



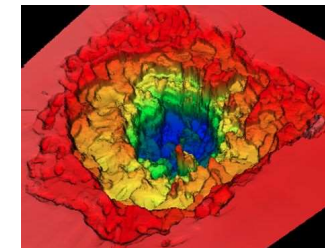
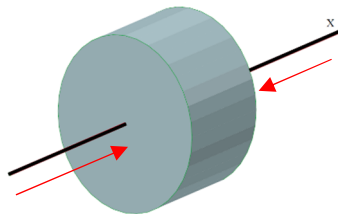
# ON HIGH FIELD PLASMONICS AND A POTENTIAL NUCLEAR FUSION APPLICATION

NORBERT KROO

(on behalf of the NAPLIFE project)

WIGNER PHYSICS RESEARCH CENTER and  
HUNGARIAN ACADEMY OF SCIENCES

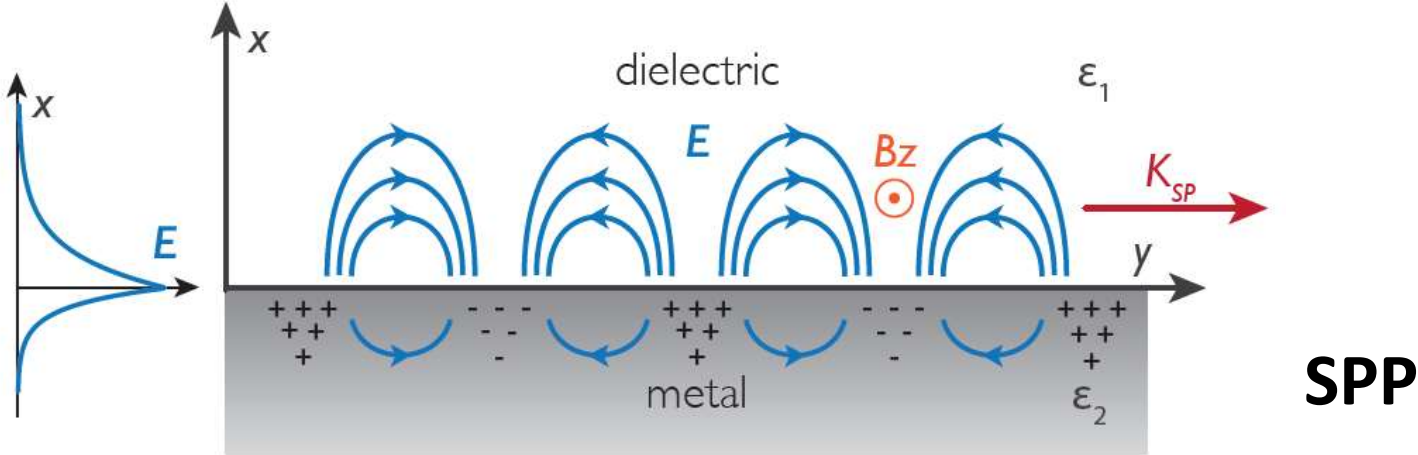
*Motto: Only those, who are prepared to go  
too far, can know how far they may go.*



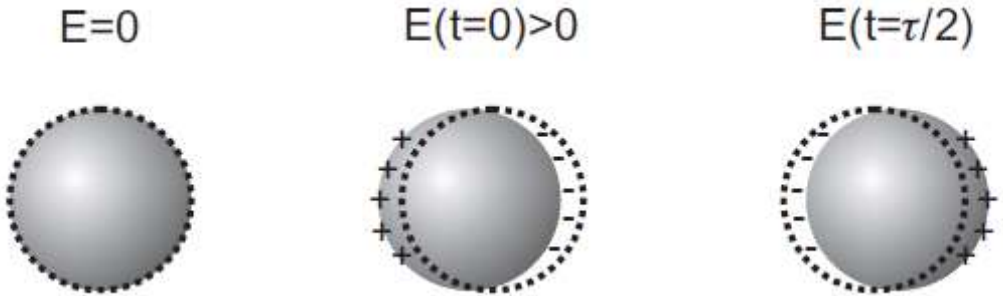
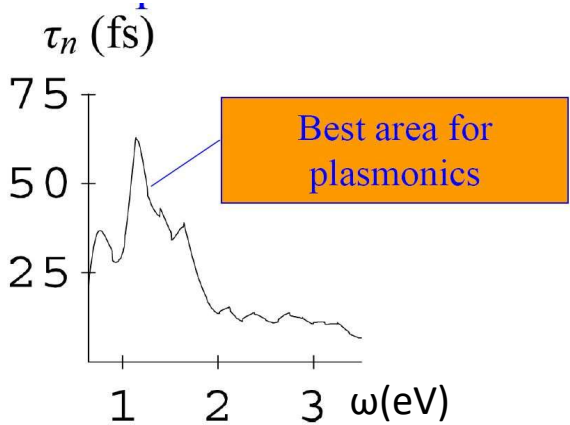
CASPER, 07.21.2023

# PLASMONICS AND HIGH FIELDS APPLICATIONS

Special type of the optical near field



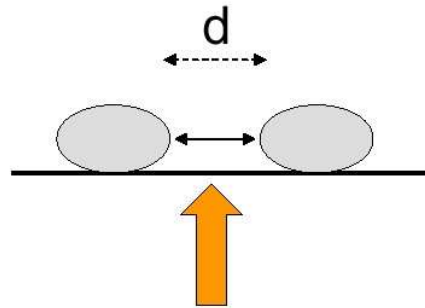
**SPP**



**LSP**

Ti:Sa laser:  $\lambda=800\text{nm}$  ( $\sim 1.55\text{eV}$ ) ;  $t_{(SPP)} \sim 20\text{fs}$

# LSPP



$$d \ll \lambda$$

“Near-field coupling”

⇒ Resonator coupling

$$d \gg \lambda$$

“Dipole-dipole coupling”

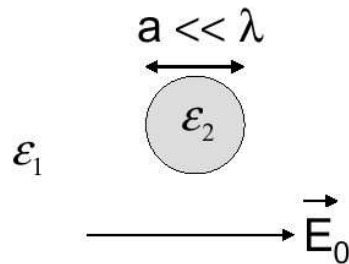
⇒ Interferences

If  $a \ll \lambda \Rightarrow$  dipole :

$$p = \epsilon_2 \alpha E_0$$

with:

$$\alpha = 4\pi a^3 \frac{\epsilon_2 - \epsilon_1}{\epsilon_2 + 2\epsilon_1}$$



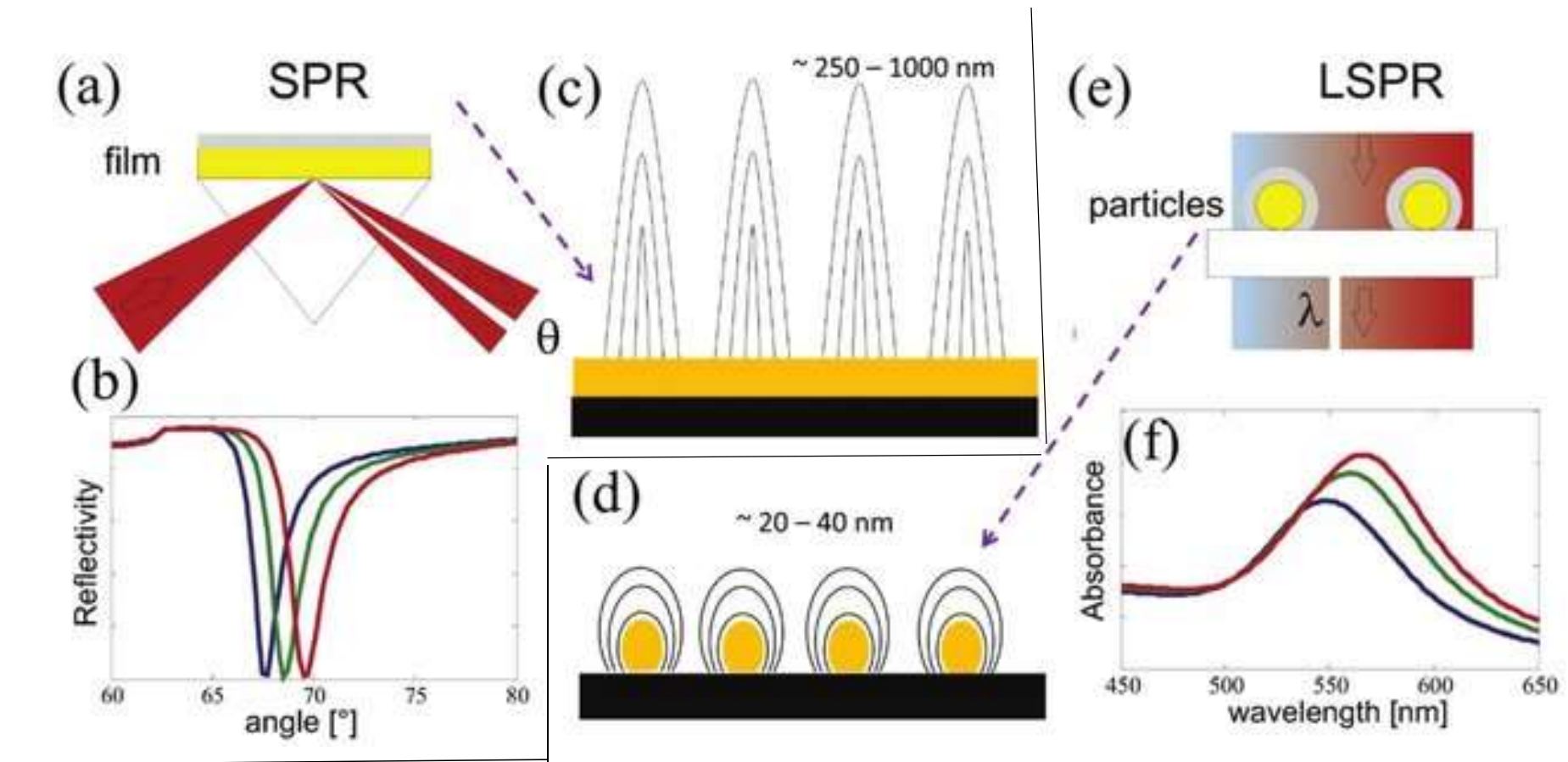
Resonance when  $\epsilon_2(\omega) = -2 \times \epsilon_1(\omega)$

⇒ Enhanced absorption

⇒ Enhancement of the near-field & scattering

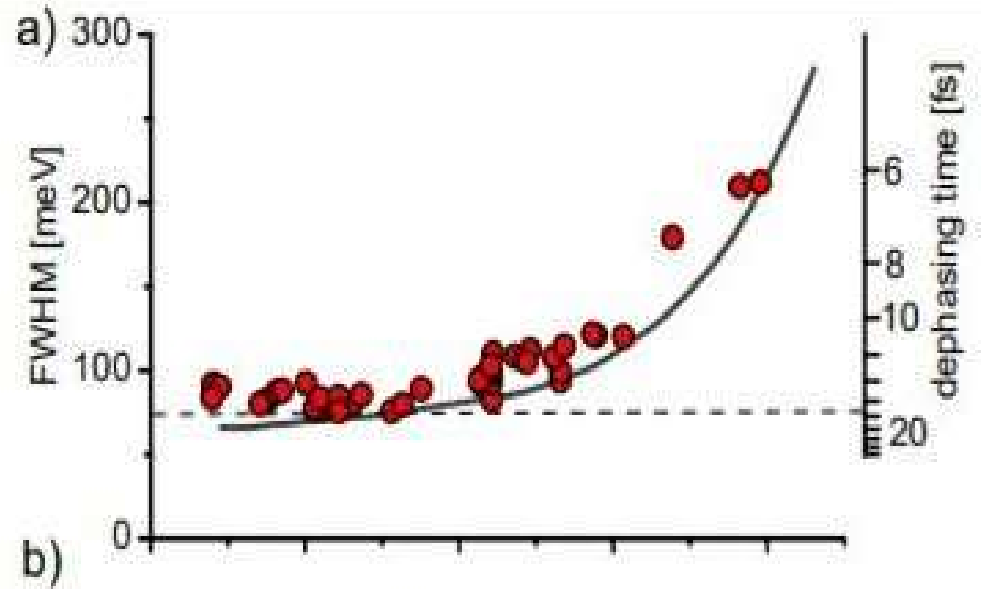
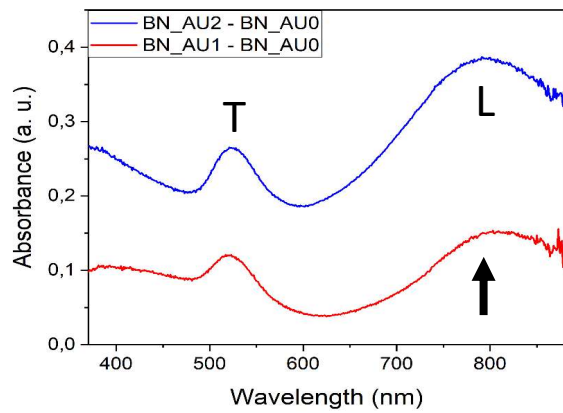
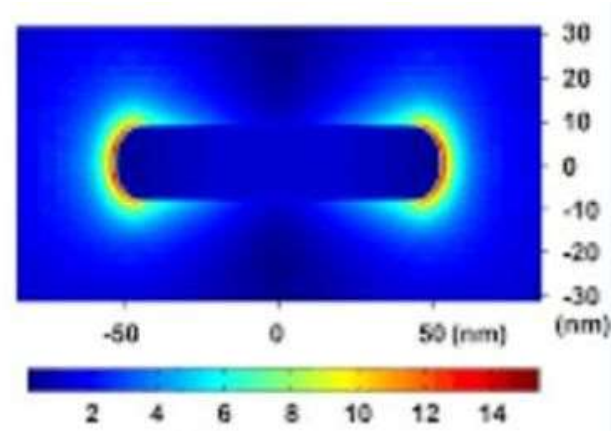
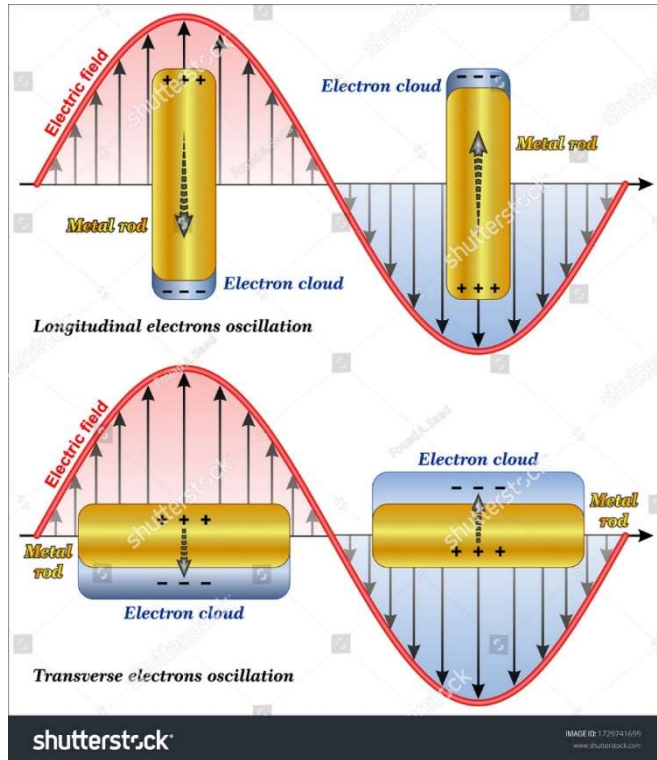
# LOCALIZED PLASMONS (LSPP) UP TO $10^{20}$ W/cm<sup>2</sup>)

(The basic difference between SPP-s and LSPP-s)

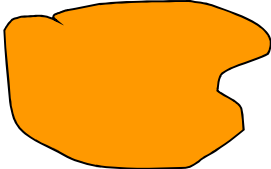



- LSPP: - NO PENETRATION INTO THE PLASMONIC MATERIAL (e.g. metal)  
- SMALLER PENETRATION INTO THE DIELECTRIC /VACUUM  
-NO DISPERSION  
-BROADER RESONANCE

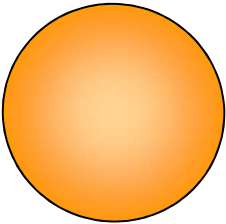
# Nanorod: Transverse and longitudinal modes!

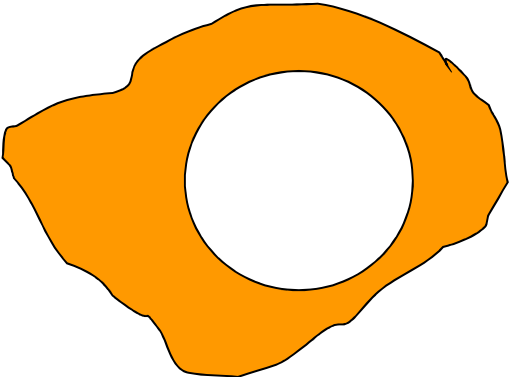


# PLASMONREONANCE OF NANOPARTICLES SHAPE AND- ÉS SIZEDEPENDENT!

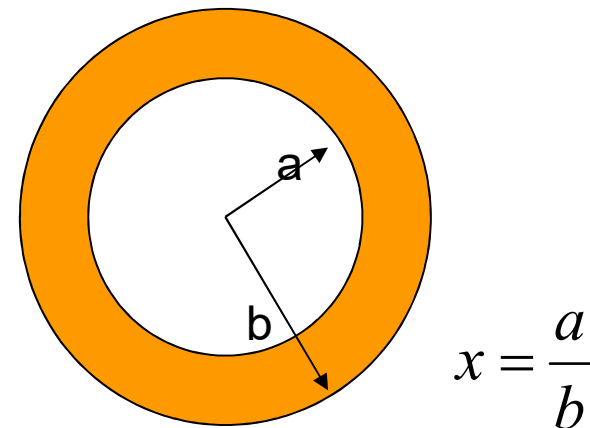
$\omega_B = \sqrt{\frac{4\pi e^2 n}{m_e}}$ 

 Bulk

$\omega_{surf} = \frac{\omega_B}{\sqrt{2}}$ 

 Surface

$\omega_{S,l} = \omega_B \sqrt{\frac{l}{2l+1}}$ 

 Sphere

Üreg
 
 $\omega_{C,l} = \omega_B \sqrt{\frac{l+1}{2l+1}}$

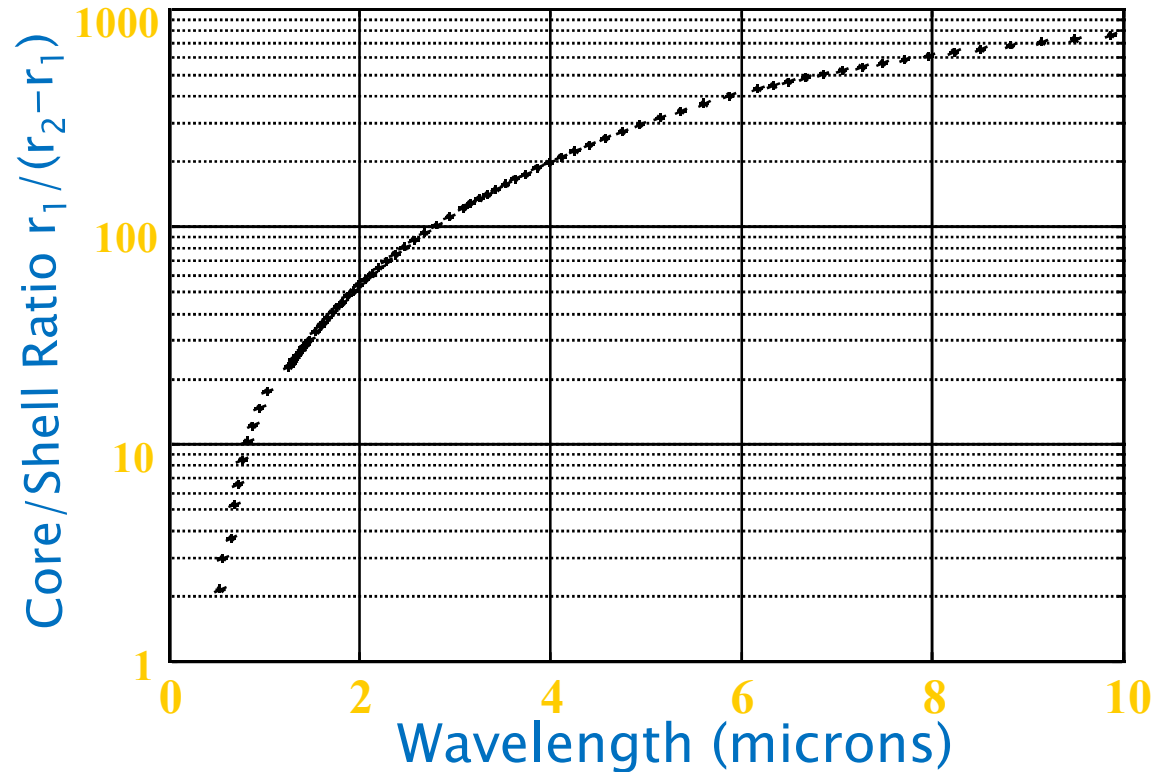
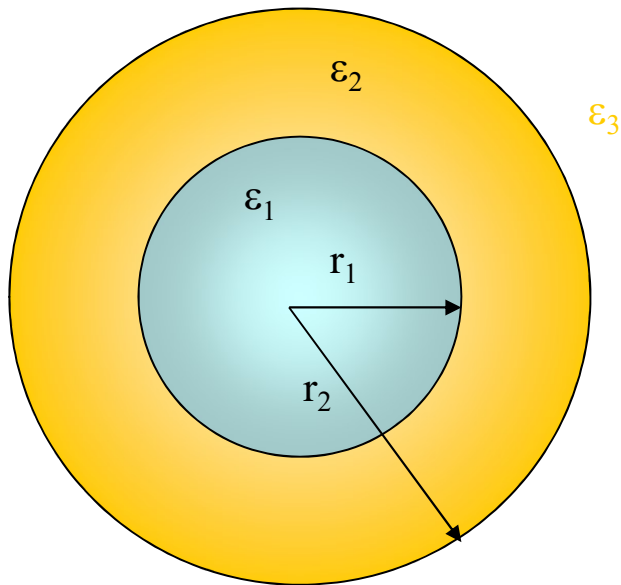
Nanoshell:



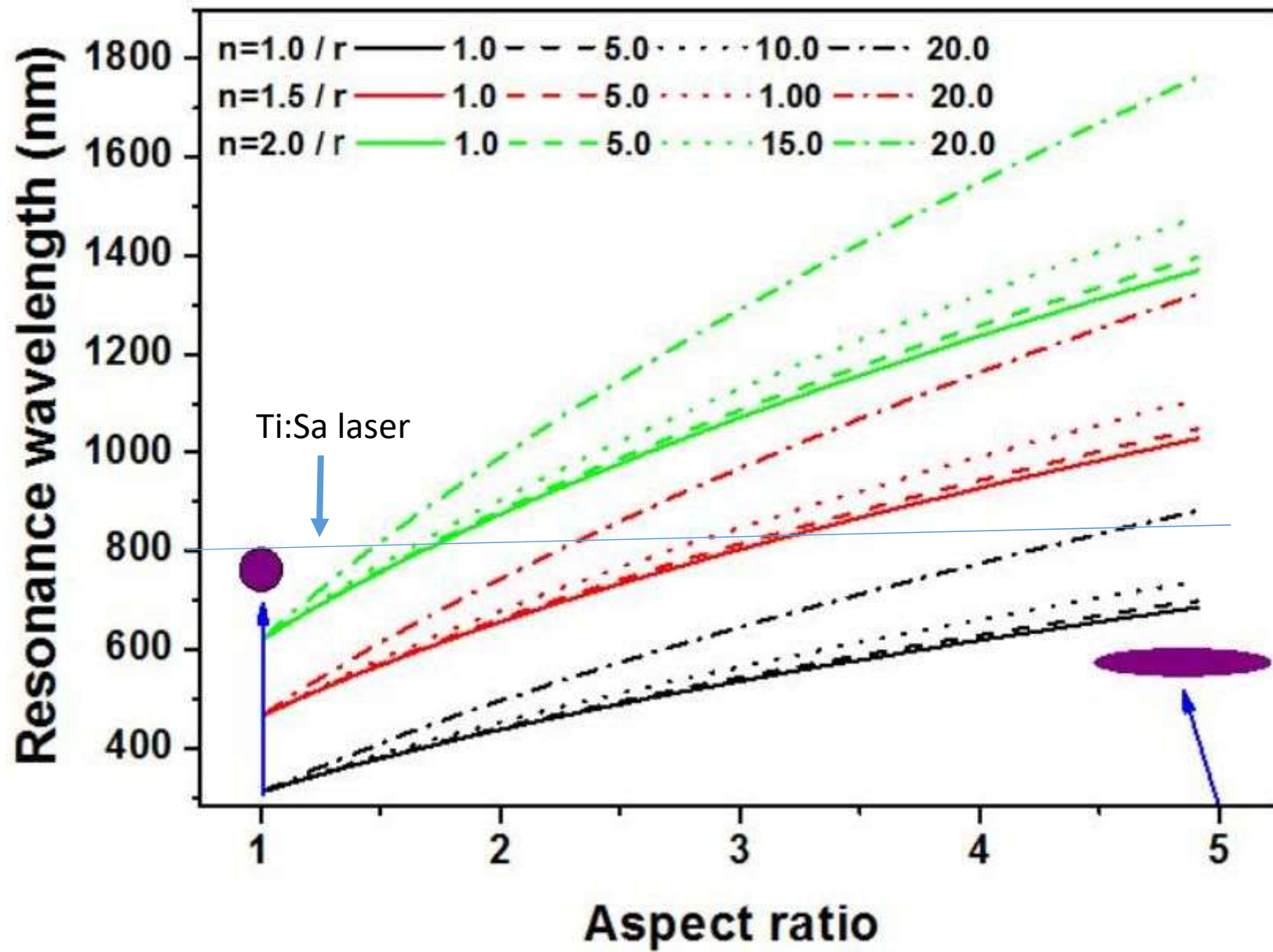
$$\omega_{l\pm}^2 = \frac{\omega_B^2}{2} \left[ 1 \pm \frac{1}{2l+1} \sqrt{1 + 4l(l+1)x^{2l+1}} \right]$$

**Nanoshell plasmon resonance depends on the x ratio .**

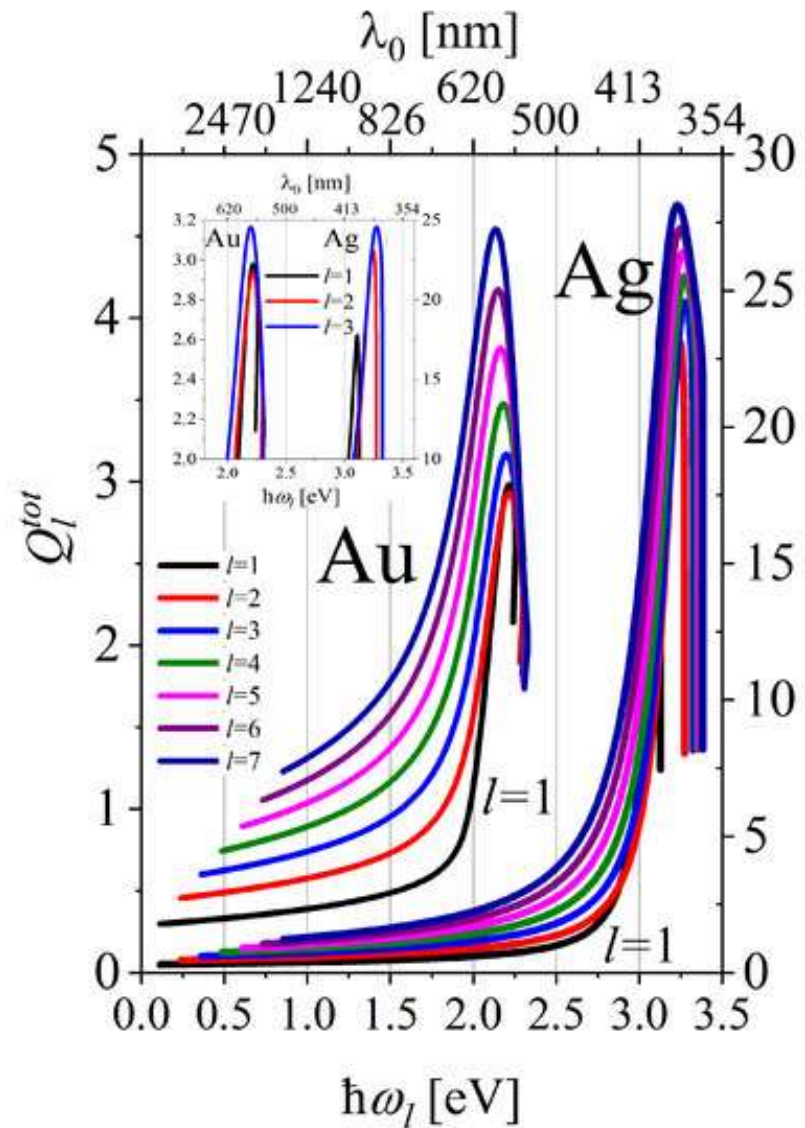
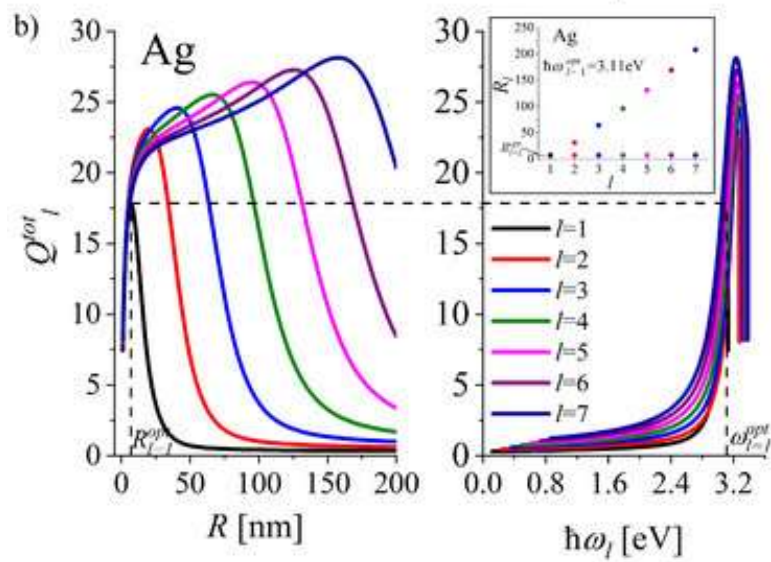
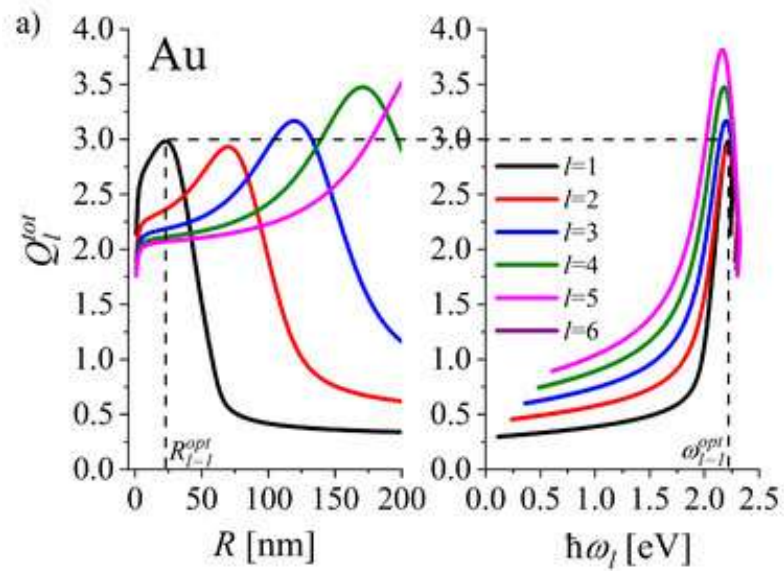
# Spectral tuning range nanoshell plasmon rezonance



1.24 ← electron volts → 0.124  
300 ← THz → 30  
10,000 ←  $\text{cm}^{-1}$  → 1,000



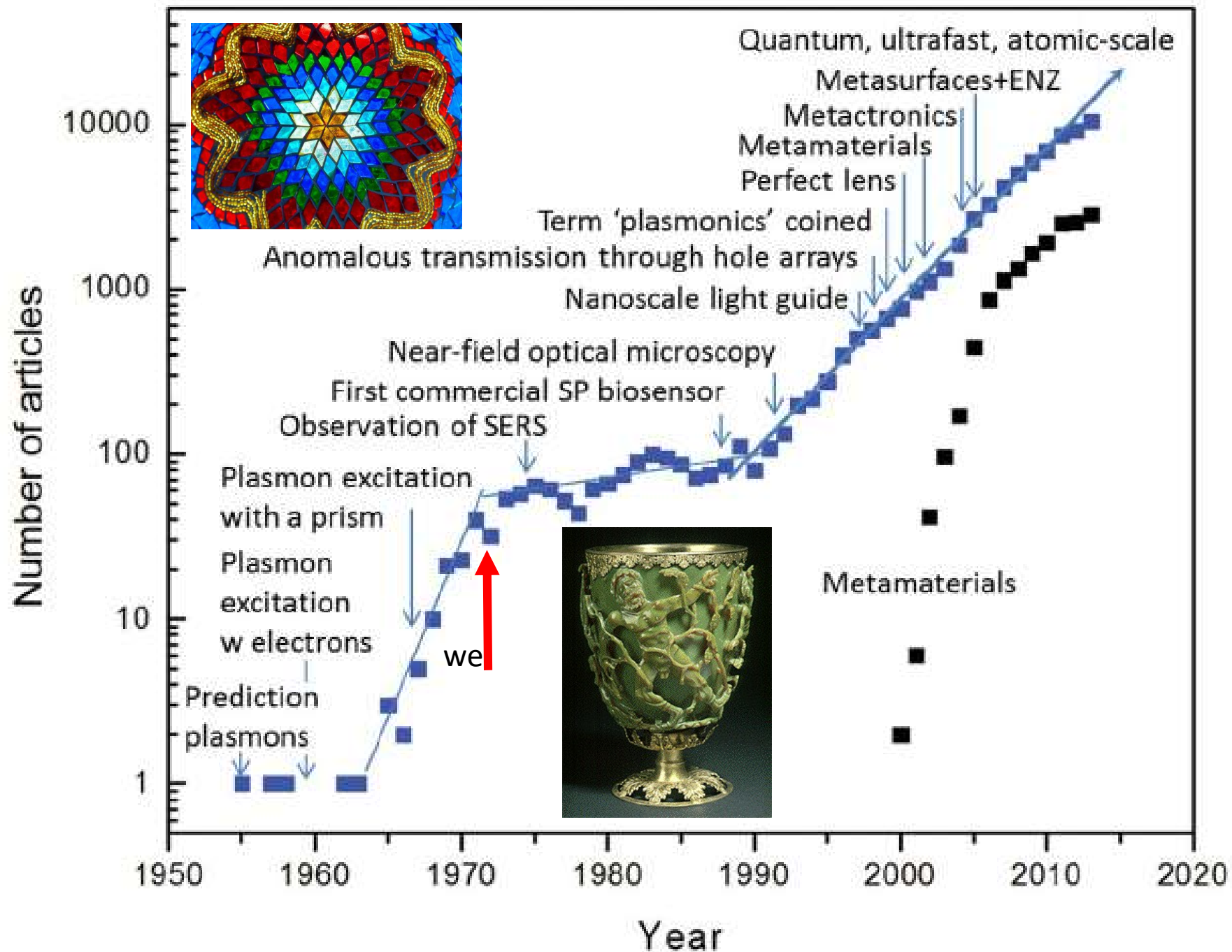






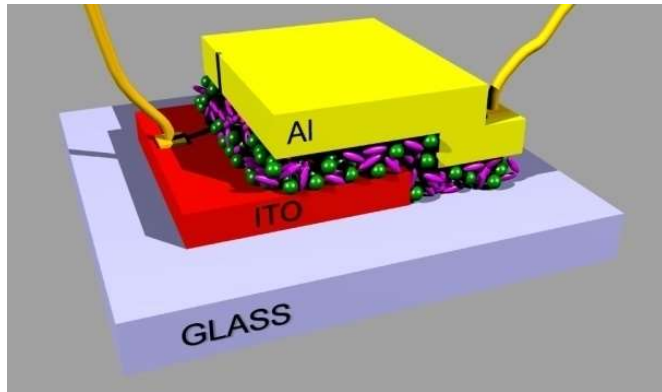
***SPP and LSPP are a „NEW TYPE OF LIGHT“,  
they are***

1. BOUND TO THE (METAL) SURFACE (SPP and LSPP)
2. HAVE SPECIFIC DISPERSION PROPERTIES (LSPP no dispersion)
3. THE DIFFRACTION LIMIT DOES NOT APPLY,
4. CAN BE SQUEEZED TO NANOSIZED VOLUMES
5. SCREEN POSITIVE (e.g. proton) PARTICLE FIELDS
6. IS CORRELATED MOTION OF A HIGH NUMBER OF CONDUCTION  
ELECTRONS
7. COULD REPRESENT VERY HIGH ELECTRIC FIELDS (hot spots)
8. MAY BE THE SOURCE OF DIFFERENT NONLINEAR PROCESSES
9. SHOW NON-CLASSICAL PROPERTIES

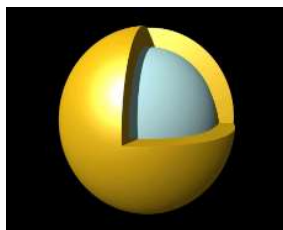
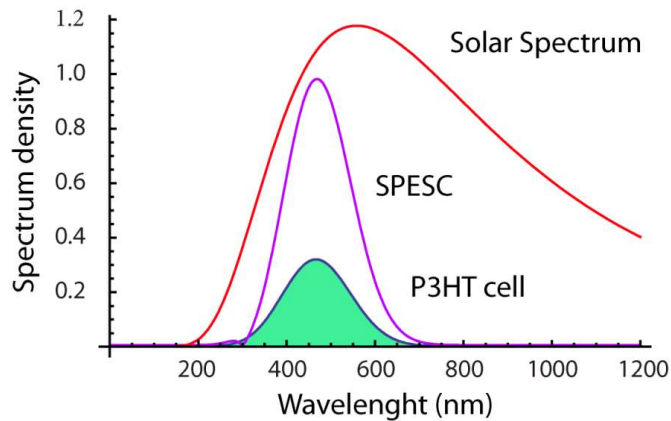
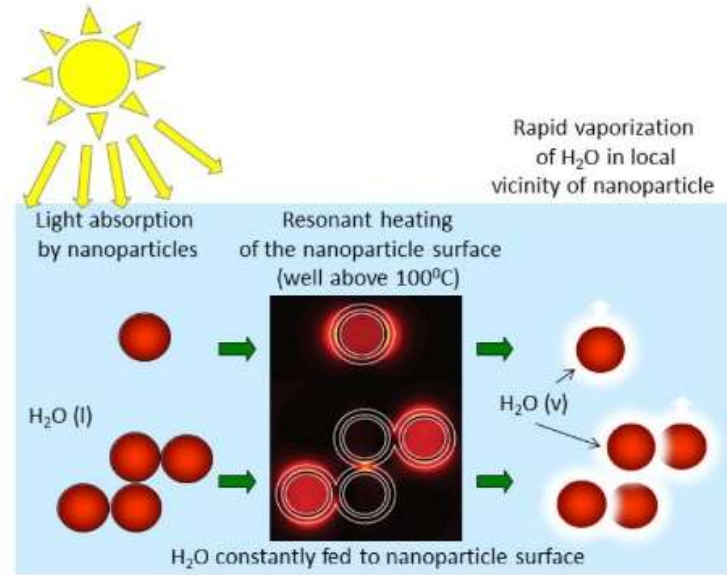


# Some potential new energy technologies

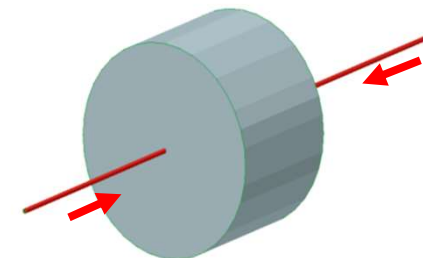
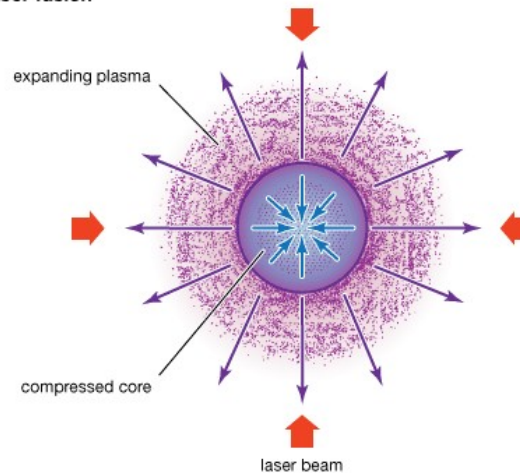
(involving nanotechnologies)



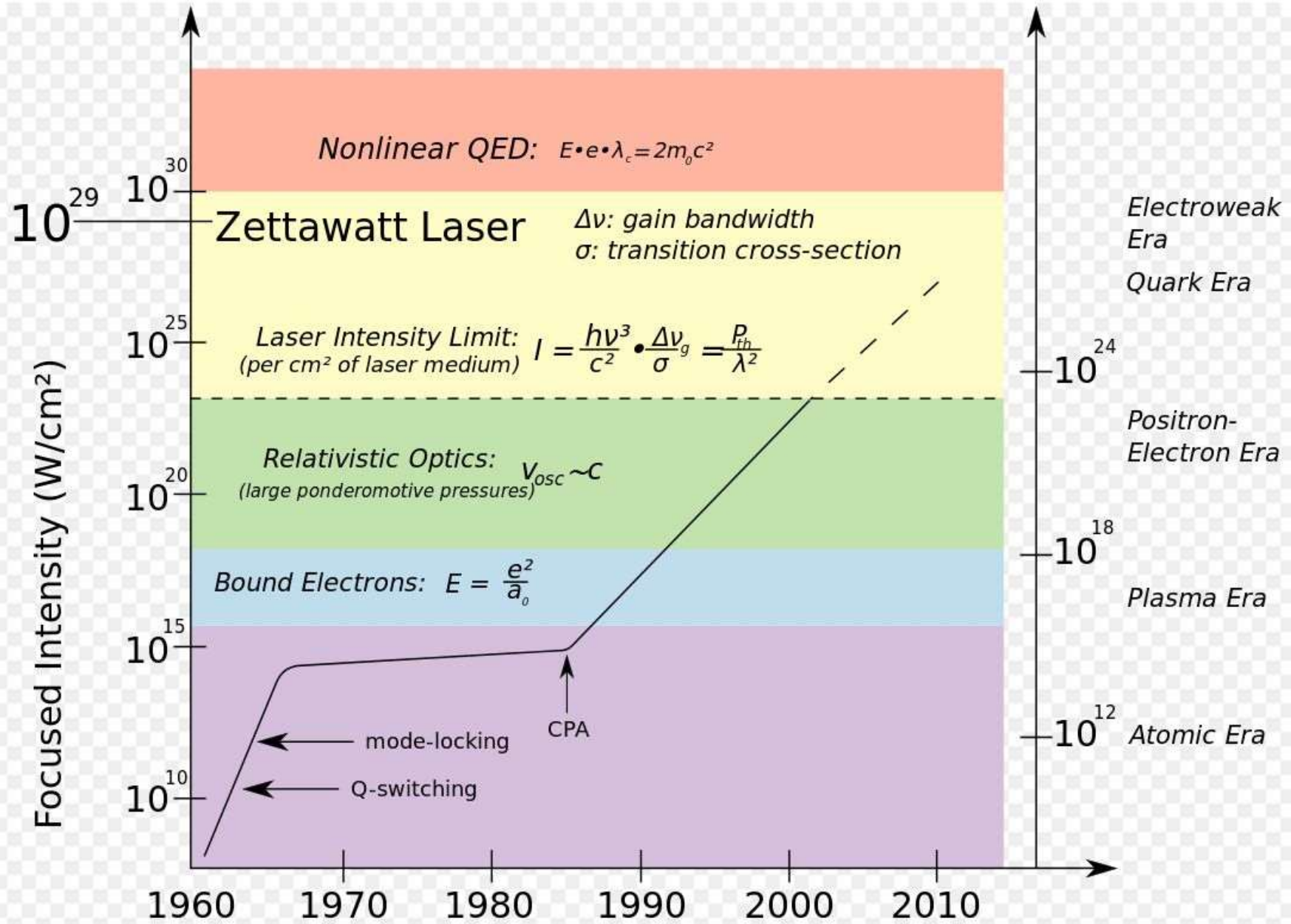
**P3HT Cell**  
 efficiency = 6%  
**SPESC (P3HT)**  
 efficiency = 17.5%



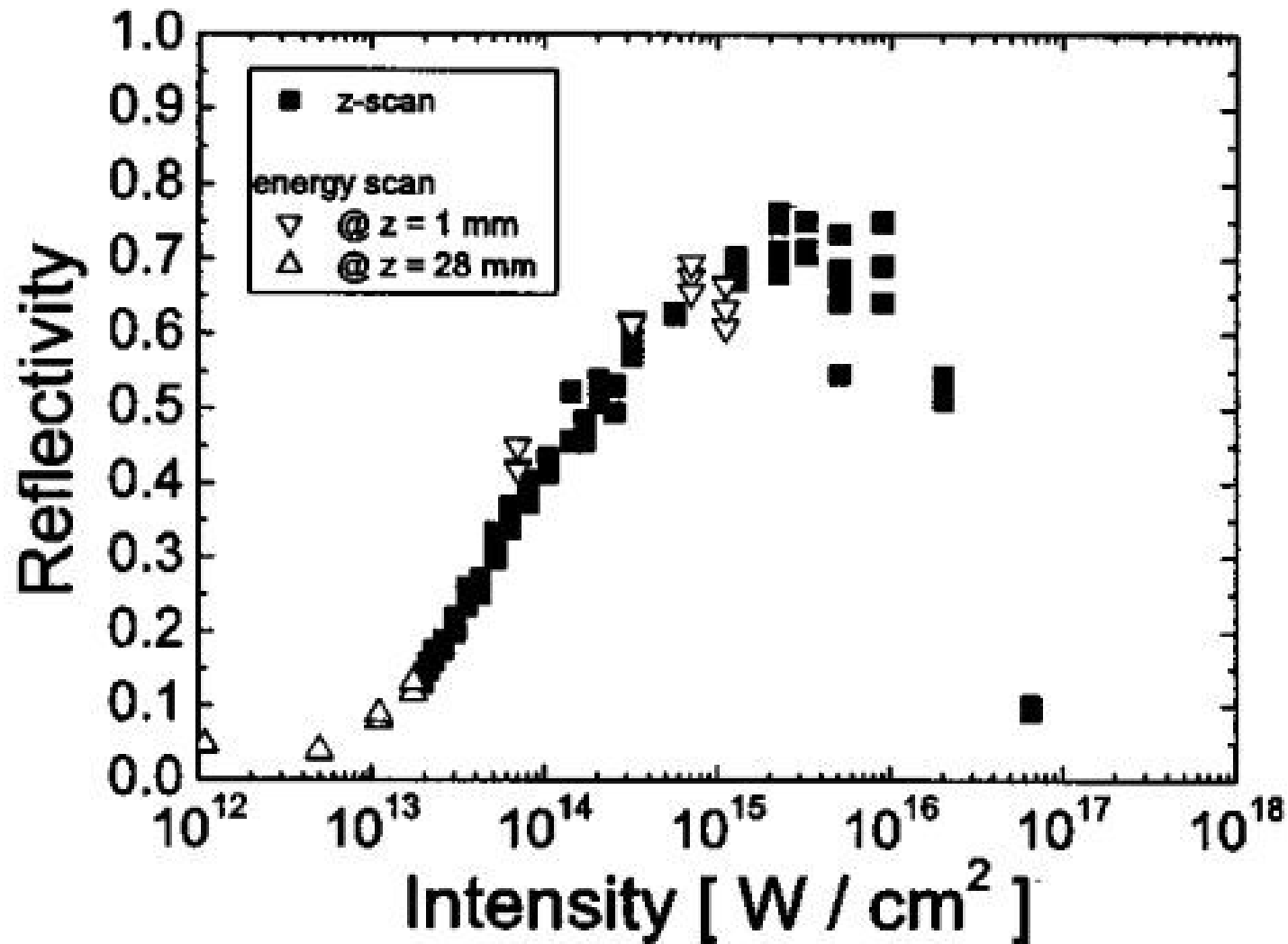
**Laser fusion**



Toward our proposal

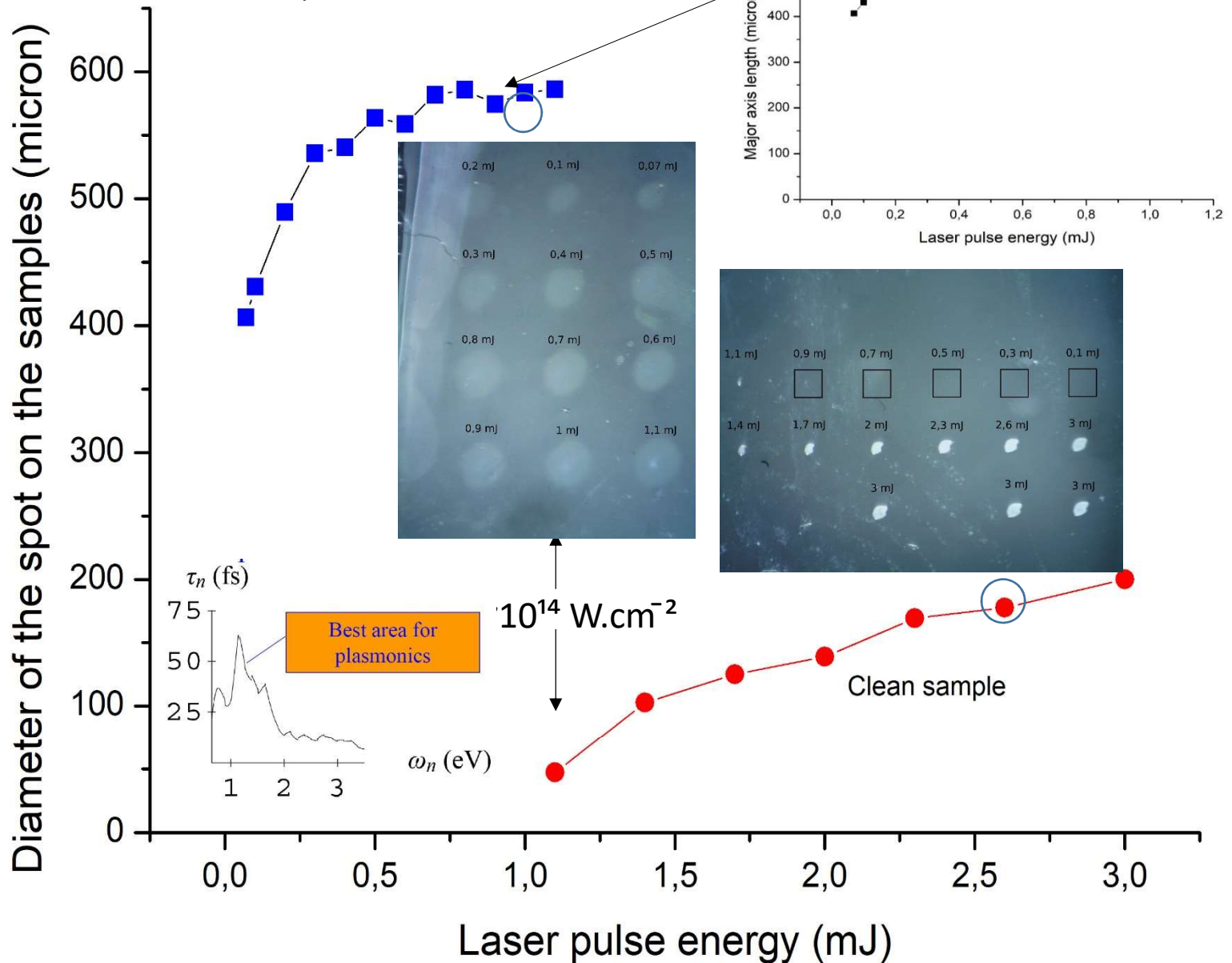


## PLASMA MIRROR REFLECTIVITY



Ch.Ziener et al : J.Appl.Phys. 93,768 (2013)

Laser pulse length: 300 fs  
 Ti:Sa laser:  $\lambda=800\text{nm}$ ,  $\sim 1.55\text{eV}$



Giant plasmonic amplification; the laserlight reaches the nanoantennas;

# LIGHT PENETRATION INTO THE TARGET

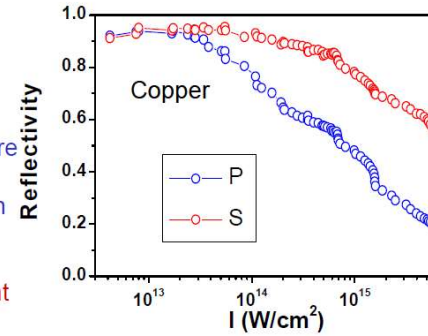
## Plasma absorption

$A = 1 - R$

$I < 3 \times 10^{13} \text{ W cm}^{-2}$ , A is almost polarization independent & obeys Fresnel laws, as IB is dominant

• at higher intensities, there is a clear polarization dependence of absorption

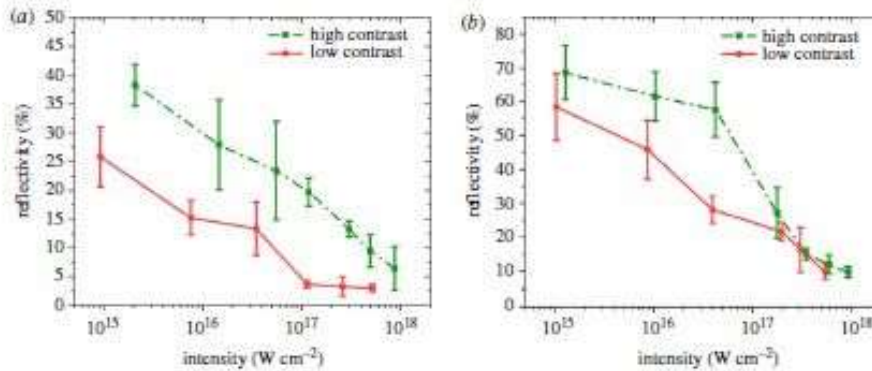
• the difference in absorption should account for extra absorption mechanisms, which are polarization dependent



R vs I at 45°

TIFR data

G.R.Kumar: [www.tifr.res.in/~uphill](http://www.tifr.res.in/~uphill)



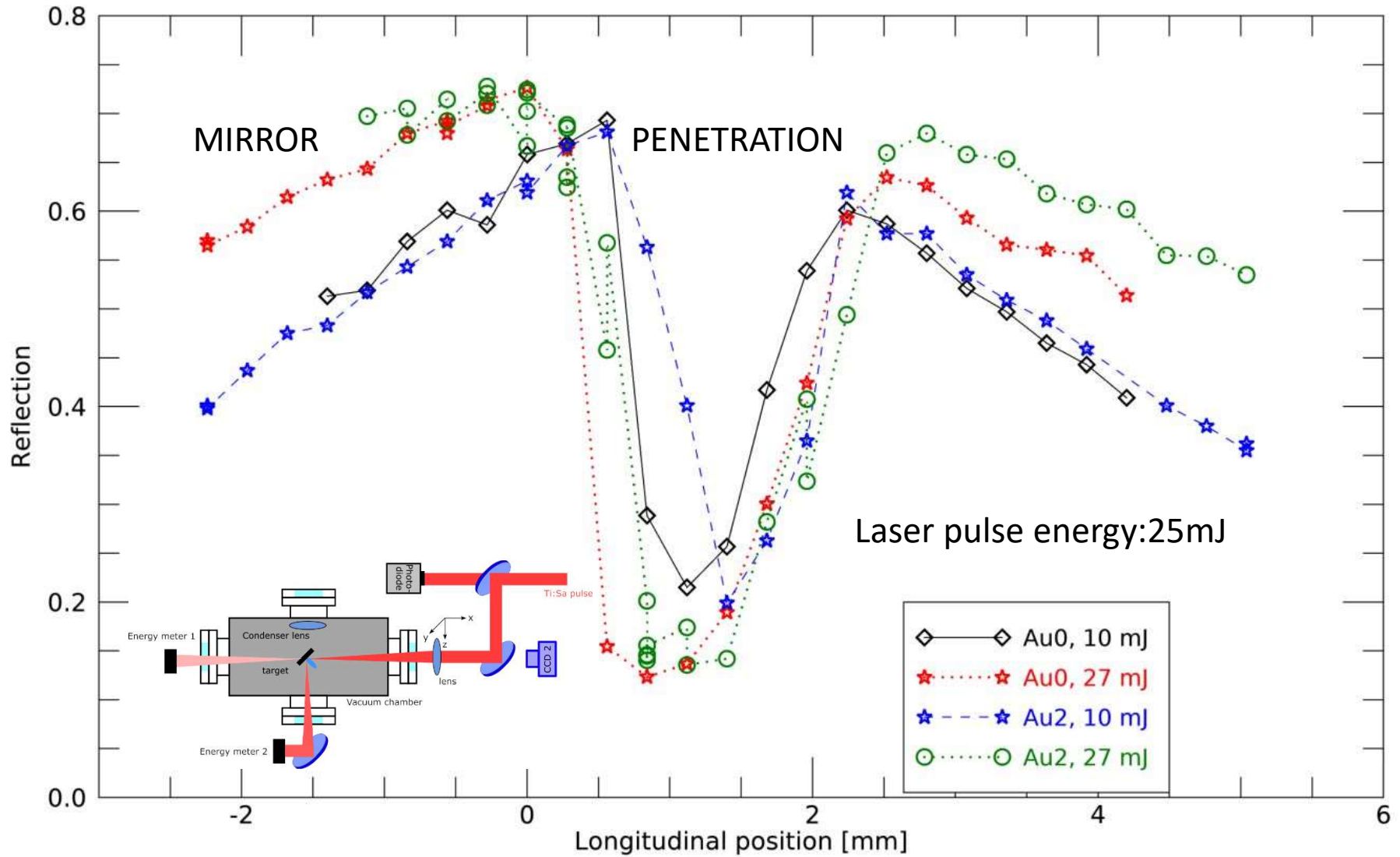
**Figure 2.** Measured reflectivity from (a) gold and (b) boron targets. The red circles (solid line) correspond to the case of low and the green squares (dotted line) the high-contrast case.

Zs.Kovács, I.B.Földes:  
Phyl.Trans.R.Soc A378  
20200043 (2020)



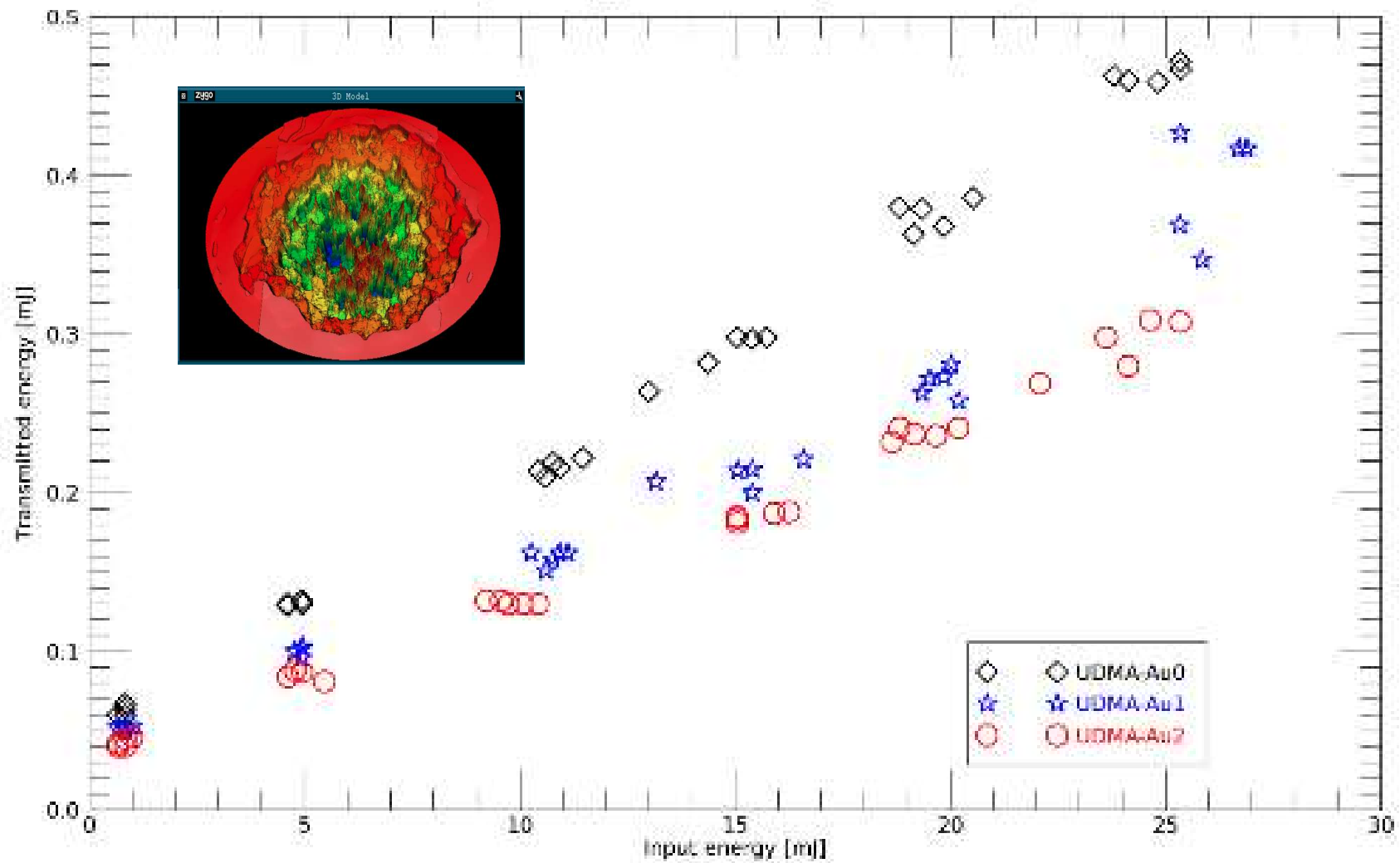


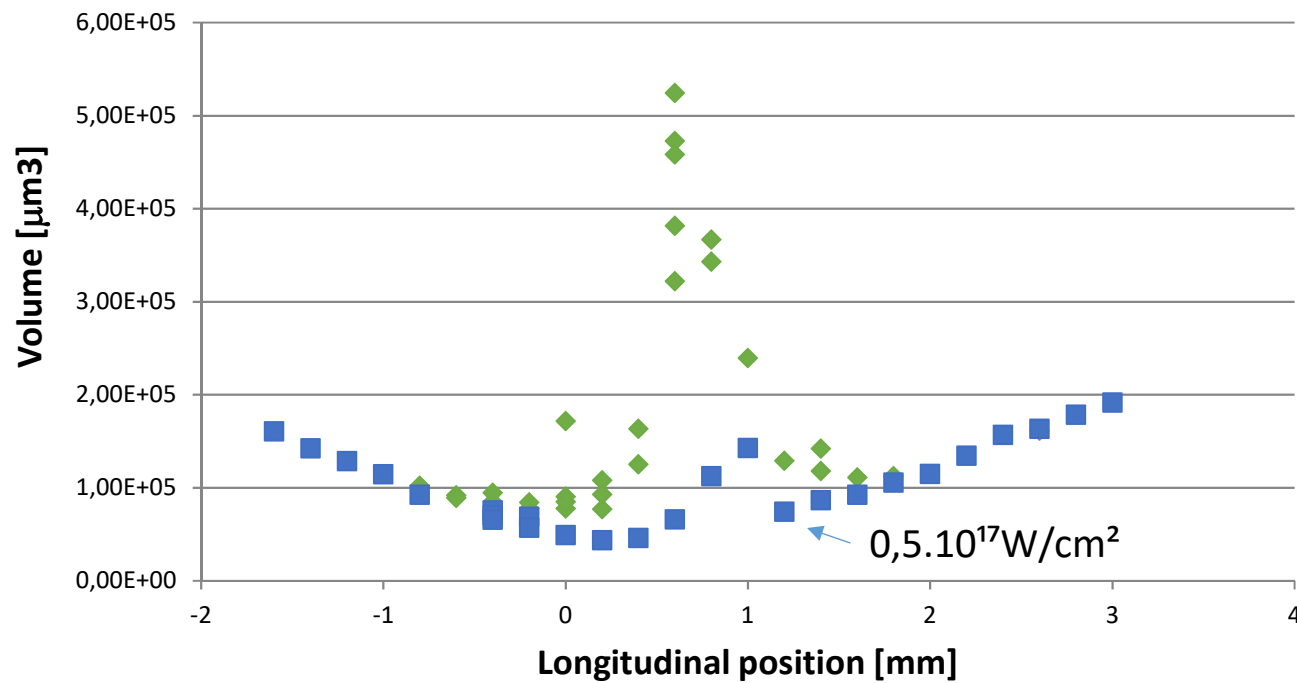
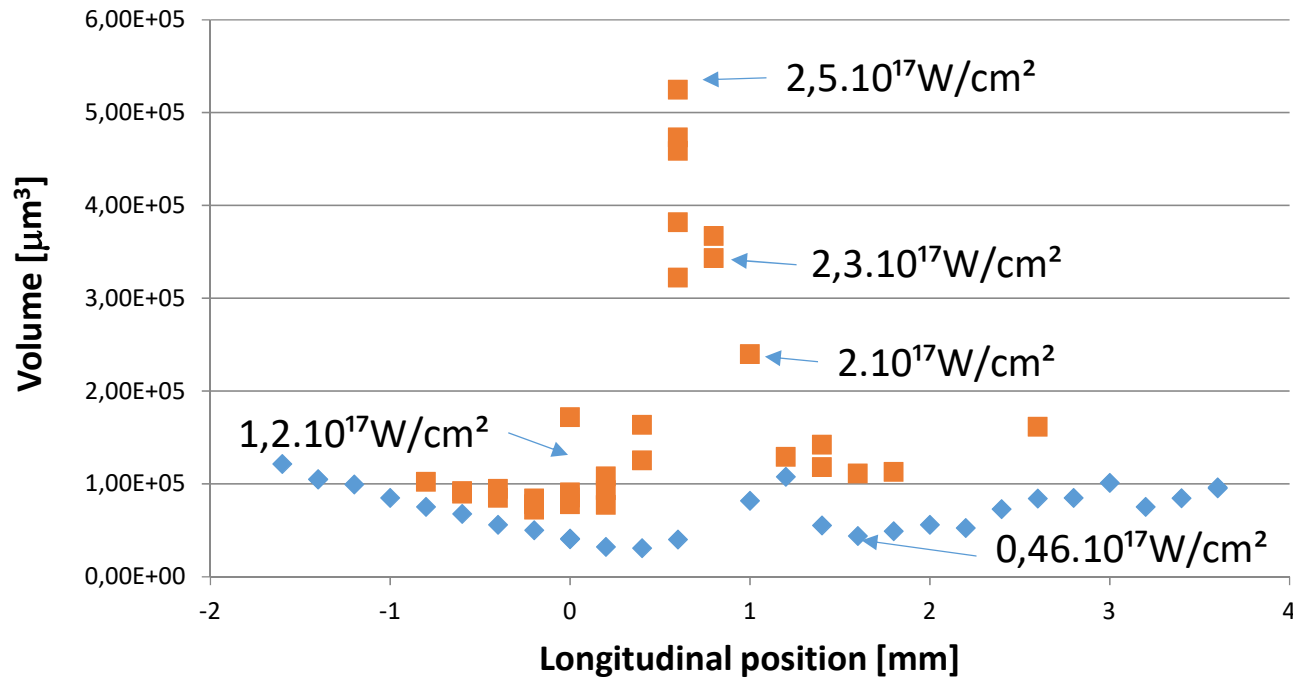
Reflection vs. focusing on Au0 and Au2 samples (2022.02.20.)



Sample: 160µm thick transparent polymer (UDMA).

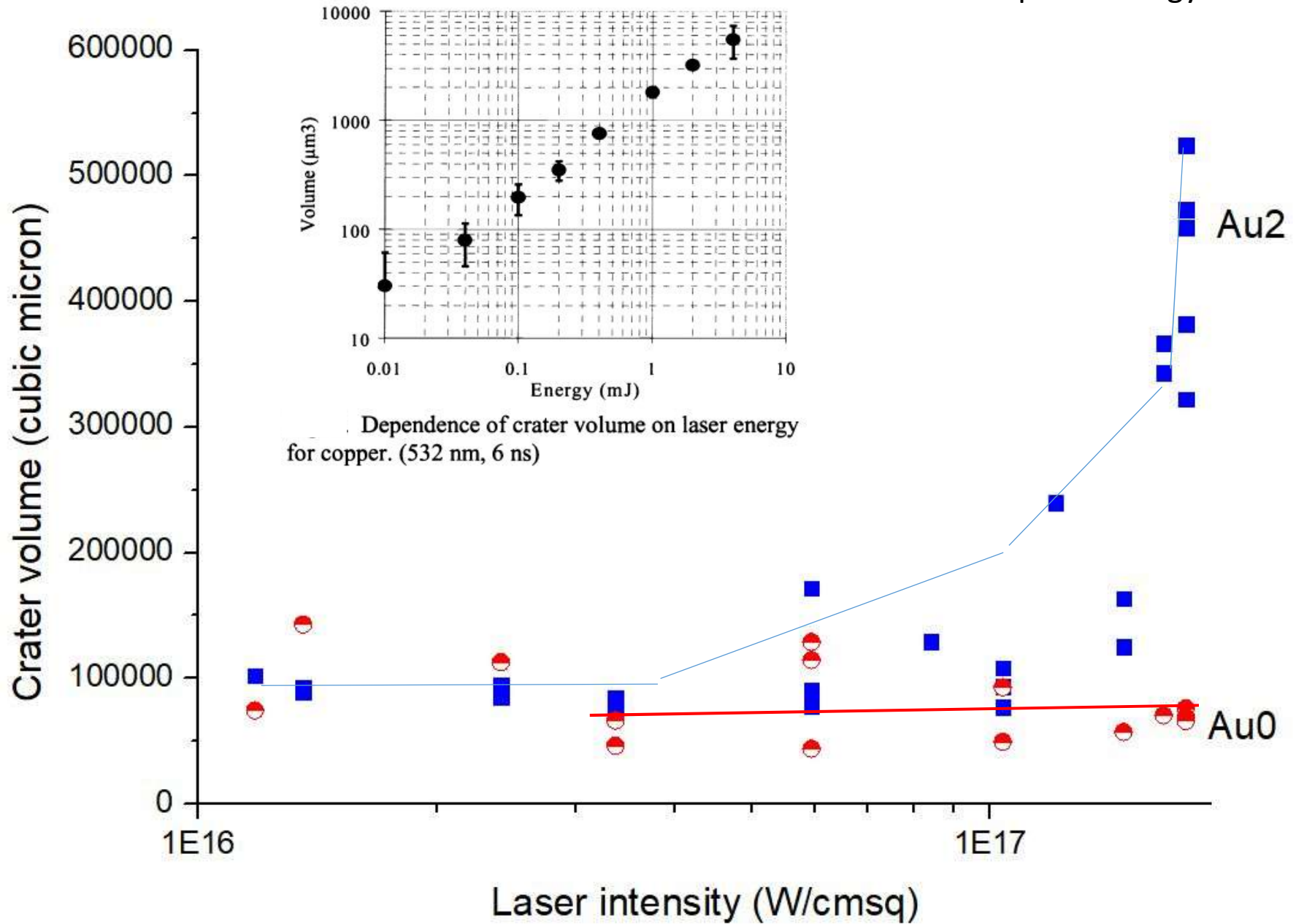
Laser pulse energy transmitted through the UDMA samples





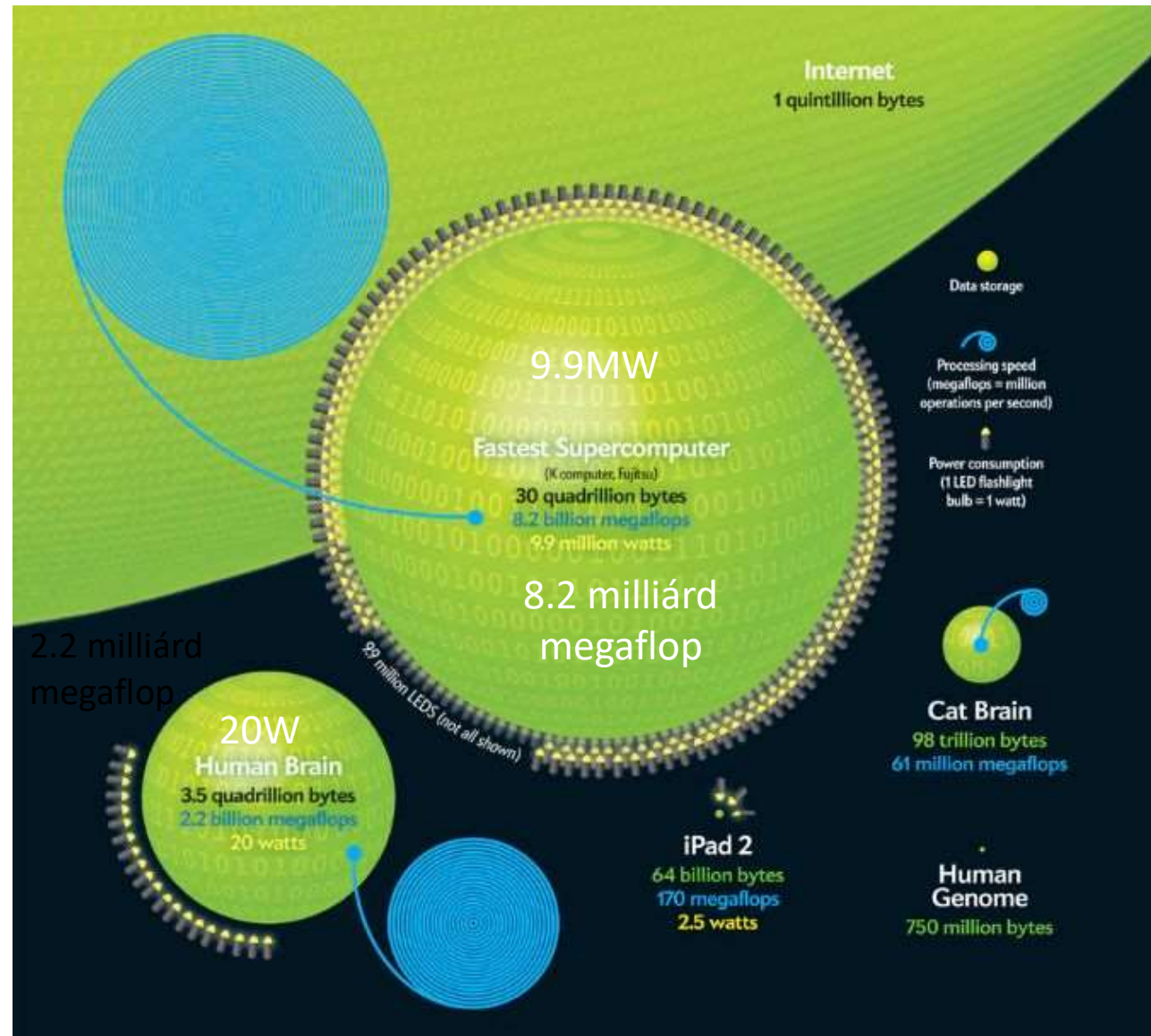
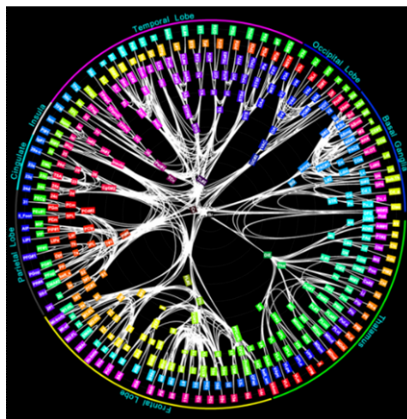
**HIGH FIELD  
PLASMONICS  
WORKS!**

Laser pulse energy: 25mJ



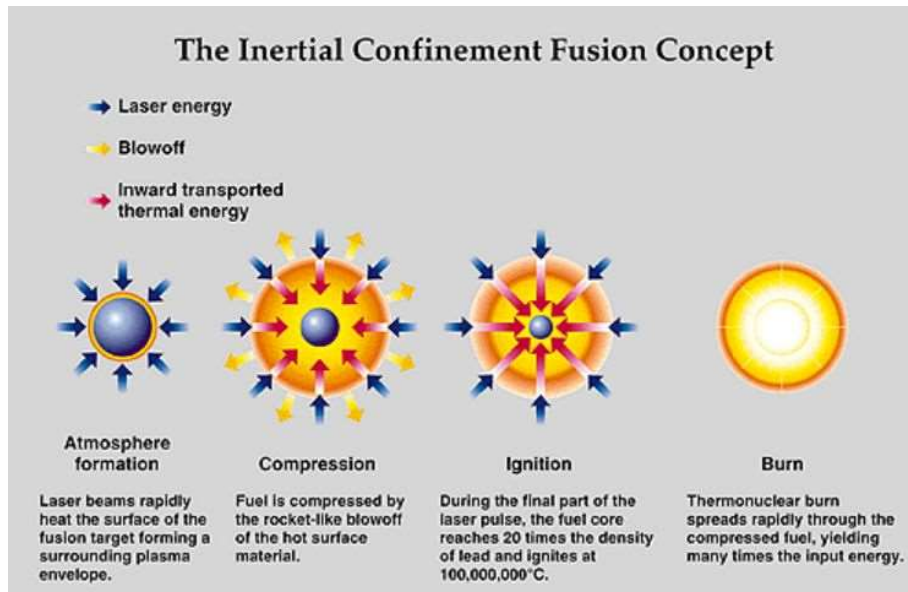
# COPYING NATURE!

## HUMAN BRAIN VS SUPER COMPUTER



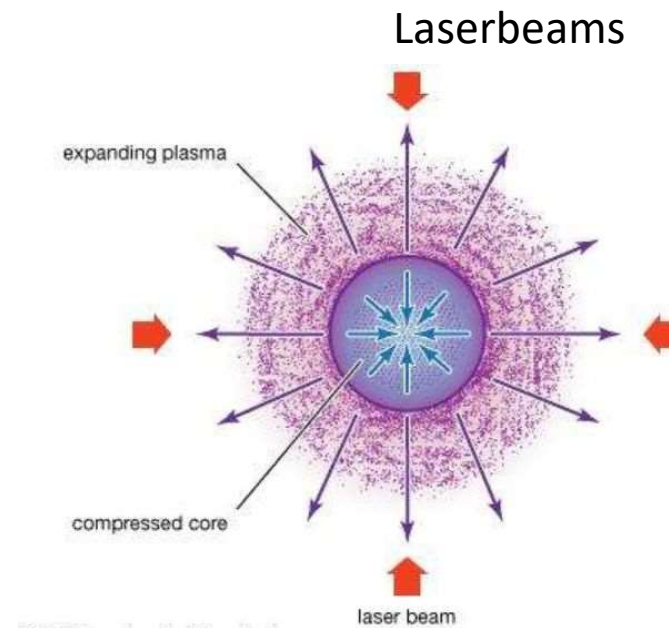
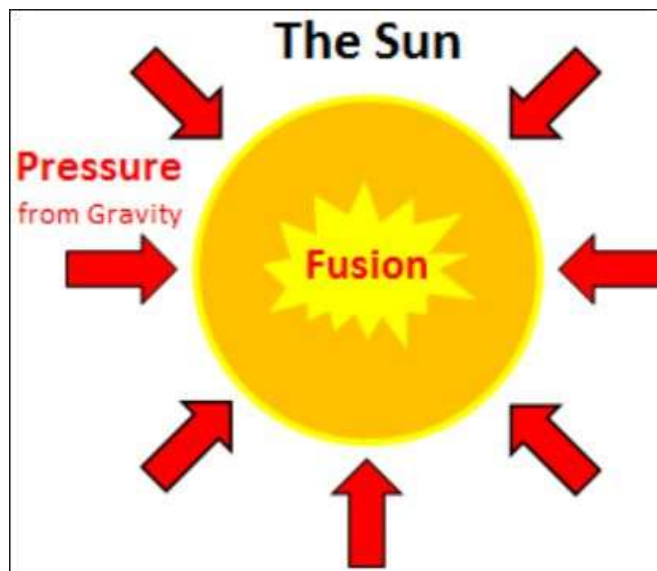
(IBM TrueNorth chip simulation of 1million neurons and 256 million synapses , 10 thousand times less energy) [Memristors\(nanorods\)](#)

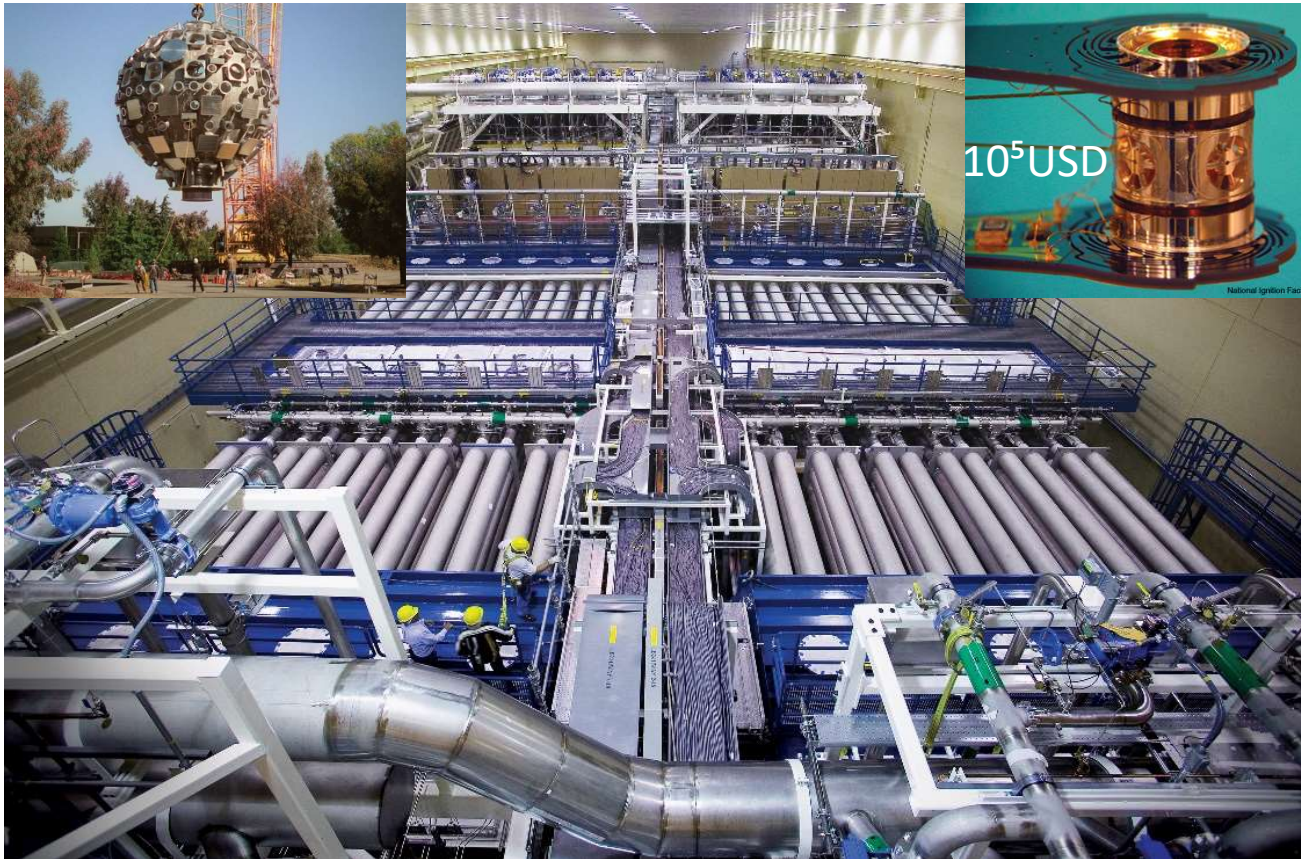
# No2 application (LSPP)



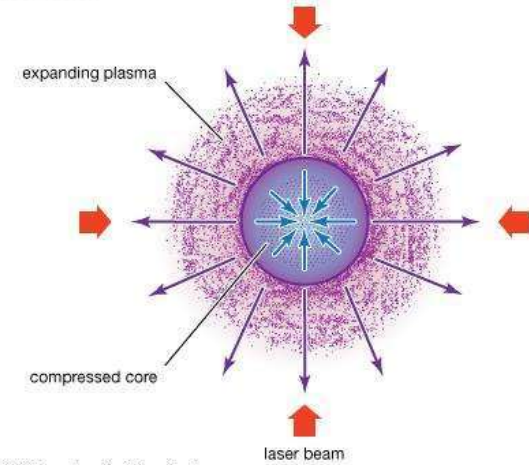
The most successful technologies imitate nature

$$E = mc^2$$





Laser fusion



© 2009 Encyclopædia Britannica, Inc.

## Problems of inertial fusion

Long laser pulses ( $\sim 50\text{ns}$ )  
 Raileigh-Taylor instability  
 Complicated target construction  
 Enormous laser energy (400/1.8MJ)

- High requirements on irradiation symmetry
- Insufficient laser repetition rate
- Very precise injection system is needed
- The target position has to be tracked in order to ensure required irradiation precision

**TO COMBINE 2 DIFFERENT (e.g. fusion and nano-) TECHNOLOGIES TO REACH FUSION AT THESE ULTRAHIGH EM FIELDS?**

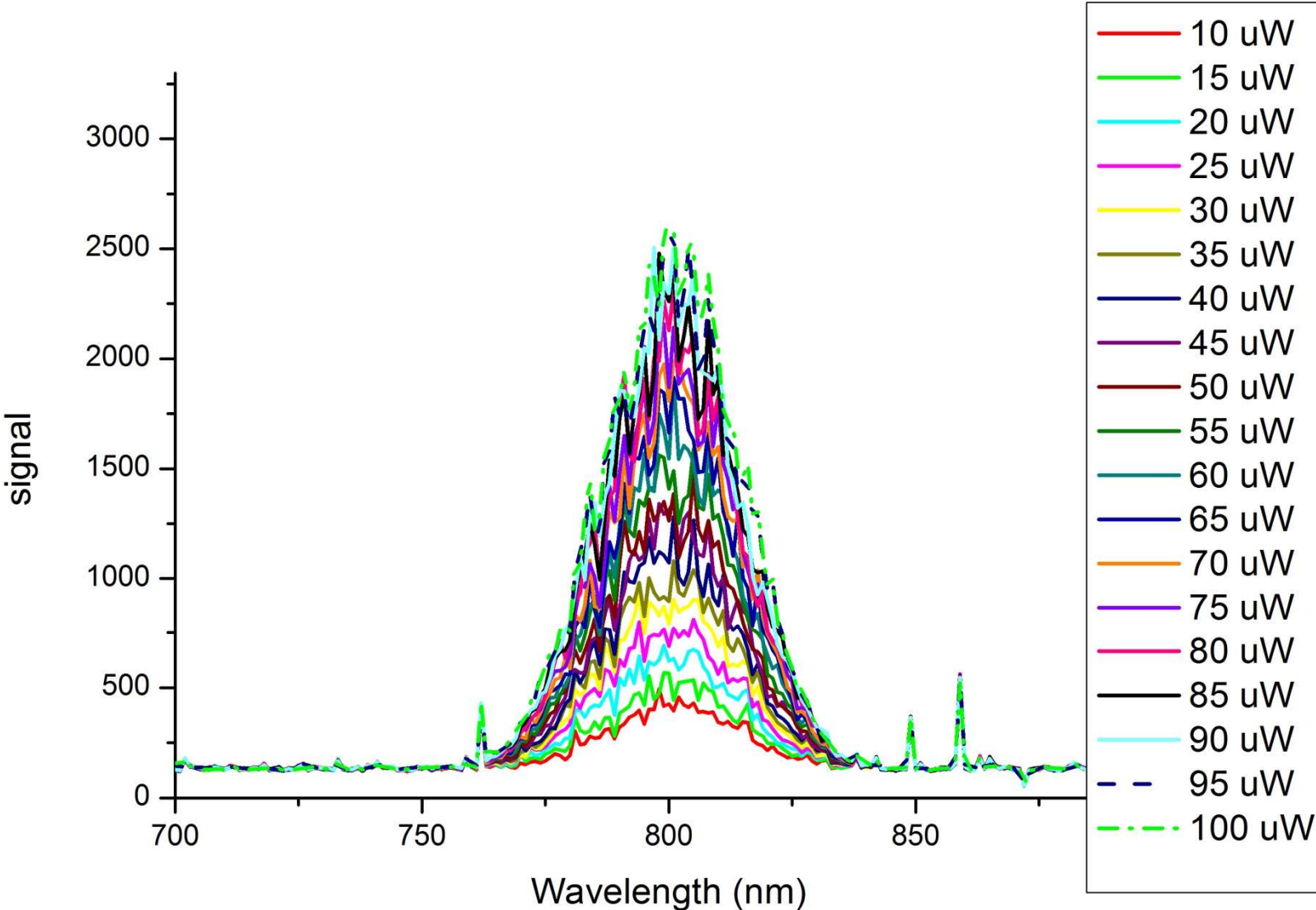
# OUR PROPOSAL: COMBINE PLASMONICS WITH NUCLEAR FUSION TECHNOLOGY

SOME POTENTIALLY EXPLOITABLE HIGH FIELD PLASMONIC EFFECTS:

1. Go for localized surface plasmon polaritons (LSPP)
2. Lifetime of LSPP-s is in the few ten femtosecond range. We may get high intensity laser pulses in this time-domain and the plasma instabilities disappear.
3. High electron densities and EM fields can be obtained in small (nanosized) volumes on resonant plasmonic nanoparticles (hot spots).
4. The near field of plasmons screen the repulsive field of positively charged (e.g. protons) particles and so they may fuse more easily. So do the ponderomotive forces.
5. The large number of conduction electrons move in the plasmonic excitations in correlated way and their momentum may be in sum transferred in high exciting fields to positive particles, moving together with them, further increasing the probability of nuclear fusion.
6. With these short pulses we do not need many beams, like in the NIF, the target can be a thin film, illuminated only by 2 beams from opposite directions, and the same energy density may be achieved in the whole thickness of the target sample, and this may



# Ti:Sa LASER PULSE SPECTRA AT DIFFERENT INTENSITIES



# ILLUSTRATION OF THE RESULT OF THE SCREENING EFFECT



HOT SPOTS

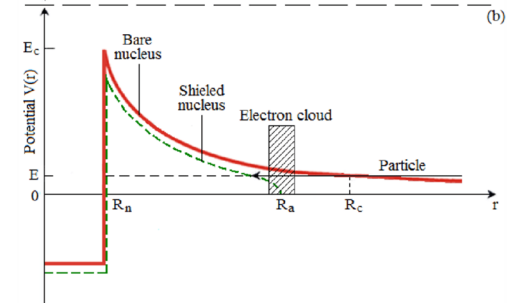
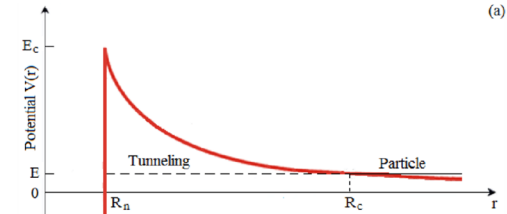
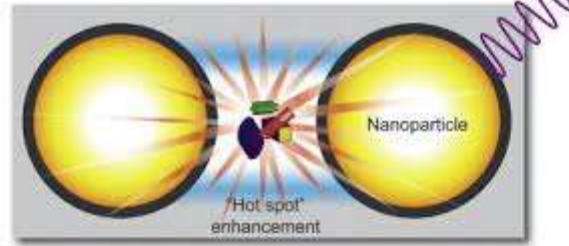
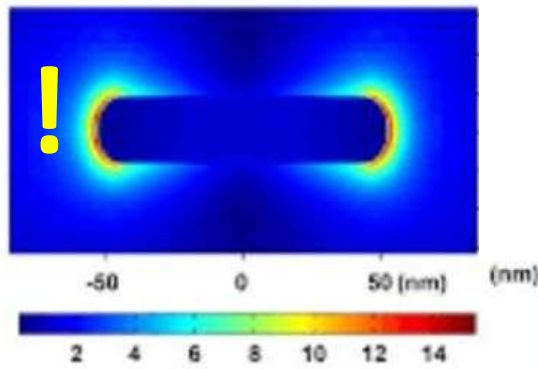
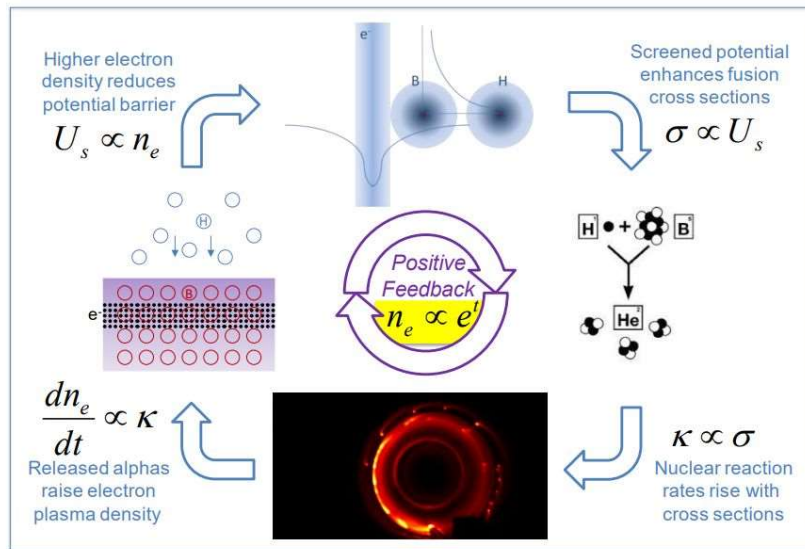


Illustration with the  $H+B \rightarrow 3He$  reaction



A.Y.Wong and C-C.Shinh:UCLA 2019

Ponderomotive screening

$$F_p = (e^2/4m_e\omega^2) \cdot \text{grad}(E^2)$$

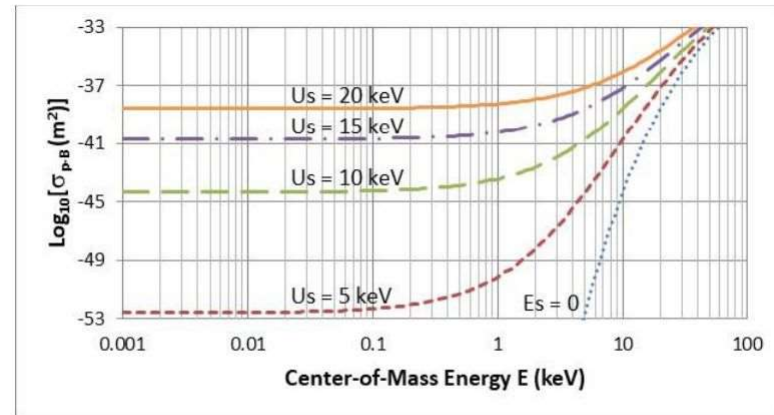
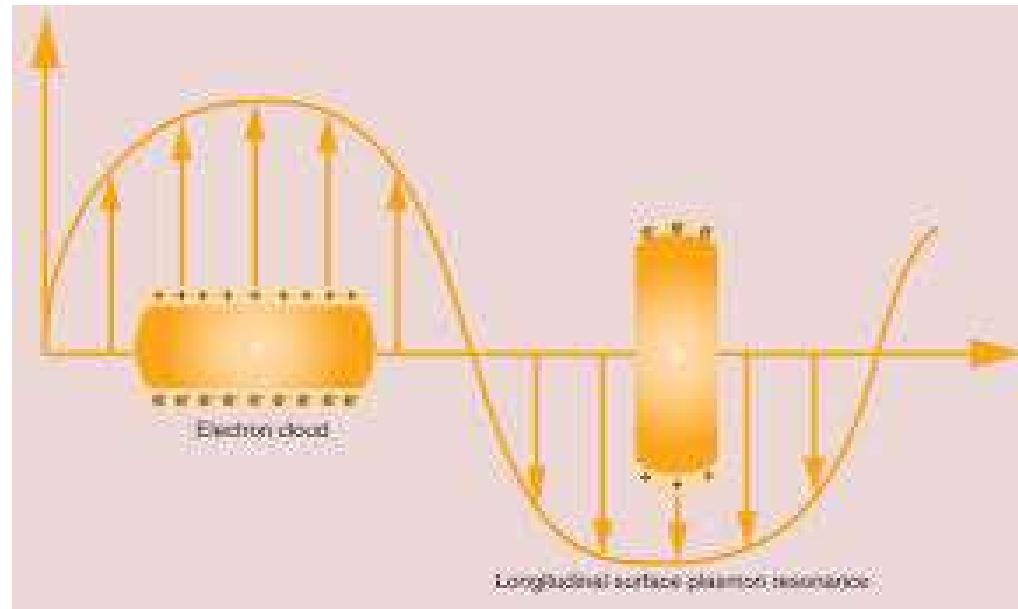
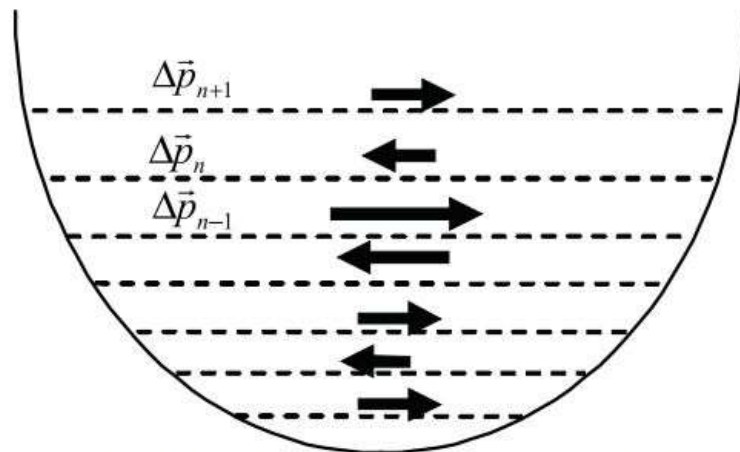


Figure 1:  $p\text{-}^{11}\text{B}$  cross section as function of particle energy for the screening electron densities up to  $E_s = 20\text{keV}$ . The cross section near  $E = 10\text{eV}$  grows over 14 orders of magnitude (from  $10^{-53}$  to  $10^{-39}\text{m}^2$ ) over the range of 5 to 20keV.

# Demonstration of the correlated state



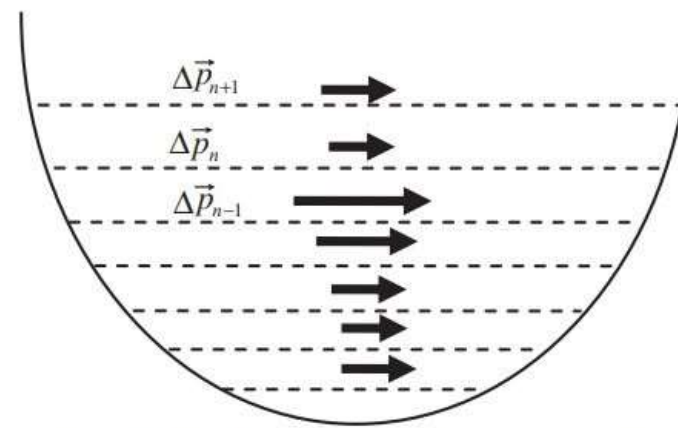
**Uncorrelated state**



Small fluctuation of a total momentum



**Correlated state**

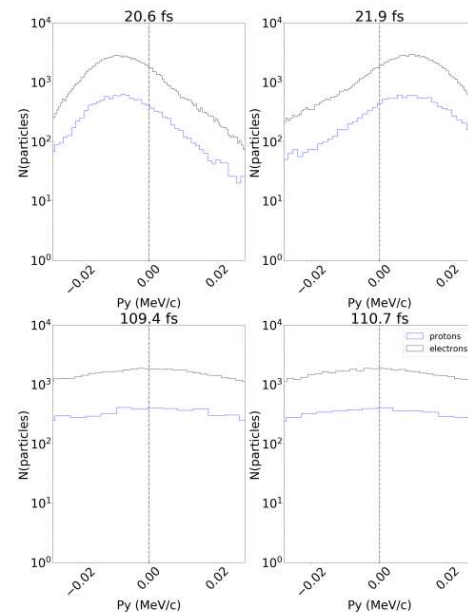
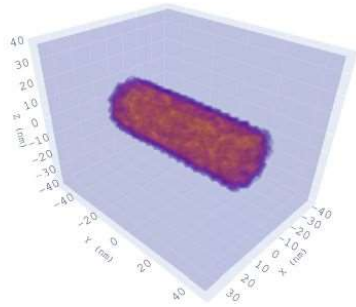


Giant fluctuation of a total momentum

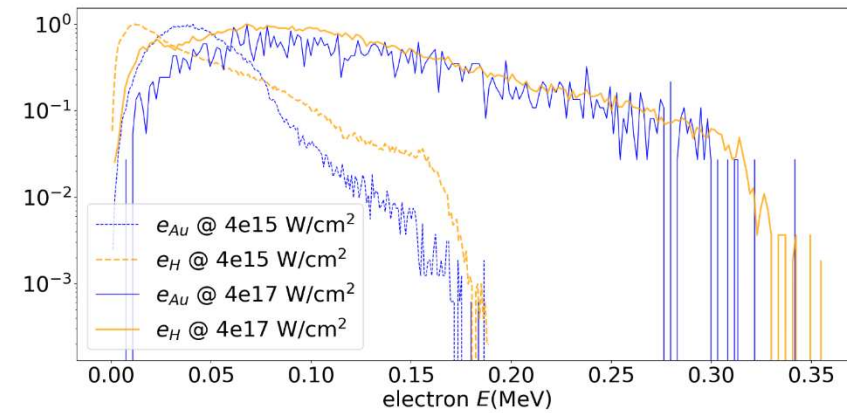
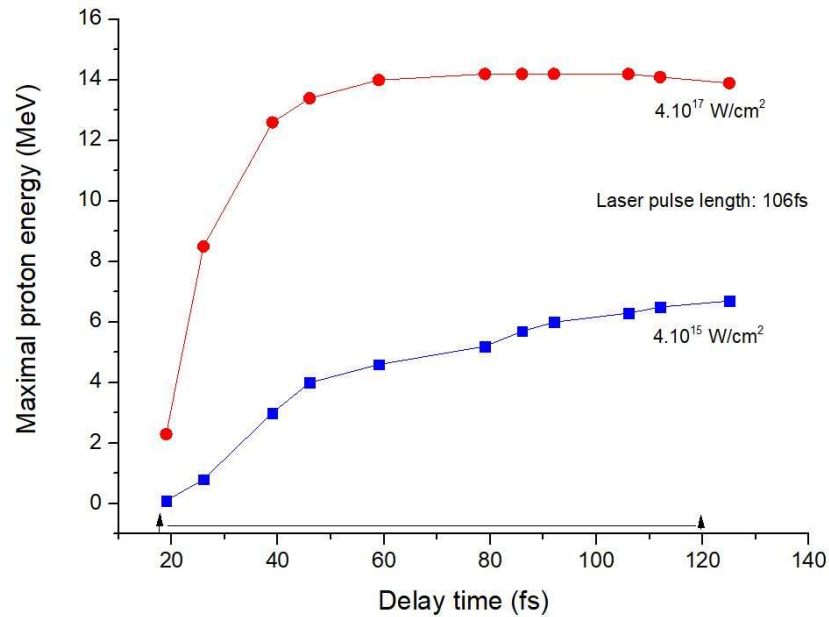
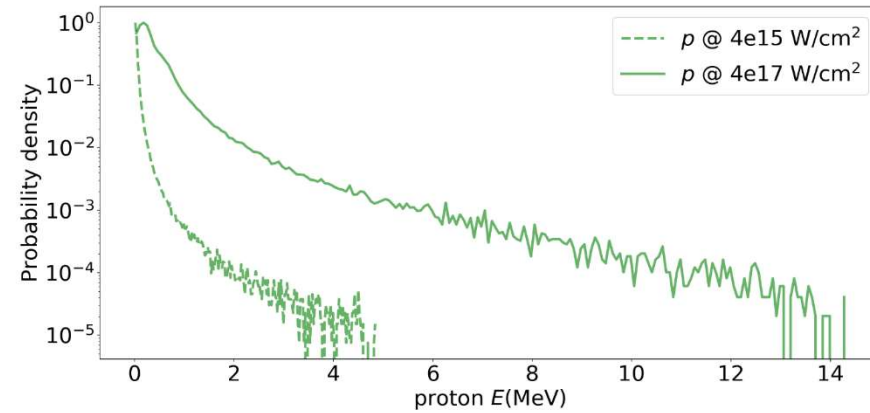


# SIMULATION OF PROTON AND ELECTRON ENERGIES AT A SINGLE NANOROD

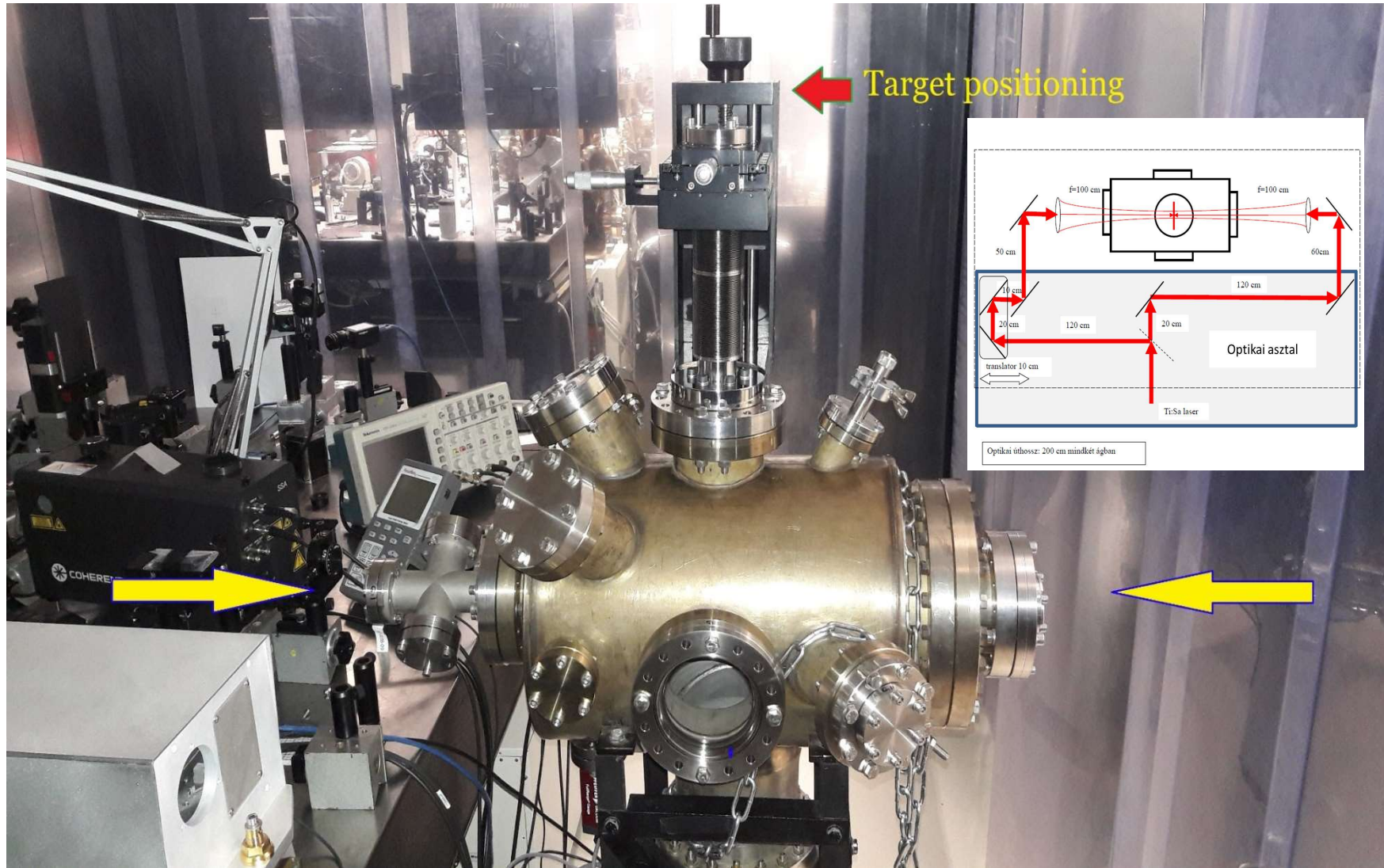
Nanorod inside a PIC simulation box

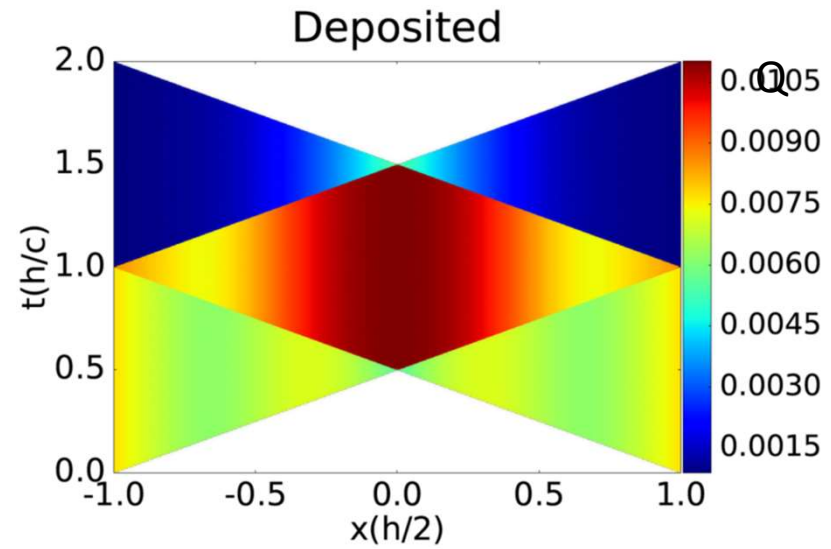
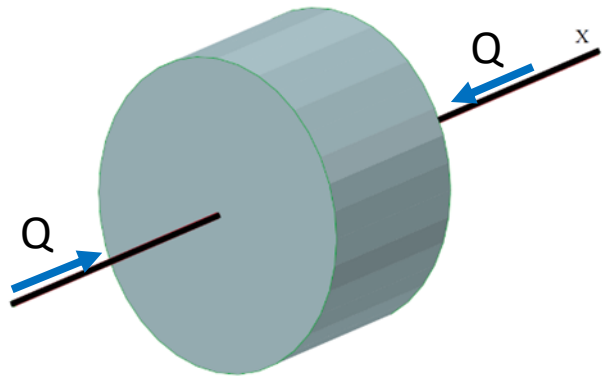


59.67 fs

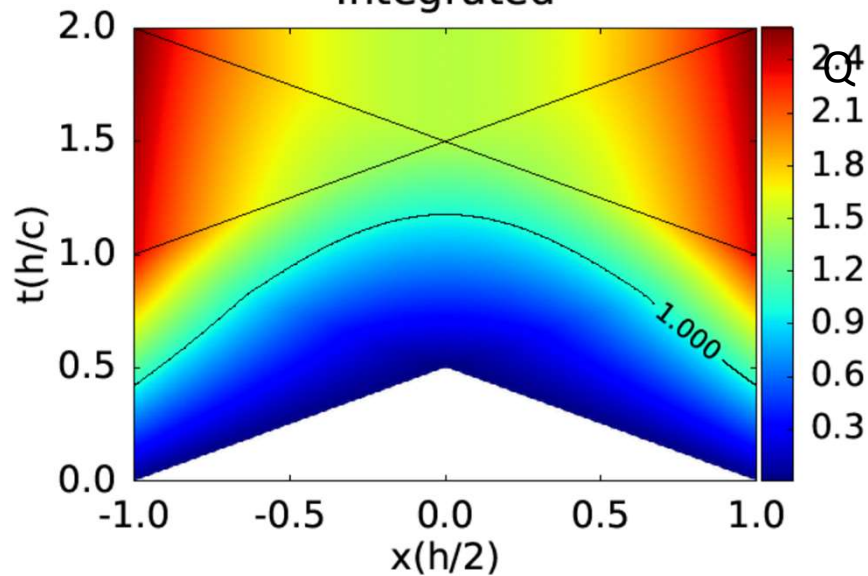


# The future: Two-sided irradiation

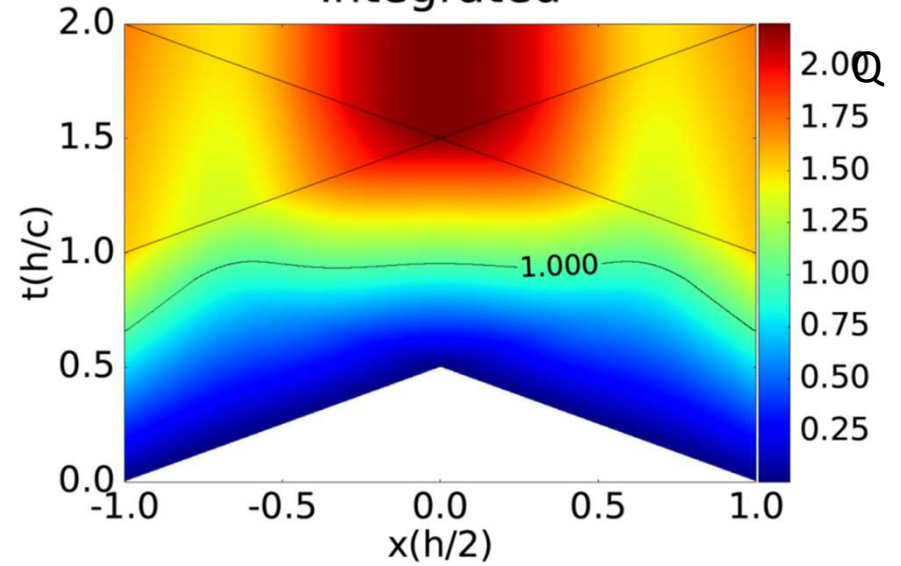




without nanoparticles  
Integrated



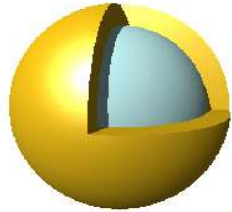
with nanoparticles  
Integrated



Some preliminary results with one side illumination

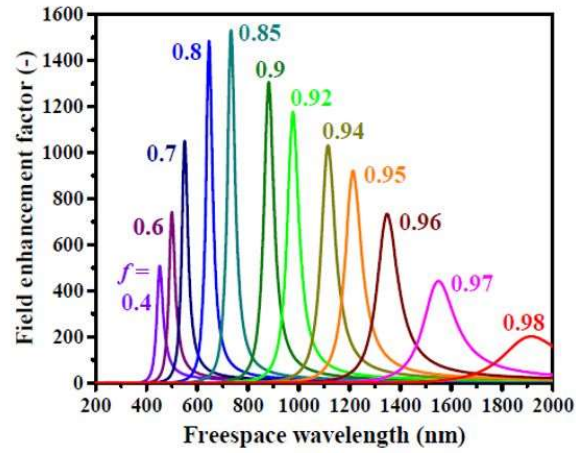


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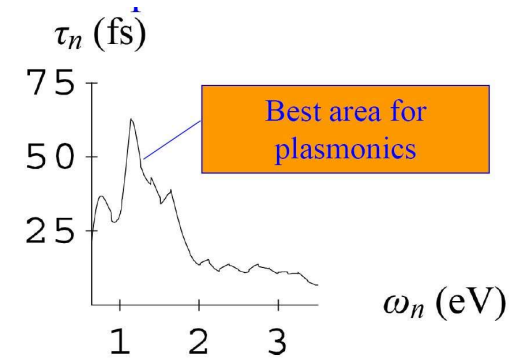


NANOSHELL  
( $n \times 10 \text{ nm}$ )

NANOROD ( $\sim 85 \times 25 \text{ nm}$ )



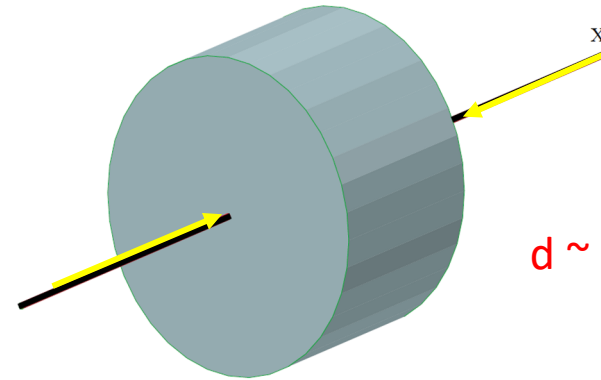
$\lambda = 800 \text{ nm}$



←

$n \cdot 10 \mu$

NANOPARTICLES IN  
THE FUSION MATERIAL

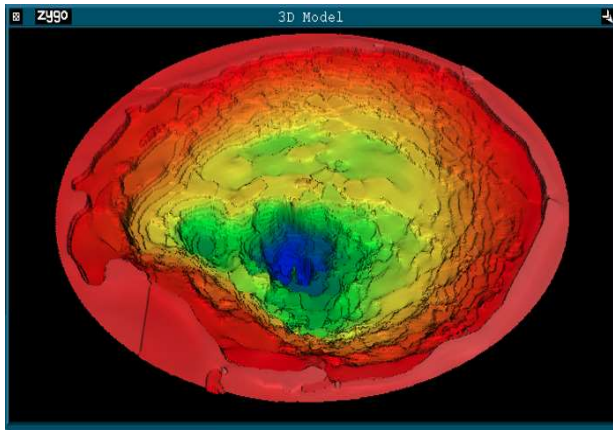


$d \sim n \cdot 10 \mu$

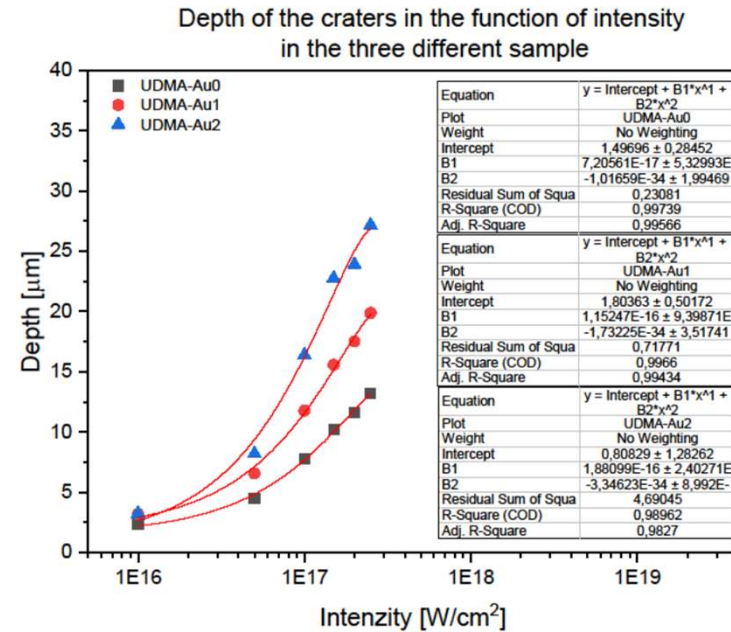
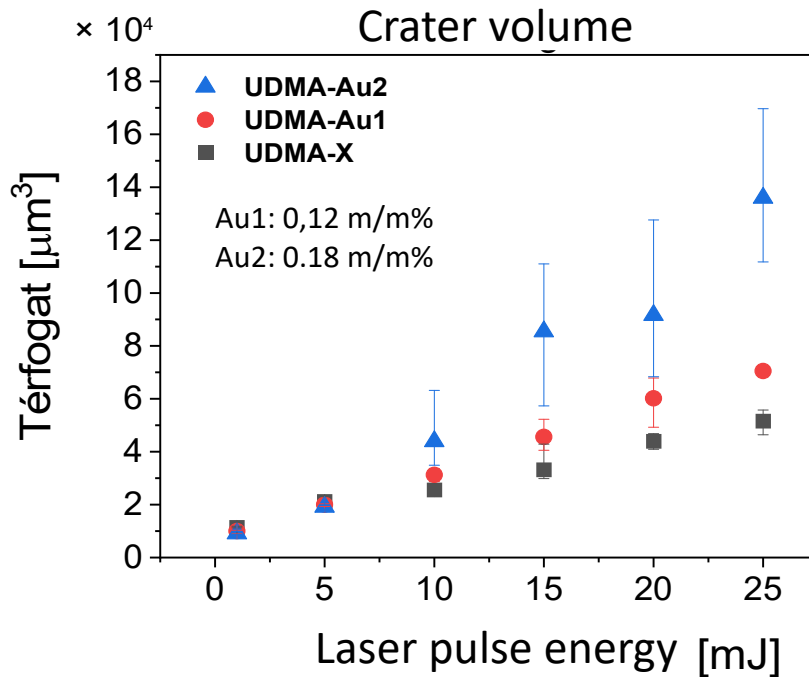
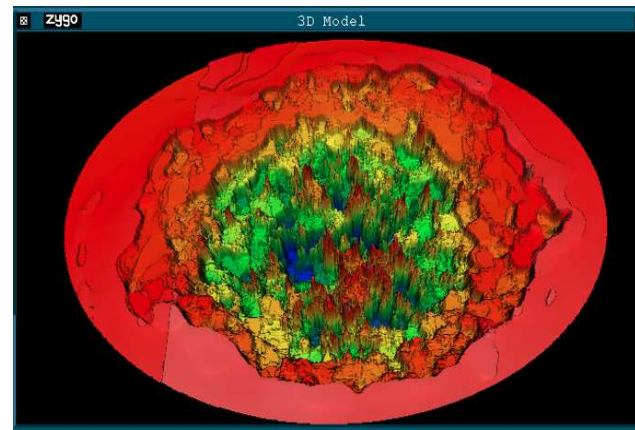
FEMTOSECOND LASER PULSES;  
 HIGH REPETITION FREQUENCY;  
 LIGHT SPEED: NO TIME FOR INSTABILITIES;  
 ONLY TWO BEAMS;  
 TIMELIKE VOLUME IGNITION

# 1. DIAGNOZIS (crater volume)

Volume:  $V_0$



Volume  
max.  $3.5V_0$



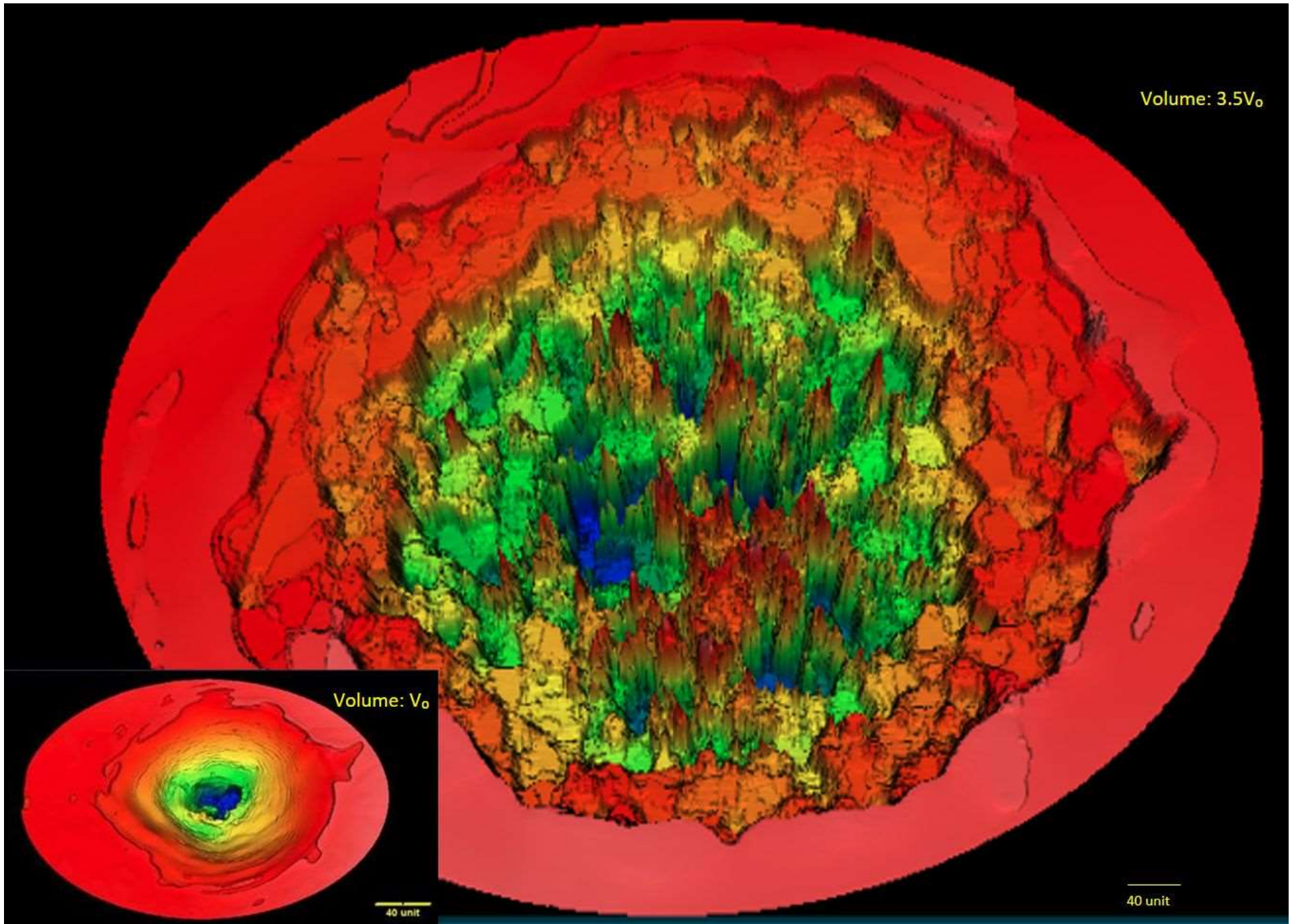


Volume:  $3.5V_0$

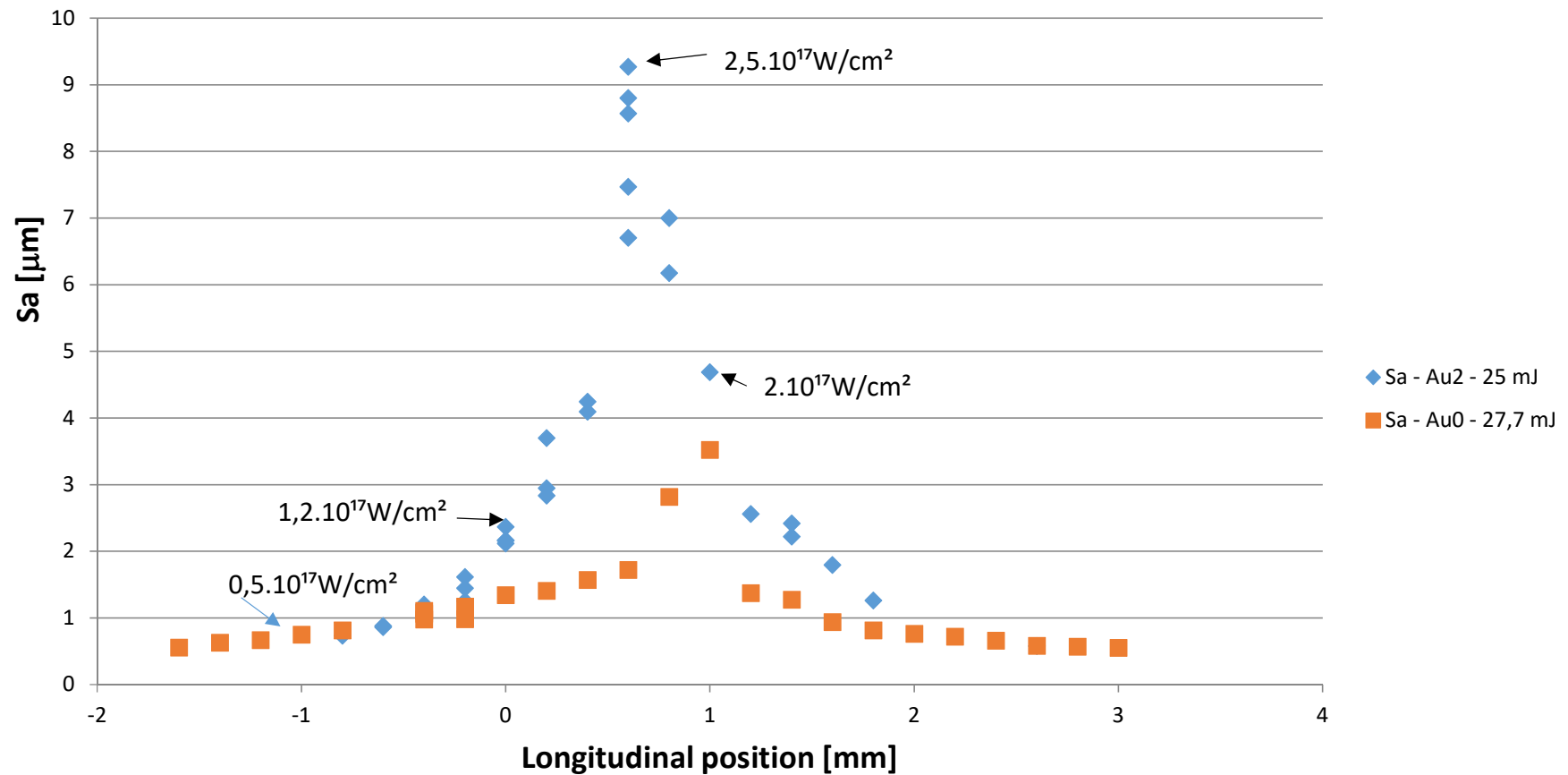
Volume:  $V_0$

40 unit

40 unit

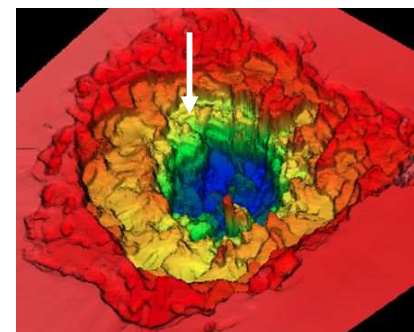
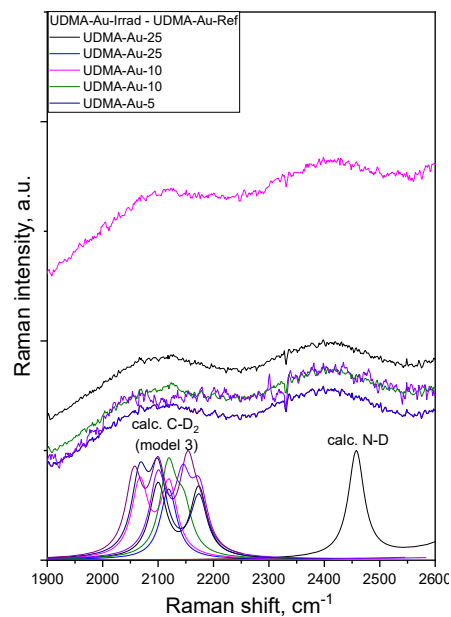
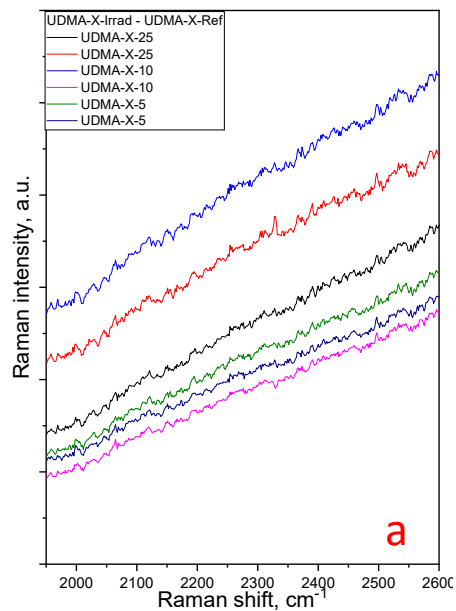


Surface roughness as function of the longitudinal position  
of the Au2 vs. Au0 samples  
Energy of the impulse: 27,7 mJ (Au0) and 25 mJ (Au2)

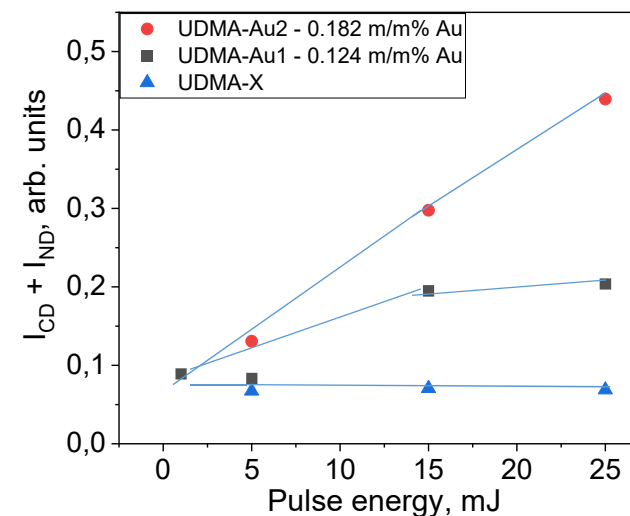
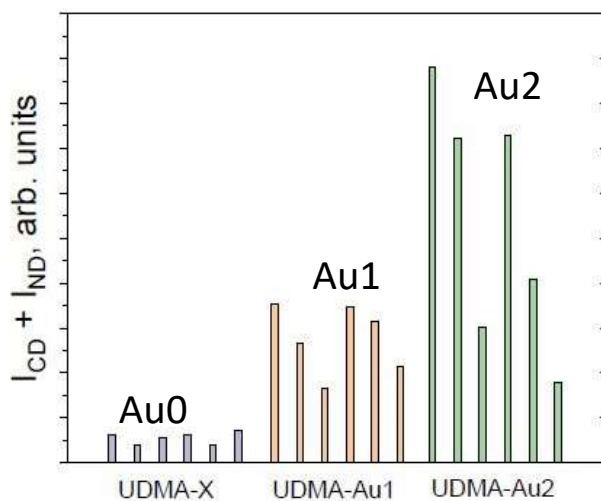
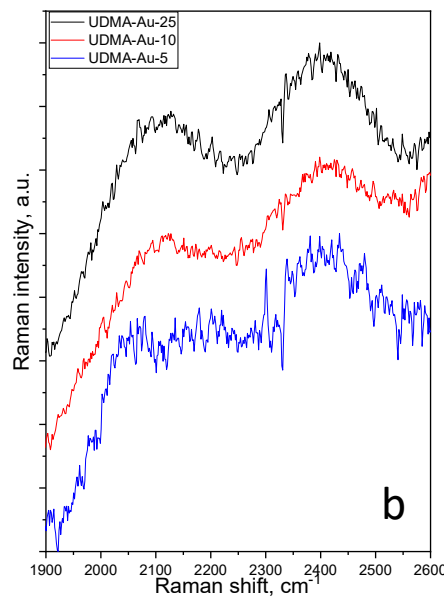
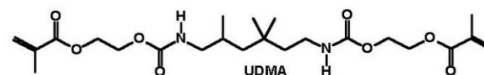


## 2. Diagnosis : Raman scattering from the crater surface

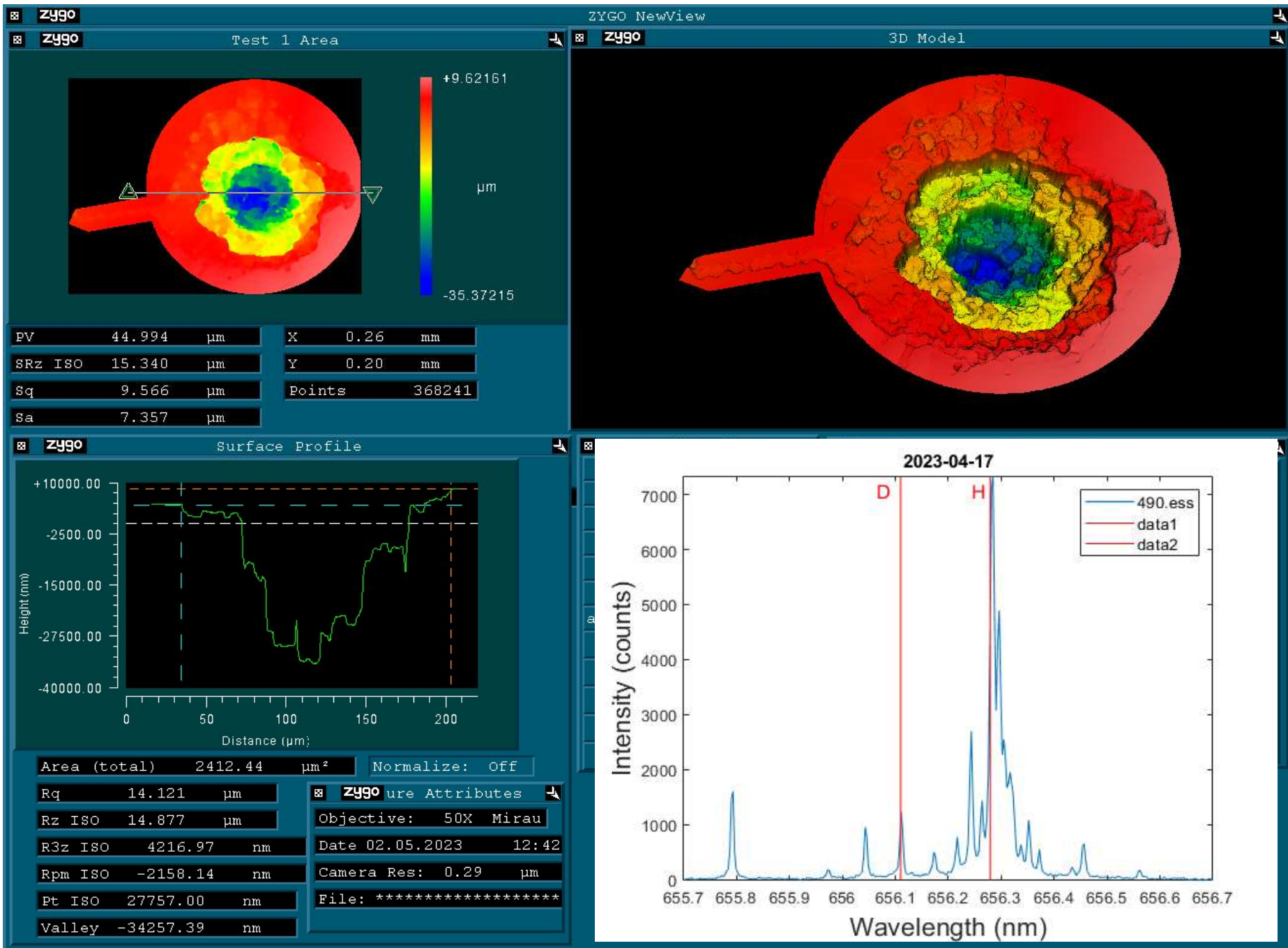
arXiv2210.00619(2022), submitted to  
Advanced Optical Materials



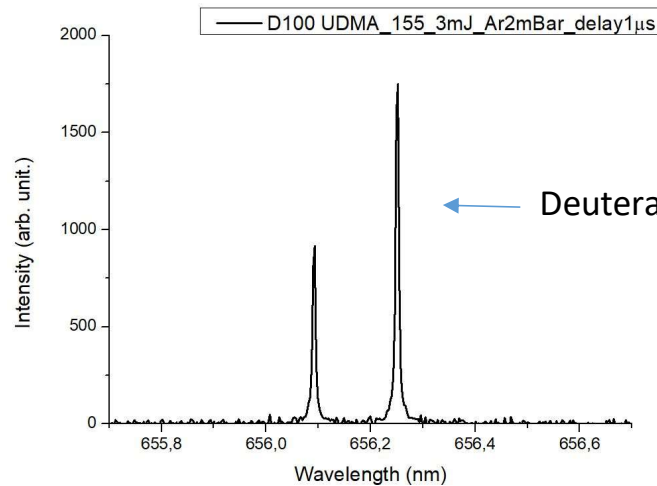
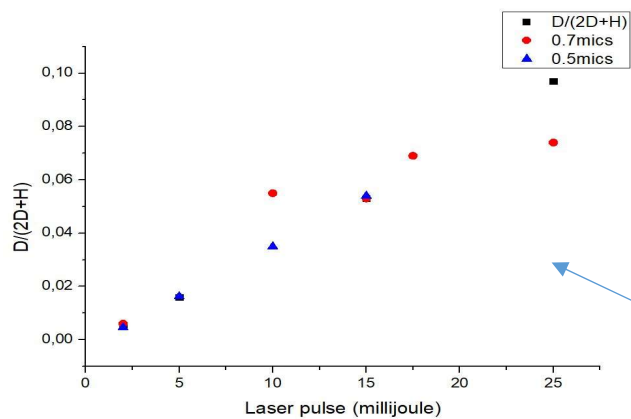
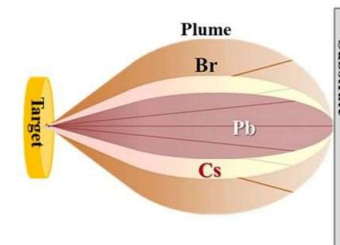
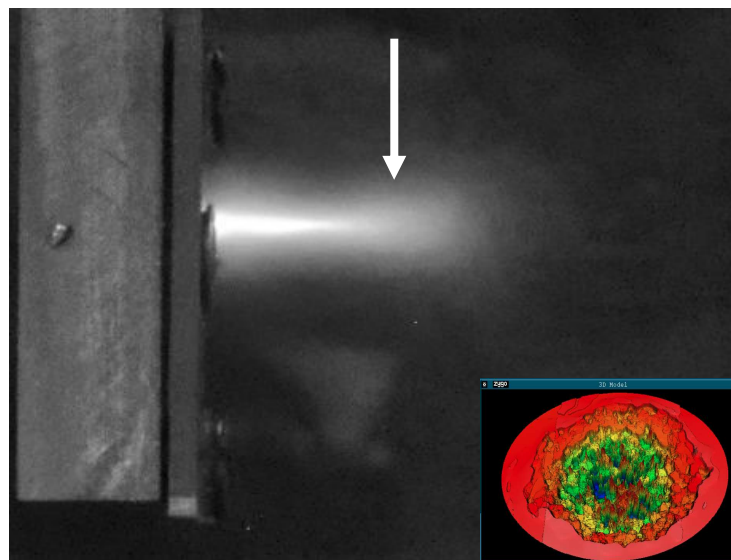
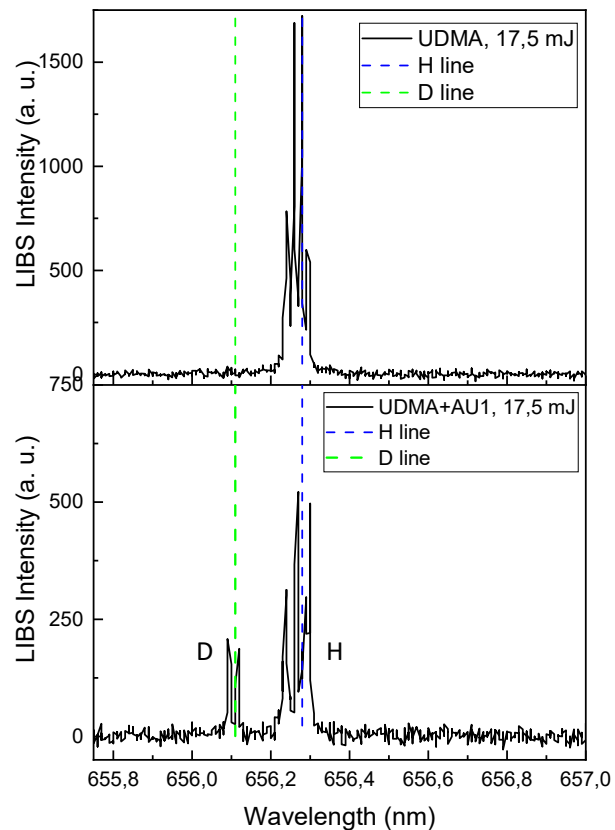
$I_{\text{laser}} > 10^{16} \text{ W/cm}^2$







## SOME RESULTS OF THE H<sup>α</sup> AND D<sup>α</sup> SPECTRAL LINES

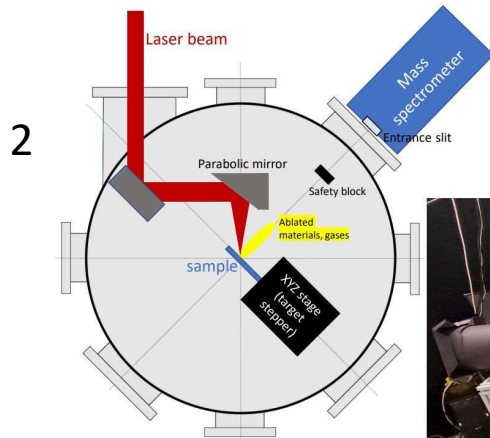
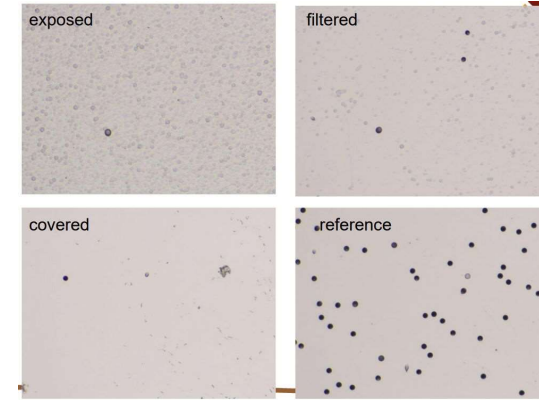
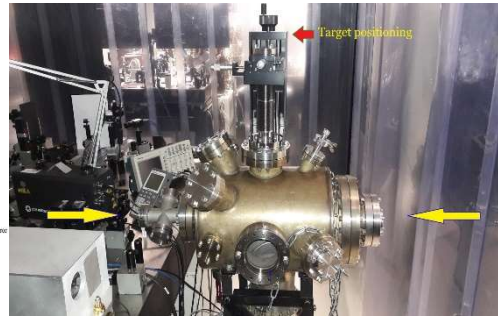
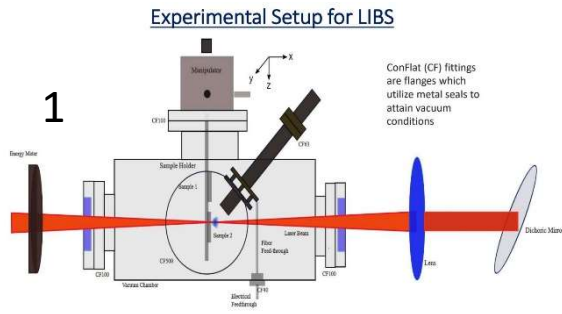


2D+H The total number of H atoms before the transmutation process  
 Number of D atoms in the case of 17.5mJ laser pulses :  $\sim 1.76 \times 10^{15}$

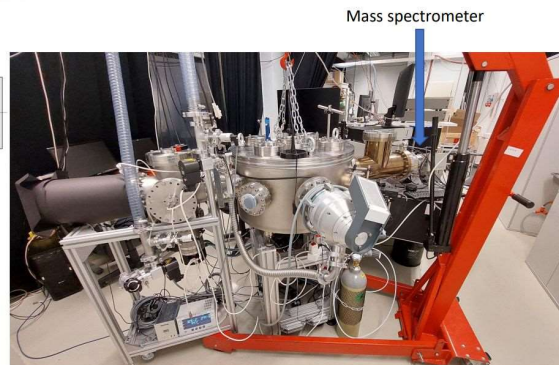
# IN WORK BUT NOT YET CONCLUDING TECHNIQUES:

1. Atomic optical spectroscopy,
2. Mass spectrometry.
3. Nuclear detection techniques.

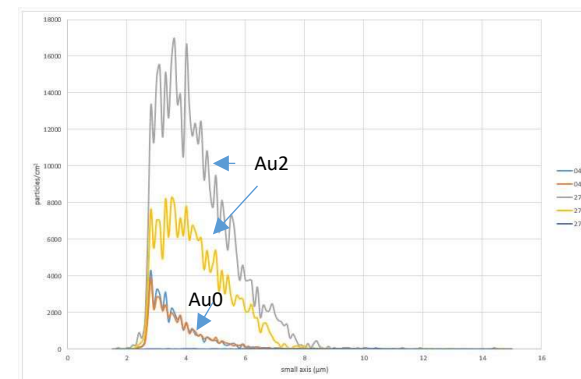
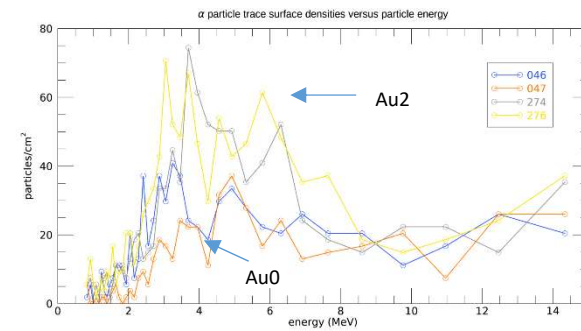
## 3: CR39 film, 1 laser shot



Experimental arrangement



Boron nitride seeded samples



## Extreme Light Infrastructure – Szeged, HU



ELI-ALPS Szeged:  
EU Extreme Light Infrastructure  
Also attosec laser pulses

2PW extreme energies;  
10Hz/1Hz,  $<10\text{fs}$ , 30 J,  $10^{12}$  contrast



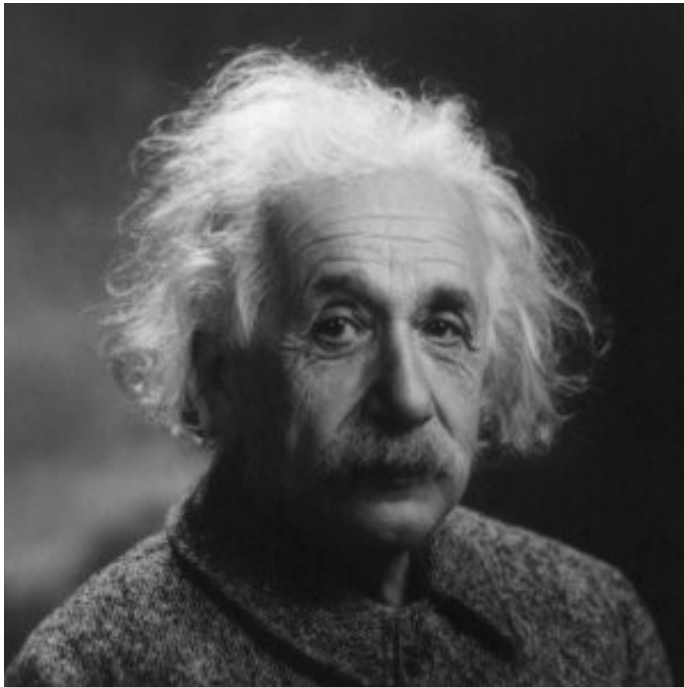
## SUMMARY:

- Localized plasmons (LSPP) differ from the propagating ones with positive consequences.
- Properties are shape and material dependent (Au and Ag resonances in the visible spectrum)
- At high laser intensities no plasma mirror effect.
- Nanoparticles are effective at high laser intensities. Field amplification (hot spots).
  - Optimal lifetime for femtosecond lasers (e.g. Ti:Sa, 800nm)
  - Simplified geometry.
  - Screening (near field and ponderomotive effect)
  - Correlated momentum transfer
- Energy production (crater volume up to 8 times larger with Au nanorods) and  $H \longrightarrow D$  transmutation (fusion) as the explanation, indicated by Raman scattering on the C-D and N-D vibrations
- Supportive modelling results.
- Still several open questions.



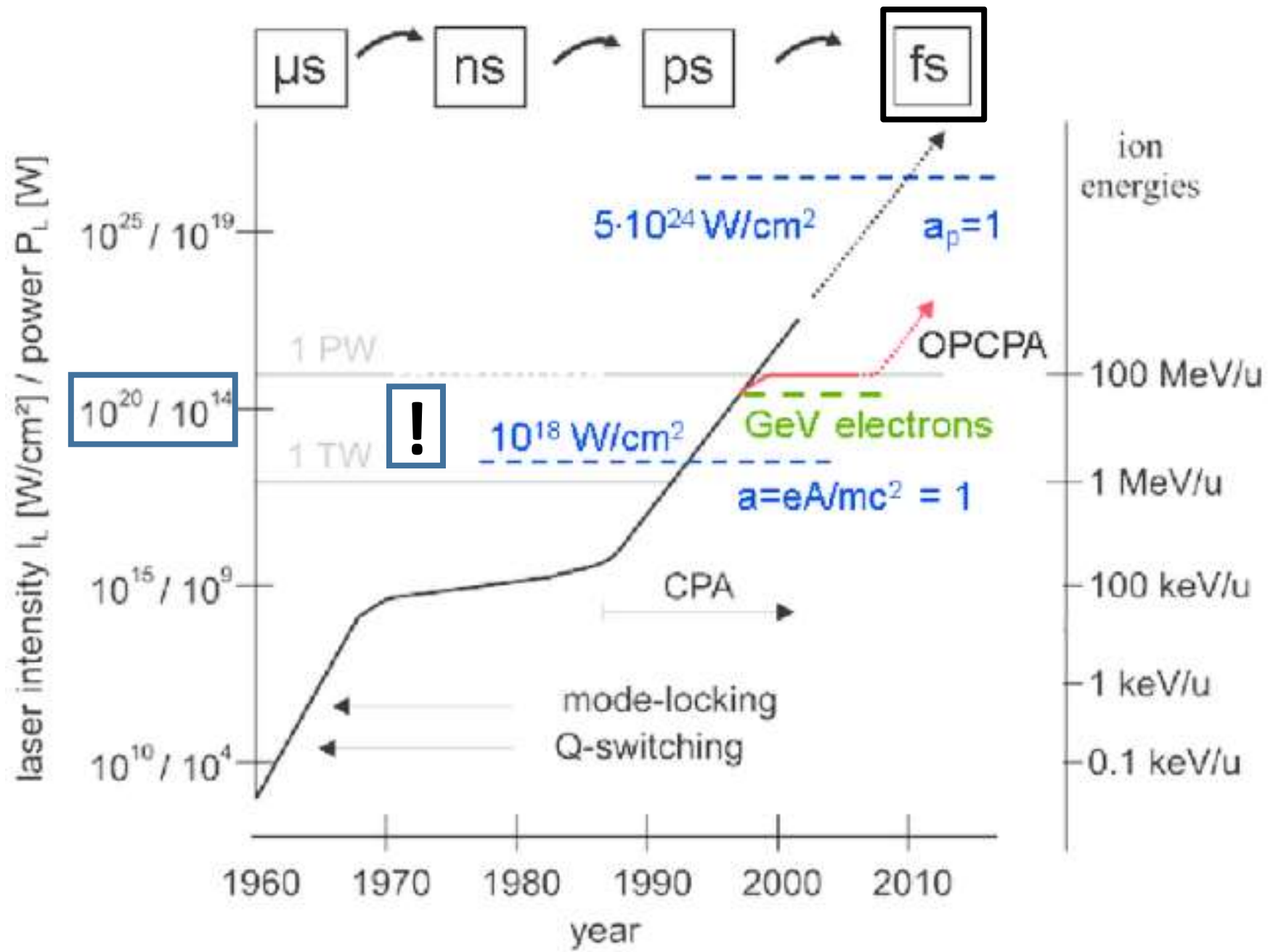
# THE ADVICE OF ALBERT EINSTEIN FOR THE FUTURE:

THE PROBLEMS WE ARE FACING TODAY CAN NOT BE  
SOLVED WITH THE SAME WAY OF THINKING BY WHICH WE  
CREATED THEM.

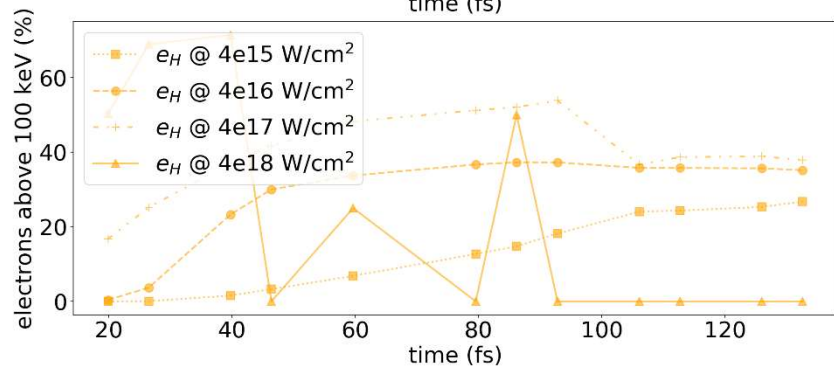
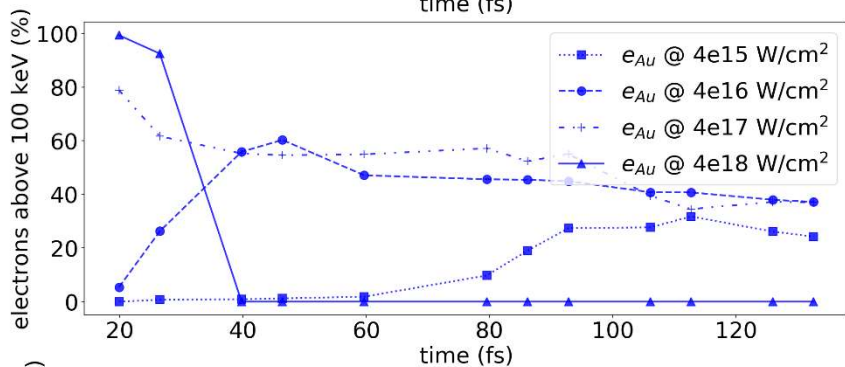
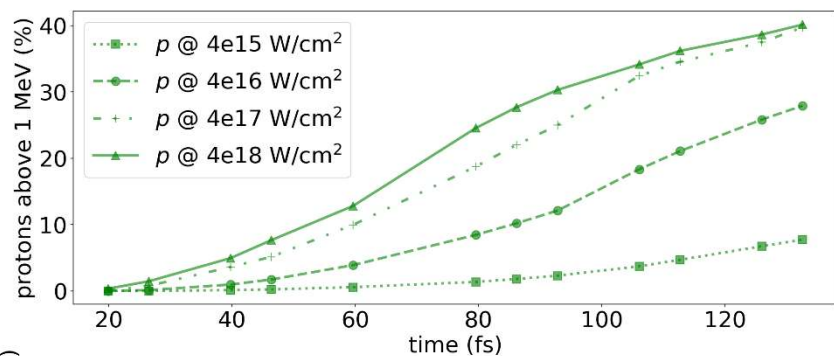


THIS IS WHY CREATIVE  
SCIENTIFIC THINKING IS  
THE KEY TO OUR FUTURE

THANKS FOR YOUR  
ATTENTION



Fraction of particles at energies



Fraction of particles at energies

