



LU ASI zinātniskais seminārs ar vieslektoru

Dag Hanstorp

Trīs desmitgades lāzera spektroskopijas

the simple bending formula. Groups of rods (containing 10-22 rods) were given different periods of etching, and the depth of material removed from the surface of the rods was calculated for each group.

The variation of the mean breaking strength of these groups of rods, with depth of material removed from the surface, is shown in Fig. 1. Also shown on Fig. 1 are the 95 per cent confidence limits on the mean strength and the highest strength value recorded in each group of rods. Fig. 2 is a histogram comparing the distribution of breaking stresses for a group of rods which have been etched for 40 min. with that for unetched rods. The maximum strengths

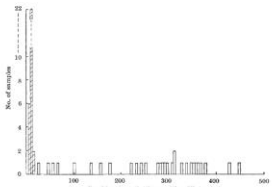


Fig. 1. Frequency of breaking strength at a loading of breaking strength for etched (unetched) and unetched rods. The average breaking strength for 40 min. etching removed from surface = 4.5×10^{-2} cm.

Behavior and Townes¹ have proposed a technique for the generation of very monochromatic radiation in the infra-red optical region of the spectrum using an alkali vapor as the active medium. Townes² and his associates³ have discussed proposals involving electron-excited gaseous systems. In this laboratory an optical pumping technique has been successfully applied to a fluorescent rod resulting in the attainment of negative temperatures and stimulated optical emission at a wave-length of 6943 Å.; the active material used was ruby (chromium in corundum).

A simplified energy-level diagram for triply ionized chromium in this crystal is shown in Fig. 3. When the material is irradiated with energy of wave-length of about 5200 Å., at a wave-length of about 5200 Å., chromium ions are excited to the 3P_2 state and then quickly lose some of their excitation energy through non-radiative transitions to the 3P_1 state⁴. This state then slowly decays by spontaneously emitting a sharp doublet, the components of which are at 6943 Å. and 6929 Å. (Fig. 2a). Under very intense excitation, the population of this metastable state (3P_1) can become greater than that of the ground state; this is the condition for negative temperatures and consequently amplification via stimulated emission.

To demonstrate the above effect a ruby crystal of 1 cm. dimensions coated on two parallel faces with silver was irradiated by a high-power flash lamp;

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494 NATURE August 6, 1960 vol. 192



Fig. 1. Energy-level diagram of Cr^{3+} in corundum, showing transitions produced.

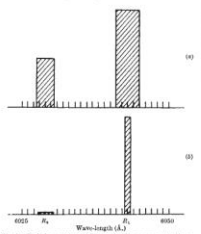
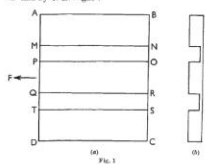


Fig. 2. Emission spectrum of ruby, a, low-power excitation; b, high-power excitation; c, high-power excitation.

have shown that the behaviour of the metal in these circumstances is in many ways simpler and more informative than that exhibited by wires or rods under tension.

From the point of view of the industrial study of creep, the method has the disadvantage that the specimens are somewhat troublesome to prepare and measure. To get over this difficulty I have devised a method in which the specimen has the form shown in Fig. 1. In the plate $ABCD$ (Fig. 1a) are cut rectangular grooves $MNOP$, $QRST$, as shown in cross-section in Fig. 1b. The metal $ABCD$ is securely held, and a force F applied to the metal $PQRS$ in a direction parallel to the grooves. Under these conditions the shear stress distribution in the rectangular plates $MNOP$, $QRST$ is not strictly uniform, as it is in the disk method, which is clear from the fact that the shear stress over the free ends must be zero. The distribution has been worked out by m^3 and by C. E. Inglis⁵.



Mr. D. B. Gilding has been working under my direction on the best form of the plate and has established that if the ratio of MN to NO is in the region of 7, the results on creep obtained with this disposition described correspond closely to those obtained by the method of Andrade and Ziffert. It seems possible that the new method may be of use in further investigations of creep and may also have applications to the problem of fatigue.

E. N. DA C. ANDRADE
Department of Metallurgy and Technology,
Imperial College of Science and Technology,
London, S.W.7.

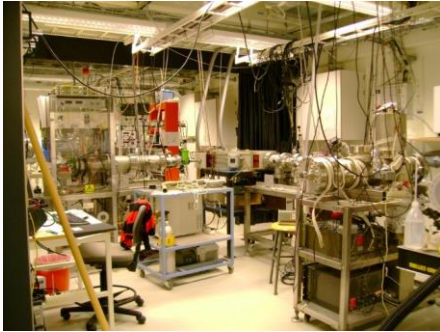
¹ Ashfield, R. S. *ibid.*, and Townes, C. H., *Proc. Roy. Soc. A*, **258**, 3 (1957); **264**, 121 (1959).
² Ashfield, R. S. *ibid.*, **264**, 121 (1959).
³ Ashfield, R. S. *ibid.*, **264**, 121 (1959).
⁴ Townes, C. H., *Proc. Roy. Soc. A*, **264**, 121 (1959).
⁵ Inglis, C. E., *Proc. Roy. Soc. A*, **264**, 121 (1959).
⁶ Inglis, C. E., *Proc. Roy. Soc. A*, **264**, 121 (1959).



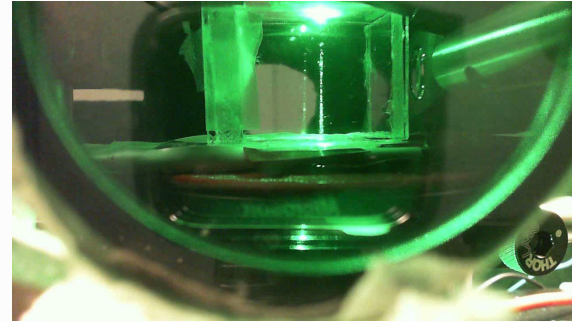
T. H. Maiman Nature 187, 493-494 (1960)

My research interests

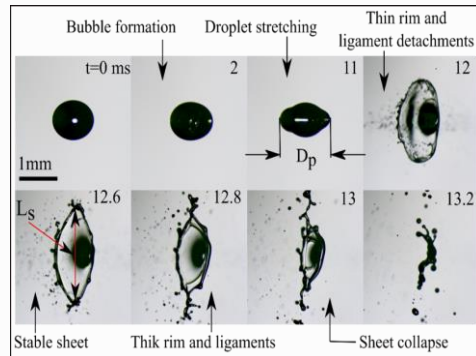
Spectroscopy of negative ions



Optical manipulation

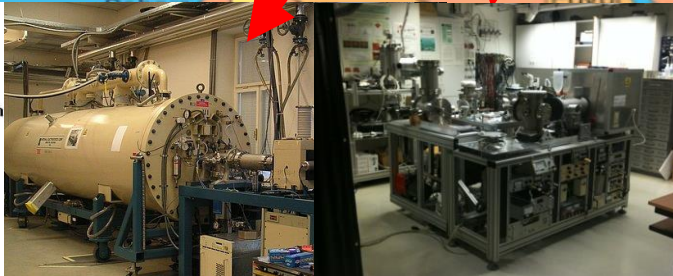
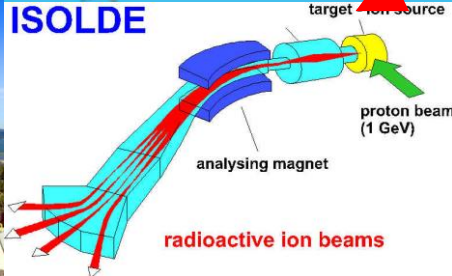
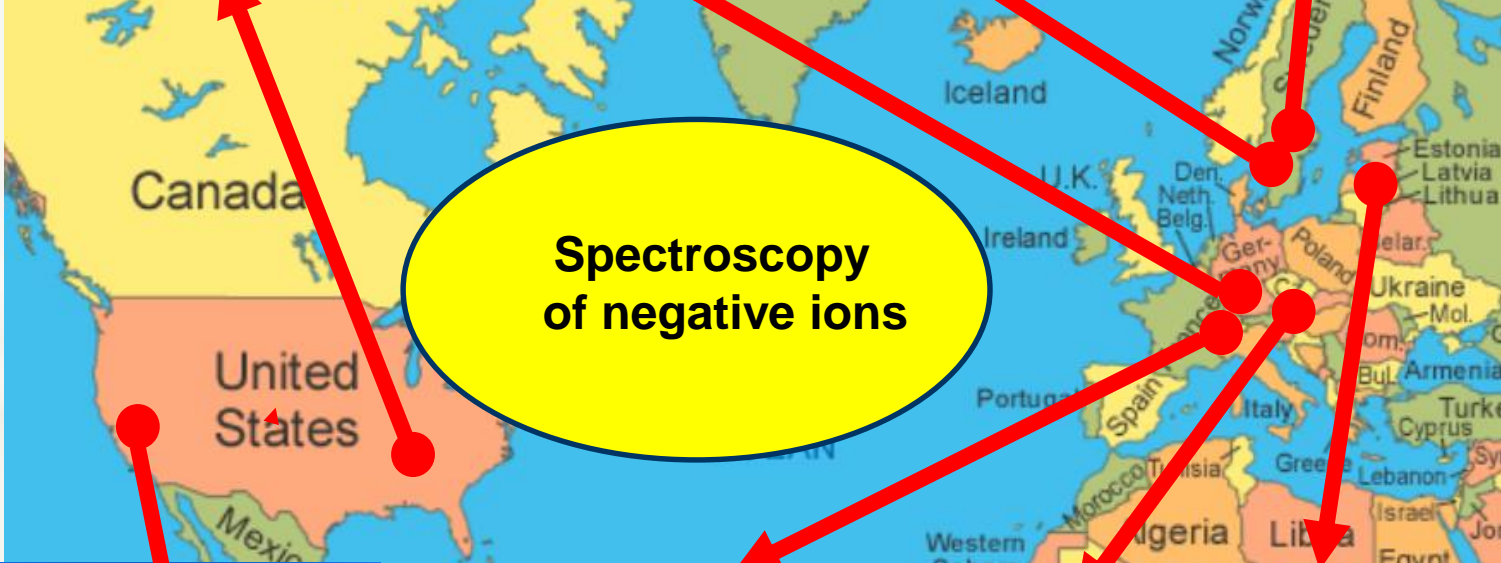


Femtosecond spectroscopy



Physics Education Research





Negative ions

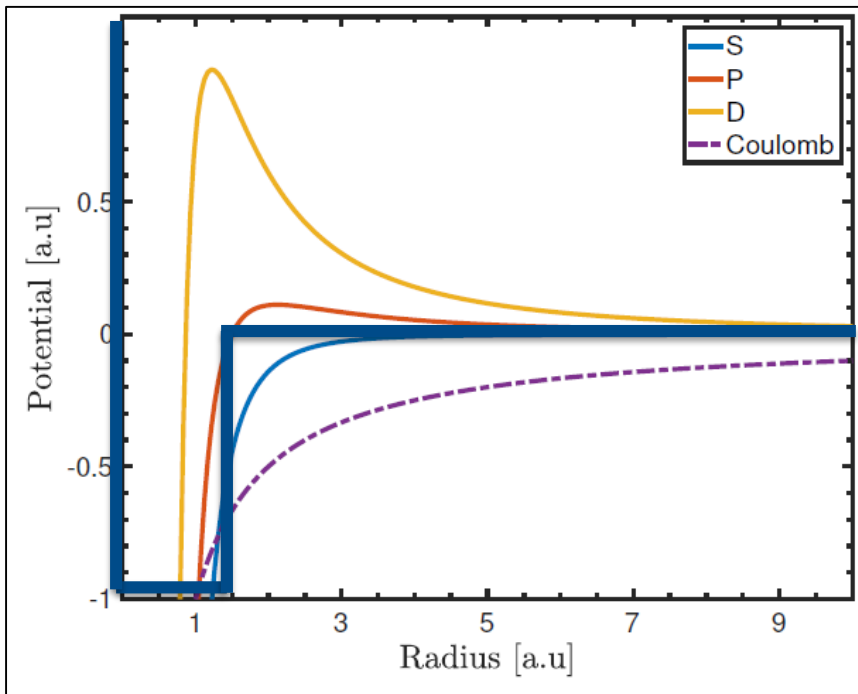


$$H|\Psi\rangle = \left[\frac{\hbar^2}{2\mu} \Delta + V(r) \right] |\Psi\rangle = E|\Psi\rangle$$

$$H = H_e + H_{ee} + V(r_i) = -\frac{\hbar^2}{2m_e} \sum_i \Delta_i + \sum_{i,j,i \neq j} \frac{e^2}{4\pi\epsilon_0 r_{ij}} - \frac{Ze^2}{4\pi\epsilon_0 r_i}$$

$$V_C(r) = -\frac{e^2}{4\pi\epsilon_0 r}$$

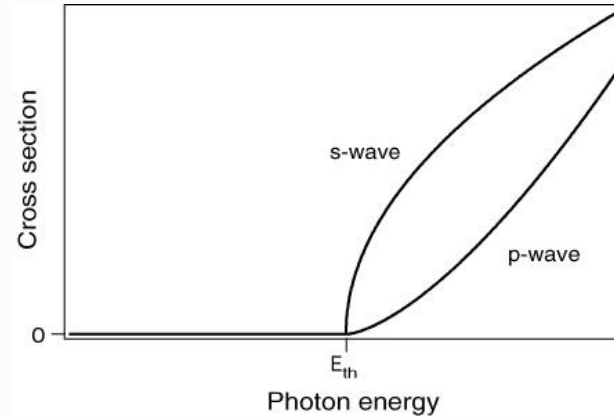
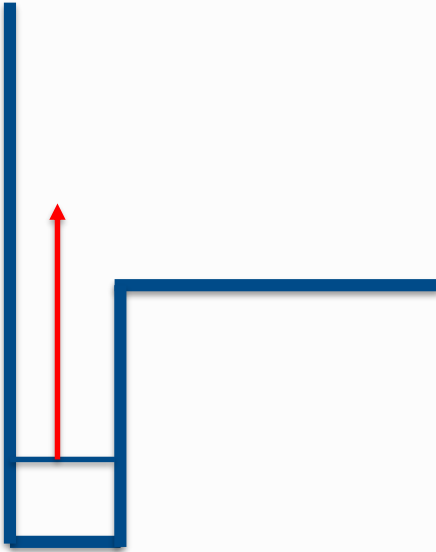
$$V(r) = \frac{l(l+1)}{2r^2} - \frac{\alpha_D}{2r^4}$$

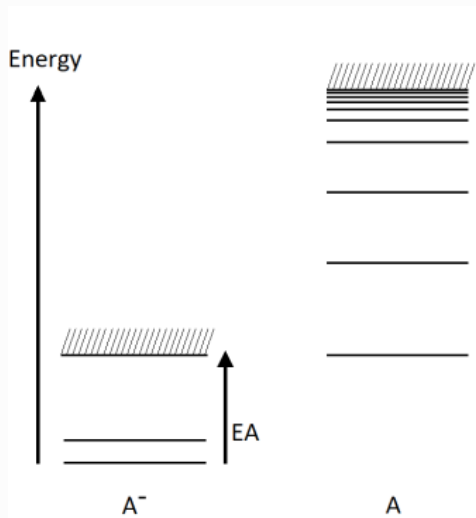


Photodetachment: $hf + A^- \rightarrow A + e^-$

Wigner threshold law: $\sigma(E) = (E_\gamma - E_{th})^{l+\frac{1}{2}}$

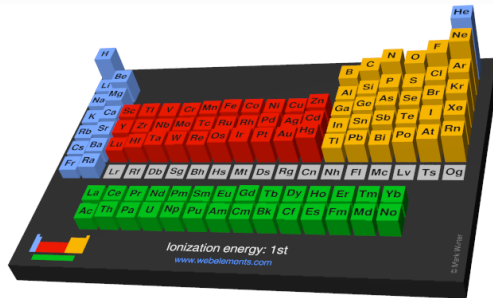
l angular momentum of emitted electron



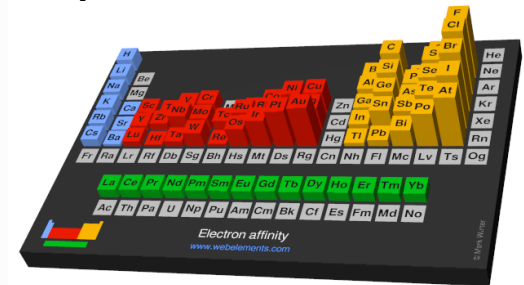


- **Electron affinity (EA)** is the binding energy of the additional electron (in the order of $\sim 1\text{eV}$)
- **Electron correlation dominated** binding allows probing of theories beyond Hartree-Fock approximation
- **Almost no bound states** with opposite parity existing (except La, Os, Ce, Th and U)

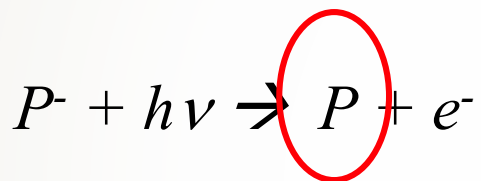
Periodic table of the a) ionization potentials



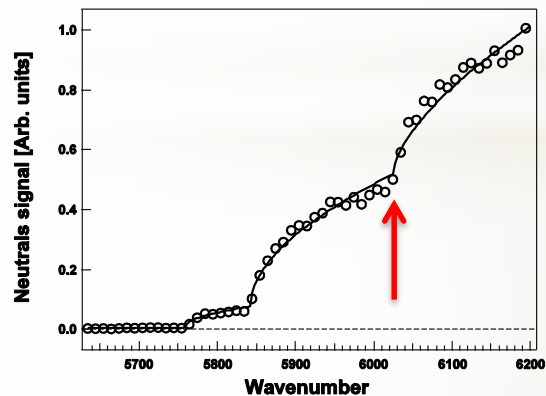
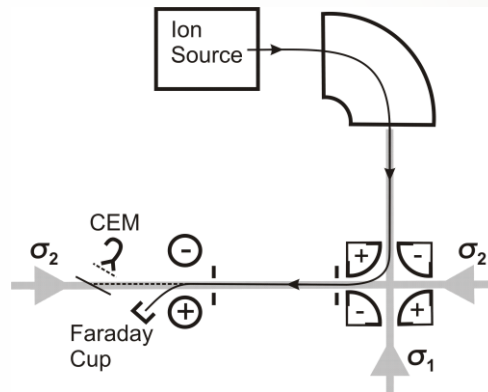
b) electron affinities



Determination of electron affinities:



$$\sigma^P = k (E - E_{EA})^{l+1/2}$$



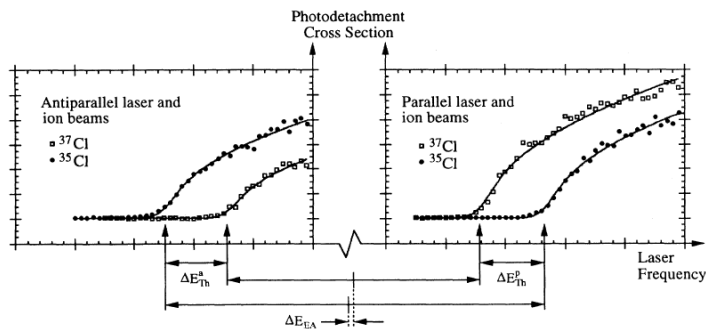
$$EA = 746.68(6) \text{ meV}$$

Isotope shift in the electron affinity of chlorine

U. Berzins^{*}, M. Gustafsson, D. Hanstorp, A. Klinkmüller, U. Ljungblad, and A.-M. Mårtensson-Pendrill
Department of Physics, Chalmers University of Technology, S-412 96 Göteborg, Sweden
and Göteborg University, S-412 96 Göteborg, Sweden
 (Received 14 July 1994)

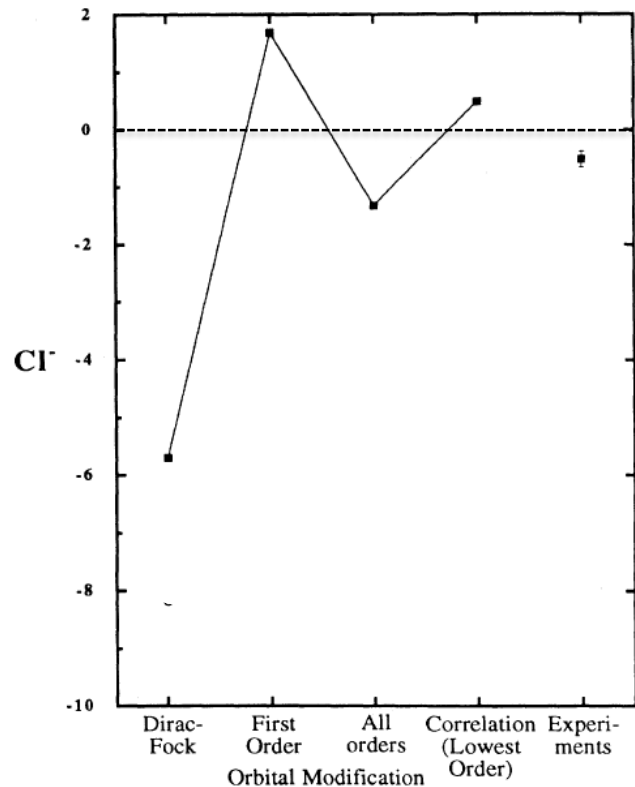
The specific mass shift in the electron affinity between ^{35}Cl and ^{37}Cl has been determined by tunable-laser photodetachment spectroscopy to be $-0.51(14)$ GHz. The isotope shift was observed as a difference in the onset of the photodetachment process for the two isotopes. In addition, the electron affinity of Cl was found to be $29138.59(22)$ cm^{-1} , giving a factor of 2 improvement in the accuracy over earlier measurements. Many-body calculations including lowest-order correlation effects demonstrate the sensitivity of the specific mass shift and show that the inclusion of higher-order correlation effects would be necessary for a quantitative description.

PACS number(s): 35.10.Hn, 32.80.Fb, 31.30.Gs



$$\delta\nu_{\text{SMS}} = -0.51(14) \text{ GHz}$$

Specific Mass Shift (GHz)



Isotope shift on the chlorine electron affinity revisited by an MCHF/CI approach

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Abstract

Today, the electron affinity is experimentally well known for most of the elements and is a useful guideline for developing *ab initio* computational methods. However, the measurements of isotope shifts on the electron affinity are limited by both resolution and sensitivity. In this context, theory is of great help to further our knowledge and understanding of atomic structures, even though correlation plays a dominant role in negative ions' properties and, particularly, in the calculation of the specific mass shift contribution. This study solves the longstanding discrepancy between calculated and measured specific mass shifts on the electron affinity of chlorine (Berzinsh *et al* 1995 *Phys. Rev. A* **51** 231).

(Some figures may appear in colour only in the online journal)

	SMS	MS	FS	RIS	IS
				This work	
HF	-1.348	-0.607	-0.003(22)	-1.351(22)	-0.610(22)
val. FC-MCHF	-0.674	+0.067	-0.002(20)	-0.676(20)	+0.065(20)
val. MCHF	-0.495	+0.246	-0.003(21)	-0.497(21)	+0.244(21)
final results	-0.535(51)	+0.206(51)	-0.003(22)	-0.538(72)	+0.203(72)
				Berzinsh <i>et al</i> [7]	
Exp.				-0.51(14)	+0.22(14)
DF	-1.3	-0.6	+0.014(14)	-1.3	-0.6
MB low corr.	+0.50	+1.24	+0.014(14)	+0.51(2)	+1.26(2)

DESIREE – Double Electrostatic Ion Ring Experiment



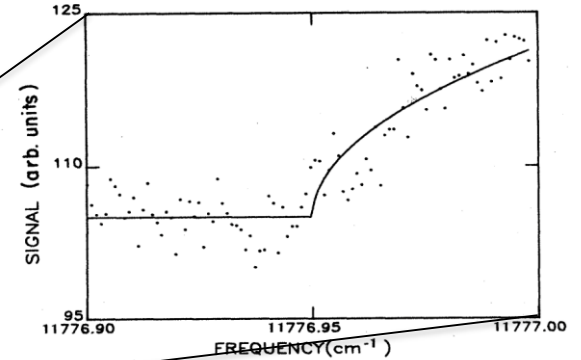
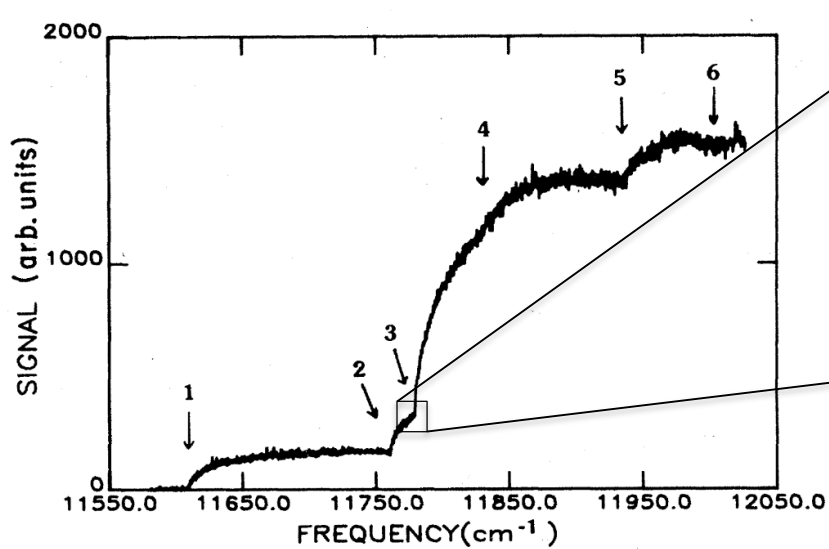
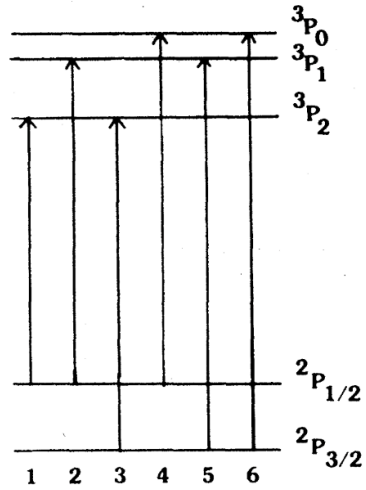
mu

eV

s

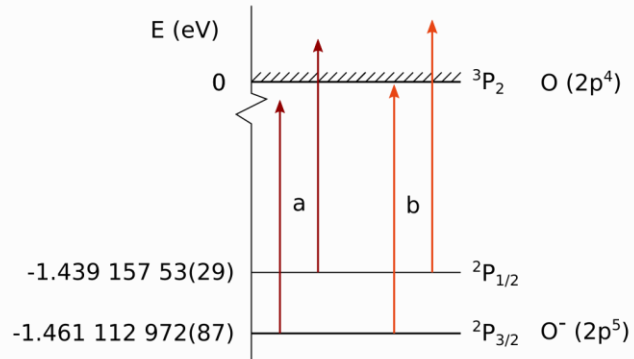
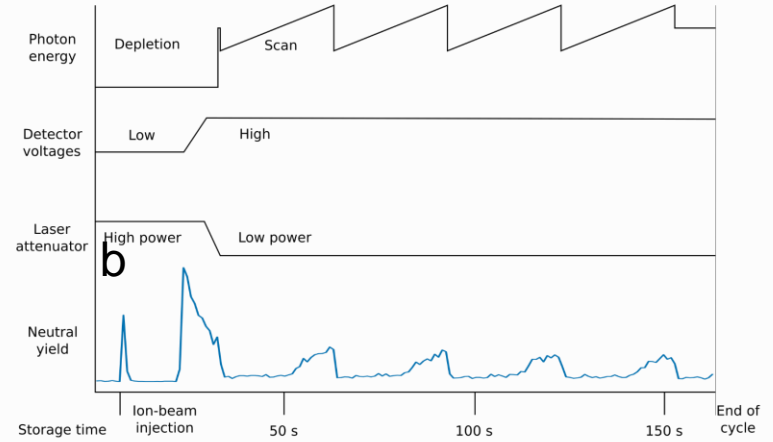
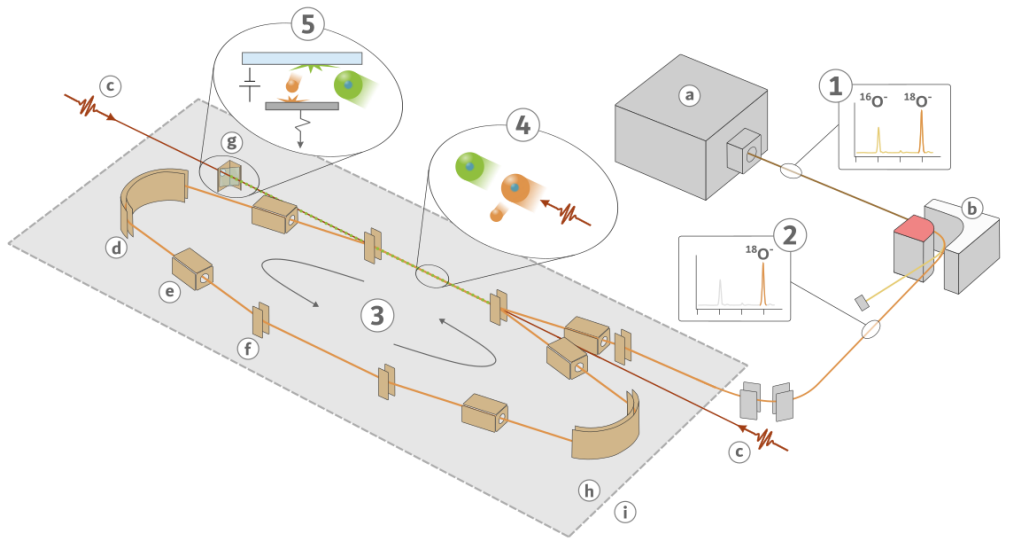
RO

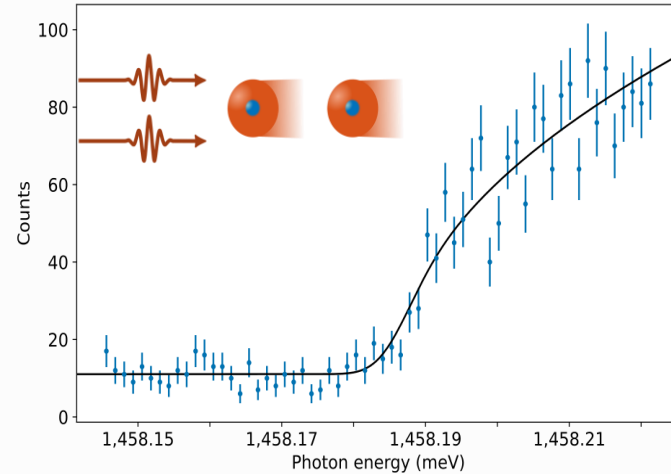
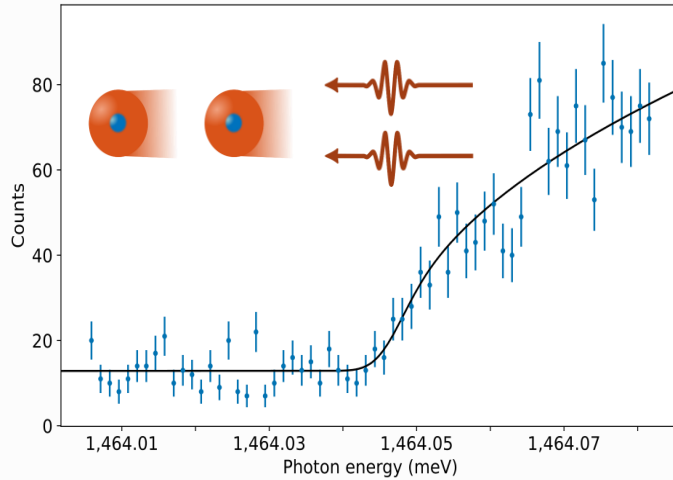
A Precision measurement of the EA of O^-



$$S/B = 15/100 = 1/7$$

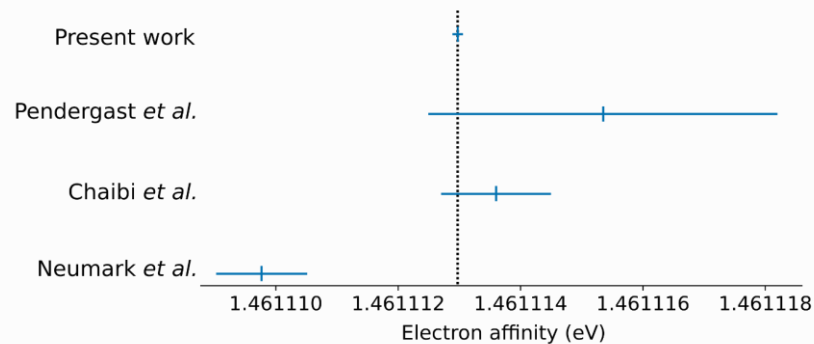
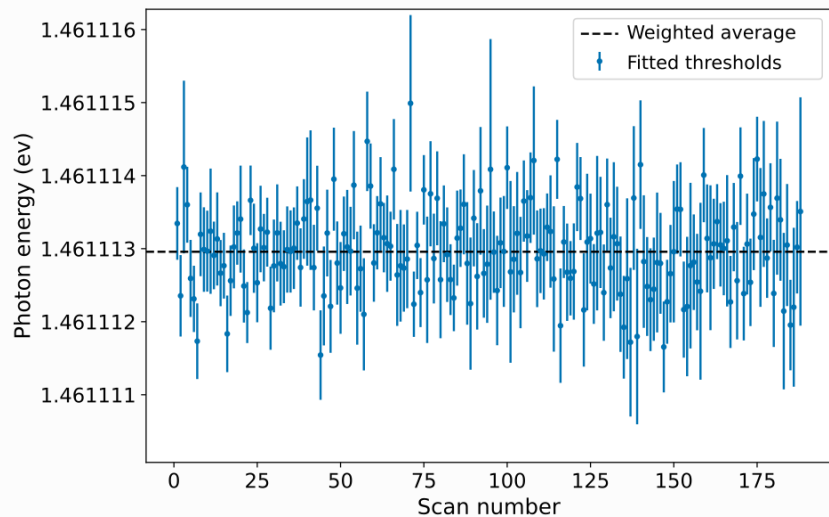
Neumark, D. M., et al. Physical Review A 32.3 (1985): 1890





$S/B = 80/15 = 5/1 \rightarrow 35$ times improvement

$$\sqrt{E_{EA}^p E_{EA}^a} = \sqrt{\frac{1+v/c}{\sqrt{1-v^2/c^2}} E_{EA} \frac{1-v/c}{\sqrt{1-v^2/c^2}} E_{EA}} = \sqrt{E_{EA} E_{EA}} = E_{EA}$$



Final result: $E_{EA} = 1.461\,112\,972\,(87)\text{ eV}$

- 10-fold precision improvement
- Theoretical results vary between ~1.45-1.47 eV

¹⁸⁻¹⁶O isotope shift

- Previous IS experimental value: -9.2(2.2) μeV
C. Blondel, Physical Review A 64, 052504 (2001).

Previous Theoretical value: -7.104 μeV
Godefroid and C. F. Fischer, Phys. Rev. A 60, R2637 (1999).

$$E_{EA}({}^{16}\text{O}) = 1.461\,112\,972\,(87)\text{ eV}$$

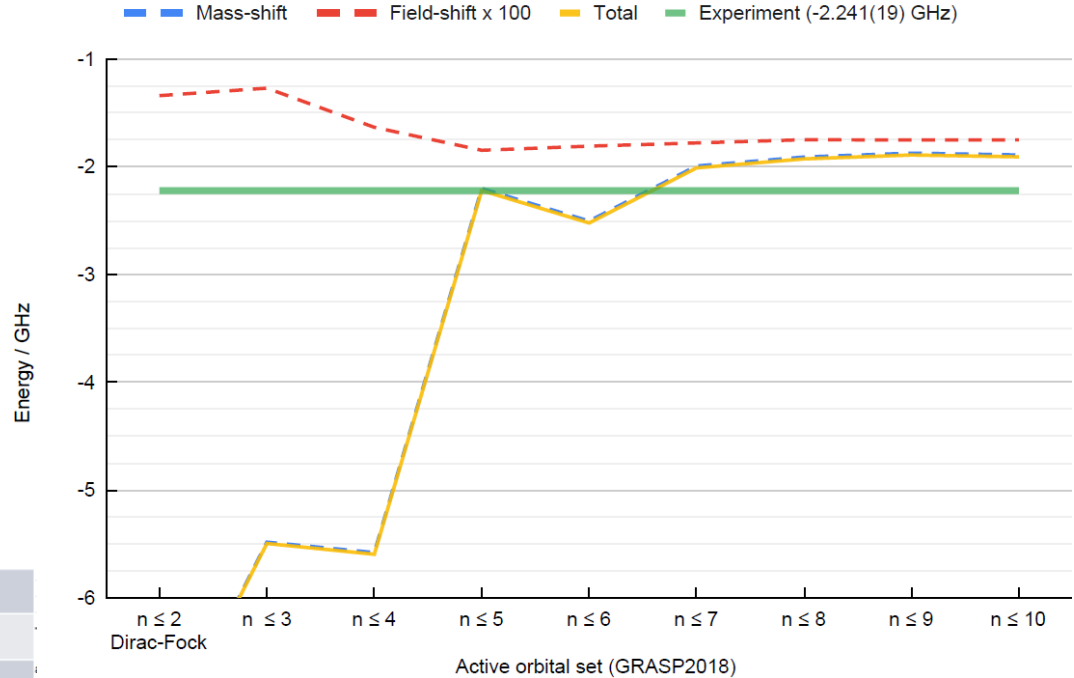
$$E_{EA}({}^{18}\text{O}) = 1.461\,103\,706\,(67)\text{ eV}$$

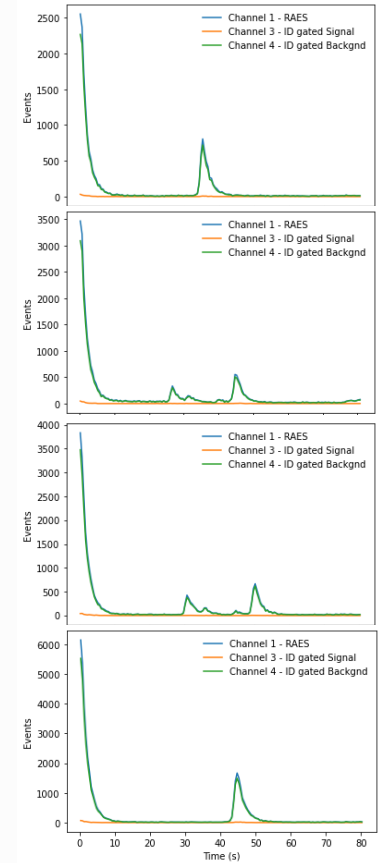
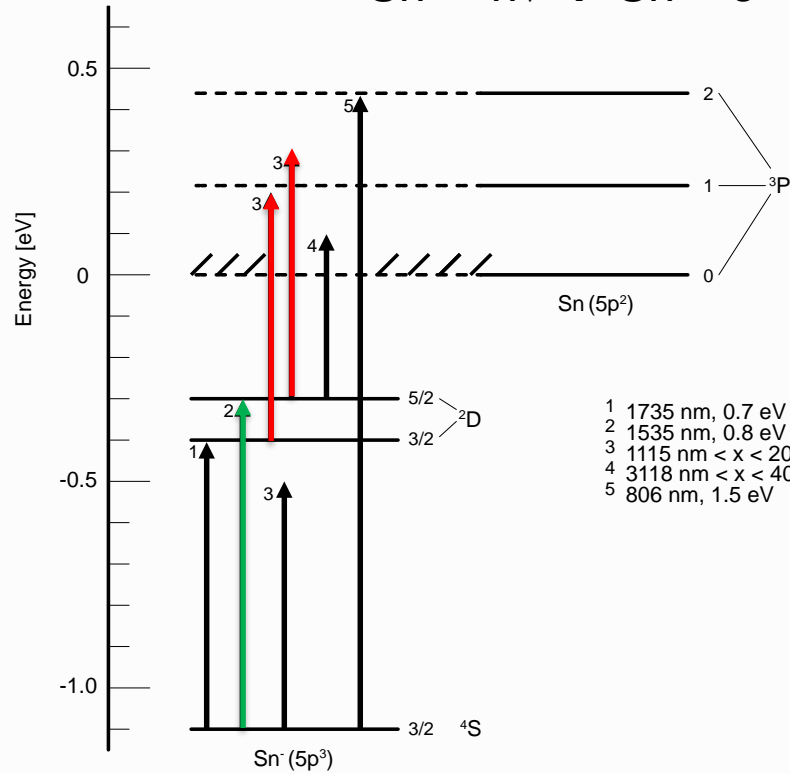
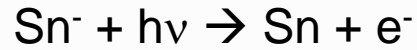
$$IS = -0.000\,009\,267(11)\text{ eV}$$

Theory by Jon Grumer

- **Atomic structure:** MCDHF + RCI using latest dev. version of GRASP*, correlation model inspired by Godefroid and Froese Fischer***
- **total IS on EA** = $-7.9 \mu\text{eV}$
Mass shift dominates completely
tricky to converge due to counteracting normal and specific mass shifts

		year	IS (μeV)
Blondel	Exp.	2001	9.2 (2.2)
Godefroid	Theory	1999	7.104
Kristiansson	Exp.	2022	9.267(11)
Grumer	Theory	2023	7.90





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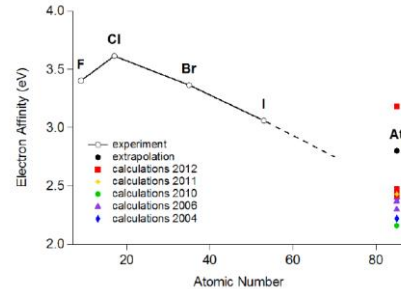
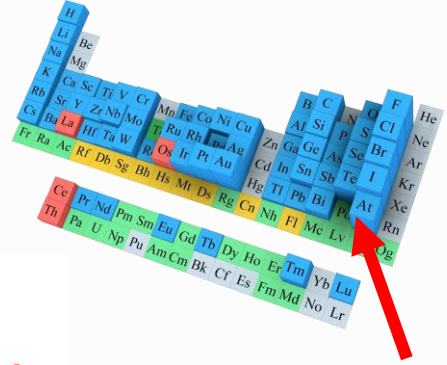
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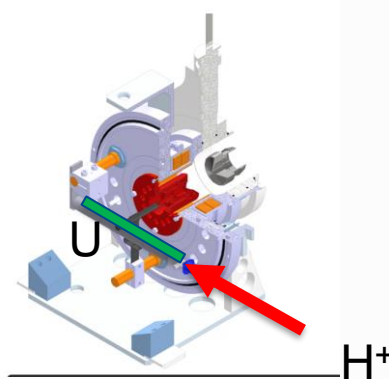
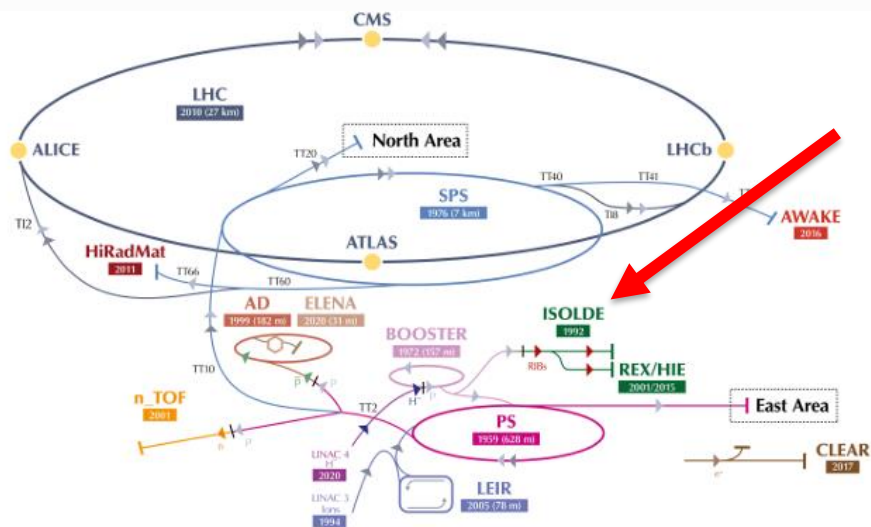
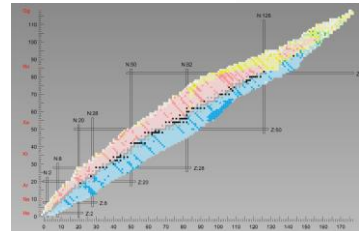
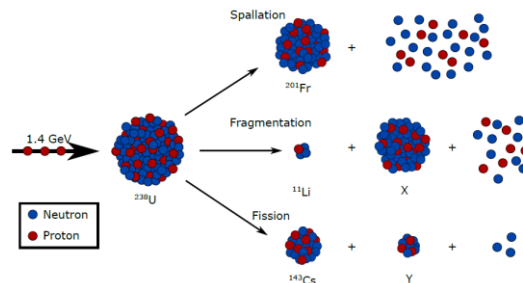
The electron affinity of Astatine

- Least abundant element on earth
- 70 mg in the crust of the earth (1 atom per 100 kg mass)
- Decays through α -decay
- Small knowledge about its chemical and physical properties
- Used in cancer treatment Targeted Alfa Therapy (TAT) (suitable lifetime and energy, non-toxic, non-radioactive daughters)



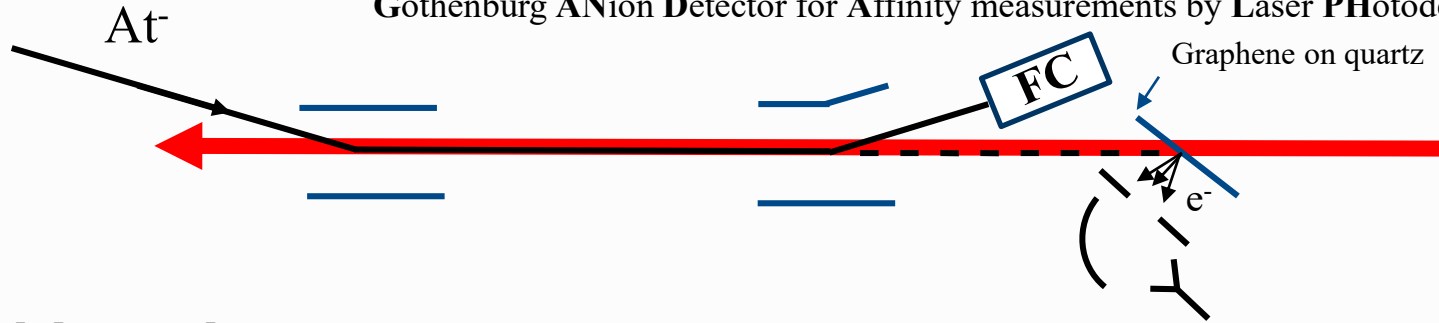


ISOLDE



GANDALPH

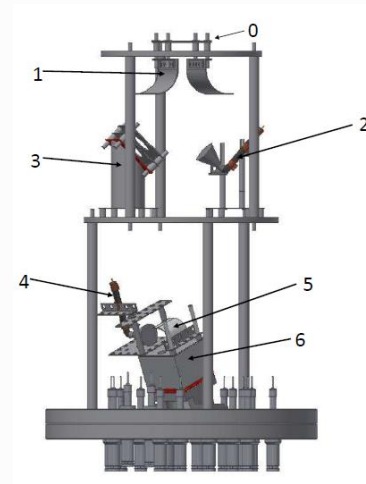
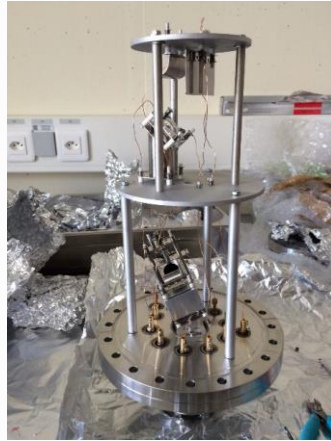
Gothenburg ANion Detector for Affinity measurements by Laser PHotodetachment



For each laserpuls:

Signal:
0.01 atom

Background:
 10^{14} photons



Drawing: Annie Ringvall-Moberg

RESULTS

Experiment: EA= 2.415 78 (5) eV

Theory EA = 2.414 (16) eV

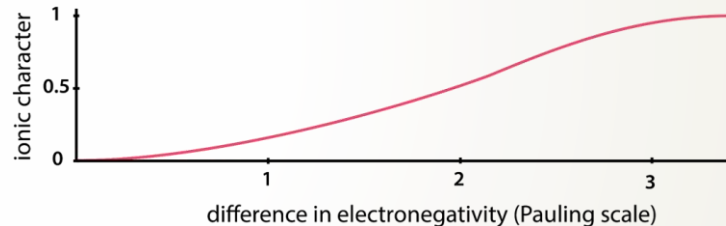
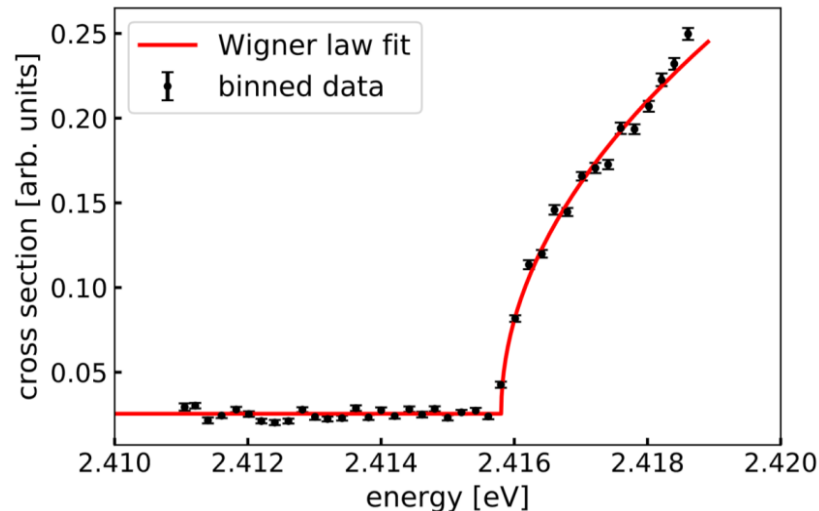
By Anastasia Anastasia Borschevsky and co-workers: DIRAC15 program package using the single reference coupled-cluster approach in the framework of the Dirac- Coulomb Hamiltonian (DC-CCSD(T))

Rothe et al. Nat. Commun. 4, 1835 (2013).

IP(At) = 9.317 51(8) eV



Electronegativity = $(IP + EA)/2 = 5.866 65 \text{ eV}$

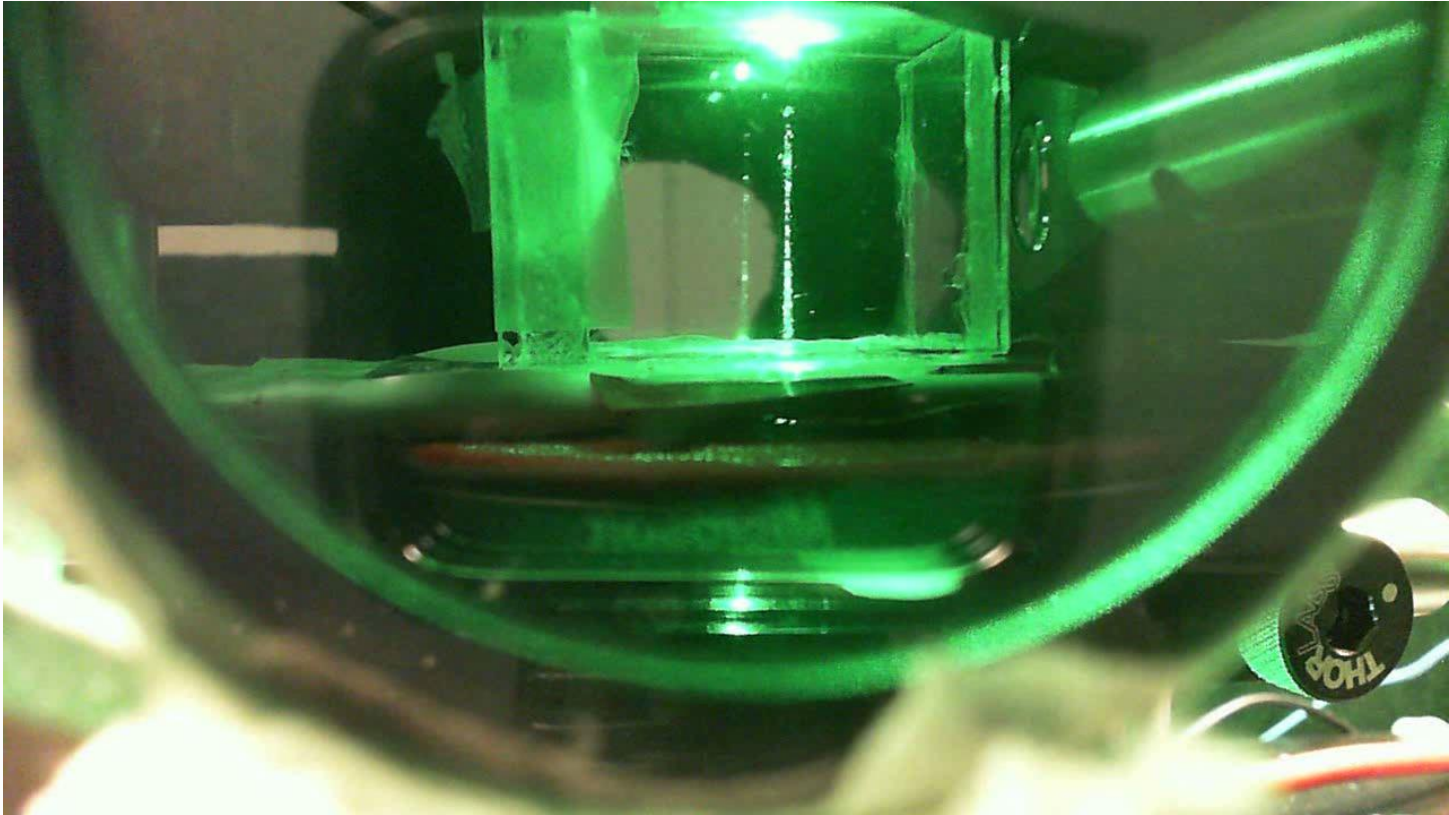


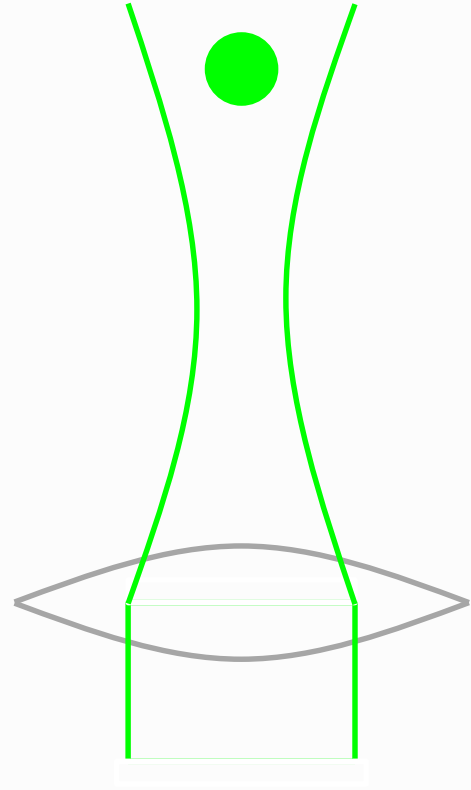
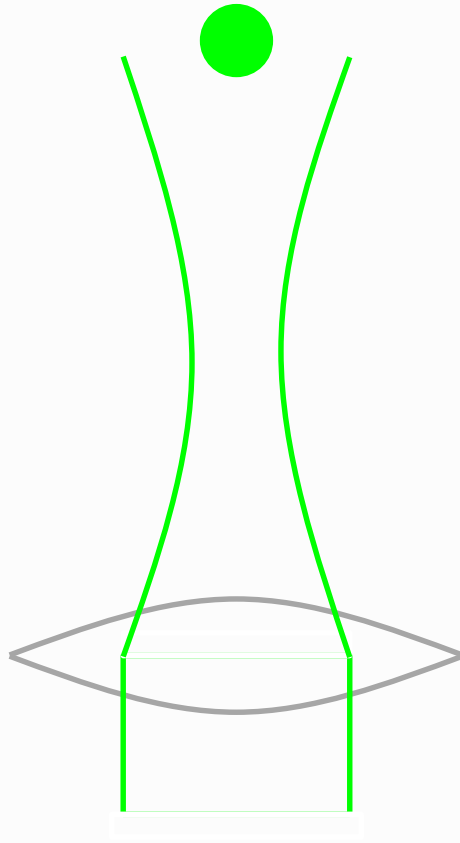
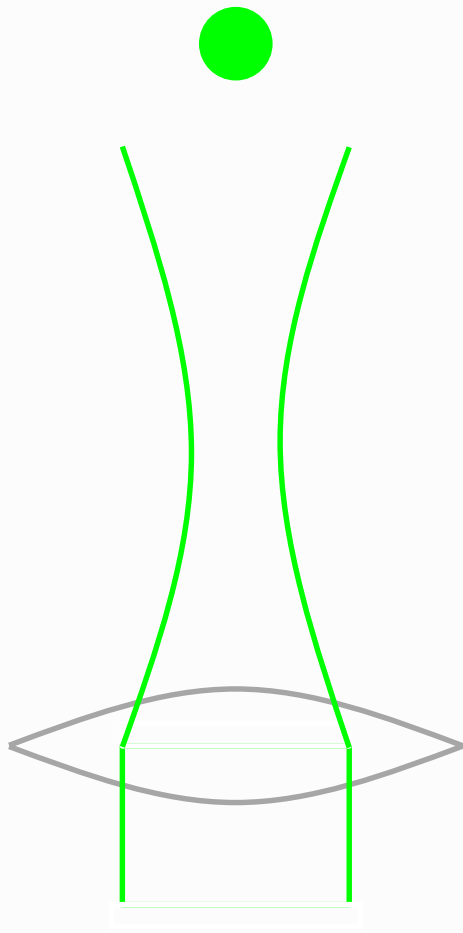
¶ EUROPEAN ORGANIZATION FOR NUCLEAR RESEARCH ¶

Proposal to the ISOLDE and Neutron Time-of-Flight Committee ¶

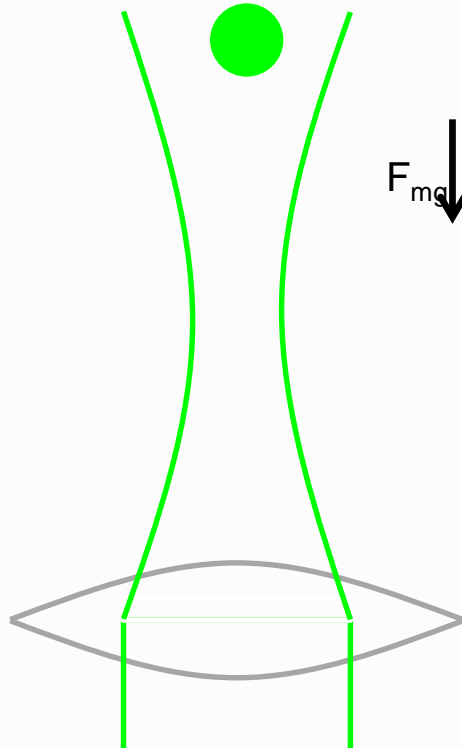
¶ Measurement of shifts in the electron affinities of chlorine isotopes ¶

¶
Dag Hanstorp¹, Jakob Welander¹, David Leimbach¹, Annie Ringvall-Moberg^{1,2}, Michel Godefroid³,
Per Jönsson⁴, Jörgen Ekman⁴, Tomas Brage⁵, Klaus Wendt⁶, Reinhard Heinke⁶, Oliver Forstner⁷,
Yuan Liu⁸, Ronald Garcia-Ruiz⁹, Shane Wilkins⁹, Adam Vernon⁹, Cory Binnersley⁹, Kieran Flanagan⁹,
Gerda Neyens¹⁰, Agi Koszorus¹⁰, Kara Lynch², Sebastian Rothe², Tim Giles², Katerina Chrysalidis^{2,6},
Pierre Larmonier², Valentin Fedosseev² and Bruce Marsh². ¶

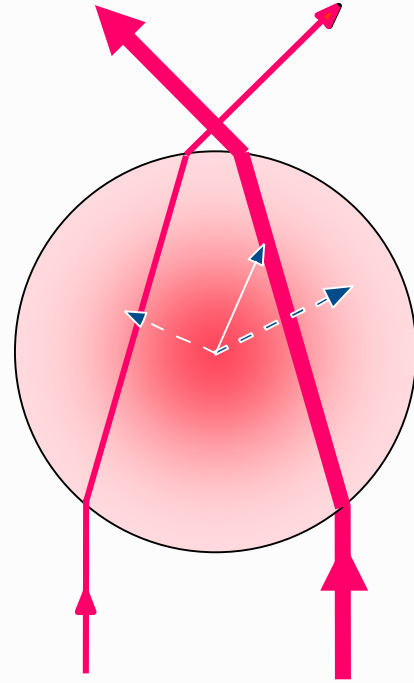




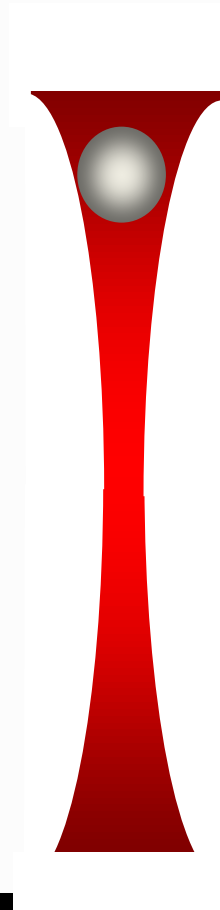
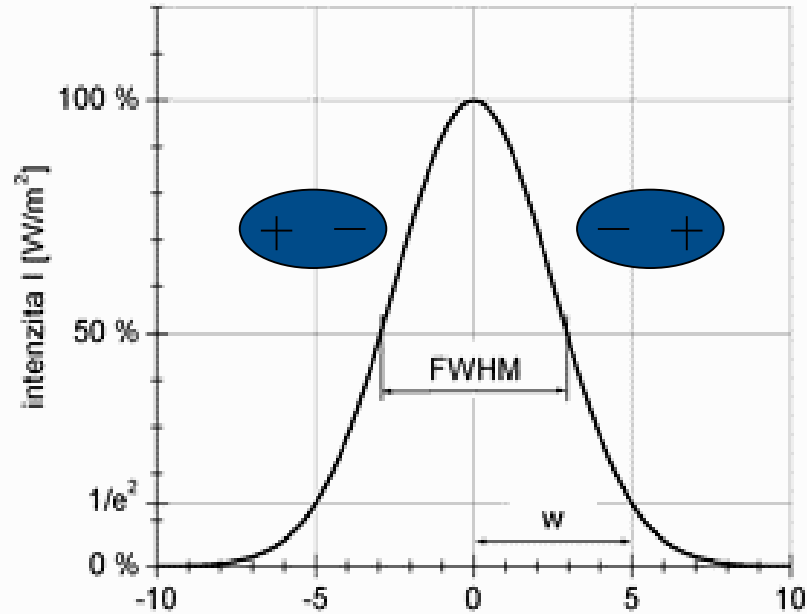
OPTICAL LEVITATION



F_{mg} F_{rad}



Trapping in Wave optic



Optical levitation

VOLUME 24, NUMBER 4

PHYSICAL REVIEW LETTERS

26 JANUARY 1970

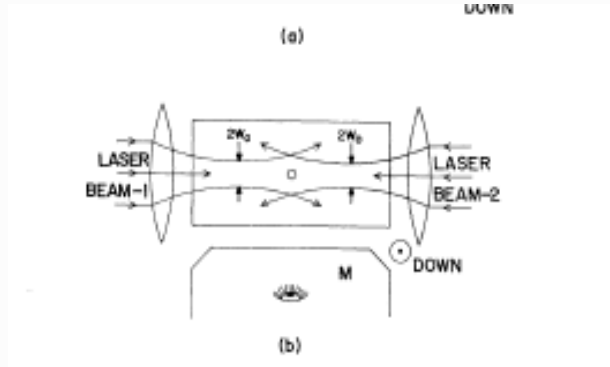
ACCELERATION AND TRAPPING OF PARTICLES BY RADIATION PRESSURE

A. Ashkin

Bell Telephone Laboratories, Holmdel, New Jersey 07733

(Received 3 December 1969)

Abstract: The acceleration and trapping of particles by radiation pressure is described.



APPLIED PHYSICS LETTERS

VOLUME 19, NUMBER 8

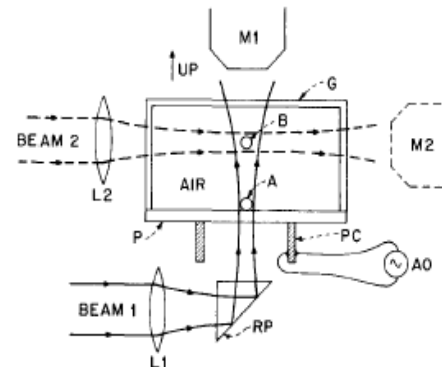
15 OCTOBER 1971

Optical Levitation by Radiation Pressure

A. Ashkin and J. M. Dziedzic

Bell Telephone Laboratories, Holmdel, New Jersey 07733

(Received 14 June 1971; in final form 13 August 1971)



OPTICAL MANIPULATION - ARTHUR ASHKIN

- 1970 – Optical levitation

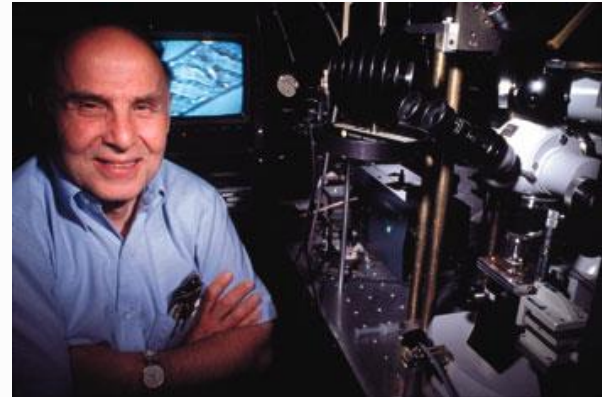
Ashkin&J.M.Dziedzic, App.Phys.Lett.**19**,283(1971)

- 1986 – The optical tweezers -trapped living

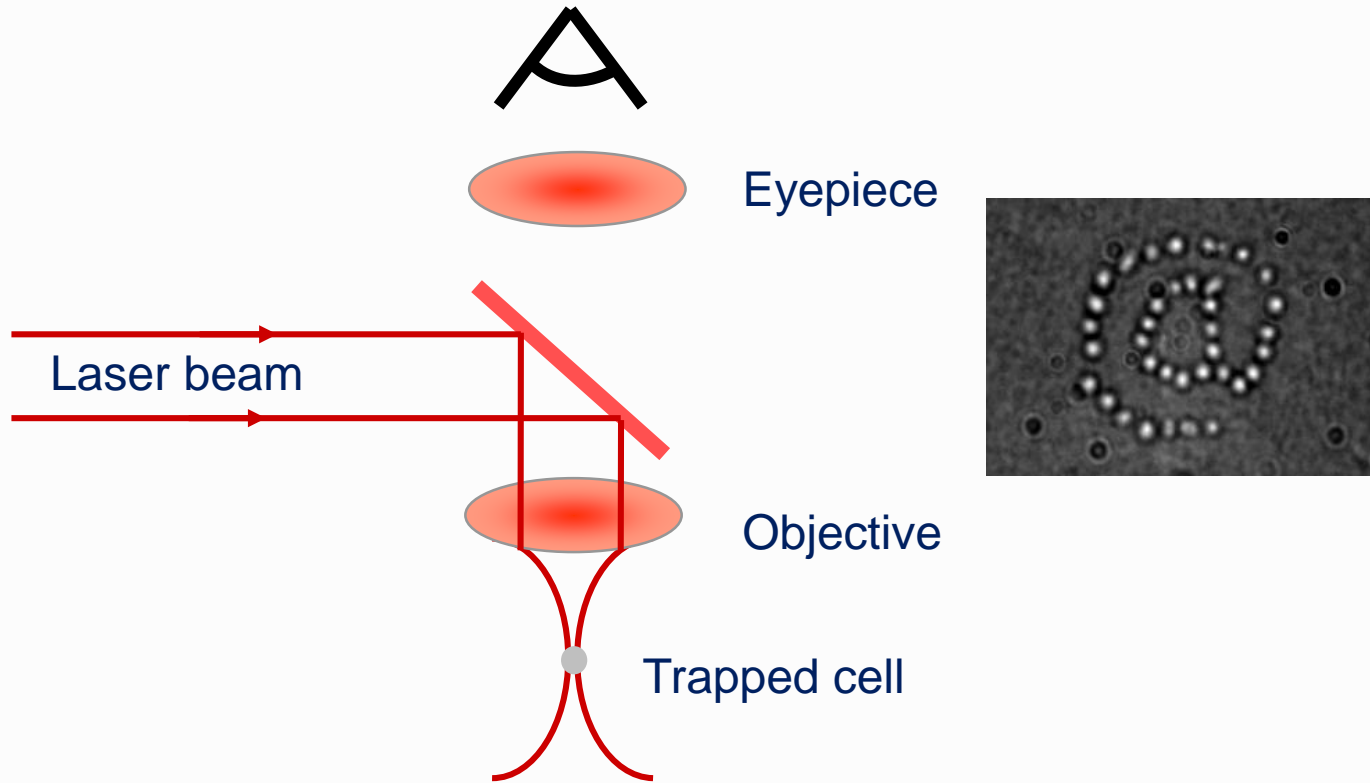
A. Ashkin et al., Optics Letters. **11**, 288 (1986)

- Nobel prize in physics 2018

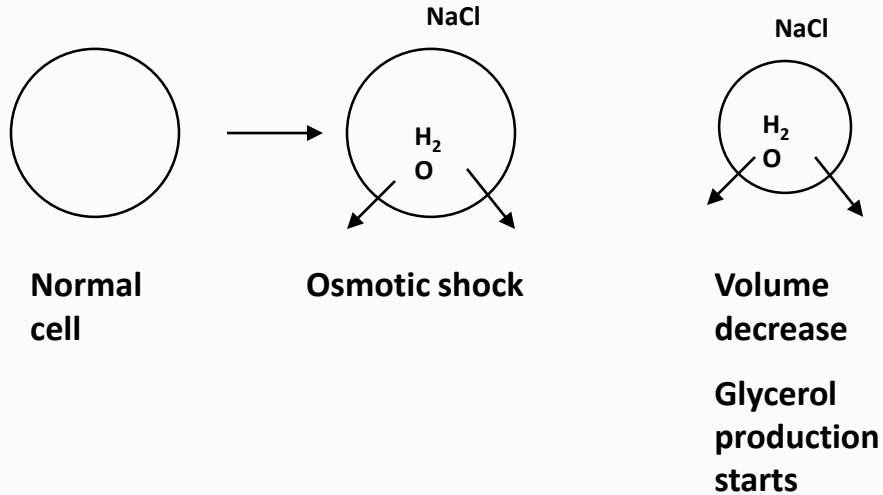
“for the optical tweezers and their application to biological systems.”



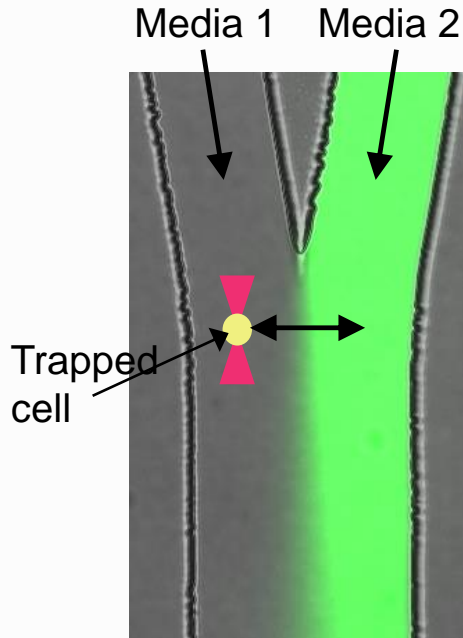
The optical tweezers



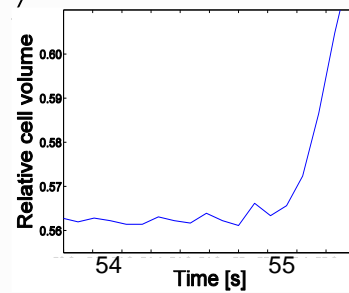
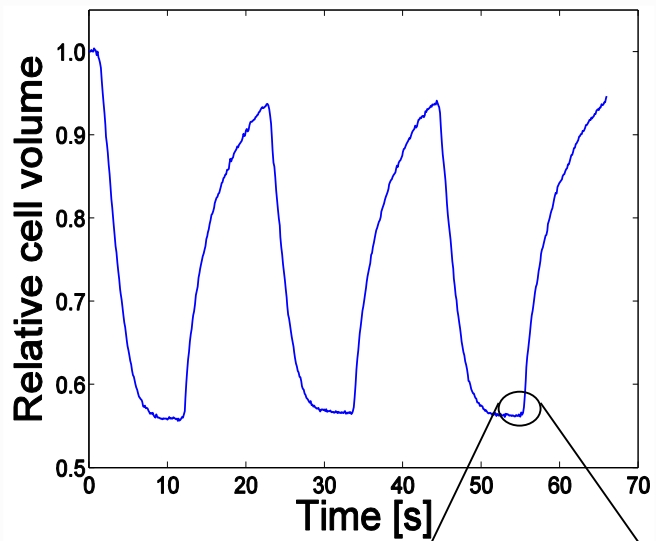
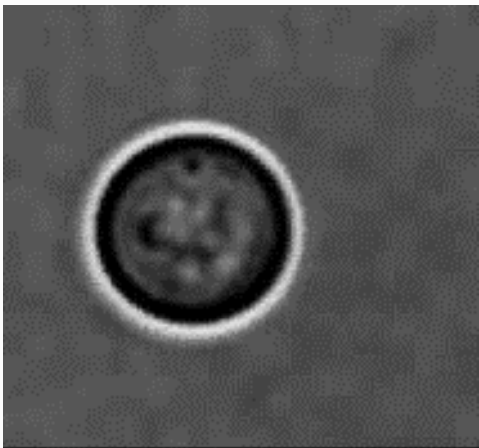
Salt stress of a single yeast cell



Salt stress of a single yeast cell



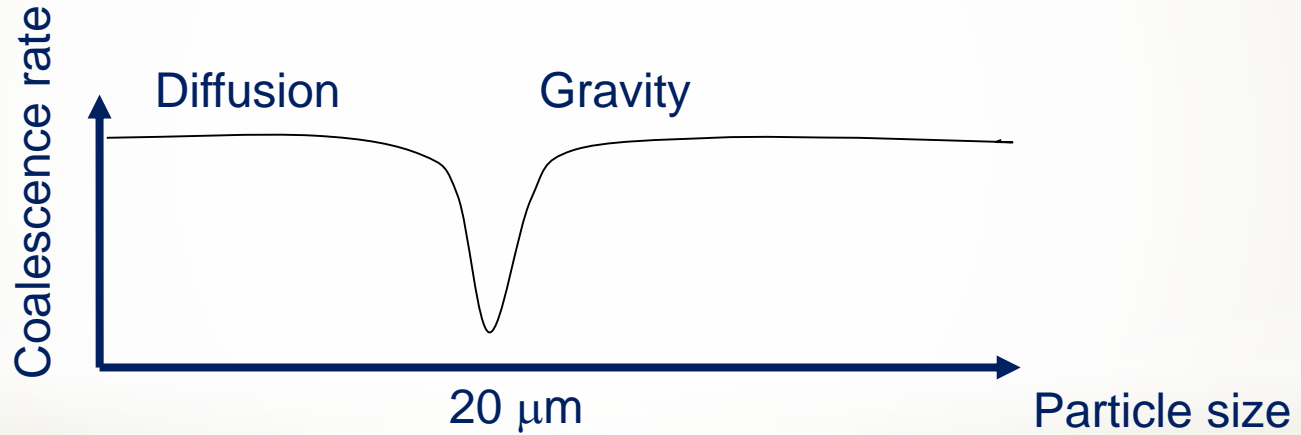
In cooperation with group of Stefan Hohmann
At Cell and Molecular Biology, Göteborg University



How are raindrops created?

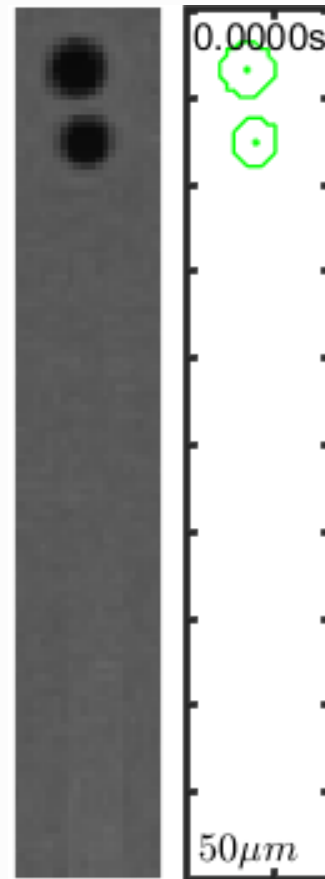
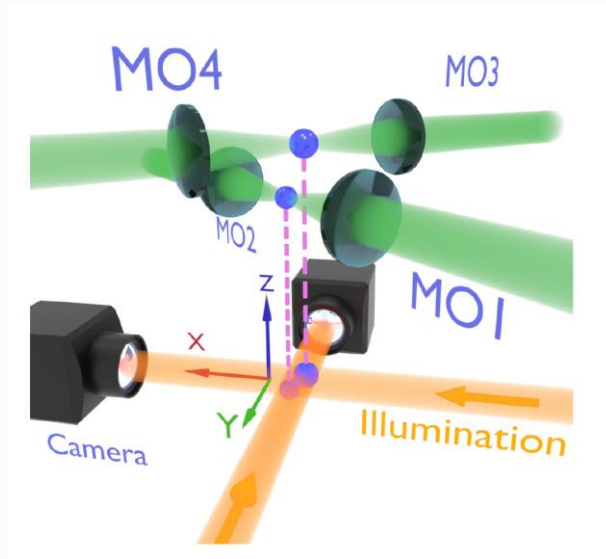
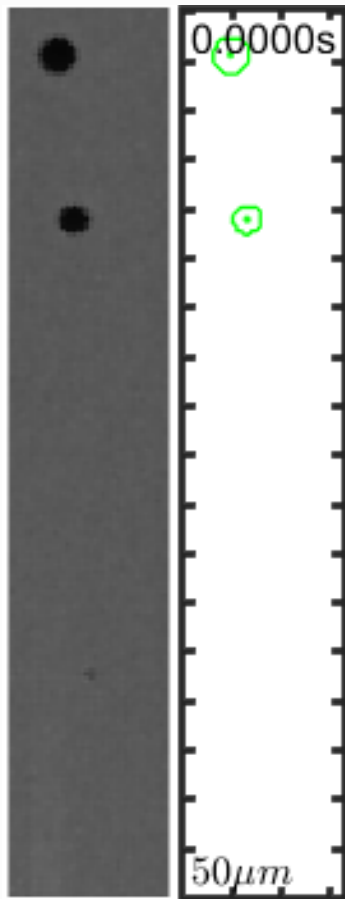


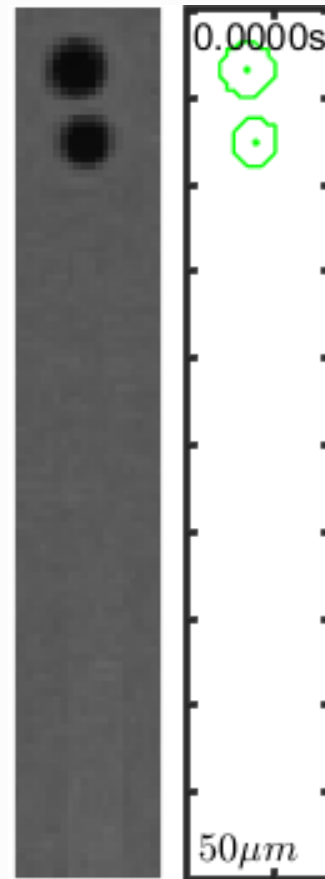
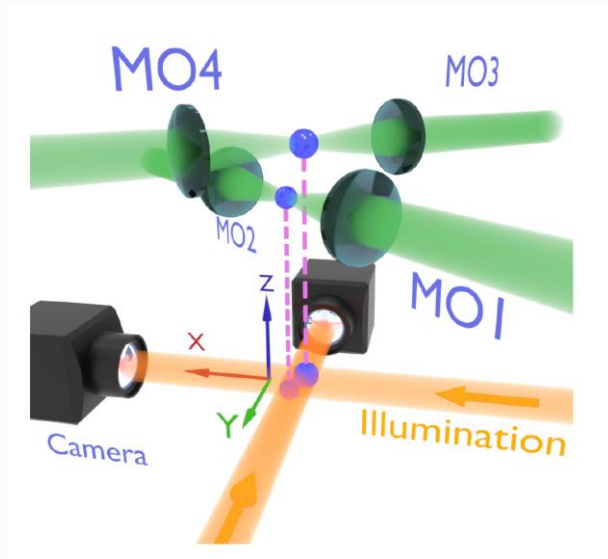
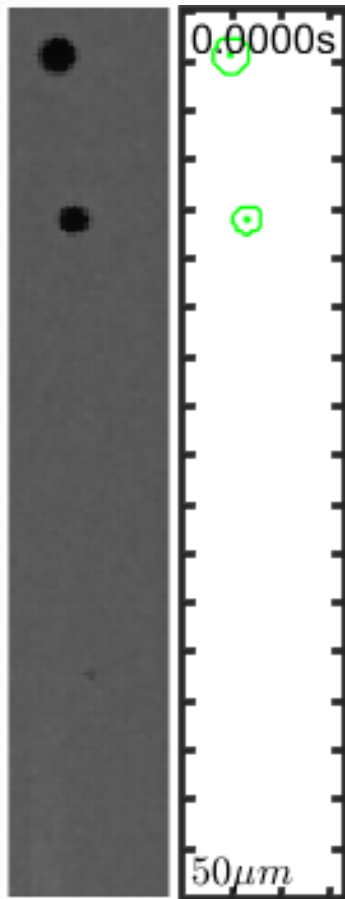
The bottleneck problem:

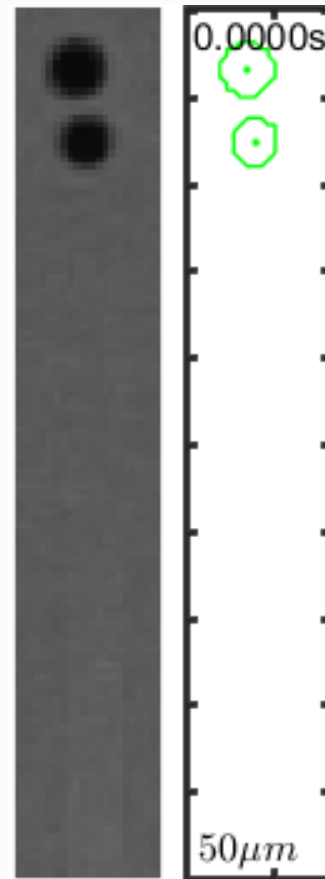
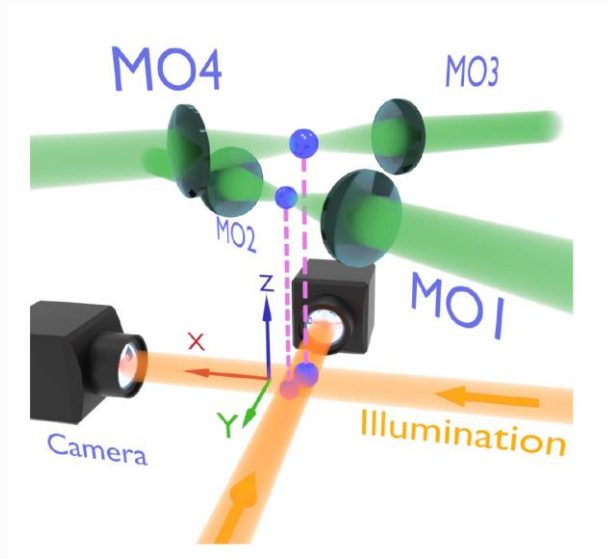
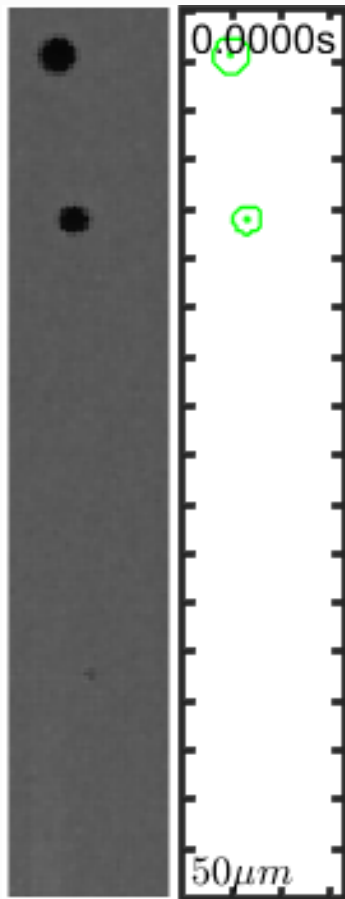


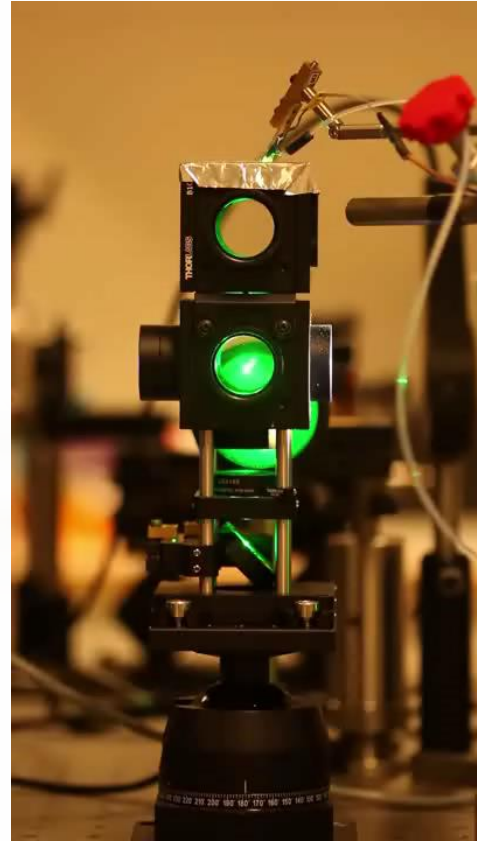
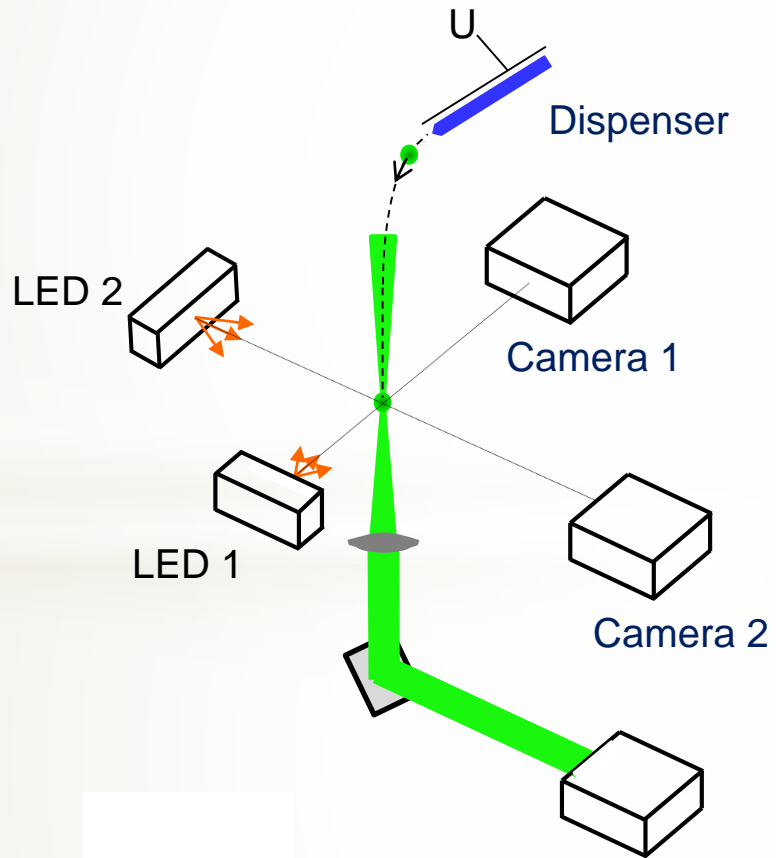
A similar problems occurs in planet formation

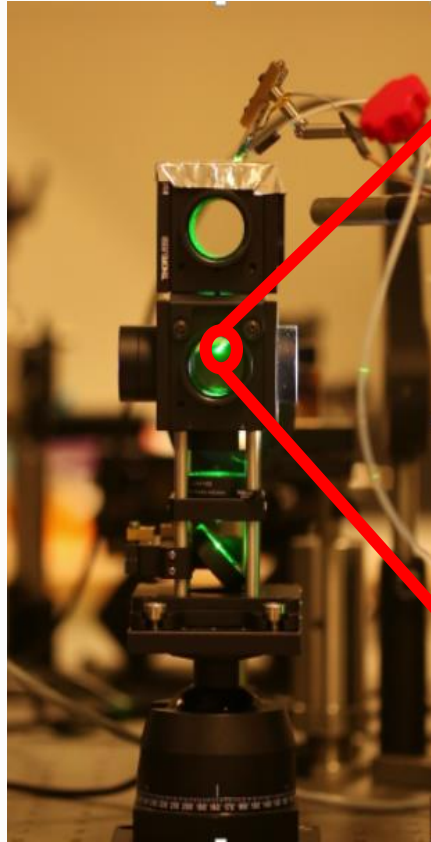
Research project "Bottlenecks for particle growth in turbulent aerosols" from the Knut and Alice Wallenberg Foundation, coordinated by Bernhard Mehlig



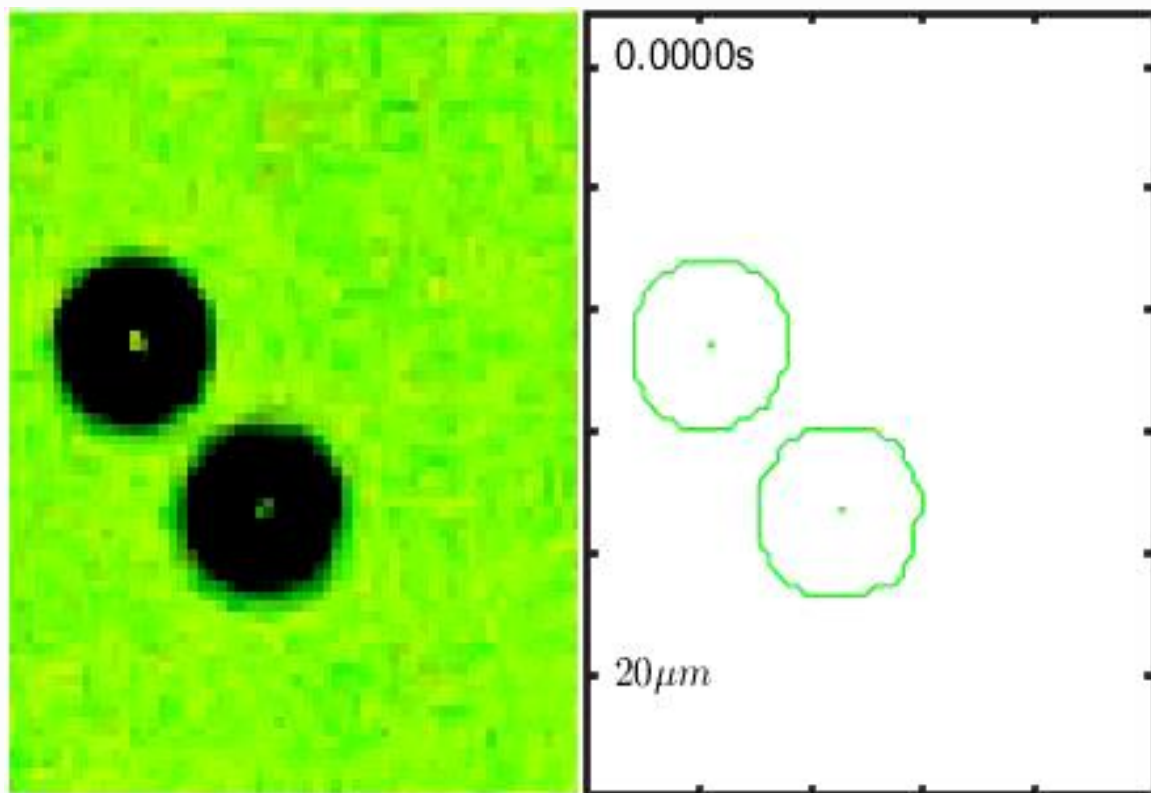


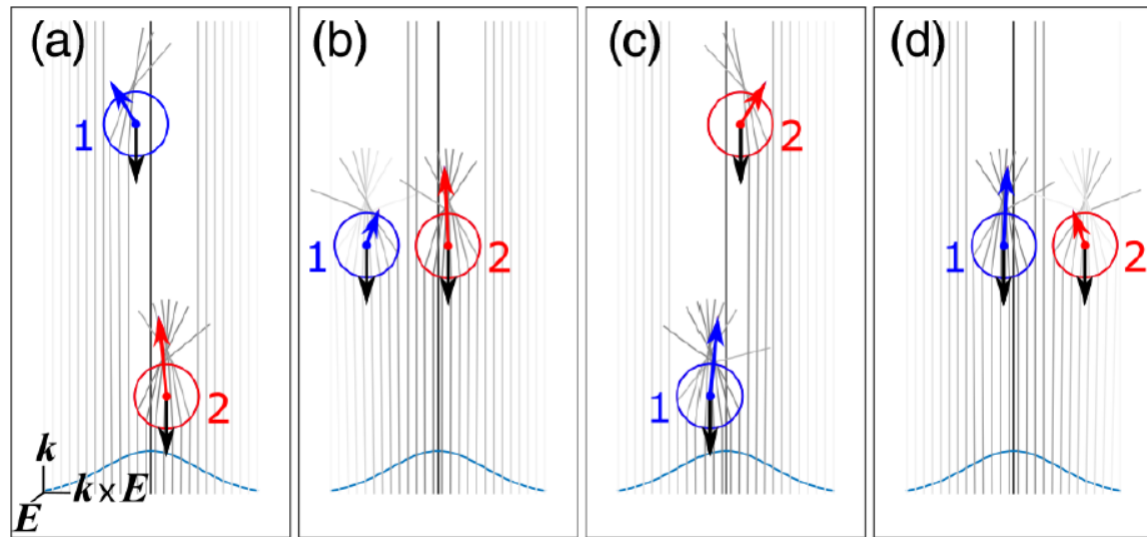






$\bar{30} \mu\text{m}$
600mW laser





NUMERICAL MODEL

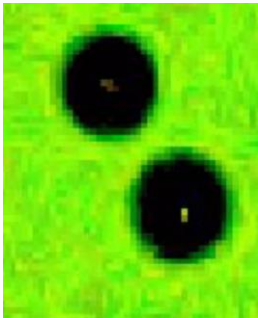
- Gravitation
- Hydrodynamic forces (Stoke)
- Coulumb forces
- Optical forces
 - Ray optics regime
 - Fresnels law´s
 - Momentum of photon

$$m \frac{dv^{(i)}}{dt} = F_G^{(i)} + F_H^{(i)} + F_Q^{(i)} + F_O^{(i)}, \quad (i = 1, 2).$$

$$F_G^{(i)} = -\frac{1}{6} \pi \rho g D^3 \hat{k},$$

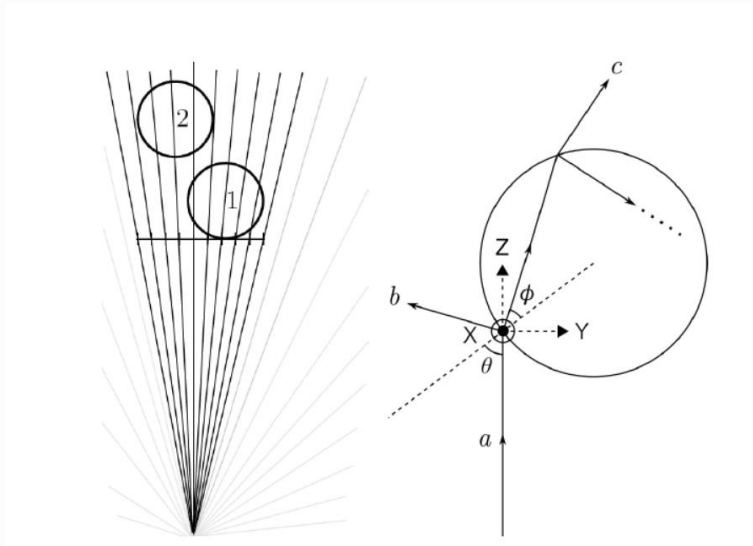
$$F_H^{(i)} = 3\pi\eta D \left[-v^{(i)} + \sum_{j \neq i}^2 \frac{3D}{8r_{ij}} \left(I + \frac{r_{ij} r_{ij}}{r_{ij}^2} \right) \cdot v^{(j)} \right]$$

$$F_Q^{(i)} = \sum_{j \neq i}^2 \frac{F_Q r_{ij}}{r_{ij}} \left[\frac{D^2}{r_{ij}^2} \right].$$

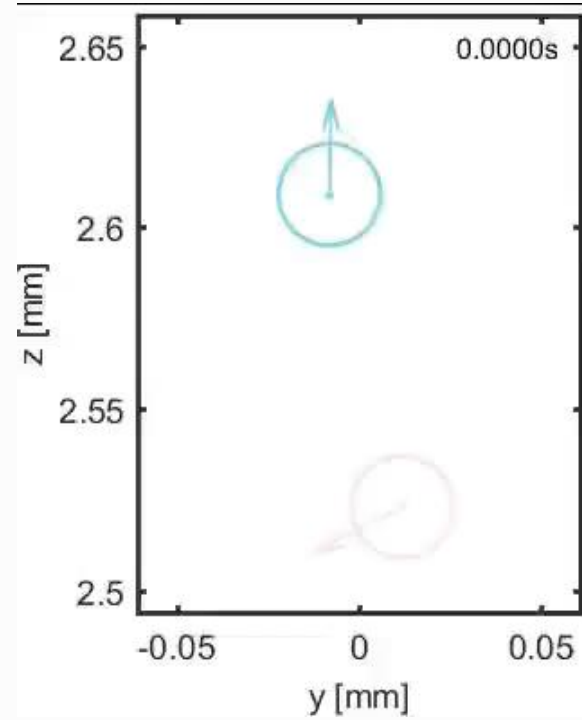
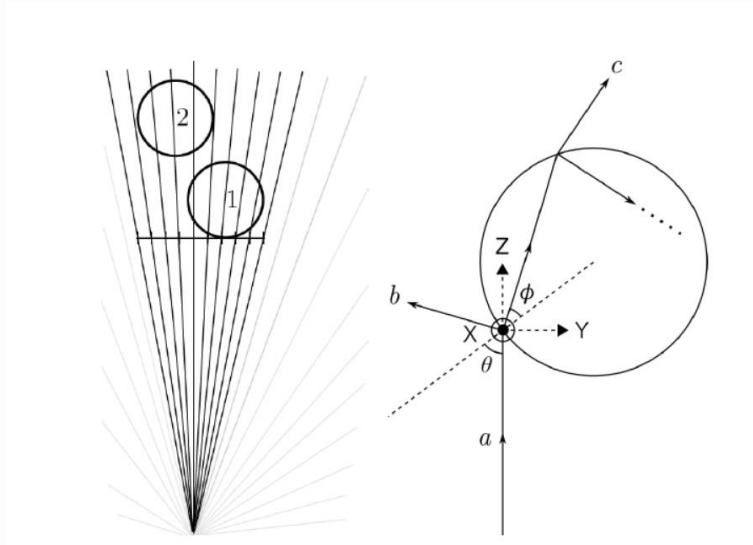


Forces	Expression	Magnitude (N)
Optical	$F_O = qPD^2/(cw^2)$	3.3×10^{-10}
Gravity	$F_G = \pi\rho g D^3/6$	1.4×10^{-10}
Hydrodynamic	$F_H = 9\pi\eta Dv_T/8$	5.3×10^{-11}
Electrostatic	$F_Q = Q^2/(4\pi\epsilon D^2)$	2.9×10^{-11}

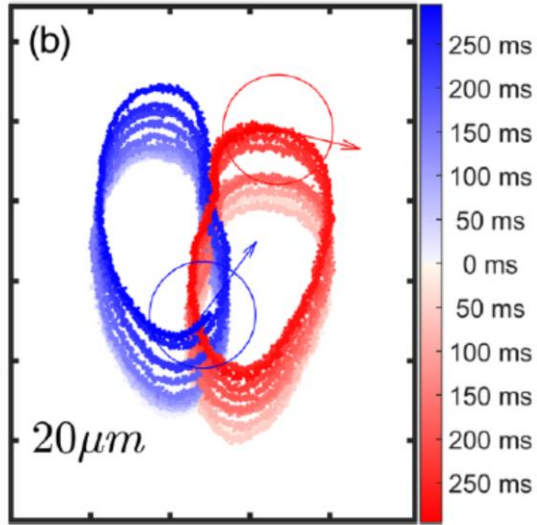
NUMERICAL MODEL



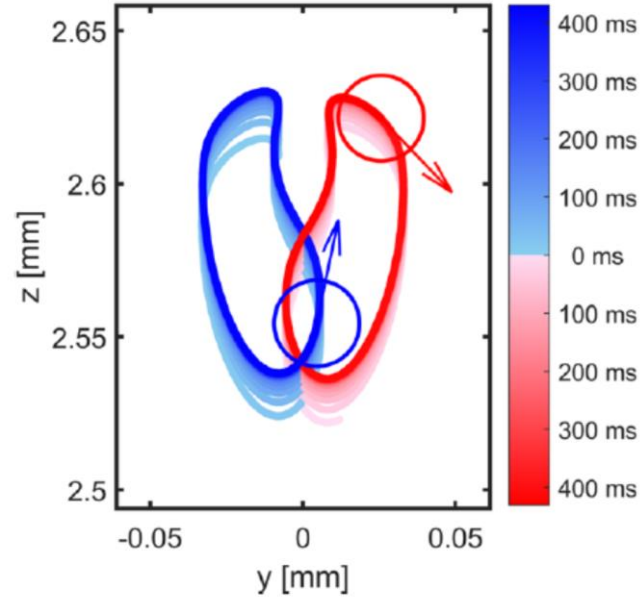
NUMERICAL MODEL



EXPERIMENT



MODEL



Juggling with Light

Albert J. Bae*

Max Planck Institute for Dynamics and Self-Organization, 37077 Goettingen, Germany

Dag Hanstorp and Kelken Chang†

Department of Physics, University of Gothenburg, 412 96 Gothenburg, Sweden

 (Received 11 October 2018; published 1 February 2019)



Juggling with Light

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Max Planck Institute for Dynamics and Self-Organization, 37077 Goettingen, Germany

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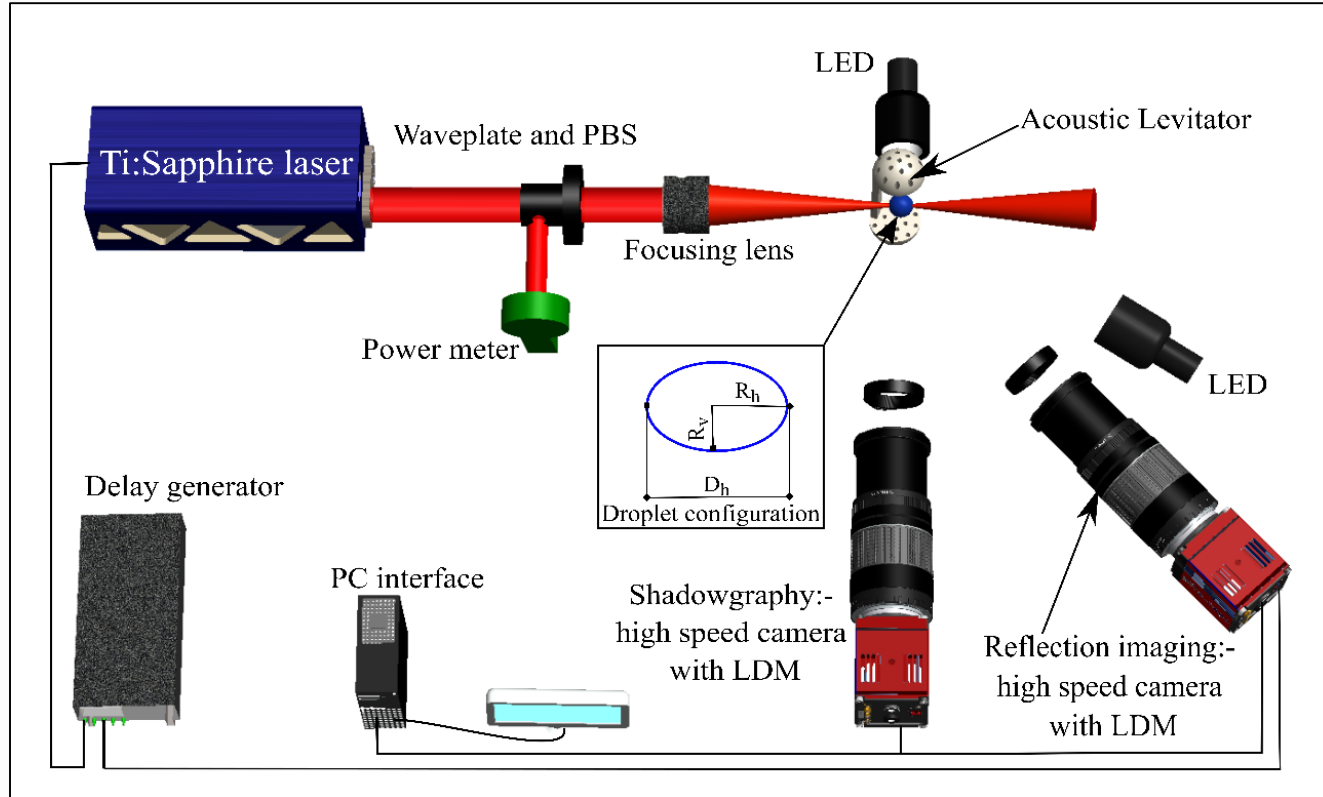
Department of Physics, University of Gothenburg, 412 96 Gothenburg, Sweden

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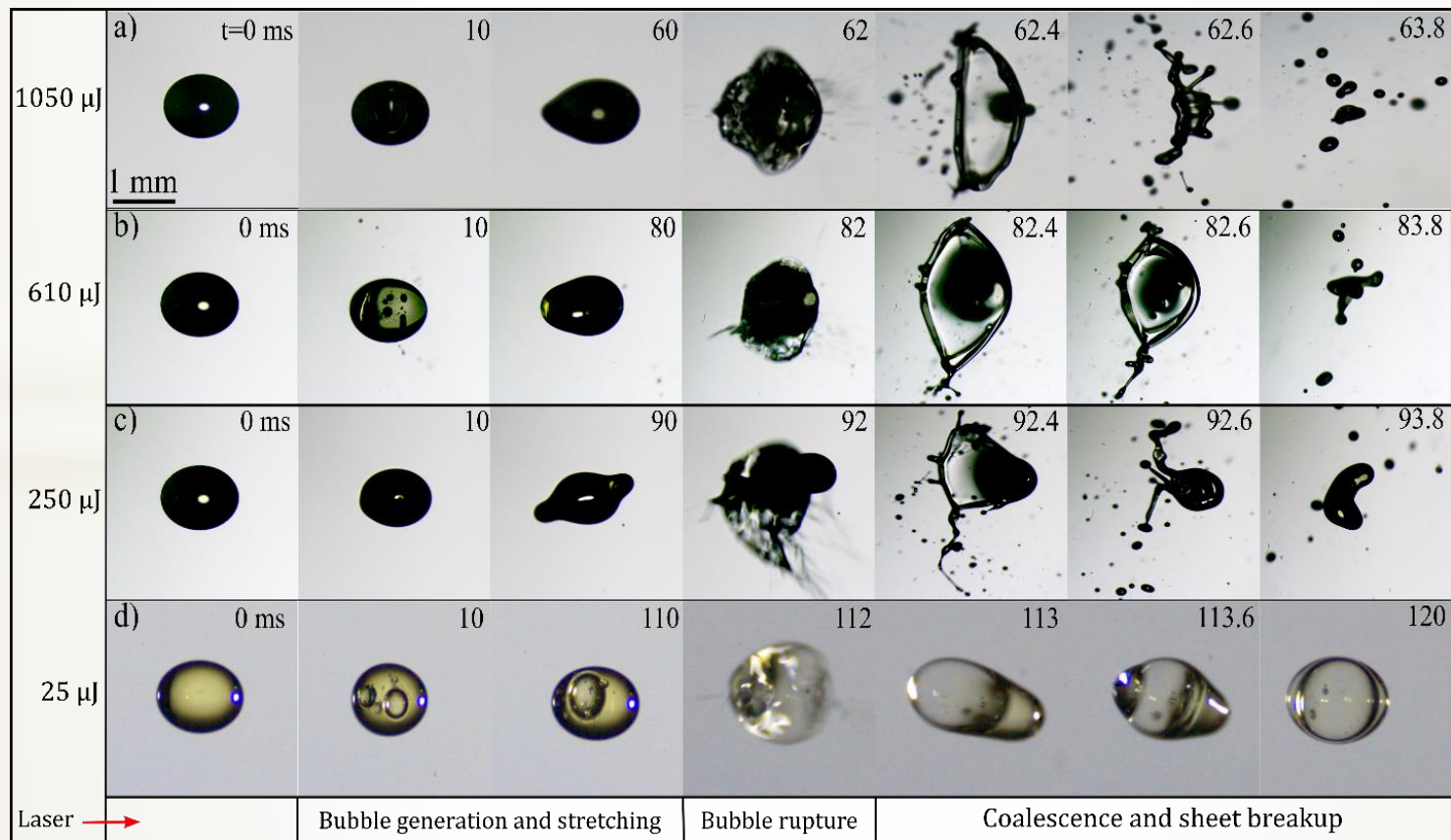
Bubble dynamics and atomization of acoustically levitated biofuel droplets using femtosecond laser pulses

Vishal S. Jagadale¹, Devendra Deshmukh¹, Dag Hanstorp^{2, *} & Yogeshwar Nath Mishra^{1,2,3}



Bubble dynamics and atomization of acoustically levitated biofuel droplets using femtosecond laser pulses

Vishal S. Jagadale¹, Devendra Deshmukh¹, Dag Hanstorp^{2, *} & Yogeshwar Nath Mishra^{1,2,3}



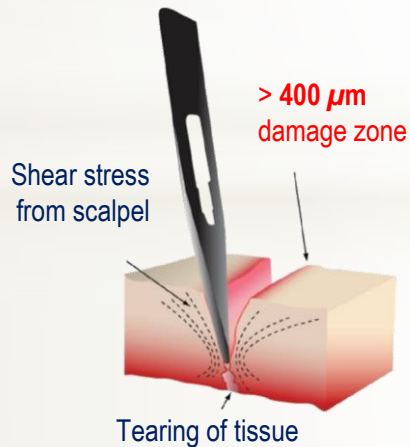
Femtosecond laser bone sectioning

Master project, Department of Physics

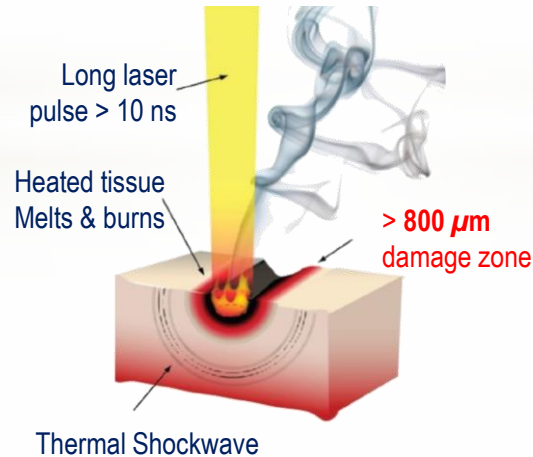
Sahlgrenska hospital + GU physics

Motivation: Sectioning of bones in osteology to access the bone marrow. This is of great interest in medical researcher e.g. in investigations of leukemia for an analysis of cells in the bone marrow using optical methods.

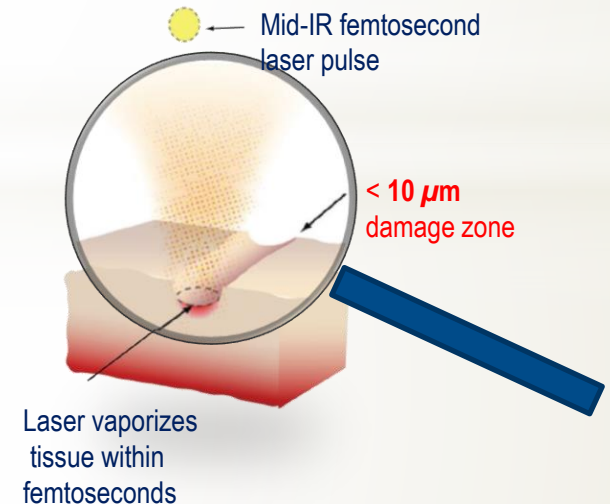
Today: a) Surgical Scalpel



b) Conventional Medical Laser



Solution: Femtosecond laser sectioning



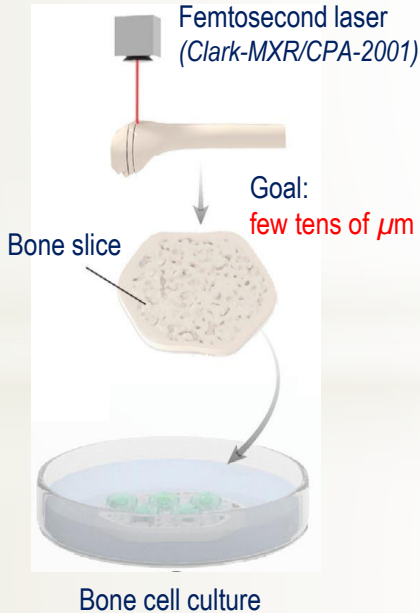
☀ Almost impossible for fresh bones (living cells cannot be preserved)

Femtosecond laser bone sectioning

Master project, Department of Physics

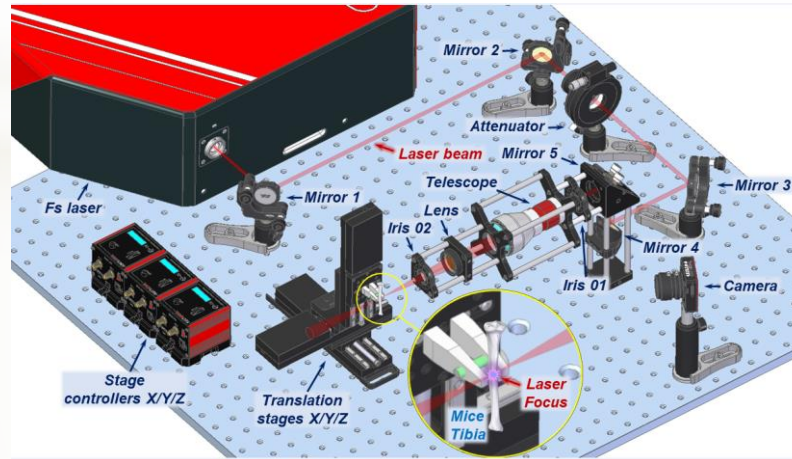
Sahlgrenska hospital + GU physics

Concept:



Project goal: Build a femtosecond laser bone sectioning platform at Laserlab Göteborg that can produce

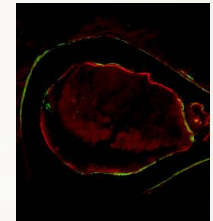
- A) Smooth surfaces
- B) Thin slices (tens of micrometers, i.e. a fraction of the thickness of a human hair)



Preliminary results: Mice bone microtome sections with trabecular bones



300 μm slice
PFA-fixed tibia



100 μm slice
Pre-stained bone

Number of students:

- One in physics
- Two in physics
- One in physics and one in medicine

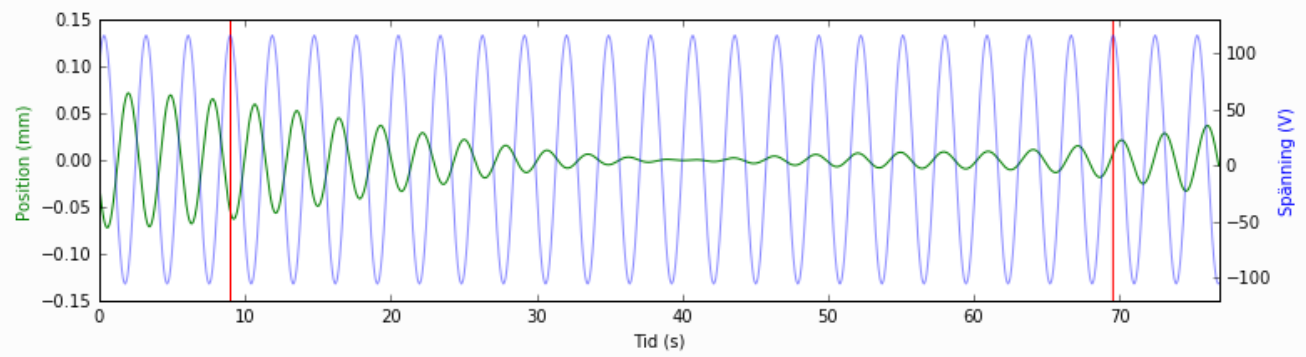
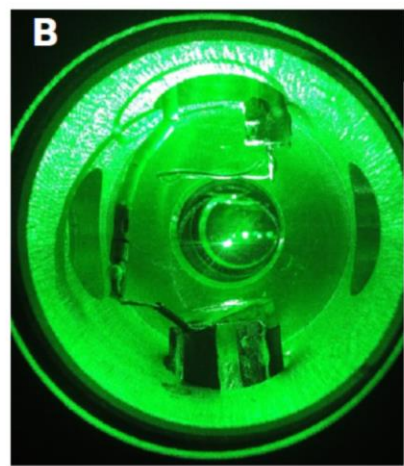
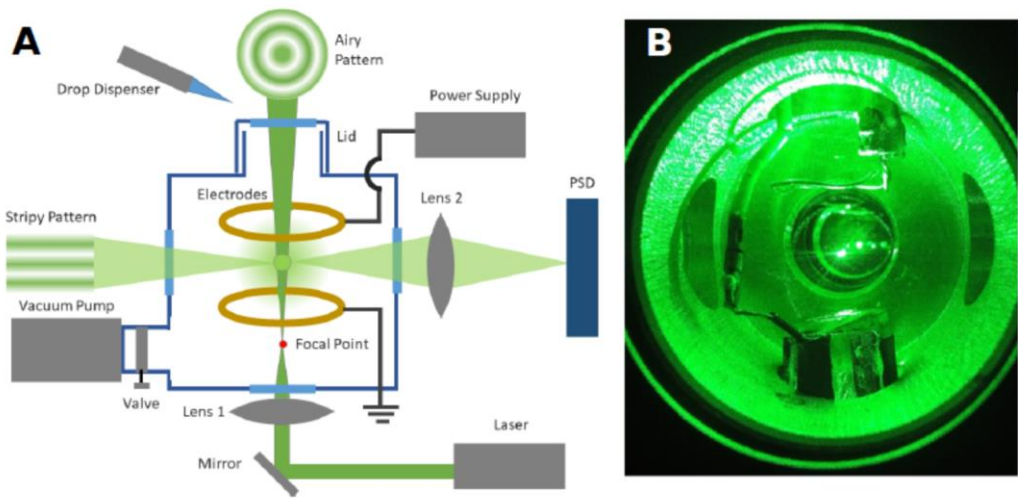
Prior knowledge:

- Optics
- Mechanics
- Electronics/computer control/programming
- Physics
- or Medicine

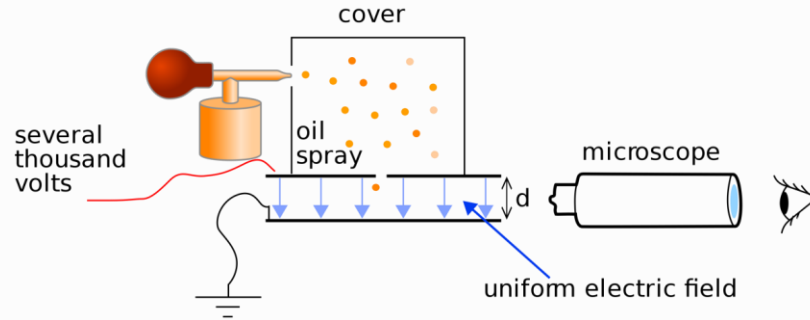
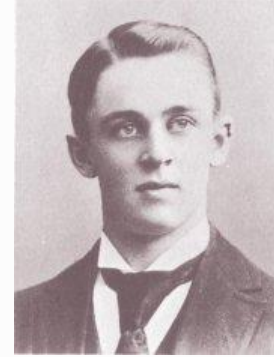
Contacts:

Prof. Dag Hanstorp (dag.hanstorp@gu.se)

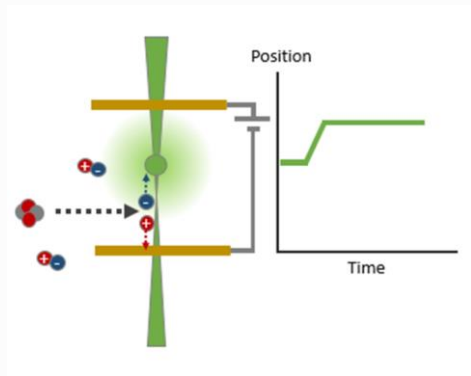
Dr. Di Lu (di.lu@physics.gu.se)

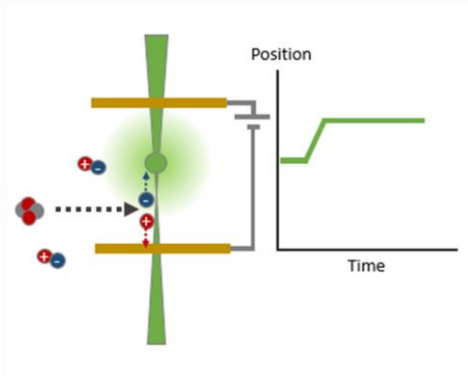
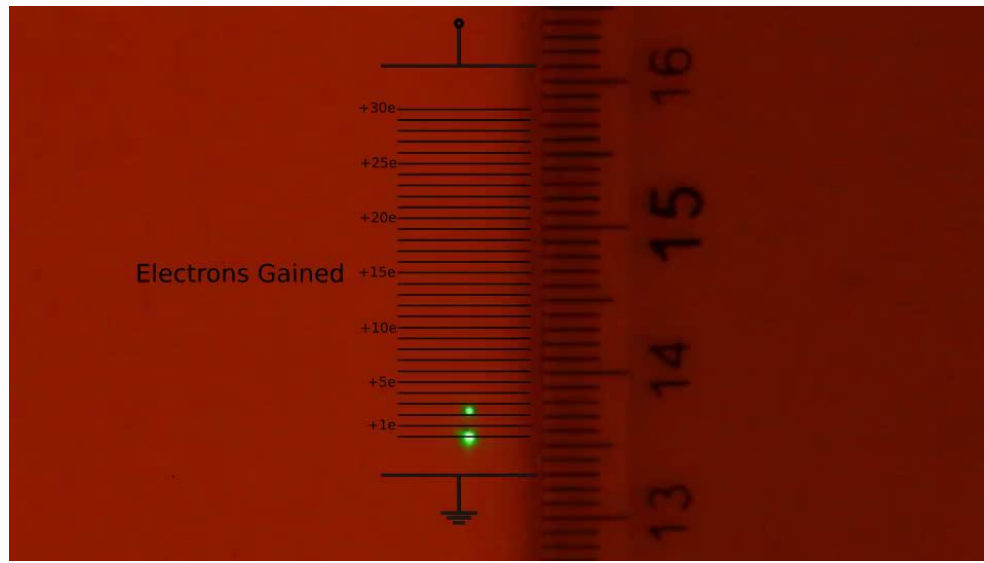


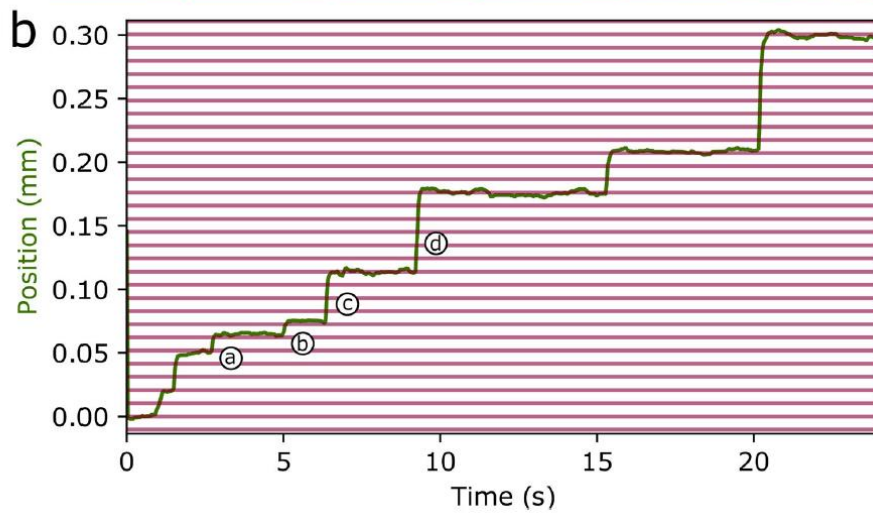
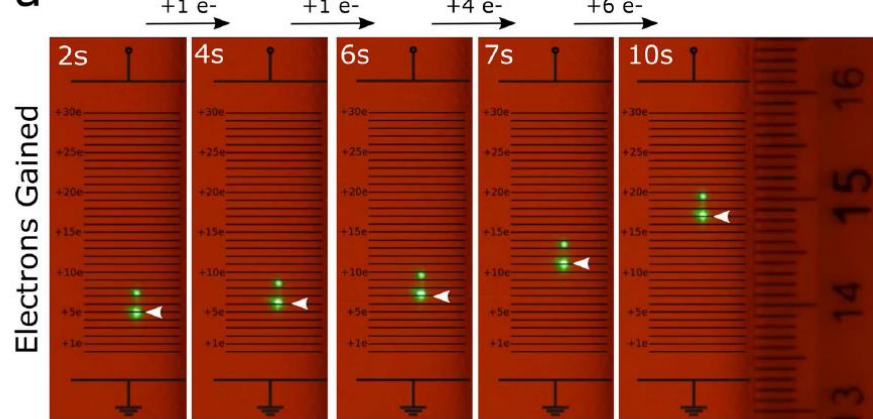
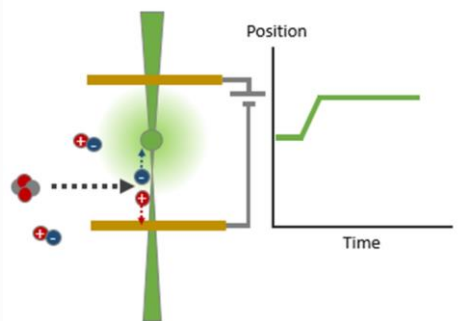
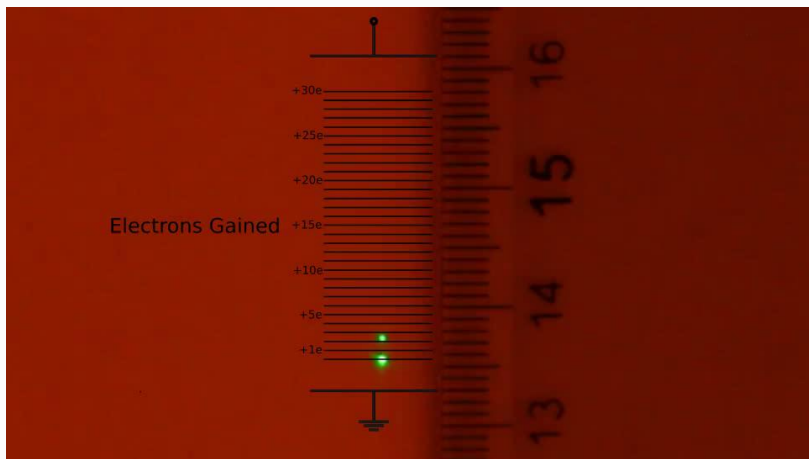
Millikan's experiment



On the Elementary Electric charge and the Avogadro







ACKNOWLEDGEMENT





Paldies parjūsu uzmanību!

Conclusion and outlook

- High resolution EA measurements → Beamtime approved at DESIREE for Si⁻ in September (Julia Karls)
- Isotope shifts → Beamtime approved at CERN for studies of a IS in the chain ³⁸Cl- ⁴³Cl
- Lifetimes studies → Beamtimes at DESIREE approved for studies of Th⁻ and Sn⁻
- EA of Radioactive elements → Beamtime for Po⁻ approved at CERN

Acknowledgement

nature communications



Article

<https://doi.org/10.1038/s41467-022-33438-y>

High-precision electron affinity of oxygen

Received: 10 May 2022

Accepted: 19 September 2022

Published online: 07 October 2022

Check for updates

Moa K. Kristiansson¹, Kiattichart Chartkunchand^{1,2}, Gustav Eklund¹, Odd M. Hole¹, Emma K. Anderson³, Nathalie de Ruelle¹, Magdalena Kamińska¹, Najeeb Punnakayathil¹, José E. Navarro-Navarrete¹, Stefan Sigurdsson¹, Jon Grumer⁴, Ansgar Simonsson¹, Mikael Björkhage¹, Stefan Rosén¹, Peter Reinhed¹, Mikael Blom¹, Anders Källberg¹, John D. Alexander¹, Henrik Cederquist¹, Henning Zettergren¹, Henning T. Schmidt¹ & Dag Hanstorp⁵



ARTICLE

<https://doi.org/10.1038/s41467-020-17599-2> OPEN

Check for updates

The electron affinity of astatine

David Leimbach^{1,2,3,9}, Julia Karls², Yangyang Guo⁴, Rizwan Ahmed⁵, Jochen Ballof^{1,6}, Lars Bengtsson⁷, Ferran Boix Pamies¹, Anastasia Borschevsky⁴, Katerina Chrysalidis^{1,3}, Ephraim Eliav⁷, Dmitry Fedorov⁸, Valentin Fodosseev¹, Oliver Forstner^{9,10}, Nicolas Galland¹¹, Ronald Fernando Garcia Ruiz^{11,2}, Camilo Granados¹, Reinhard Heinke³, Karl Johnston¹, Agota Koszorus¹³, Ulli Köster¹⁴, Moa K. Kristiansson¹⁵, Yuan Liu¹⁶, Bruce Marsh¹, Pavel Molkanov⁸, Lukáš F. Pašteka¹⁷, João Pedro Ramos²⁰, Eric Renaut¹¹, Mikael Reponen¹⁸, Annie Ringvall-Moberg¹², Ralf Erik Rossel¹, Dominik Studer³, Adam Vernon¹⁹, Jessica Warbinek^{2,3}, Jakob Welander², Klaus Wendt³, Shane Wilkins¹, Dag Hanstorp² & Sebastian Rothe¹



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Thank you!