

Optical and atomic force microscopy: from single molecules to living cells

Frédéric EGHIAIAN, Post-doctoral researcher
(INSERM U1006, Director Simon Scheuring)

My career...

Master/Ph.D (2001-2005, Paris)

Structural and biochemical studies of the sheep prion protein

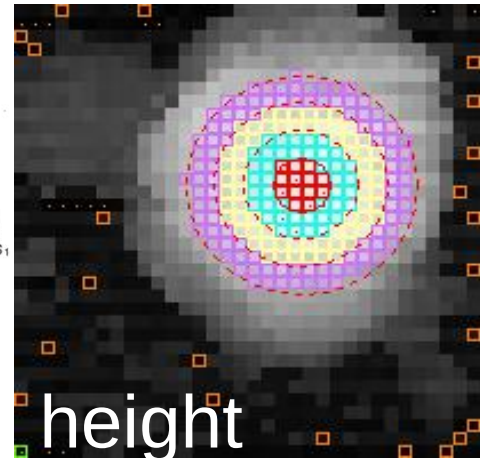
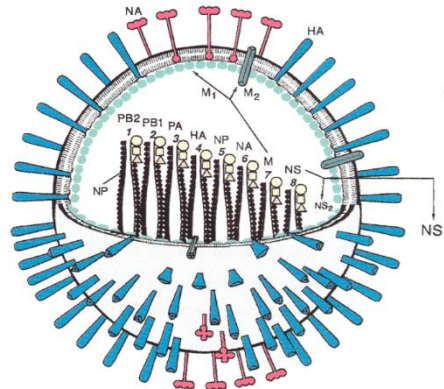


Eghiaian et al, PNAS 2004

Eghiaian et al, PNAS 2007

Post-doc (2005-2009, London, 2009-2012 Göttingen):

Single molecule studies of Influenza and ribosomes



Li et al, BJ 2011

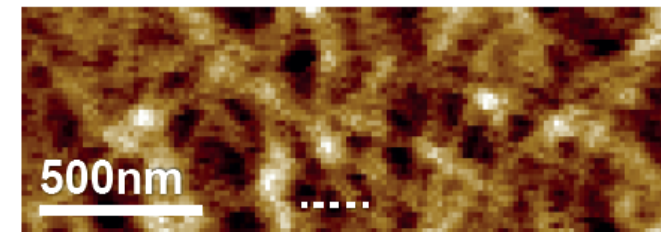
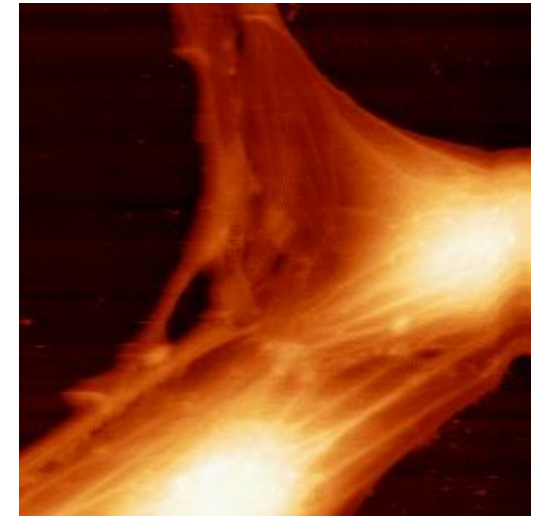
Eghiaian et al. JBC, 2012

Li et al., BJ 2014

Eghiaian et al, in preparation

Post-doc (INSERM/AMU U1006, Marseille, 2013-2016):

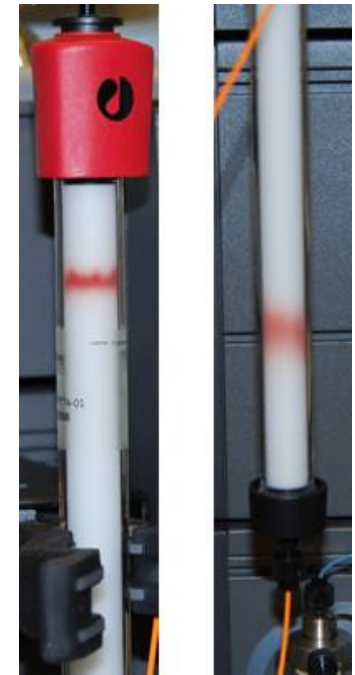
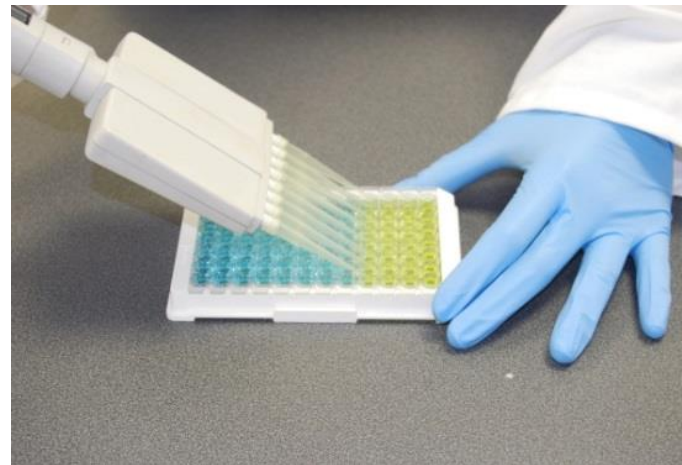
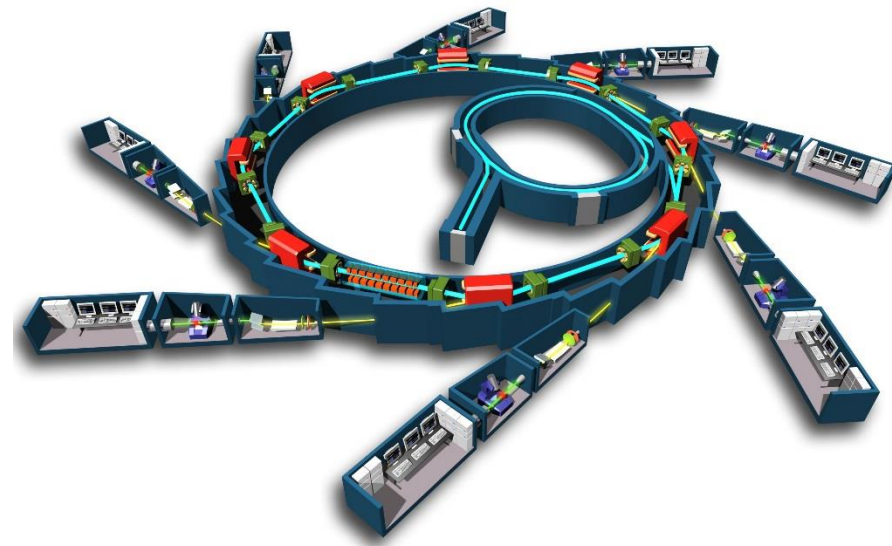
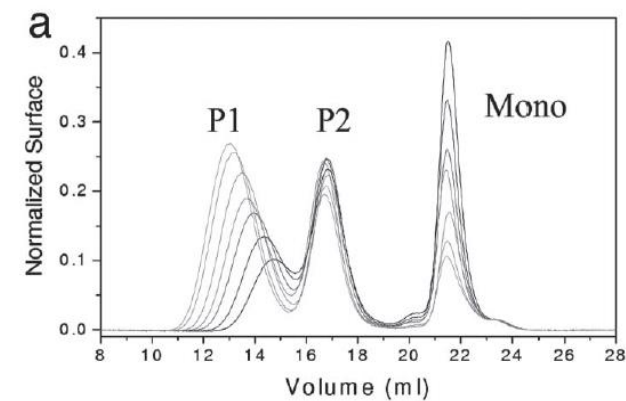
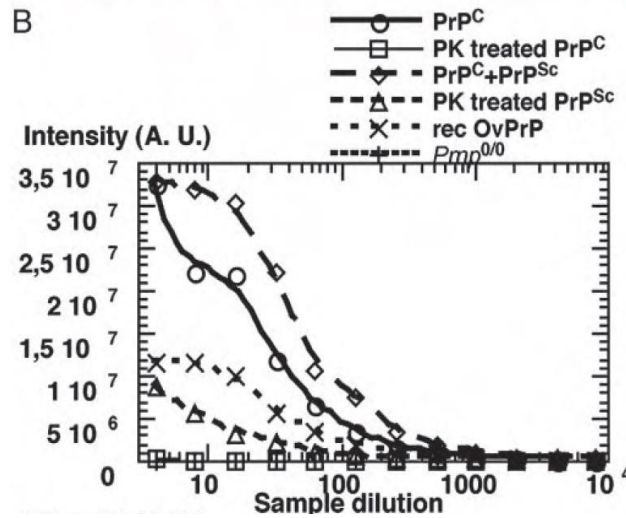
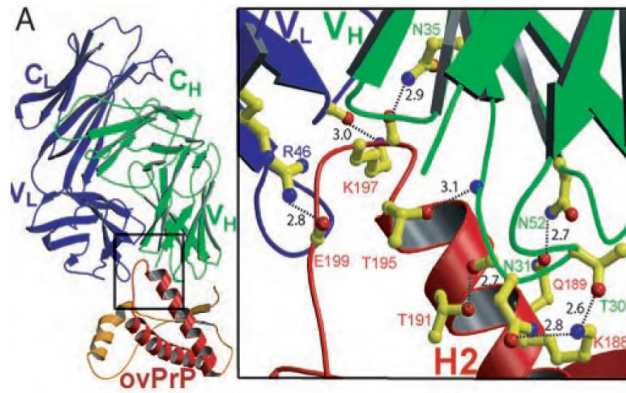
AFM of actin dynamics and endocytosis in living cells



Eghiaian et al, Biophys J 2015

Rigato et al, ACS Nano 2015

Eghiaian et al, in preparation



Eghiaian et al, PNAS 2004

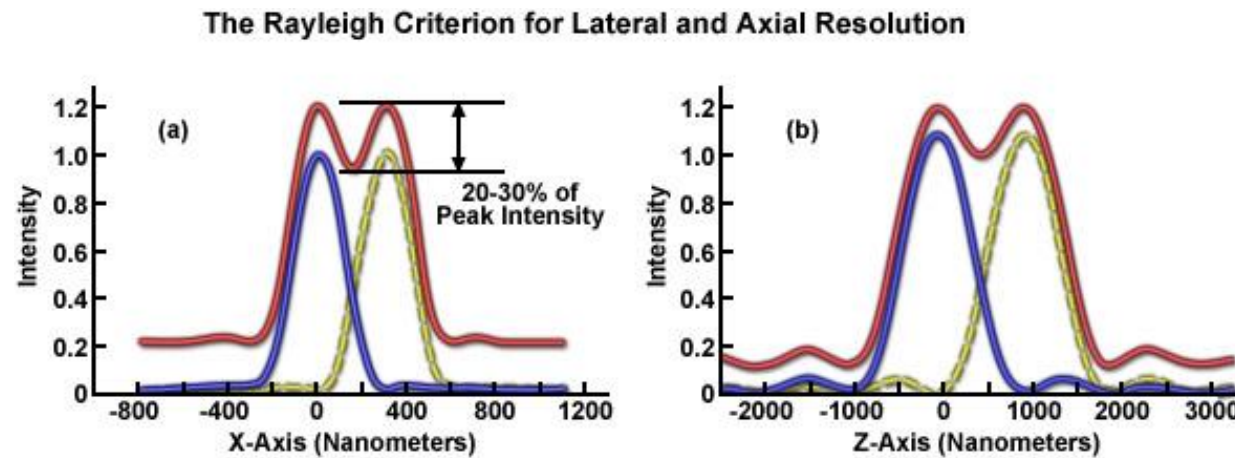
Eghiaian et al, PNAS 2007

How can one see dynamics at the single molecule level ?

Single molecule
fluorescence ???

Challenges of single molecule fluorescence

- Contrast: A single fluorophore is not very bright and bleaches fast...
- Resolution: Lateral and Axial resolutions in microscopy are limited !



$$\text{Abbe Resolution}_{x,y} = \lambda / 2NA$$

$$\text{Abbe Resolution}_z = 2\lambda / NA^2$$

$$\text{Rayleigh Resolution}_{x,y} = 0.61\lambda / NA$$

$$\text{Sparrow Resolution}_{x,y} = 0.47\lambda / NA$$

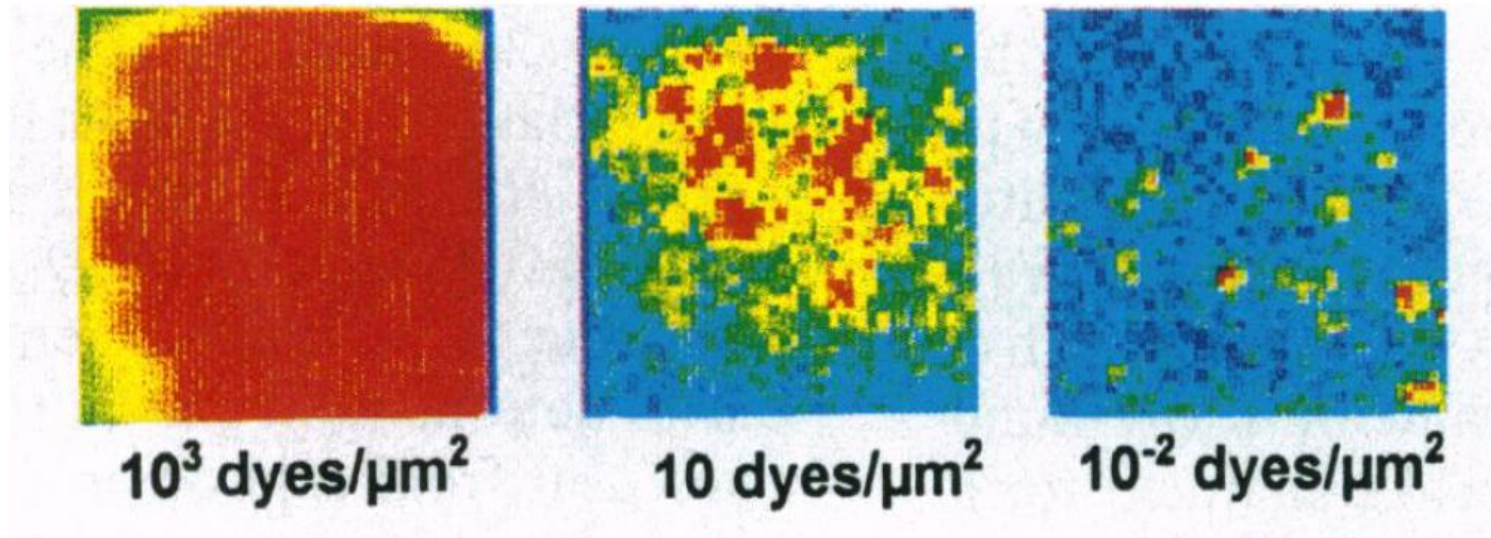
Detecting single molecules by fluorescence microscopy

Diffusion of single fluorescent lipids in planar (glass-supported) lipid bilayers

Proc. Natl. Acad. Sci. USA
Vol. 93, pp. 2926–2929, April 1996
Biophysics

Imaging of single molecule diffusion

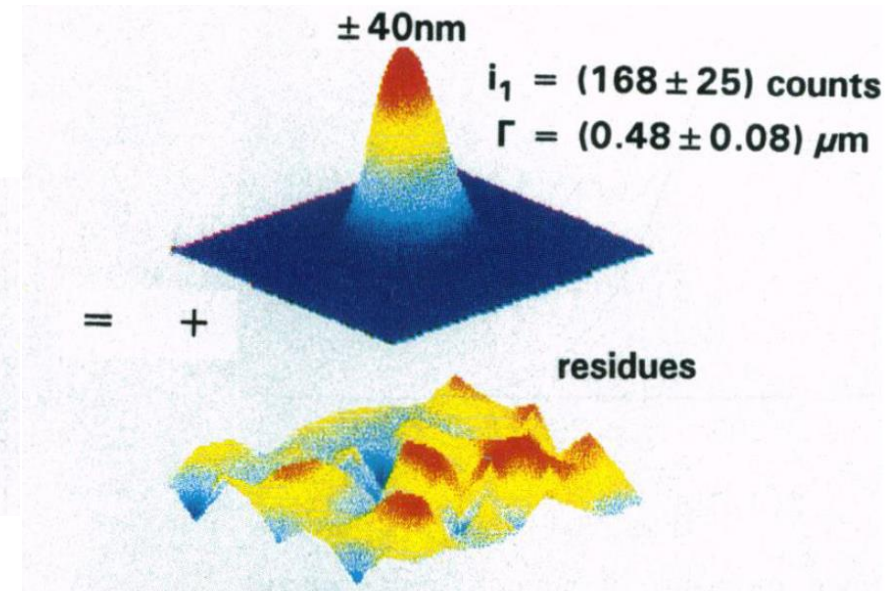
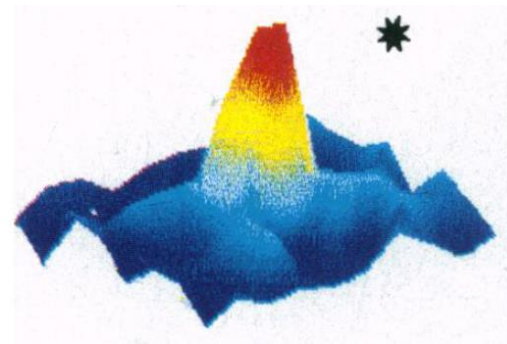
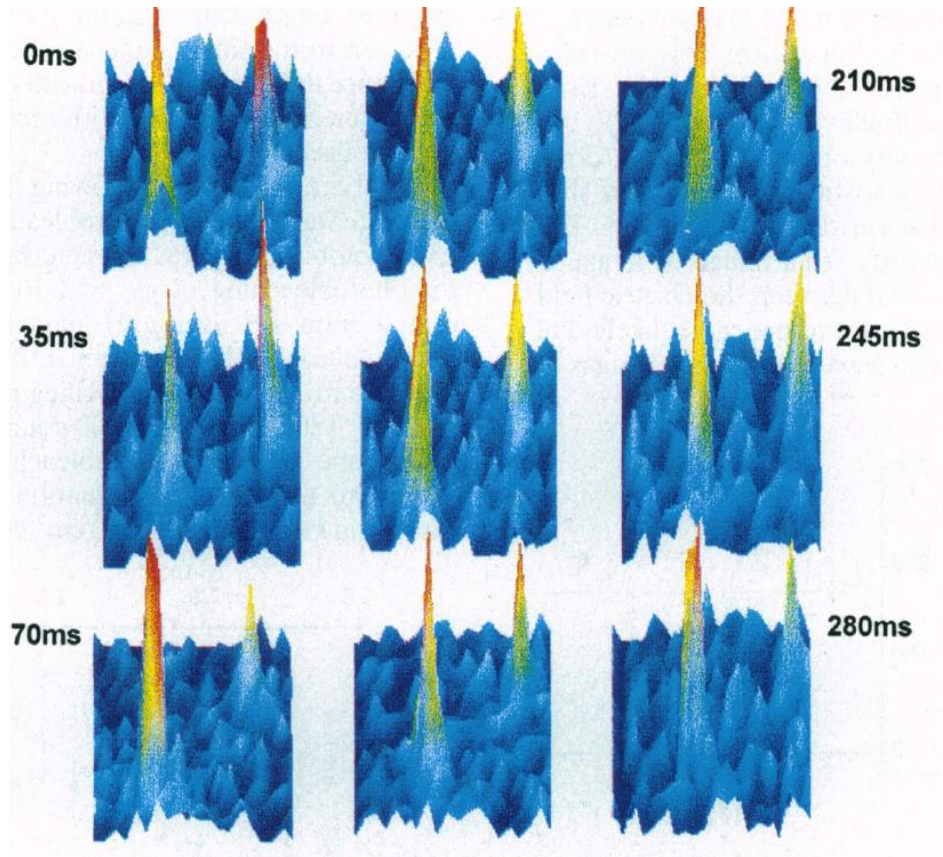
TH. SCHMIDT, G. J. SCHÜTZ, W. BAUMGARTNER, H. J. GRUBER, AND H. SCHINDLER



Reduce molecular density of fluorophores: single fluorophores resolved

High illumination power $\rightarrow$ High Signal-to-noise ratio

Detecting single molecules by fluorescence microscopy



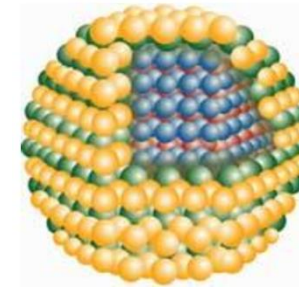
How to increase signal-to-noise ?

- Decrease background fluorescence
- Increase brightness of fluorophores
 - Inorganic nanoclusters
 - Quantum dots
 - Novel small organic fluorescent dyes

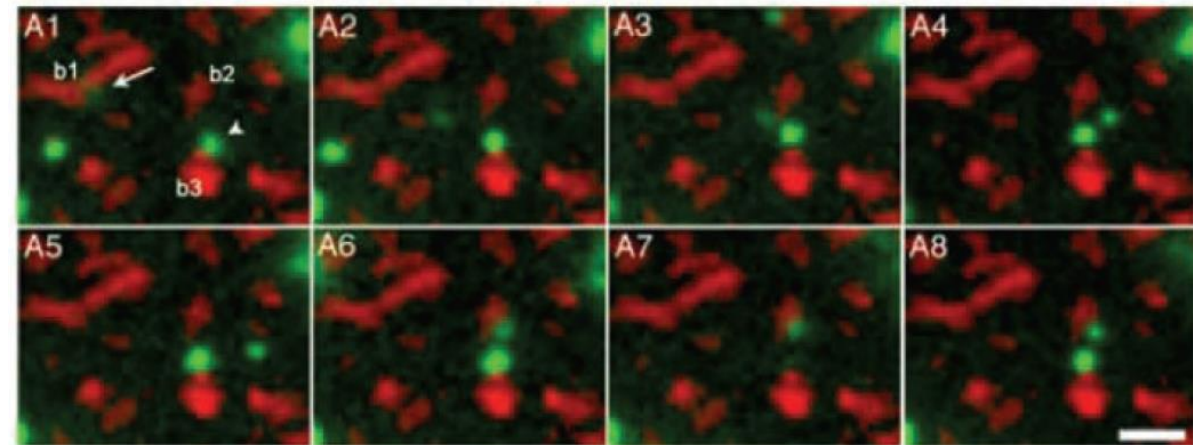
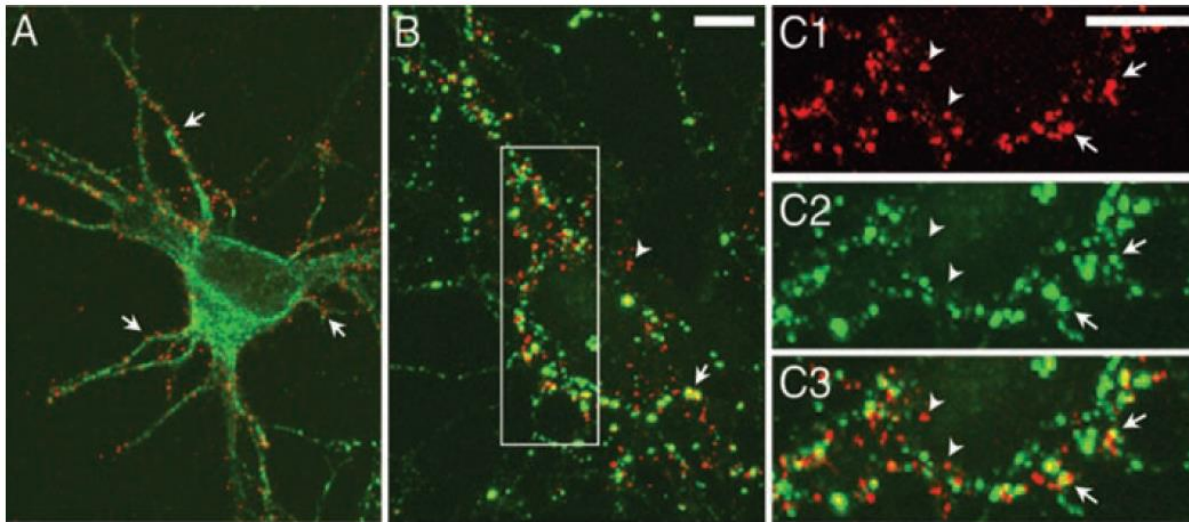
(Not-so-) New dyes: Quantum dots, inorganic nanoclusters...

Diffusion Dynamics of Glycine Receptors Revealed by Single-Quantum Dot Tracking

Maxime Dahan *et al.*
Science **302**, 442 (2003);
DOI: 10.1126/science.1088525



10-100nm !!!

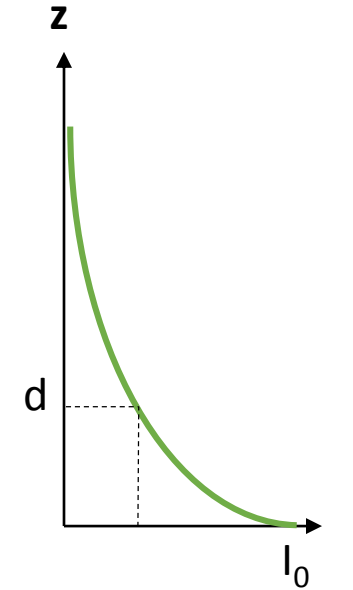
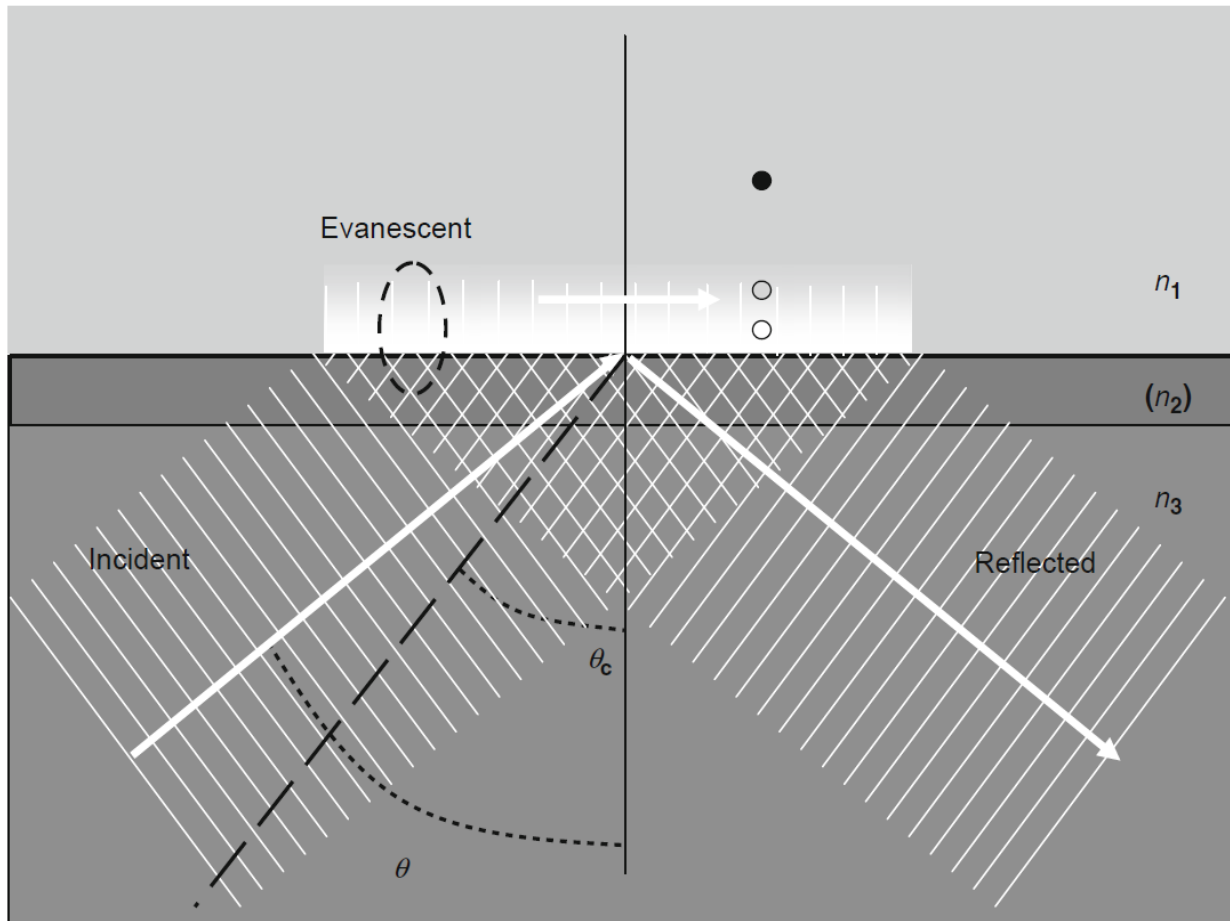


BLINKING !!!

How to increase signal-to-noise ?

- Decrease background fluorescence
- Increase brightness of fluorophores
 - Inorganic nanoclusters
 - Quantum dots

Total Internal Reflection: evanescent wave



$$\theta_c = \sin^{-1} \left(\frac{n_1}{n_3} \right)$$

$$I(z) = I(0)e^{-z/d}$$

$$d = \frac{\lambda_0}{4\pi} (n_3^2 \sin^2 \theta - n_1^2)^{-1/2} = \frac{\lambda_0}{4\pi n_3} (\sin^2 \theta - \sin^2 \theta_c)^{-1/2}$$

Advantages of TIRF-M

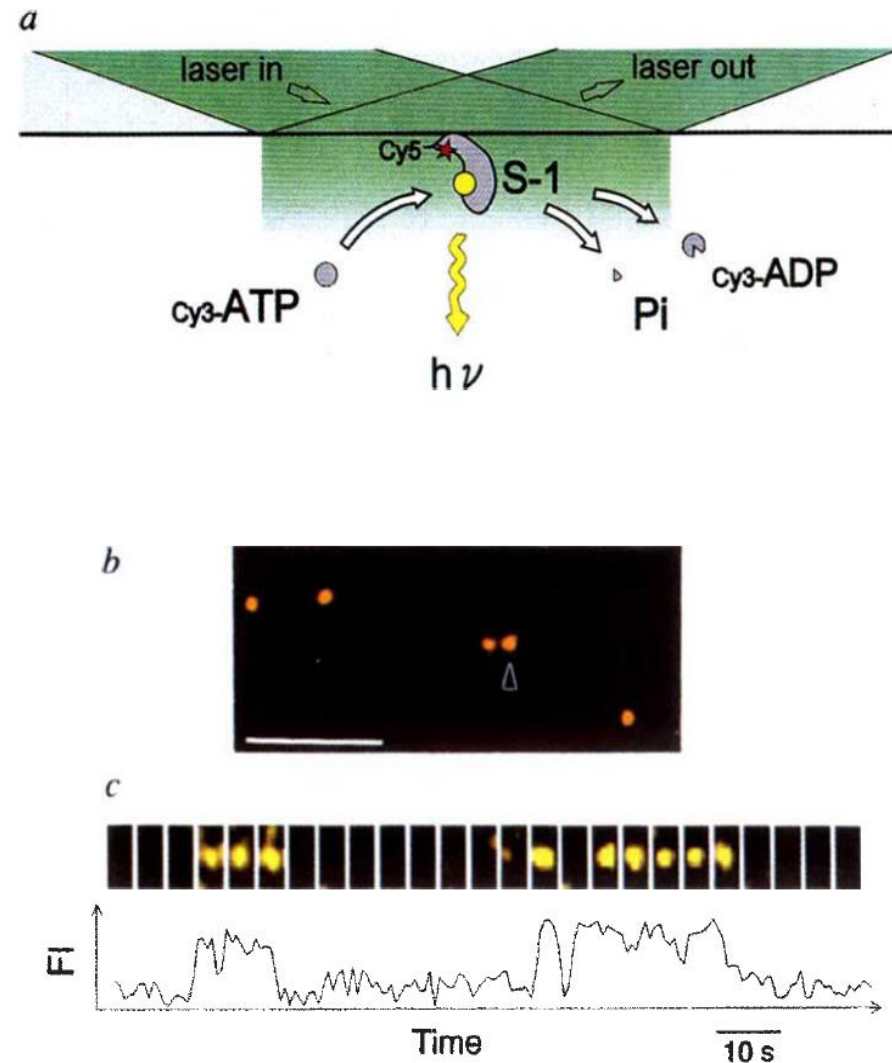
- Restricted excitation volume (close to surface): better signal to noise
- Gain in excitation light intensity
- No loss/gain in resolution

TIRF-M: single molecule imaging

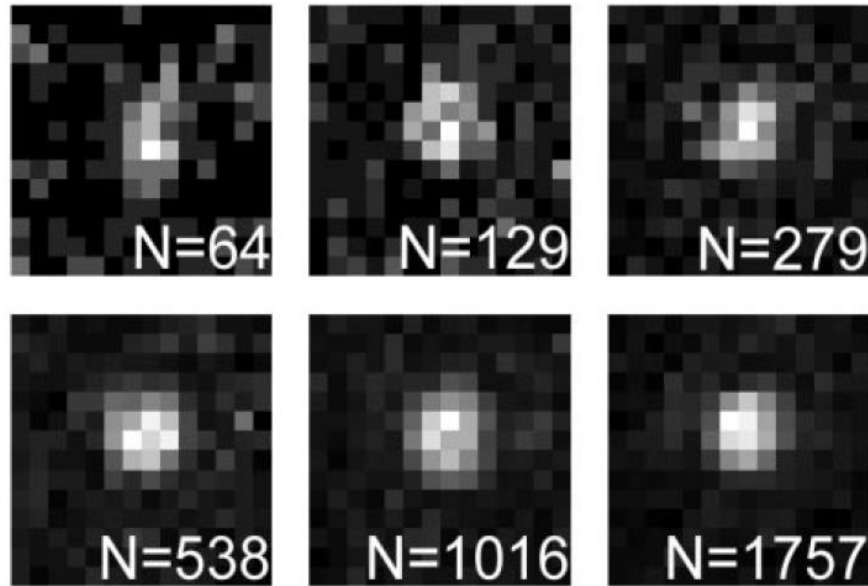
Imaging of single fluorescent molecules and individual ATP turnovers by single myosin molecules in aqueous solution

Takashi Funatsu*, Yoshie Harada*,
Makio Tokunaga*, Kiwamu Salto*
& Toshio Yanagida*[†]

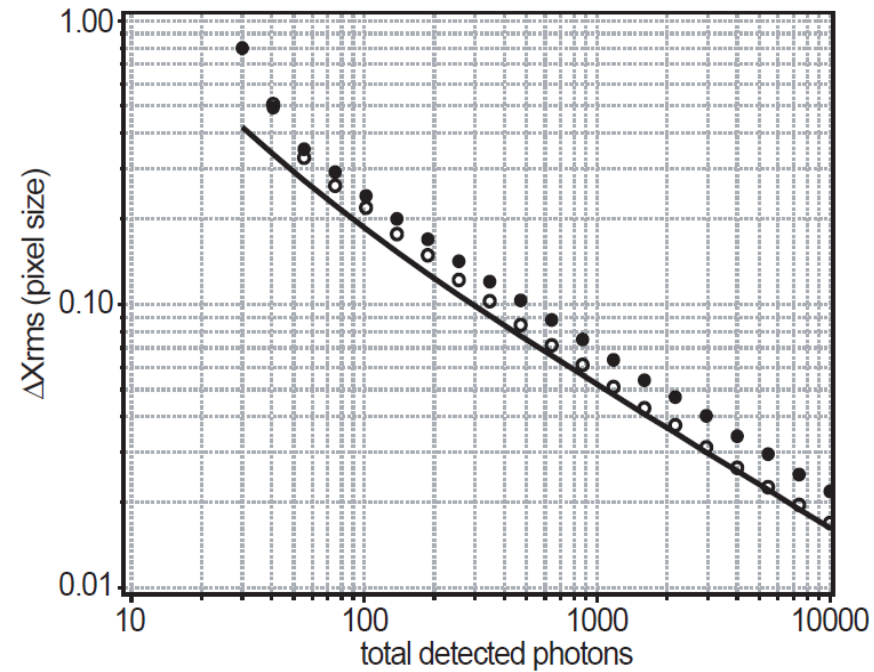
NATURE · VOL 374 · 6 APRIL 1995



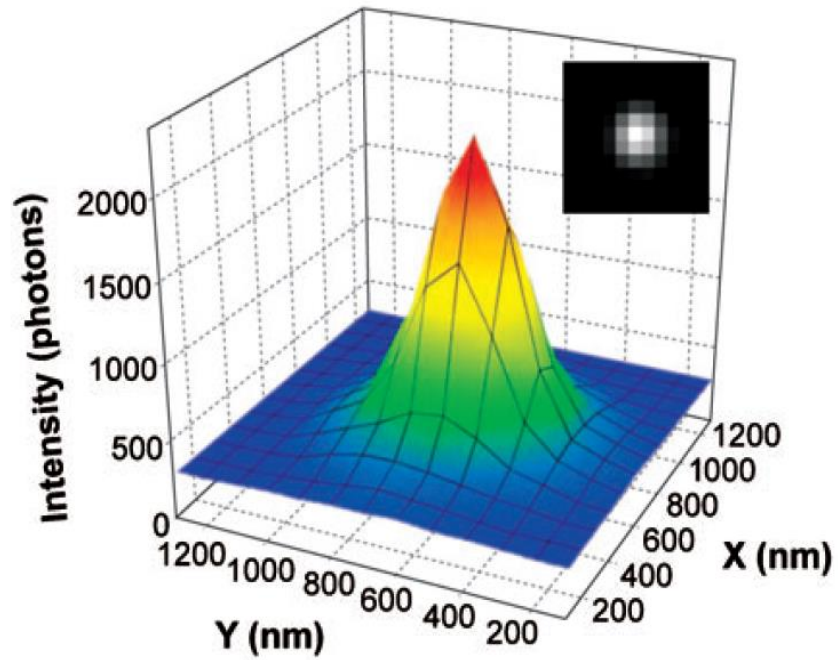
Increased positional accuracy



$$\langle \Delta x^2 \rangle = s^2 / N$$

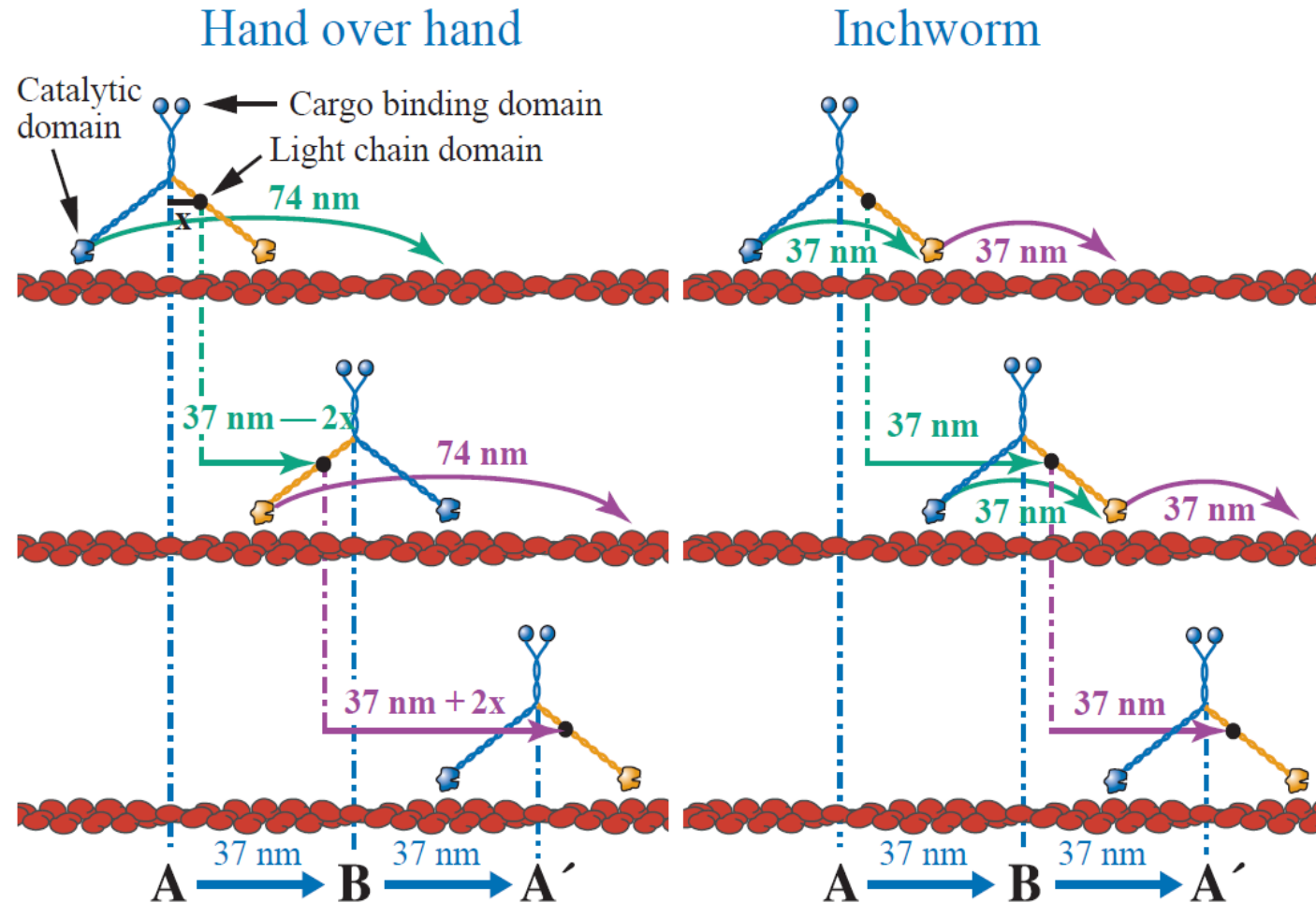


Increased positional accuracy

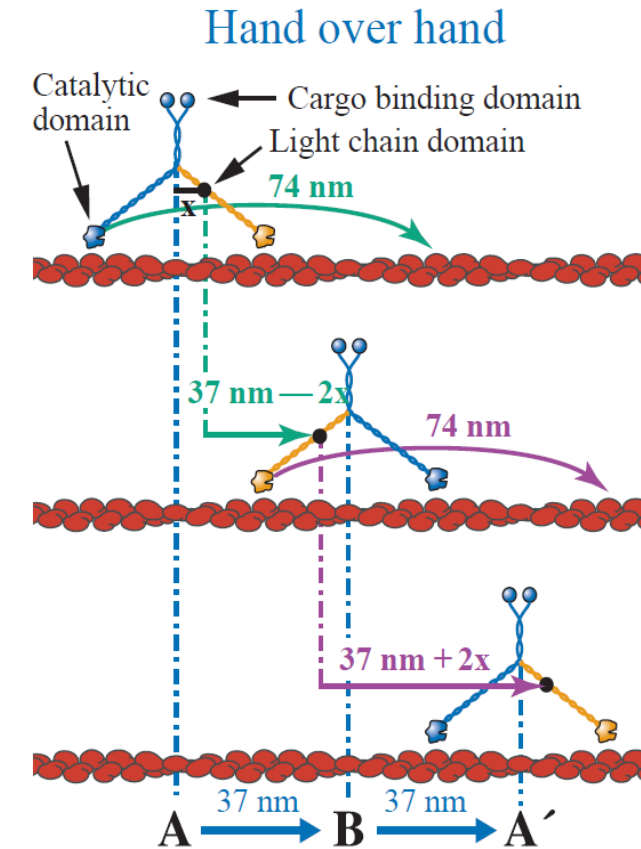
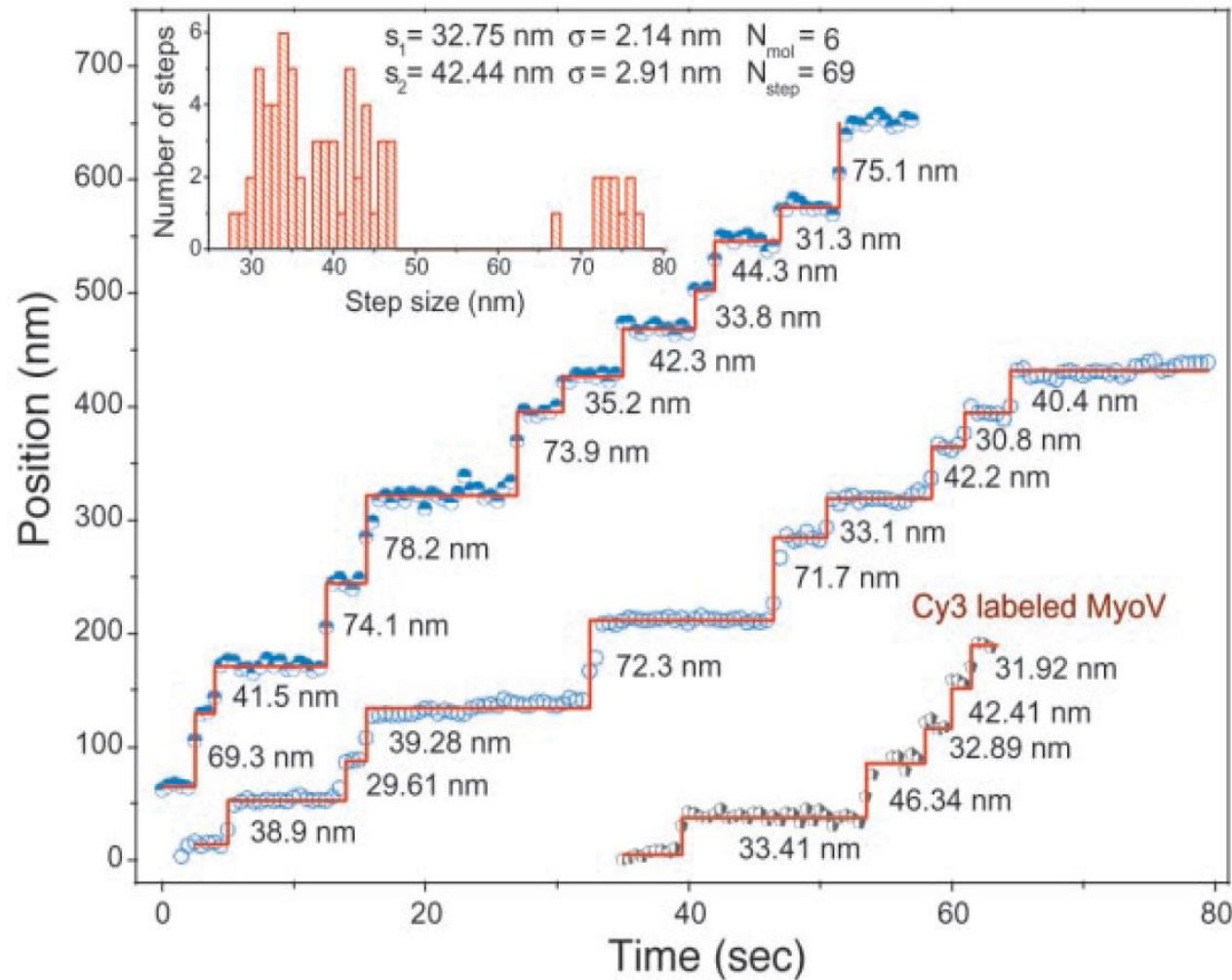


- 2D Gauss fit of point-spread function (spot)
- Standard error of the mean (spot center)=positional accuracy
- 1nm positional accuracy achieved

What is the step-size of the Myosin V motor ?

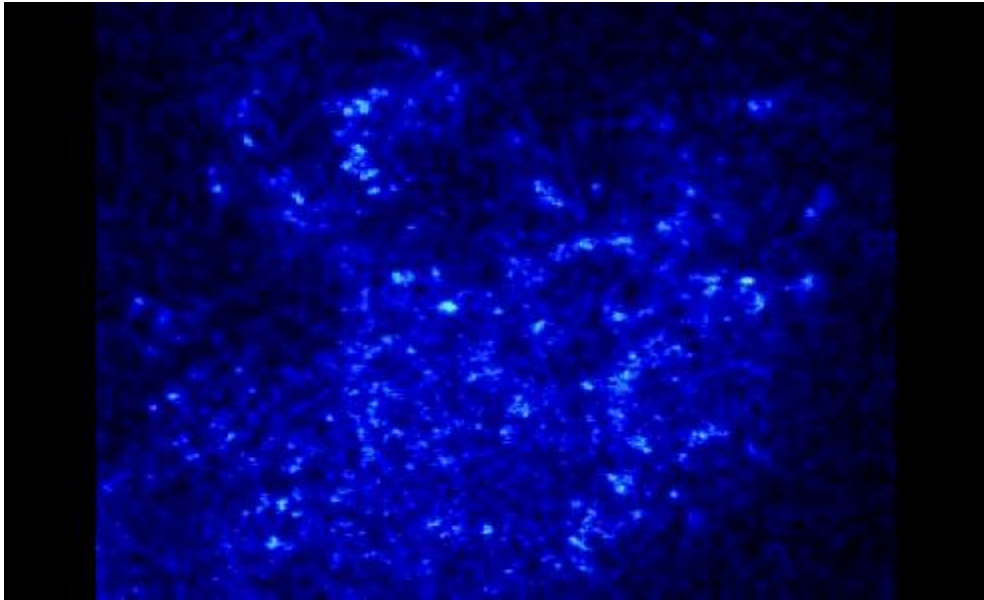


FIONA confirms hand over hand mechanism

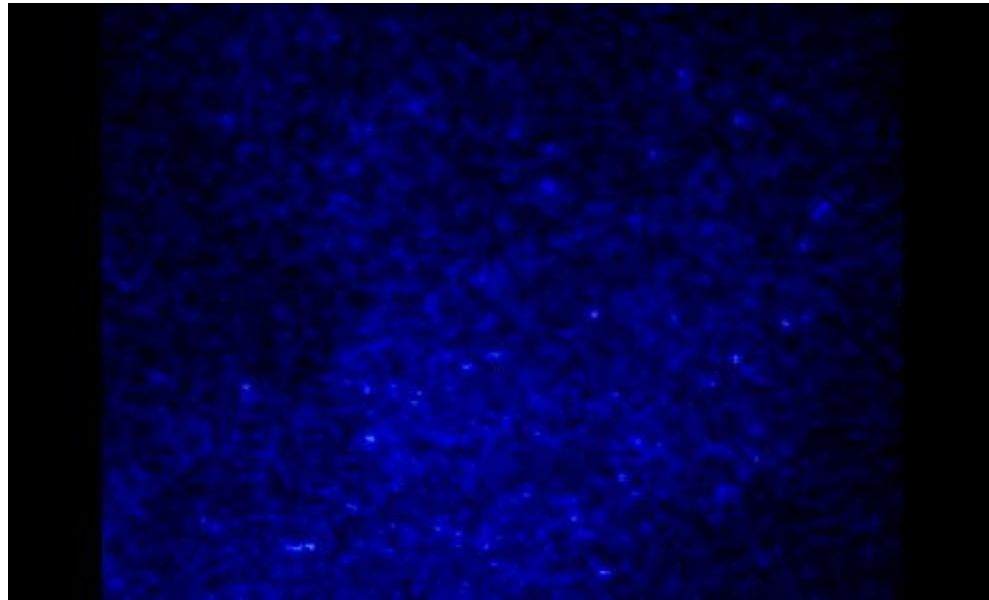


TIRF-M in living cells: Photobleaching

HA-GFP in live HUVECs (low density,
1000 molecules in the image)



After a few seconds....



PALM: Photo-Activation Localization Microscopy

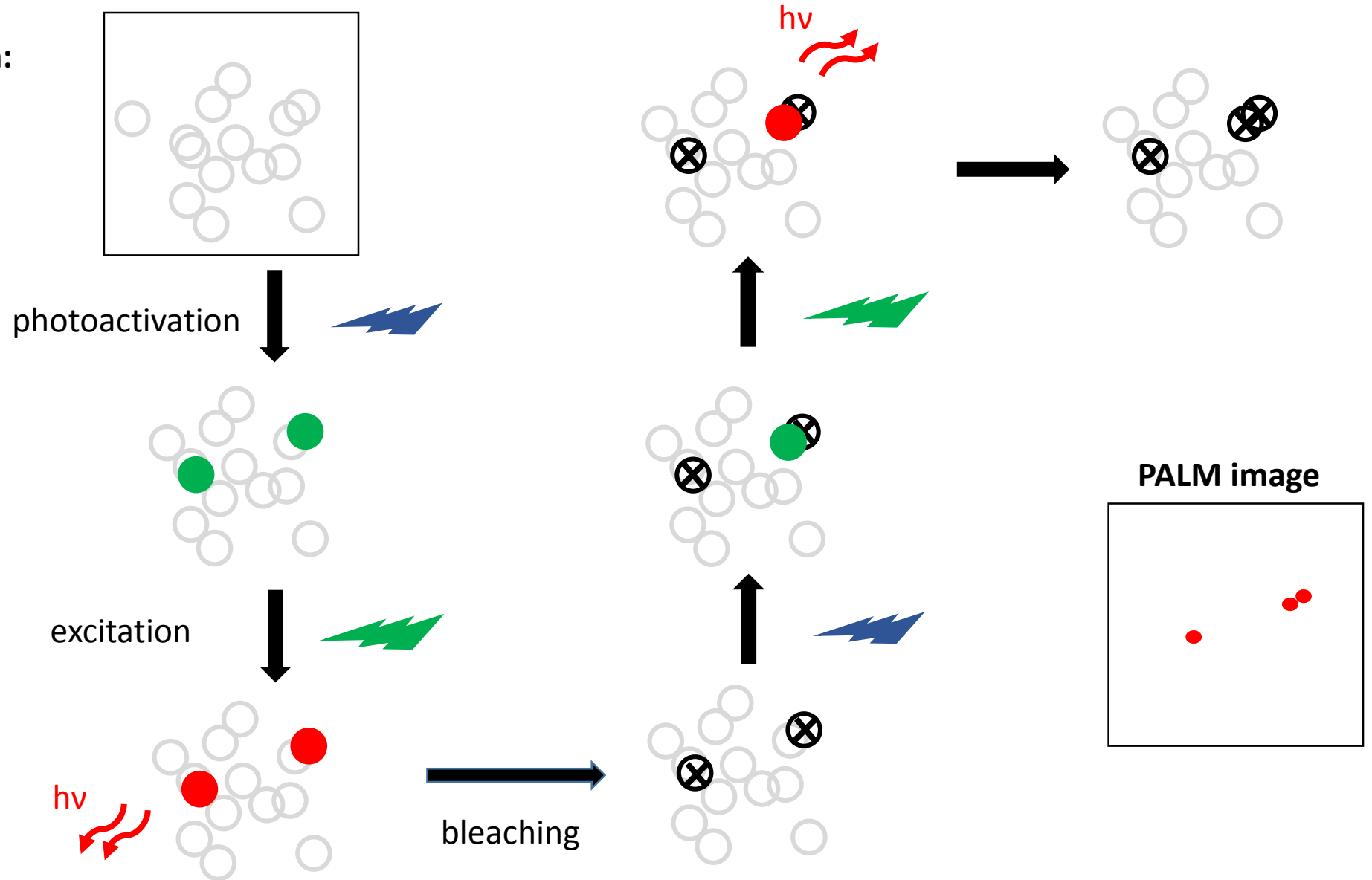
Photo-activable protein:

Eos (=GFP)

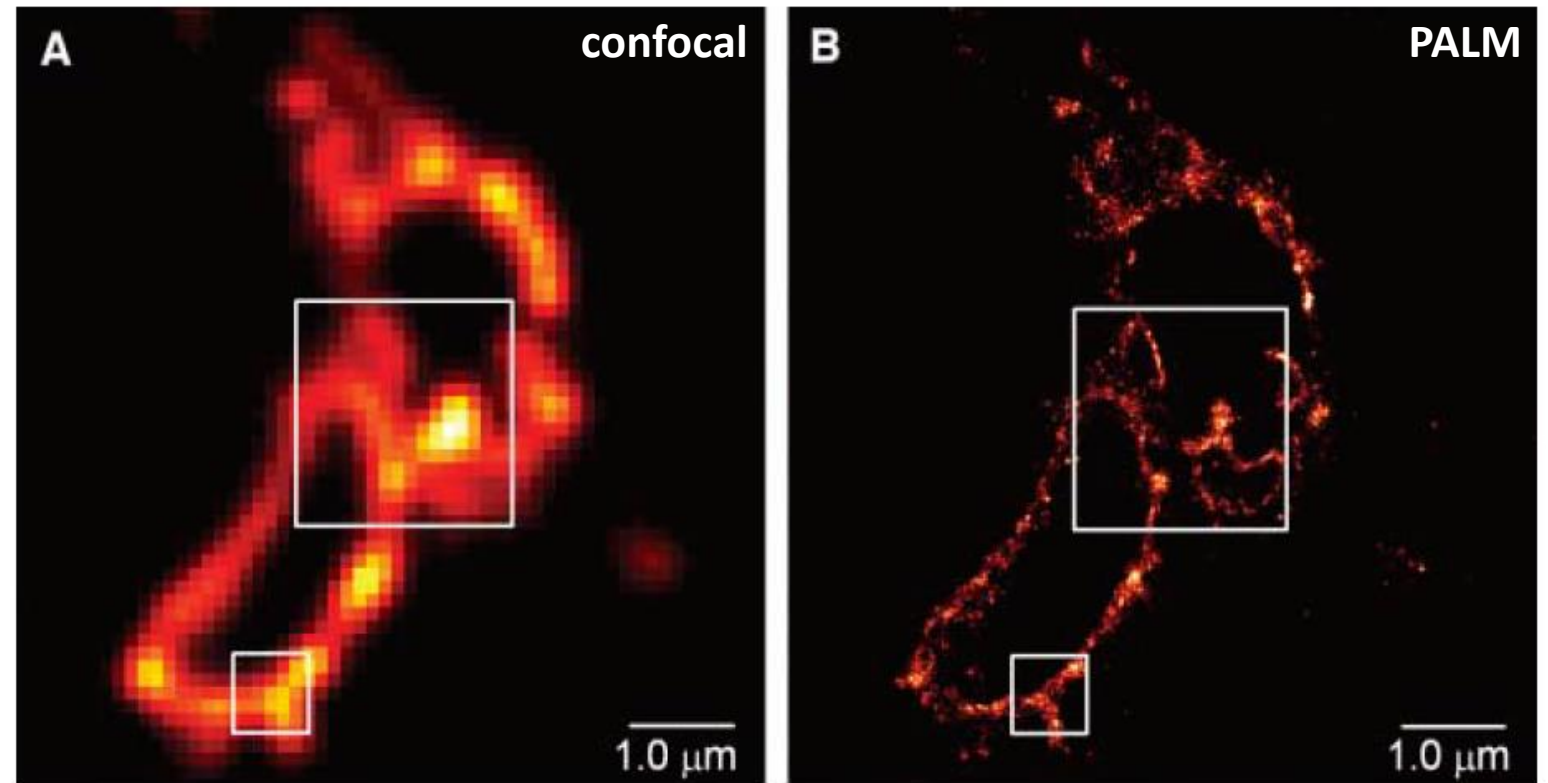
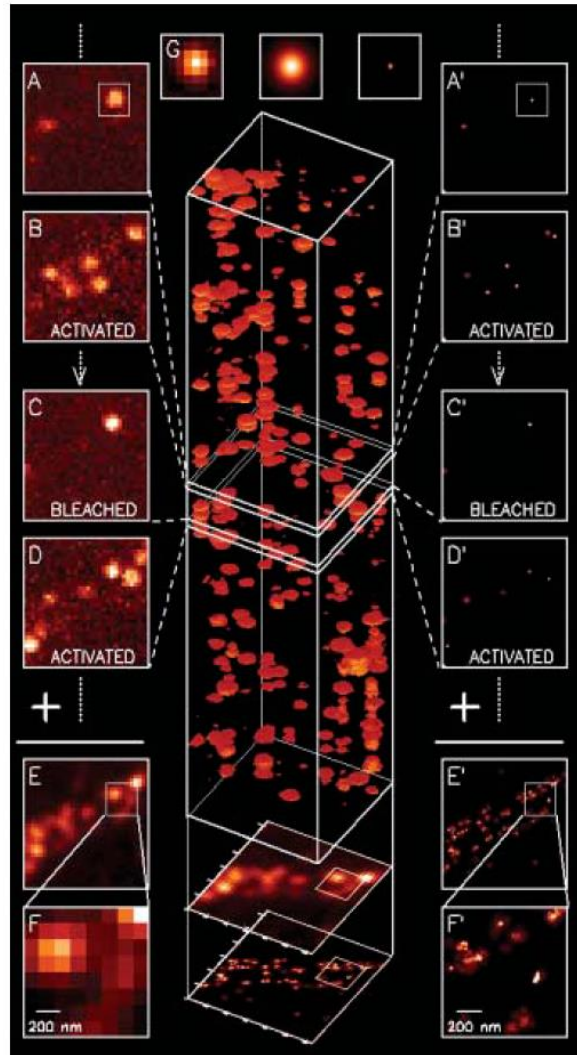
Dronpa

Kaede

...



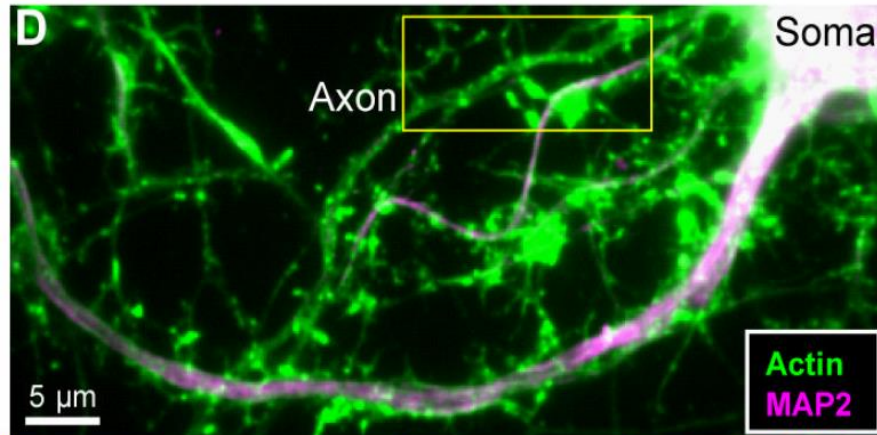
PALM: Photo-Activation Localization Microscopy



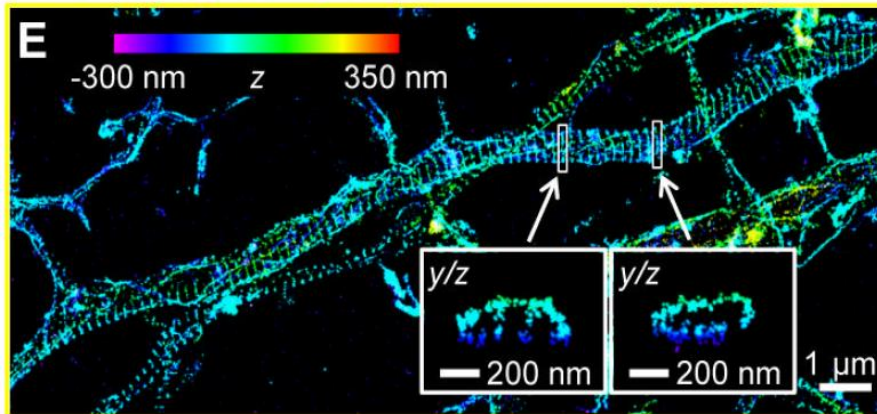
Super-resolution fluorescence microscopies in 2015

STORM/PALM/BALM/CALM...

epifluorescence

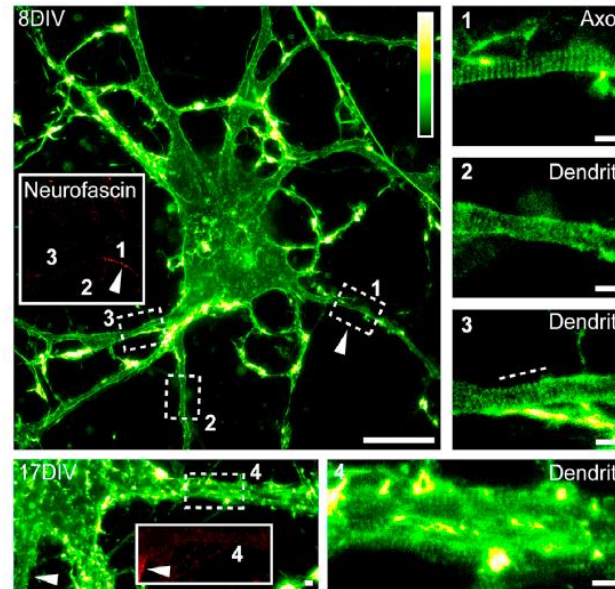
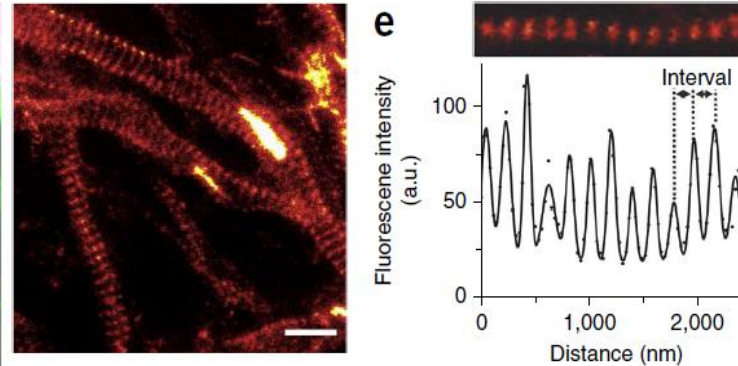


STORM



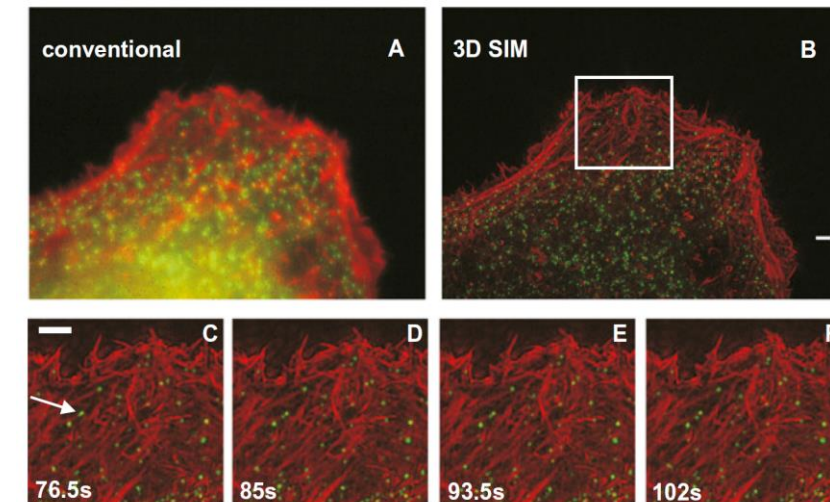
Xu et al, Science 2014

STED



Lukinavicius, Nature Methods 2015
D'Este, Cell Reports 2015

Structured illumination



Fiolka et al, PNAS 2012

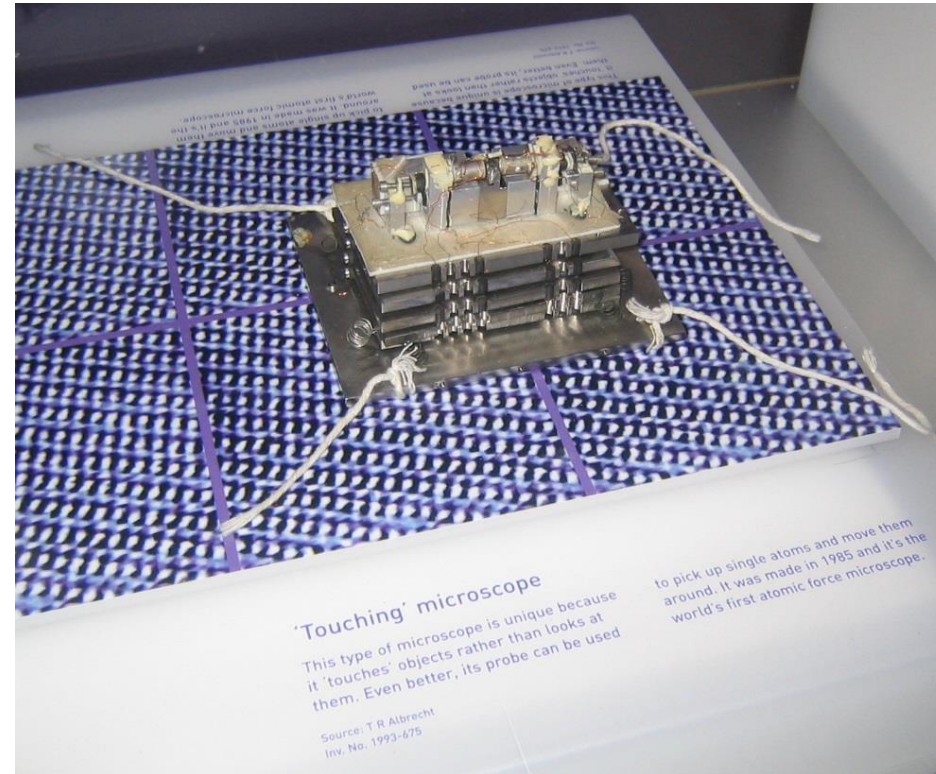
Pitfalls of optical microscopy

- **Requires a reporter fluorescent molecule**
 - technical limitation (chemical labelling, molecular biology)
 - potential changes in functionality (misfolding, steric hindrance, loss of a functional site)
- **Limited lateral (10nm) resolution, axial (20nm) and positional accuracy (1nm)**
 - Co-localization $\neq$ direct interaction
- **One can only see what is labelled**
 - Maximum number of simultaneously detected fluorophores is drastically limited
- **Radiation damage**
 - Fast photobleaching of the fluorophores
 - High power illumination leads to toxicity

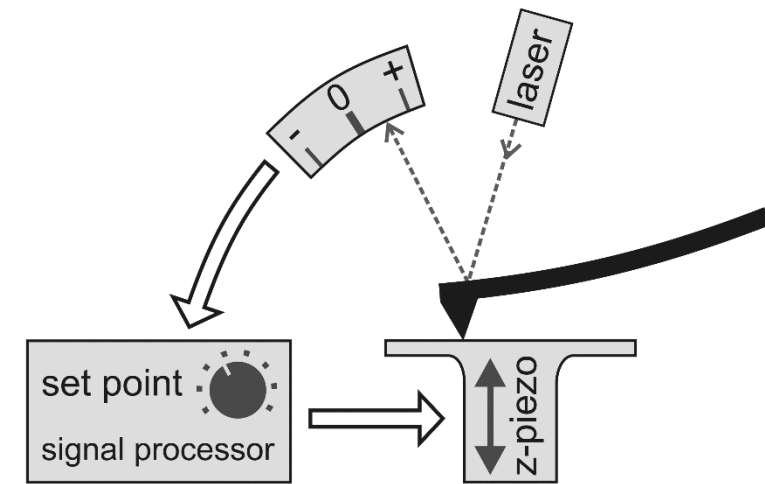
ATOMIC FORCE MICROSCOPY

→ “...The only method that can self-sufficiently perform nanometer-resolution imaging AND nano-manipulation of the sample with pN force resolution (...) in liquid and at room temperature...”

(Eghiaian et al, FEBS Letters 2014)

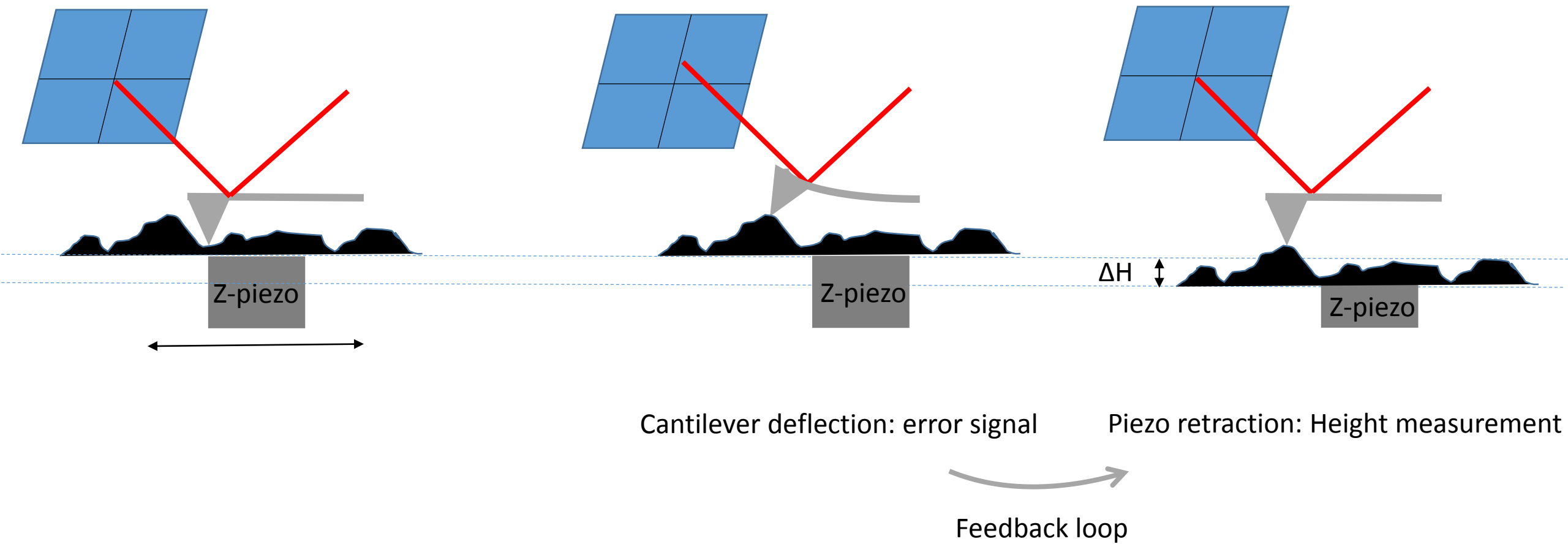


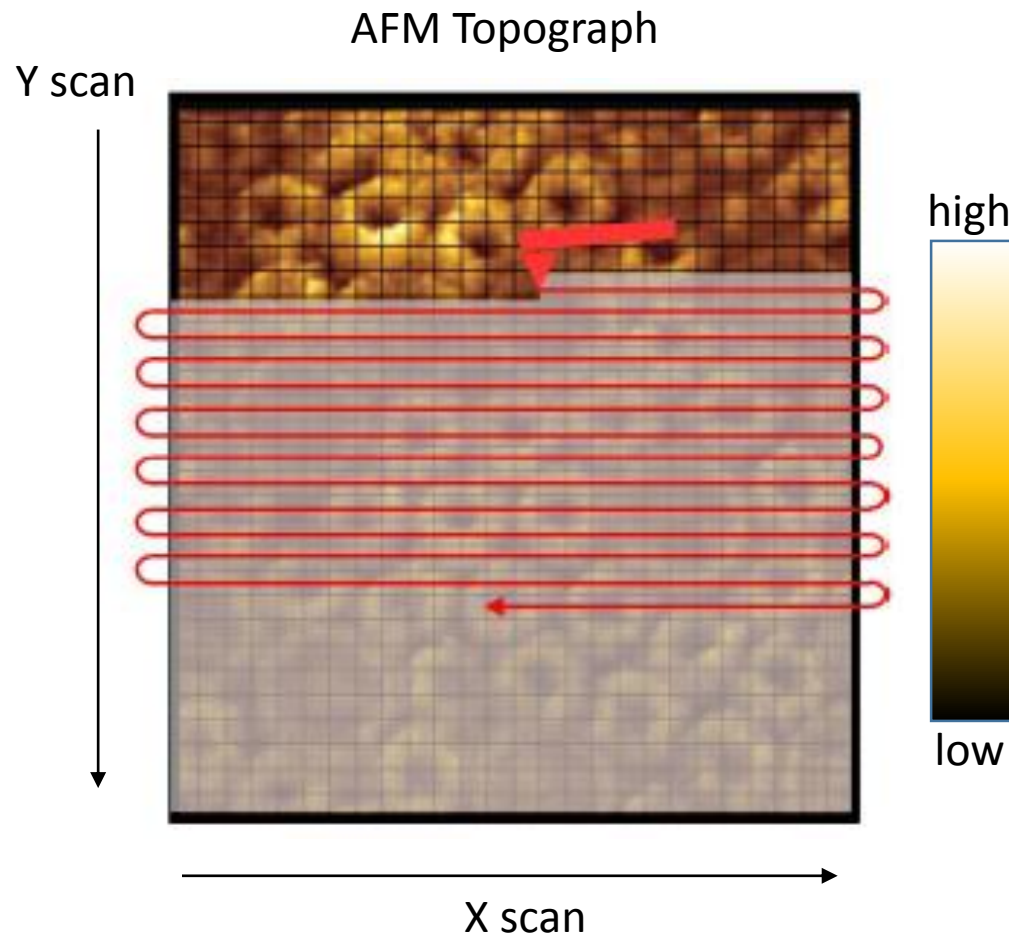
AFM basics



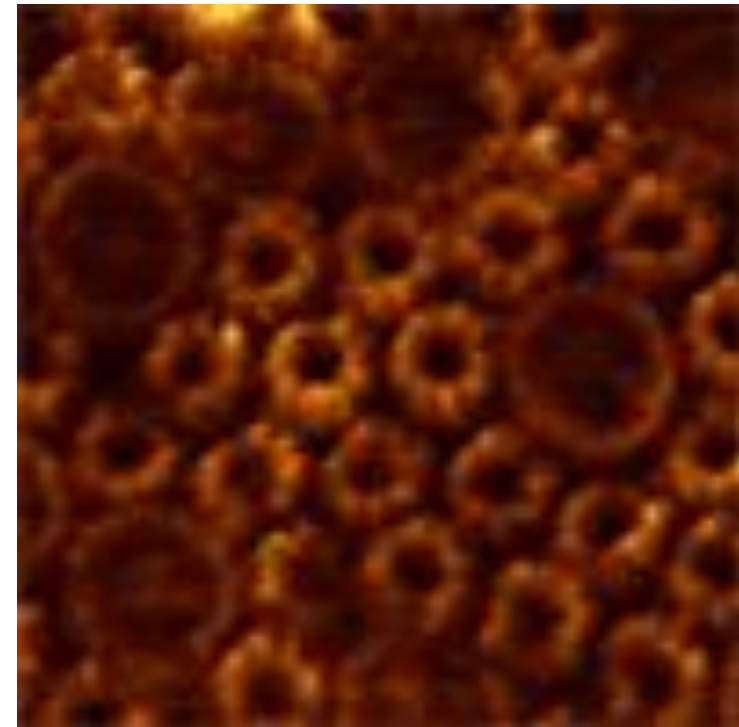
Eghiaian & Schaap, Methods Mol Biol 2011

Basic AFM: contact mode



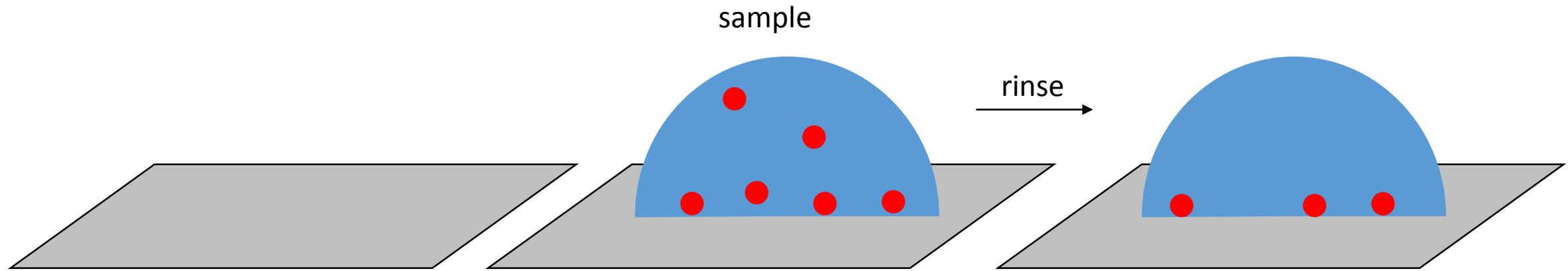


Native *Rhodospirillum*
Photometricum chromatophore



Scheuring & Sturgis, Science 2005

Sample preparation: surface adsorption of a sample



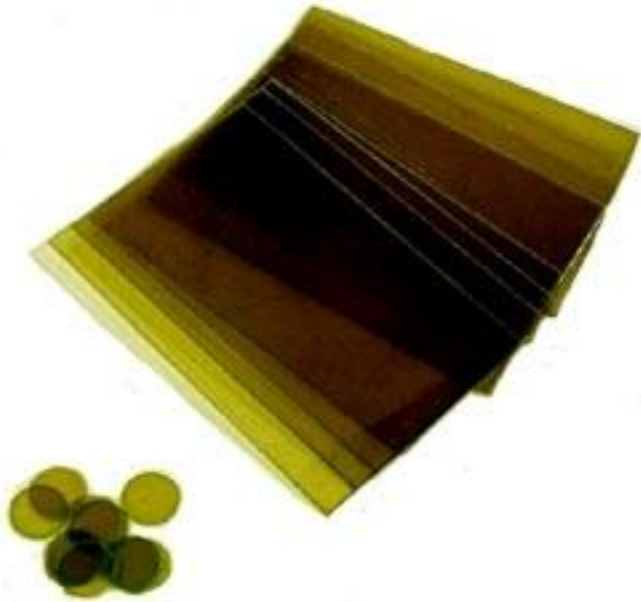
substrate

Unspecific adsorption

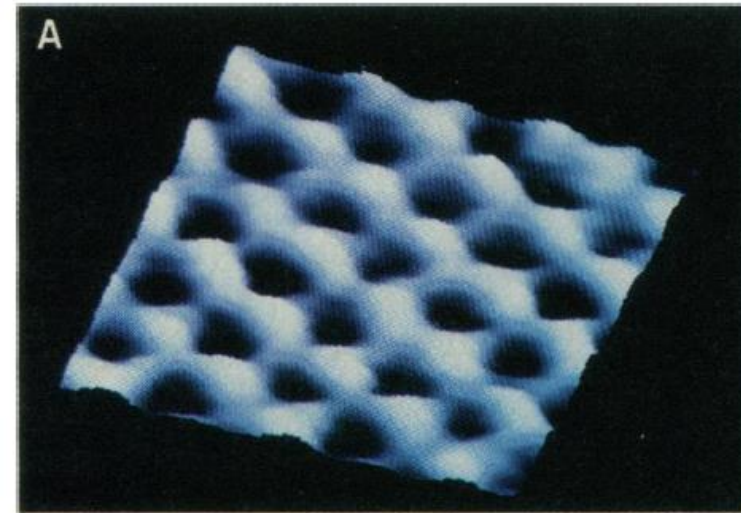
- Divalent cations help
- Surface functionalization (silanes, polylysine, patterning, covalent immobilization)
- planar bilayers (membrane proteins, 2D crystals...)

Sample preparation: mica as substrate

mica



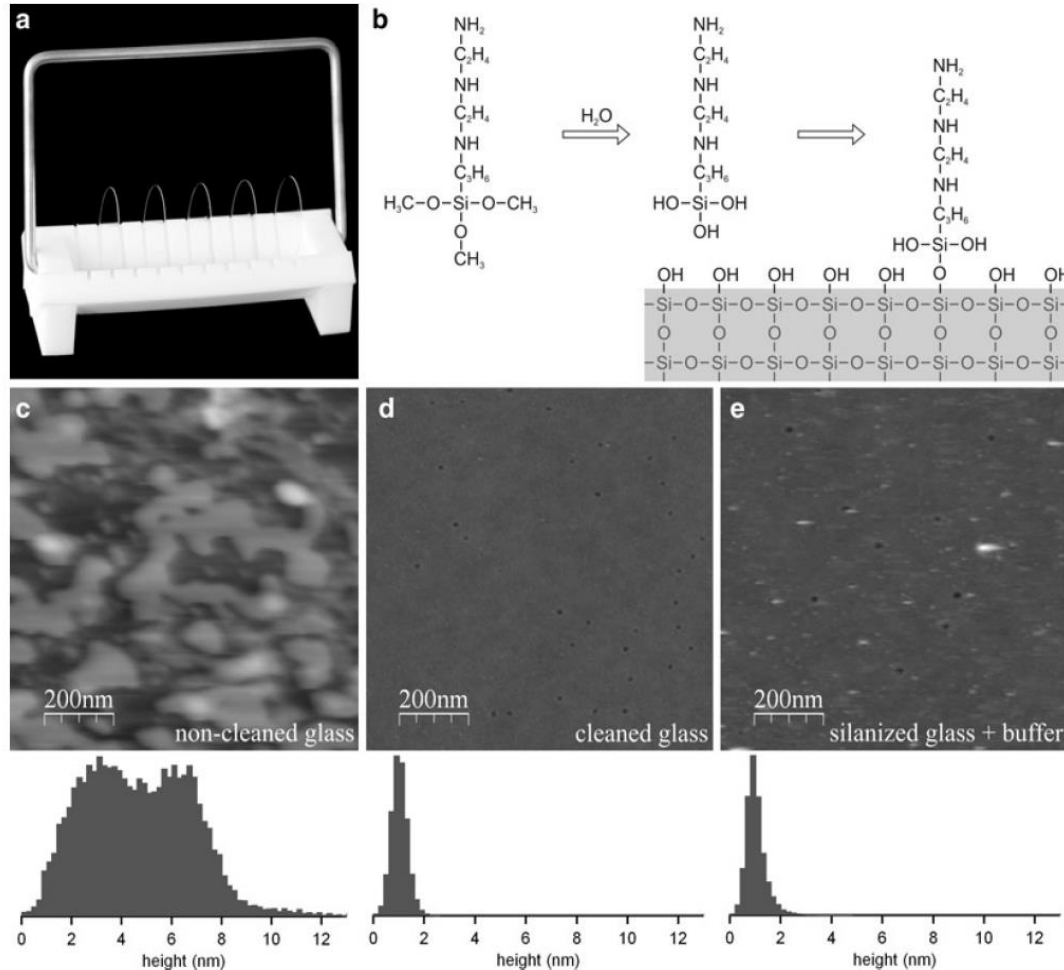
26x26 Å



Drake et al, Science 1989

Sample preparation: many other substrates

glass (bare or functionalized: silane...)



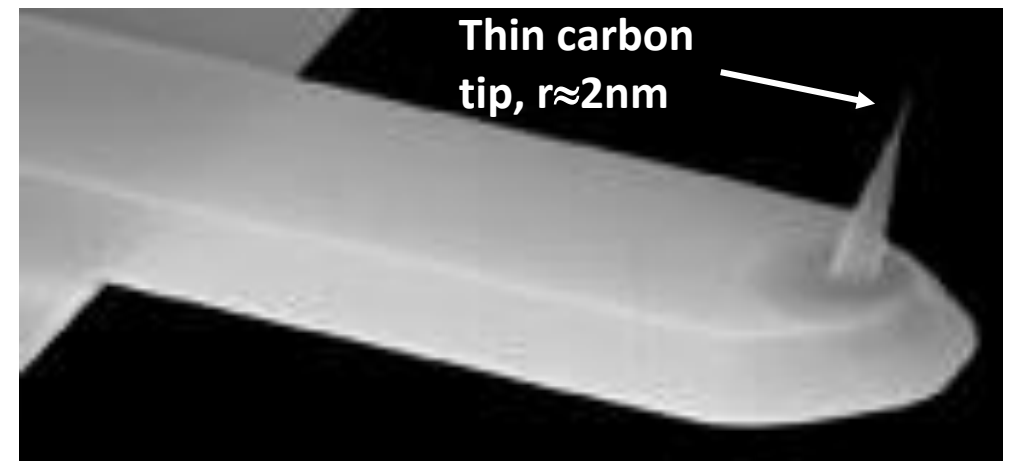
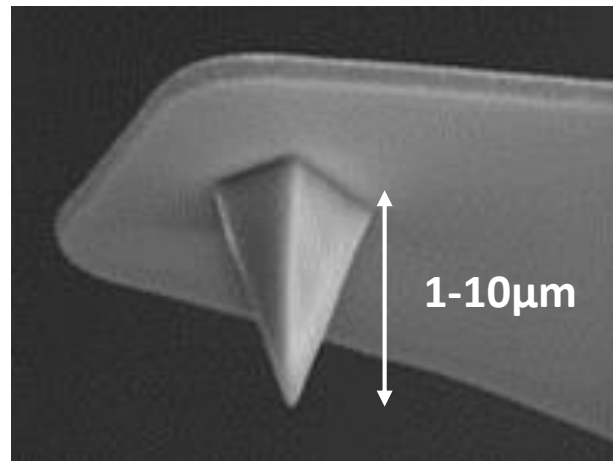
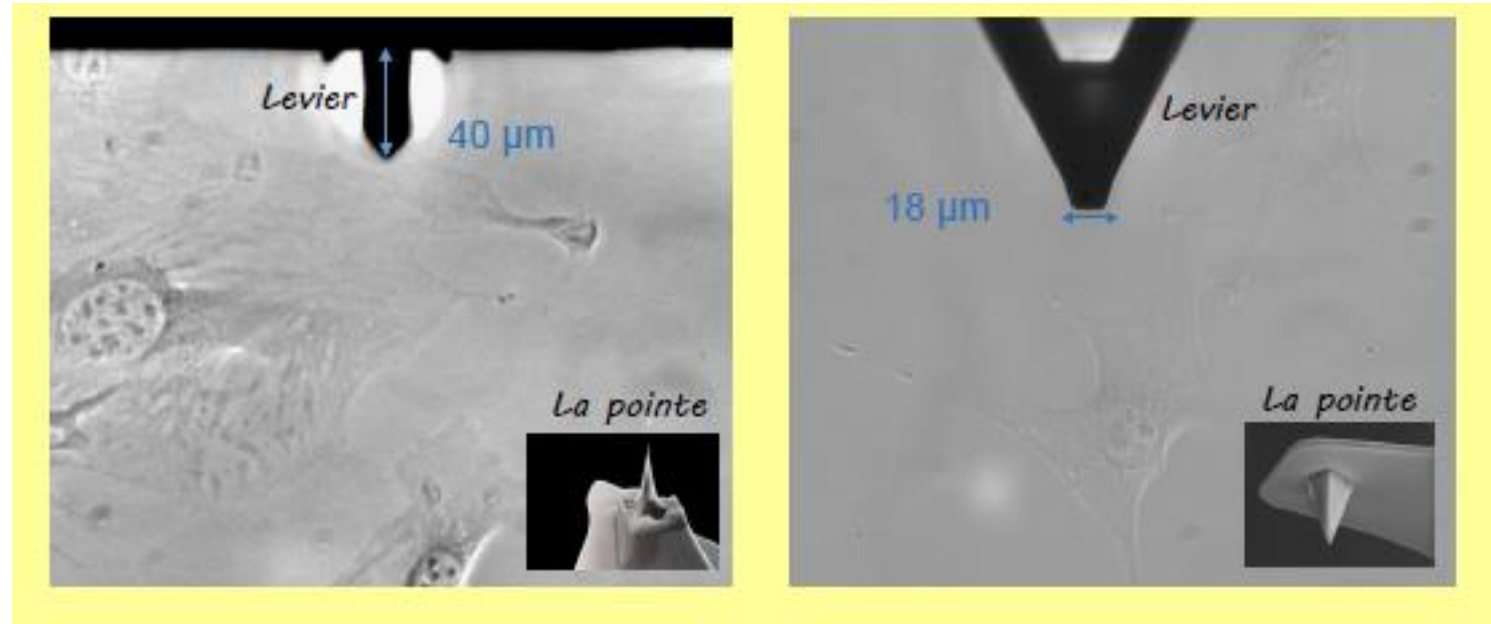
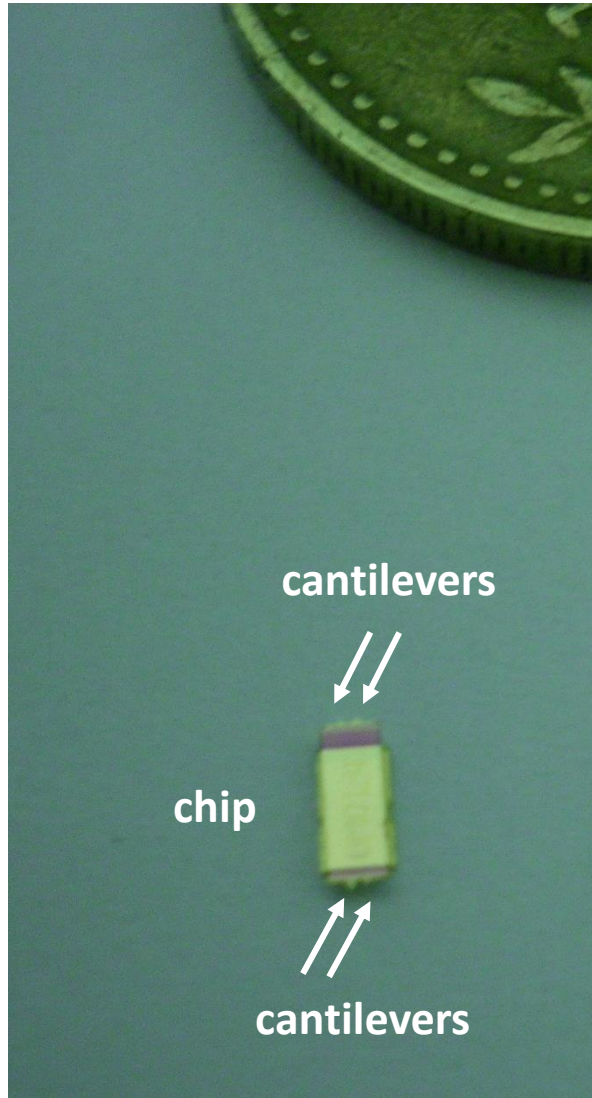
HOPG (highly ordered polygraphite)



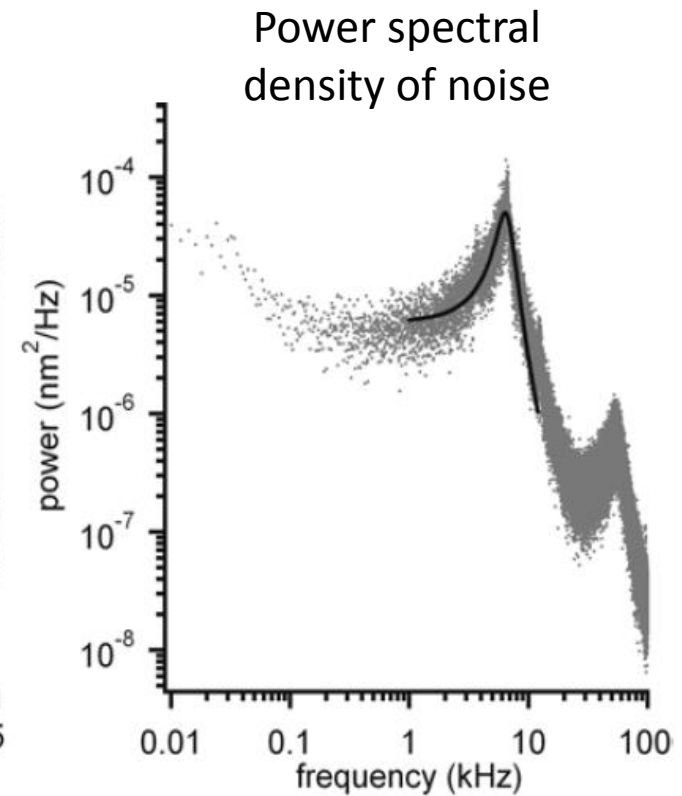
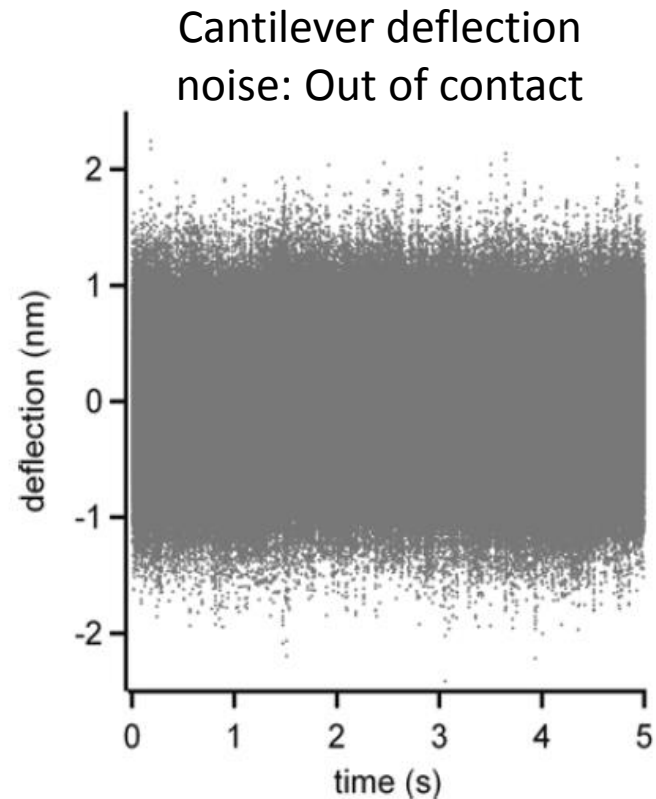
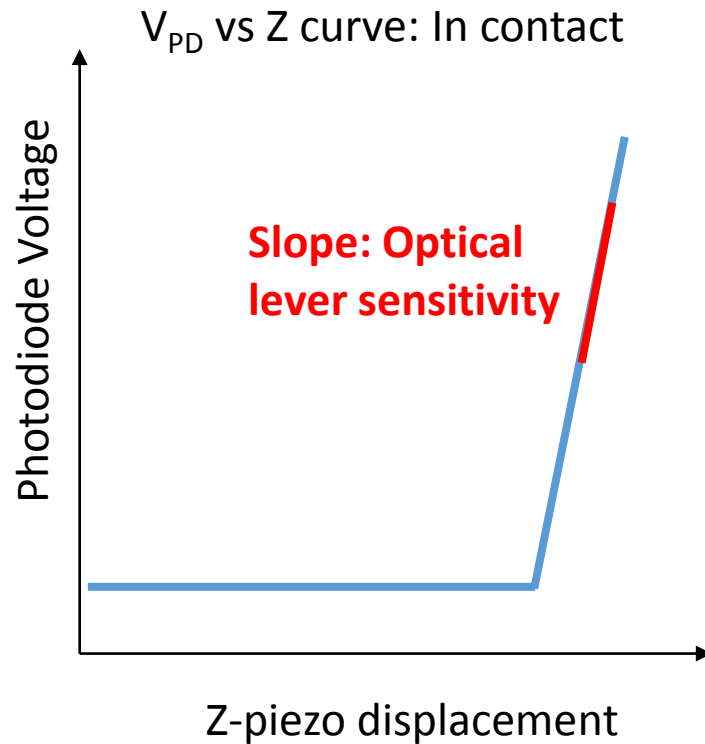
Silicon, silicon oxide...



Cantilevers



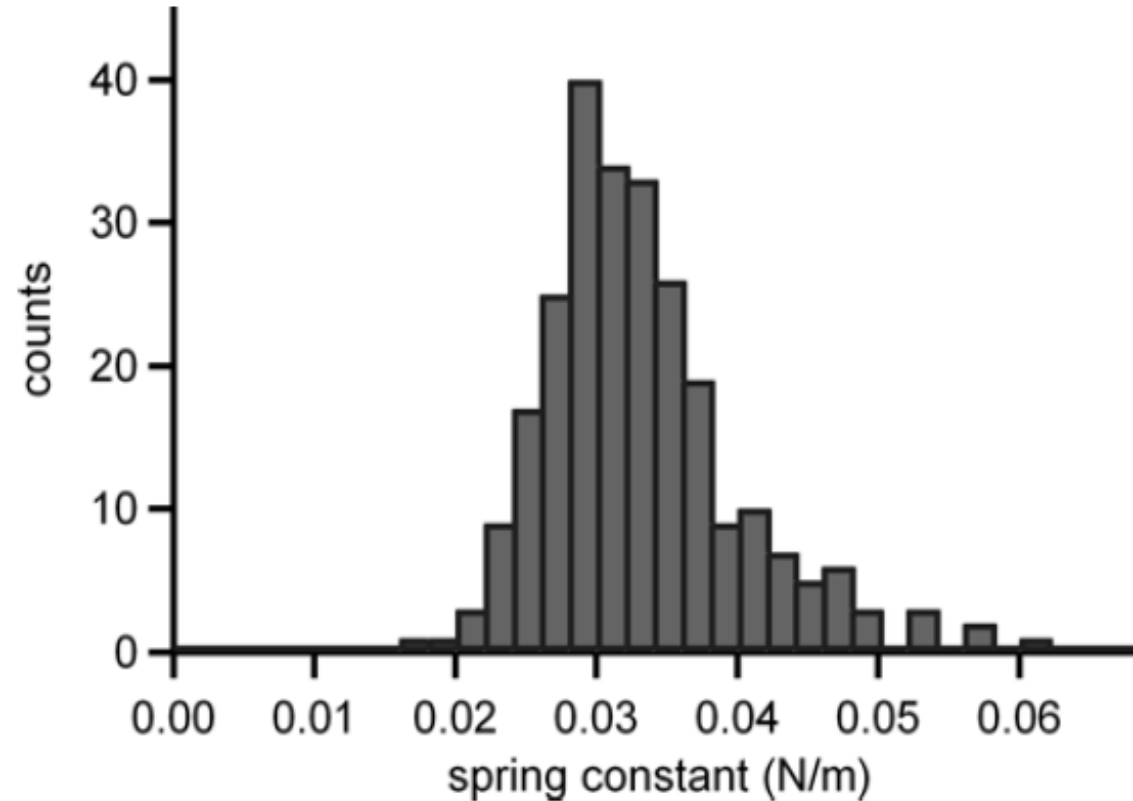
Cantilever calibration



$$S_x(f) = \frac{A}{f} + B + \frac{S_0 f_0^4}{Q^2 (f_0^2 - f^2)^2 + (f_0 f)^2}$$

$$k = \frac{2K_B T Q}{\pi S_0 f_0}$$

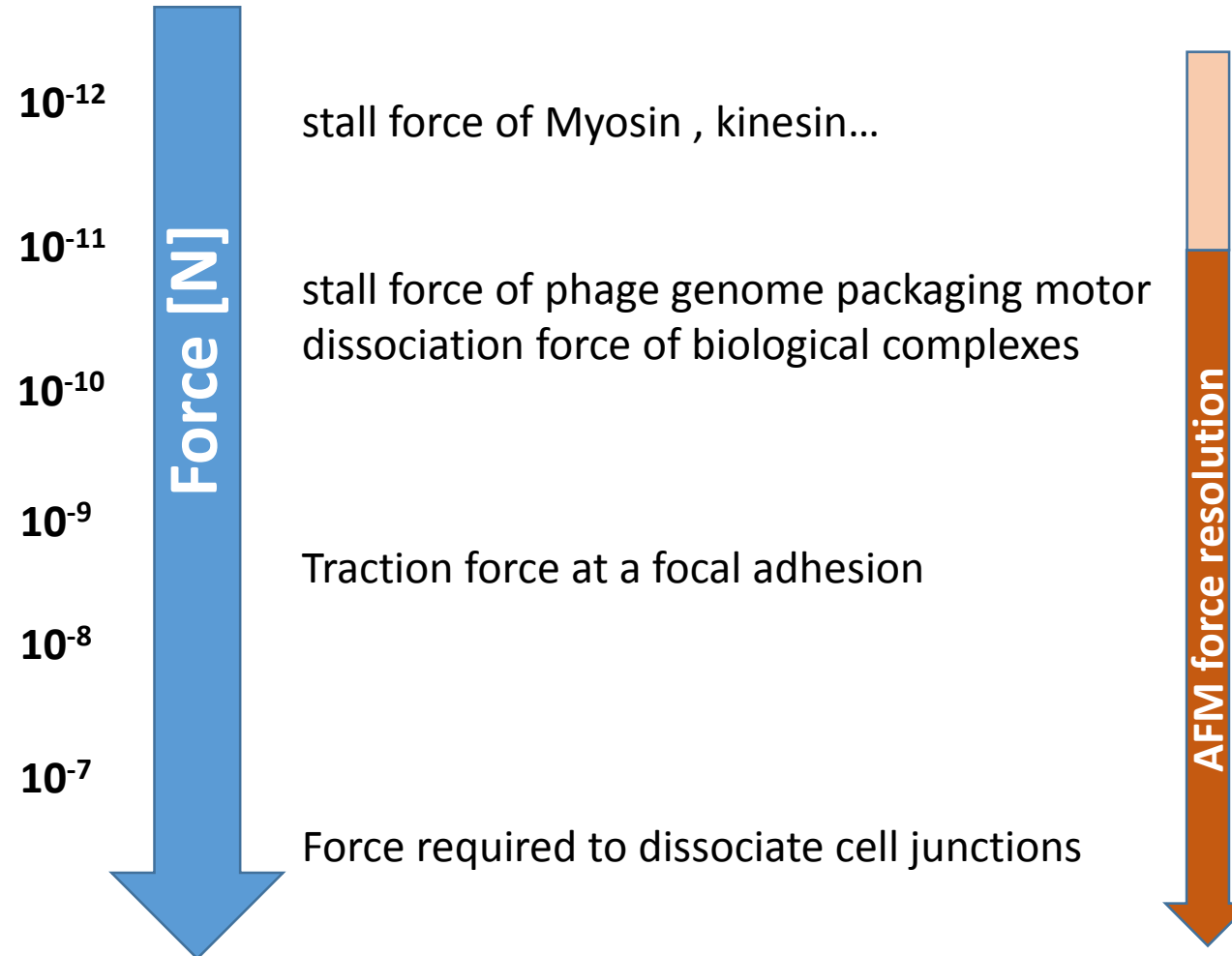
Cantilever calibration



Nominal: 30 pN/nm

**-> spring constant
calibration is important !**

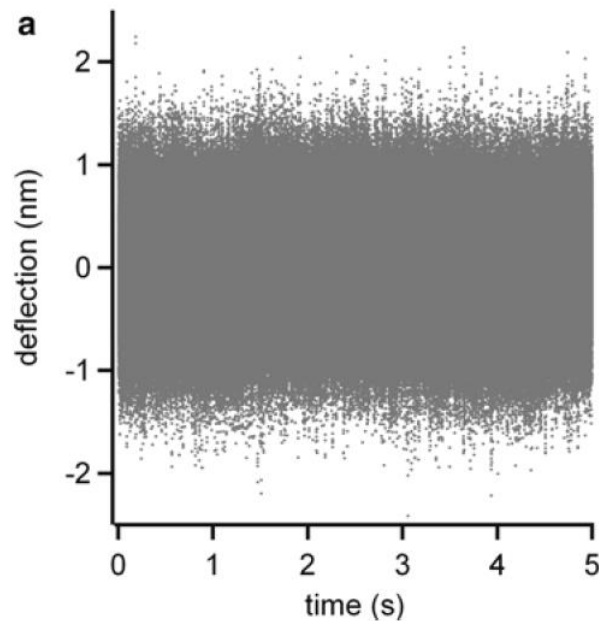
Forces in AFM-Biology



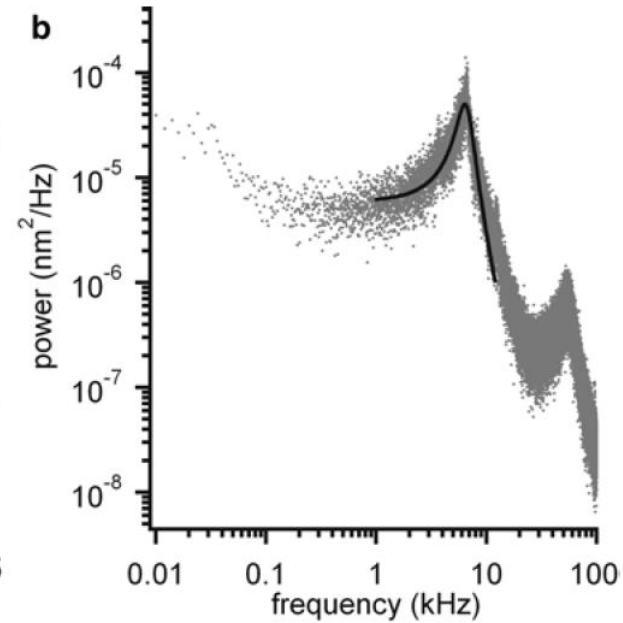
Cantilever theory: force noise

Deflection of a cantilever in liquid

Time-domain

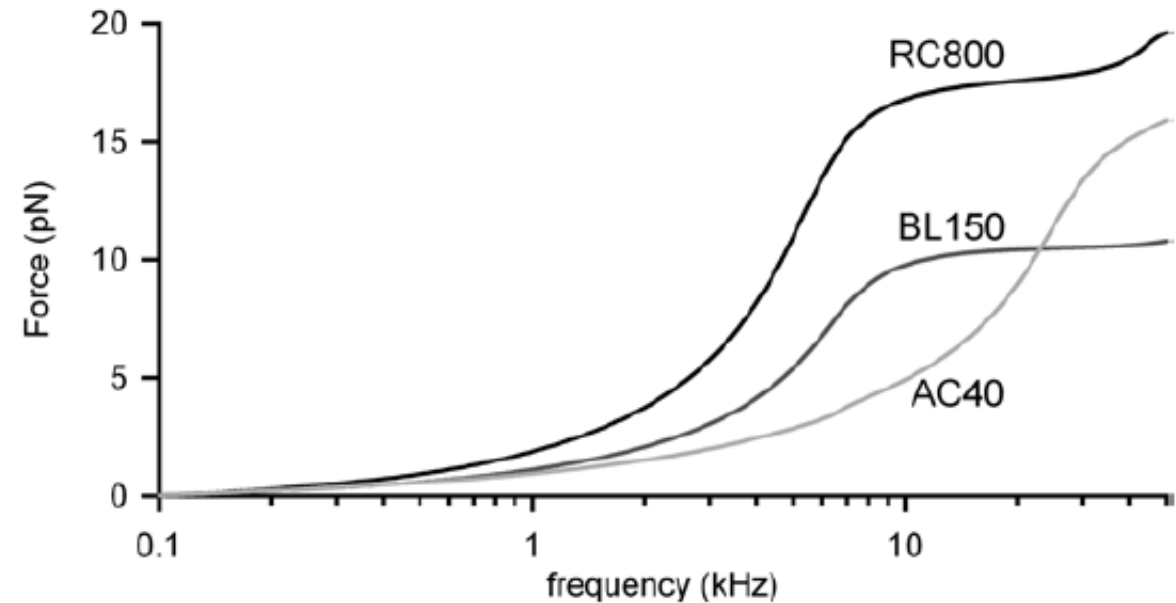


Power Spectrum (PSD)



Force noise of cantilevers in liquid

Integrated PSD * Spring constant



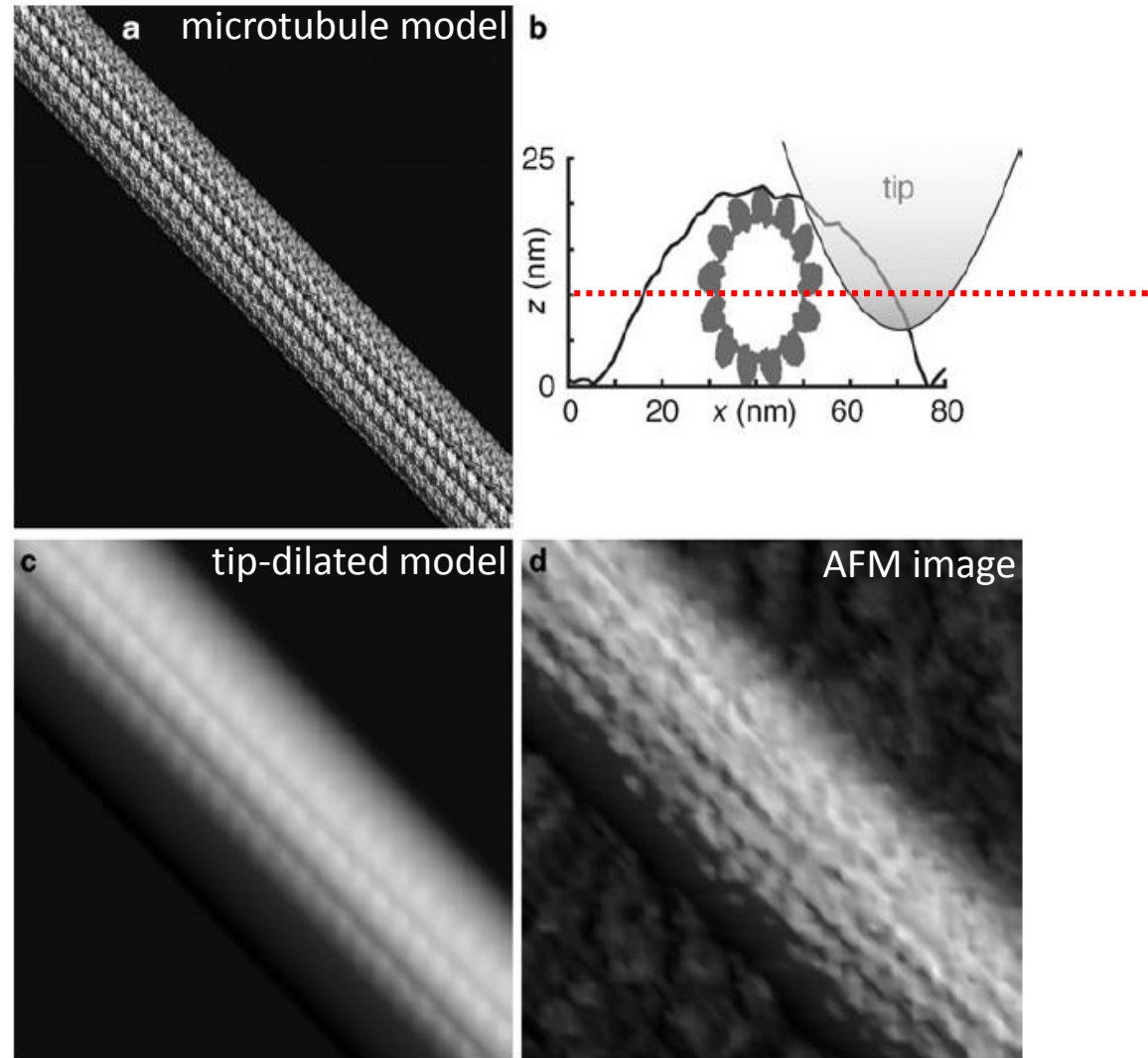
Force limit

$$\Delta F = \sqrt{4k_B T \Delta f \beta}$$

Δf : Bandwidth of experiment

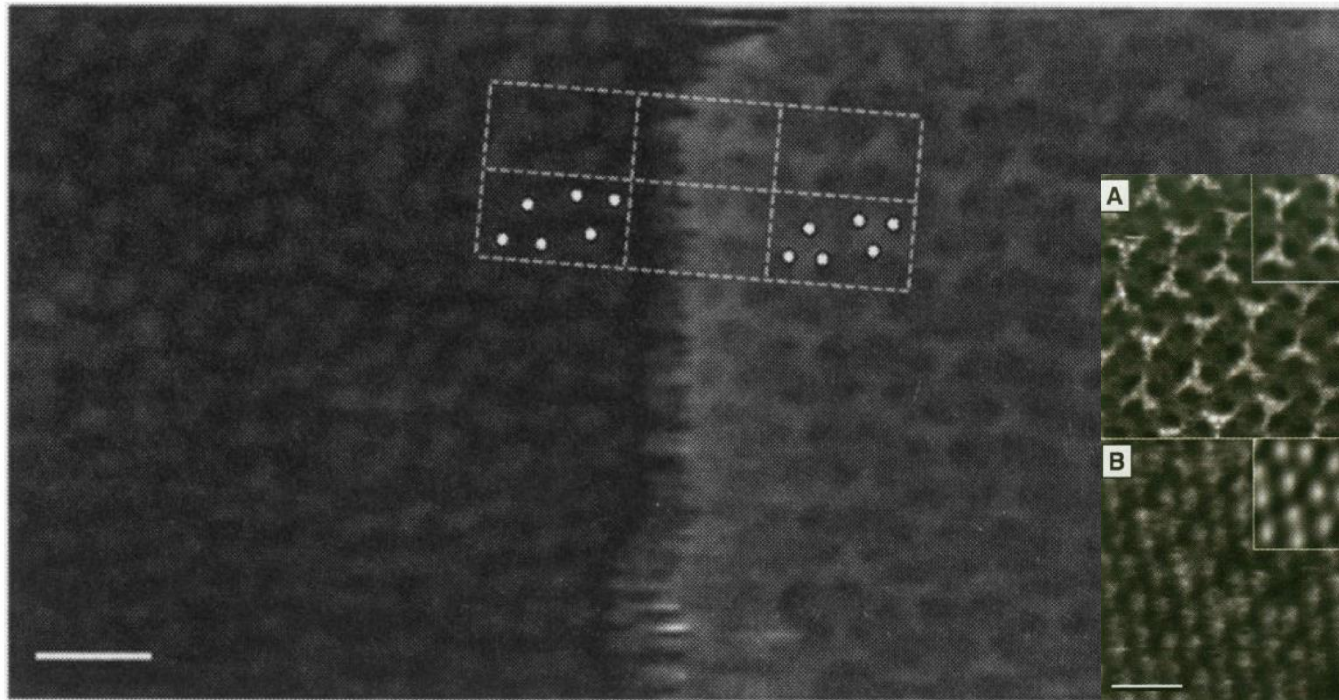
β : drag

Resolution limit: tip sample dilation



Top half
information is
more relevant

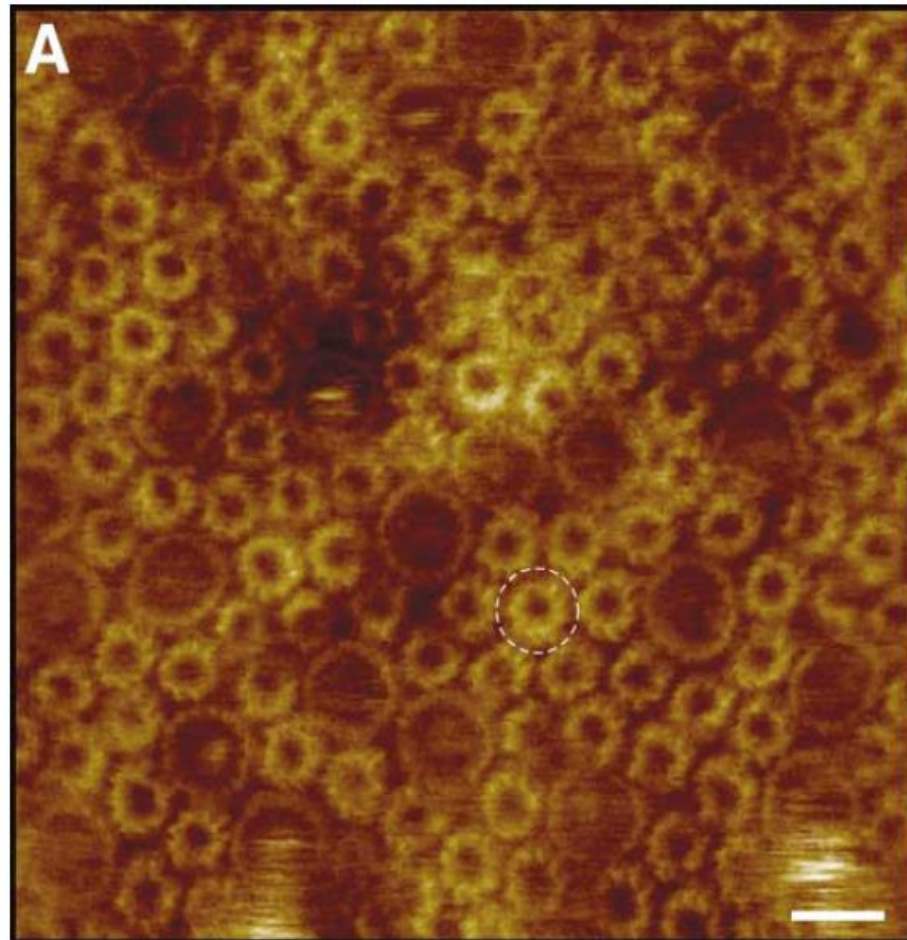
Membrane protein imaging by AFM



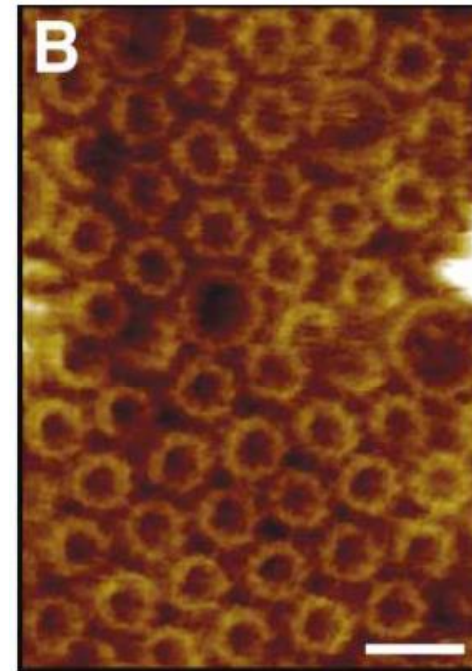
Membrane protein imaging by AFM

Native *Rhodospirillum Rubrum* chromatophore

High light conditions

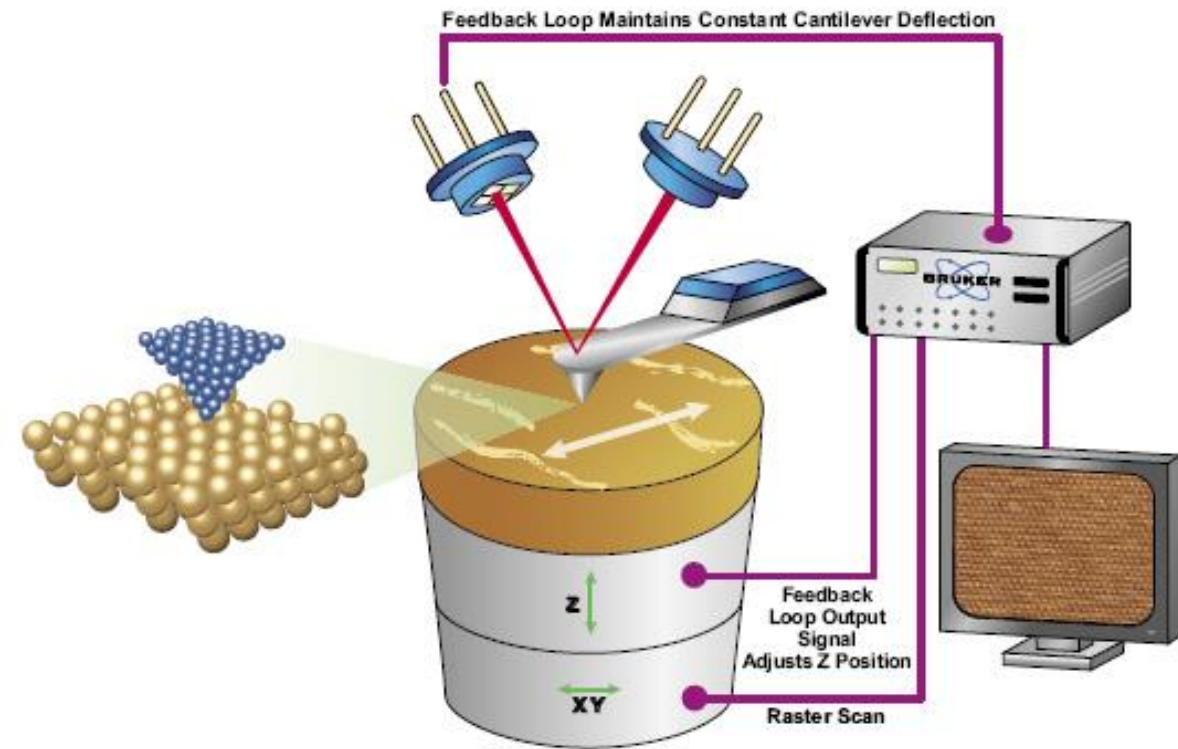


Low light conditions

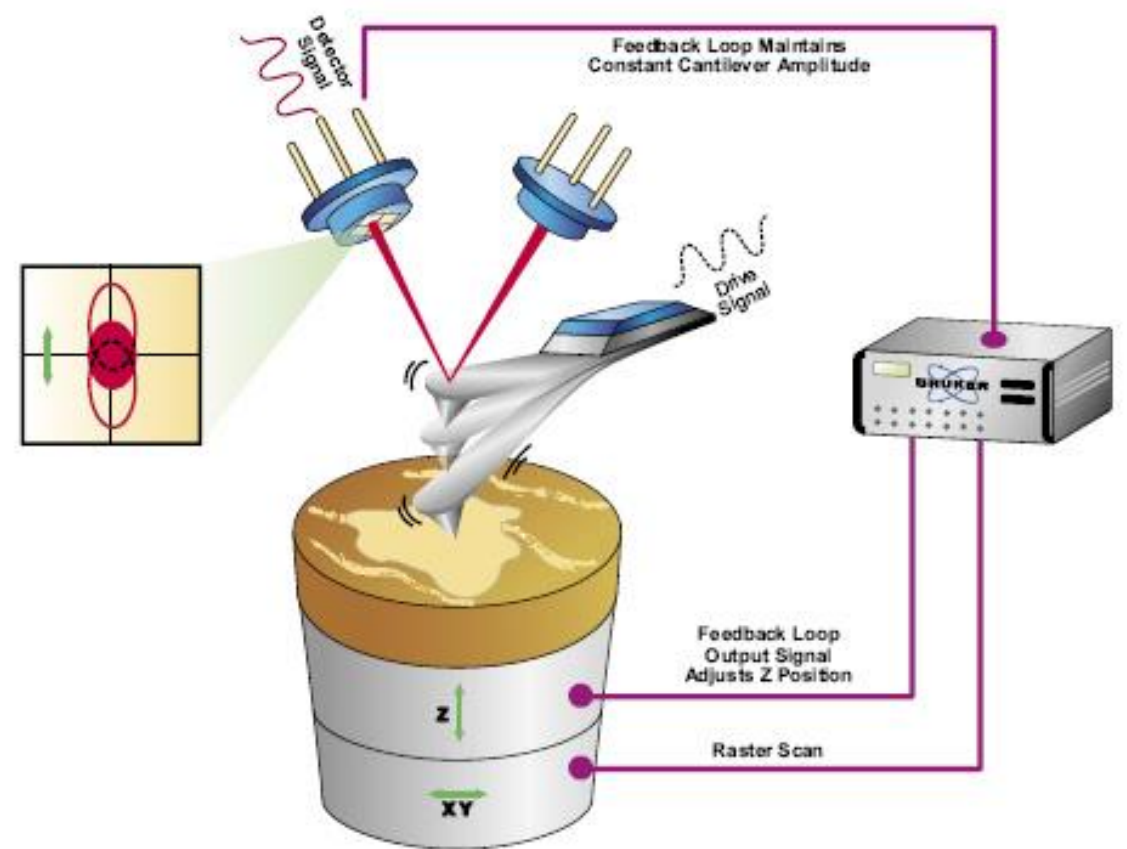


Intermittent contact mode

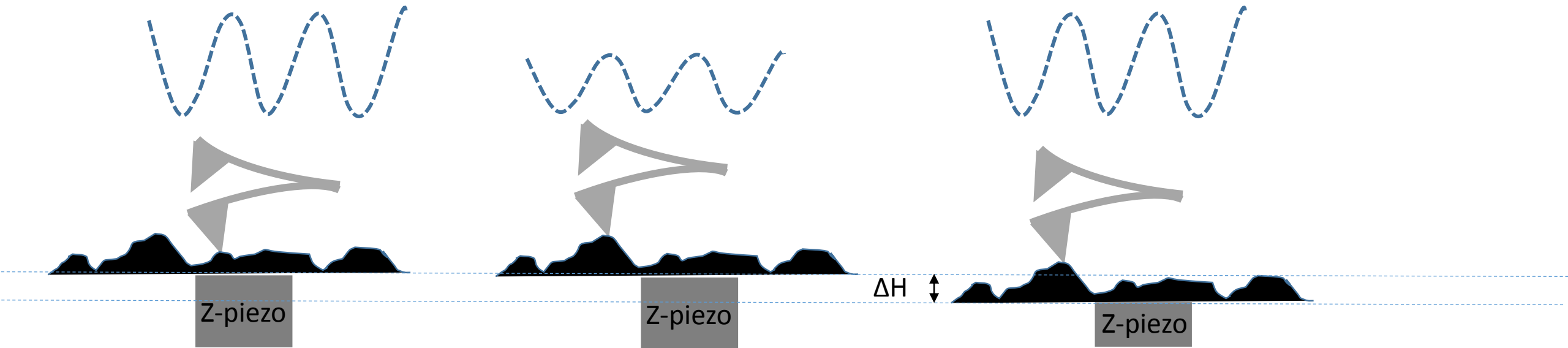
Contact mode (feedback on deflection)



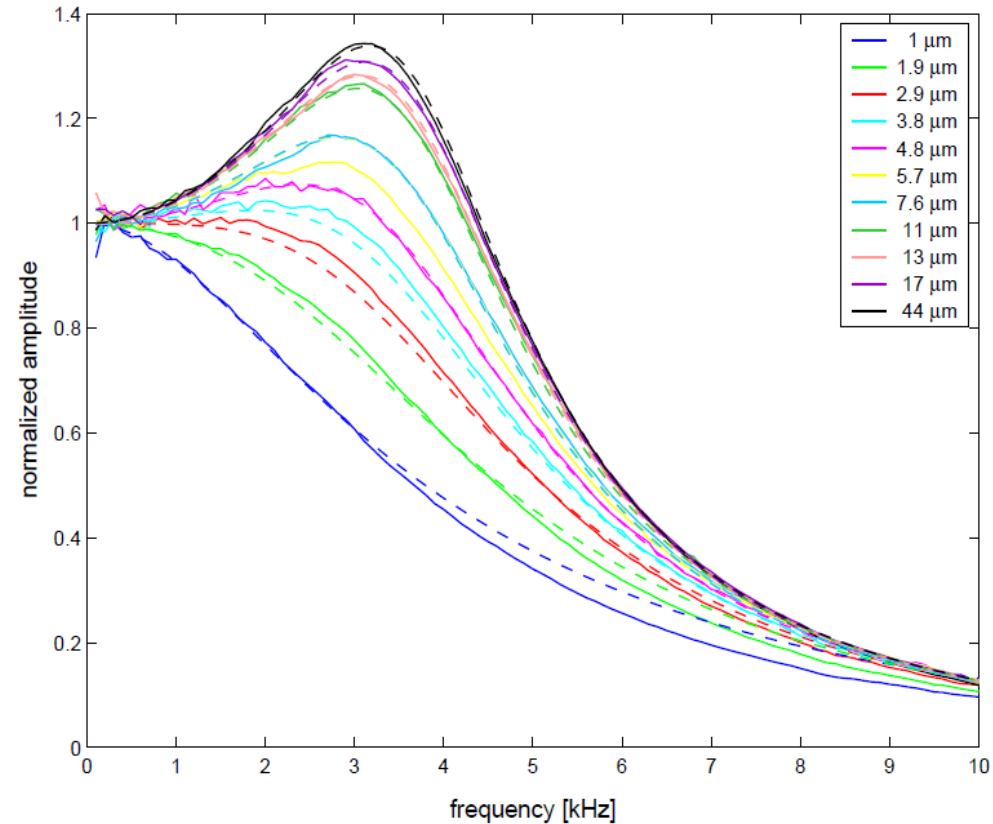
Tapping mode (feedback on amplitude)



Intermittent contact mode



Non-contact mode



Rankl et al, Ultramicroscopy 2004



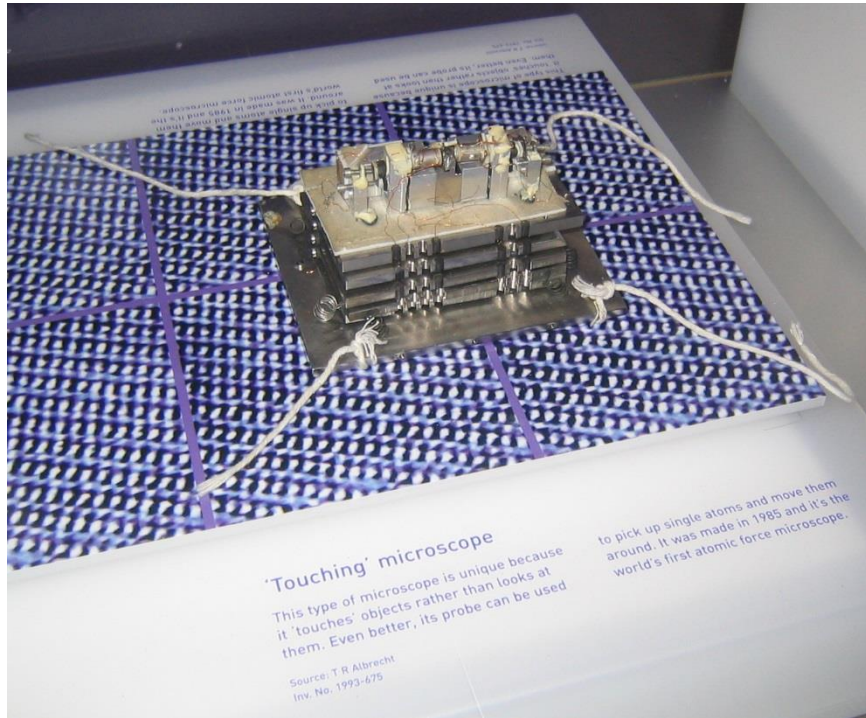
Feedback on frequency ???



frequency modulation AFM !
(non-contact mode)

Evolution of the atomic force microscope

1985



AFM prototype !
(contact mode)

1995



Veeco Nanoscope (now Bruker)

Contact, intermittent contact

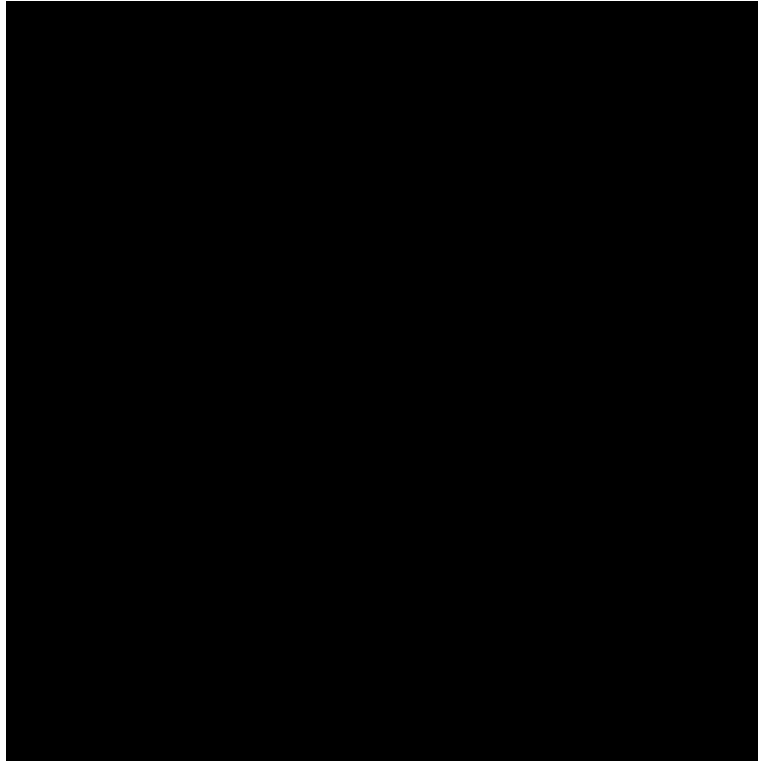
2005



JPK Nanowizard

Contact, force mapping,
intermittent contact...

Speed limit...



To break the speed limit...

- Increase scanner speed
- Increase cantilever resonance frequency (make smaller)
- Make feedback loop faster



Cantilever theory

Spring constant

$$k = E \cdot t^3 \cdot w / 4 L^3$$

E: Young's modulus

T: thickness

W: width

L: length

Resonance frequency

$$f_0 = \sqrt{k/m}$$

m: cantilever mass

THE Breakthrough: High-speed AFM

2010

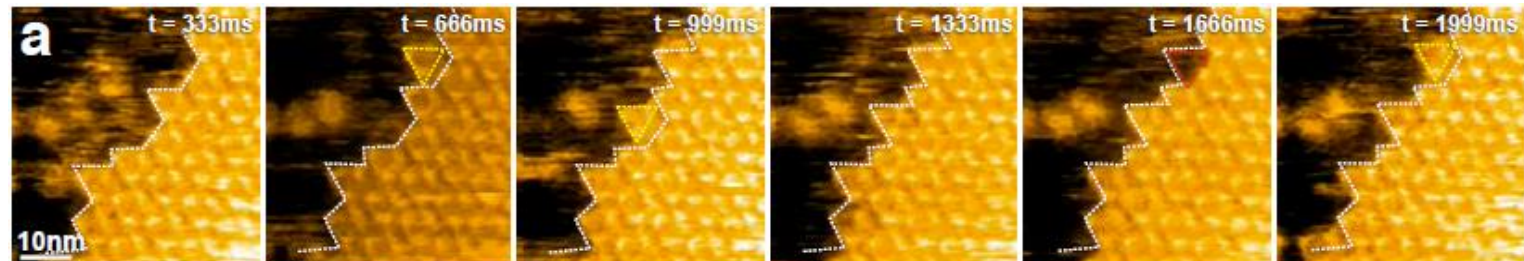


Pr. Toshio Ando

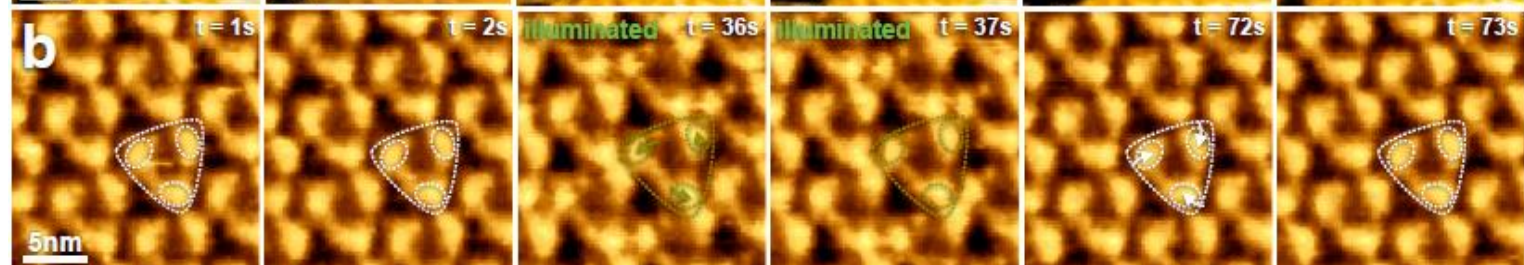


Dynamic imaging of biological membranes by HS-AFM

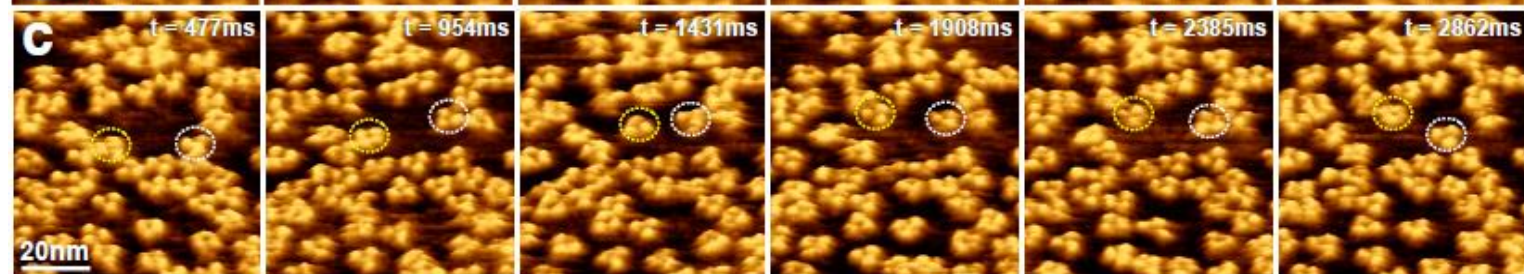
Purple membrane 2D
crystal formation



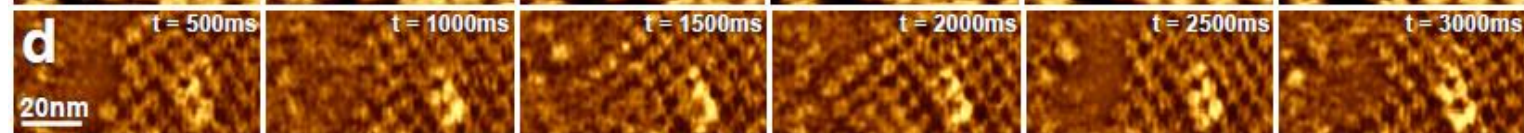
Bacteriorhodopsin
light response



Omp-F protein-protein
interactions in native
membranes



AQP0 assemblies

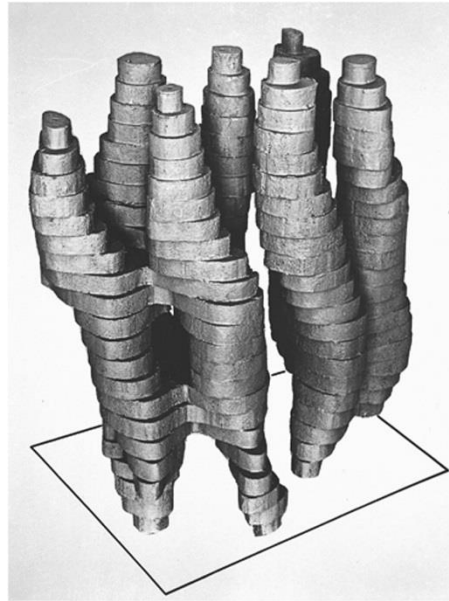


Dynamic imaging of biological membranes by HS-AFM

Bacteriorhodopsin from the purple membrane of *Halobacterium Salinarum*

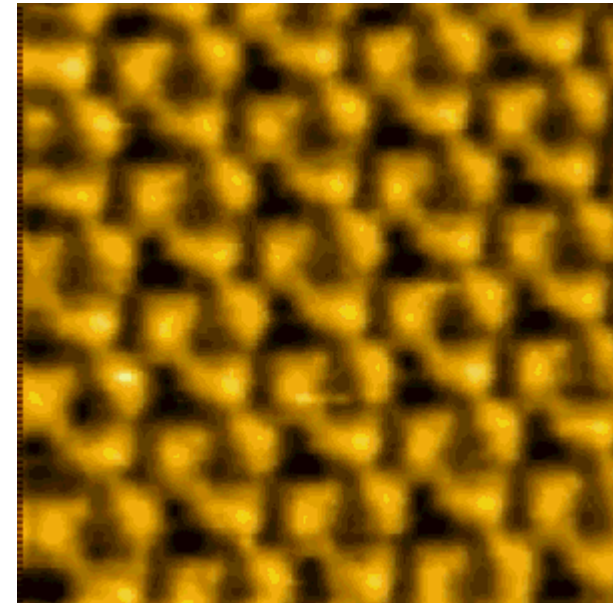
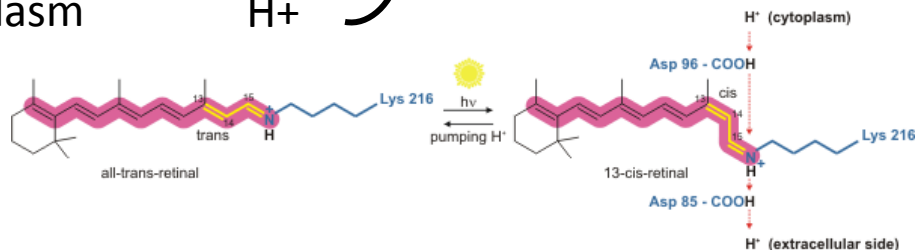
Extracellular side

H⁺



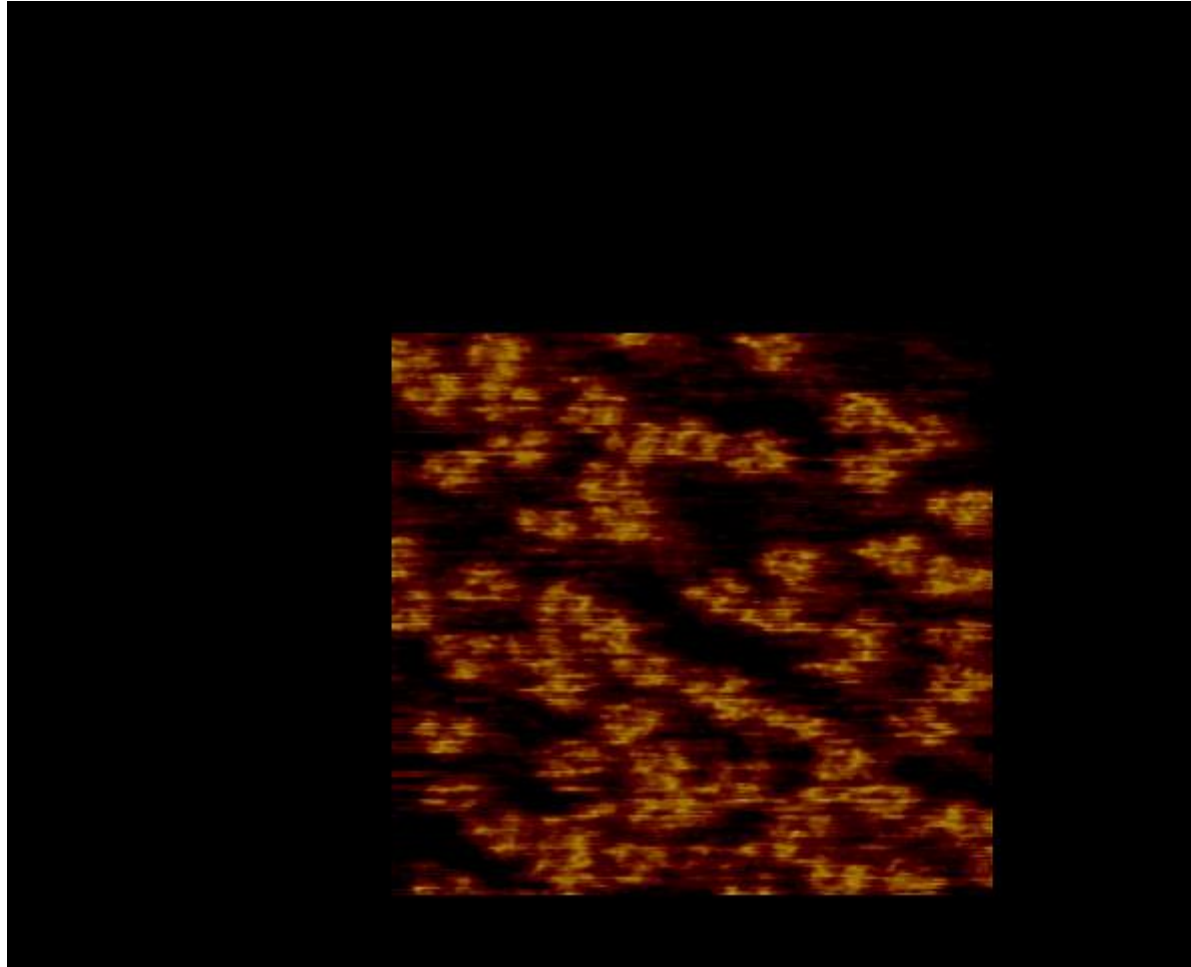
cytoplasm

H⁺

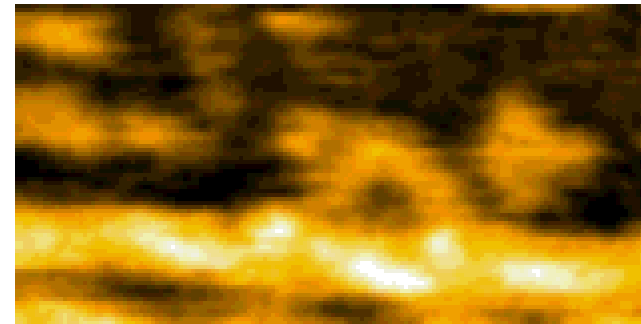
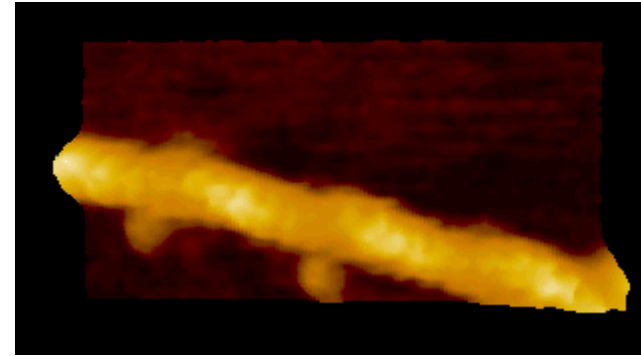
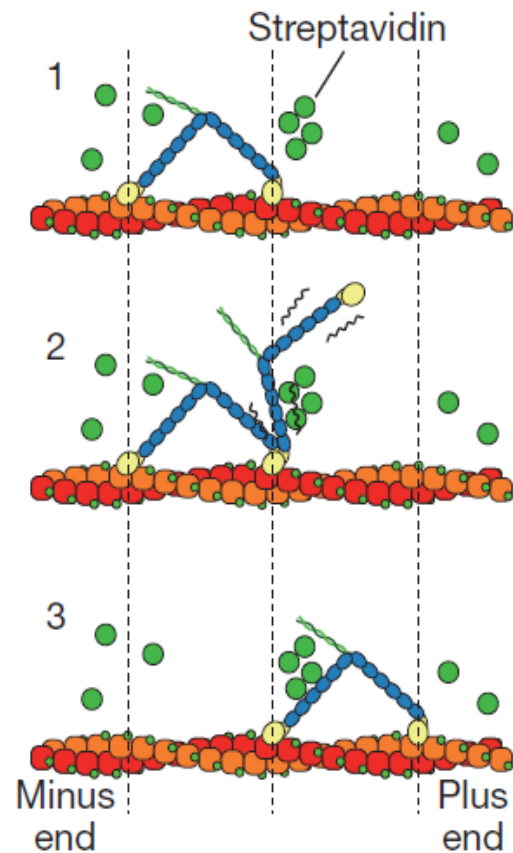
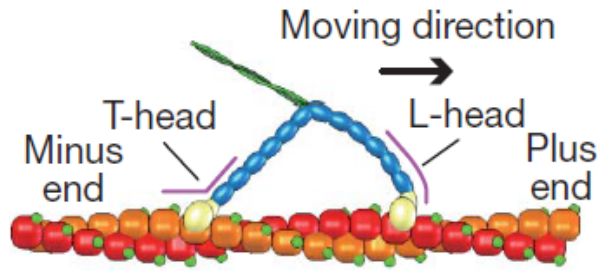


Henderson R, Unwin PN., *Nature* 1975
Shibata et al, *Nature Nanotechnology* 2010

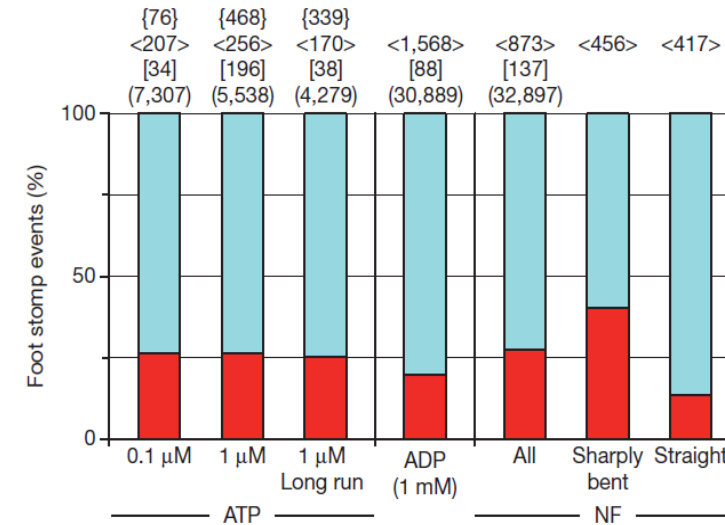
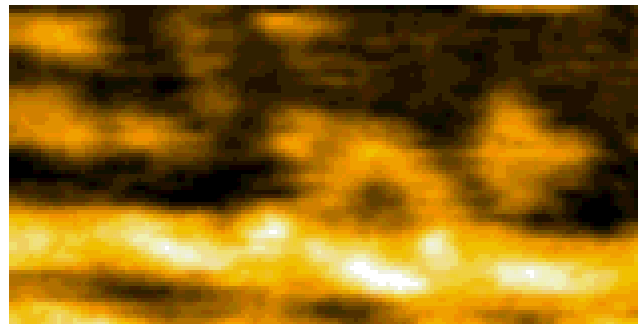
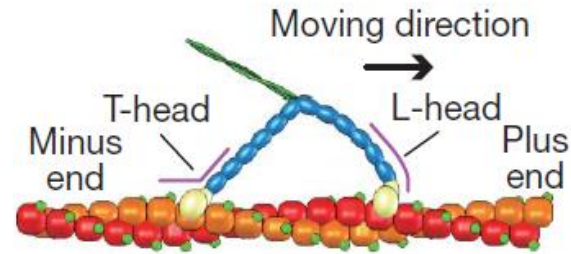
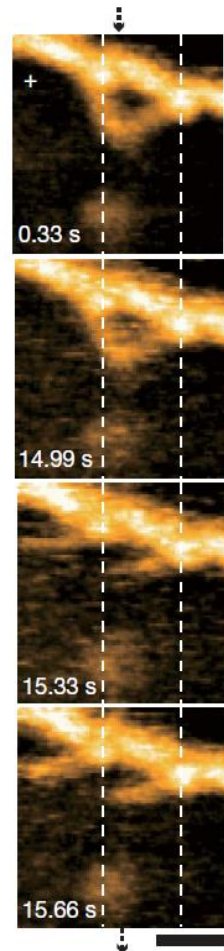
Dynamic imaging of biological membranes by HS-AFM



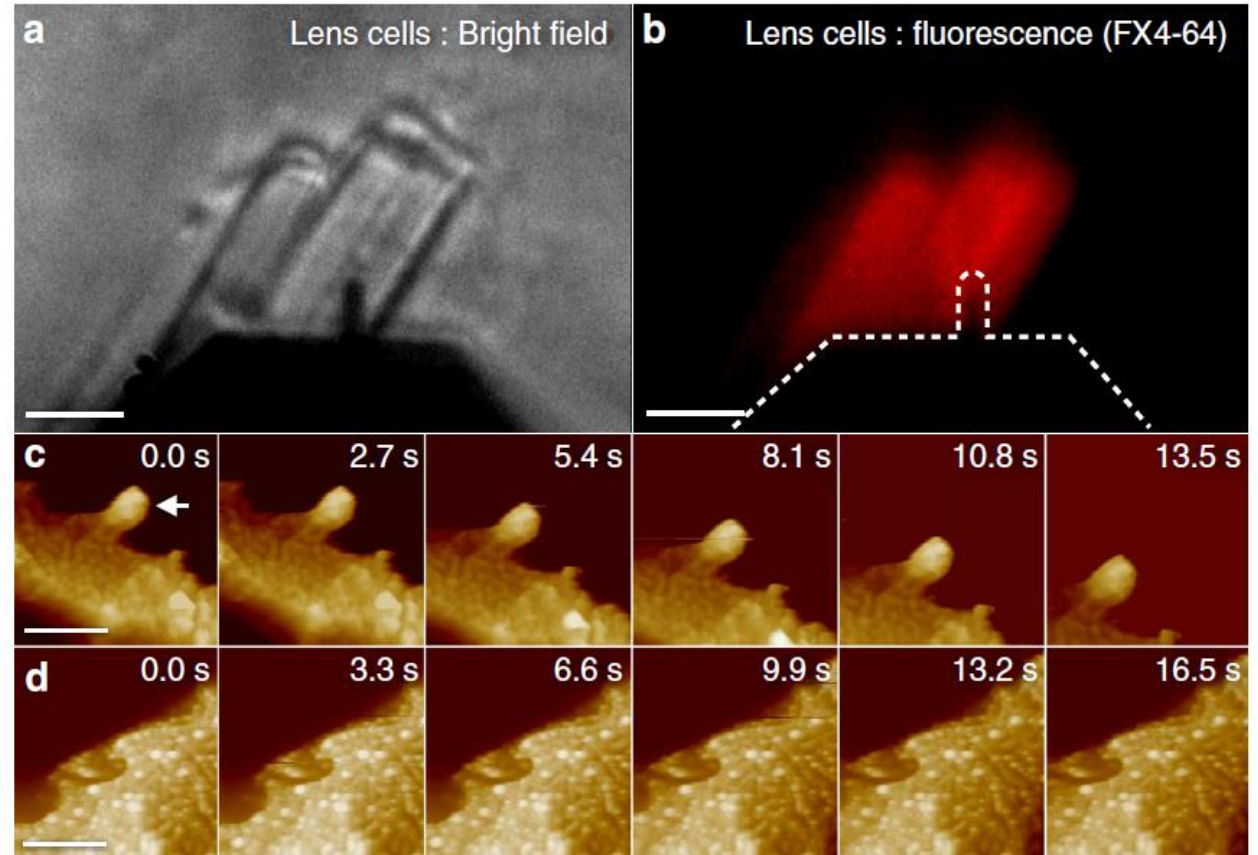
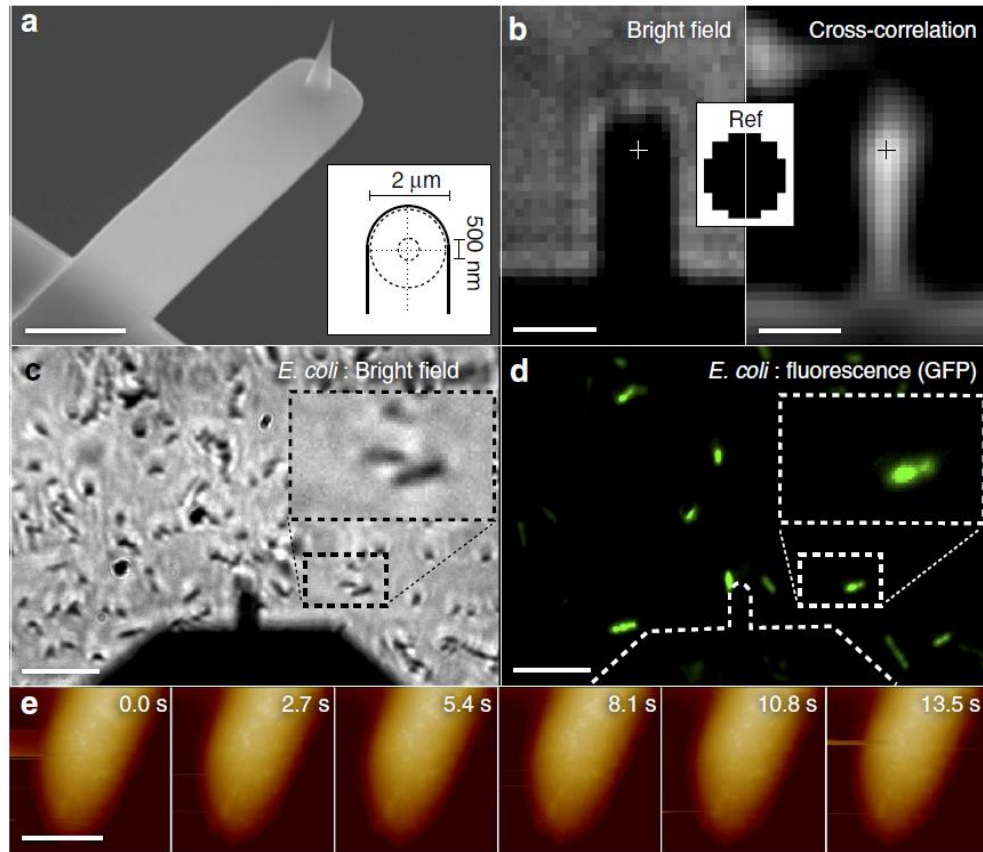
Myosin V walk on actin seen by HS-AFM



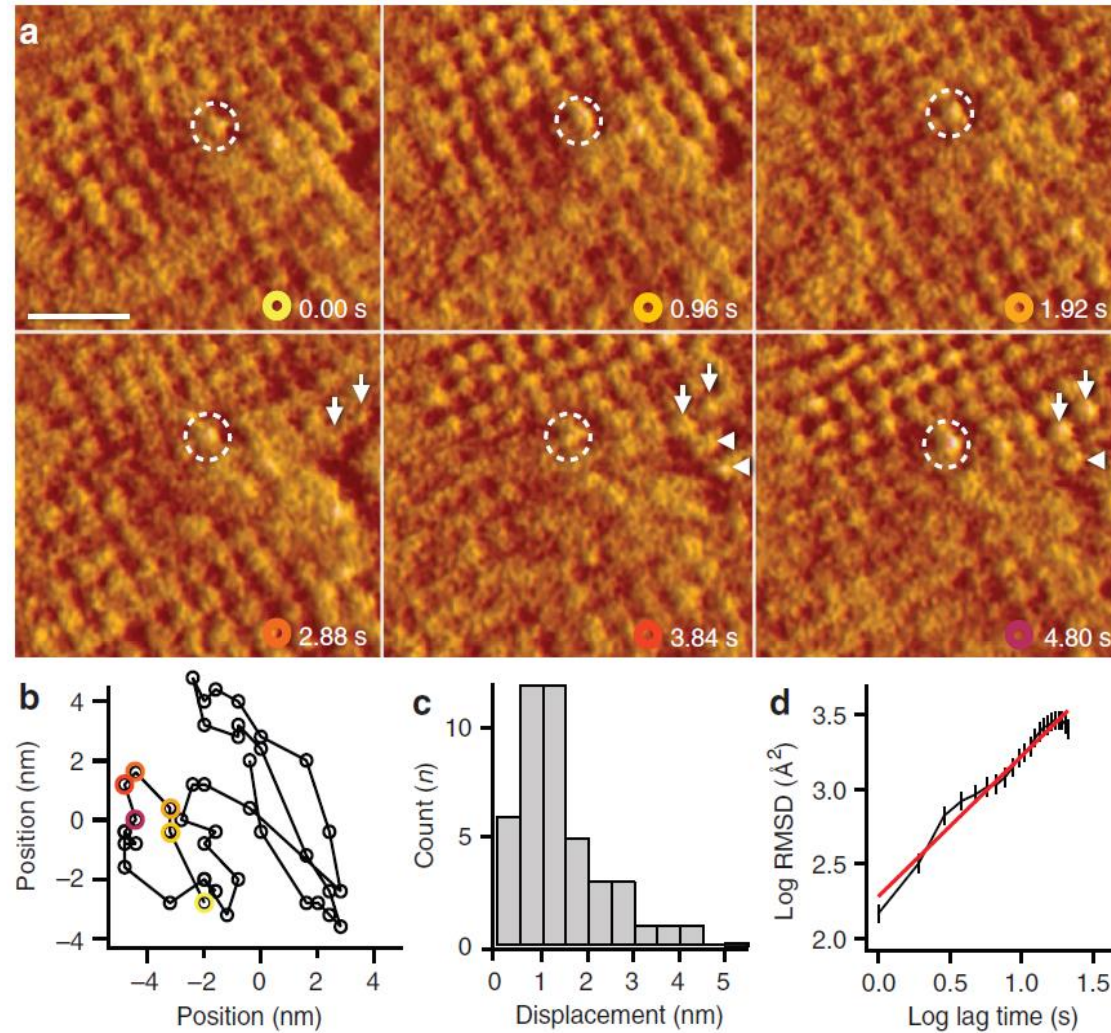
Myosin V walk is driven intramolecular tension release



HS-AFM+OM: Imaging of lens cells

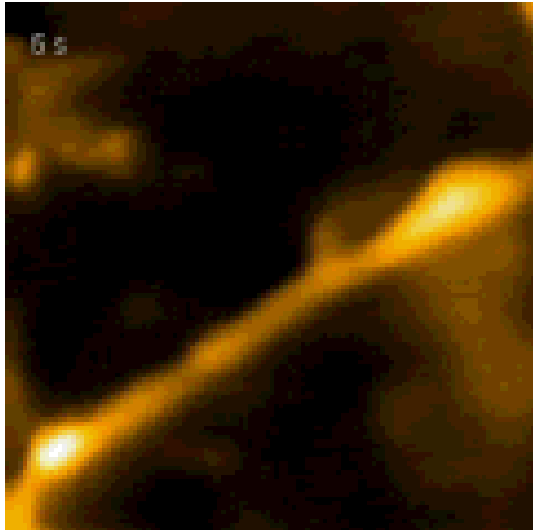


HS-AFM+OM: Characterization of lens cells junctions



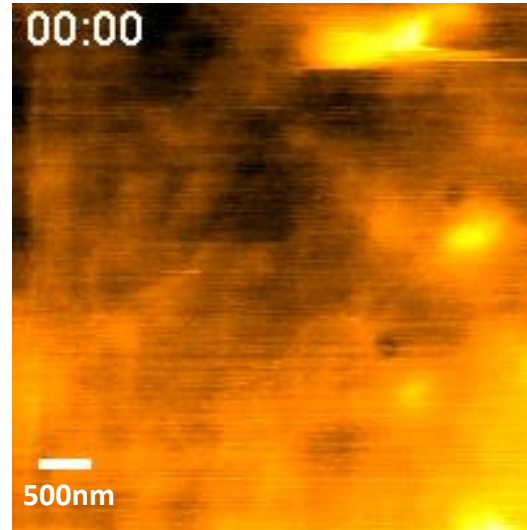
AFM imaging on (and inside !) cells

Cell morphological change
(Dendrite extension)



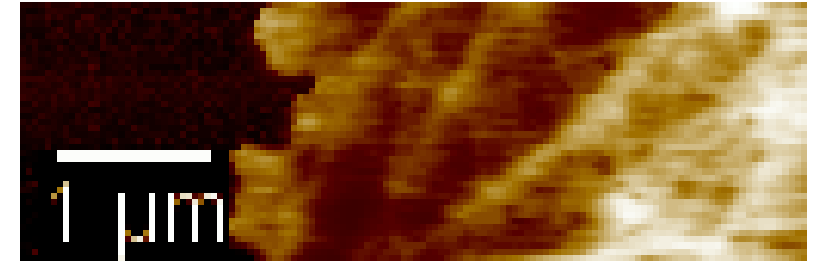
Shibata et al, Sci Rep 2015

Endocytosis/Exocytosis



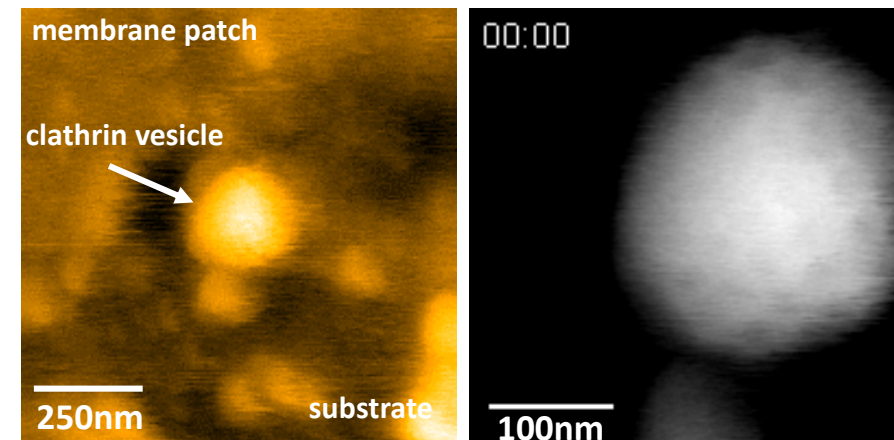
Watanabe et al, Rev Sci Instr 2013

Actin cortex dynamics (16s/image)

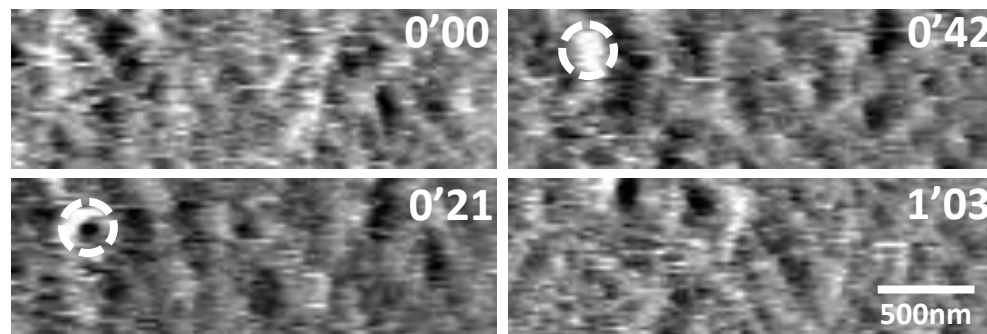


Eghiaian et al, Biophys J 2015

Endocytosis from inside the cell ?



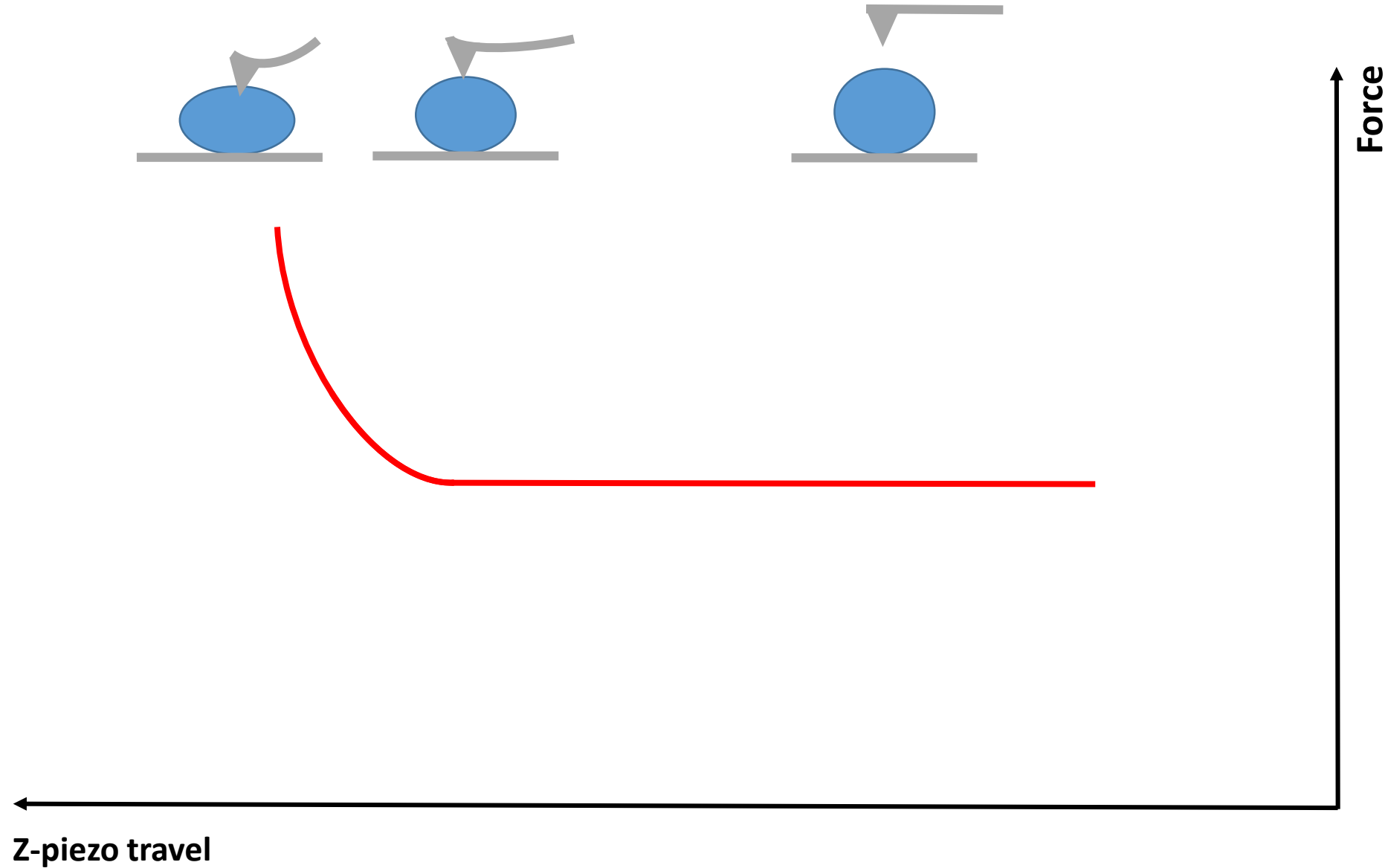
Eghiaian, unpublished



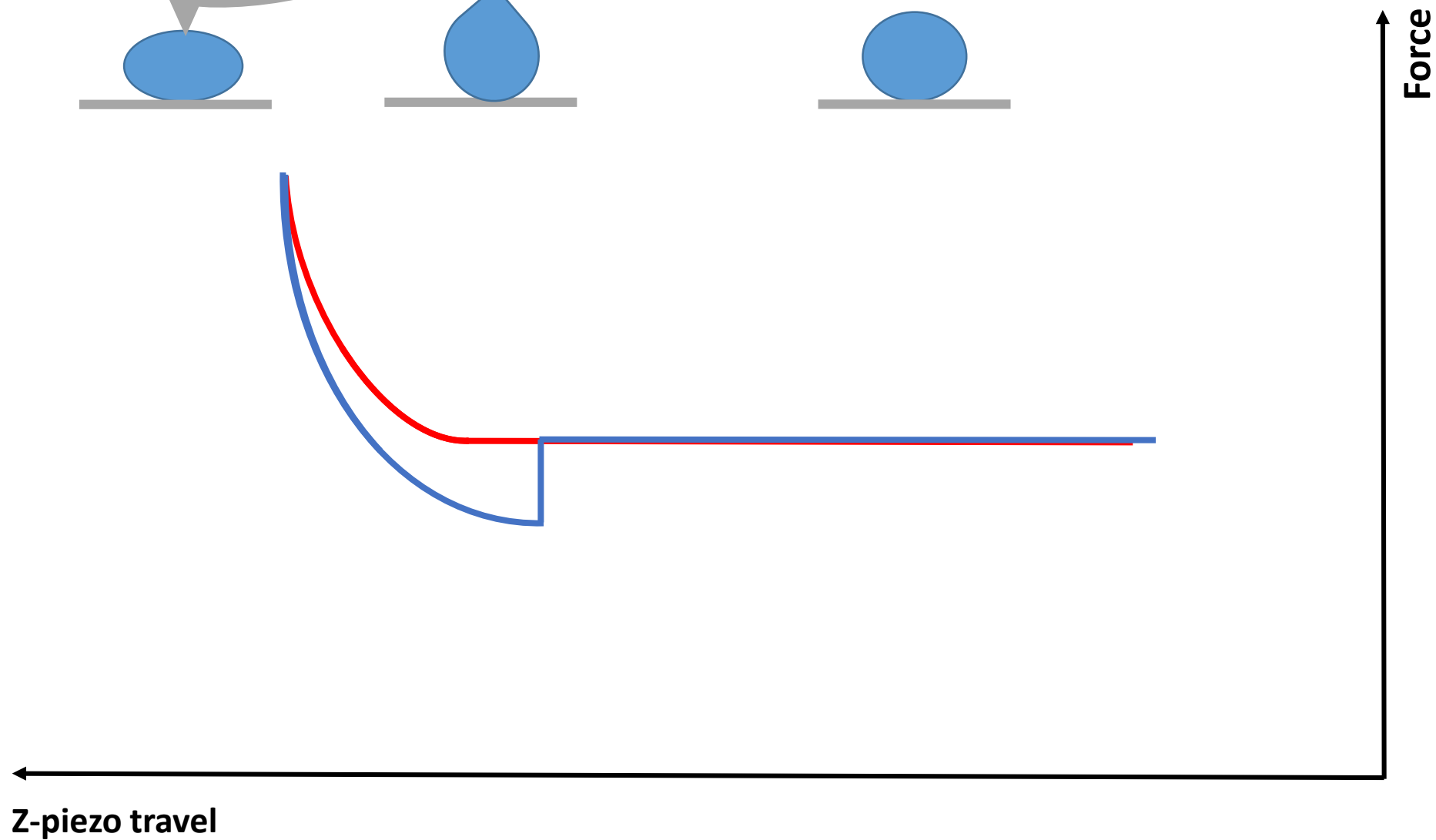
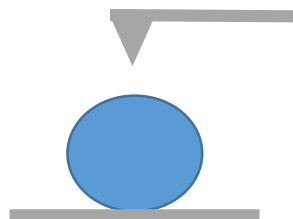
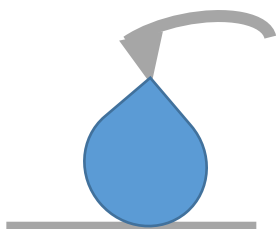
Eghiaian, unpublished

FORCE MEASUREMENTS BY AFM

Force spectroscopy



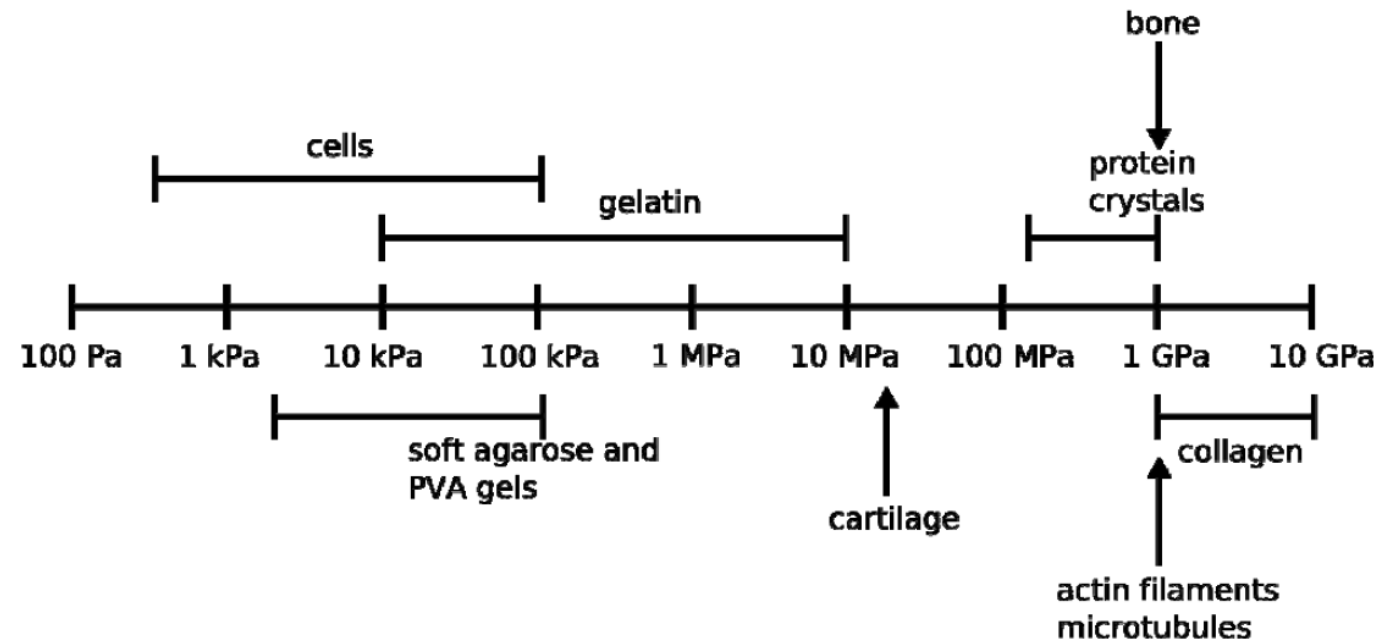
Force spectroscopy



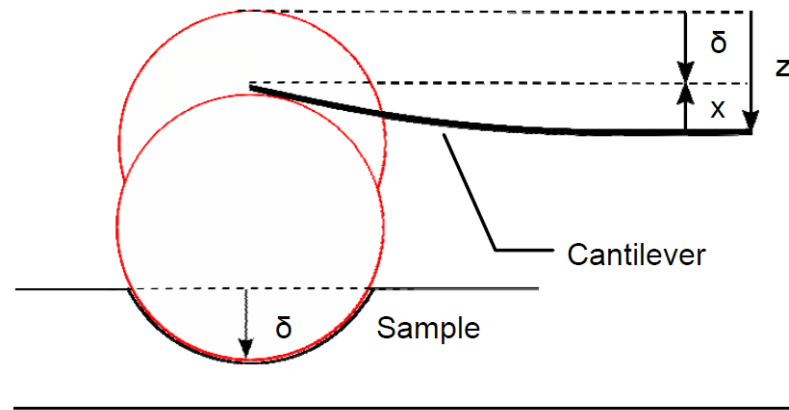
Determining Young's moduli of materials

$$E \equiv \frac{\text{tensile stress}}{\text{extensional strain}} = \frac{\sigma}{\varepsilon} = \frac{F/A_0}{\Delta L/L_0} = \frac{FL_0}{A_0\Delta L}$$

$$F = \left(\frac{EA_0}{L_0} \right) \Delta L = kx$$



Determining Young's moduli of materials: Hertz model



Parabolic

$$F = \frac{4\sqrt{R_c}}{3} \frac{E}{1-\nu^2} \delta^{3/2}$$

R_c = radius of tip curvature



Spherical

$$F = \frac{E}{1-\nu^2} \left[\frac{a^2 + R^2}{2} \ln \frac{R+a}{R-a} - aR \right]$$

$$\delta = \frac{a}{2} \ln \frac{R+a}{R-a}$$

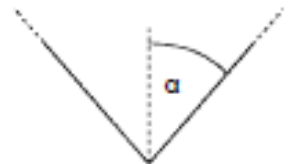
R = radius of the sphere

Conical

$$F = \frac{E}{1-\nu^2} \frac{2 \tan \alpha}{\pi} \delta^2$$

$$a = \frac{2 \tan \alpha}{\pi} \delta$$

α = semi-opening angle of the cone

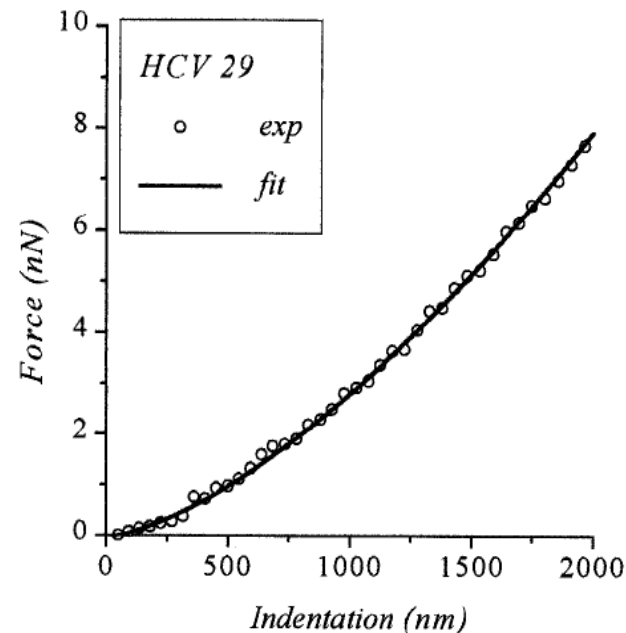
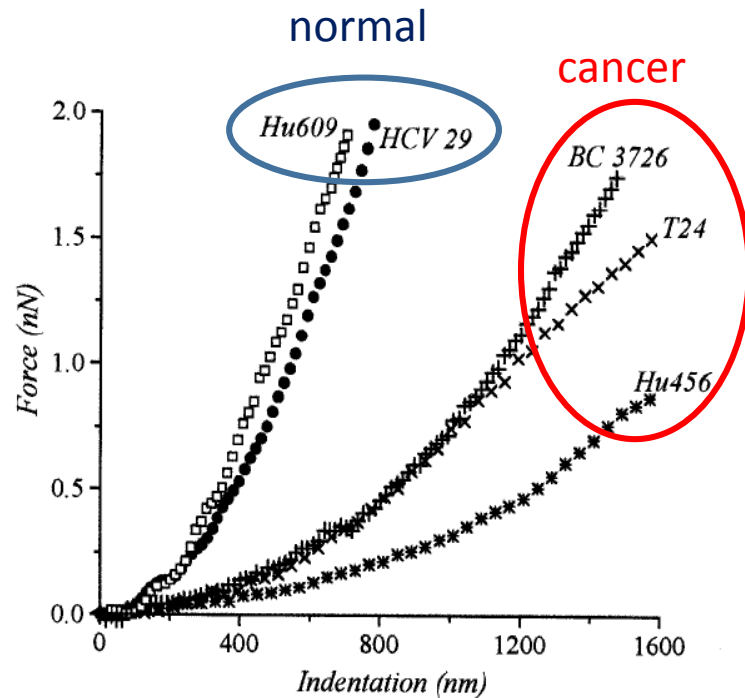


AFM study of cell mechanics

M. Lekka · P. Laidler · D. Gil · J. Lekki · Z. Stachura
A.Z. Hryniewicz

Elasticity of normal and cancerous human bladder cells studied by scanning force microscopy

Eur Biophys J (1999) 28: 312–316



Cell line	Young's modulus (kPa) $\mu_{\text{cell}}=0$	Young's modulus (kPa) $\mu_{\text{cell}}=0.5$	Number of analysed force curves
Hu609	12.9 (± 4.8)	9.7 (± 3.6)	325
HCV 29	10.0 (± 4.6)	7.5 (± 3.6)	121
BC 3726	1.4 (± 0.7)	1.0 (± 0.6)	214
T24	1.0 (± 0.5)	0.8 (± 0.4)	201
Hu456	0.4 (± 0.3)	0.3 (± 0.2)	10

^a The total number of measured cells is about 20 cells for each line

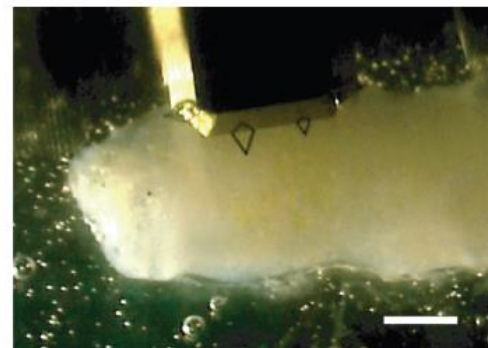
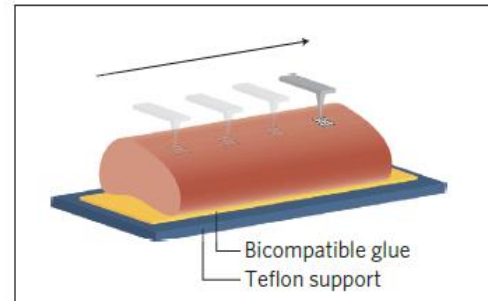
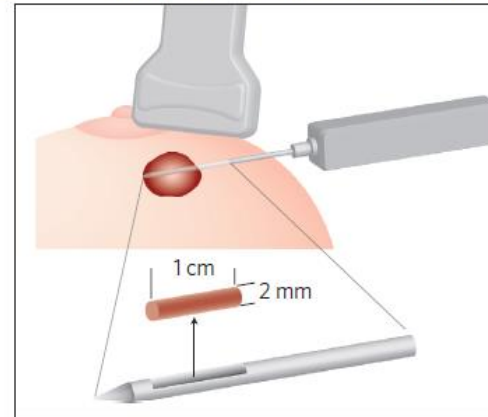
AFM as a Diagnostic tool ???



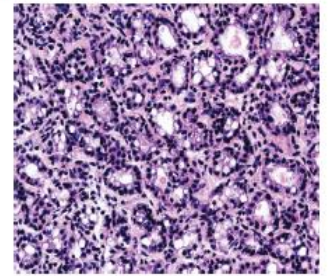
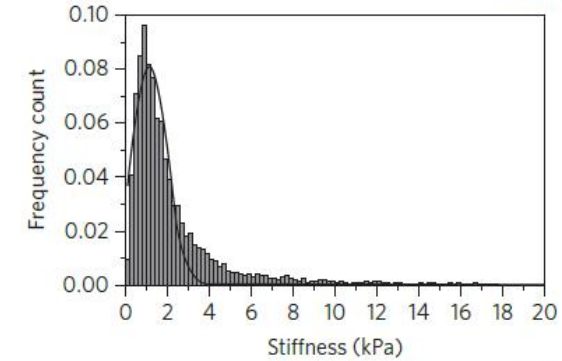
The nanomechanical signature of breast cancer

Marija Plodinec^{1,2}, Marko Loparic^{1,2†}, Christophe A. Monnier^{1†}, Ellen C. Obermann^{3†},
Rosanna Zanetti-Dallenbach^{4†}, Philipp Oertle¹, Janne T. Hyotyla¹, Ueli Aebi², Mohamed Bentires-Alj⁵,
Roderick Y. H. Lim^{1*} and Cora-Ann Schoenenberger²

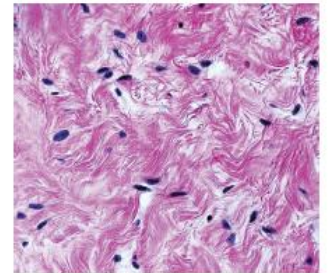
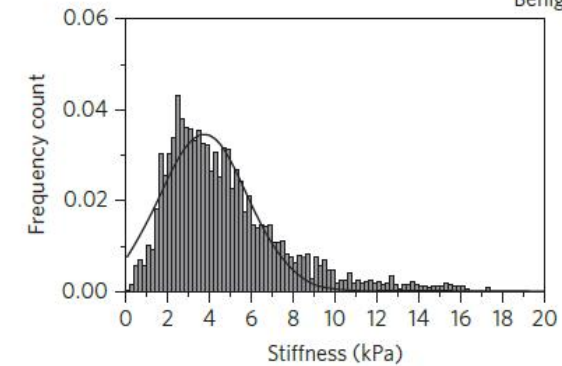
Testing human breast biopsies by IT-AFM



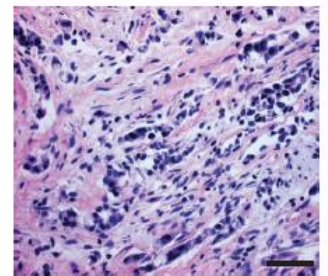
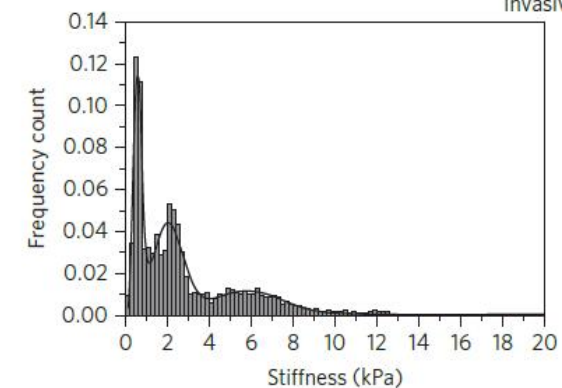
Normal tissue



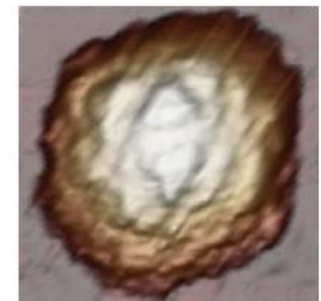
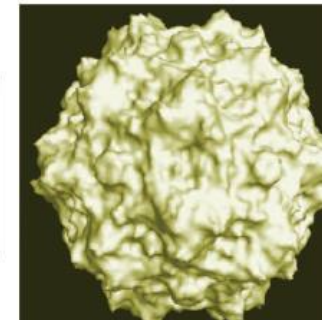
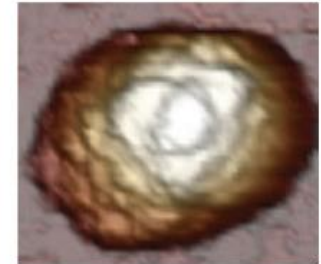
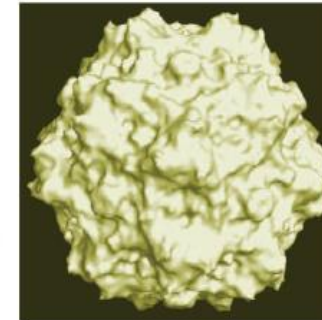
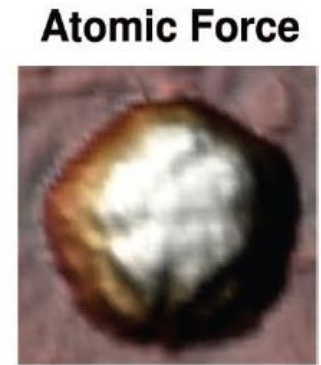
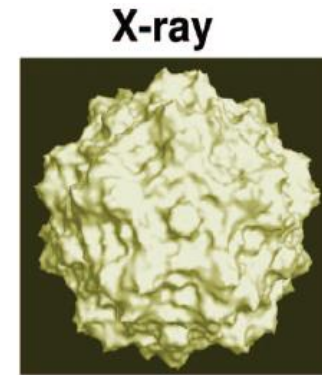
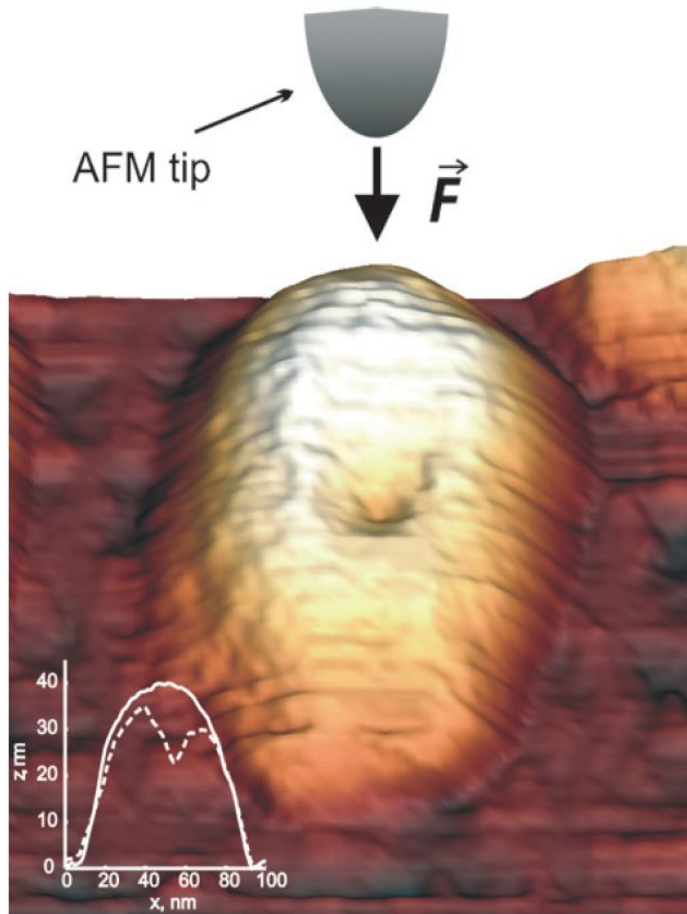
Benign lesion



Invasive cancer

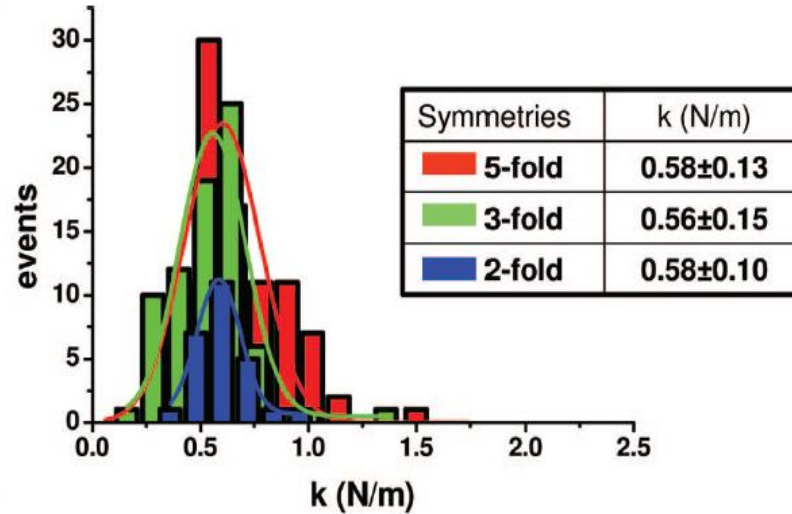
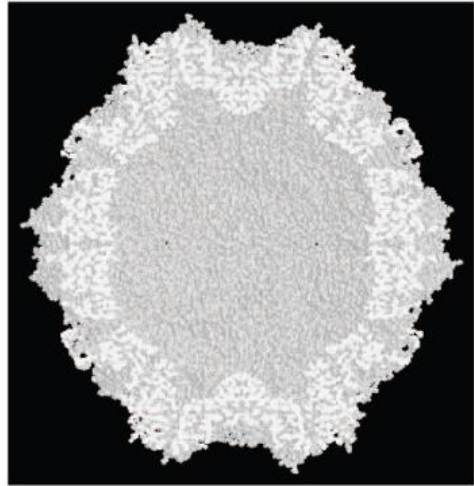


AFM studies of virus mechanics

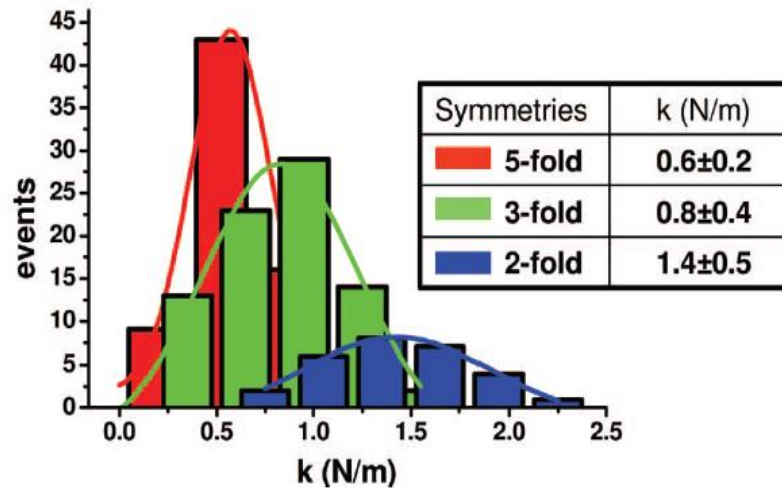
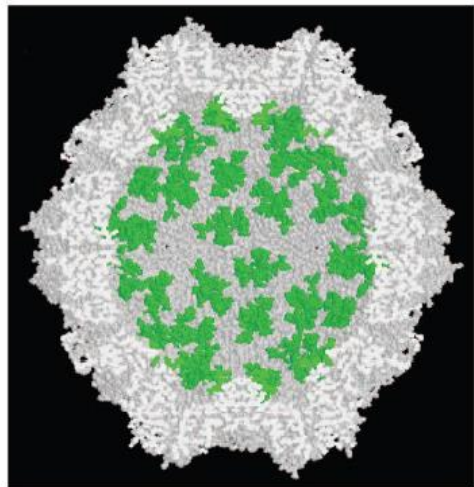


AFM studies of virus mechanics (DNA reinforcement)

Empty capsid

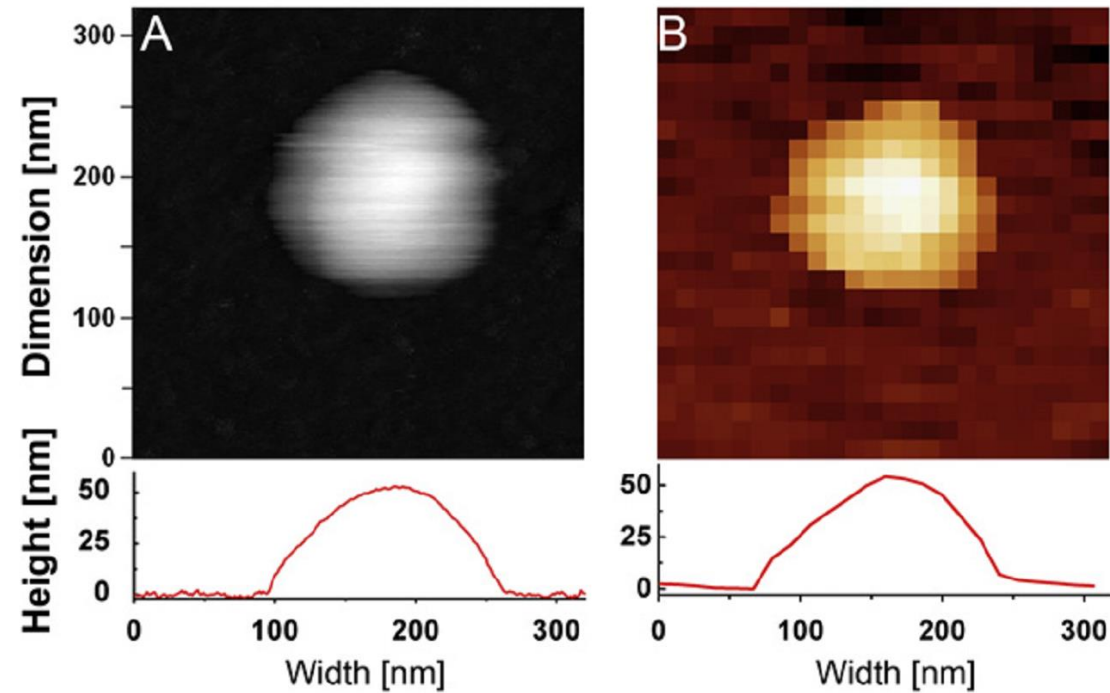
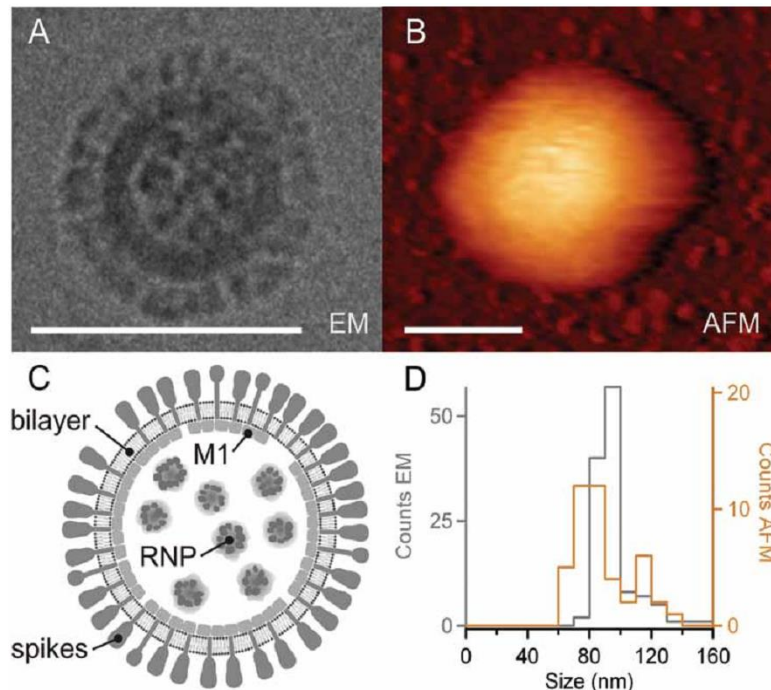
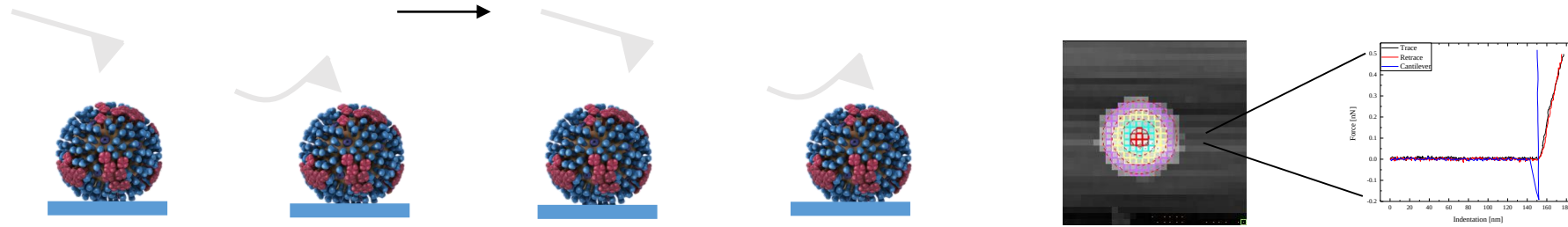


Virion (with DNA)

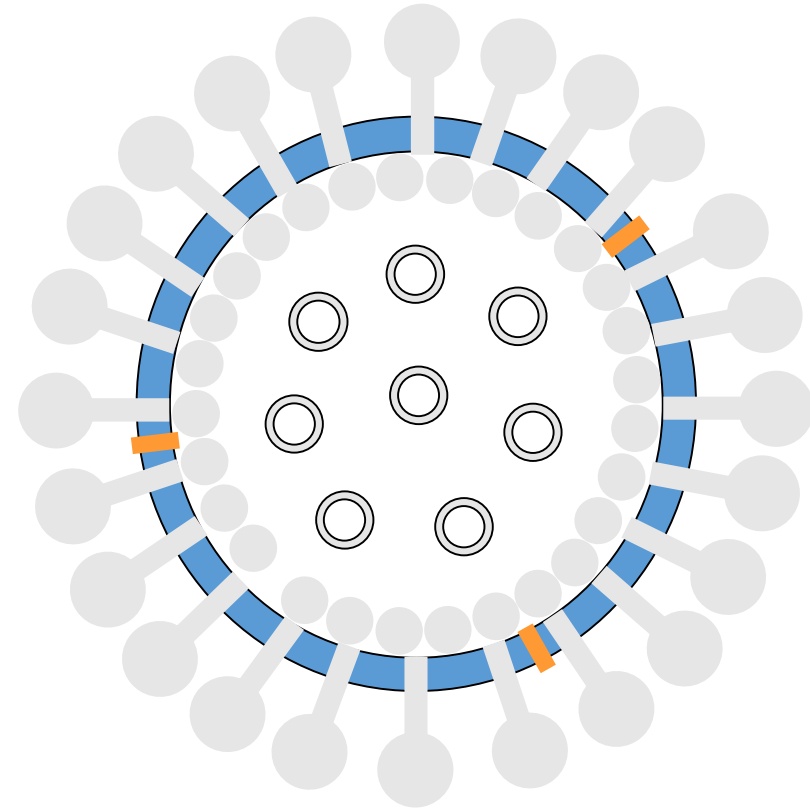
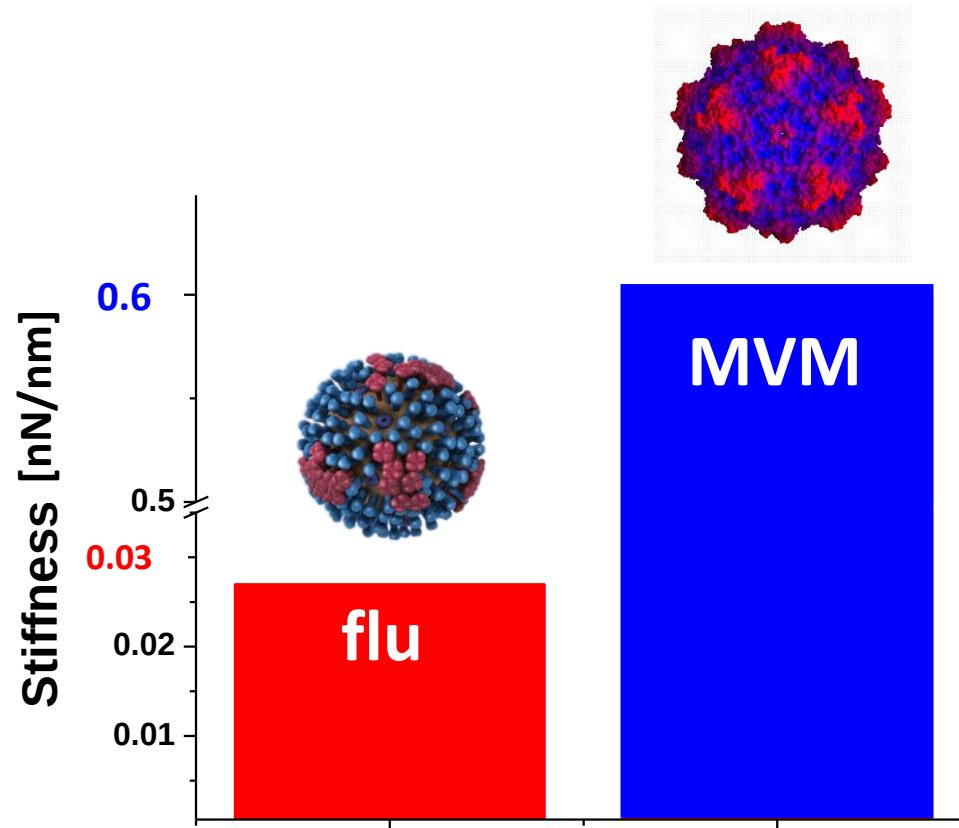


Force mapping of flu virions

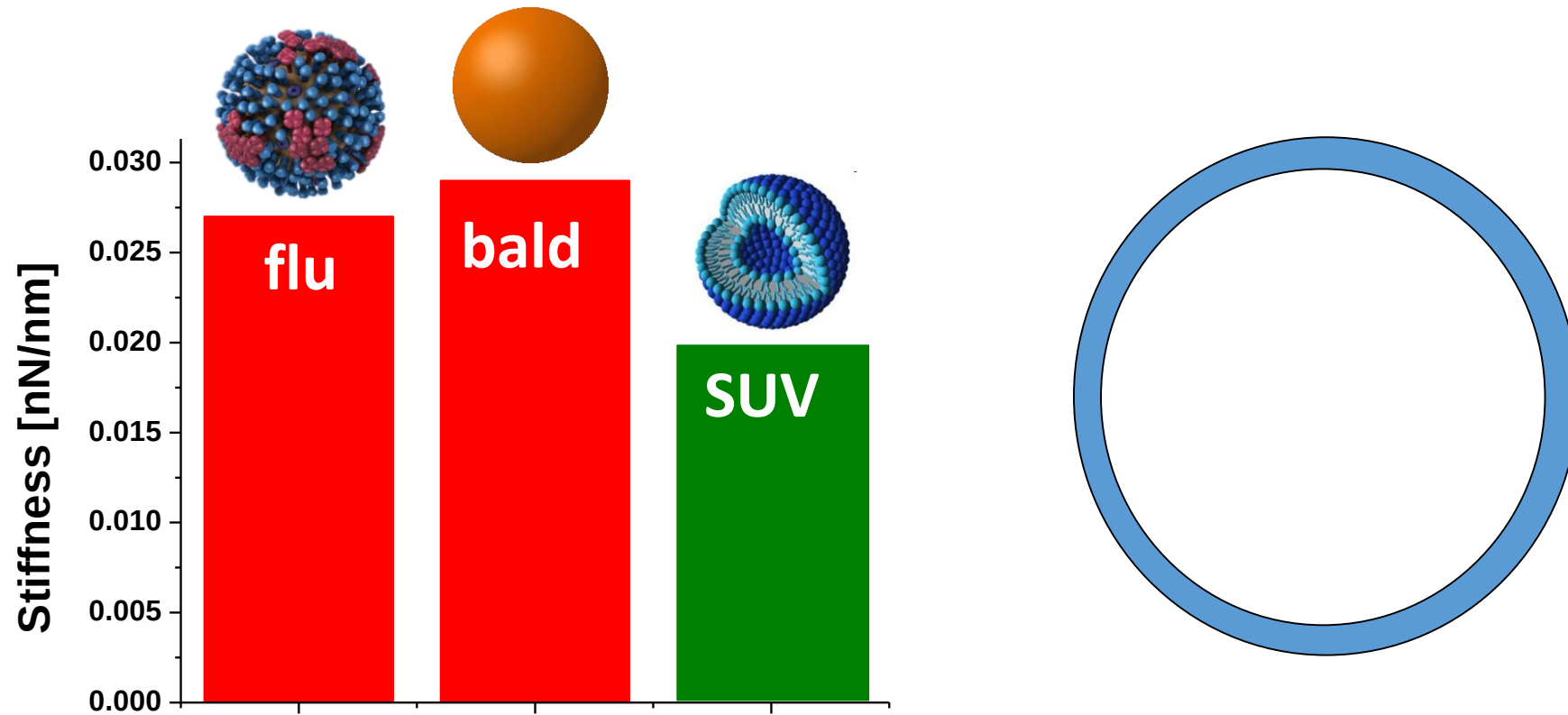
Force mapping



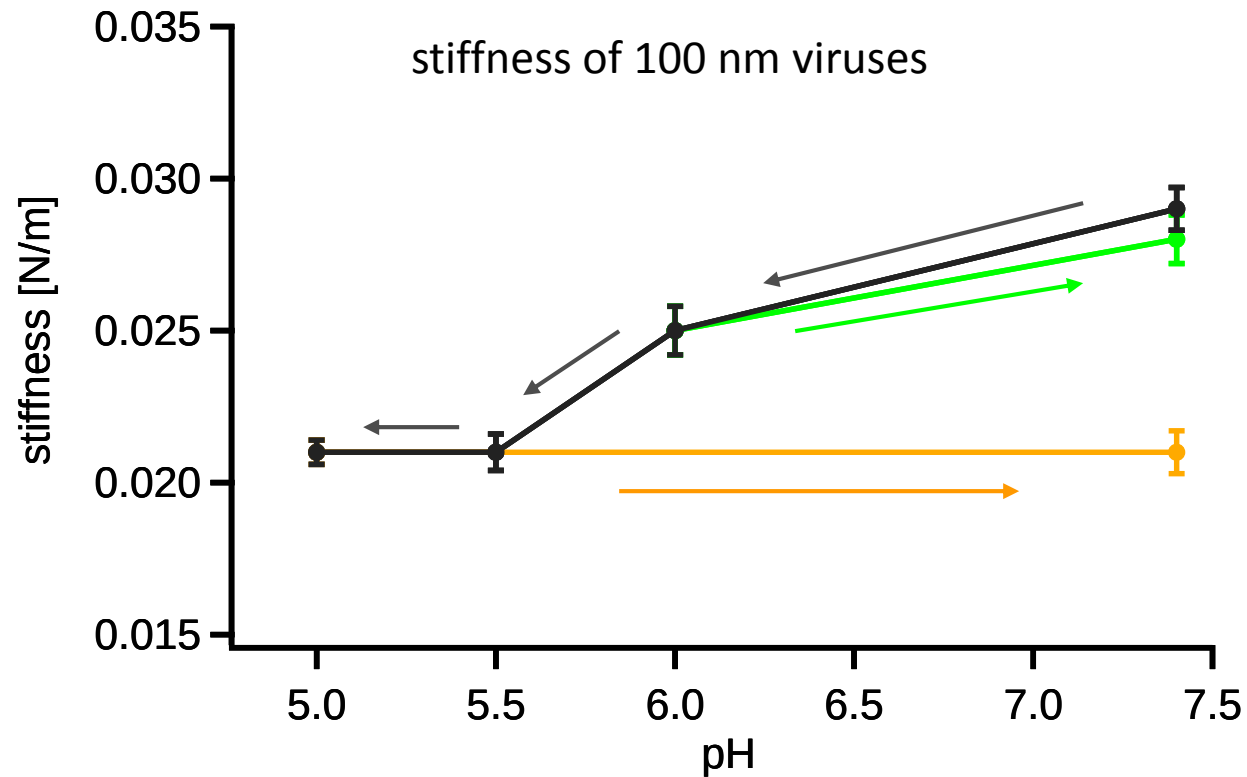
AFM studies of influenza virus mechanics



Lipids define mechanical response



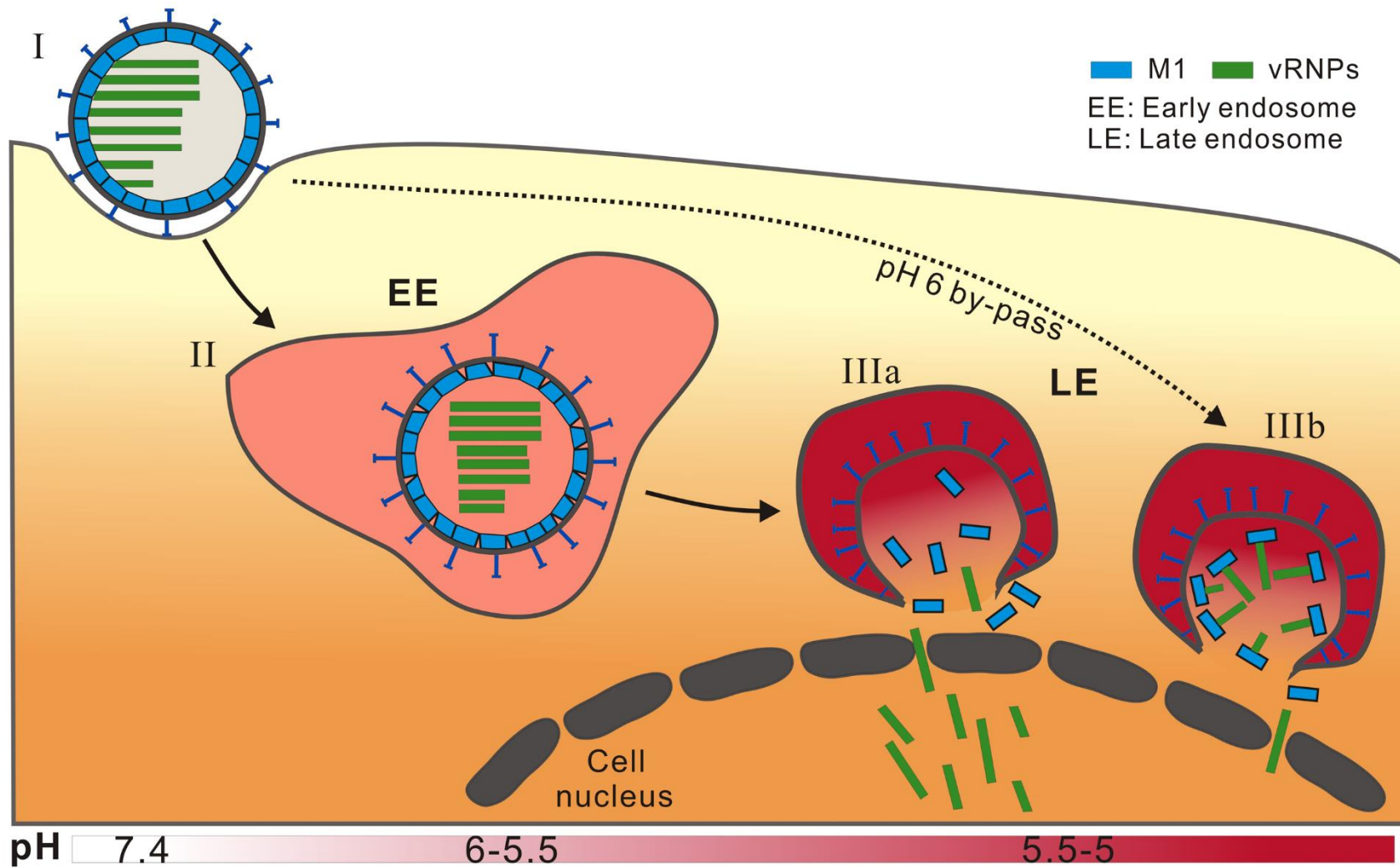
pH-dependent viral stiffness



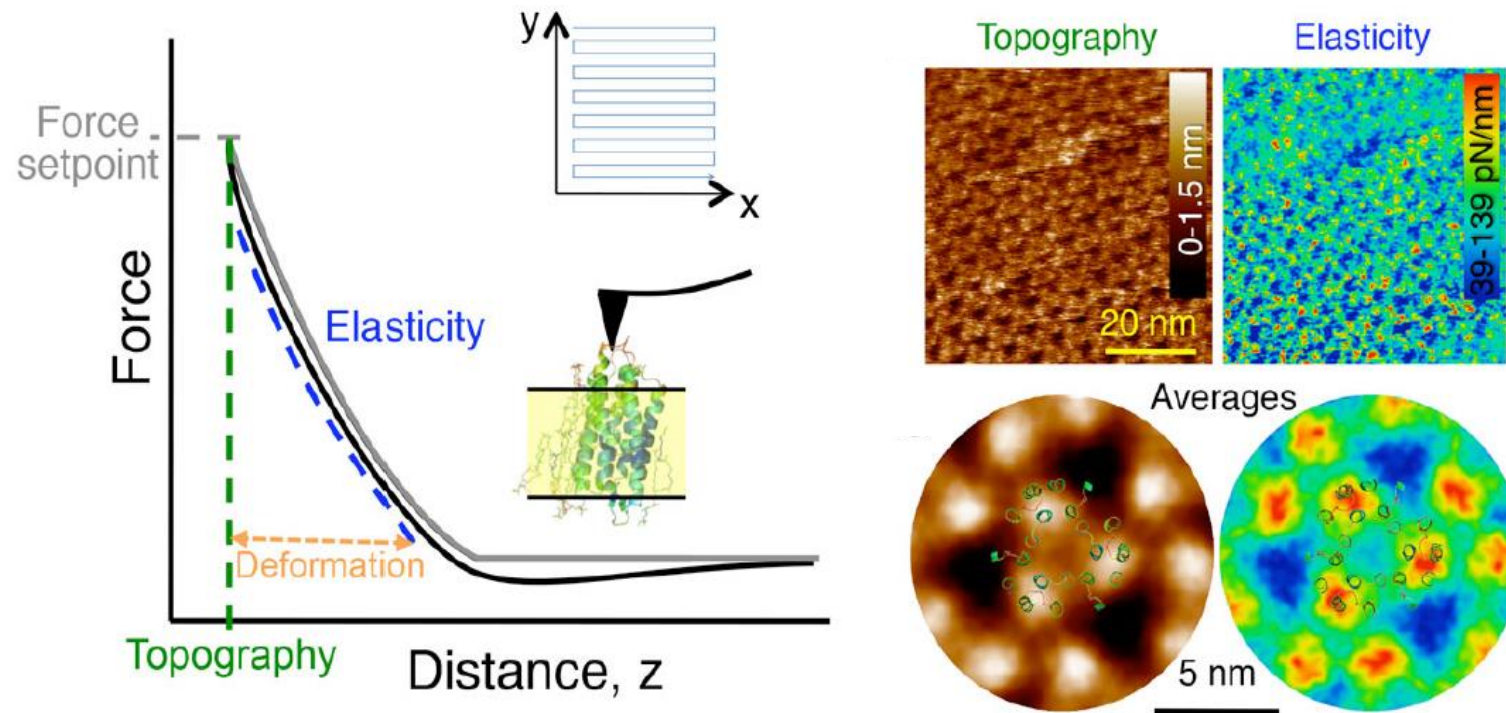
protein capsid disassembly reduced stiffness only by 30 %

irreversible below pH 6

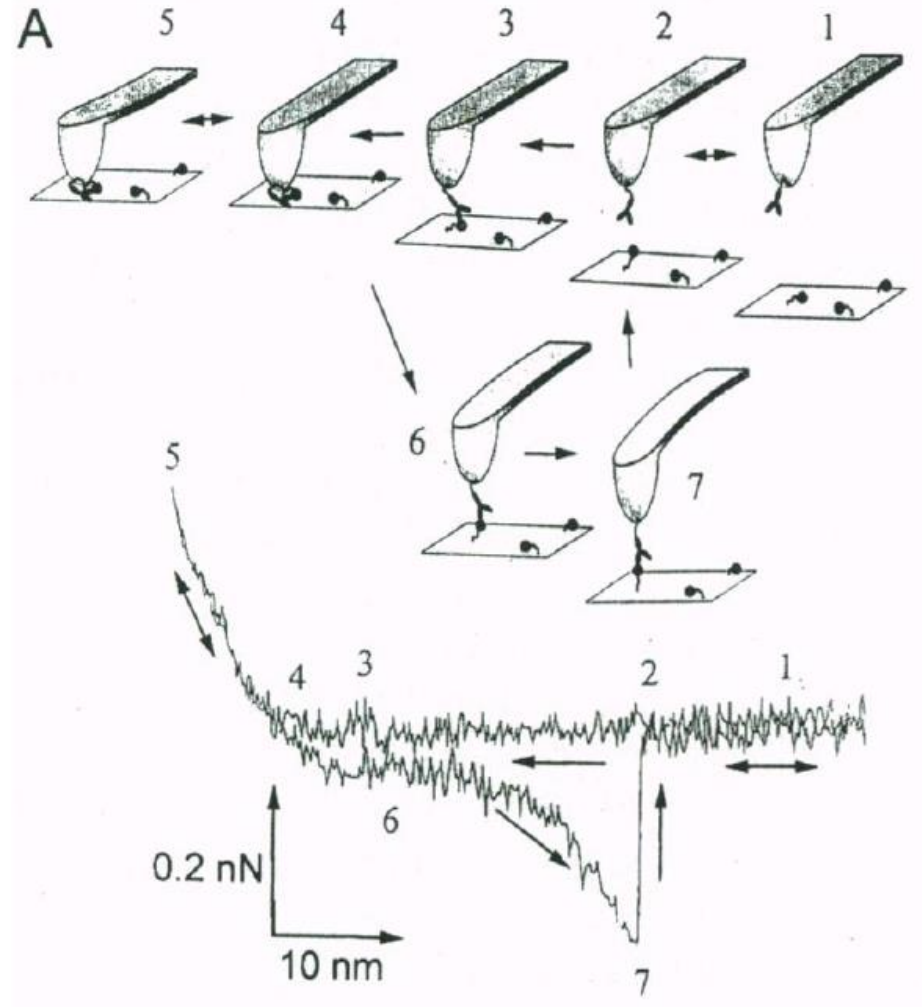
what determines the remaining 70 % of stiffness?



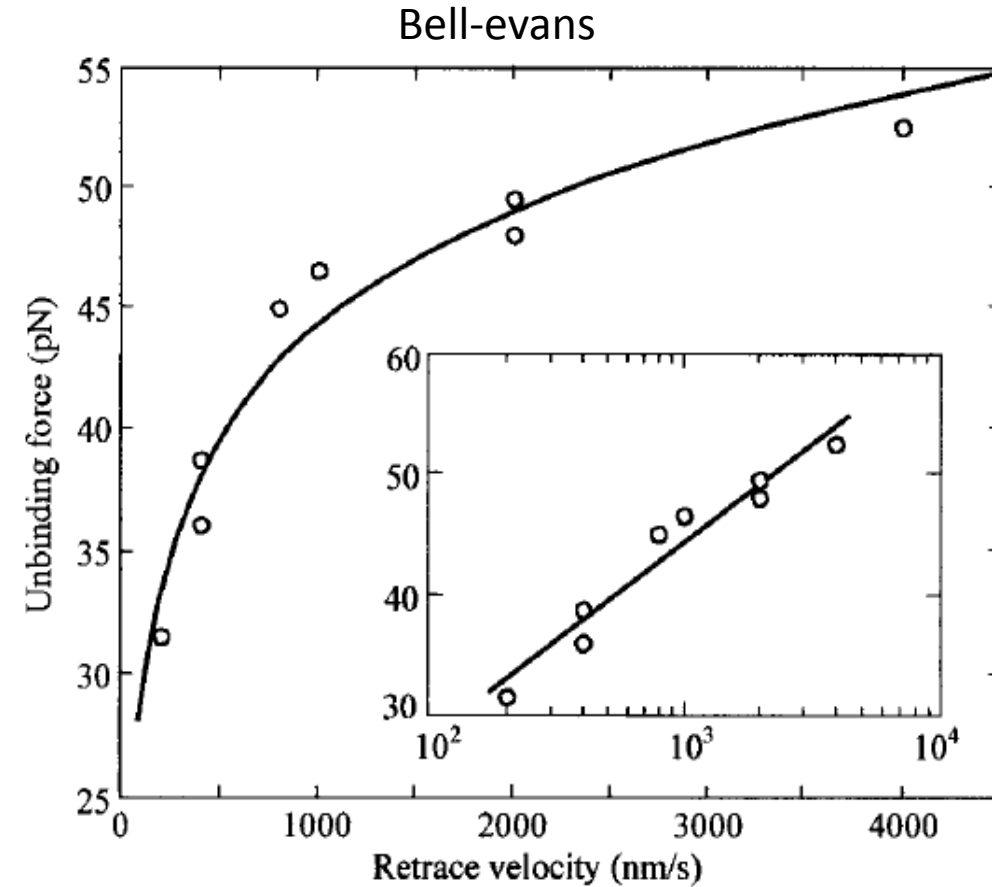
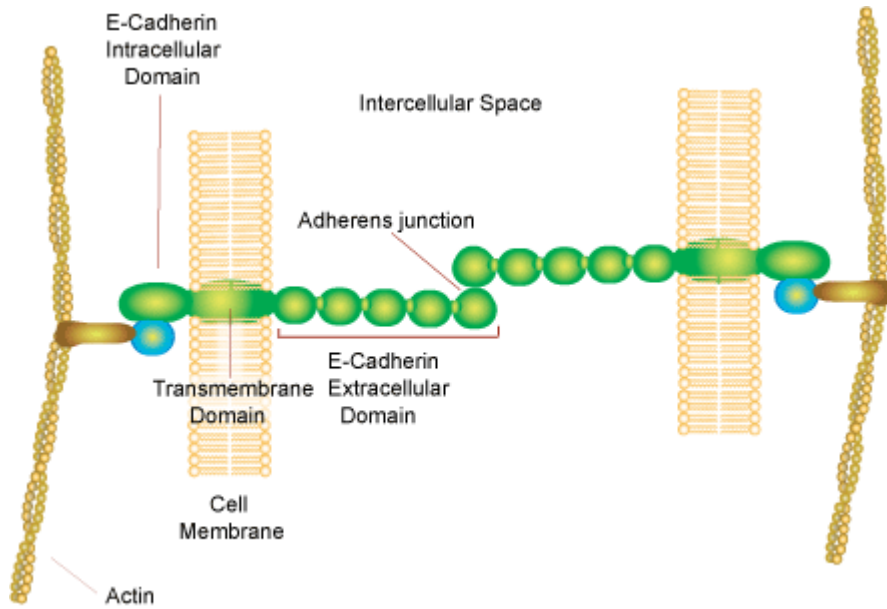
Mechanical measurements on single proteins



Single pair interaction



Single pair interaction



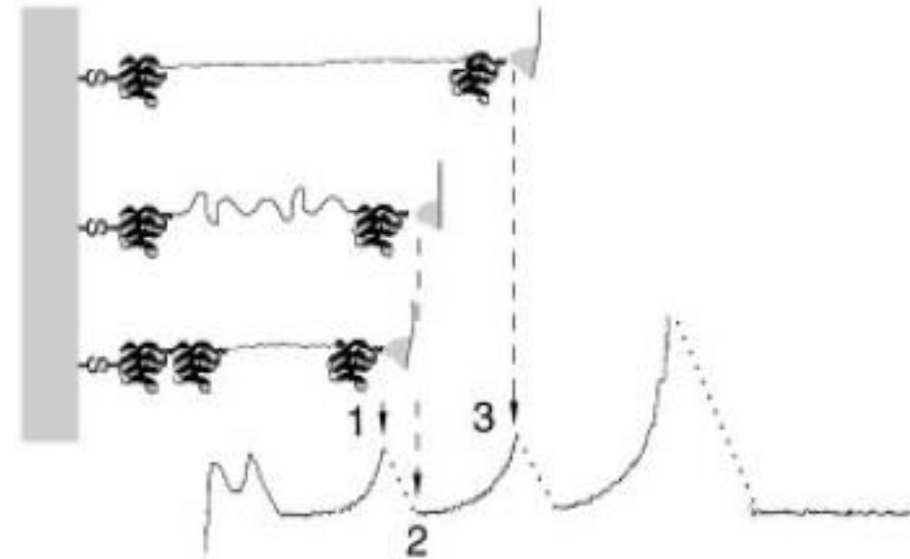
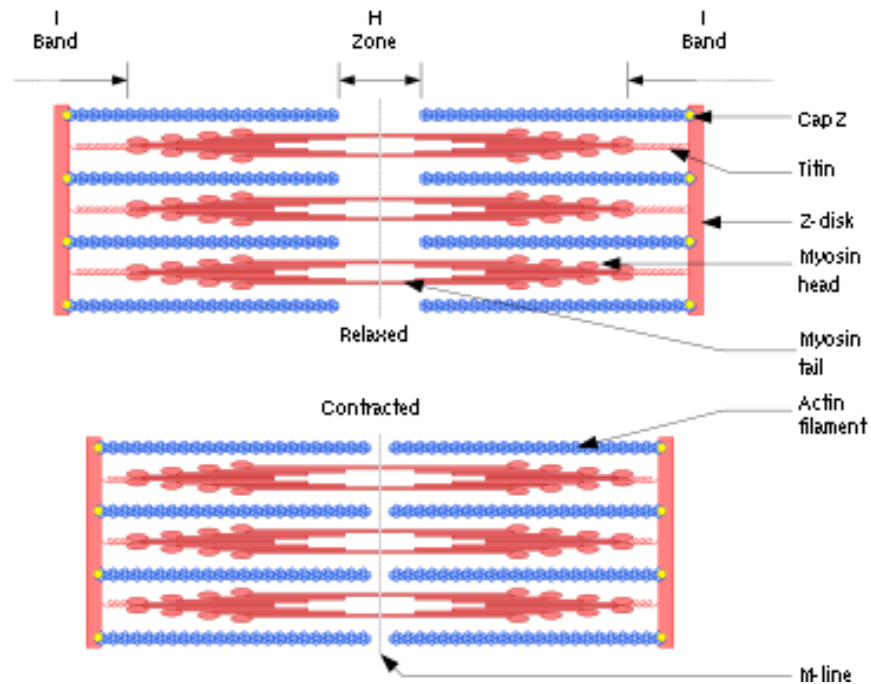
$$\tau(f_u) = \tau_0 \exp(-l_r f_u / k_B T)$$

Baumgartner et al, PNAS 2000

Bell, Science 1977

Evans & Ritchie, BJ 1997

Mechanical characterization of muscle Titin



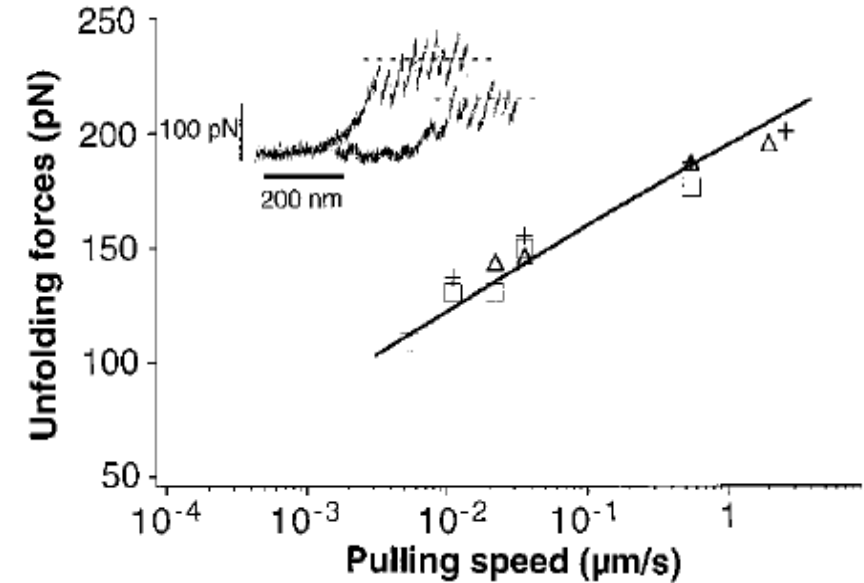
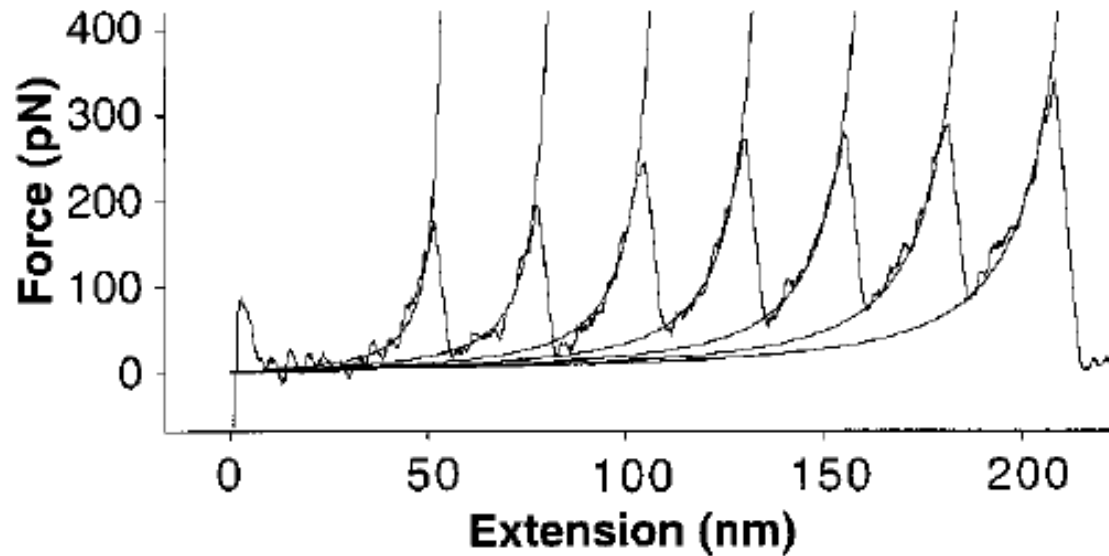
<http://en.wikipedia.org/wiki/Titin>

Single molecule unfolding

Reversible Unfolding of Individual Titin Immunoglobulin Domains by AFM

Matthias Rief *et al.*

Science **276**, 1109 (1997);

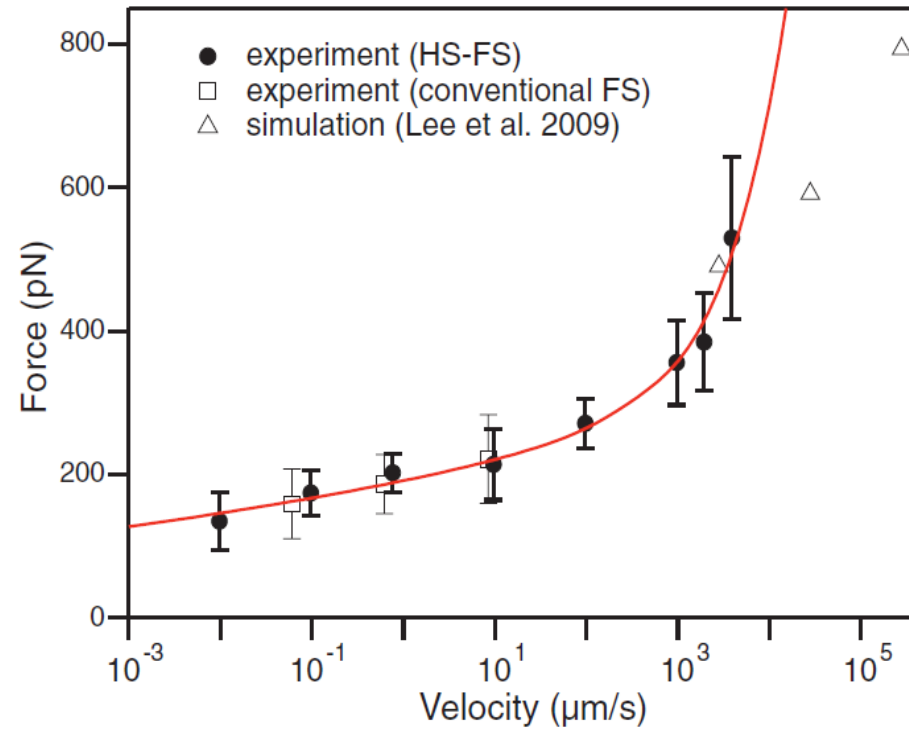
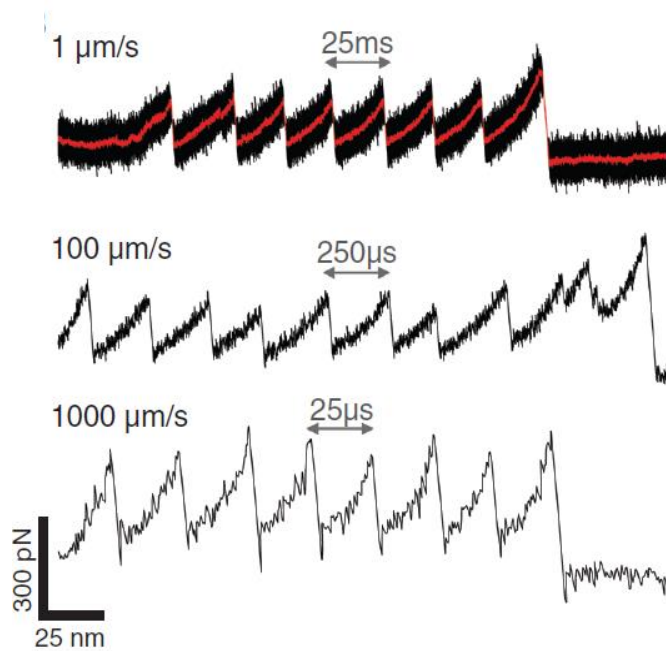


$$F(x) = (kT/b) [0.25(1 - x/L)^{-2} - 0.25 + x/L]$$

L: contour length of the polymer chain

$$\alpha = \alpha_0 \exp(F\Delta x/kT)$$

High-speed single molecule unfolding





Credits

U1006 Marseille

Simon SCHEURING

Anna Francesca Rigato

Zohreh Sedaghat

Humboldt Uni Berlin

Andreas HERRMANN

Christian SIEBEN

Kai LUDWIG

Chris Höfer

Salvatore Chianta

Universität Göttingen

Iwan SCHAAP

Sai LI

Columbia (NYC, J Frank's lab)

Amédée des Georges

NIMR Mill Hill (UK)

Claudia VEIGEL

CNRS Gif-sur-Yvette/INRA Jouy en Josas

Marcel KNOSSOW (thesis director)

Human REZAEI (unofficial co-director)

Institut Pourquoi

Stéphanie LESCEU

Recommended readings

- Mechanics of Motor Proteins and the Cytoskeleton (J Howard)
- Ando et al, Chem Rev 2014, Filming biomolecular processes by high-speed atomic force microscopy
- Mechanics of the Cell (David Boal)

