

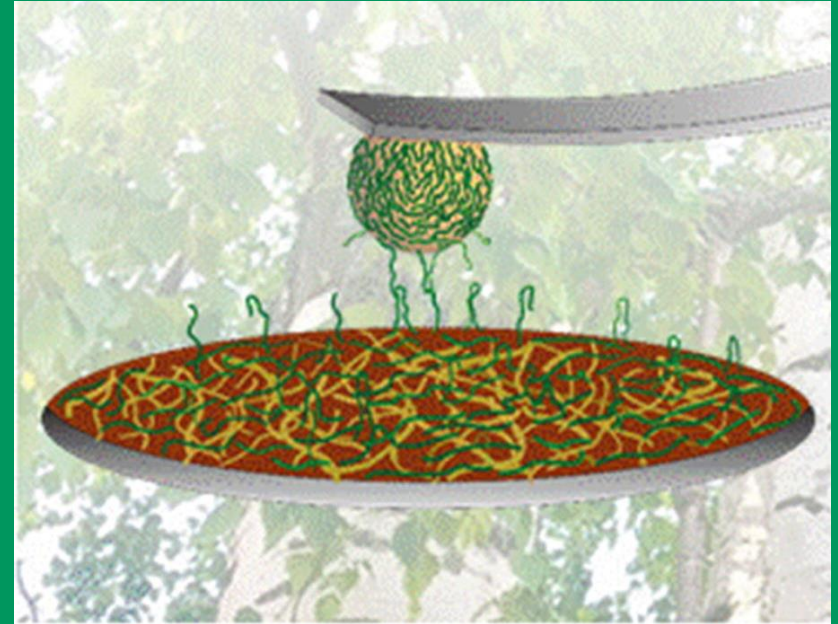
Atomic force microscopy II: Colloidal Probe

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4.3.2022



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Engineering



Learning outcomes

After this lecture you

- **Understand why is it useful to measure direct surface forces and friction forces**
- **Know the main requirements to get reliable results**
- **You are familiar with various force measuring strategies**
 - Ranging from chemical force spectroscopy to single cell force spectroscopy – focus on colloidal probe microscopy

Why measure forces?

Have you measured surface forces using AFM?
Have you measured interactions in any other way?
Are surface forces relevant in your research?

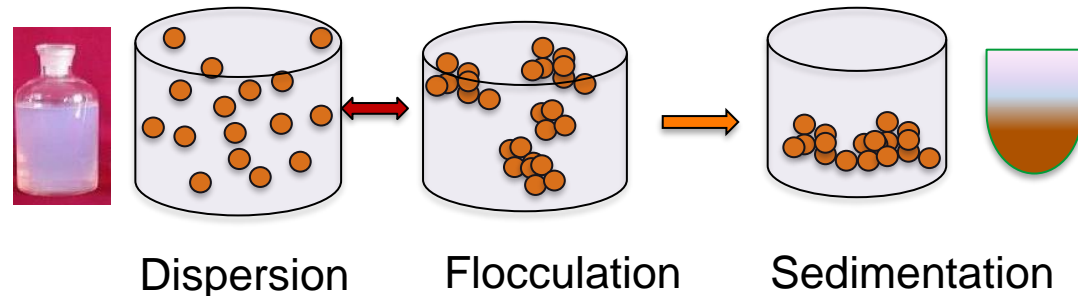
Nano vs. colloidal vs. macroscopic

- In nanoparticle dispersions the surface forces between particles determine if the particles aggregate or stay well dispersed

We can predict (and affect) flocculation and stability of dispersions.

Our knowledge of interaction of e.g. charged, hydrophobic or polymer coated particles in aqueous solutions are based on careful surface force measurements.

For more complex systems measurements are needed to increase our understanding.



The basic principle of force measurements

The base of a spring is moved by known amount ΔD_0 .

Due to forces between the surfaces the spring deflects ΔD_s while the surface separation changes by ΔD .

$$\Delta D_s = \Delta D_0 - \Delta D$$

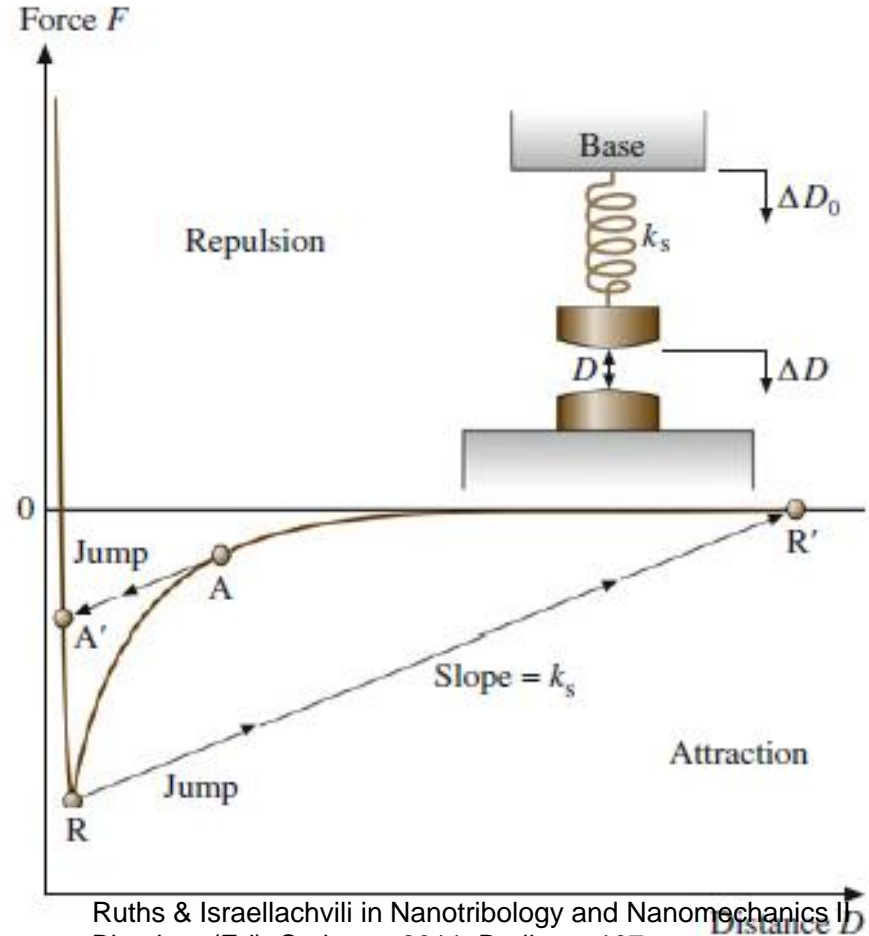
The difference in force, between the initial and final separation is given by

$$\Delta F = k_s \Delta D_s$$

Where k_s is the spring constant

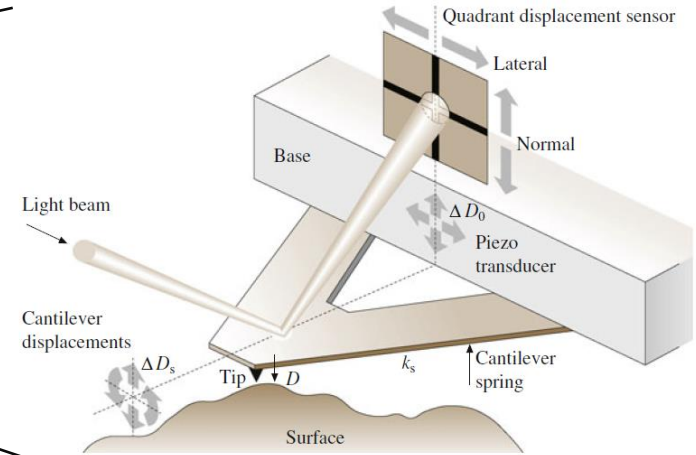
By measuring ΔF at various distances from zero force to hard wall contact the whole force curve (law) can be constructed.

The force law $F(D)$



Ruths & Israellachvili in Nanotribology and Nanomechanics I
Bhushan (Ed), Springer, 2011, Berlin, pp107.

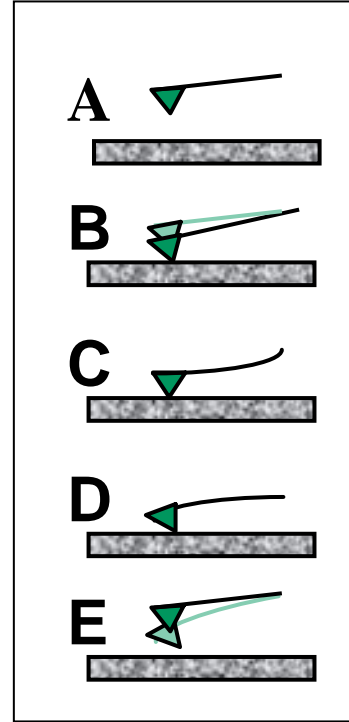
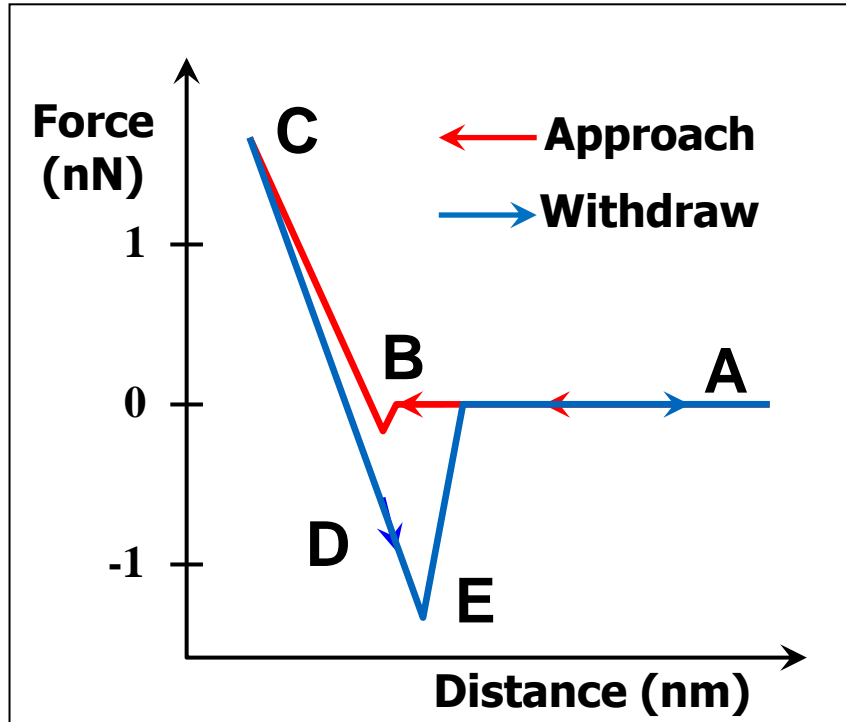
Atomic force microscopy



Ruths & Israellachvili in Nanotribology and Nanomechanics II Bhushan (Ed), Springer, 2011, Berlin, pp107.

Not just imaging – Also direct quantification of forces between surfaces

Principle of force spectroscopy



Direct surface force measurements – Information on specific and non-specific interactions



 Chemical force microscopy (CFM)

COO-

 Single-molecule force spectroscopy (SMFS)

 Colloidal probe microscopy (CPM)

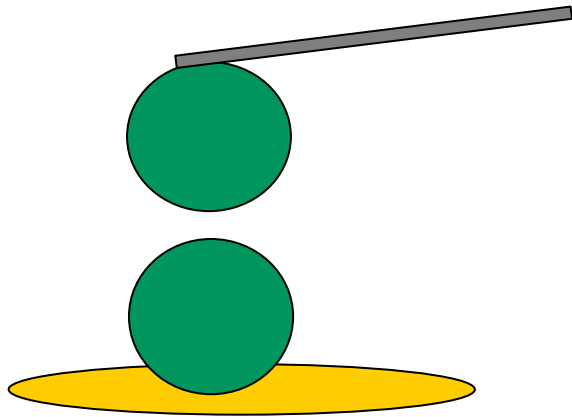
Cellulose
sphere

 Single-cell force spectroscopy (SCFS)

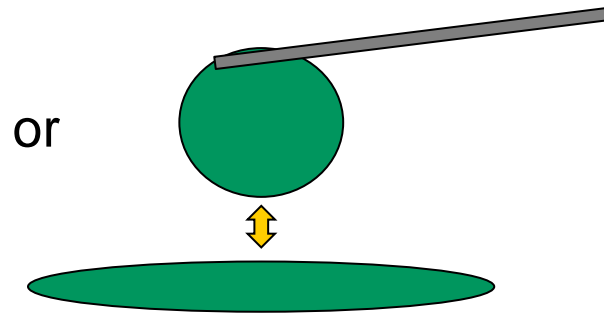
Colloidal probe microscopy

Well-defined geometry facilitates correlation to theory

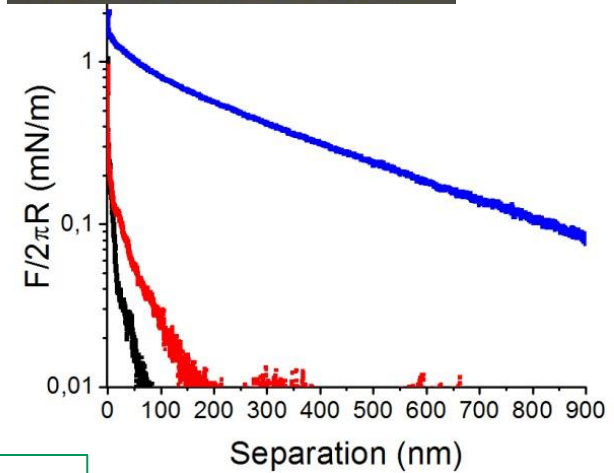
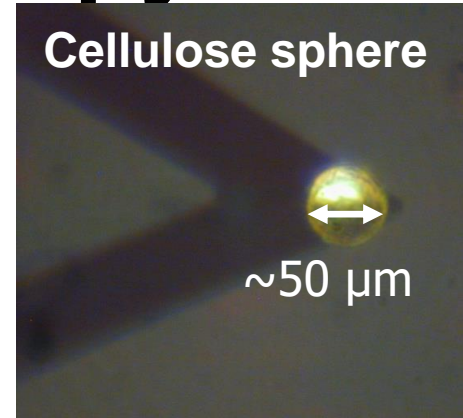
Sphere against sphere



Sphere against plane



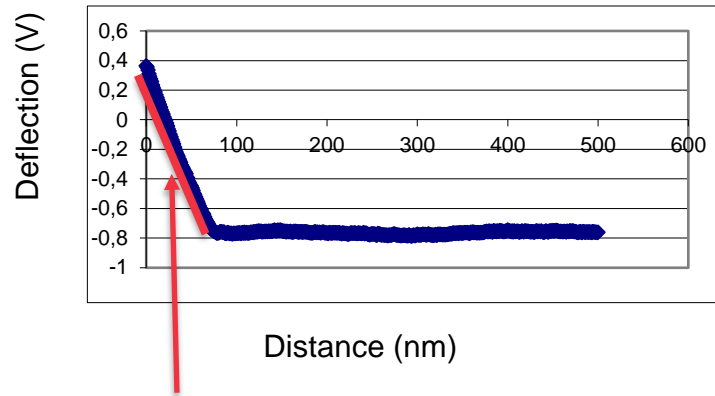
or



Very simplified: From the shape of the force curve we can say what forces are important in the system: electrostatic, steric, hydrophobic,...

AFM force spectroscopy in practice

Raw data = deflection as a function of relative distance



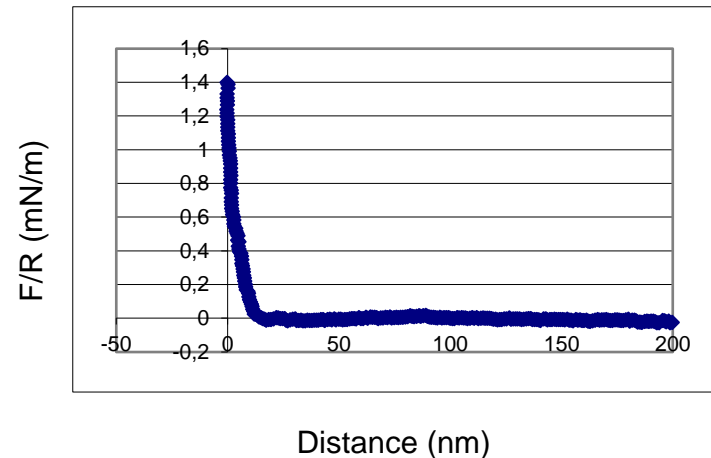
Sensitivity, spring constant, radius of sphere(s) has to be determined



For soft materials the sensitivity has to be determined on a hard substrate

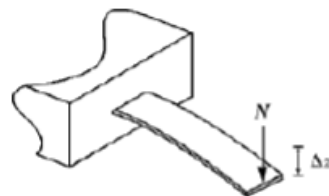
How do you get from raw data to force as a function of distance between probe and substrate?

Force as a function of distance between probe and substrate



Determination of normal spring constant

$$\text{spring constant } k_z = \frac{N}{\Delta z} \quad \text{Force/deflection}$$
$$(k_s = \frac{F}{\Delta D})$$

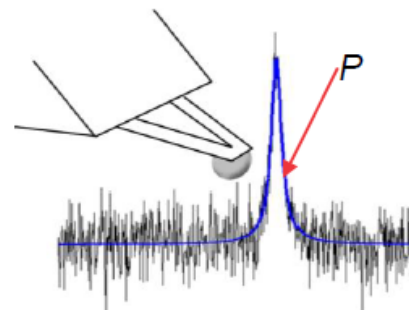


The thermal noise method

Measures the resonance frequency peak determined from the analysis of the thermal fluctuations of the cantilevers

$$\text{spring constant } k_z = \frac{kT}{P}$$

k = Boltzmann's constant, T = Temperature, P area of the power spectrum of thermal fluctuation



<https://doi.org/10.1016/j.colsurfa.2013.11.018>

The Sader method

In addition to the thermal fluctuation and sensitivity, the dimensions of the cantilever needs to be known

$$k_z = 0.1906\rho b^2 L Q_f \omega_f^2 \Gamma_i^f (\omega_f)$$

ρ is the density of the fluid, b and L are the width and length of the cantilever, respectively, Q_f is the quality factor and Γ_i^f is the imaginary component

A!

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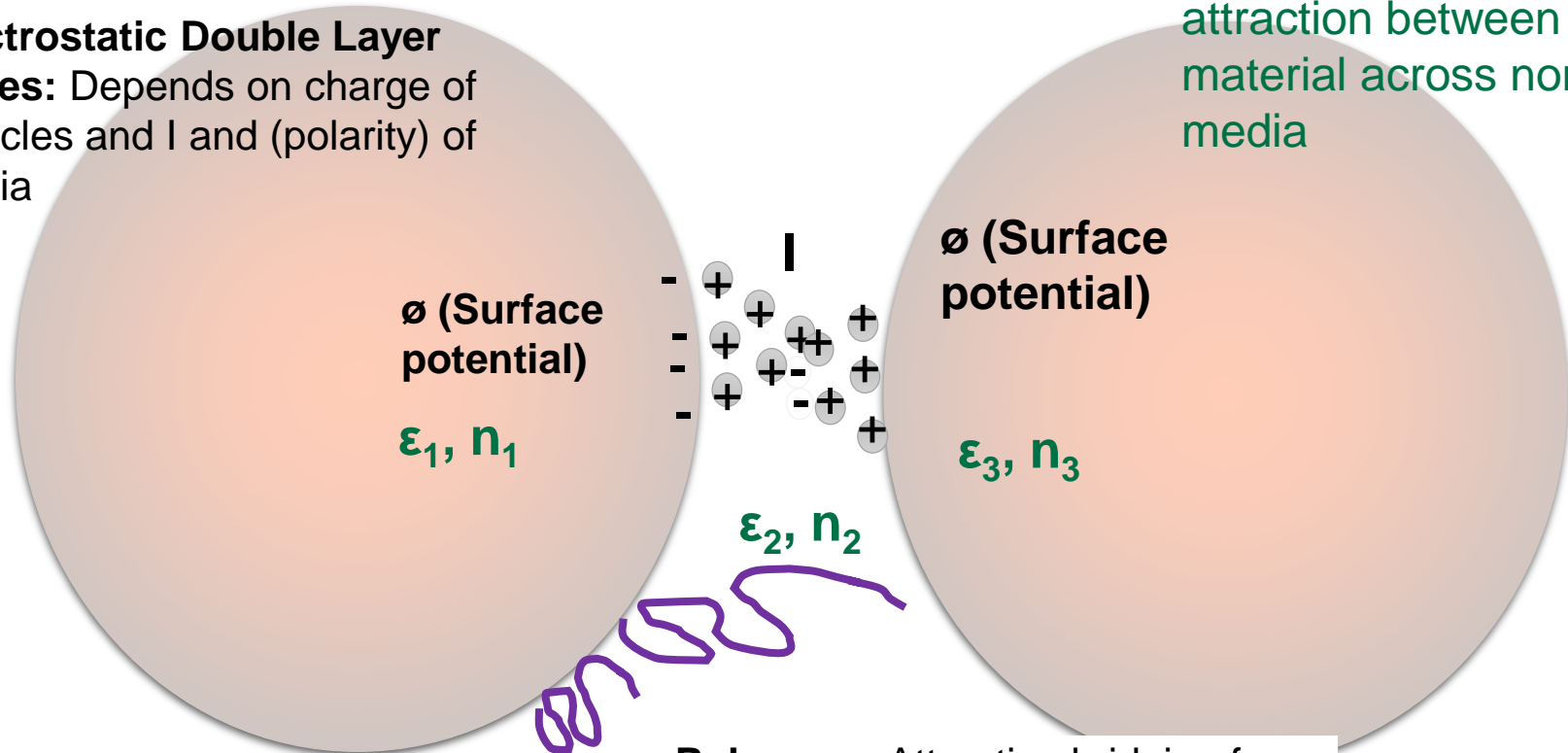
Surface forces: *A very simplified* overview

Surface forces mind map

vdW forces: Depends on polarity of particles and media. Highest attraction between polar material across nonpolar media

Electrostatic Double Layer

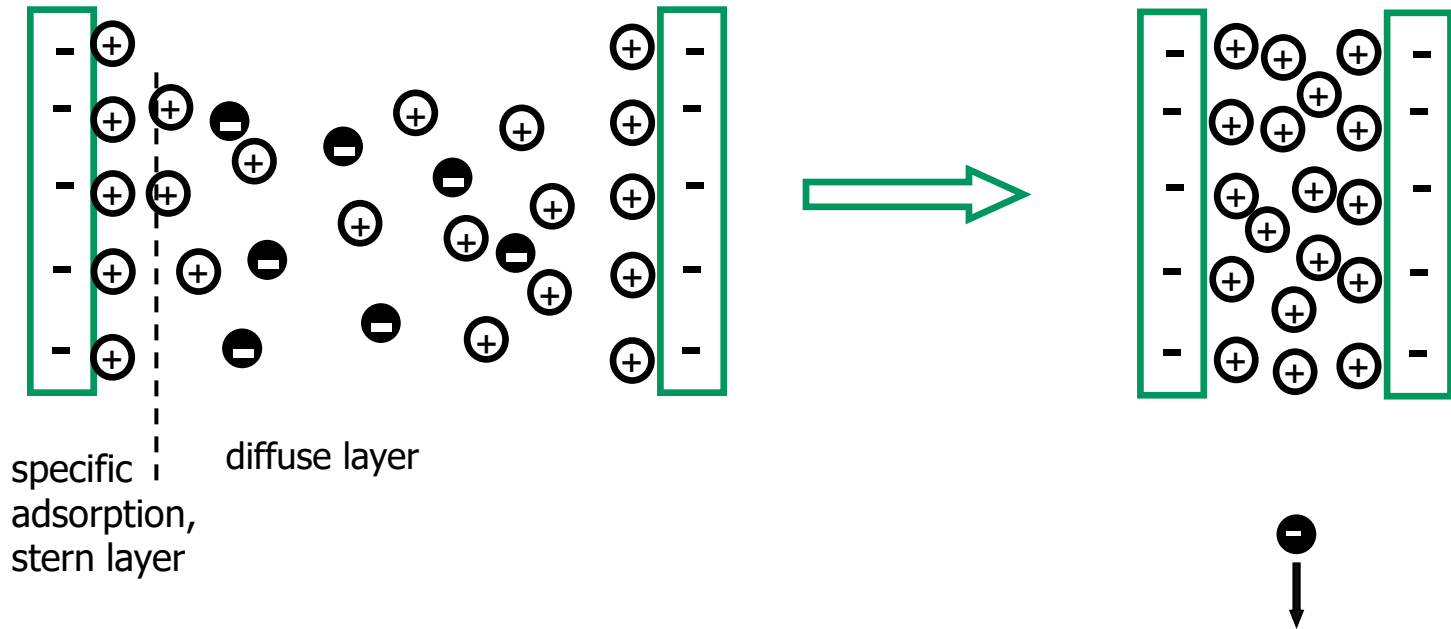
forces: Depends on charge of particles and I and (polarity) of media



Polymers: Attractive bridging force or steric repulsion. Depends on coverage and interactions between polymer and solvent.

Double layer force – interaction of (like) charged particles

surface potential, Φ_0



specific adsorption, stern layer

diffuse layer

A cloud of ions is found close to a charged surface. Counterions are enriched, co-ions are depleted. The double layer is related to the osmotic pressure due to the overlap of ion clouds

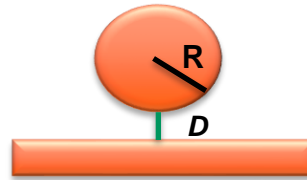
DLVO theory - summary

DLVO-theory

- Derjaguin, Landau, Verwey and Overbeek
- $F_{\text{tot}} = F_{\text{van der Waals}} + F_{\text{electrostatic}}$ (F = force)

A_H = Hamaker constant, a measure of the “polarity difference” between particles and media

$$F_{\text{vdW}} = -A_H R / 12D^2$$



$$F_{DL} \approx 2\pi\epsilon_0\epsilon_r R\kappa\phi^2 \exp(-\kappa D)$$

Debye length, $\kappa^{-1} = \sqrt{\frac{\epsilon_0\epsilon_r kT}{2e^2 I N_A}}$

R = radius of sphere

D = distance between surfaces

k = Boltzmann constant

T = temperature

ε = static dielectric constants for the three media

A_H = Hamaker constant for the system

ε_0 = vacuum permittivity

ϕ = Surface potential

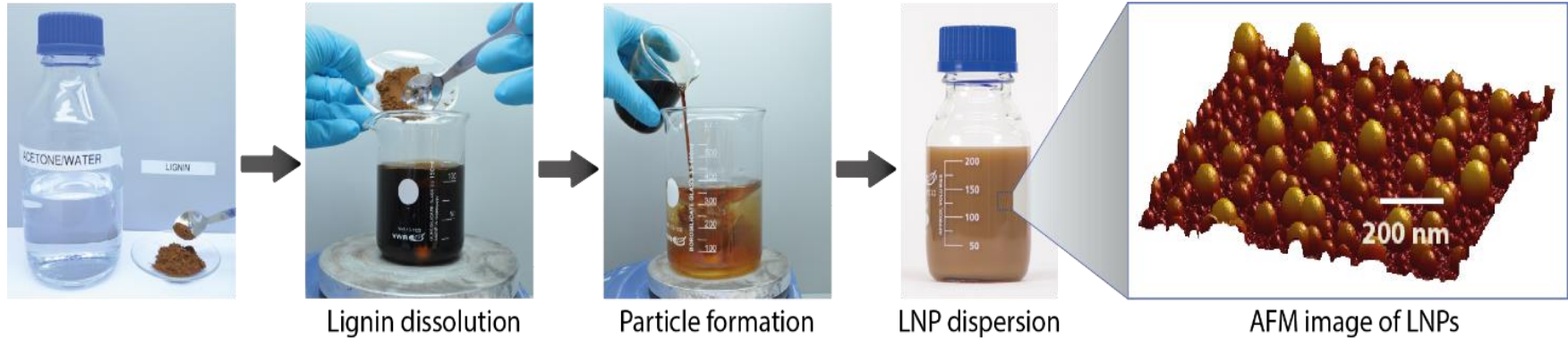
N_A = Avogadro's constant

I = ionic strength $I = \frac{1}{2} \sum_i z_i^2 c_{0,i}$

z = valency of ion

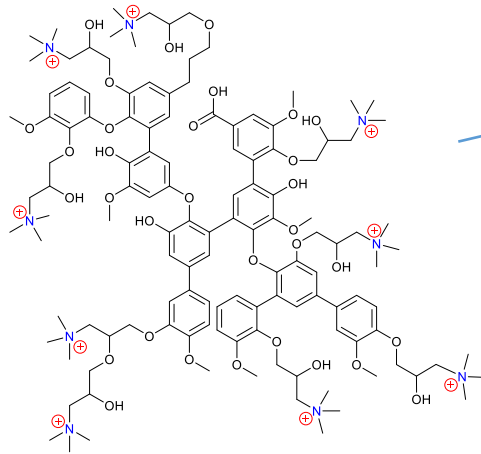
C_0 = concentration of ion i

Colloidal lignin particle (CLP) preparation via nanoprecipitation



Produces **stable** aqueous dispersion of **spherical** lignin nanoparticles (diameter~100 nm)

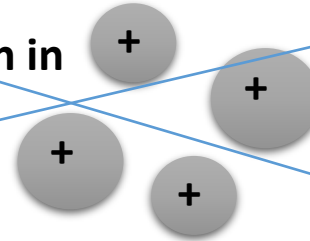
How to make cationic lignin particles?



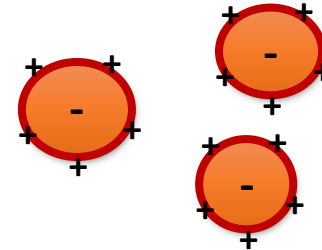
Cationic lignin

Reaction with
glycidyltrimethylammonium
chloride

~~Direct precipitation in
antisolvent~~

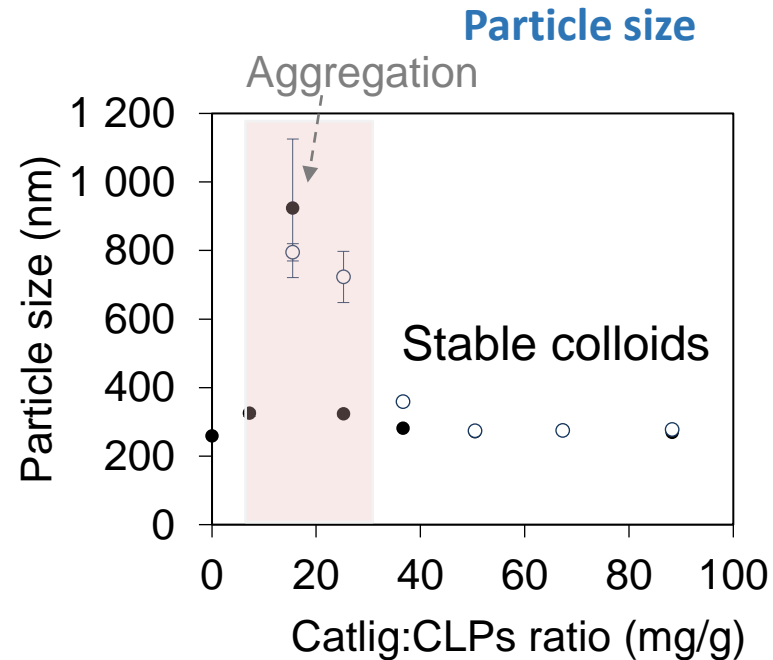
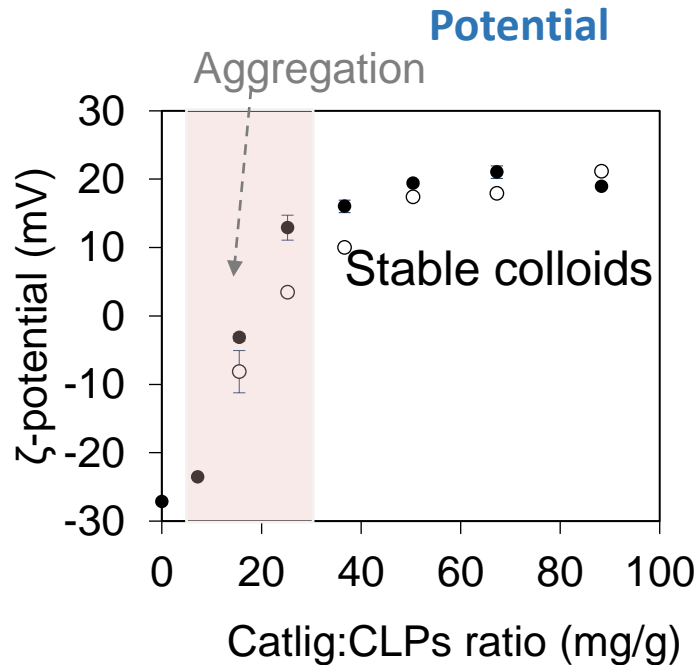


Adsorption to
anionic colloidal
lignin particles



- Stable
- High cationic charge possible
- Minimum amount of nonrenewable chemicals

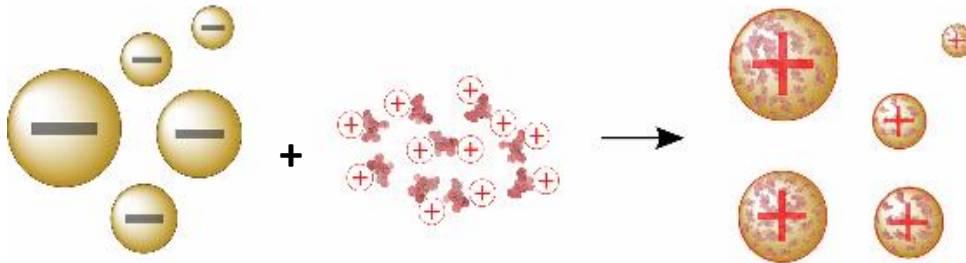
Dispersion stability of cationic lignin particles



Explanation



**Low surface coverage
→ Charge
neutralization**



**High surface coverage
→ Stable cationic
particles,
Overcompensation**

Derjaguin approximation

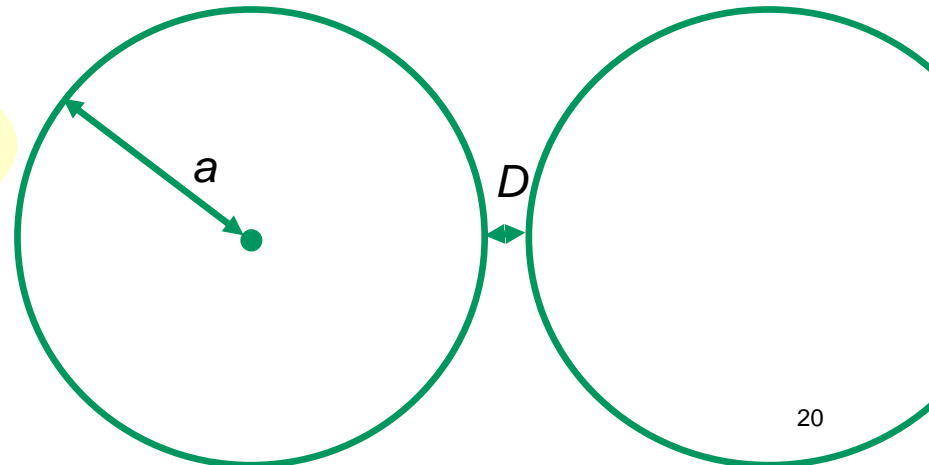
Why do we normalize with the radius of the sphere(s)?

The interaction energy between two flat surfaces =

$$W(D) = \frac{F_{spheres}(D)}{2\pi \left(\frac{a_1 a_2}{a_1 + a_2} \right)} = \frac{F_{cylinders}(D) \sin \theta}{2\pi \sqrt{a_1 a_2}} = \frac{F_{sphere+flat.surf}(D)}{2\pi}$$

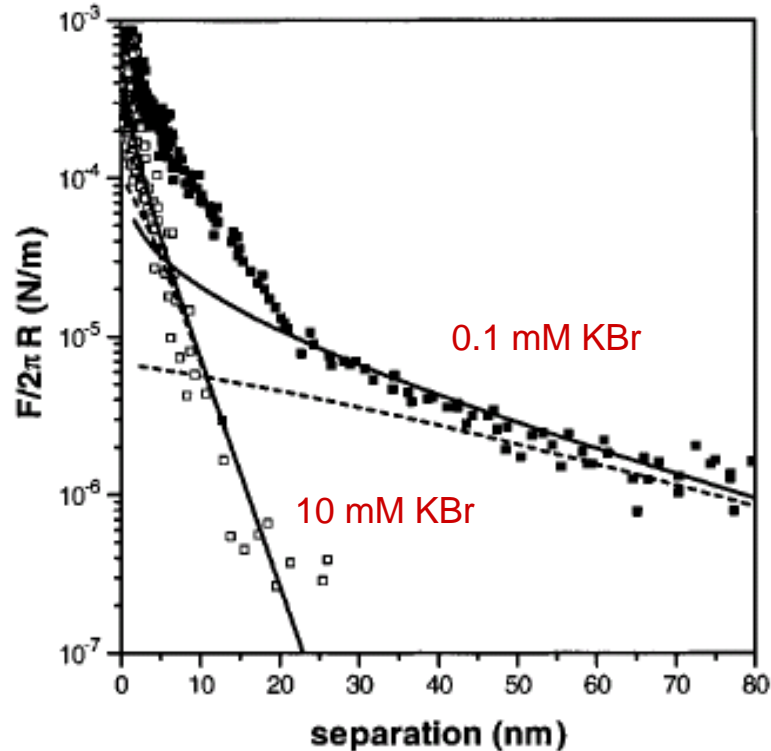
- Valid if D (distance between the surfaces) $\ll a$ (radius)

Normalisation makes it possible to compare measurements

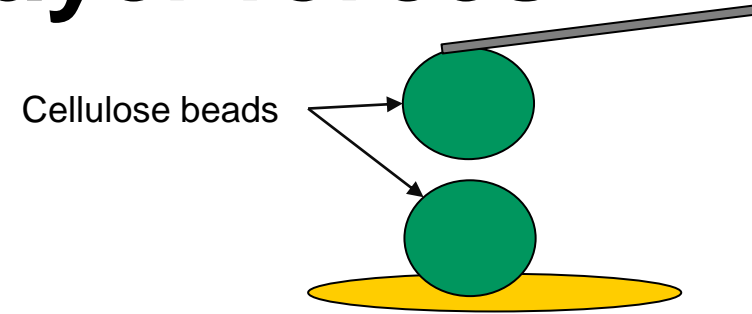


Examples

Electrostatic double layer forces



Carambassis and Rutland Langmuir, 1999



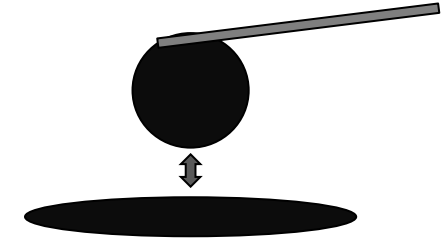
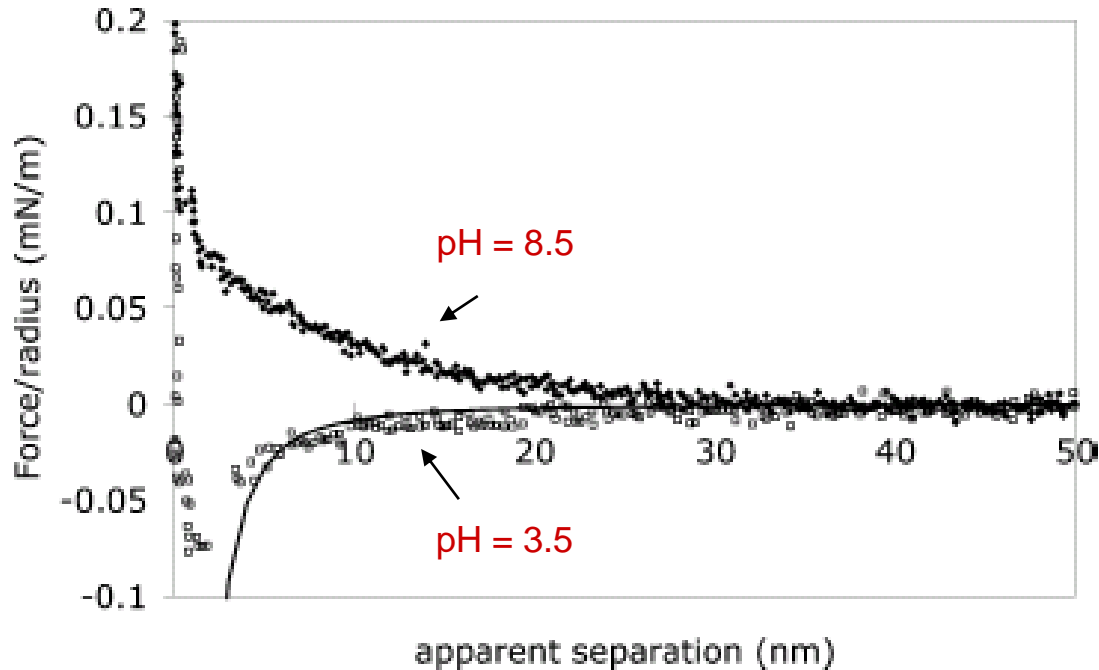
Electrostatic and steric forces were observed

pH about 7

Why do we not observe van der Waals attraction at short distances?

Steric repulsion at short separation

Van der Waals forces between cellulose surfaces



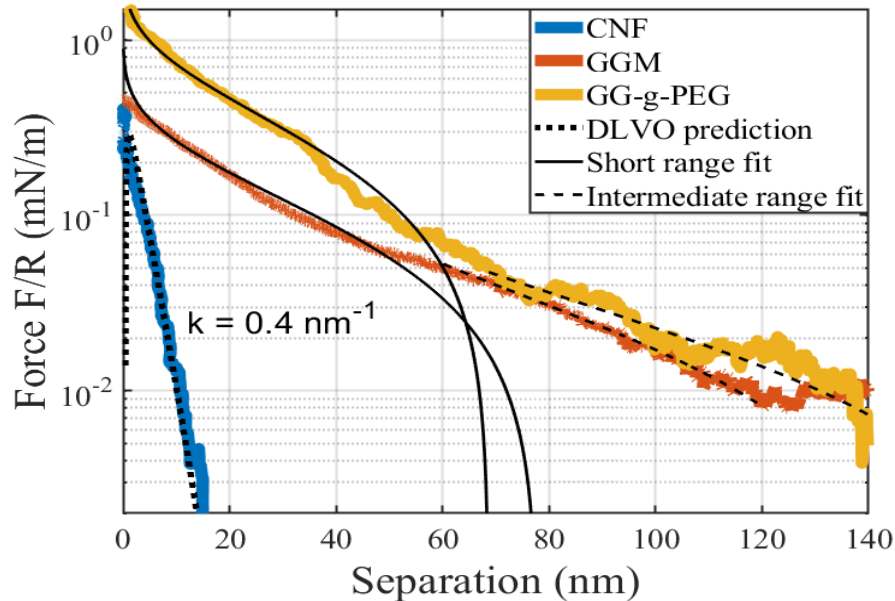
One cellulose sphere
against a cellulose film,
1 mM NaCl

Why purely attractive at low
pH and repulsive forces at
higher pH?

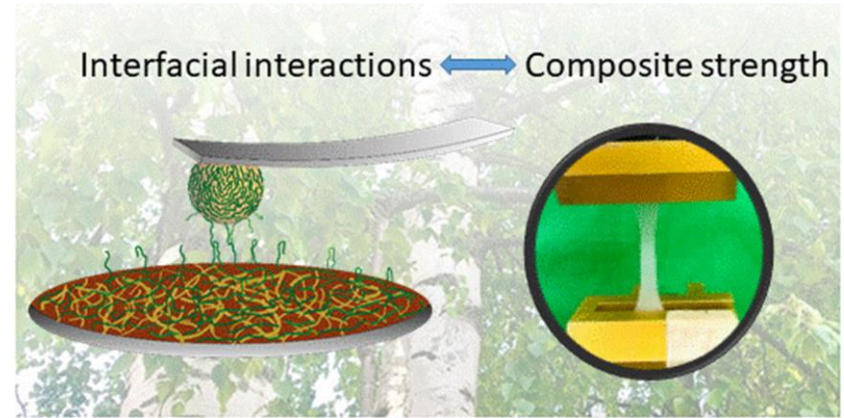
Notley et al., Langmuir 2006

<https://pubs.acs.org/doi/10.1021/la052886w>

Polysaccharides adsorbed onto cellulose – Steric forces



Brush length of adsorbed polymer layer and correlation to mechanical properties



CNF –cellulose bead before adsorption:

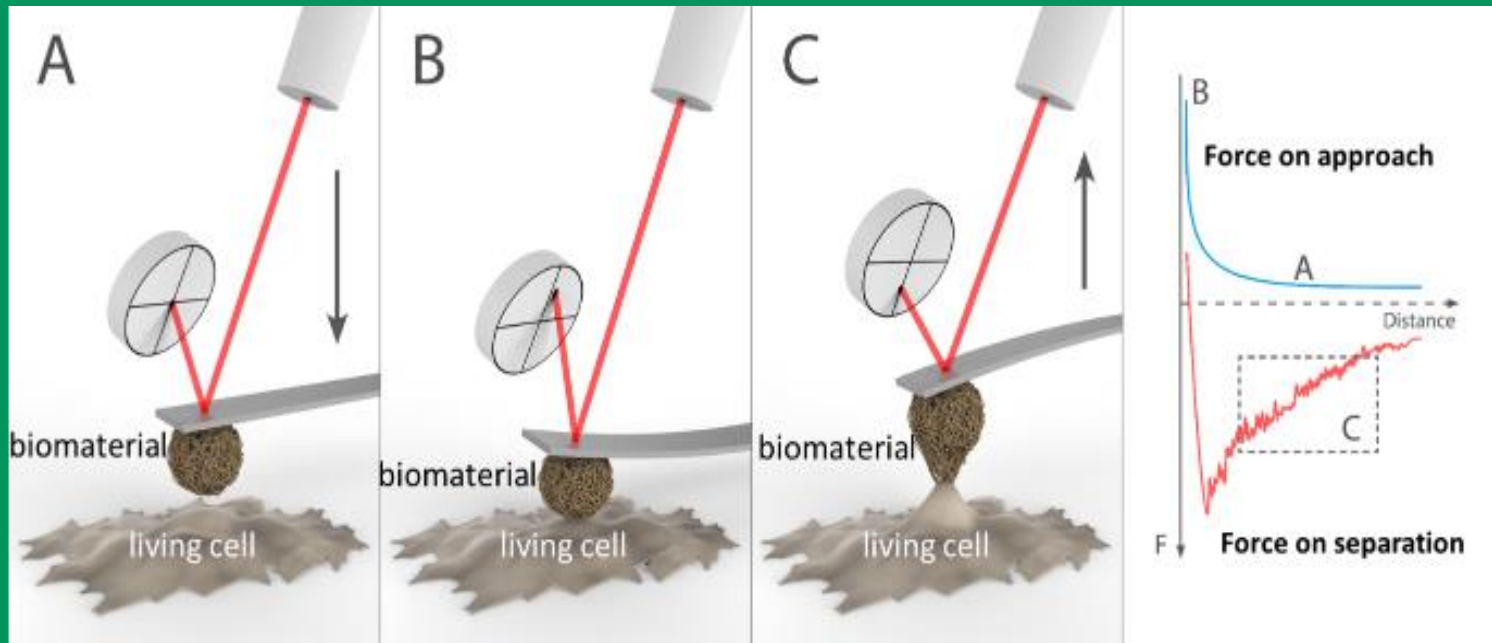
DLVO prediction ok

For the other systems: mainly steric forces due to adsorbed polysaccharides

Some reflections

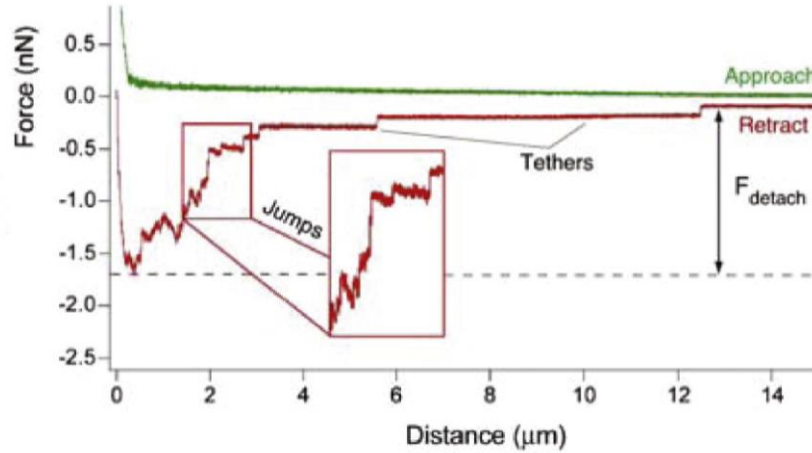
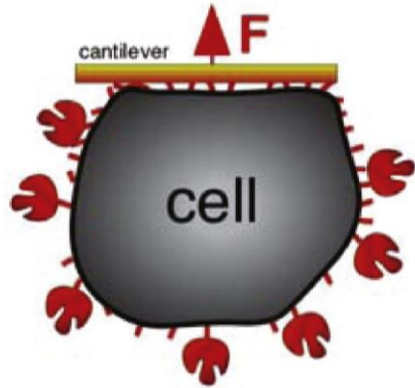
- **Force curve and effect of ionic strength tells us what forces dominate**
- **We can determine surface potential via fitting to DLVO theory**
- **We can estimate structure of adsorbed polymer layer, like brush length or brush vs mushroom structure via fitting to polymer brush model**

Interactions between living cells and biomaterials

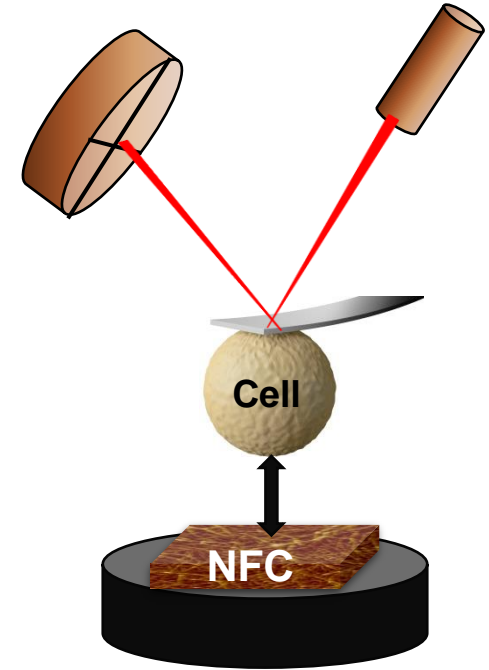


Commonly used method: Single cell force spectroscopy

(c)



Current Opinion in Biotechnology



Muller et al <https://doi.org/10.1016/j.copbio.2009.02.005>

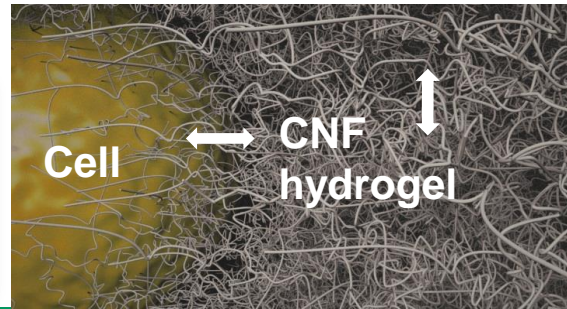
SCFS gives understanding of:
Molecular mechanism of cell adhesion
Specific and non specific interactions

Forces between stem cells and biomaterials

Challenge: stem cells cannot survive alone -> Colloidal probe approach

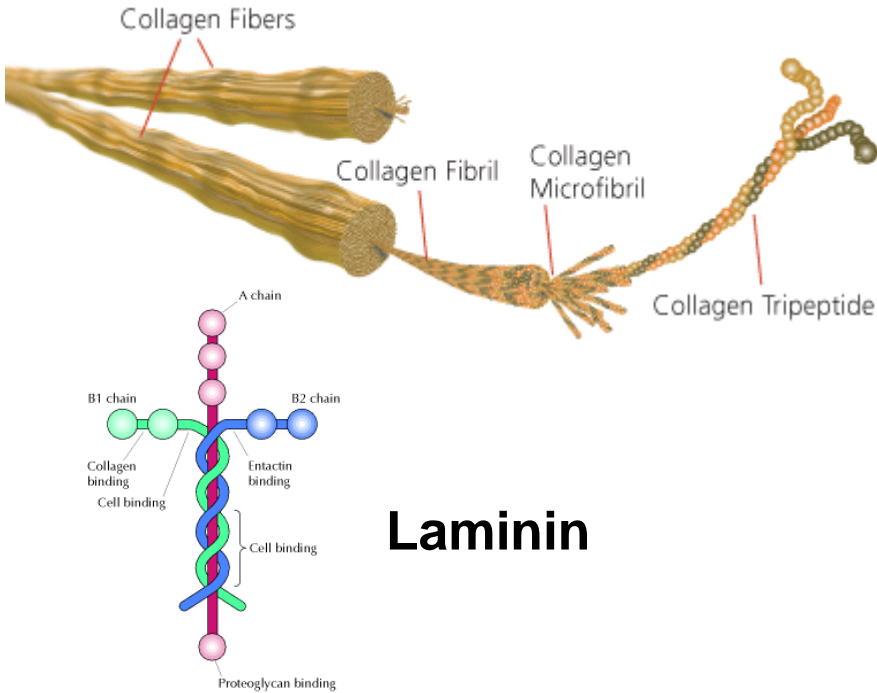
- To give tools for better material design for tissue engineering
- To correlate direct interactions forces between living cells and biomaterials with phenomena found during cell culture *in vitro*

Cellulose nanofibrils, CNF



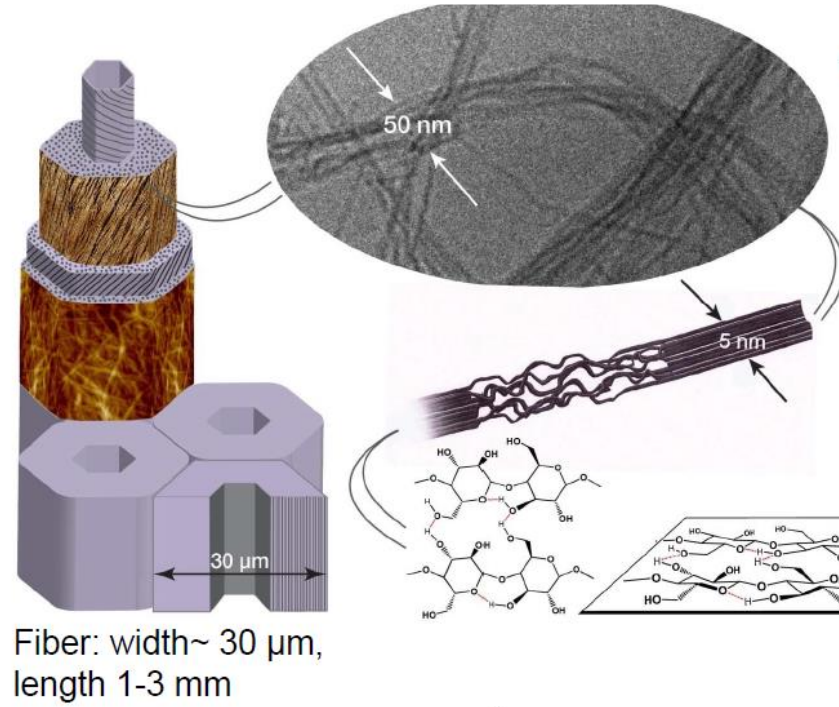
Choice of biomaterials

Collagen I and IV



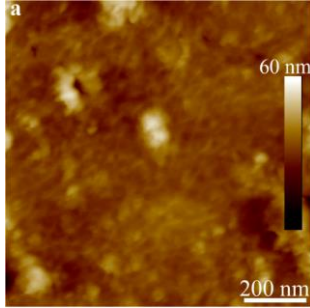
Laminin

Cellulose nanofibrils from wood

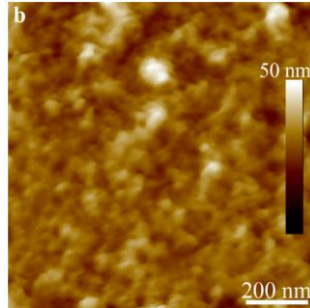


The system

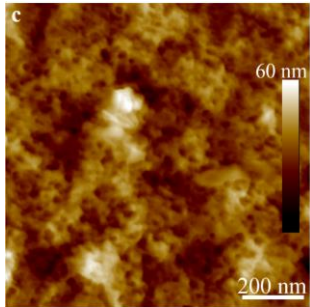
Biomaterials coated on probe



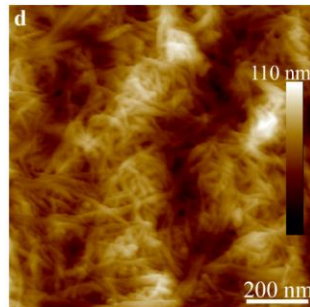
Collagen I



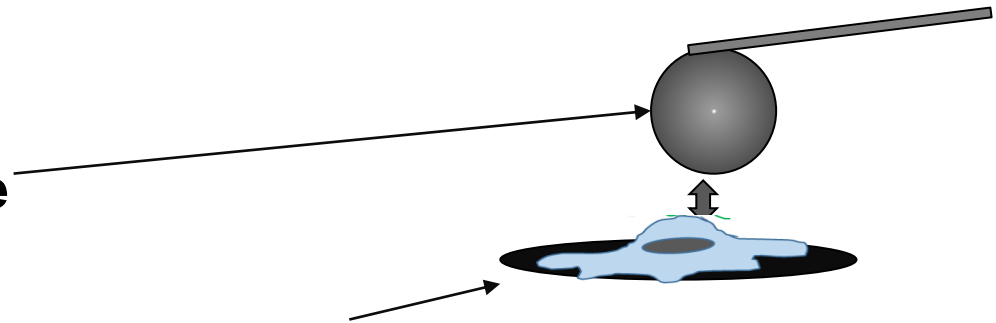
Collagen IV



Laminin 521

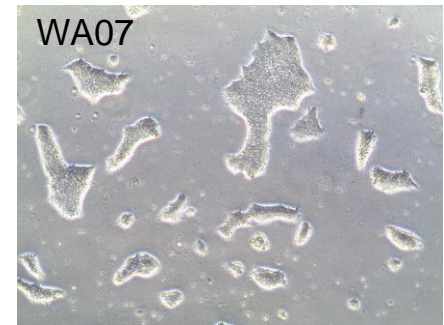
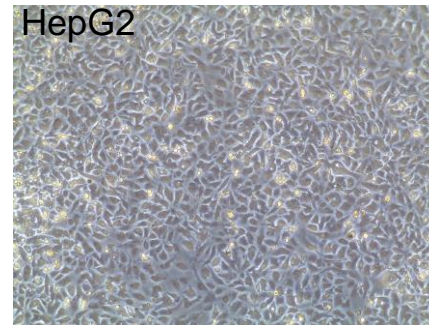


CNF (Growdex[®])



Cell types

WA07 (human embryonic stem cell line)
HepG2 (human hepatocellular carcinoma cells)

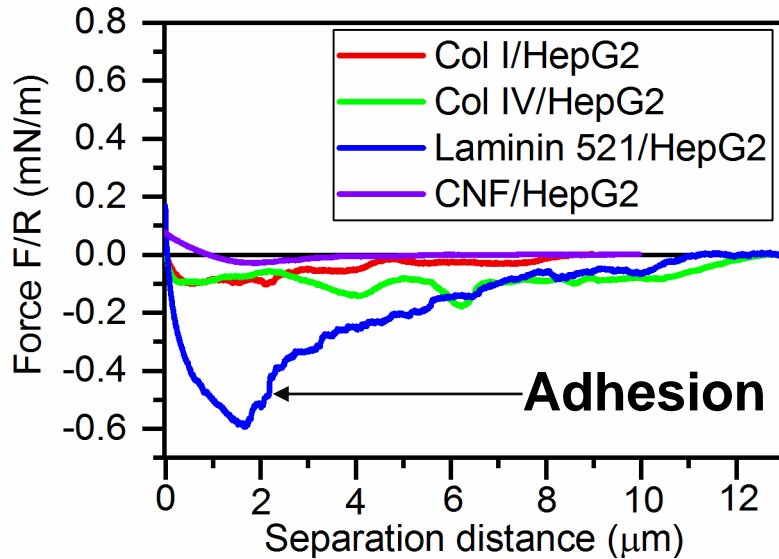


Interactions between *living* cells and biomaterials – Effect of cell line

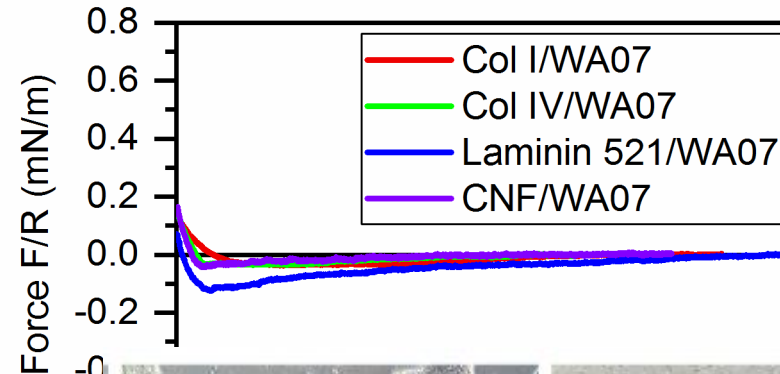


30s in contact
Retraction curves

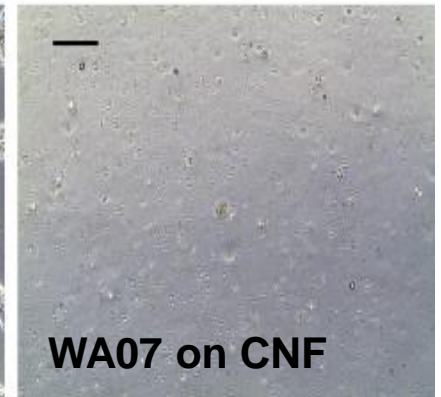
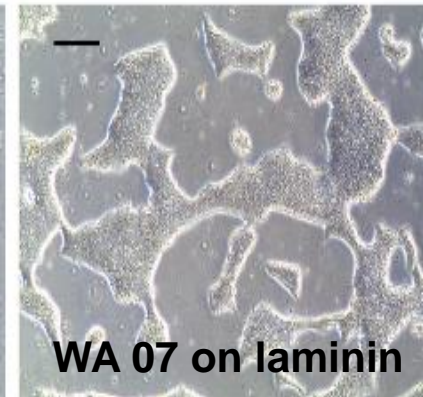
HepG2 (carcinoma cells)



WA07 (stem cells)



Very long ranged adhesion
Stem cells have weaker interaction
than commonly used cancer cells

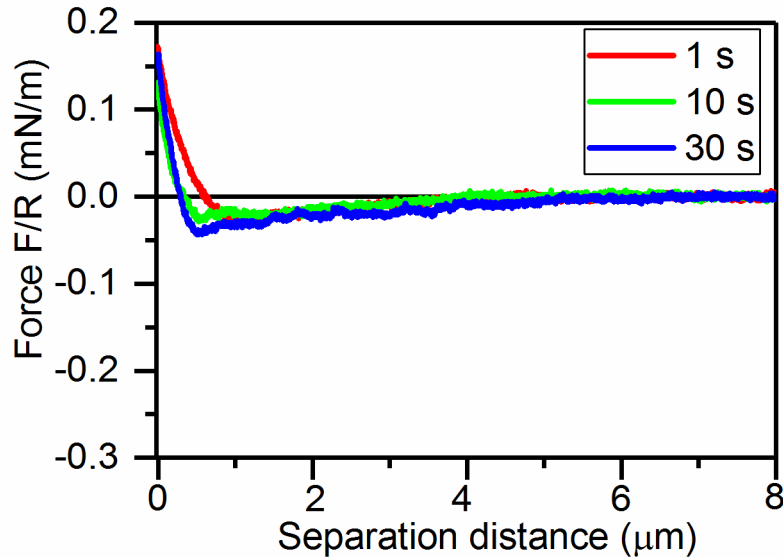


Effect of time in contact

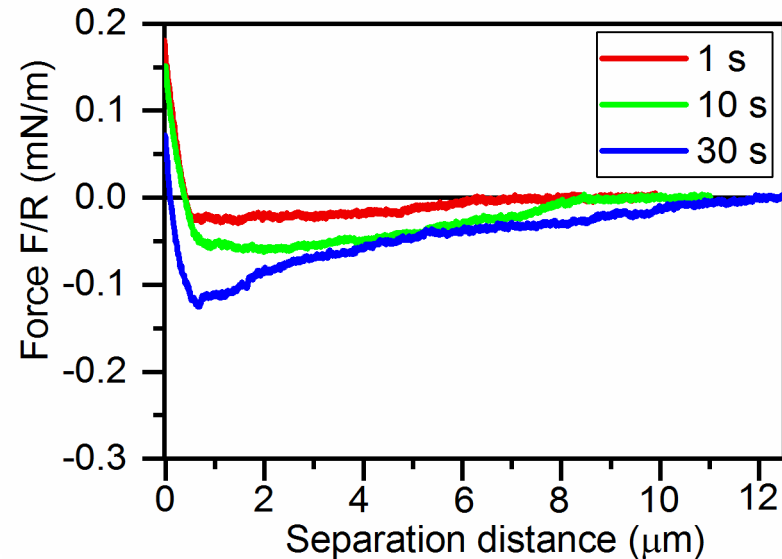


Retraction curves

Living WA07/CNF



Living WA07/Laminin 521



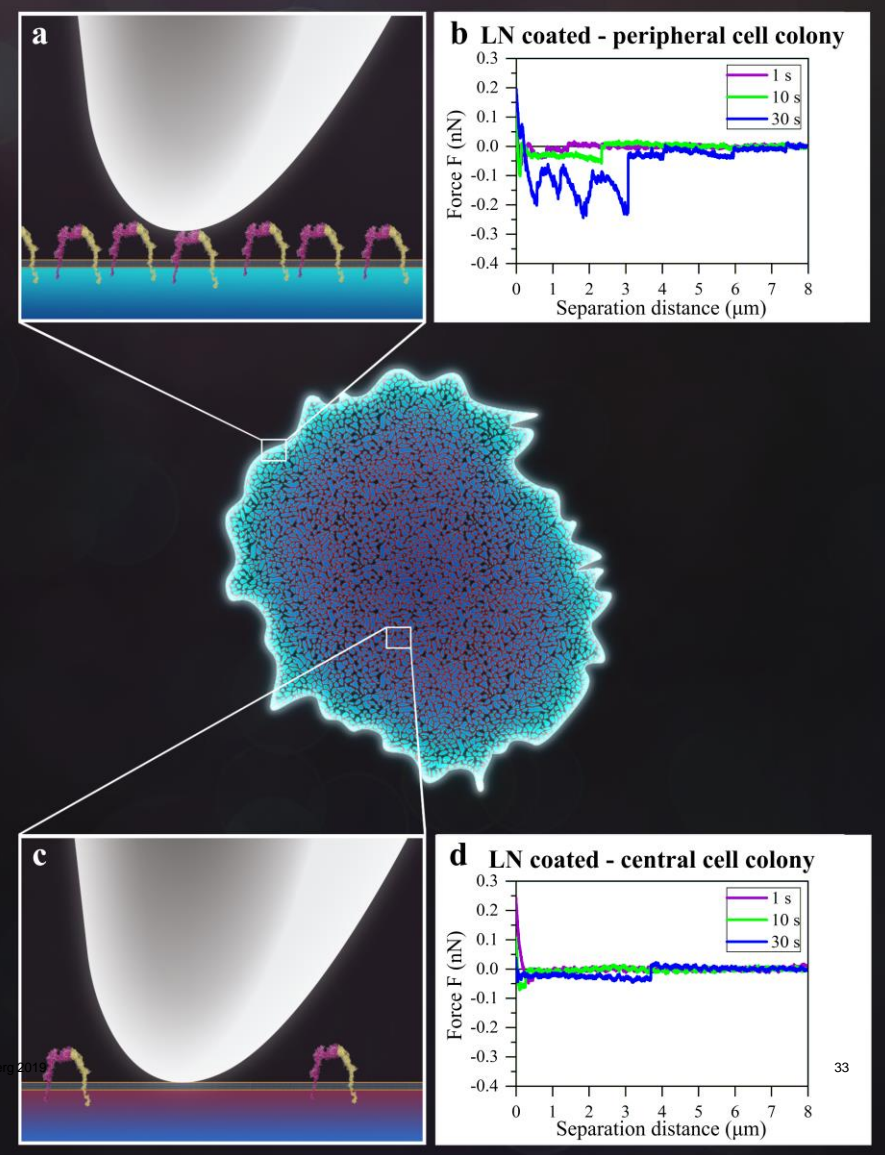
Specific vs nonspecific forces

Living cells on the substrate
(human pluripotent stem cells)

Well defined tip coated with laminin

Strong interaction between
activated integrins at the cell
membrane and the laminin

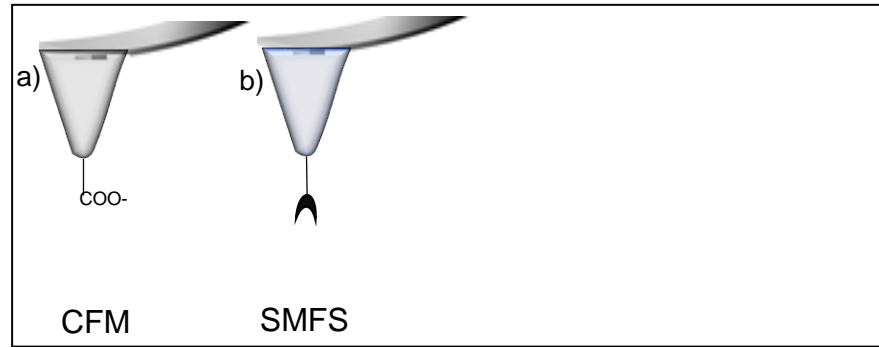
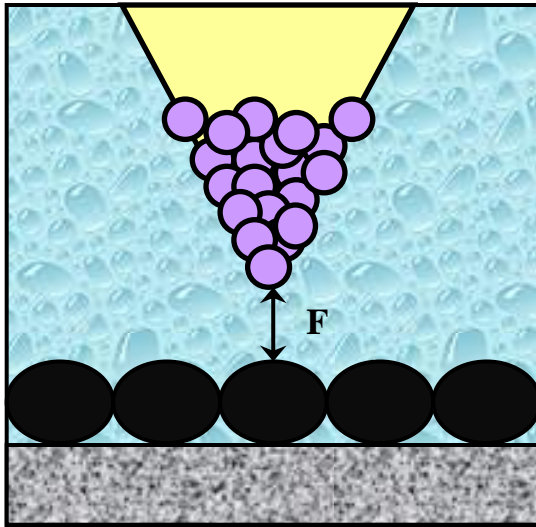
Different force curve profile -> the
integrins are more concentrated at
the periphery of cell colonies



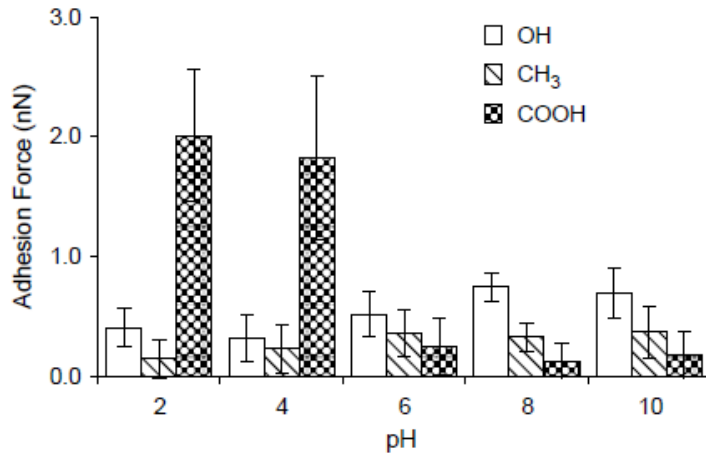
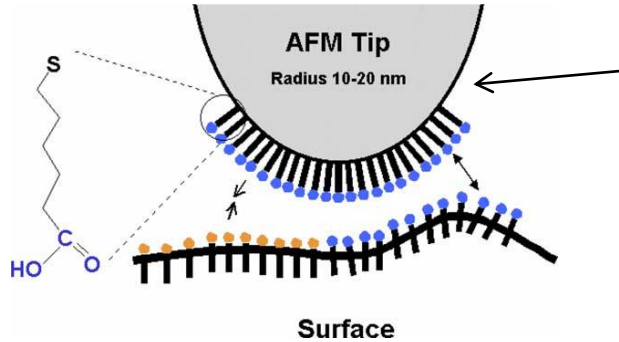
Take home message from previous example

- Colloidal probe method suitable for delicate cells
- Specific and unspecific interactions can be distinguished
- Good correlation between phenomena found during in vitro cell culturing and direct force measurements
- Cell – biomaterial interactions can be controlled with proteins

Chemical Force Microscopy and Single Molecule Force Spectroscopy



Chemical force spectroscopy: Example interactions with cellulose pulp fibre surface

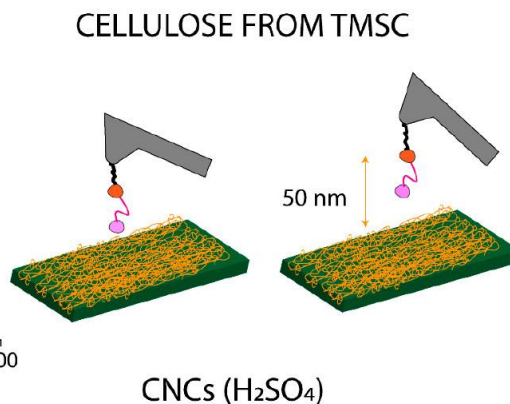
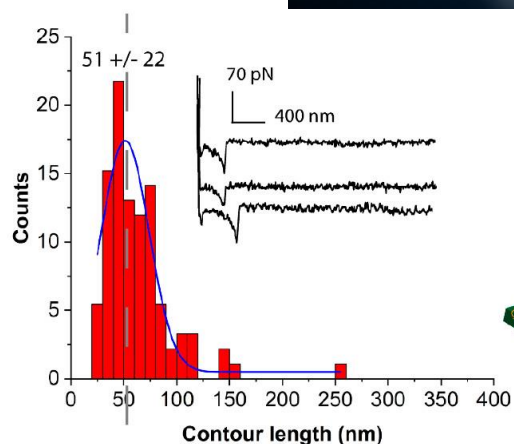
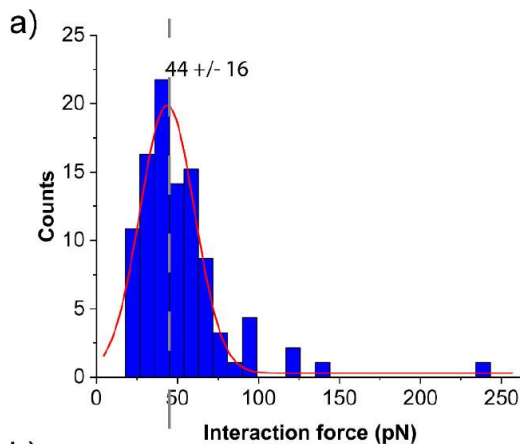
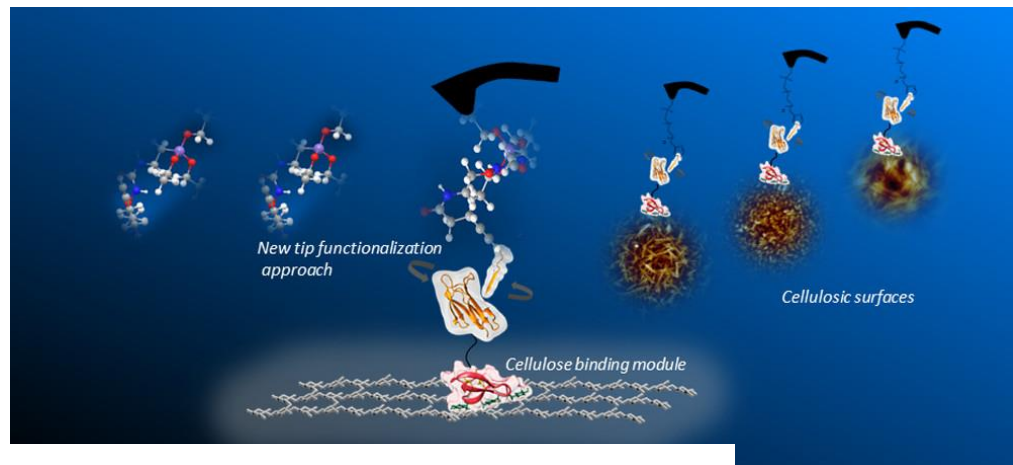


- Highest adhesion between COOH and cellulose
- pH dependence
- Almost no adhesion between CH₃ and cellulose

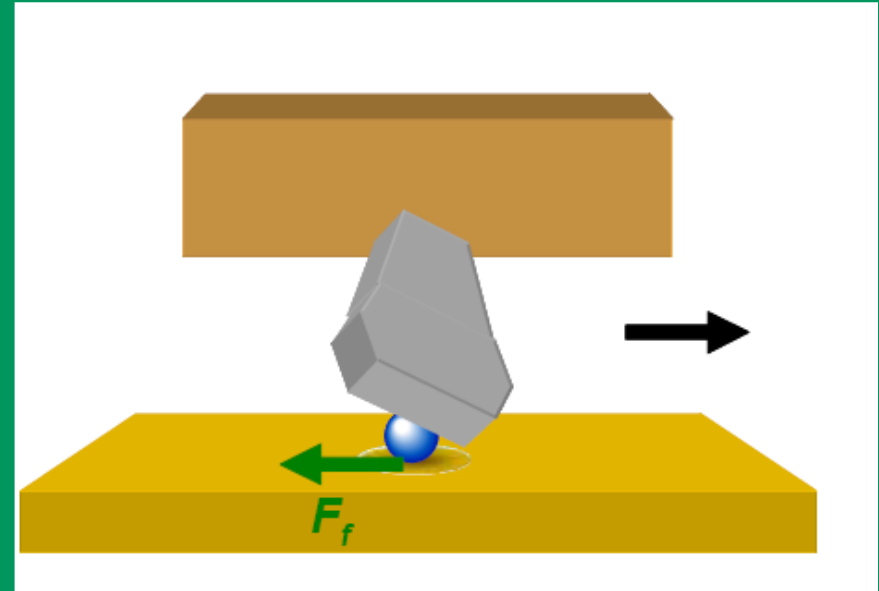
Bastidas et al Carbohydrate
Polymers 62(2005) 369

Binding forces of cellulose binding modules on cellulosic nanomaterials

Effect of cellulosic material on the binding force between CBM and cellulose



Friction force measurements



Friction – what is it?

One should distinguish between two different regimes:

Hydrodynamic (liquid) friction

- the substrates are separated by a thick (> 0.01 mm) liquid film
- friction mainly determined by viscosity of liquid lubricant

Boundary lubrication

- the substrates are separated by a thin (a few atomic diameters) lubricating film
- also dry friction

Friction is the resistance to motion during sliding or rolling of a solid body against another.

The force acting in the direction opposite to the direction of motion is called friction force

Friction



Amontons law: F (friction force) = μL

μ = friction coefficient, L = load

$F_1 = F_2$ i.e. no dependence on contact area!

What about surface roughness??



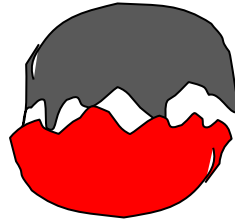
Since friction usually is affected by roughness we need to seek an explanation which involves adhesion.

This requires that surface area is important **BUT** Amontons law tells us that friction depends only on load

Is there a load – area relationship?

The real contact area is usually much smaller than the geometrical area

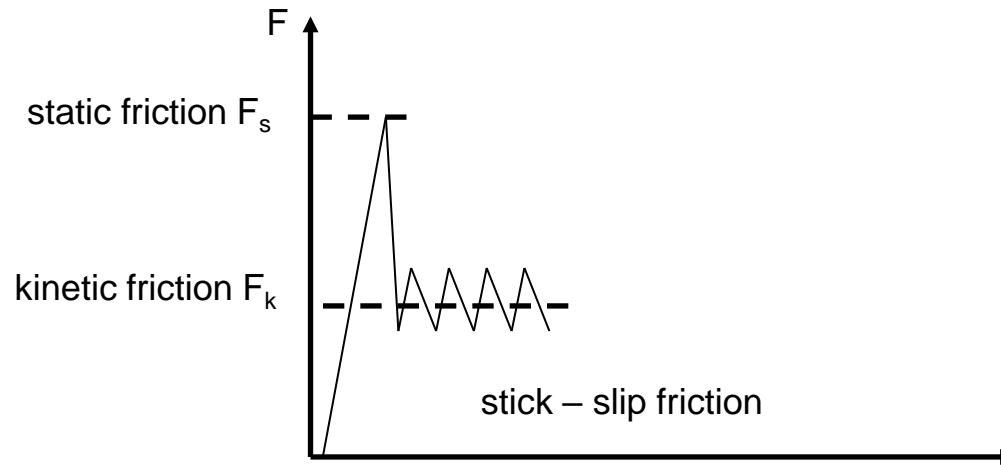
For soft samples the real area is dependent on load => Amontons law



**A fundamental understanding of adhesion and friction requires an understanding of the mechanisms on the atomic/molecular scale
=>Friction force measurements with AFM or SFA**

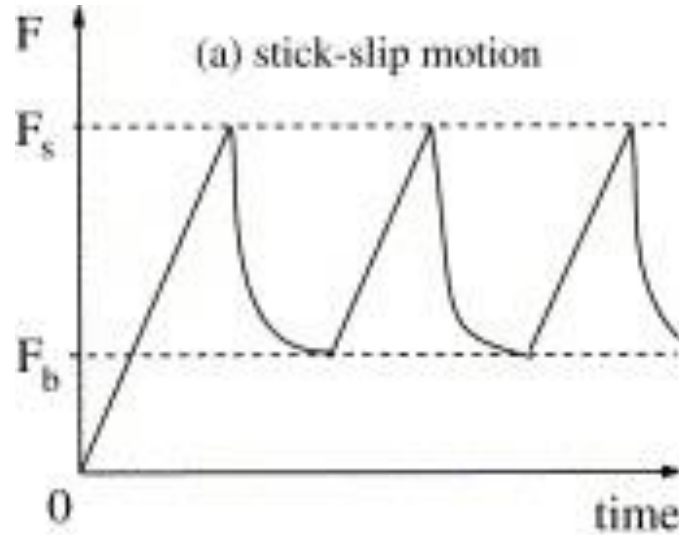
Kinetic versus static friction

friction forces

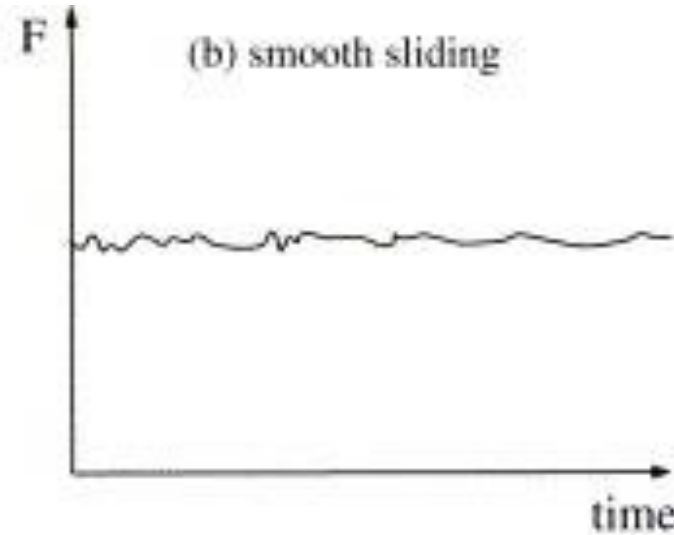


The static friction force is always larger than the kinetic friction force

Stick-slip vs. smooth sliding

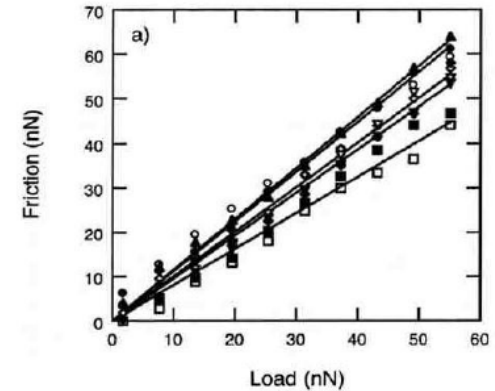
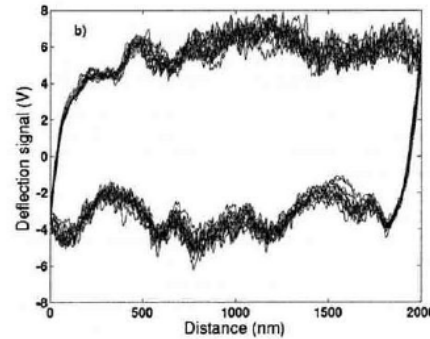
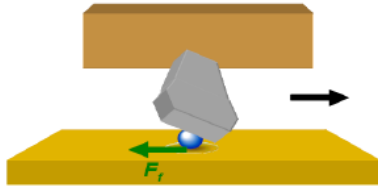


Observed for soft systems and/or low velocities



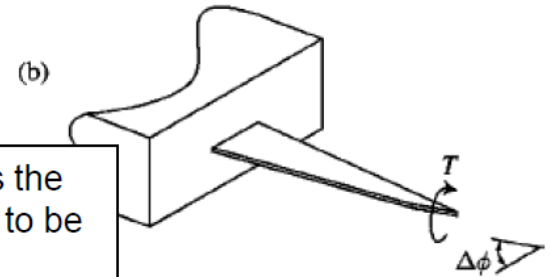
Observed for stiff surfaces and or high velocities

Friction force measurements using AFM



- Friction loops at different loads are measured
- Friction as a function of load
- The slope \rightarrow Friction coefficient (μ)

For friction force measurements the torsional spring constant needs to be determined



Example

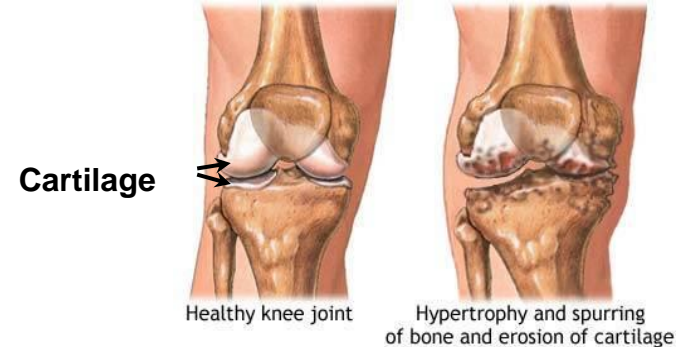
Bioinspired lubricating films of cellulose nanofibrils and hyaluronic acid

Background

Articular cartilage, a flexible tissue that protect the bones in the joints, has a **limited capacity for self-repair**.

Main requirements:

- Able to withstand high applied loads
- High resistance to wear
- Low friction coefficient (lubrication)
- Biocompatible



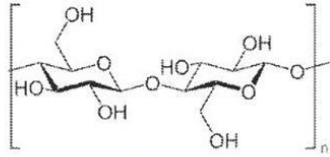
Cartilage damage and cartilage-related diseases:

Europe: about **25 million people** suffer from osteoarthritis; **90 million EUR/year**.

Approach

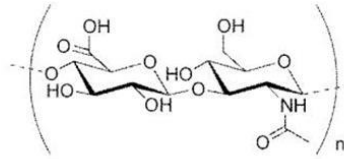
- **Hyaluronic acid (HA) is a natural lubricating polymer present in the joints.**
- **Hyaluronic acid (HA) has low mechanical properties**
- **CNF films have very good mechanical properties but high friction**

**Combining the mechanical strength of
CNF with the lubricating effect of HA**



CNF

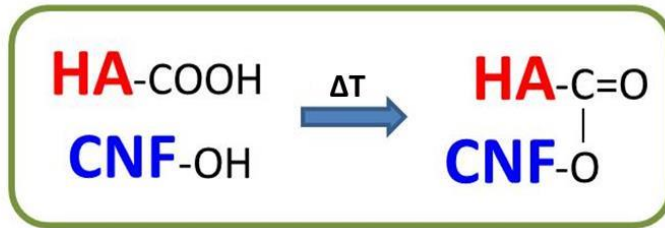
Cellulose nanofibrils



HA

Hyaluronic acid

HA attached to CNF films by esterification reaction between hydroxyl and carboxyl groups



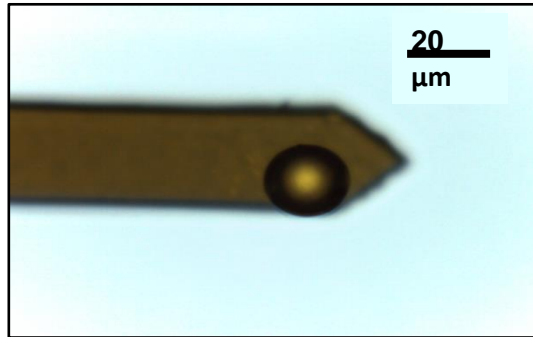
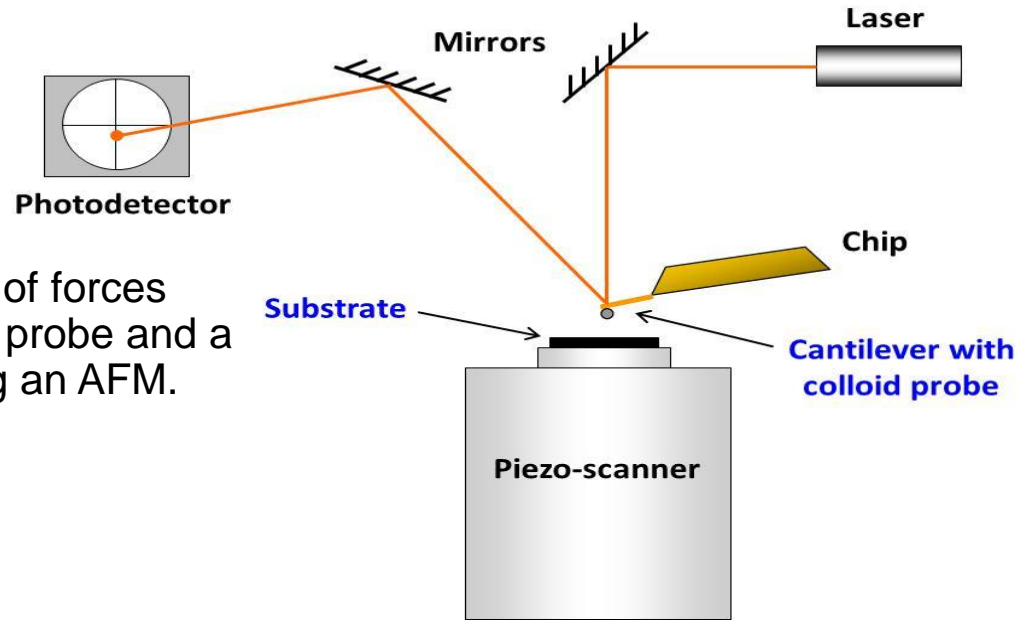
**Objective:
Durable lubricating layer**

Atomic force microscope (AFM) and colloid probe technique

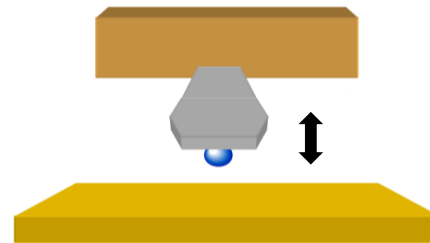


AFM

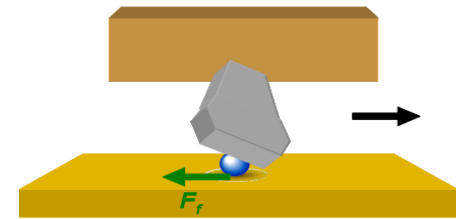
Measurement of forces between a colloid probe and a substrate using an AFM.



Glass colloid probe



Surface force measurement



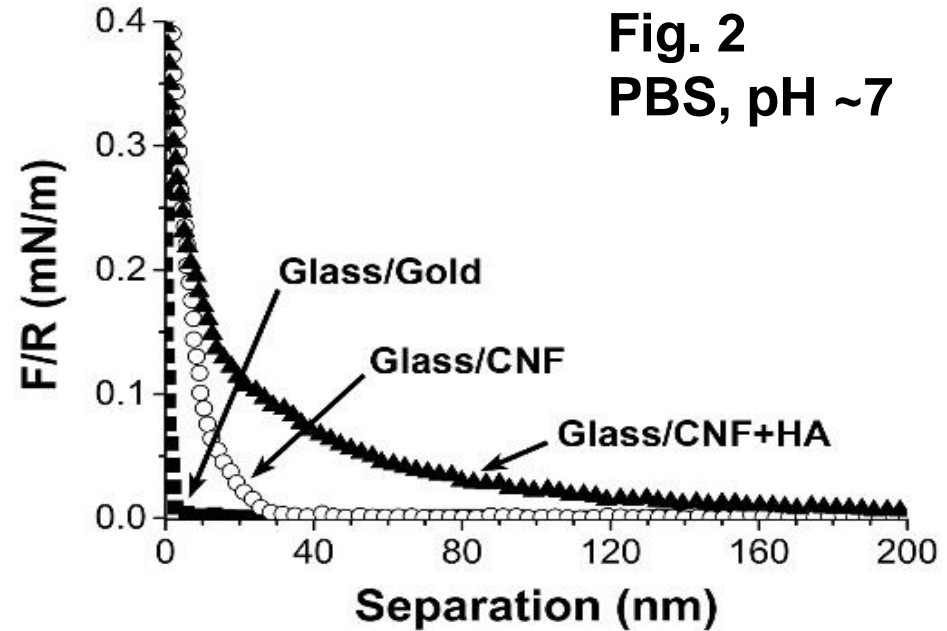
Friction force measurement

Normal forces – What forces are present?

Van der Waals Forces? **Yes**
Electrostatic Double-Layer Forces? **Yes**
Steric forces? **Yes**

Increase in steric repulsion

What is the reason for the steric repulsion?



Effect of ionic strength

Hydrated layer, high repulsion
→ Low friction

Phosphate buffered saline (PBS): 10 mM Na_2HPO_4 , 1.8 mM KH_2PO_4 , 137 mM NaCl, 2.7 mM KCl **High I**

Phosphate buffer (PB): 10 mM Na_2HPO_4 , 1.8 mM

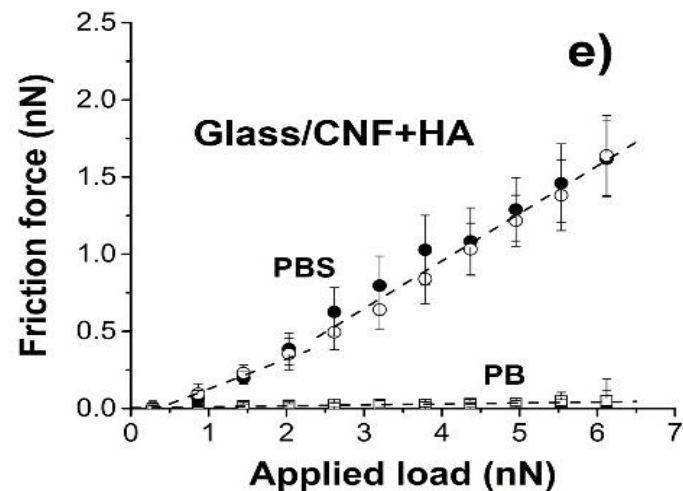
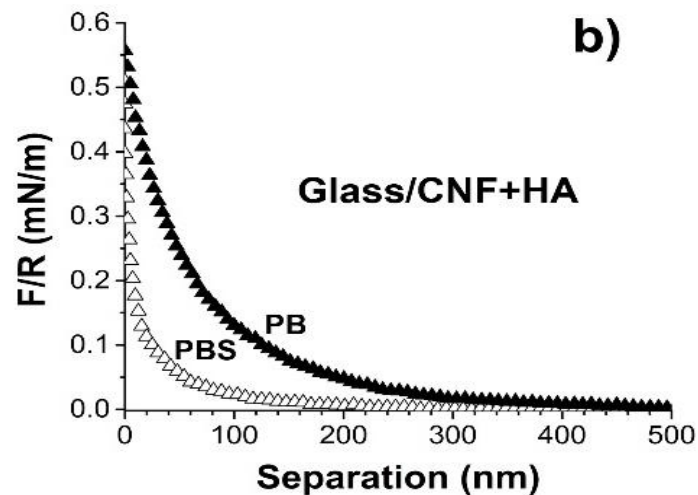
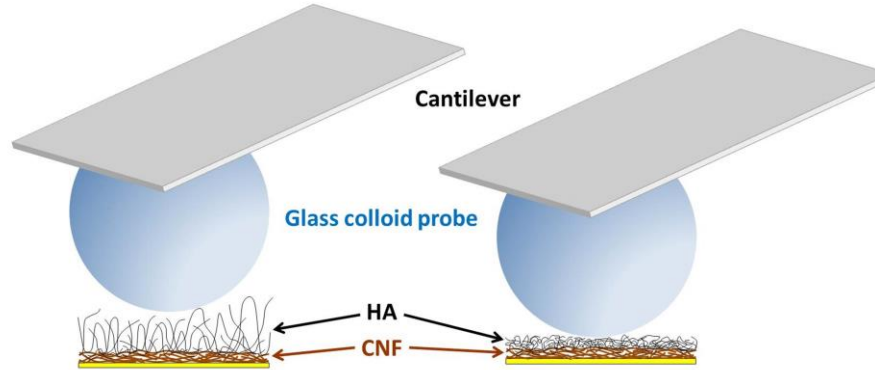


Fig 3.

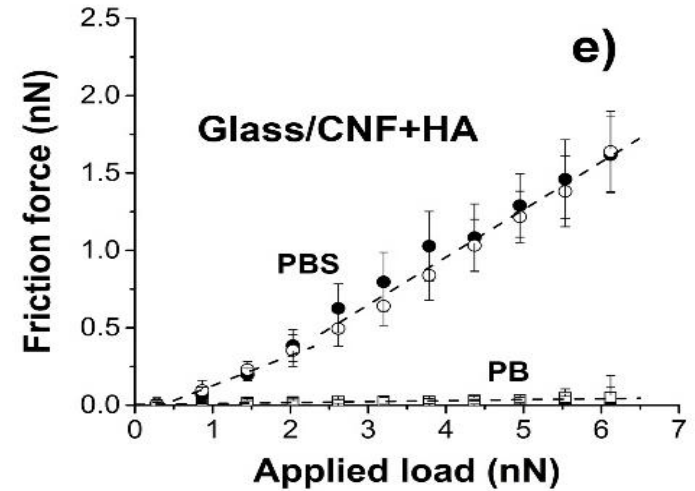
[dx.doi.org/10.1016/j.colsurfb.2015.11.047](https://doi.org/10.1016/j.colsurfb.2015.11.047)⁵¹

Effect of ionic strength II

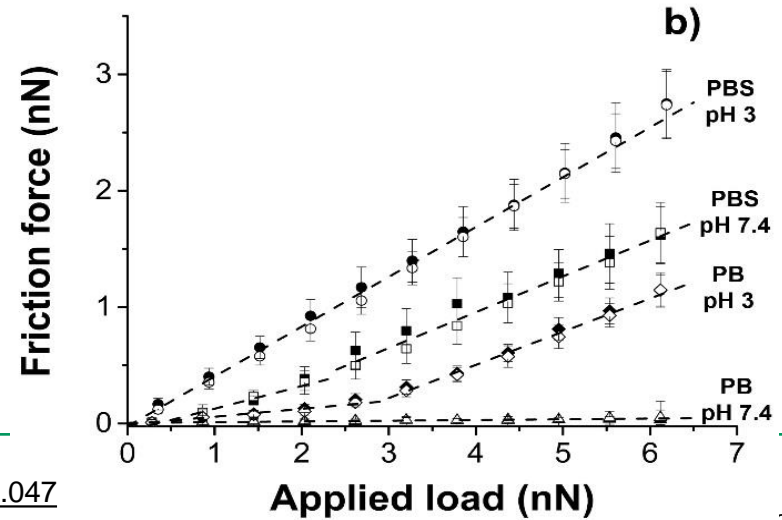
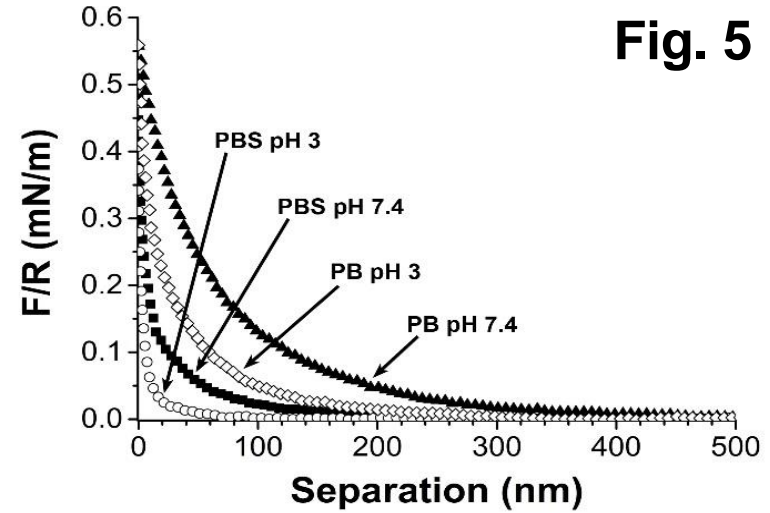


Low I
Hydrated fibril and polymer layers
Very low friction

High I
Collapsed polymer layer
Higher friction



Effect of pH



Reflections on the previous example

Lubrication was achieved by attachment of polymers

Hydrated polymer layer

Extended polymer chains (good solvent)

Similar approach of surface modification can be used for:

- **Steric stabilization of nanoparticles**
- **Antifouling surfaces (grafting of PEG chains)**
- **In composites for better alignment of reinforcing fibers**

Take-home message

- **Interfacial properties are decisive in nanomaterials**
- **You gain understanding and can optimize your systems by doing direct surface force measurements**
- **Consider:** Differences and advantages between the different ways of modifying the tip
 - *Chemical force microscopy (CFM)*
 - *Single-molecule force spectroscopy (SMFS)*
 - *Colloidal probe microscopy (CPM)*
 - *Single cell force spectroscopy (SCFS)*