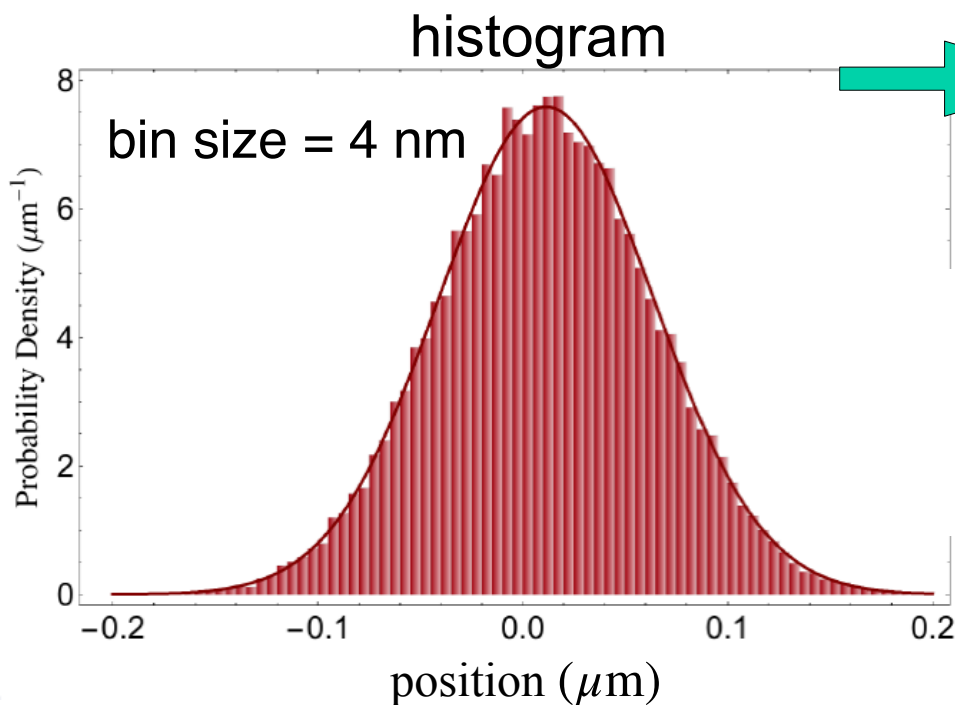
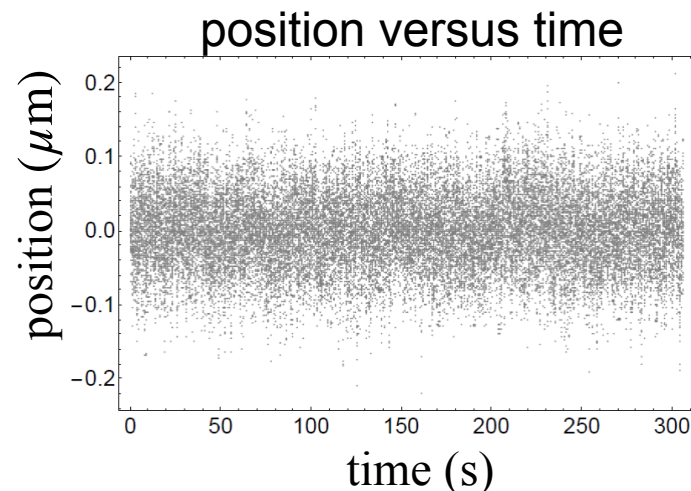
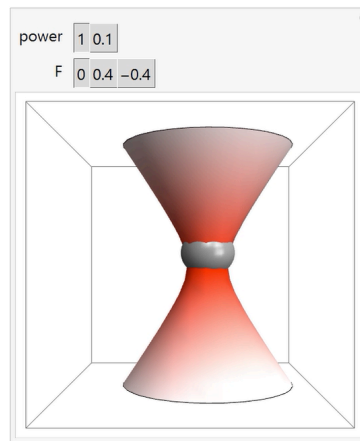


Outline

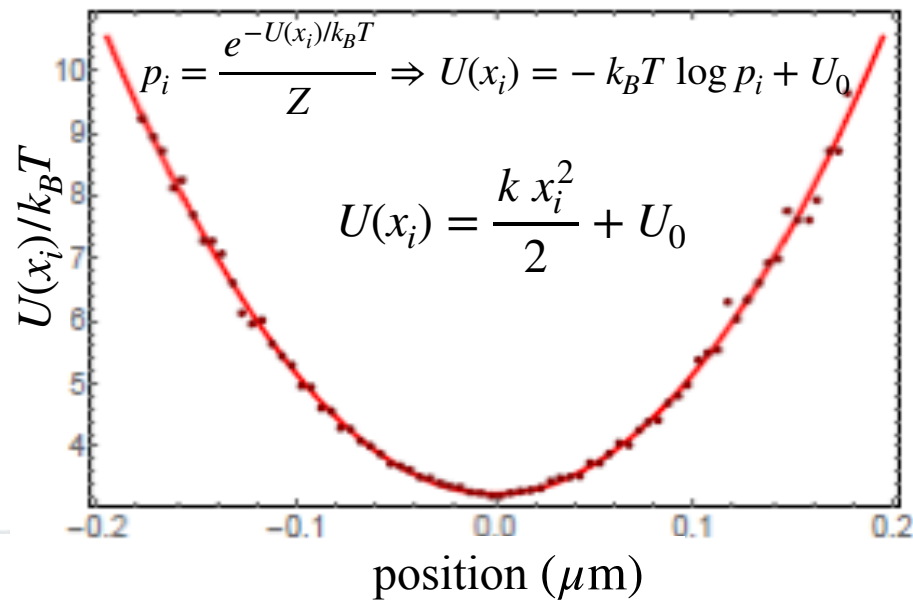
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measuring the fluctuations

Isolated trapped sphere:
Brownian fluctuations at
thermal equilibrium

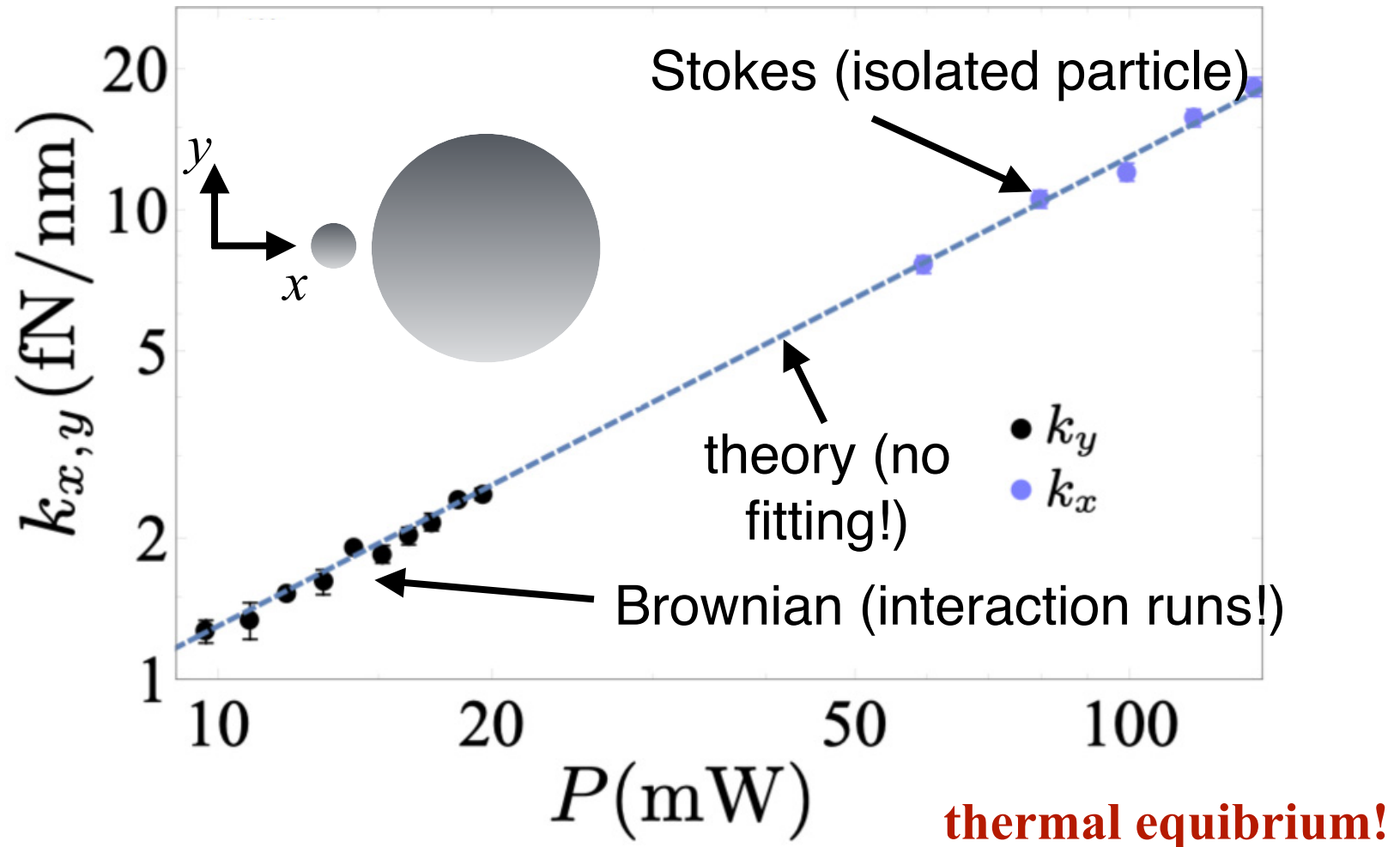


probability
distribution p_i $\lim_{N \rightarrow \infty} \frac{N_i}{\sum_{i=1}^N N_i} = p_i$



Brownian and Stokes calibrations

Since the trapping beam is circularly polarized, we expect $k_x = k_y$



measuring the interaction energy from fluctuations

each run: 5000 frames

sampling time = 100 ms

correlation time @ 1 micron: $\tau = \gamma_{\text{drag}}/k = 100$ ms

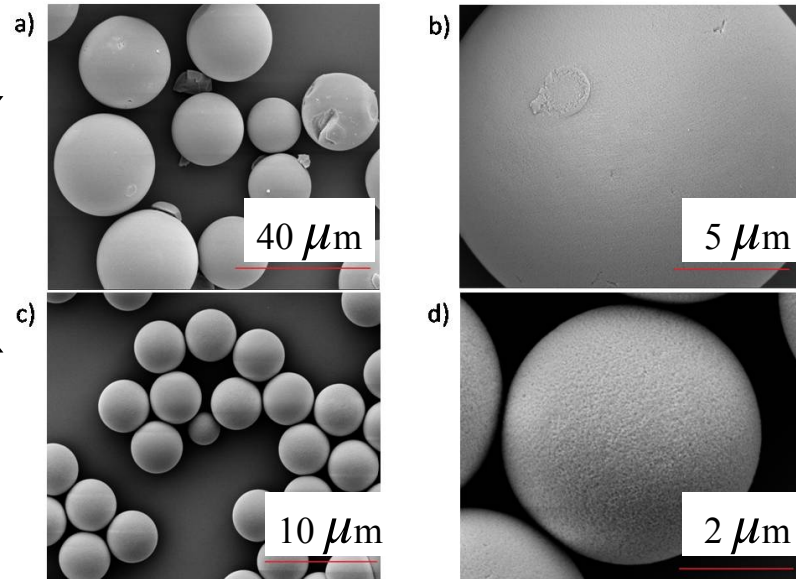
exposure time = 1 ms

HRSEM images of silica microspheres

Radii:

$$R_1 = (11,74 \pm 0,09) \mu\text{m}$$

$$R_2 = (2,35 \pm 0,02) \mu\text{m}$$



measuring the interaction energy from fluctuations

Low salt concentration:
Debye length

$$\lambda_D \sim 30 \text{ nm}$$

- * several runs
- * subtract optical potential

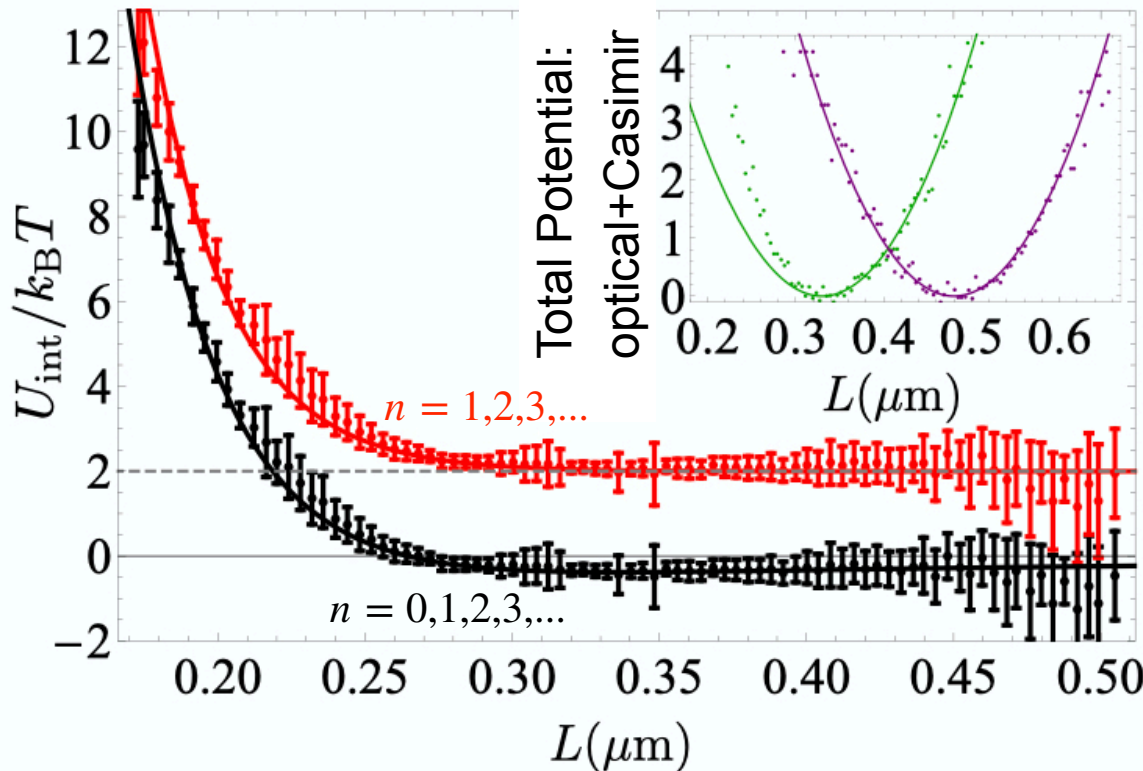
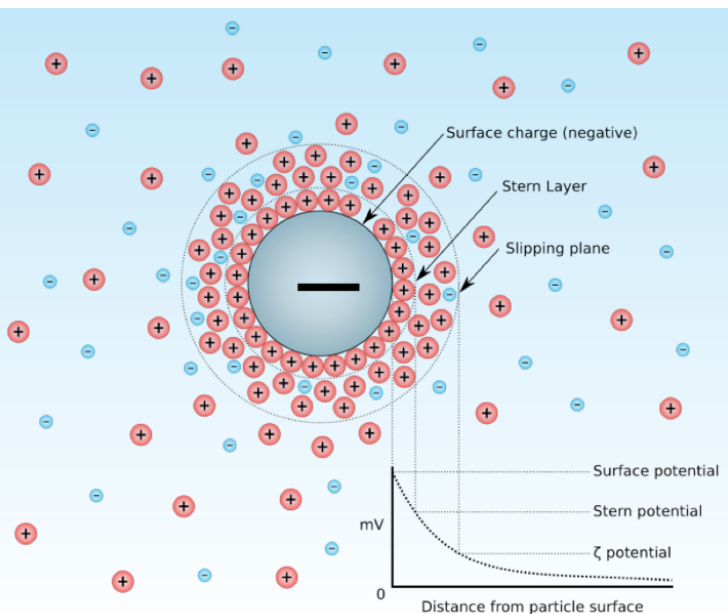


TABLE II: Parameters employed for the curve fit of the measured interaction energy: charge density σ and Debye screening length λ_D . In addition to the double-layer interaction energy (3), we also consider the Casimir interaction either with or without the zero-frequency TM contribution.

Casimir model	σ (mC/m^2)	λ_D (nm)
$n = 1, 2, \dots$	-1.7 ± 0.3	25.0 ± 0.9
$n = 0, 1, 2, \dots$	-0.8 ± 0.1	29.3 ± 0.6

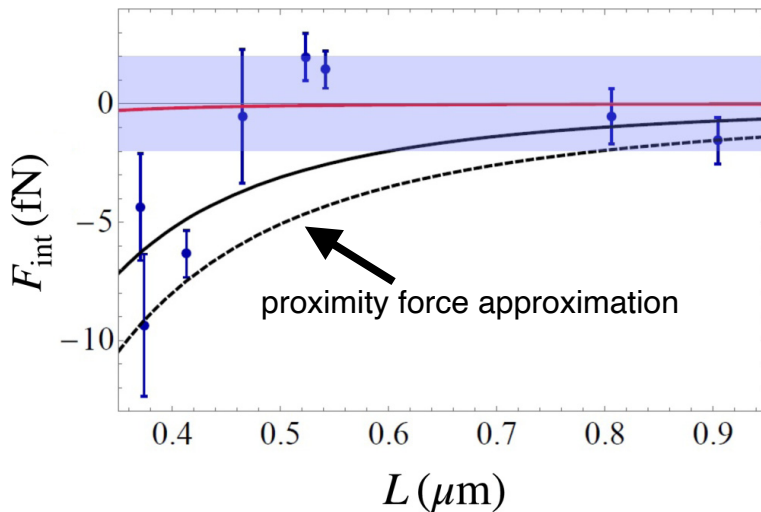
measuring the interaction energy from fluctuations

High salt concentration (0.22 M):
Debye screening length $\lambda_D = 0.64$ nm

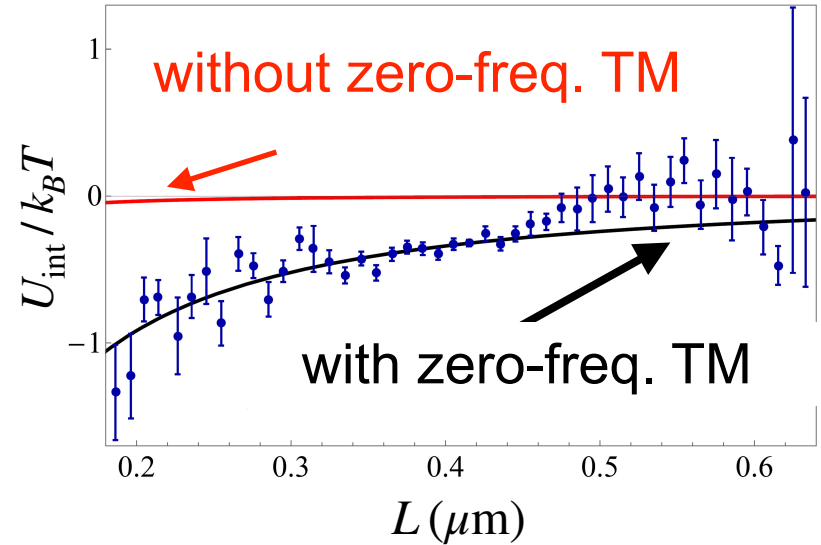
- * several runs
- * subtract optical potential

no fitting!!

Casimir force



Casimir potential versus distance



$$\left. \begin{array}{l} \text{shaded region} \\ \text{red line} \end{array} \right\} \sigma_x \sim 2 \text{ nm} \Rightarrow \sigma_F \sim 2 \text{ fN}$$

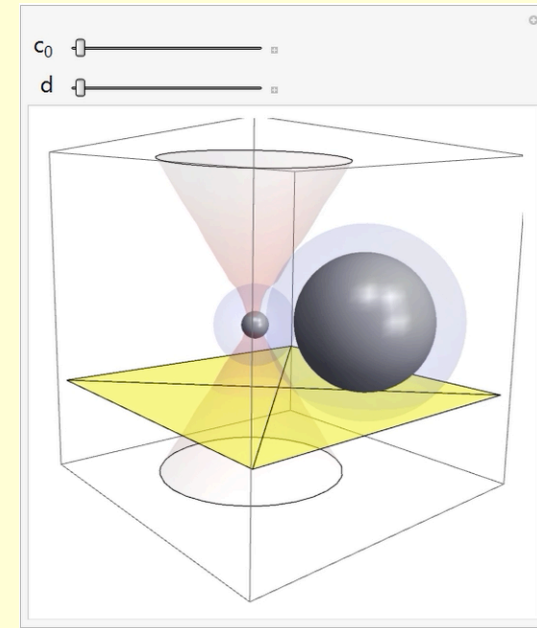
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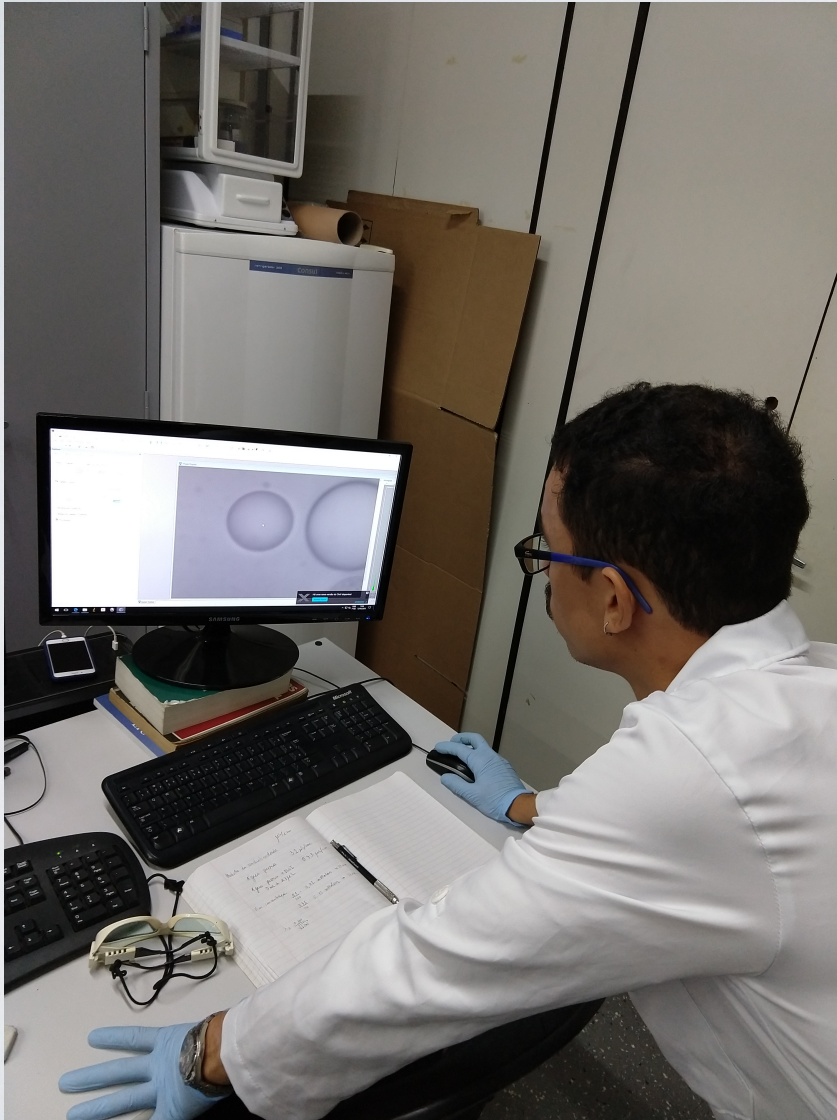
Conclusion

Casimir interaction between dielectric microspheres in aqueous solution: **contribution from transverse magnetic polarization at zero frequency**

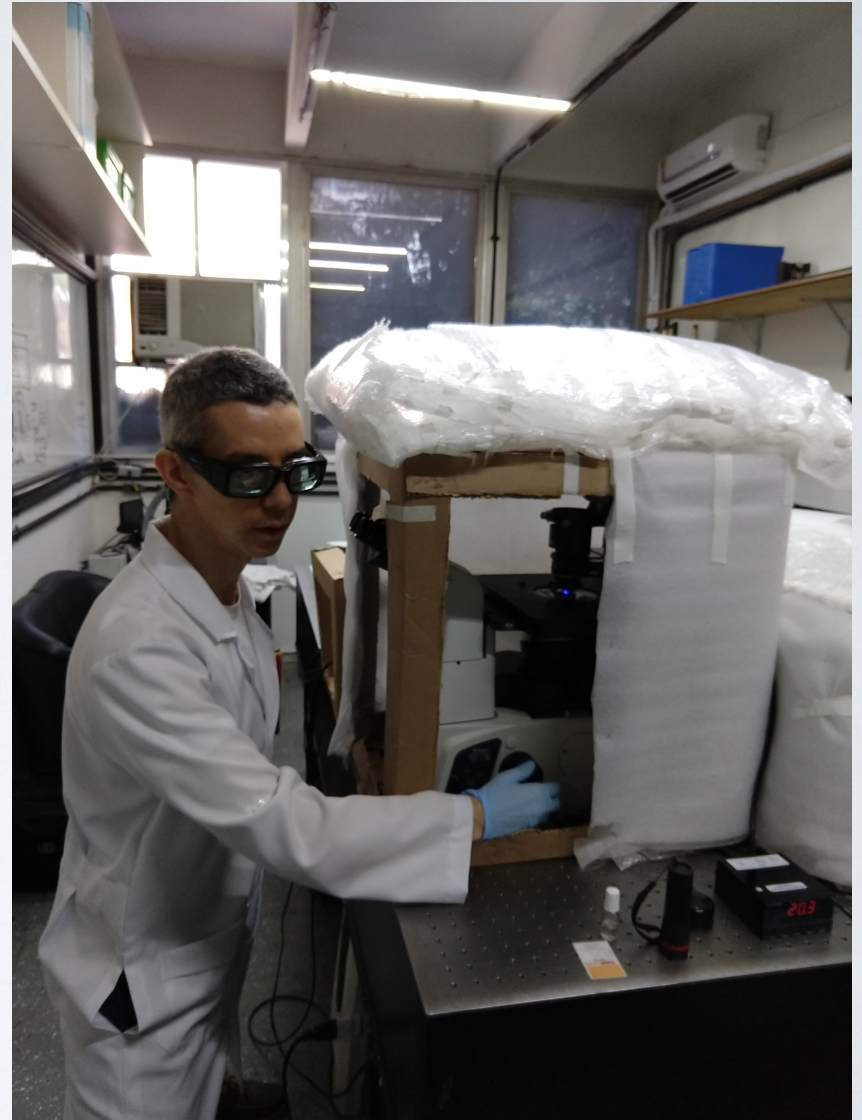
- unscreened !
- interaction is of a longer range than previously thought
- zero frequency dominant at distances > 100 nm
- universal function of geometric aspect ratios
- interaction energy $\sim k_B T$ at $L/R_{\text{eff}} \sim 0.1$
- measured with optical tweezers for silica microspheres in the distance range 200 - 500 nm



Measuring the Casimir force



Luis Pires



Diney Ether Jr

Optical Tweezers UFRJ



Luis Pires

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Nathan Viana

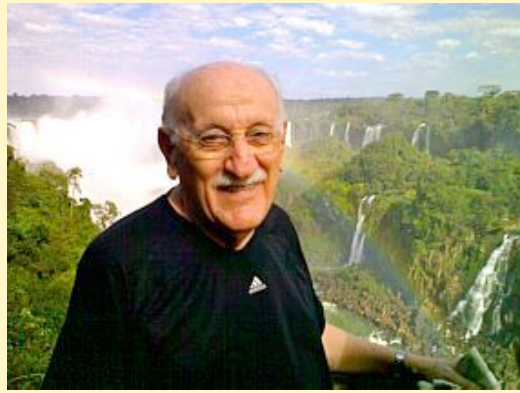
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Thank you!

Mie theory of image formation

aligning the sphere centers: image contains info about z

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Nonparaxial Mie Theory of Image Formation in Optical Microscopes and Characterization of Colloidal Particles

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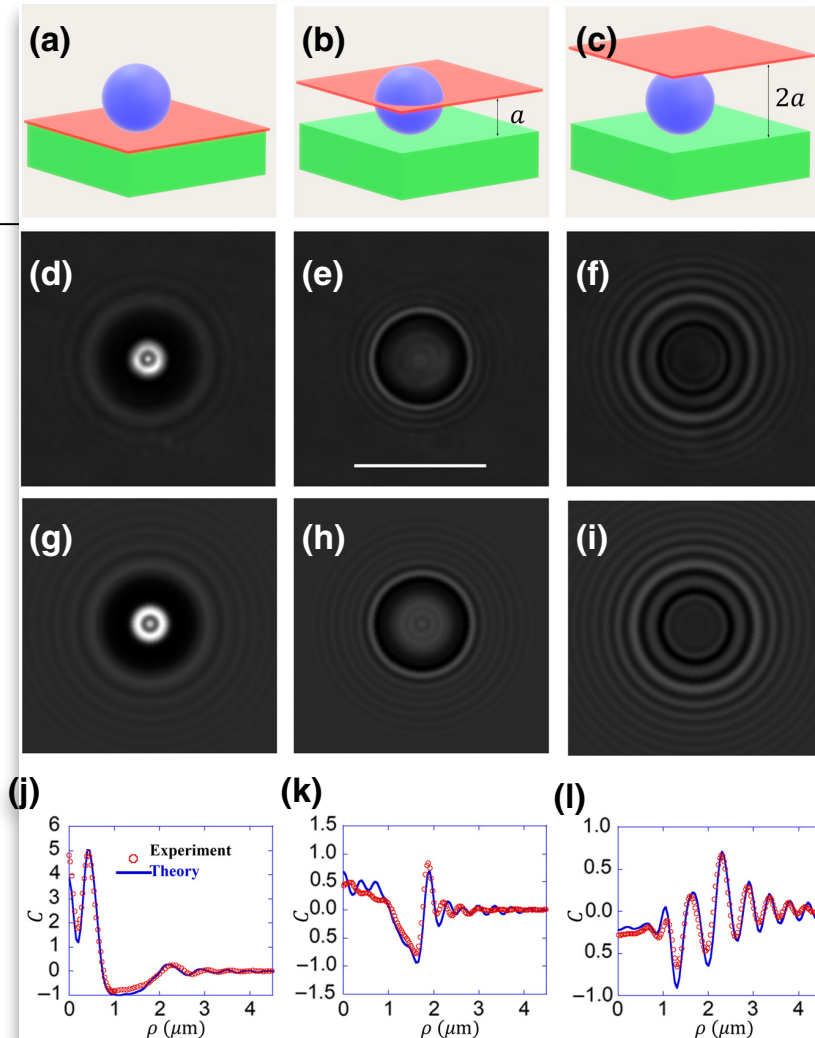
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sensitivity of the position measurement

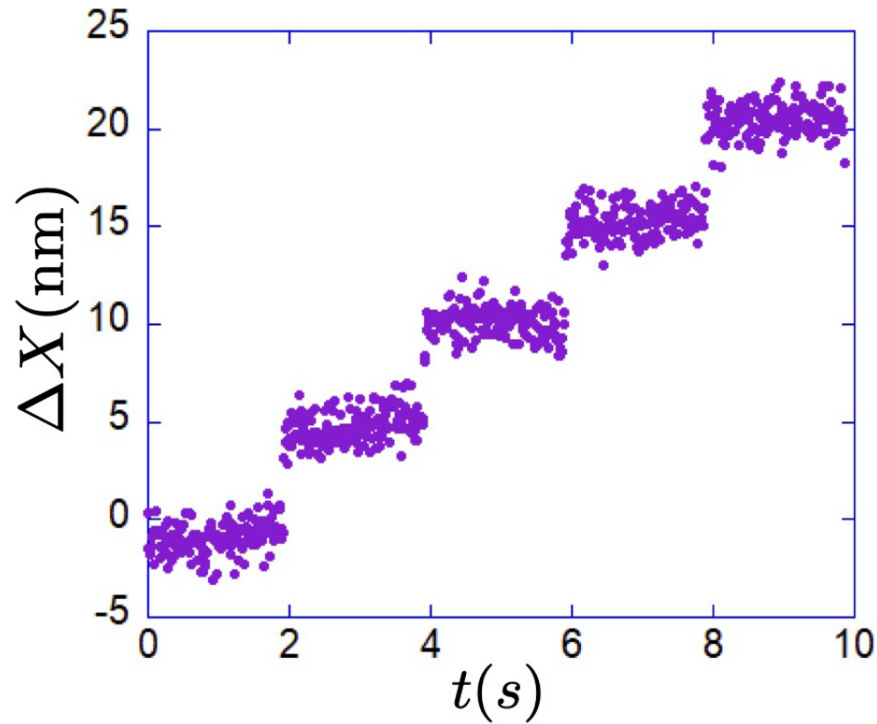


FIG. 7. Microsphere position versus time. A silica microsphere is adhered to the cover slip, which is driven laterally by 5 nm every 2 s with the help of a piezoelectric nanopositioning system.

Screening of the Casimir force

Numerical example

Polystyrene - aqueous solution - Polystyrene $T = 293$ K

Lorentz model for water and polystyrene - parameters taken from P.J. van Zwol and G. Palasantzas, PRA **81**, 062502 (2010)

$$\text{Hamaker constant : } H = -12\pi L^2 \frac{\mathcal{F}_{PP}}{A}$$

