

Designing Platinum features to mimic protein channels

Stefano Pagliara

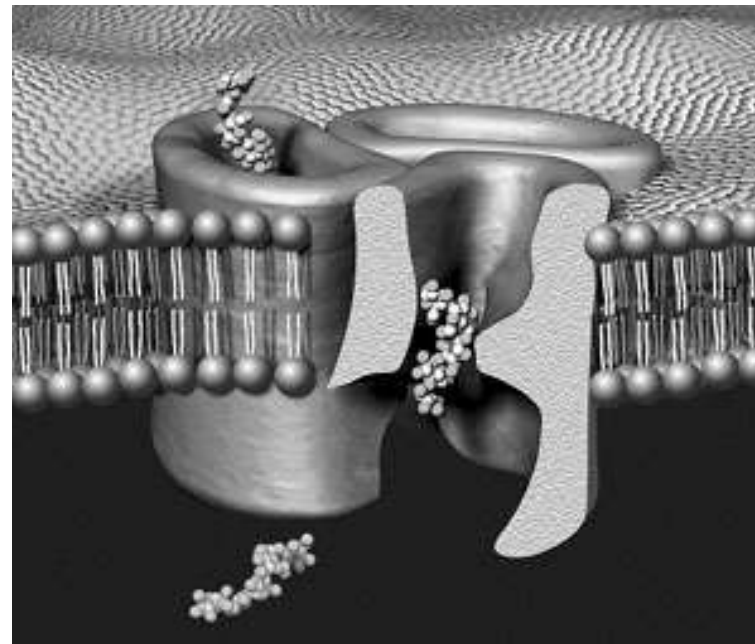
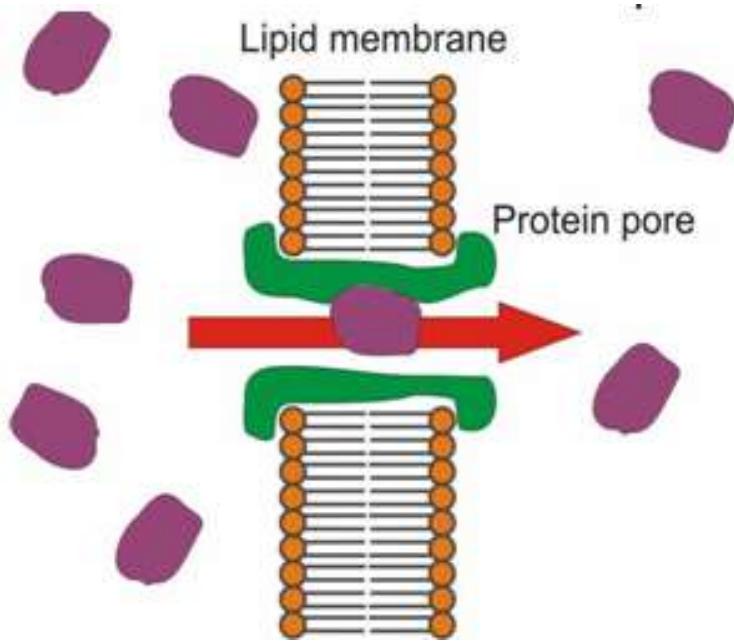
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Outline

- Channel-facilitated membrane transport
- Lab-on-a-chip fabrication technologies
- Chip fabrication exploiting Platinum features
- Single particle control
- Conclusions and outlook

Channel-facilitated membrane transport

Transport of solutes across living system membranes occurs through membrane protein forming water-filled channels



Maltose
translocating
the
maltoporin
channel in *E.*
Coli

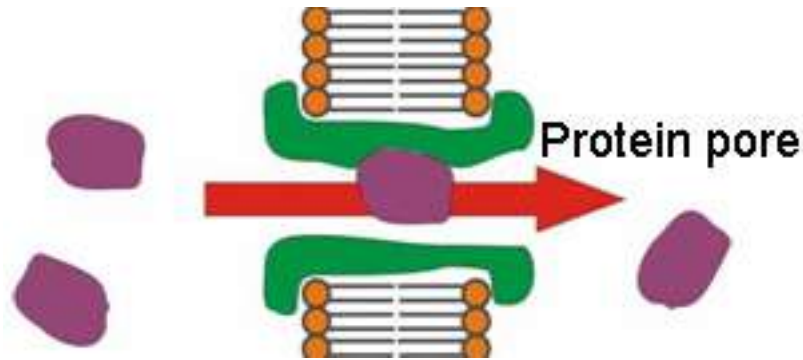
Understanding the
transport mechanism



more efficient drugs

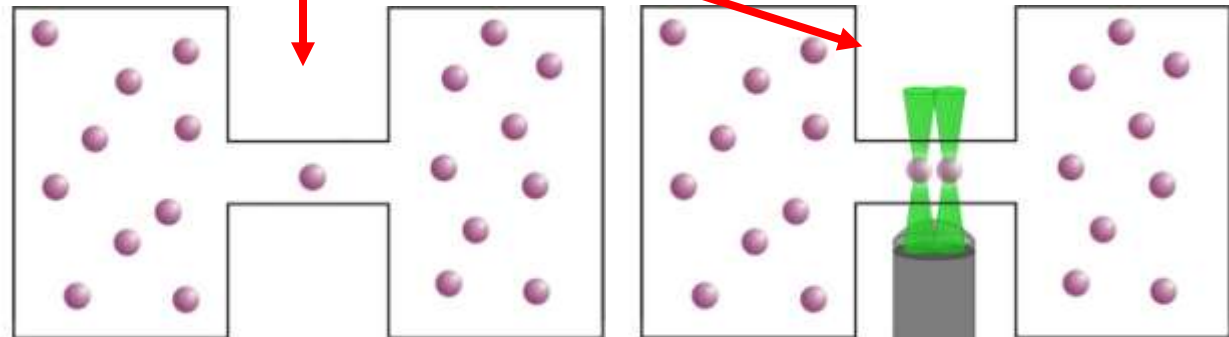
Model system

Nanoscale
inaccessible
natural system



Microscale
controllable
model system

Colloids

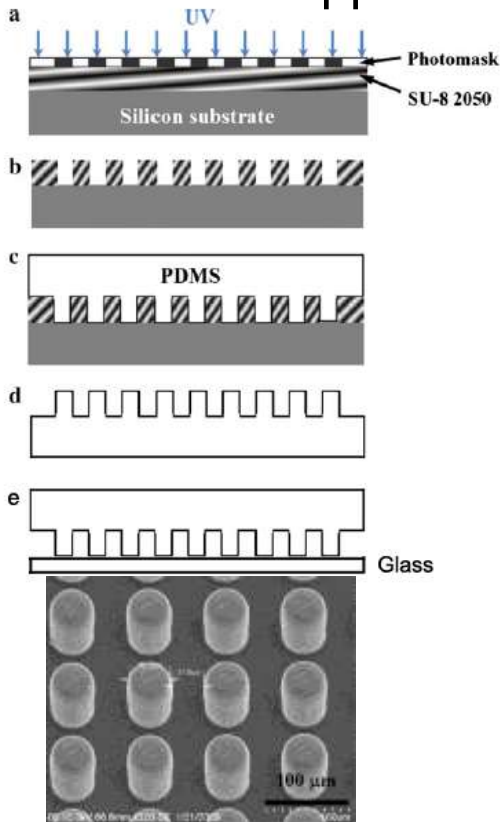


Microfluidics

Holographic optical tweezers

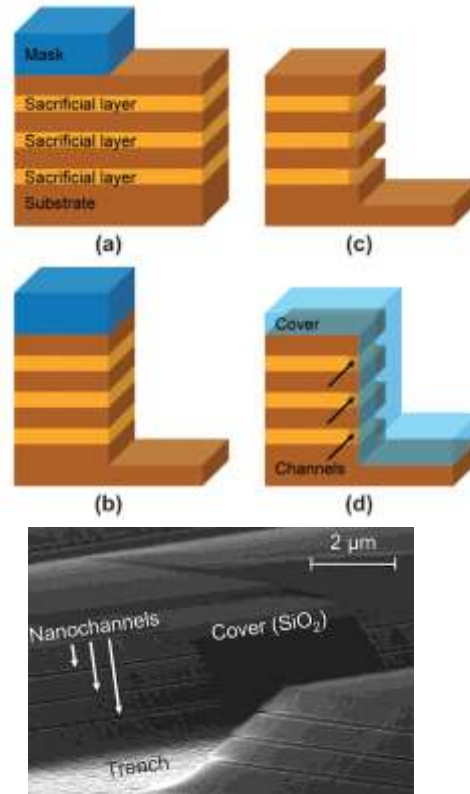
Lab-on-a-chip fabrication technologies

Conventional approach



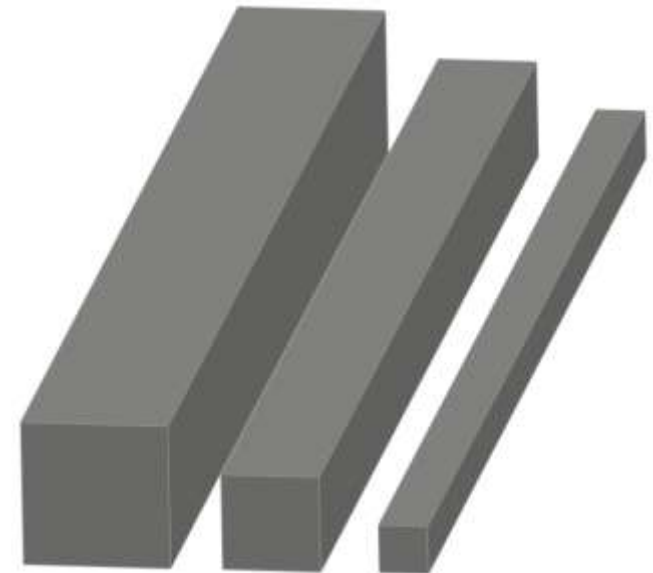
Microscale planar features

Alternative approach



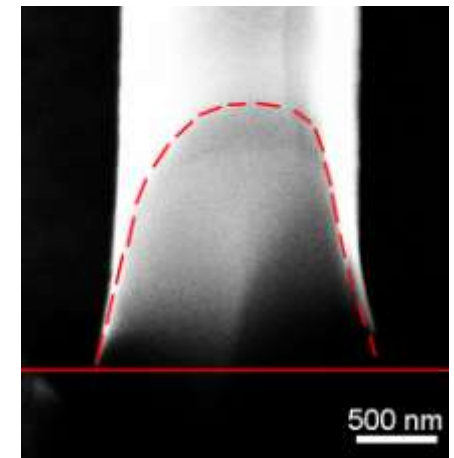
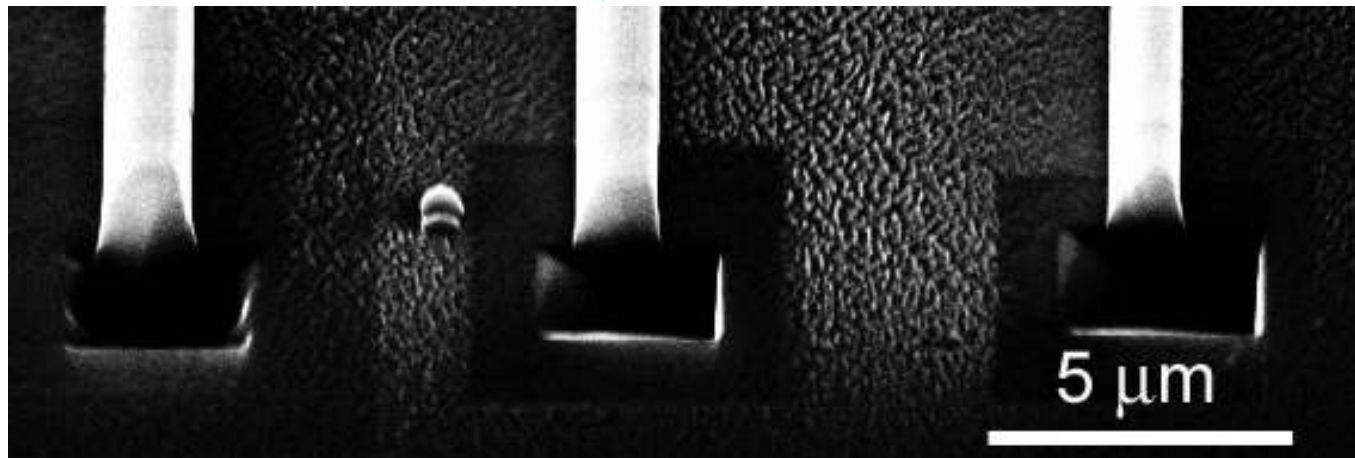
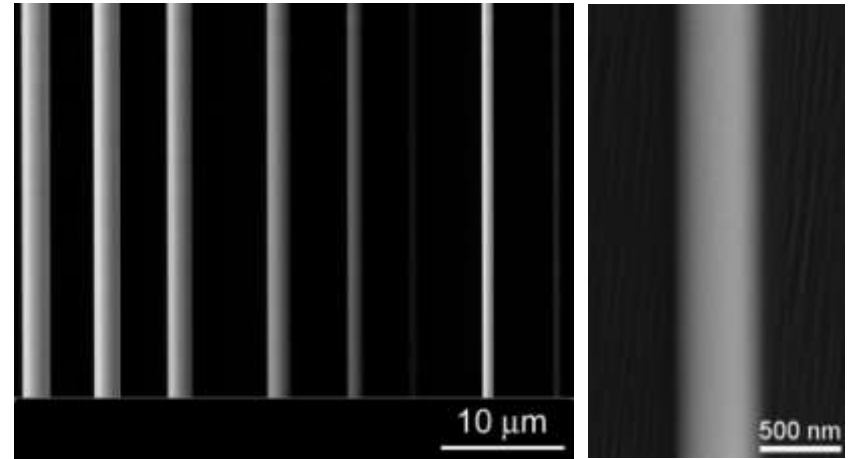
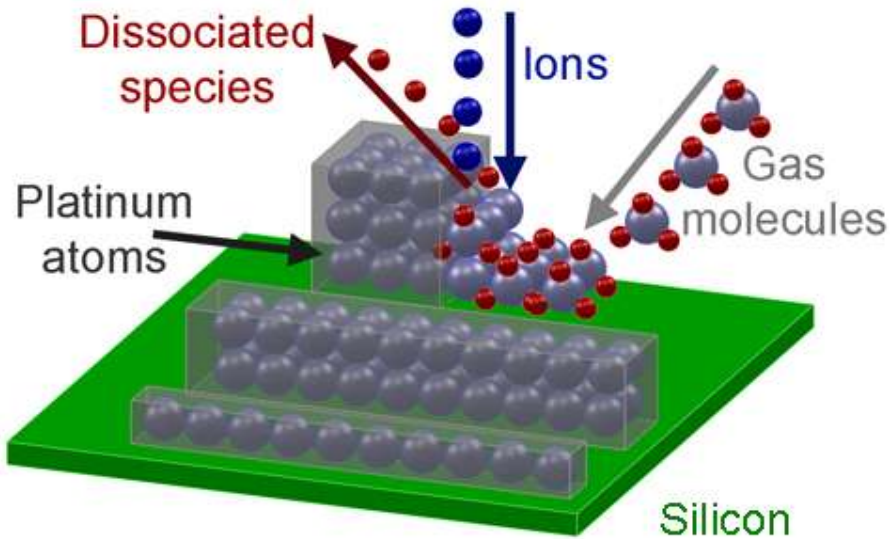
Nanoscale planar features

Challenge: 3D control

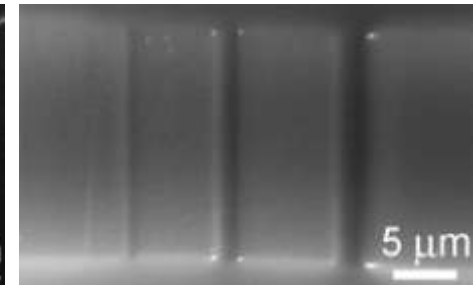
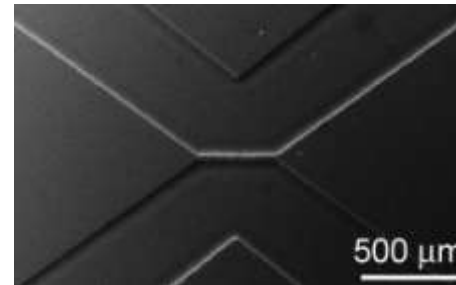
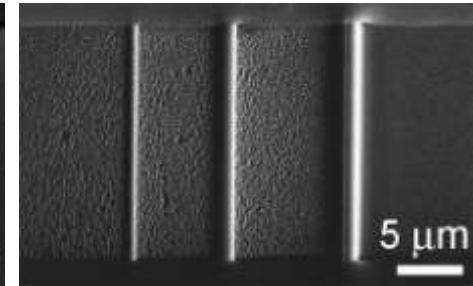
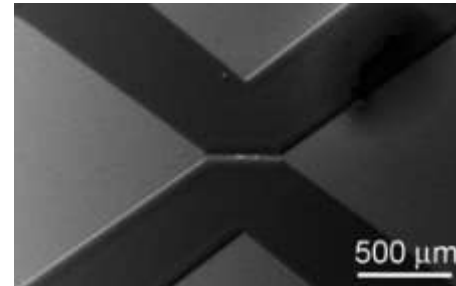
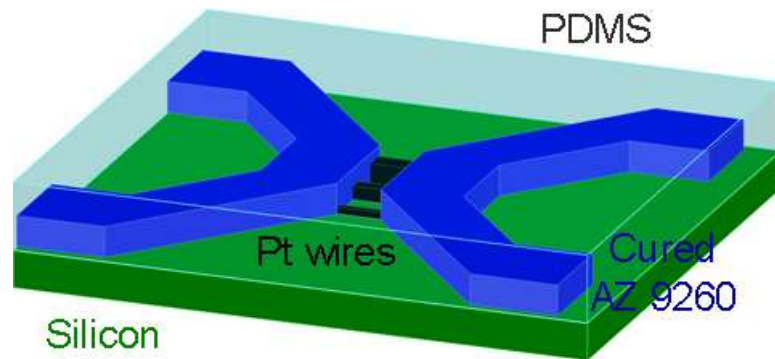
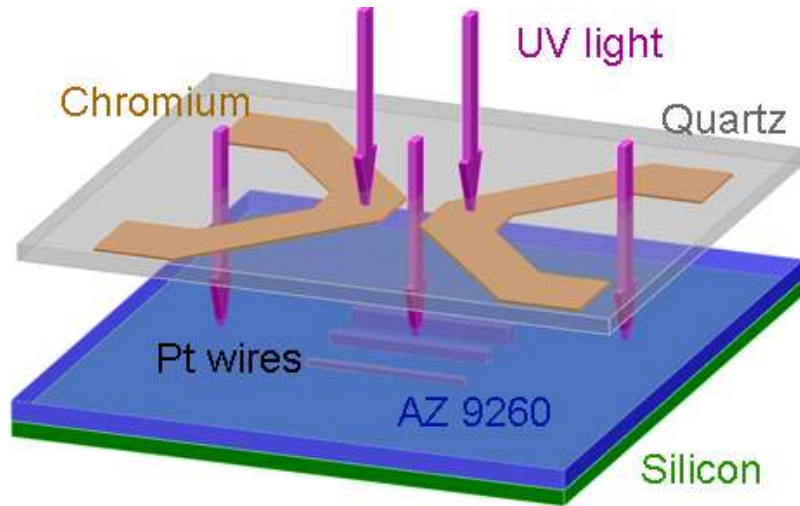


Nanoscale features
with variable thickness

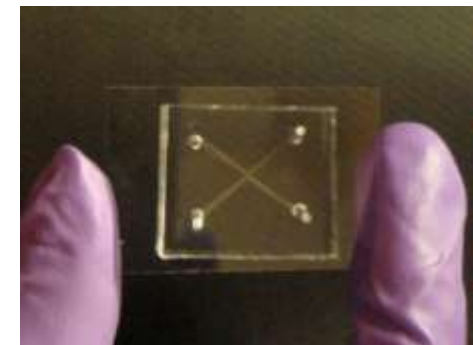
Chip fabrication exploiting focused ion beam



Chip fabrication. Photolithography and replica molding

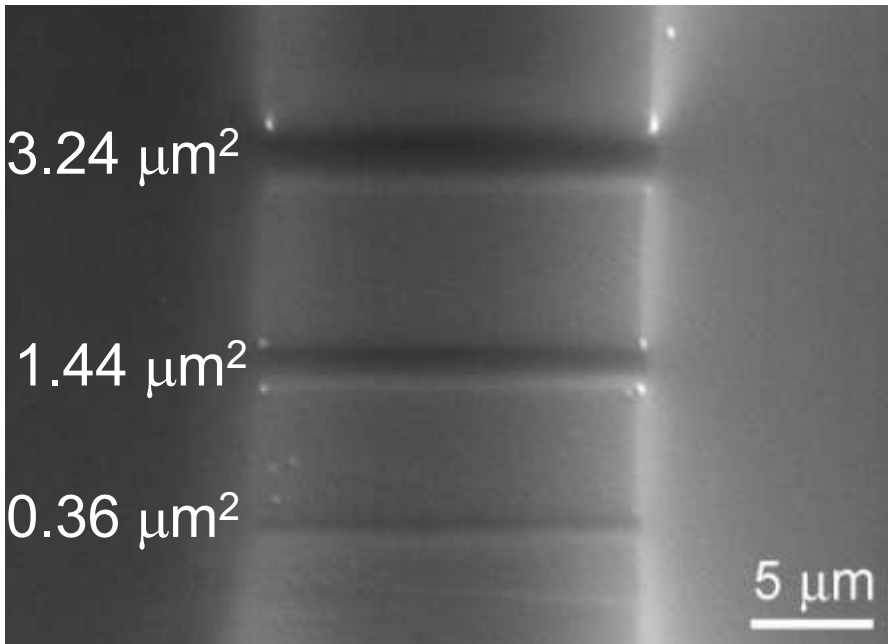


Final device:
patterned PDMS
chip bonded to a
glass slide

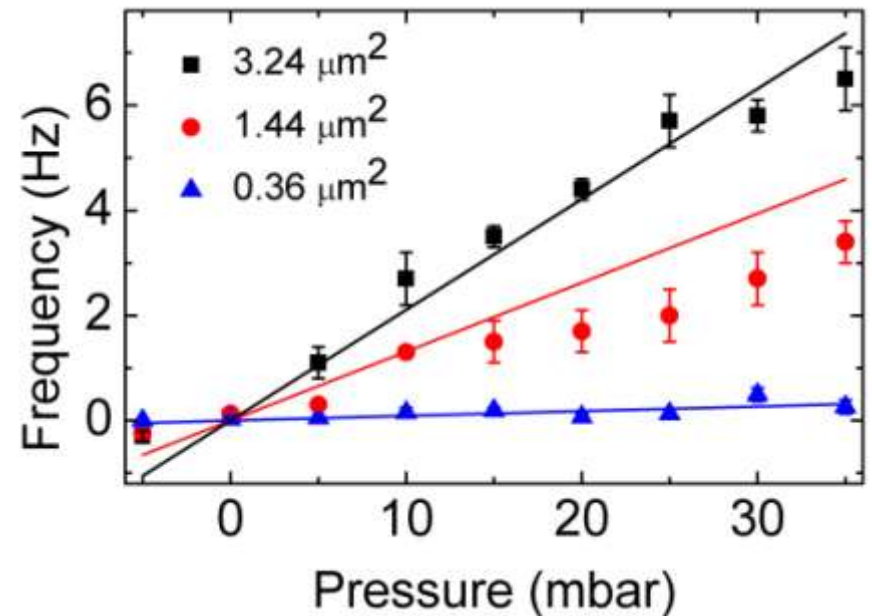


Particle control by pressure gradient

Flow control of 300 nm particles through an array of microchannels with different cross section via computerized pressure-based flow control system



Poiseuille flow $f \propto \Delta P$

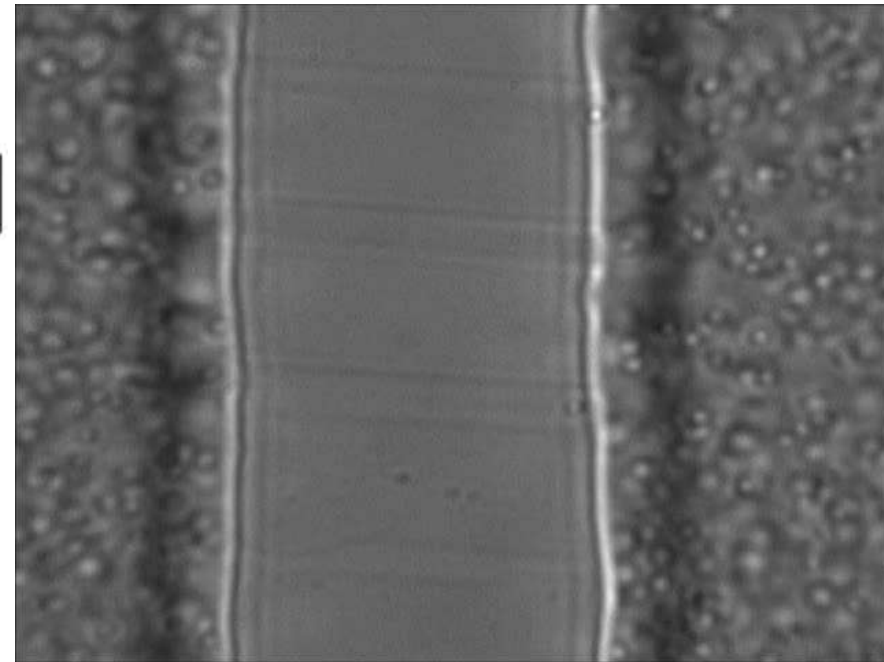
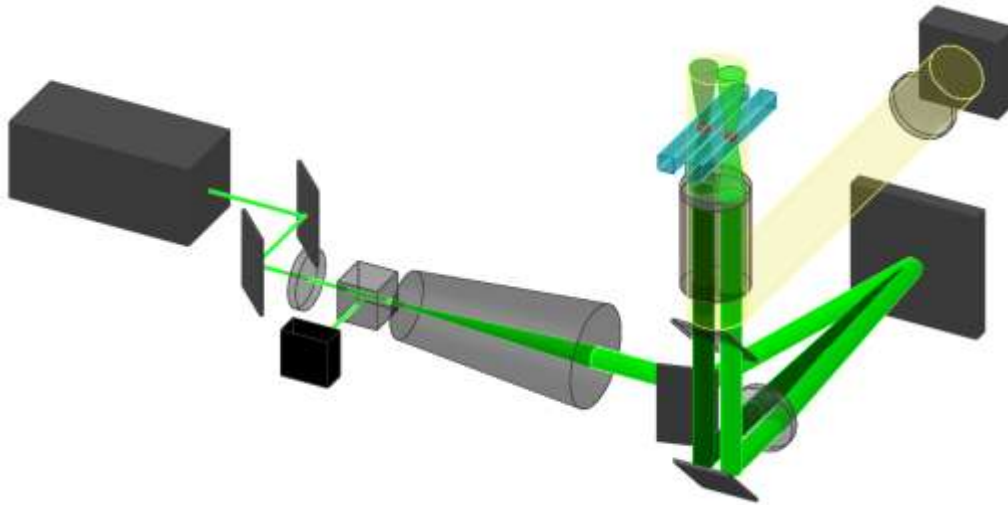


Particle manipulation with holographic optical tweezers (HOT)

Use of a spatial light modulator to split a single laser beam into many optical traps that can be independently positioned in 3D

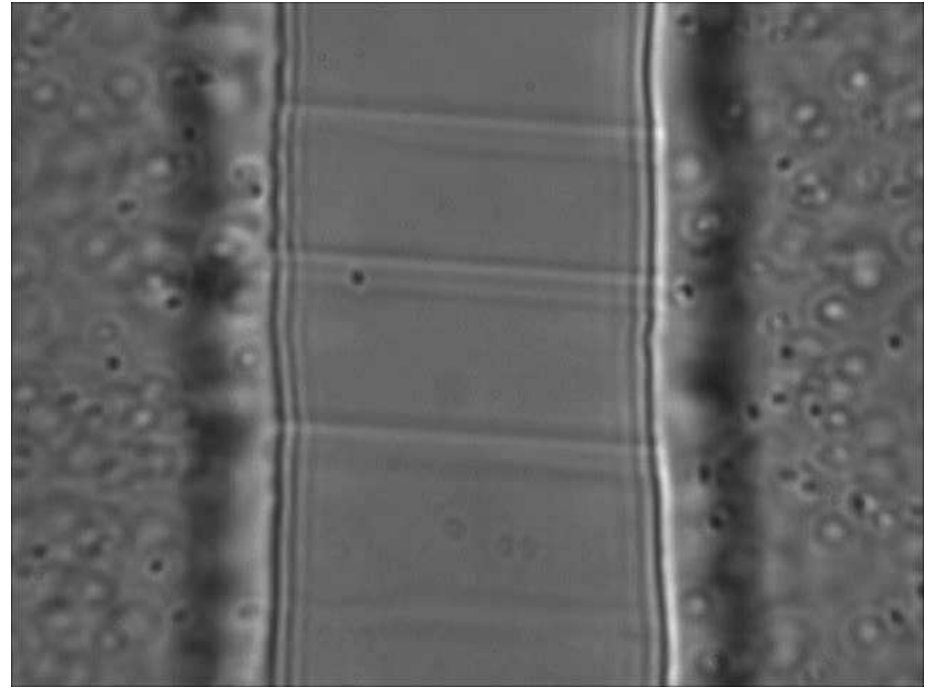


Manipulation of different particles in different channels

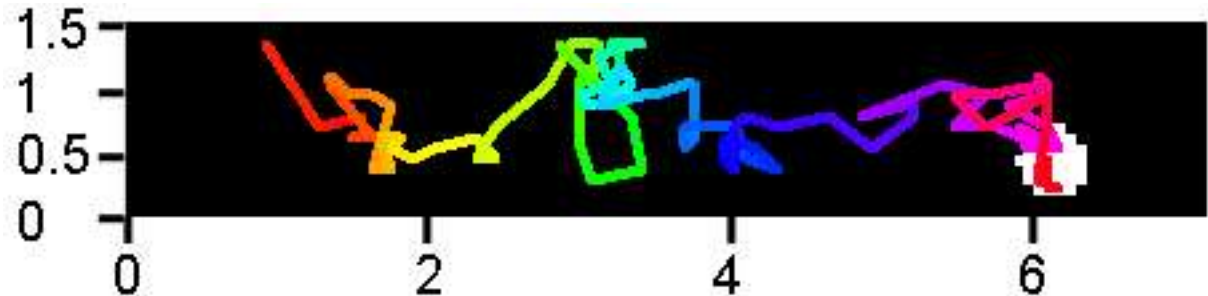


Particle tracking

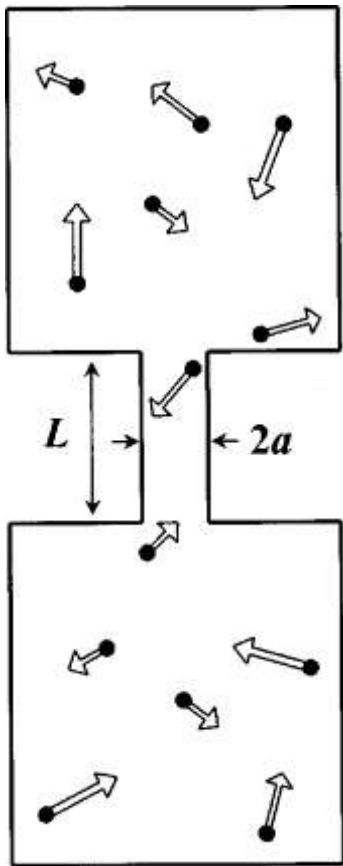
Tracking single particles
diffusing through the
different microfluidic
channels



Particles are isolated from
the background and
tracked by a custom-
made program based
on Labview



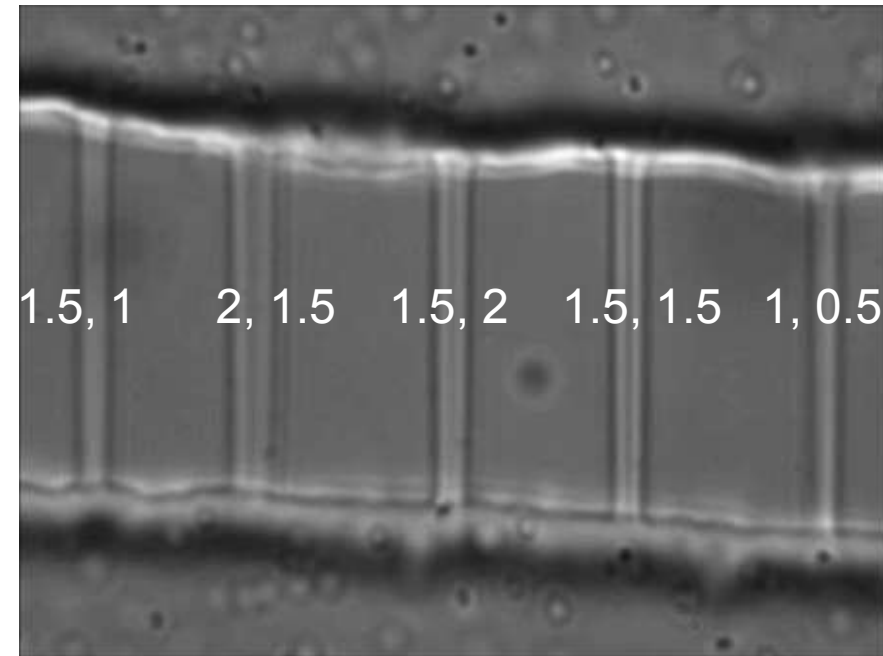
Particle diffusion through long and narrow microchannels



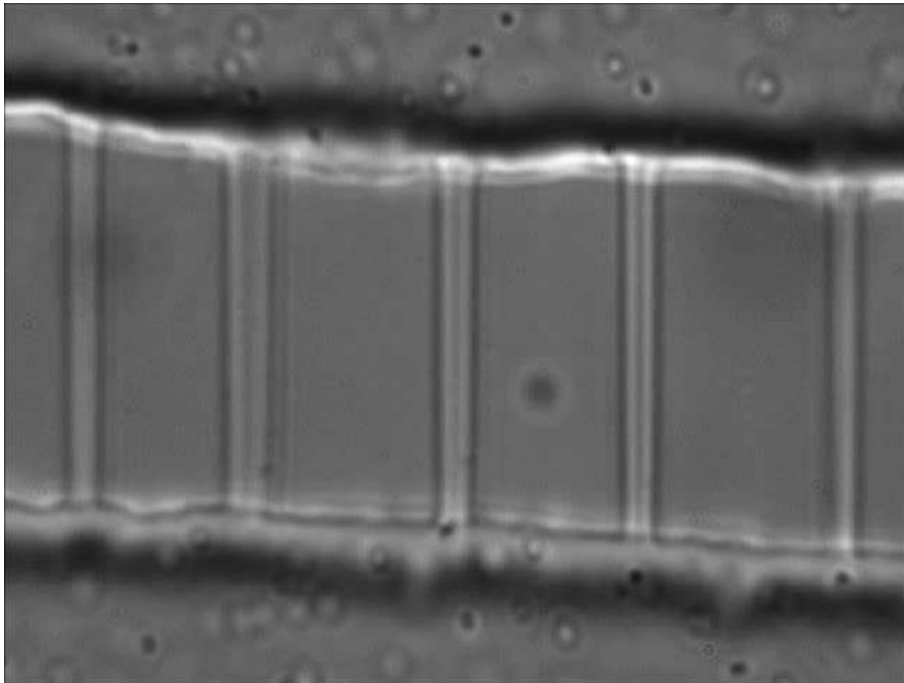
The microfluidic chip is filled with carboxyl-functionalized polystyrene particles with mean diameter of 500 nm dispersed in a 0.2 % solids (w/v) 5 mM KCl suspension

27 particles per $10 \mu\text{m}^3$

The microscopic reservoirs are connected by five parallel channels with different width and thickness:



Particle diffusion through arrays of microchannels

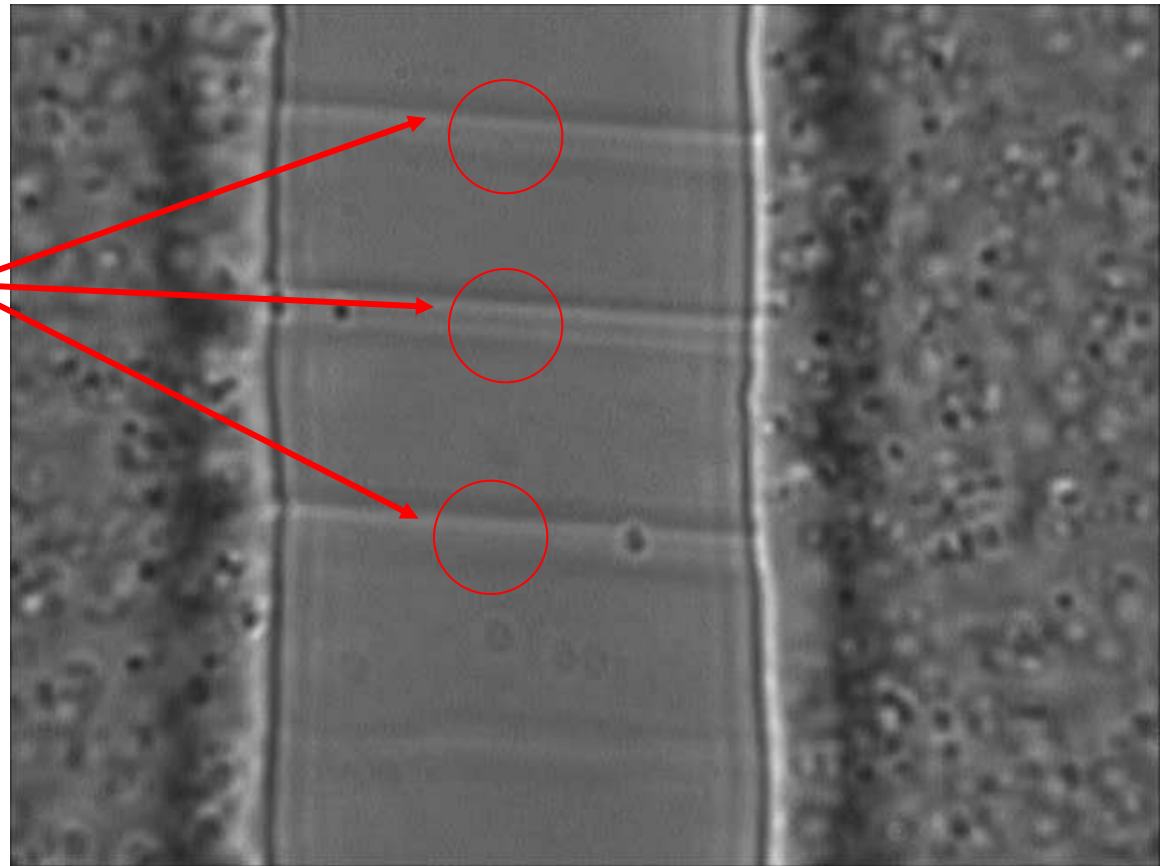


Width	1.5	2	1.5	1.5	1
Thick.	1	1.5	2	1.5	0.5
Pred. Att.	1.5	2.5	2.5	1.5	0
Top Att.	2	2	3	1	0
Bot. Att.	1	3	1	1	0

Particle diffusion with binding sites

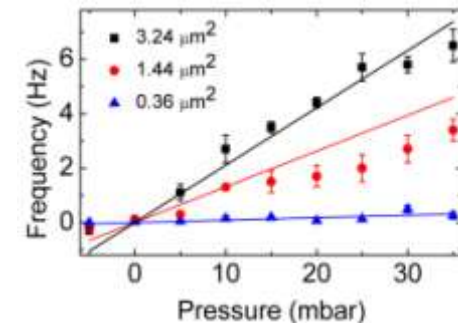
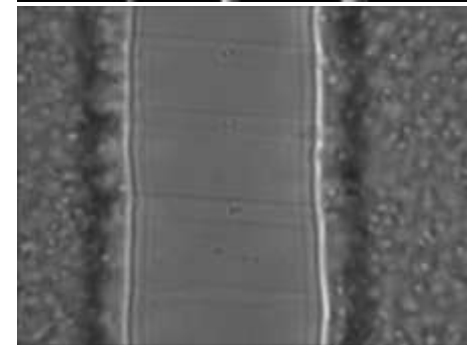
Looking for the optimal binding potential in protein channels

3 optical traps with different intensity placed in the middle of 3 different channels with the same dimension



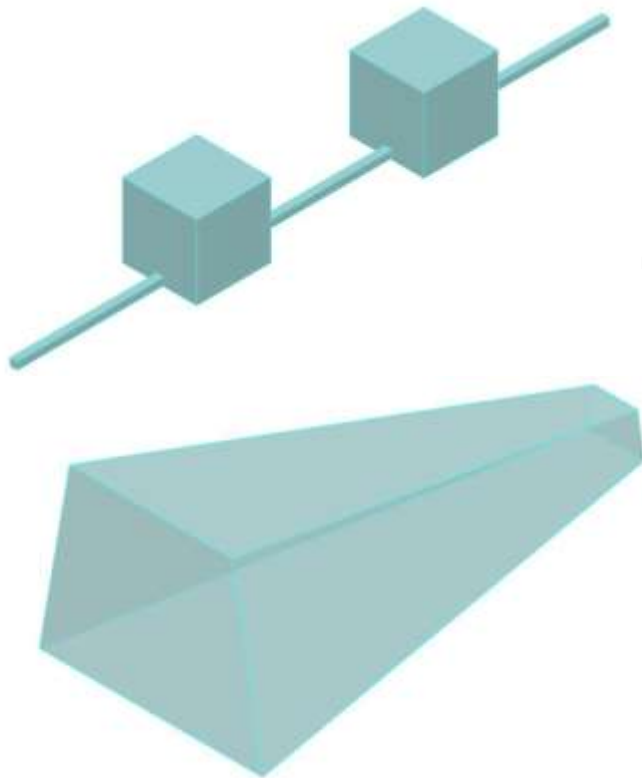
Conclusions

- Parallel channels with variable height and width
- Particle diffusion inside microfluidic channels
- Particle control through holographic optical tweezers and pressure gradient

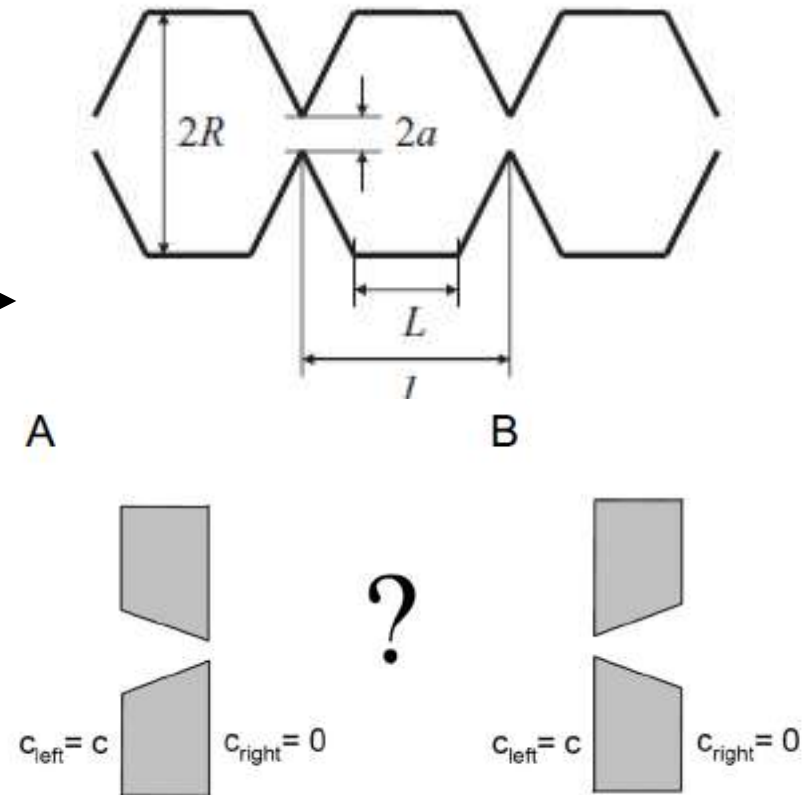


Outlook

FIB deposition of complex 3D structures



Entropic effects in particle transport



Acknowledgments

Dr. U. F. Keyser

C. Chimerele, O. Otto

Dr. R. Langford

Dr. D. G. A. L. Aarts (Department of Chemistry, University of Oxford)

Prof. S. M. Bezrukov (Laboratory of physical and structural biology, NIH)

Thank you for the attention!