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Indirect nanoplasmonic sensing.

B. Demirdjian, I. Ozerov, A. Ranguis, F. Bedu, C. R. Henry

Aix-Marseille University, CNRS, CINaM UMR 7325, 13288 Marseille, France



SCIENTIFIC BACKGROUND

Development of chemical and biological nanosensors based on the extraordinary optical properties of noble-metal nanoparticles.

Nanosensors based on localized surface plasmon resonance (LSPR) are sensitive to small local changes in refractive index at the surface of metal nanoparticles during the adsorption/desorption of molecules.

→ These changes induce a shift in the wavelength of the LSPR response.

Variations of the LSPR peak extinction is linked to the collective oscillation of electrons in the metal. The minimum of reflection corresponds to a maximum of absorption.

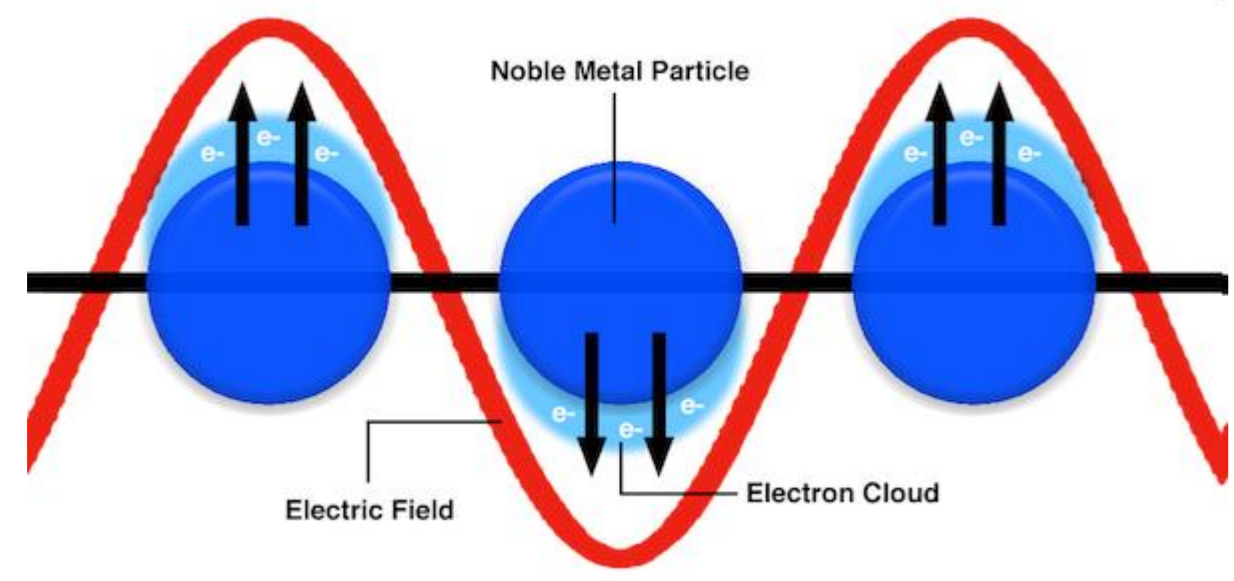
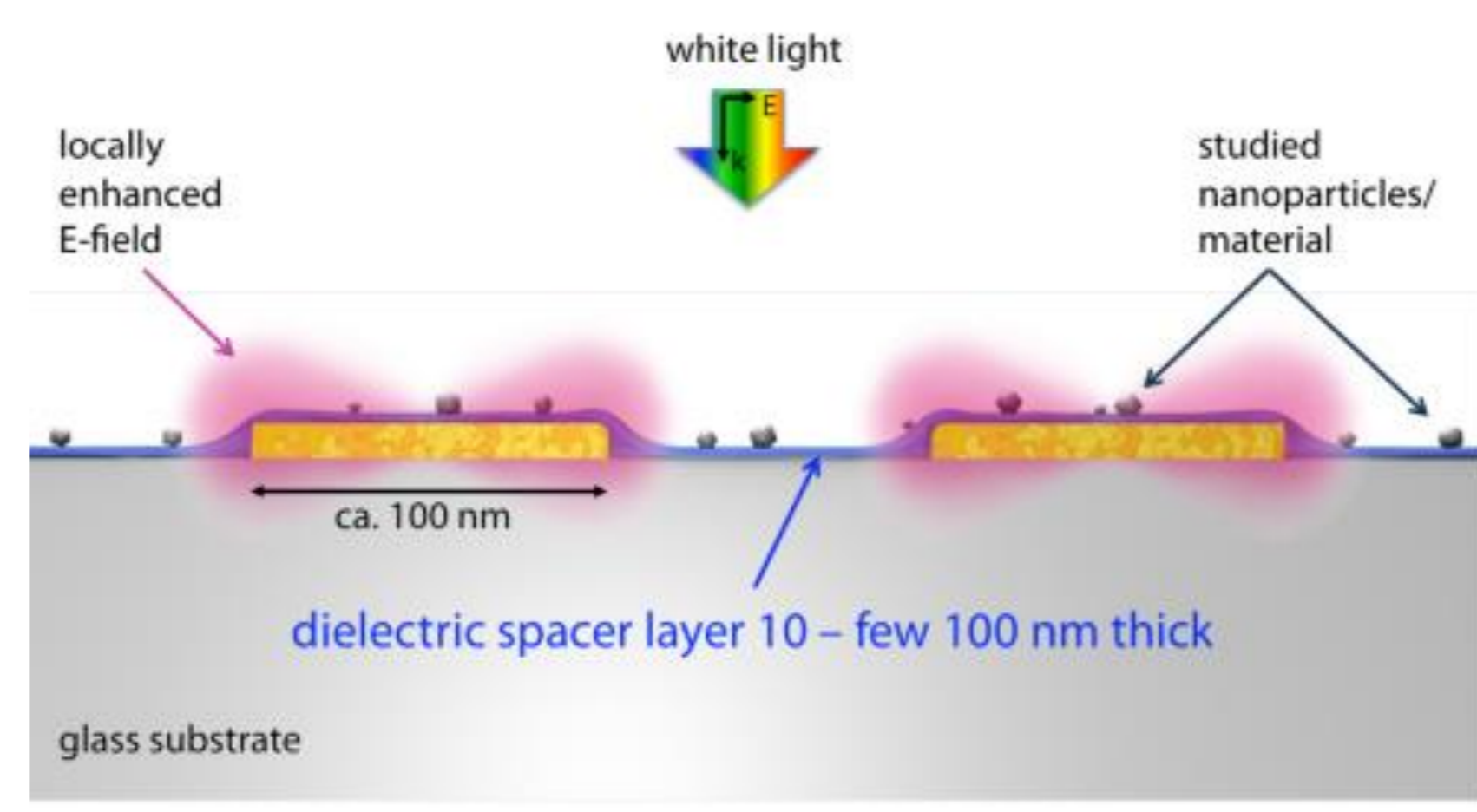


Figure: <http://simslab.uwaterloo.ca/research/biomems-sensors.php>

INDIRECT NANOPLASMONIC SENSING (INPS)

Gas adsorption on nanoparticles modify their surface dielectric properties

→ a shift in the wavelength of the LSPR response of the underlying gold disk detector.



<http://www.insplon.com/technology/indirect-nanoplasmonic-sensing>

GOLD NANODISKS FABRICATION: EBL PROCESS

Sample holder = borosilicate glass window, e = 1 mm, Ø = 25,4 mm
Cleaning: acetone + US, then isopropanol + US, EDI rinsing, oxygen plasma oven at 150°C (300 W during 10 min)

PMMA spin-coating (resin 950 K at 4 %, speed : 4000 rpm, e = 270 nm annealing 10 min at 170°C)

Gold layer deposition (5 nm) to remove the charges (Edwards 306)

The gold film is irradiated with an electron beam (Raith PIONEER) Area of 1 x 1 mm²

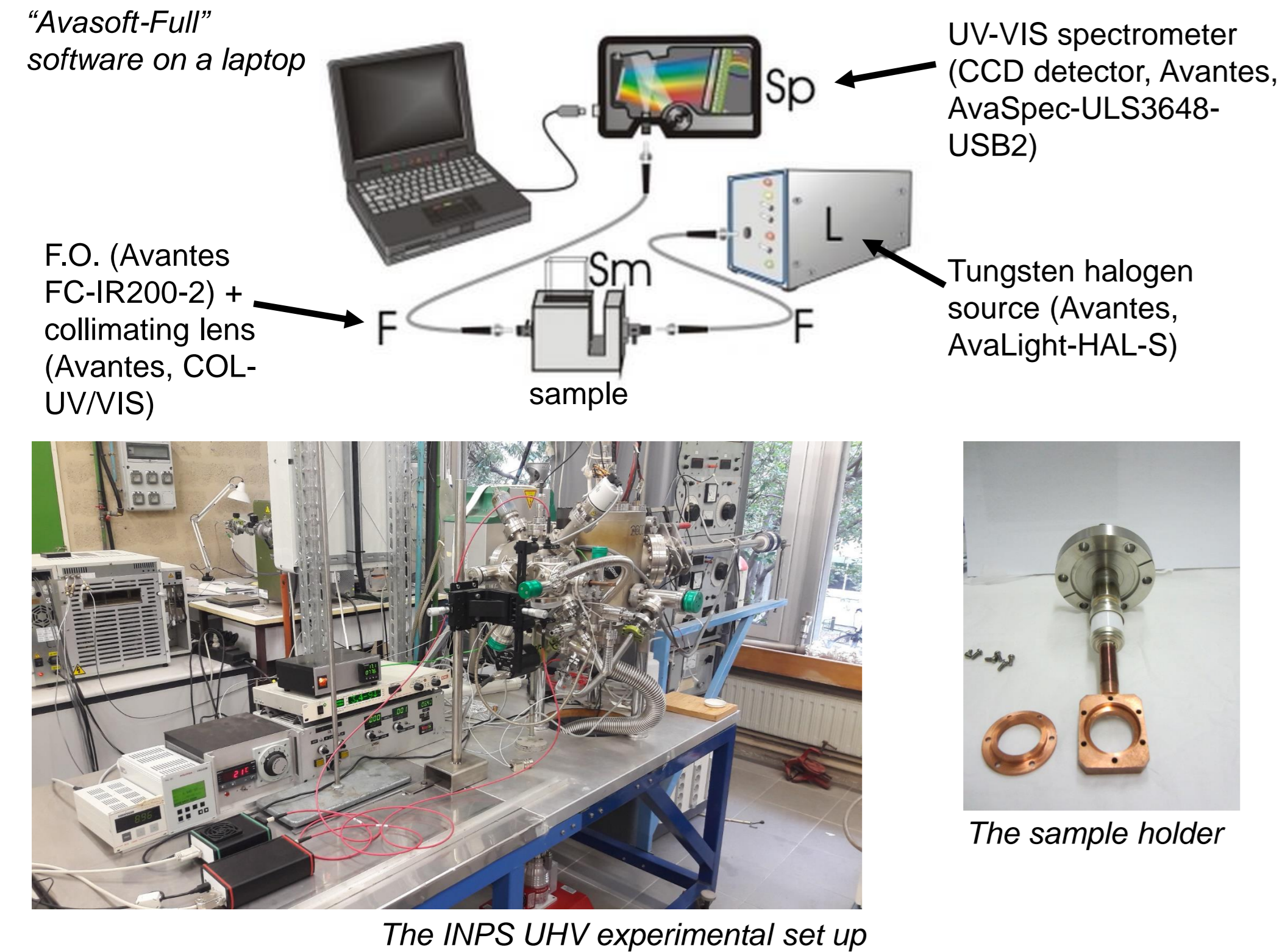
Revelation in acetone (Au is removed) then with MIBK/IPA 1 : 3 during 45 s Then with IPA during 45 s (→ holes in the resin)

Cr and Au evaporation within the PMMA resin (Edwards 306) e_{Cr} = 2 nm, e_{Au} = 20 nm

Lift-off of the resin (acetone + US)

→ Precise control of the shape, the size and the pitch of gold nanodisks

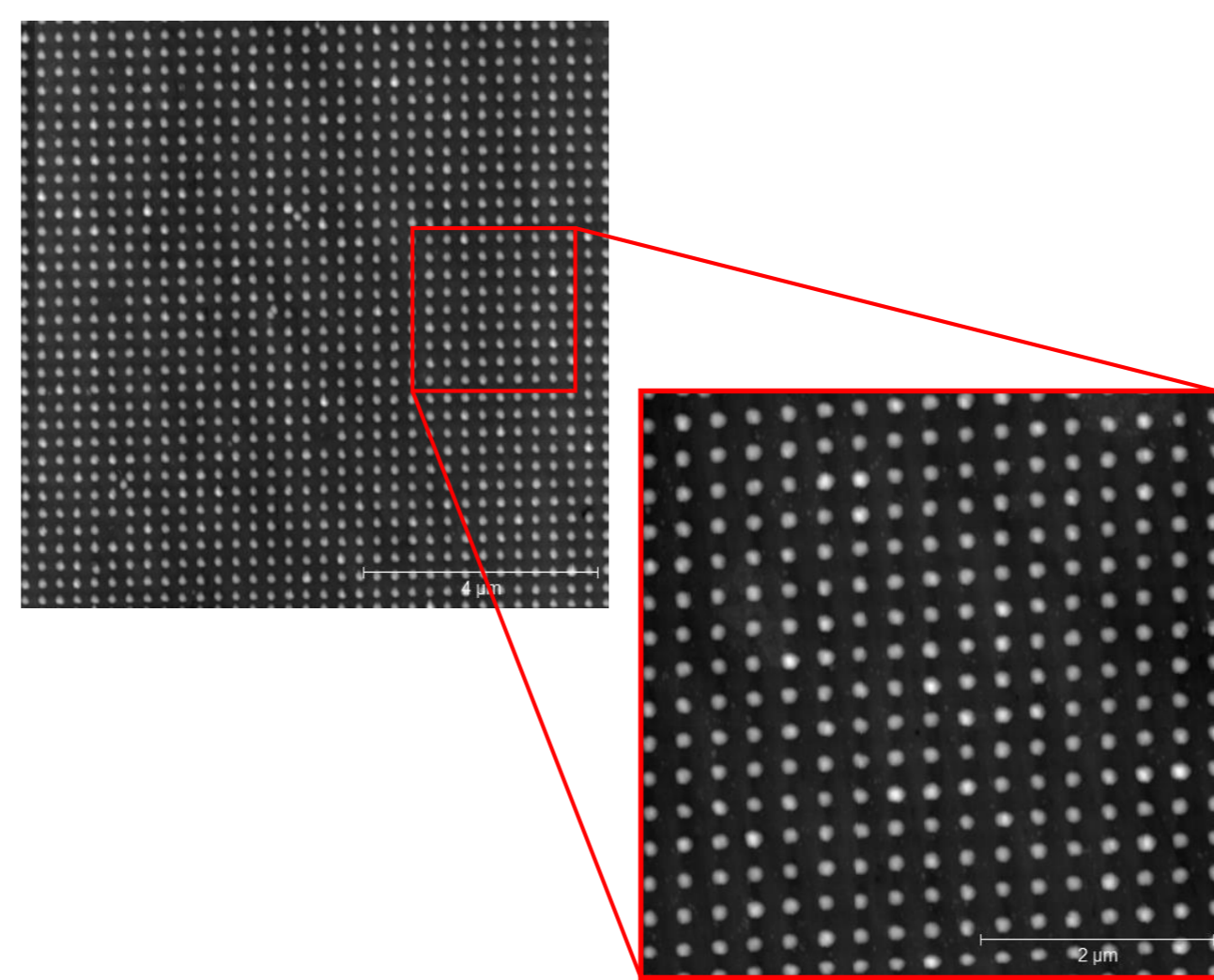
LSPR MEASUREMENTS: EXPERIMENTAL SET-UP



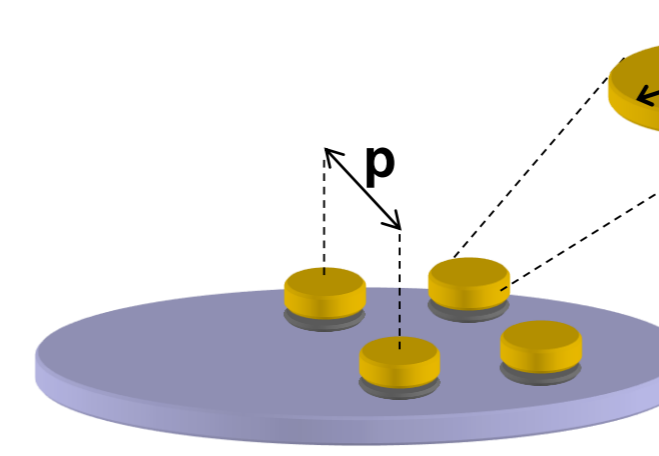
The INPS UHV experimental set up

OPTIMISATION OF THE GOLD NANODISKS PARAMETERS

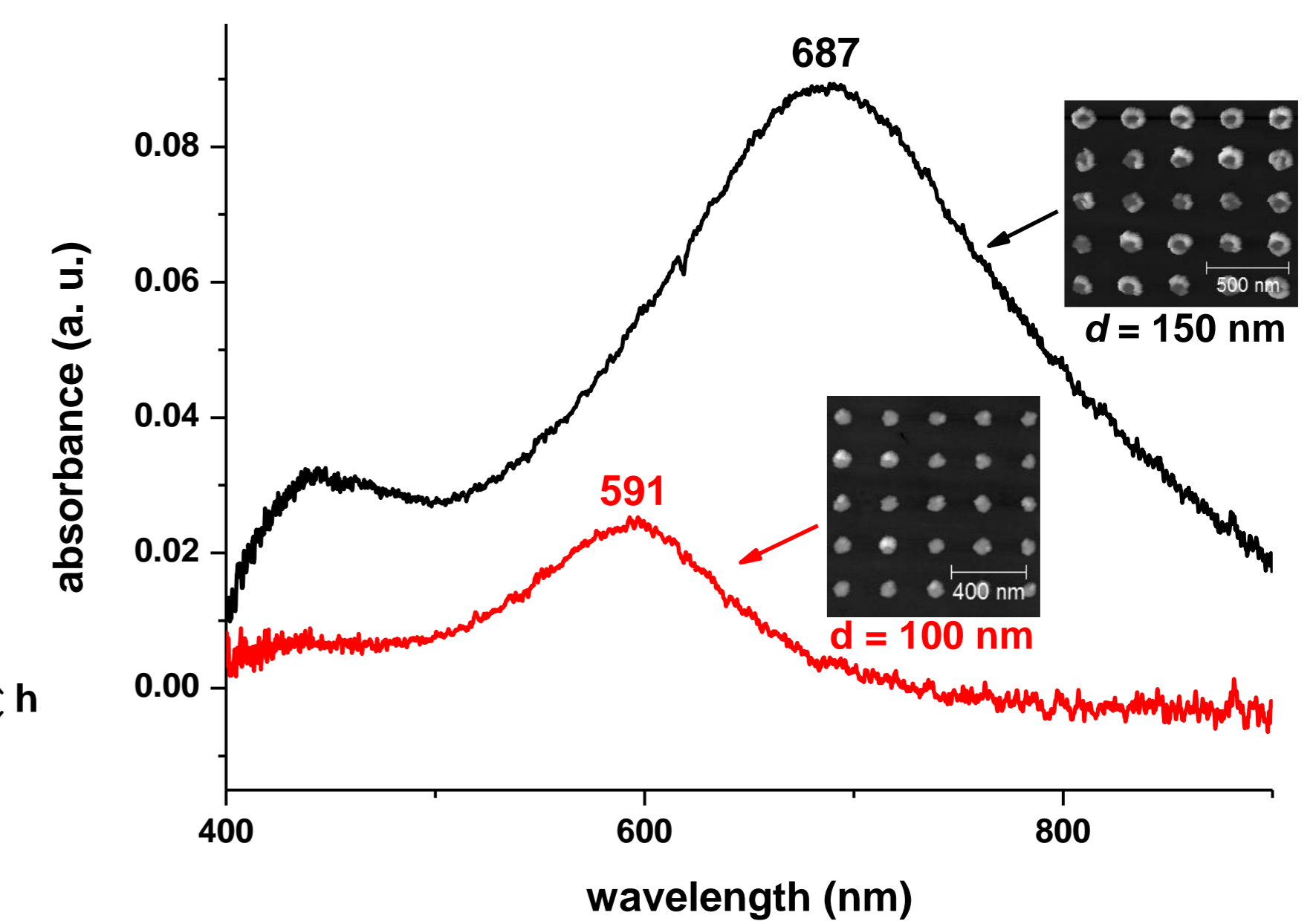
FDTD simulations → best theoretical LSPR S/N signals obtained with the biggest aspect ratio ie the biggest diameter (for a given h)



<p> = 300 nm
<h> = 29 nm,
d = 100, 150 nm



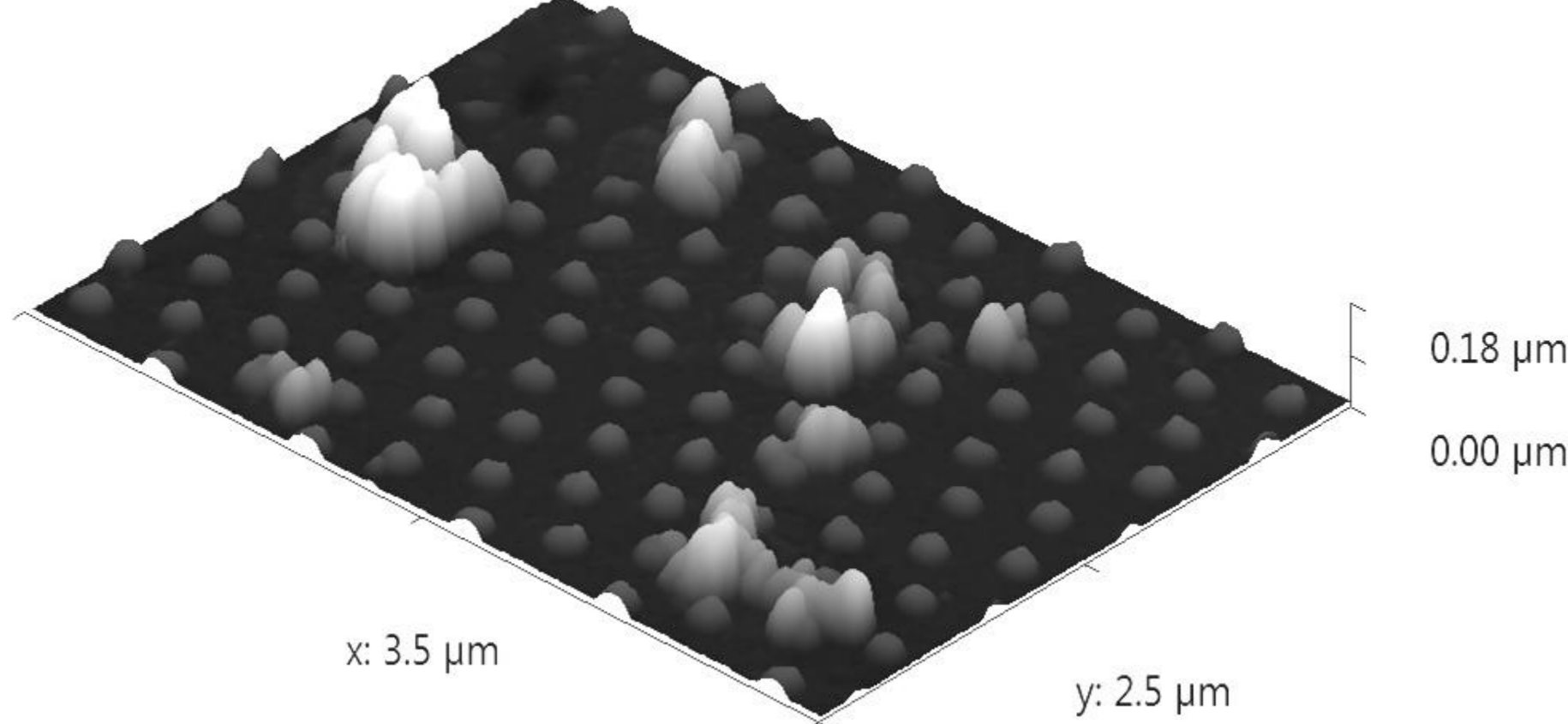
Tapping mode topographical AFM images of gold nanodisks on borosilicate window (images obtained with a PSIA apparatus XE-100)



Experimental LSPR responses for gold nanodisks deposited onto a borosilicate glass window vs the diameter, d. Tapping mode topographical AFM images of gold nanodisks with d = 100 and 150 nm. Images obtained with a PSIA XE-100.

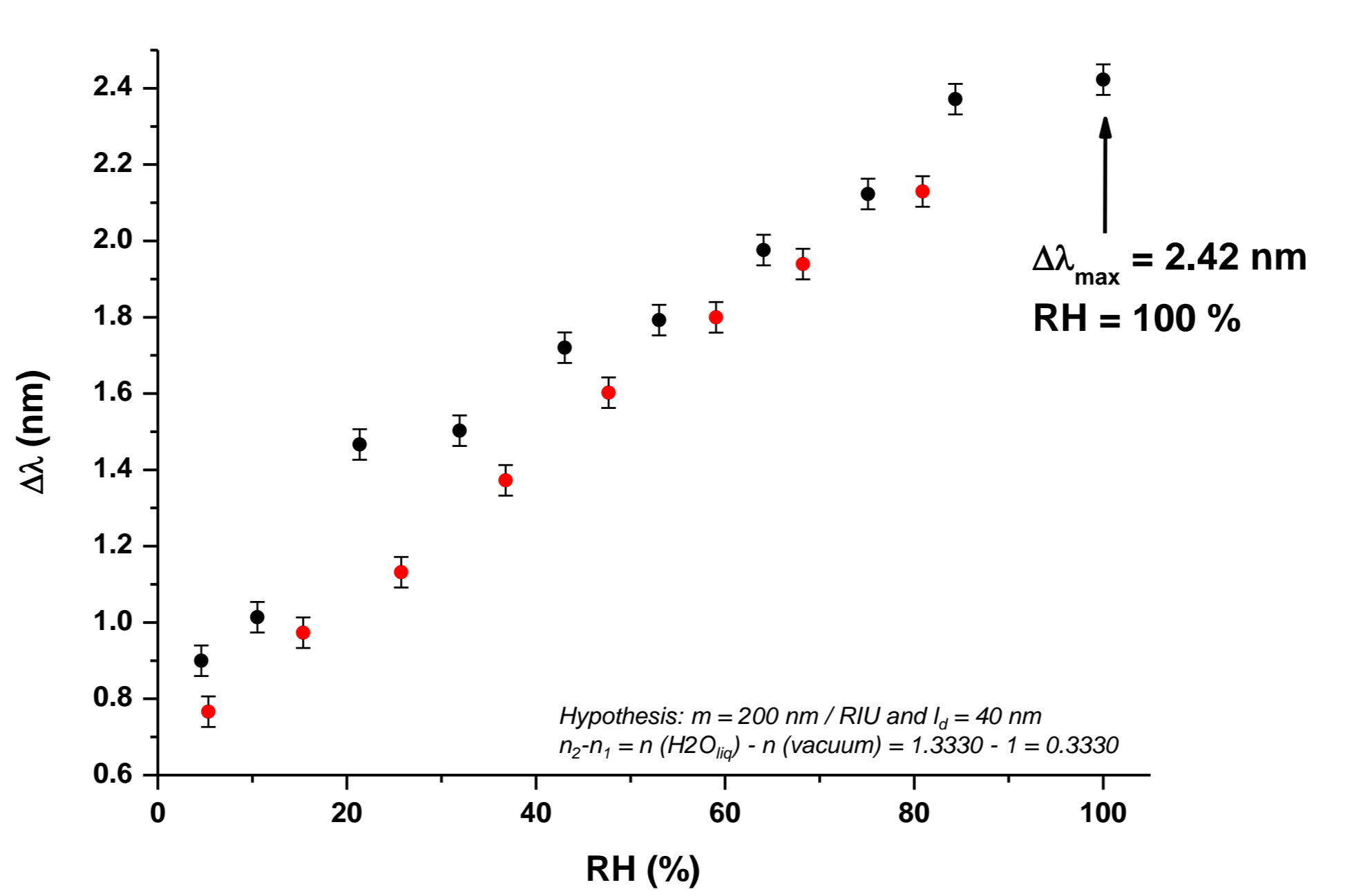
SOOT DEPOSITION ON GOLD NANODISKS: DROPLET DEPOSITION

Soot particles are involved in several atmospheric processes (ice nuclei, contrails formation, radiative forcing, chemical reactions...), it is important to well characterize the water/soot interaction mechanism



Tapping mode topographical AFM image of AEC soot particles deposited on gold nanodisks (covered by a SiO₂ layer). Image obtained with a PSIA apparatus (XE-100).

REFERENCE : WATER ADSORPTION ON SiO₂ / GOLD DISKS



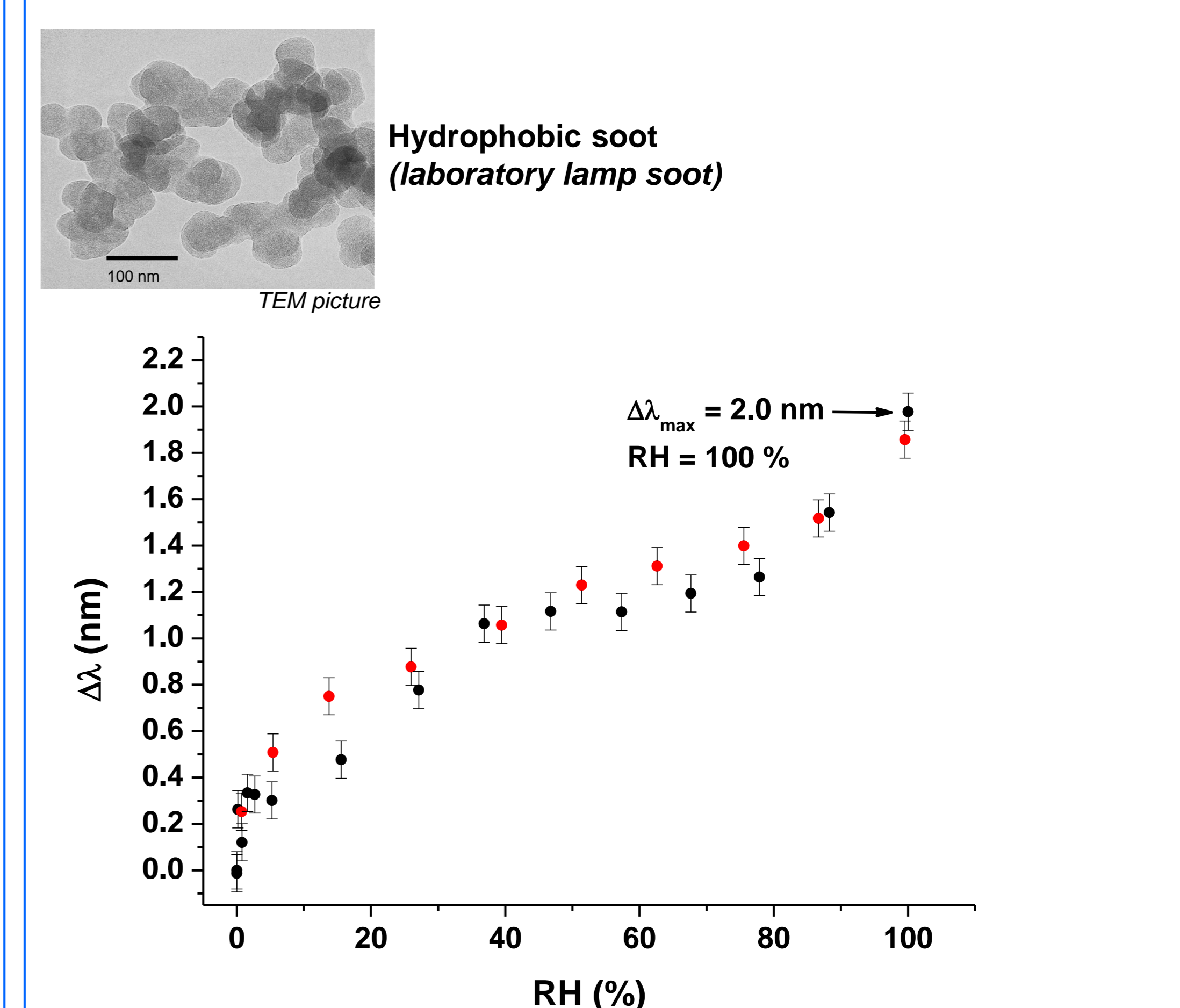
Evolution of the shift $\Delta\lambda$ of the LSPR response versus the relative humidity inside the reactor at room temperature during the adsorption of water vapor on the gold nanodisks covered by a SiO₂ layer. Black dots are obtained during the adsorption, and red dots are obtained during the desorption. Error bars are indicated.

$\Delta\lambda$ is almost linear vs the relative humidity, at RH = 100%, $\Delta\lambda = \Delta\lambda_{\max} = 2.42$ nm.
→ water layer thickness d = 0.67 nm (~ 2.2 water ML) Equation (1)

Good reversibility of the adsorption and desorption curves (H₂O physisorption and no chemisorption of water molecules on SiO₂)

Highly sensitive sensor: accuracy of the wavelength measurement in the LSPR response is 0.04 nm → detection of about 2/100 of water ML !

WATER ADSORPTION ON HYDROPHOBIC SOOT PARTICLES



Evolution of the shift $\Delta\lambda$ of the LSPR response versus the relative humidity inside the reactor at room temperature during the adsorption of water vapor on soot particles deposited on gold nanodisks covered by a SiO₂ layer. Black dots are obtained during the adsorption, and red dots are obtained during the desorption. Error bars are indicated.

RESPONSE OF THE LSPR NANOSENSORS

The response of LSPR nanosensors follows a simple model described by the group of Campbell:

$$\Delta\lambda = m (n_2 - n_1) [1 - \exp(-2d/l_d)] \quad (1)$$

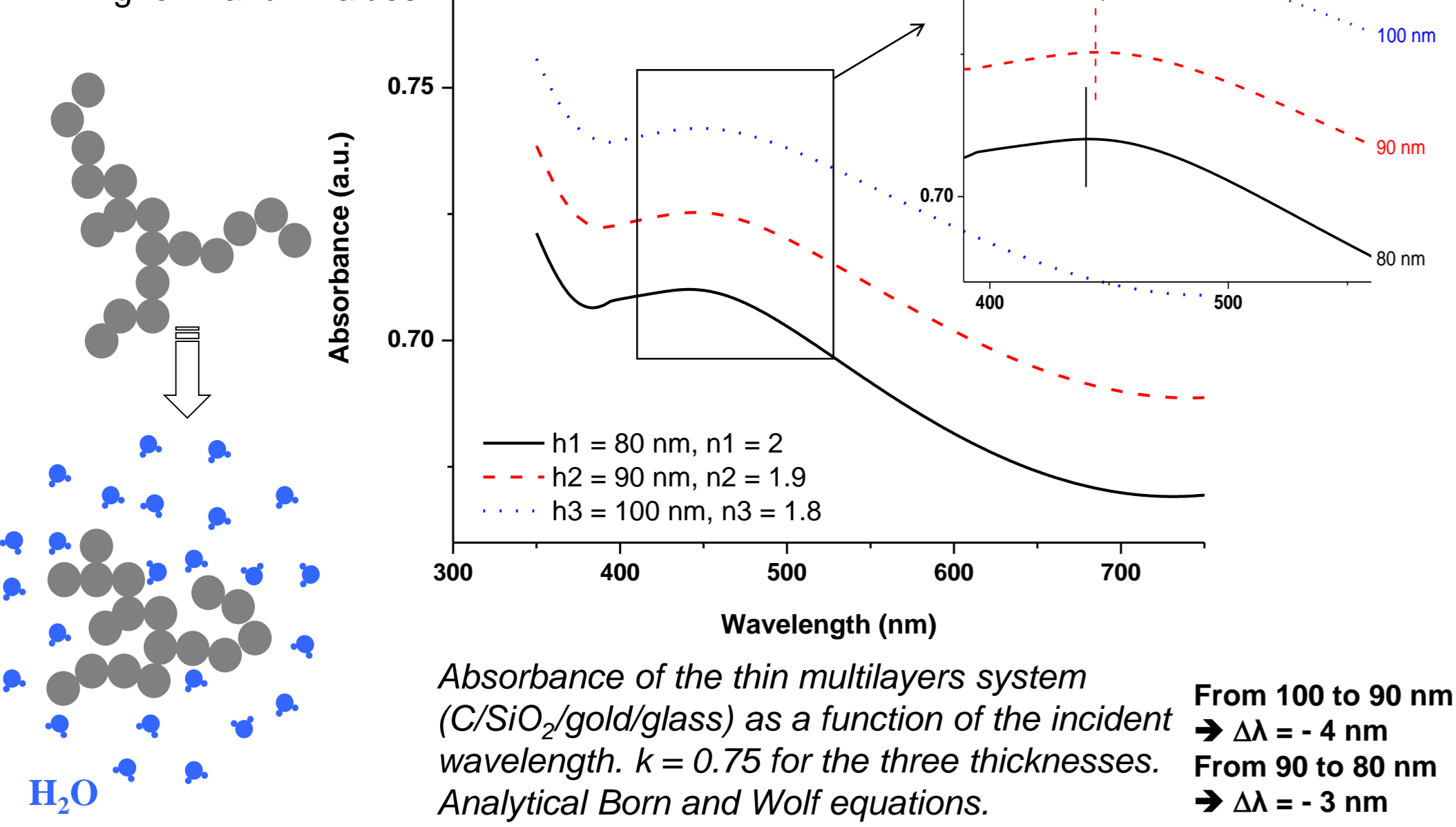
$\Delta\lambda$: wavelength shift
m: sensitivity of the refractive index (RI)
 n_2 and n_1 : RIs of different surrounding media
d: effective thickness of the adsorbate layer
 l_d : characteristic decay length of the evanescent electromagnetic field.

BLUE SHIFT ? → MODELISATION (A. KARAPETYAN)

Hydrophilic soot aggregates collapse into + dense structures during RH ↑ (Mikhailov):

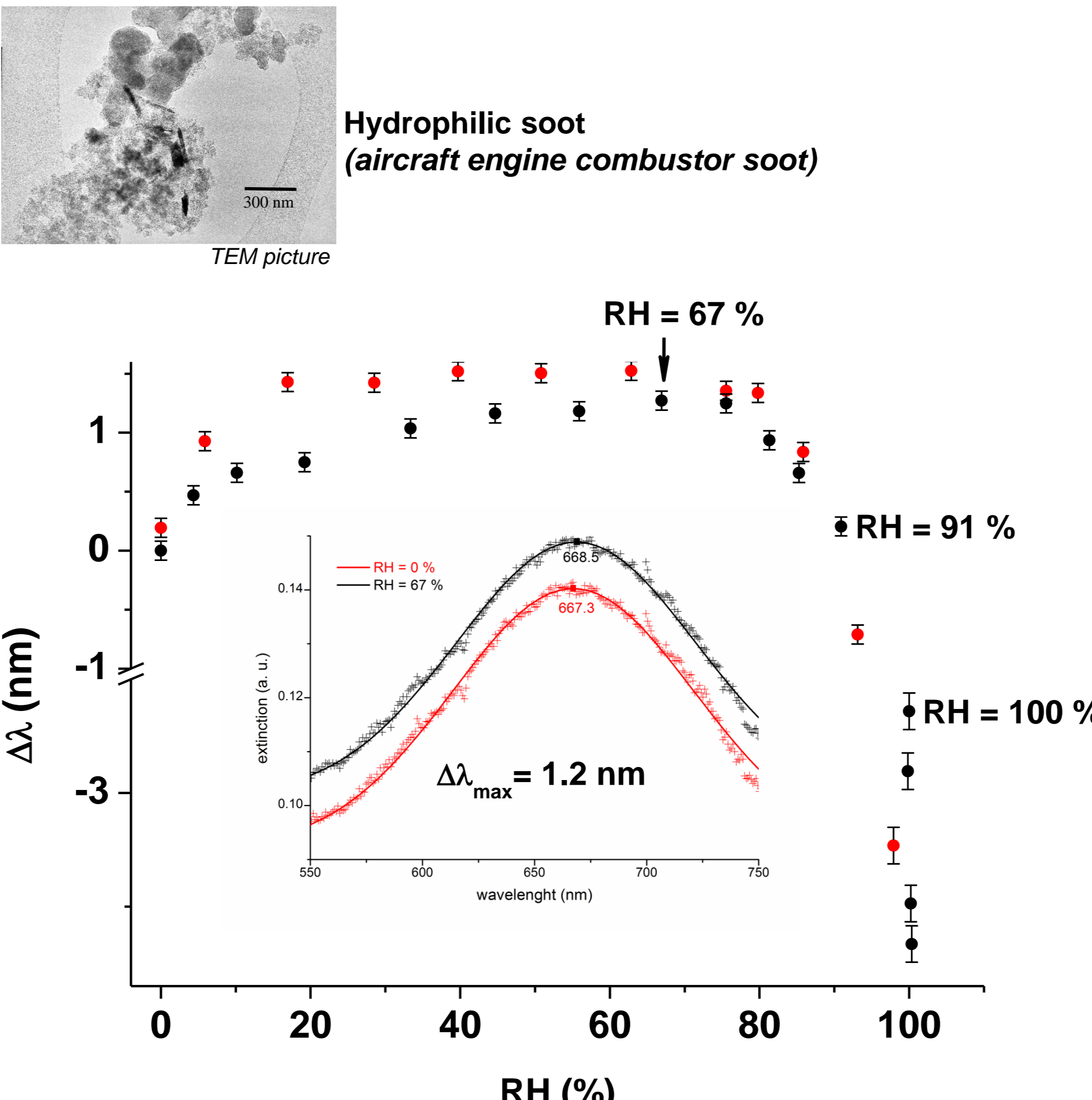
→ RH = 100 % soot aggregates are more compact

- higher mass fractal dimension Df
- smaller gyration radius
- higher n and k values



Absorbance of the thin multilayers system (C/SiO₂/gold/glass) as a function of the incident wavelength. k = 0.75 for the three thicknesses. Analytical Born and Wolf equations.

WATER ADSORPTION ON HYDROPHILIC SOOT PARTICLES



Evolution of the shift $\Delta\lambda$ of the LSPR response versus the relative humidity inside the reactor at room temperature during the adsorption of water vapor on soot particles deposited on gold nanodisks covered by a SiO₂ layer. Black dots are obtained during the adsorption, and red dots are obtained during the desorption. Error bars are indicated.

→ Peculiar and reversible blue-shift !

Conclusions

- High sensitive water sensor (detection of a few hundredths of H₂O ML)
- Detect morphology changes of soot aggregates
- Large (P, T) domains of studies
- Non destructive probe

Outlooks

- Quantitative measurements with mass spectrometry
- Continuing FDTD simulations
- Chemical reactivity gas / nanoparticles:

•Catalysis
Metallic nanoparticles (Pt, Pd, ...) + CO, O₂ + H₂, NO_x

•Environmental heterogeneous reactions
Ice nucleation on soot particles
Photochemistry and reactivity NO₂/soot → O₃

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